

HANDBOOK FOR 2021

FACULTY of HEALTH SCIENCES

DEPARTMENT of BIOMEDICAL and CLINICAL TECHNOLOGY

The above department offers two programmes: Biomedical Technology/Medical Laboratory Science & Clinical Technology

This handbook offers information on both programmes.

WHAT IS A UNIVERSITY OF TECHNOLOGY?

A university of technology is characterized by being research informed rather than research driven where the focus is on strategic and applied research that can be translated into professional practice. Furthermore, research output is commercialized thus providing a source of income for the institution. Learning programmes, in which the emphasis on technological capability is as important as cognitive skills, are developed around graduate profiles as defined by industry and the professions.

NOTE TO ALL REGISTERED STUDENTS

Your registration is in accordance with all current rules of the Institution. If, for whatever reason, you do not register consecutively for every year/semester of your programme, your existing registration contract with the Institution will cease. Your re-registration anytime thereafter will be at the discretion of the institution and, if permitted, will be in accordance with the rules applicable at that time.

IMPORTANT NOTICES

The rules in this departmental handbook must be read in conjunction with the General Rules (G Rules) contained in the DUT General Handbook for Students as well as the relevant subject Study Guides.

Your attention is specifically drawn to Rule G1 (8), and to the process of dealing with students' appeals.

FACULTY of HEALTH SCIENCES FACULTY VISION, MISSION, GOALS & VALUES

(November 2012 for 2013-2017)

Vision:

"Leading Transformative and Innovative Health Sciences Education"

Mission Statement:

- "Developing Holistic Professionals responsive to Healthcare needs" through Excellence in:
- Teaching and Learning
- Research, Innovation and Engagement
- Fostering Entrepreneurship

Values

Professionalism

(To work within regulatory frameworks of professional conduct. To maintain and develop professional expertise and good work ethic.)

Integrity

(To conduct ourselves with strong moral principles. To be honest and authentic. To do what is ethical and just.)

Ubuntu

(To treat people with respect, fairness, courtesy, politeness and kindness.)

Transparency

(To conduct ourselves with openness and honesty through shared governance.)

Accountability

(To accept responsibility for one's actions.)

Goals

The Faculty aims to:

- Respond to the National health human resource and industry needs within the health sector.
- 2. Ensure the offering of entrepreneurial and leadership skills as a core component of all programmes within the Faculty of Health Sciences.
- 3. Continue to develop community-based projects to foster social responsibility through collaborative projects between programmes.
- 4. Enhance established quality management frameworks to support teaching and learning.
- 5. Develop applied research responsive to community and industry needs.
- Develop mechanisms for the dissemination and application of research outcomes to inform teaching and learning, assessment, community engagement and further research.
- 7. Improve research participation and output through increased post-graduate student enrolment, publications and establishment of research groups.
- 8. Enable the generation of third-stream income through research and innovation (patents / artifacts) in order to supplement existing sources of income for the next five years.
- 9. Attract and retain diverse quality staff, while promoting advancement of individual potential.
- 10. Position DUT Health Sciences nationally

Values

The Faculty is guided by the following core values:

- I Transparency, openness, honesty, and shared governance
- 2 Professional and personal respect for others
- 3 Educational relevance, equity and transformation (curriculum, access and success)
- 4 Loyalty, accountability, dignity and trust

DEPARTMENTAL MISSION & GOALS

The above department offers two programmes:

Biomedical Technology and Clinical Technology

Vision:

Globally recognized for Medical Laboratory and Clinical Technology Science Education

Mission:

"Develop Critical, Investigative Professionals for Diagnosis and Disease Management"

Through

- Teaching and Learning
- Research
- Community and Industry Engagement
- Entrepreneurship

Values

Professionalism

(To conduct oneself within established standards and norms. To demonstrate professional skills and behaviour)

Integrity

(To be honest and trustworthy. To be ethical and fair in critical analysis and reporting.)

Accountability

(To be answerable for one's actions. To be accountable to our society. To be committed)

Patients' Lives Matter

Graduate attributes:

- Use a range of information technologies to identify, gather and disseminate information.
- Engage in the generation of new knowledge in their specialist professional disciplines and academic fields which will be investigated and recorded scientifically.
- 3. Work independently, identify, critically analyse and solve problems in their professional, individual and societal environments
- 4. Lead and effectively manage team members in an organisation and within their communities.
- Be aware of cultural diversity and show respect to indigenous knowledge, cultures and values
- 6. Think critically and have excellent decision making skills including awareness of personal strengths and limitations.
- 7. Communicate effectively within the health care and educational environment, using visual, mathematical and/or language skills in the modes of oral and or written presentation
- 8. Use science and technology effectively and critically, showing responsibility towards the environment and health of others
- Participate as responsible citizens in the life of local, national and global communities

Goals

The department aims to:

- 1. Provide quality teaching, learning and support to students
- 2. Respond to national human resource and industry needs
- 3. Provide excellent professional value-driven education, promote entrepreneurship and leadership skills.
- 4. Produce graduates that are independent thinkers functioning within a team
- 5. Foster professional and ethical conduct
- 6. Keep abreast with current and future technological trends
- Enhance the quality management frameworks to support teaching, learning, assessment and research.
- 8. Encourage research responsive to community and health needs
- 9. Position the Department of Biomedical and Clinical Technology nationally and internationally.
- 10. Attract and retain diverse quality staff while promoting advancement of individual potential
- Maintain relationships within the institution, relevant professional bodies, industry, educational institutions, alumni and other stakeholders.
- 12. Foster national and international collaboration and partnerships
- 13. Strive for excellence and success
- 14. Embrace an attitude of life-long learning with the aim to improve professional clinical practice through research

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I. DEPARTMENTAL & FACULTY CONTACT DETAILS

All departmental enquiries to:

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 Tel No:
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 (031) 373 5295

 Email:
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Location of Department: MB 2-9 ML Sultan Campus

All Faculty enquiries to:

Faculty Officer: Miss FT Mayisela
Tel No: (031) 373 2701

Email: thembim@dut.ac.za

Location: Health Faculty Office, Gate 8,

Steve Biko Road, Mansfield Site

Area, Ritson Campus

Executive Dean: Prof A Ross
Executive Dean's Secretary Mrs Bilkish Khan

Tel No: (031) 373 2704 Fax No: 0866740237

Email: bilkishk@dut.ac.za

Location: Executive Dean's Office, Gate 8,

Steve Biko Road, Mansfield Site

Area, Ritson Campus

2. **DEPARTMENTAL STAFF**

Staff NAME AND QUALIFICATION

Head of Department Dr J N Mbatha PhD: Medical Micro (UKZN)

Senior Lecturers Mrs B T Mkhize, MTech: Biomed Tech (DUT)

Dr M J Mohapi, PhD (Health Sciences); MED (Higher

Education)

Dr P Pillay, PhD (UKZN)

Dr S C Benjamin¹ DTech: Clin Tech (DUT) Dr D R Prakaschandra, PhD (Cardiology) (UKZN)

Lecturer Mr M E Memela, MTech: Clin Tech (DUT)

Miss T S Ndlovu, MTech: Biomed Tech (DUT) Mr D Govender, NHD: Med Tech (MLST)

Mr C Sydney2, M Med Sc (UKZN)

Mr DC Mdluli (MSc Med; BTech: Clin Tech)

Senior Lab Technician

Mrs Y Pillay, Comp Prog (MLST)

Laboratory Technicians Mr J Mbuyazi, ND: Pharmaceutical Marketing (MLST)

Ms T C Qangule, ND: Med Tech Micro (Pen Tech)

Mr D Reddy, Cytotechnician, (CTCMIAC)

Laboratory Assistant
Departmental Secretary

Miss H Ramphal, ND: OMT (DUT)
Mrs B G Nene, BTech: OMT (DUT)

Head of Programme : Clinical Technology

² Head of Programme : Biomedical Technology and Medical Laboratory Science

3. DEPARTMENTAL INFORMATION & RULES

3.1 PROGRAMMES OFFERED BY THE DEPARTMENT

This department offers two programmes, namely:

- Biomedical Technology/Medical laboratory Science
- Clinical Technology

3.2. Qualifications offered by the department

Learning programmes are offered in this Department which will, upon successful completion, lead to the award of the following qualifications:

Qualification	Qualification Code	SAQA NLRD Number	Important Dates	
Biomedical ⁻	Technology Programme			
ND: Biomedical Technology*	NDBMTI	1895	Teach- out date: 2021	
ND: Biomedical Technology (ECP)*	NDBMFI		Teach- out date: 2021	
BTech: Biomedical Technology*	BTBMT1/BTBMT2	1899	Teach- out date: 2021	
Master of Health Sciences in Medical Laboratory Science	MHMLSI	96822	Not applicable	
Doctor of Medical Laboratory Science	DRMLSI	96805	Not applicable	
BHSc in Medical Laboratory Science		101689		
Clinical Te	chnology Programme			
ND: Clinical Technology*	NDCLTI	1879	Teach- out date: 2021	
ND: Clinical Technology (ECP)*	NDCLF2		Phasing out date: 2017	
BTech: Clinical Technology*	BTCLT1/BTCLT2	1889	Phasing out date: 2019	
Masters of Health Sciences in Clinical Technology	MHCLTI	96956	Not applicable	
Doctor of Medical Clinical Sciences	DRMCSI	96809		
BHSc in Clinical Technology		96409		

^{*} Qualification phased out, NO INTAKE.

3.3. DEPARTMENTAL INFORMATION

3.3.1. Academic Integrity

Please refer to the General Rules pertaining to the academic integrity G13 (1)(o). These will be enforced wherever necessary to safeguard the worthiness of our qualifications, and the integrity of the Faculty of Health Sciences at the DUT.

3.3.2. Code of Conduct for Students

In addition to the General Rules pertaining to Student Conduct SR3 (3), a professional code of conduct pertaining to behaviour, appearance, personal hygiene and dress shall apply to all students registered sessions with the Faculty of Health Sciences, at all times.

3.3.3. Uniforms

Students must adhere to instructions regarding specific dress code required during practical sessions and/ hospital visits. All students are required to wear laboratory coats on top of their own clothing and closed shoes during practical and some practical sessions may also need students to wear masks or goggles and gloves.

3.3.4. Attendance

Students are encouraged to achieve 100% attendance for all planned academic activities as these are designed to provide optimal support for the required competency. Where absence is unavoidable, the student must timeously advise the department of the reason. Only exceptional reasons will be accepted for absence from guest lectures, industry or field trips. Poor attendance records may lead to penalties as per programme rules. Where absence impacts on assessment, please refer to Section 3.4. (Departmental Rules) below.

3.3.5. Health and Safety

Students must adhere to all Health and Safety regulations both while at DUT and in Work Integrated Learning (WIL) placements. Failure to do so will be treated as a breach of discipline.

3.3.6. Registration with the Professional Board

As a Student: Within two weeks of registration with the Department, students are required to register as Student Medical Technologists / Medical Laboratory Scientist or Student Clinical Technologists with the Health Professions Council of South Africa as determined in the regulations set out in the Allied Health Service Professions Act, 1982 (Act 63 of 1982) (Regulation R629, Government Gazette No 11221 of 31 March 1988).

As a Graduate (Biomedical Technology/Medical Laboratory Science)

A graduate, on successful completion of the qualification and the required internship, and after passing a competency assessment to satisfy the requirements of the Professional Board for Medical Technology, may register as a qualified Biomedical Technologist or Medical Laboratory Scientist (as applicable) with the Health Professionals Council of South Africa (HPCSA). After registration with the HPCSA, graduates may work in government, private health care laboratories and research laboratories. Unregistered Biomedical Technologists/Medical Laboratory Science may work in non-diagnostic laboratories. To practice independently as a Biomedical Technologist/Medical Laboratory Scientist, two years post-registration experience is required.

As a Graduate (Clinical Technology):

A graduate, on successful completion of the qualification and after having satisfied the requirements of the Professional Board for Radiography and Clinical Technology, may register as a qualified Clinical Technologist (as applicable) with the HPCSA.

3.3.7. Student appeals:

Rule G1 (8) in the DUT General Handbook apply.

3.4. DEPARTMENTAL RULES

3.4.1 Special Tests and condonement

No summative assessments will be condoned. Summative means all assessment marks that contribute to the final mark of a subject, but not including examinations for the purpose of this rule.

- If a student misses a summative written or oral or practical test, for
 reasons of illness, a special test may be granted if the student provides a
 valid medical certificate specifying the nature and duration of the illness,
 and a declaration that for health reasons it was impossible for the
 student to sit for the test. This certificate must be submitted to the
 lecturer or head of programme, no later than one week after the date of
 the missed test.
- If a student misses a summative written or oral or practical test, for
 reasons other than illness, a special test may be granted if the student
 provides a valid declaration that for unavoidable reasons it was
 impossible for the student to sit for the test. This declaration must be
 submitted to the programme coordinator, no later than one week after
 the date of the missed test.
- In addition, a special test may be granted to students with borderline academic results. The special test which may take the form of an oral test, may be set at the end of the period of registration, and may include a wider scope of work than the original test.
- Any student who misses an assessment and who does not qualify for a special test, and any student who qualifies for a special test but fails to write it, shall be allocated a zero mark for the missed assessment. A student who qualifies for a special test granted for borderline academic results, but fails to write it, or achieves lower than their original results, shall be allocated their original results.

3.4.2 Student Appeals

Rule G1 (8) of the general handbook applies.

SECTION A: BIOMEDICAL TECHNOLOGY PROGRAMME

4. NATIONAL DIPLOMA: BIOMEDICAL TECHNOLOGY (NDBMT1)

4.1. PROGRAMME INFORMATION

Biomedical Technology is a profession of highly knowledgeable and skilled individuals who perform clinical laboratory tests on patient samples. The services offered by Biomedical Technologists are an important component of patient health care, as the results obtained from these laboratory tests are a vital tool in the diagnosis, treatment and prevention of disease. The qualifying student will be able to organize and perform laboratory operations in clinical diagnostic laboratories and related fields in compliance with statutory requirements for ethics, safety and quality assurance. Supervisory, management and research skills are developed.

4.1.1 Duration of the programme

Students in Biomedical Technology/Medical Laboratory Science must attend formal lectures and practical sessions at the Durban University of Technology in all modules for the duration of their studies. The minimum study period is three years, including a six (6) months experiential learning component which occurs in the sixth semester.

Successful applicants for study towards a ND: Biomedical Technology will be accepted into a three-year minimum or an extended, four-year programme of study which comprises of theoretical and practical learning.

4.1.2 Assessment and Moderation

Most subjects in this programme have main and supplementary final examinations. Certain subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

4.1.3 Registration with the Professional Board

As a Student: Within two weeks of registration with the Department, students are required to register as Student Medical Technologist with the Health Professions Council of South Africa as determined in the regulations set out in the Allied Health Service Professions Act, 1982 (Act 63 of 1982) (Regulation R629, Government Gazette No 11221 of 31 March 1988).

As a Graduate

A graduate, upon successful completion of the qualification and the required internship, and having passed all assessment to satisfy the requirements of the Professional Board for Medical Technology, may register as a qualified Biomedical Technologist (will be phased out in 2019) or a Medical Laboratory Scientist (as applicable) with the HPCSA. After registration with the HPCSA,

graduates may work in government, private health care laboratories and research laboratories. Unregistered Biomedical Technologists may work in non-diagnostic laboratories. To practice independently as a Biomedical Technologist, two years post-registration experience is required.

4.1.4 Work Integrated Learning Rules

The WIL component includes a six (6) months placement which occurs in the sixth semester. This is a compulsory component of the programme. The student must be registered at the Durban University of Technology for the duration of this period. The student must comply with the rules and regulations as set out in the Medical Technology laboratory where placed.

4.2 Learning Programme Structure

Code	Subjects	Year of Study	NQF Level	Nated Credits	Pre-req Code
IMET 101	Introduction to Medical Technology	1	5	0.050	None
CSTA101	Calculation and Statistics	la	5	0.100	None
CHMB102	Chemistry	la	5	0.125	None
PYSC105	Physics	la	5	0.100	None
BIOA202	Biochemistry2	Ιb	5	0.125	None
IMMU202	Immunology2	Ιb	5	0.125	None
ANPHII4	Anatomy & Physiology(Module A)	la	5	0.125	None
ANPH124	Anatomy & Physiology(Module B)	Ιb	5	0.125	None
PAPH201	Pathophysiology 2	Ιb	5	0.125	None
BLTT201	Blood Transfusion Technology 2	2a	6	0.125	IMMU202
CEPA 101	Cellular Pathology I	2a	6	0.125	ANPHII4 ANPHI24
CPAT101	Chemical Pathology I	2a	6	0.125	BIOA202, CHMB102
MCGYI0I	Microbiology I	2a	6	0.125	
HAEM203	Haematology 2	2b	6	0.125	BLTT201, PAPH201
CEPA201	Cellular Pathology 2	2b	6	0.125	CEPA101 PAPH201
CPAT202	Chemical Pathology 2	2b	6	0.125	CPATI01 PAPH201
MCGY203	Microbiology 2	2b	6	0.125	MCGY101 PAPH201
HAEM 303	Haematology 3	3a	6	0.125	HAEM203
CEPA 301	Cellular Pathology 3	3a	6	0.125	CEPA201
CPAT303	Chemical Pathology 3	3a	6	0.125	CPAT202
MCGY301	Microbiology 3	3a	6	0.125	MCGY203
LABP301	Laboratory Practice 3	3b	6	0.500	

^{*}A pre-reg means this subject must be passed prior to registration (prerequisite)

a denotes first semester, b denotes second semester

4.3 PROGRAMME RULES

4.3.1 Minimum admission requirements.

In addition to Rule G7, the minimum admission requirement for a student who registers for the Bachelor are:

National Senior Certificate (NSC) with a Bachelor Degree endorsement and must include the following subjects at the stated ratings.

Compulsory Subjects	NSC Rating
English	3
Life Orientation	4
Mathematics	4
Life Science	4
Physical Science	4
And one 20 credit subject	3

Senior Certificate (SC) with matriculation exemption and must include the following subjects at the stated ratings:

COMPULSORY SUBJECTS	HG	SG
Mathematics	D	С
Physical Sciences	D	С
Biology / Life Sciences / Physiology	D	С

Admission requirements based upon Work Experience, Age and Maturity and RPL

The DUT general rules G7 (3) and G7 (8) respectively, will apply.

Admission of International students

The DUT's Admission's Policy for International Students and general rules G4 and G7 (5), apply.

4.3.2 Selection Criteria

In accordance with Rule G5, acceptance into the programme is limited to 30 places. As more qualifying applications are received than can be accommodated, the following selection process will determine placement in the programme:

- All applicants must apply through the Central Applications Office (CAO).
- Initial shortlisting for selection is based on the applicant's academic performance in Grade 12 (Grade 11, or Grade 12 trial marks, will be used for current matriculants).
- Shortlisted students will be invited to undergo placement testing.
- Applicants who pass the placement tests are invited for an interview.
- Provisional acceptance is given to selected applicants awaiting National Senior Certificate (NSC) results. If the final Grade 12 NSC results do not meet the minimum entrance requirements, this provisional acceptance will be withdrawn.
- Final selection for placement will be based on results in the SC / NSC and DUT placement tests as well as on recommendations from the interview panel.

Assessment	Weighting (%)
Results of the Senior Certificate or National Senior Certificate	30%
Placement Testing	35%
Interview Score	35%

4.3.3 Pass Requirements

Notwithstanding the DUT pass requirements (G14 and G15), and those detailed as follows, students are encouraged to apply themselves to their learning, and strive for the best academic results possible in order to adequately prepare themselves for their future careers, and to maximize possible employment opportunities.

- A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Biomedical Technology.
- Promotion to semester 3 of study requires a pass in at least 50% of the
 previous level subjects, i.e. year 1 subjects; notwithstanding
 prerequisites and co-requisites. Students who have passed less than 50%
 of their subjects in a level are considered to be not making satisfactory
 academic progress.
- Promotion to semester 4 of study requires a pass in at least 50% of semester 3 subjects; notwithstanding prerequisites. Students who have passed less than 50% of their subjects in a level are considered to be not making satisfactory academic progress.

- Promotion to semester 5 of study requires a pass in at least 50% of the
 previous level subjects, i.e. semester 4 subjects; notwithstanding
 prerequisites. Students who have passed less than 50% of their subjects
 in a level are considered to be not making satisfactory academic
 progress.
- Prior to commencing with Laboratory Practice 3, a student must have passed all Semester 1 to Semester 4 subjects, and must have obtained a sub minimum of 40 % for: Chemical pathology 3, Cellular pathology 3, Haematology 3 and Microbiology 3.

4.3.4 Re-registration Rules

Rule G16 applies

4.3.5 Exclusion Rules

In addition to Rule G17 the following departmental rule applies:

A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Biomedical Technology. Deregistration from any subject is subject to the provisions of Rule G6 (2).

4.3.6 Interruption of Studies

In accordance with Rule G21A (b), the minimum duration for this programme will be three (3) years of registered study and the maximum duration will be five (5) years of registered study, including any periods of work-integrated learning (WIL). Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

5. NATIONAL DIPLOMA: BIOMEDICAL TECHNOLOGY: EXTENDED CURRICULUM PROGRAMME (NDBMF1) [Phasing out]

5.1. PROGRAMME INFORMATION

Biomedical Technology is a profession of highly knowledgeable and skilled individuals who perform clinical laboratory tests on patient samples. The service offered by Biomedical Technologists is an important component of patient health care, as the results obtained from these laboratory tests are a vital tool in the diagnosis, treatment and prevention of disease. The qualifying student will

be able to organize and perform laboratory operations in clinical diagnostic laboratories and related fields in compliance with statutory requirements for ethics, safety and quality assurance. Supervisory, management and research skills are developed.

5.1.1 Duration of the Programme

Successful applicants for study towards a ND: Biomedical Technology will be accepted into an extended, four-year minimum programme of study. This extended curriculum has been designed in order to enhance student development and to improve the student's chances of successful completion. Students in Biomedical Technology must attend formal lectures and practical sessions at the Durban University of Technology in all subjects for the duration of their studies. The minimum study period for the ND: Biomedical Technology (ECP) is four years, including a six (6) months experiential learning component.

5.1.3 Assessment and Moderation

Most subjects in this programme have main and supplementary final examinations. Certain subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

5.1.4 Registration with the Professional Board

As a Student: Within two weeks of registration with the Department, students are required to register as Student Medical Technologists with the Health Professions Council of South Africa as determined in the regulations set out in the Allied Health Service Professions Act, 1982 (Act 63 of 1982) (Regulation R629, Government Gazette No 11221 of 31 March 1988).

As a Graduate

A graduate, upon successful completion of the qualification and the required internship, and having passed a competency assessment to satisfy the requirements of the Professional Board for Medical Technology, may register as a qualified Biomedical Technologist (as applicable) with the HPCSA. After registration with the HPCSA, graduates may work in government, private health care laboratories and research laboratories. Unregistered Biomedical Technologists may work in non-diagnostic laboratories. To practice independently as a Biomedical Technologist, two years post-registration experience is required.

5.1.5 Work Integrated Learning Rules

The WIL component includes a six (6) months placement which occurs in the eighth semester. This is a compulsory component of the programme. The

student must be registered at the Durban University of Technology for the duration of this period. The student must comply with the rules and regulations as set out in the Medical Technology laboratory where placed.

5.2 LEARNING PROGRAMME STRUCTURE

NATIONAL DIPLOMA: BIOMEDICAL TECHNOLOGY: EXTENDED CURRICULUM PROGRAMME (NDBMFI)

Code	Subjects	Year of Study	NQF Level	Nated Credits	Pre-req Code
FCMR101	Foundation Chemistry	la	5	0.100	none
FPHY101	Foundation Physics	la	5	0.100	none
FLBT101	Laboratory Techniques	2a	5	0.175	none
FBIO202	Foundation Biochemistry	2a	5	0.063	none
FIMM202	Foundation Immunology	2a	5	0.062	none
IMET101	Introduction to Medical Technology	I	5	0.050	none
CSTA101	Calculation and Statistics	Ib	5	0.100	none
CHMY101	Chemistry	Ib	5	0.125	FCMR101
PYSC105	Physics	Ib	5	0.100	FPHY101
BIOA202	Biochemistry2	2b	5	0.062	FBIO202
IMMU202	Immunology2	2b	5	0.063	FIMM202
ANPH114	Anatomy & Physiology(Module A)	2a	5	0.125	none
ANPH124	Anatomy & Physiology(Module B)	2b	5	0.125	none
PAPH201	Pathophysiology 2	2b	5	0.075	none
BLTT201	Blood Transfusion Technology 2	3a	6	0.100	IMMU202
CEPA101	Cellular Pathology I	3a	6	0.100	ANPHII4, ANPHI24,
CPATI01	Chemical Pathology I	3a	6	0.100	BIOA202, CHMB102
MCGY101	Microbiology I	3a	6	0.100	
HAEM203	Haematology 2	3b	6	0.100	BLTT201, PAPH201
CEPA201	Cellular Pathology 2	3b	6	0.100	CEPA 101 PAPH201
CPAT202	Chemical Pathology 2	3b	6	0.100	CPATI0I PAPH20I
MCGY203	Microbiology 2	3b	6	0.100	MCGY101 PAPH201
HAEM303	Haematology 3	4a	6	0.100	HAEM203
CEPA301	Cellular Pathology 3	4a	6	0.100	CEPA201
CPAT303	Chemical Pathology 3	4a	6	0.100	CPAT202
MCGY301	Microbiology 3	4a	6	0.100	MCGY203
LABP301	Laboratory Practice 3	4 b	6	0.475	nil

^{*}A pre-req means this subject must be passed prior to registration (prerequisite)

5.3 PROGRAMME RULES

5.3.1 Minimum Admission Requirements

In addition to Rule G7, the minimum admission requirement for a student

a denotes first semester, b denotes second semester

who registers for the National Diploma: Biomedical Technology are: National Senior Certificate (NSC) with a Bachelor Degree endorsement and must include the following subjects at the stated ratings.

Compulsory Subjects	NSC Rating
English	3
Life Orientation	4
Mathematics	4
Life Science	4
Physical Science	4
And one 20 Credit Subject	3

Senior Certificate (SC) with matriculation exemption and must include the following subjects at the stated ratings:

Compulsory Subjects	HG	SG
Mathematics	D	С
Physical Sciences	D	С
Biology / Life Sciences / Physiology	D	С

Admission requirements based on work experience, age & maturity; and recognition of prior earning (RPL).

The DUT general rules G7 (3) and G7 (8) respectively, will apply.

Admission of international students

The DUT's Admission's Policy for International Students and general rules G4 and G7 (5), apply.

5.3.2 Selection Criteria

In accordance with Rule G5, acceptance into the ECP programme is limited to 15 places.

As more qualifying applications are received than can be accommodated, the following selection process will determine placement in the programme:

- All applicants must apply through the Central Applications Office (CAO).
- Initial shortlisting for selection is based on the applicant's academic performance in Grade 12 (Grade 11, or Grade 12 trial marks, will be used for current matriculants).
- Shortlisted students will be invited to undergo placement testing.
- Applicants who pass the placement tests are invited for an interview.
- Provisional acceptance is given to selected applicants awaiting National Senior Certificate (NSC) results. If the final Grade 12 NSC results do not meet the minimum entrance requirements, this provisional acceptance will be withdrawn.
- Final selection for placement will be based on results in the SC / NSC and DUT placement tests as well as on recommendations from the interview panel.

Assessment	Weighting (%)
Results of the Senior Certificate or National Senior Certificate	30%

Placement Testing	35%
Interview Score	35%

5.3.3 Pass Requirements

Notwithstanding the DUT pass requirements (G14 and G15), and those detailed as follows, students are encouraged to apply themselves to their learning, and strive for the best academic results possible in order to adequately prepare themselves for their future careers, and to maximize possible employment opportunities.

- A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Biomedical Technology.
- Promotion to semester 3 of study requires a pass in Foundation
 Chemistry and Foundation Physics and at least 1 mainstream subject of
 the previous level, i.e. Introduction to Medical Technology, Calculations
 and Statistics, Physics 1 or Chemistry 1. Students who have passed less
 than 50% of their subjects in a level are considered not to be making
 satisfactory academic progress.
- Promotion to semester 4 of study requires a pass in Foundation Immunology, Foundation Biochemistry and Laboratory Techniques, and all year I subjects. Students who have passed less than 50% of their subjects in a level are considered not to be making satisfactory academic progress.
- Promotion to semester 5 of study requires a pass in at least 50% of the
 previous level subjects, i.e. semester 4 subjects. (Prerequisites have to be
 satisfied). Students who have passed less than 50 % of their subjects in a level
 are considered not to be making satisfactory academic progress.
- Promotion to semester 6 of study requires a pass in at least 50% of the
 previous level subjects, i.e. semester 5 subjects; notwithstanding prerequisites.
 Students who have passed less than 50% of their subjects in a level are
 considered to be not making satisfactory academic progress.
- Promotion to semester 7 of study requires a pass in at least 50% of the
 previous level subjects, i.e. semester 6 subjects; notwithstanding prerequisites.
 Students who have passed less than 50% of their subjects in a level are
 considered to be not making satisfactory academic progress.
- Prior to commencing with Laboratory Practice 3, a student must have passed all Semester 1 to Semester 4 subjects, and must have obtained a sub minimum of 40% for: Chemical pathology 3, Cellular pathology 3, Haematology 3 and Microbiology 3.

5.3.4 Re-registration Rules

Rule G16 applies

5.3.5 Exclusion Rules

In addition to Rule G17, the following departmental rule applies:

- A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Biomedical Technology.
- Deregistration from any subject is subject to the provisions of Rule G6 (2).

5.3.6 Interruption of Studies

In accordance with Rule G21A (b), the minimum duration for this programme will be four (4) years of registered study and the maximum duration will be five (5) years of registered study, including any periods of WIL. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

6. BACHELOR OF TECHNOLOGY: BIOMEDICAL TECHNOLOGY (BTBMT2) [Phasing out]

6.1 PROGRAMME INFORMATION

The qualifying Student will be able to organize and perform laboratory operations in clinical diagnostic laboratories and related fields in compliance with statutory requirements for ethics, safety and quality assurance.

Supervisory, management and research skills are developed. They will be able to integrate laboratory tests and results with pathophysiological conditions. Students will be able to conduct research grounded in a deep knowledge of their area of specialization. Management skills are developed with a view to encouraging entrepreneurial development and business management.

After registration with the HPCSA, they may work in government, private and research laboratories. To practice independently as a Medical Technologist, two (2) years post-registration experience is required. Unregistered Biomedical Technologists may work in non-diagnostic laboratories.

Assessment and Moderation

Most subjects in this programme have main and supplementary final examinations. Certain subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

6.2. **Learning Programme Structure for the** Bachelor of Technology. Biomedical Technology (BTBMT2)

Code	Subjects	Year of Study	NQF Level	NATED Credits
RMTQ 201	Research Methods and Techniques	1	7	0.125
MOLE401	Molecular Biology IV	1	7	0.250
LABM 201	Laboratory Management	2	7	0.125
IPAT401	Integrated Pathophysiology IV	2	7	0.250
RPBM101	Research Project	2	7	0.250

6.3 PROGRAMME RULES

6.3.1 Minimum Admission Requirements & Selection Criteria

In addition to Rule G23(1), G3, G4 and G7, students applying for this qualification must be in possession of a ND: Biomedical Technology or National Diploma: Medical Technology and proof of registration with the HPCSA in the Medical Technology category or have granted status or advanced standing according to rule G10. Applicants with a ND: Medical Technology have to demonstrate competence in the fundamentals of Biochemistry to the satisfaction of the department. Additional credits may have to be taken if this competence is not demonstrated

In accordance with Rule G5, acceptance into the programme is limited to 20 places and entry to the BTech programme is not automatic. As more qualifying applications are received than can be accommodated, the following selection criteria will determine entry into the programme, with the 20 highest ranking candidates gaining entry into the programme:

- Submission of BTech application forms by due date.
- Applicant's academic performance in the ND: Biomedical Technology see ranking criteria below.
- Workplace experience (post National Diploma)

6.3.2 THE RANKING CRITERIA

I. Average marks of the final year of the National Diploma					
2. Years to complete the	National Diploma qualifica	ntion			
Minimum duration	Minimum duration	Minimum duration	Minimum duration		
+ 3yrs	+ 2 yrs	+ l yr			
0	I	3	5		
3. Workplace experience post National Diploma					
0-I year	I-3 years	3-5years	> 5years		
0		3	5		

An applicant's ranking criteria is determined by the total points score obtained by the addition of the scores obtained in the individual ranking criteria, as shown in the **example** in the table below

Critoria	Ranking score (points)
Average marks of the final year	60
National Diploma completed in minimum duration	5
Workplace experience (Diploma just completed)	0

Total 65

To gain access into the BTech programme, a student must have a minimum of 60 points.

(w.e.f. 28/08/2014)

6.3.3 Pass Requirements

In addition to Rule G14 and G15, the following rules apply. Students are encouraged to apply themselves to their studies, and strive for the best academic results possible in order to adequately prepare themselves for their future careers.

6.3.4 Re-registration Rules

Rule G16 applies.

6.3.5 Exclusion Rules

Rule G17 applies.

6.3.6 Interruption of Studies

In accordance with Rule G23A, the minimum duration for this programme will be one (I) year of registered study and the maximum duration will be two (2) years of registered study. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

7. BACHELOR OF HEALTH SCIENCES IN MEDICAL LABORATORY SCIENCE 7.1. PROGRAMME INFORMATION

The Bachelor of Health Sciences in Medical Laboratory Sciences is a professional degree with a minimum number of 480 SAQA credits and is offered at NQF level 8 of the HEQSF. Whilst the majority of the modules are core, some of them are generic in nature and these are offered by both the Faculty of Health Sciences and the institution at large. At each level of study the student has an opportunity to choose from at least two of the elective modules and students will also register for research modules.

Medical laboratory Science is a profession of highly knowledgeable and skilled individuals who perform diagnostic tests on patient samples in a clinical laboratory and are skilled to conduct research. The service offered by Medical Laboratory Scientists is an important component of patient health care, as the results obtained from these laboratory tests are a vital tool in the diagnosis, treatment and prevention of disease. The qualifying student will be able to organize and perform laboratory operations in clinical diagnostic laboratories and related fields in compliance with statutory requirements for ethics, safety and quality assurance. Supervisory, management and research skills are developed.

7.1.1 Duration of the Programme

Successful applicants for study towards a BHSc: Medical Laboratory Science will be accepted into a four-year minimum programme of study. This four year degree level 8 curriculum has been designed in order to enhance student development produce a holistic, diagnostic and research grounded graduate who will directly articulate to the Master's degree.

Students in Medical Laboratory Science must attend formal lectures and practical sessions at the Durban University of Technology in all modules for the duration of their studies. The minimum study period for the BHSc: Medical Laboratory Sciences is four years, including a six (6) months of work integrated learning component and one year clinical training in a Clinical Diagnostic Laboratory. The maximum period of study for this four year degree is six years.

7.1.2 Assessment and Moderation

Most modules in this programme have main and supplementary final examinations. Certain modules in this programme do not have a final examination. The results for these modules are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each module at the back of this handbook. Moderation follows the DUT requirements.

7.1.3 Registration with the Professional Board

As a Student: Within two weeks of registration with the Department, students are required to register as Student Medical Laboratory Scientist with the Health Professions Council of South Africa as determined in the regulations set out in the Allied Health Service Professions Act, 1982 (Act 63 of 1982) (Regulation R629, Government Gazette No 11221 of 31 March 1988).

As a Graduate

A graduate, upon successful completion of the qualification and the required industry exposure, and having passed a final competency assessment in the fourth year to satisfy the requirements of the Professional Board for Medical Laboratory Science, may register as a qualified Medical Laboratory Scientist (as applicable) with the HPCSA. After registration with the HPCSA, graduates may work in government, private health care laboratories and research laboratories. Unregistered Medical Laboratory Scientist may work in non-diagnostic laboratories. To practice independently as a Medical Laboratory scientist, two years post-registration experience is required.

7.1.4 Work Integrated Learning Rules

The WIL component includes a six (6) months placement which occurs in the sixth semester and a one year placement which occurs in the seventh and eighth semesters. This is a compulsory component of the programme. The

student must be registered at the Durban University of Technology for the duration of this period. The student must comply with the rules and regulations as set out in the diagnostic laboratory where placed.

7.2 Learning Programme Structure: Bachelor of Health Sciences in Medical Laboratory Science

Module code	Module Title	Year of Study	HEQSF level	HEQSF Credit	Period of Study	HEMIS credits	Pre- requisite
CMTR101	Chemistry	I	5	16	a	0.111	
PHISTIT	Physics (Module 1)	1	5	8	a	0.057	
PHIS121	Physics (Module 2)	I	5	8	b	0.057	
FMLS101	Fundamentals of Medical Laboratory Science	I	5	12	a	0.086	
STTS101	Statistics	I	5	8	I p	0.051	
ANPA102	Anatomy and Physiology 1A	I	5	12	a	0.086	
ANPB102	Anatomy and Physiology 1B	I	5	12	l p	0.086	
CBIO101	Cell Biology	I	5	16	a	0.112	
IMLG101	Immunology	I	5	16	a	0.111	
CSTN101	Cornerstone 101	I	5	12	a	0.094	
VWKPI0I CLDVI0I	Values in the workplace Cultural Diversity	-	5	8	a	0.067	
EVAH101 IGSH101	Environmental Awareness for healthcare Practitioners Issues of Gender & Society within Health care	1	5	12	1	0.082	
CLCM101	Clinical Chemistry I	2	6	16	2 a	0.107	Cell Biology
MMCR101	Medical Microbiology I	2	6	8	2 a	0.053	Anatomy & Physiology
MDMA201	Medical Microbiology IIA	2	7	16	2 b	0.106	Medical Microbiology I
HMTL101	Haematology I	2	6	16	2 b	0.107	Immunology
IMHT101	Immunohaematology I	2	6	16	2	0.106	Immunology
HPTH101	Histopathology I	2	6	16	2 b	0.106	Anatomy & Physiology
CYTLI0I	Cytology I	2	6	16	2 b	0.106	Anatomy & Physiology
MLCB101	Molecular Biology	2	6	8	2 a	0.053	Cell Biology
FPTH101	Fundamentals of Pathology	2	6	8	2	0.054	Anatomy & Physiology
SYSP101	Systemic Pathophysiology	2	6	8	2 b	0.054	Anatomy & Physiology

TENEI01	The entrepreneurial edge	2	6	8	32 a	0.067	
GENV101	The global environment						
EQDVI01	Equality and diversity						
CLCM201	Clinical Chemistry II	3	7	16	3 a	0.138	Clinical
							Chemistry I
MDMB201	Medical Microbiology IIB	3	7	16	3 a	0.138	Medical
1 101 10201	riedical riici obiology iib	3	,	10	J	0.130	Microbiology
							2A
HMTL201	Haematology II	3	7	16	3 a	0.138	Haematology
HI11 LZ01	maematology II	3	/	16	3 ª	0.136	Haematology
	<u> </u>		_				<u> </u>
CYTL201	Cytology II	3	7	16	3 a	0.138	Cytology I
CLLP101	Clinical Laboratory	3	7	16	3 a	0.139	All year I
	Practice I						and year 2
							modules
PMTG101	Principles of management	3	7	8	3 ь	0.068	
RSJS101	Restorative justice	3	7	8	3 a	0.069	
EDÚT101	Educational Techniques	3	7	12	3 a	0.103	
ETMH101	Ethics and Medical Law					******	
PRRSIOI	Principles of Research	3	7	8	3 ь	0.069	
RPJA101	Research Project Module	4	8	20	4 a	0.167	Principles of
KrjATUT	A Research Project Module	4	٥	20	4 "	0.167	
BBIBLOL	A		_		4.	0.120	Research
RPJB101	Research Project Module	4	8	16	4 b	0.139	Principles of
	В						Research
IPPA101	Integrated	4	8	12	4 a	0.089	Clinical
	Pathophysiology Module						Chemistry 2
	Α						Medical
							Microbiology
							2
							Haematology
							2
							Cytology 2
IPPB102	Integrated	4	8	8	4 b	0.086	Clinical
	Pathophysiology Module B						Chemistry 2
	i acropinysiology i rodaic 2						Cytology 2 2
							Haematology
							2
							Medical
							Microbiology
							٠,
							2
LDTMIOL	1 . M	4	8	12	4.	0.106	D : : 1 (
LBTM101	Laboratory Management	4	8	12	4 a	0.106	Principles of
							management
	Clinical Laboratory	4	8			0.433	
	Practice 2: includes the						
	following specialisation						
	options from I - 10						
	below (the student will						
	have to select one of						
	these advanced						
	specialization modules at						
	52 credits):						
	32 ci cuits).						
1					ĺ		1
1					ĺ		1
1					ĺ		1
1					ĺ		1
1					ĺ		1
					ĺ		1
					ĺ		1
					ĺ		1
1					ĺ		1
CPHA101	Clinical Pathology Module	4	8	28	4 a		Clinical
1							

	A					Chemistry 2
						Medical
						Microbiology
						2 Haematology
						2
						Cytology
CPHB101	Clinical Pathology Module	4	8	24	4 b	Clinical
	В	-				Chemistry 2
						Medical
						Microbiology
						2
						Haematology 2
						Cytology 2
CLCA301	Clinical Chemistry IIIA	4	8	28	4 a	Clinical
CLCB301	Clinical Chemistry IIIB	4	8	24	4 b	Chemistry 2
	J	-				Medical
						Microbiology
						2
						Haematology
						2
MDMA301	Medical Microbiology IIIA	4	8	28	4 a	Cytology Clinical
MDMB301	Medical Microbiology IIIB	4	8	24	4 b	Chemistry 2
110110001	ricalcal File oblology IIID	•	Ĭ			Medical
						Microbiology
						2
						Haematology
						Compleme 2
CYTA301	Cytology IIIA	4	8	28	4 a	Cytology 2 Clinical
CYTB301	Cytology IIIB	4	8	24	4 b	Chemistry 2
C112301	Cytology IIIB	•	Ĭ			Medical
						Microbiology
						2
						Haematology
						2 Complemy 2
HMTA301	Haematology IIIA	4	8	28	4 a	Cytology 2 Clinical
HMTB301	Haematology IIB	4	8	24	4 b	Chemistry 2
111112301	nacmacology no	•	Ĭ			Medical
						Microbiology
						2
						Haematology
						2
HISA201	Histopathology IIA	4	8	28	4 a	Cytology 2 Clinical
HISAZUI	i iistopaulology IIA	7	°	20	1 "	Chemistry 2
			1			Medical
						Microbiology
						2
			1			Haematology
						2
HISB201	Histopathology IIP	4	8	24	4 b	Cytology 2 Clinical
IHMA201	Histopathology IIB Immunohaematology IIA	4	8	28	4a	Chemistry 2
11 11/1/201	minunonaematology IIA	T		20	'-	Medical
1			1			Microbiology
			1			2
			1			Haematology
			1			2
						Cytology 2

IHMB201	Immunohaematology IIB	4	8	24	4 b	Clinical
						Chemistry 2
						Medical
						Microbiology
						2
						Haematology
						2
						Cytology 2

^{*}A pre-req means this subject must be passed prior to registration (prerequisite) a denotes first semester, b denotes second semester

7.3 Programme Rules

7.3.1 Minimum Admission Requirements

In addition to Rule G7, the minimum entrance requirement is a National Senior Certificate (NSC) valid for entry into a Bachelor's Degree endorsement and must include the following at the stated minimum ratings below:

NSC REQUIREMENTS	SENIOR CERTIFICATE REQUIREMENTS		
Compulsory subjects	NSC Rating	Compulsory subjects	SC Symbol
English (Home language) OR English (1st additional language)	4	English HG	D
Mathematics	4	Mathematics HG	D
Life Sciences	4	Biology HG	D
Physical Sciences	4	Physical Science HG	D
And two other 20 credit subjects of which only one may be a language	3		

In addition to Rule G7, the minimum entrance requirement for a holder of a valid National Certificate (Vocational) for entry into a Bachelor's Degree must include the following subjects as the stated minimum ratings below:

Compulsory Subjects	NC (V)		
English	60%		
Mathematics	60%		
Physical Sciences	70%		
Life sciences	70%		
Four other subjects, only one of which may be a language	60%		

7.3.2 Minimum Admission Requirements in respect of Work Experience, Age, Maturity, RPL and International Students

The DUT general rules G7(3) and G7(8) respectively will apply.

The DUT's Admissions Policy for International Students and General Rules G4 and G7 (5) will apply.

7.3.3 Selection Criteria

All applicants must apply through the Central Applications Office (CAO). In accordance with Rule G5, acceptance into the programme is limited. Since more

In accordance with Rule G5, acceptance into the programme is limited. Since more applications are received than can be accommodated, the following selection process will apply:

- Initial shortlisting for selection is based on the applicant's academic performance in Grade 12 (Grade 11, or Grade 12 trial marks, will be used for current grade 12 learners).
- Applicants obtaining more than 23 points in their matriculation examination stand a better chance of selection.
- The point scores for each National Senior Certificate (NSC) subject or the Senior Certificate (SC) results is obtained by using the table below:

Senior Certificate (SC)

Symbol	Α	В	С	D	Е	F
Higher Grade	8	7	6	5	4	3
Standard Grade	6	5	4	3	2	ı

National Senior Certificate (NSC)

8	=	90 – 99%
7	=	80 – 89%
6	=	70 – 79%
5	=	60 – 69%
4	=	50 – 59%
3	=	40 – 49%
2	=	30 – 39%
ı	=	0 –29%

No points are allocated for ten (10) credit subjects.

- Applicants who meet the minimum departmental admission requirements for the Bachelor of Health Sciences in Medical Laboratory Science will be ranked according to the points scored in Grade 11 and Grade 12 and may be invited to participate in the selection process.
- The percentage weighting assigned to each of these scores will be as follows:

Assessment	Weighting
Results of the Senior Certificate or National Senior Certificate	60%
Interview Score	40%

- Selected applicants will be placed into either the four-year degree or an Extended Curriculum Programme.
- Provisional acceptance is given to selected applicants awaiting (NSC) and National Certificate (Vocational) results. If the final Grade 12 NSC/ National Certificate (Vocational) results do not meet the minimum entrance requirements, this provisional acceptance will be withdrawn.

7.3.4 Pass Requirements

Notwithstanding the DUT pass requirements (G14 and G15), and those detailed as follows, students are encouraged to apply themselves to their learning, and strive for the best academic results possible in order to adequately prepare themselves for their future careers, and to maximize possible employment opportunities.

- In addition to the DUT General Rule G17*, a first year student who fails six or more of the modules with an average of less than 40% in the failed modules during that year is not permitted to re-register for the Bachelor of Health Sciences in Medical Laboratory Science programme. A student who fails 6 modules with an average of 40% in the failed modules, is not precluded from proceeding to the second semester. De-registration from any module is subject to the provisions of Rule G6 (2)*.
- Promotion to semester 3 of study requires a pass in at least 50% of the
 previous level subjects, i.e. year 1 subjects; notwithstanding
 prerequisites and co-requisites. Students who have passed less than 50%
 of their modules in a level are considered to be not making satisfactory
 academic progress.
- Promotion to semester 4 of study requires a pass in at least 50% of semester 3 modules; notwithstanding prerequisites. Students who have passed less than 50% of their modules in a level are considered to be not making satisfactory academic progress.
- Promotion to semester 5 of study requires a pass in at least 50% of the
 previous level subjects, i.e. semester 4 subjects; notwithstanding
 prerequisites. Students who have passed less than 50% of their subjects
 in a level are considered to be not making satisfactory academic
 progress.
- Prior to commencing with Clinical Laboratory Practice I, a student must have passed all Semester I to Semester 4 subjects, and must have obtained a sub minimum of 40% for any of the following modules:

- Chemical pathology 2, Cytology 2, Haematology 2, Medical Microbiology 2B, Histopathology 2 and Immunohaematology 2.
- Promotion to semester 7 and 8 requires successful completion of all semester 1 to 6 modules.

7.3.5 Re-registration Rules

Rule G16 applies

7.3.6 Exclusion Rules

In addition to Rule G17, the following departmental rule applies:

- A first year student who fails six or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: BHSc: Medical Laboratory Science.
- Deregistration from any subject is subject to the provisions of Rule G6 (2).

7.3.7 Interruption of Studies

In accordance with Rule G21A (b), the minimum duration for this programme will be four (4) years of registered study and the maximum duration will be five (5) years of registered study, including any periods of WIL. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

8 MASTER OF HEALTH SCIENCES IN MEDICAL LABORATORY SCIENCE (MHMLSI)

8.1 PROGRAMME INFORMATION

This full research qualification is aligned to Rule G24 and the guidelines in the Post Graduate Student Handbook.

- The Student who successfully completes this qualification will be able to apply advanced problem solving skills and critical, reflective thinking to perform independent research in a chosen field and report their findings in a dissertation that meets the accepted criteria and ethical principles for the profession. In this way they will make a contribution to the existing body of knowledge and initiate change that will help develop and advance the profession of medical technology.
- The qualifying Student will be able to conduct independent research under minimal guidance in a chosen field, and contribute to knowledge production in that field. The research problem, its justification, process and outcome is reported in a dissertation which complies with the generally accepted norms for research at that level.

8.1.1 Assessment and Moderation

In addition to Rule G24 (4), postgraduate assessment of dissertations will be aligned to Postgraduate policies and guidelines. Please refer to the General Student Handbook and the Postgraduate Student Handbook.

8.2 LEARNING PROGRAMME STRUCTURE

Code	Module	Year of Study	Assessment Type	NATED Credits	Pre-requisites	Co-requisites
MHMLSI	Dissertation	2	External Examination	1.0	None	none

8.3 PROGRAMME RULES

8.3.1 Minimum Admission Requirements

In addition to the General Handbook for Students Rule G24 (I), candidates must be in possession of a Bachelor's Degree in Biomedical Technology (NQF Level 8), or must have been granted conferment of status according to Rule G10A.

Candidates may also apply for admittance via Recognition of Prior Learning (RPL) in accordance with Rule G7 (8) and / or G10B.

8.3.2. Selection Criteria

In accordance with Rule G5, acceptance into the programme is limited and entry into the Master of Health Sciences in Medical Laboratory Practice is not automatic. Students are selected into the programme once they have completed an intention to study / a concept paper and the department has discussed the viability of the proposed topic for the Masters Qualification. The

intention to study/ concept page must include the following: Problem statement or Title of the intended study, Objectives / sub-problems / Research Questions, Rationale/motivation to do the study, Brief literature review, Brief methodology.

8.3.3 Pass Requirements

Rule G24 and the Postgraduate Student Guide apply. Students are encouraged to apply themselves to their research, and strive for the best academic results possible in order to adequately prepare themselves for their future careers.

8.3.4 Re-registration Rules

Rule G24 (2), Rule G26 (5) and the Postgraduate Student Guide apply.

8.3.5 Exclusion Rules

Rule G24 (I) (d); Rule G24 (2), and the Postgraduate Student Handbook apply.

8.3.6 Interruption of Studies

In accordance with Rule G24, the minimum duration for this programme will be one (I) year of registered study and the maximum duration will be three (3) years of registered study. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

9. DOCTOR OF MEDICAL LABORATORY SCIENCE (DRMLSI)

9.1 PROGRAMME INFORMATION

This full research qualification is aligned to Rule G25 and G26 and the guidelines in the Post Graduate Student Handbook. The purpose of this qualification is to ensure that the student who successfully completes this qualification will be able to apply advanced problem-solving skills and critical, reflective thinking to perform independent research in a chosen field and report their findings in a dissertation that meets the accepted criteria and ethical principles for the profession. In this way they will make a contribution to the existing body of knowledge and initiate change that will help develop and advance the profession of medical technology.

9.1.1 Assessment and Moderation

Post graduate assessment will be aligned to Postgraduate policies and guidelines. Rule G25 (4) and the Postgraduate Student Handbook apply.

9.2 PROGRAMME LEARNING STRUCTURE

Code	Module	Year of Study	Assessment Type	NATED Credits	Pre- requisites	Co-requisites
DRMLSI	Dissertation	3	External Examination	2.0	None	none

9.3. PROGRAMME RULES

9.3.1 Minimum Admission Requirements

In addition to Rule G25 (1), persons must be in possession of a Master's degree in Biomedical Technology (NQF 9), or have been granted status or advanced standing according to Rule G10. Please also refer to the Postgraduate Student Handbook.

9.3.2 Selection criteria

Students are selected into the programme once they have completed an intention to study and the department has discussed the viability of the proposed topic for the qualification. A sound knowledge of the fundamental principles and concepts of research and statistical methods is required.

9.3.3 Re-registration Rules

Rule G26 (5) and the Postgraduate Student Handbook apply.

9.3.4 Exclusion Rules

Rules G25 (2)(b; c(ii)) in the General Student Handbook; and the Postgraduate Student Handbook apply.

9.3.5 Interruption of Studies

In accordance with Rule G25 (2), the minimum duration for this programme will be two (2) years of registered study and the maximum duration will be four (4) years of registered study. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration. Please refer to the Postgraduate Student Handbook.

SECTION B: CLINICAL TECHNOLOGY PROGRAMMES 10 NATIONAL DIPLOMANATIONAL DIPLOMA: CLINICAL TECHNOLOGY (NDCLT1)

This programme is being phased out from 2017-2021: NO INTAKE

10.1 PROGRAMME INFORMATION

This qualification will enable the Students to acquire the necessary knowledge, skills, attitudes and values to practice as a Clinical Technologist in one of the following specialist categories: Cardiology, Cardiovascular Perfusion, Critical Care, Nephrology, Neurology, Pulmonology or Reproductive Biology. They will be able to perform procedures in one of the above seven specialist categories in order to contribute in the diagnosis and treatment of various patho-physiological conditions in conjunction with other designated health care professionals. They also perform organ system support, diagnostic, therapeutic and corrective procedures on patients using specialized health technology and techniques for the treatment of physiological dysfunction.

10.1.1 Duration of the programme

The programme consists of three years full-time study at the Durban University of Technology. The third year is composed of the Work Integrated learning (WIL) component, where a student will choose one of seven categories and study the major specialist subjects appropriate to the chosen category. The categories are as follows: Cardiology, Cardio-Vascular Perfusion, Critical Care, Nephrology, Pulmonology, Reproductive Biology and Neurophysiology.

The latter must be done at a training unit approved by the Health Professions Council of South Africa.

10.1.2 Assessment and Moderation

Some subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Other subjects do have final examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

10.1.3 Registration with the Professional Board

As a Student: On enrolment, it is mandatory that a student register as a student Clinical Technologist with the Health Professions Council of South Africa as determined in the regulations set out in the Government Gazette (No. R.1608 dated 24 July 1987).

As a Graduate: A graduate who has completed the qualification successfully, and has complied with all the conditions as set out by the HPCSA,

may register as a qualified Clinical Technologist with the Health Professions Council of South Africa in terms of the current rules for registration.

10.1.4 Work-Integrated Learning Period (WIL)

WIL will run concurrently with the specialist subjects in the third year of study, at a training unit approved by the Health Professions Council of South Africa (HPCSA). During WIL, students would be required to pass the Competency Based Test (CBT) with a minimum mark of 70%, as a Board requirement.

10.2. PROGRAMME LEARNING STRUCTURE

National Diploma: Clinical Technology

Code	Subjects	Year of Study	NQF Level	Nated Credits	SAQA credits	Pre-req Code
ANAYI0I	Anatomy I	I	5	0.250	30	None
CHMB102	Chemistry I	I	5	0.125	15	None
CAPP101	Computer Appl I	I	5	0.125	15	None
PSIO 102	Physiology I	I	5	0.250	30	None
CSTA101	Calculations & Stats	I	5	0.125	15	None
PYSC 105	Physics I	I	5	0.125	15	None
ANPH202	Anatomy & Physio 2	2	6	0.250	30	PSIO 102, ANAY 10 I
BAPO201	Biomedical Apparatus	2	6	0.250	30	None
OSPP201	Org & Systems Pathophysiology	2	6	0.250	30	PSIO102, ANAY101
PHAR201	Pharmacology 2	2	5	0.125	15	None
PYDNI0I	Psychodynamics	2	5	0.125	15	None
CPAB301	*Cardiology: Biomedical Apparatus 3	3	6	0.350	42	All level I & 2 subjects
CACP310	*Cardiology: Clinical Practice 3	3	6	0.350	42	All level 1 & 2 subjects
CCTP310	*Cardiology: Clinical Tech Practice 3	3	6	0.300	36	All level I & 2 subjects
CCBA301	*Critical Care: Biomedical Apparatus 3	3	6	0.350	42	All level 1 & 2 subjects
CCC301	*Critical Care: Clinical Practice 3	3	6	0.350	42	All level 1 & 2 subjects
CTPR301	*Critical Care: Clinical Tech. Prac. 3	3	6	0.300	36	All level 1 & 2 subjects
NEAP301	*Nephrology: Biomedical Apparatus 3	3	6	0.350	42	All level 1 & 2 subjects
NCLI301	*Nephrology: Clinical Practice 3	3	6	0.350	42	All level I & 2 subjects
NCTP301	*Nephrology: Clinical Tech. Prac. 3	3	6	0.300	36	All level I & 2 subjects
NBMA301	*Neurophysiology: Biomedical Apparatus 3	3	6	0.350	42	All level I & 2 subjects
NCLP301	*Neurophysiology: Clinical Practice 3	3	6	0.350	42	All level 1 & 2 subjects
NTPR301	*Neurophysiology: Clinical Tech. Prac. 3	3	6	0.300	36	All level 1 & 2 subjects
FBAP301	*Perfusion: Biomedical Apparatus 3	3	6	0.350	42	All level I & 2 subjects

PCTP301	*Perfusion: Clinical Practice 3	3	6	0.350	42	All level I & 2 subjects
PCTP301	*Perfusion: Clinical Tech Prac 3	3	6	0.300	36	All level 1 & 2 subjects
PBAP301	*Pulmonology: Biomedical Apparatus 3	3	6	0.350	42	All level 1 & 2 subjects
PCLP301	*Pulmonology: Clinical Practice 3	3	6	0.350	42	All level 1 & 2 subjects
PTPR301	*Pulmonology: Clinical Tech Prac 3	3	6	0.300	36	All level I & 2 subjects
RBAP301	*Reproduction: Biomedical Apparatus 3	3	6	0.350	42	All level 1 & 2 subjects
RCPR301	*Reproduction: Clinical Practice 3	3	6	0.350	42	All level 1 & 2 subjects
RTPR301	*Reproduction: Clinical Tech Prac 3	3	6	0.300	36	All level I & 2 subjects

^{*} Elective Specialist Category Subjects

10.3 PROGRAMME RULES

10.3.1 Minimum Admission Requirements

In addition to Rule G7, the minimum admission requirement for a student who registers for the National Diploma: Biomedical Technology are: National Senior Certificate (NSC) with a Bachelor Degree endorsement and must include the following subjects at the stated ratings.

Compulsory Subjects	NSC Rating
English	3
Life Orientation	4
Mathematics	4
Life Science	4
Physical Science	4
And one 20 Credit Subject	3

Senior Certificate (SC) with matriculation exemption and must include the following subjects at the stated ratings.

Compulsory Subjects	HG	SG
Mathematics	D	С
Physical Sciences	D	С
Biology / Life Sciences / Physiology	D	С

10.3.1.1 Admission requirements based on work experience, age & maturity; and recognition of prior earning (RPL).

Rules G7 (3) and G7 (8) respectively, will apply.

10.3.1.2 Admission of international students

The DUT's Admission's Policy for International Students and general rules G4 and G7 (5), apply.

10.3.2 Selection Criteria

In accordance with Rule G5, acceptance into the programme is limited to 30 places. As more qualifying applications are received than can be accommodated, the following selection process will determine placement in the programme:

- All applicants must apply through the Central Applications Office (CAO).
- Initial shortlisting for selection is based on the applicant's academic performance in Grade 12 (Grade 11, or Grade 12 trial marks, will be used for current matriculants).
- Shortlisted students will be invited to undergo placement testing.
- Applicants who pass the placement tests are invited for an interview.
- Provisional acceptance is given to selected applicants awaiting National Senior Certificate (NSC) results. If the final Grade 12 NSC results do not meet the minimum entrance requirements, this provisional acceptance will be withdrawn.

10Final selection for placement will be based on results in the SC / NSC and DUT placement tests as well as on recommendations from the interview panel.

Assessment	Weighting (%)
Results of the Senior Certificate or National Senior	30%
Certificate	
Placement Testing	35%
Interview Score	35%

10.3.3 Pass Requirements

Notwithstanding the DUT pass requirements (G14 and G15), and those detailed as follows, students are encouraged to apply themselves to their learning, and strive for the best academic results possible in order to adequately prepare themselves for their future careers, and to maximize possible employment opportunities. The General rules (G5) and in terms of Rule G7 apply to the National Diploma: Clinical technology.

10.3.4 Re-registration Rules

Rule G16 in the General Handbook applies.

10.3.5 Exclusion Rules

In addition to Rule G17, the following programme rule applies:

A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Clinical Technology. Deregistration from any subject is subject to the provisions of

Rule G6 (2).

10.3.6 Interruption of Studies

In accordance with Rule G21A (b), the minimum duration for this programme will be three (3) years of registered study and the maximum duration will be five (5) years of registered study, including any periods of WIL. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

10.3.7 Work Integrated Learning Rules (WIL)

In addition to Rule G28, the following programme rules apply:

The student must comply with the rules and regulations as set out in the Industrial Environment where placed.

Students who have not passed all first and second year subjects will not be placed for Work Integrated Learning (WIL).

(wef November 2015)

II NATIONAL DIPLOMA: CLINICAL TECHNOLOGY: EXTENDED CURRICULUM PROGRAMME (NDCLF2)

(This programme is being phased out from 2017-2021) – NO NEW STUDENTS INTAKE

III PROGRAMME INFORMATION

Successful applicants for study towards a ND: Clinical Technology will be accepted into either a three-year minimum or an extended, four-year minimum programme of study. This extended curriculum has been designed in order to enhance student development and to improve the student's chances of successful completion.

This qualification will enable the Students to acquire the necessary knowledge, skills, attitudes and values to practice as a Clinical Technologist in one of the following specialist categories: Cardiology, Cardiovascular Perfusion, Critical Care, Nephrology, Neurology, Pulmonology or Reproductive Biology. They will be able to perform procedures in one of the above seven specialist categories in order to contribute in the diagnosis and treatment of various patho-physiological conditions in conjunction with other designated health care professionals. They also perform organ system support, diagnostic, therapeutic and corrective procedures on patients using specialized health technology and techniques for the treatment of physiological dysfunction.

Students in Clinical Technology must attend formal lectures and practical sessions at the Durban University of Technology in all subjects for the duration of their studies.

II.I.I Duration of the programme

The minimum completing duration for the programme is four (4) years of full-time study and the maximum period of registration is six (6) years.

11.1.2 Assessment and Moderation

Some subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Other subjects do have final examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

11.1.3 Registration with the Professional Board

As a Student: On enrolment, it is mandatory that a student register as a student Clinical Technologist with the Health Professions Council of South Africa as determined in the regulations set out in the Government Gazette (No. R.1608 dated 24 July 1987).

As a Graduate: A graduate who has completed the qualification successfully and has complied with all the conditions as set out by the HPCSA may register as a qualified Clinical Technologist with the Health Professions Council of South Africa in terms of the current rules for registration.

11.1.4 Work-Integrated Learning Period (WIL)

The Work-Integrated Learning period will run concurrently with the specialist subjects, in the fourth year of study, at a training unit approved by the Health Professions Council of South Africa (HPCSA). During WIL students would be required to pass the Competency Based Test (CBT) with a minimum mark of 70%, as a Board requirement. The fourth year comprises the Work Integrated Learning [WIL] component, where a student will choose one of seven categories and study the major specialist subjects appropriate to the chosen category. The categories are as follows: Cardiology, Cardio-Vascular Perfusion, Critical Care, Nephrology, Pulmonology, Reproductive Biology and Neurophysiology.

The latter must be done at a training unit approved by the Health Professions Council of South Africa.

I I.2. Programme Learning Structure + Assessment column NATIONAL DIPLOMA: CLINICAL TECHNOLOGY: EXTENDED CURRICULUM PROGRAMME (NDCLF2)

Code	Subjects	Year of	NQF	Nated	Pre-req
		Study	Level	Credits	Code
FCMY101	Foundation Chemistry	I	5	0.100	
FPYC101	Foundation Physics	I	5	0.100	
ICLT101	Introduction to Clinical Technology	I	5	0.250	
CAPP101	Computer Applications I	I	5	0.135	
CHMB102	Chemistry I		5	0.08	FCMY101
PYSC105	Physics I		5	0.08	FPYC101
CSTA101	Calculation & Statistics	- 1	5	0.135	
ANAYI0I	Anatomy I	2	5	0.200	
FBAP101	Foundation Biomedical Apparatus Foundation Organs & Systems	2	5	0.2	
FO101	Pathophysiology	2	5	0.135	
PCLY101	Pharmacology I	2	5	0.035	
PSIO I 02	Physiology I	2	5	0.200	
PYDNI0I	Psychodynamics	2	5	0.135	
ANPH202	Anatomy & Physiology 2	3	6	0.200	PSIO102, ANAY101
BAPO201	Biomedical Apparatus & Procedures II	3	6	0.07	FBAP101
OSPP201	Organs & Systems Pathophysiology II	3	6	0.10	PSIO102, ANAY101 & FSOP101
PHAR201	Pharmacology II	3	5	0.100	PCLY101
CPAB301	*Cardiology: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
CACP310	*Cardiology: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects
CCTP310	*Cardiology: Clinical Tech Practice 3	4	6	0.300	All level1,2 & 3 subjects
CCBA301	*Critical Care: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
CCC301	*Critical Care: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects
CTPR301	*Critical Care: Clinical Tech. Prac. 3	4	6	0.300	All level1,2 & 3 subjects
NEAP301	*Nephrology: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
NCLI301	*Nephrology: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects
NCTP301	*Nephrology: Clinical Tech. Prac. 3	4	6	0.300	All level1,2 & 3 subjects
NBMA301	*Neurophysiology: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
NCLP301	*Neurophysiology: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects
NTPR301	*Neurophysiology: Clinical Tech. Prac. 3	4	6	0.300	All level1,2 & 3 subjects
FBAP301	*Perfusion: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
PCTP301	*Perfusion: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects
PCTP301	*Perfusion: Clinical Tech Prac 3	4	6	0.300	All level1,2 & 3 subjects
PBAP301	*Pulmonology: Biomedical Apparatus 3	4	6	0.350	All level1,2 & 3 subjects
PCLP301	*Pulmonology: Clinical Practice 3	4	6	0.350	All level1,2 & 3 subjects

PTPR301	*Pulmonology: Clinical Tech Prac 3	4	6	0.300	All level1,2	&	3
RBAP301			6	0.350	subjects All level1,2	&	3
RCPR301	*Reproduction: Clinical Practice 3	4	6	0.350	subjects All level1,2	&	3
	'				subjects All level1,2	&	3
RTPR301	*Reproduction: Clinical Tech Prac 3	4	6	0.300	subjects	~	,

11.3 PROGRAMME RULES

11.3.1 Minimum Admission Requirements

In addition to Rule G7 the minimum entrance requirement for entry into the programme of study is a National Senior Certificate (NSC) with endorsement for diploma entry with the following subjects:

Compulsory subjects	NSC Rating
English	3
Life Orientation	4
Mathematics	4
Life Science	4
Physical Science	4
And one 20-credit subject	3

The minimum requirement for holders of the Senior Certificate is a matriculation exemption with the following subjects at the stated ratings:

Compulsory Subjects	HG	SG
English	Е	D
Mathematics	D	С
Physical Sciences	D	С
Biology/Life Sciences	D	U

The DUT general rules G7 (3) and G7 (8) respectively, will apply for admission requirements based on work experience, age & maturity; and recognition of prior learning (RPL).

The DUT Admission's Policy for International Students and general rules G4 and G7 (5), apply for admission of international students.

11.3.2 Selection Criteria

In accordance with Rule G5, placement into the ECP programme is limited to 10 places. The following selection process will determine placement in the programme:

Successful applicants for study towards a ND: Clinical Technology will be

accepted into either a three-year minimum or an extended curriculum programme (four-year minimum) of study. An extended curriculum is devised in order to enhance student development and to improve the student's chances of successful completion. As more qualifying applications are received than can be accommodated, the following selection process will determine placement in the programme:

- O All applicants must apply through the Central Applications Office (CAO).
- Initial shortlisting for selection is based on the applicant's academic performance in Grade 12 (Grade 11, or Grade12 June marks, will be used for current matriculating students).
- Shortlisted students will be invited to undergo placement testing.
- Applicants who pass the placement tests may be invited for an interview.
- Provisional acceptance may be given to selected applicants awaiting National Senior Certificate (NSC) results. If the final Grade 12 NSC results do not meet the minimum entrance requirements, then provisional acceptance will be withdrawn.
- Final selection for placement will be based on results in the SC/ NSC and DUT placement tests, as well as on recommendations from the interview panel.
- Students will be ranked according to the following criteria:

Assessment	Weighting (%)
Results of the Senior Certificate or National Senior Certificate	30%
Placement Testing	35%
Interview Score	35%

11.3.3 Pass Requirements

- 1. Promotion to year 2: First year students registered in the extended curriculum program will only be eligible for subsequent registration provided that a student passes the following subjects:
 - All four Foundation subjects, i.e., Introduction to Clinical Technology, Foundation Biomedical Apparatus, Foundation Chemistry and Foundation Physics
 - Two out of the three mainstream subjects, i.e., Chemistry I, Physics I, Computer Applications I
- 2. Promotion to year 3 will only be allowed if the student passes the following subjects:
 - Anatomy I, Physiology I and Calculation and Statistics I
 - Foundation Organs and Systems Pathophysiology and Foundation Pharmacology
- 3. Promotion to year 4 will only be allowed if the student passes all 3rd year subjects

- 4. The minimum duration to complete the N Dip: Clinical Technology (Extended Curriculum Programme) is 4 years and the maximum duration is 5 years of consecutive study.
- 5. Students who do not comply with any of the rules outlined in points I to 4 above may need to apply for re-registration in the ECP Programme to the Department of Biomedical and Clinical Technology.

11.3.4 Re-registration Rules

Rule G16 in the General Handbook applies

11.3.5 Exclusion Rules

In addition to Rule G17, the following departmental rule applies:

A first year student who fails four or more subjects with a final mark of less than 40% will not be allowed to re-register for the programme: ND Clinical Technology (ECP). Deregistration from any subject is subject to the provisions of Rule G6 (2).

11.3.6 Interruption of Studies

In accordance with Rule G21A(b), the minimum duration for this programme will be four (4) years of registered study and the maximum duration will be five (5) years of registered study, including any periods of WIL. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

12. BACHELOR OF TECHNOLOGY: CLINICAL TECHNOLOGY (BTCLT 2) - NO NEW STUDENTS INTAKE

12.1 PROGRAMME INFORMATION

Completion of the qualification will enable the student to independently conduct advanced diagnostic, therapeutic, corrective procedures and organ system support on patients using specialised equipment and techniques for the treatment and/or interpretation of a diagnosis of abnormalities and disease. The individual is able to strategically manage clinical technology practice, maintain QA, perform research and train members of the health care team. The individual may be self-employed or employed by a recognised health care facility.

12.1.1 Registration with the Professional Board

A candidate who has completed the course successfully and has satisfied the requirements of the Professional Board for Clinical Technology may register

as a Graduate Clinical Technologist with the Health Professions Council of South African (HPCSA).

12.1.2 Assessment

Some subjects in this programme do not have a final examination viz: Research Methodology Clinical Technology Research Project , as well as the advanced specialist subject . The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. One subject (Principles of Management I) has a final examination. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

12.2 PROGRAMME LEARNING STRUCTURE BACHELOR OF TECHNOLOGY: CLINICAL TECHNOLOGY (BTCLT 2)

Code	Subjects	Year of Study	NQF Level	Nated Credits	Compulsory, elective or WIL
RMNC201	Research Methodology	4	7	0.250	Compulsory
PRMG101	Principles of Management	4	7	0.250	Compulsory
CLRP101	Clinical Technology Research Project	4	7	0.200	Compulsory
ACDT401	Advanced Cardiac Technology	4	7	0.300	Elective
ACRT401	Advanced Critical Care Technology	4	7	0.300	Elective
ARNT401	Advanced Renal Technology	4	7	0.300	Elective
ANPT401	Advanced Neurophysiologic Technology	4	7	0.300	Elective
APFT401	Advanced Perfusion Technology	4	7	0.300	Elective
ARST401	Advanced Respiratory Technology	4	7	0.300	Elective
ARPT401	Advanced Reproductive Technology	4	7	0.300	Elective

^{*}Elective subject

12.3 PROGRAMME RULES

12.3.1 Minimum Admission Requirements & Selection Criteria

In accordance with Rule G5, acceptance into the programme is limited to 30 places, and entry to the BTech programme is not automatic. As more qualifying applications are received than can be accommodated, the following selection criteria will determine entry into the programme, with the 30 highest ranking candidates gaining entry into the programme:

- Applicants must have completed the ND: Clinical Technology.
- Applicants are required to formally apply to the department, by the due date, to be considered for the B Tech: Clinical Technology programme.

- Applicants must submit proof of placement in a Clinical Technology training unit under the supervision of a Graduate Clinical Technologist
- Applicant's academic performance in the ND: Clinical Technology using the ranking criteria below:

THE RANKING CRITERIA

I.Average marks of the final year of the National Diploma 2.Years to complete ND: Clinical Technology							
Minimum duration	Minimum duration	Minimum duration	Minimum duration				
+ 3 years	+ 2 years	+ I year					
0	I	3	5				
3.Workplace experie	nce post National Diplor	na in an accredited train	ing unit				
0-1 year							
0 5 10 15							

 An applicant's ranking is determined by the total points score obtained by the addition of the scores obtained in the individual ranking criteria, as shown in the example in the table below:

Criteria	Ranking (points)	Score
Average final year mark in year 3 of the ND: Clinical Technology is 70%	70	
ND: Clinical Technology completed in minimum duration (3 years)	5	
Workplace experience (Diploma just completed)	0	
Total	75	•

(w.e.f. 28/08/2014)

12.3.2 Pass Requirements

In addition to Rule G14 and G15, the following rules apply. Students are encouraged to apply themselves to their studies, and strive for the best academic results possible in order to adequately prepare themselves for their future careers.

12.3.3 Re-registration Rules

Rule G16 in the General Handbook applies.

12.3.4 Exclusion Rules

Rule G17 in the General Handbook applies.

12.3.5 Interruption of Studies

In accordance with Rule G23A, the minimum duration for this programme will be one (1) year of registered study and the maximum duration will be

two (2) years of registered study. Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration.

13 BACHELOR OF HEALTH SCIENCES IN CLINICAL TECHNOLOGY

13.1 PROGRAMME INFORMATION

This qualification develops a learner to possess the necessary knowledge, skills, attitudes and values to practice as a Clinical Technologist, as a part of a multi-disciplinary team, in one of the following specialist categories: Cardiology, Cardiovascular Perfusion, Critical Care, Nephrology, Neurology, Pulmonology or Reproductive Biology. The qualifying learner will be able to independently perform diagnostic, therapeutic and corrective procedures on patients using specialised health technology and techniques for the treatment of pathophysiological conditions in a hospital-based or in a private practice setting.

This qualification will enable the learner to engage in research and contribute to the creation of new knowledge within the field. Lastly the qualification is designed to provide learners with specific clinical technology skills and competencies that are included in management and research.

The programme will be delivered full-time at DUT, with exposure to the clinical environment from first year to fourth year. The grounding for basic medical and clinical sciences will be provided in the first year, comprising of both theoretical and practical components. The theoretical component will be integrated with the practical component in the Skills Laboratory and through clinical rotational observations in the specialist categories of Clinical Technology. These clinical rotations will be undertaken at HPCSA and DUT accredited training units, and will take place on a fort-nightly basis.

The second level of study will equip the student with more complex knowledge by applying introductory concepts to understand the anatomical and physiological systems, as well as pathogenesis and progression of diseases and conditions, related to Clinical Technology.

In the 3rd level of study, the student is placed in the specific specialist category and rotates through various accredited training units up to the 4th level. Both these levels (i.e. III and IV) will employ an integrated teaching and learning approach where the student will be able to apply scientific and technological knowledge to perform diagnostic, therapeutic and life support procedures, and the evaluation thereof. The delivery of the 3rd and 4th level will be offered in both block lectures and block practical in a 50:50 ratio. The practical block will

be facilitated by DUT-appointed clinical instructors and specialist lecturers in an integrated teaching and learning approach (using e-learning, case studies, journaling, for example) to ensure that the learning outcomes are achieved, and that the quality of the delivery is maintained.

13.1.1 Duration of the programme (4 years)

In accordance with the DUT Rule G23B (2)* and Rule G23B (3)*, the minimum duration of study is four years, including any periods of clinical practice, and the maximum duration will be six years of registered study, including any periods of clinical practice.

13.1.2 Assessment and Moderation

Some subjects in this programme do not have a final examination. The results for these subjects are determined through a weighted combination of assessments. As such, there are no supplementary examinations. Other subjects do have final examinations. Students are encouraged to work steadily through the period of registration in order to achieve the highest results possible. Assessment details are listed under each subject at the back of this handbook. Moderation follows the DUT requirements.

13.1.3 Registration with the Professional Board

As a Student: On enrolment, it is mandatory that a student register as a student Clinical Technologist with the Health Professions Council of South Africa as determined in the regulations set out in the Government Gazette (No. R.1608 dated 24 July 1987).

As a Graduate: A graduate who has completed the qualification successfully and has complied with all the conditions as set out may register as a qualified Clinical Technologist with the Health Professions Council of South Africa in terms of the current rules for registration.

13.1.4 Work-Integrated Learning Period (WIL)

WIL will run concurrently with the specialist subjects, in the third year of study, at a training unit approved by the Health Professions Council of South Africa (HPCSA). During WIL students would be required to pass the Competency Based Test (CBT) with 70%, as a Board requirement.

13.2. PROGRAMME LEARNING STRUCTURE

Insert programme name

Module	Module Title	HEQSF	HEQSF	Period of	Block	Pre-	HEMIS
code		level	Credit	Study	Code	requisite module/s	credits
ICLT101	Introduction to Clinical Technology	5	8	1	21	N	0.0645
CMTR 101	Chemistry	5	16	I	21	N	0.129
PHISTIT	Physics 101	5	8	1	22	N	0.065
PHIS121	Physics 201	5	8	I	22	N	0.065
AAMYI0I	Anatomy	5	16	1	21	N	0.129
PYSLI01	Physiology	5	16	T	21	N	0.129
PTPY101	Pathophysiology I	5	8	I	22	N	0.0645
ITCT I 0 I	Instrumentation and Techniques for Clinical Technology I	5	12	I	22	N	0.0968
CSTN101	Cornerstone module	5	12	I	22	N	0.0968
ITCH101	Introduction to Technopreneurship	5	8	I	22	N	0.0645
VNVLI01	Violence and non- violence*	5	8	1	22	N	0.0645
IGSH101	Issues of Gender and Society	5	12	I	21	N	0.0968
PPDVI0I	Personal and Professional Development I	5	12	I	21	N	0.0968
AAPA101	Applied Anatomy and Physiology I a	6	12	2	21	Anatomy Physiology	0.094
AAPBI0I	Applied Anatomy and Physiology I b	6	12	2	22	Anatomy Physiology	0.094
CLTP101	Clinical Technology Practice	6	12	2	22	Introduction to Clinical Technology	0.094
ITCT201	Instrumentation and Techniques for Clinical Technology II	6	16	2	21	Instrumentati on and Techniques for Clinical Technology I	0.125
PTPY201	Pathophysiology II	6	16	2	22	Pathophysiol ogy I; Physiology	0.125
PRCL101	Pharmacology	6	16	2	21	Anatomy & Physiology	0.125
	Research Methodology I	6	16	2	22	N	0.125
HCDKI0	HIV and communicable diseases in KZN	6	8	2	21	N	0.062
EQDVI0I	Equality and Diversity	6	8	2	21	N	0.062
PPRM 101	Professional Practice & Management	6	12	2	22	N	0.094

			1	1	1		l
RMTD201	Research Methodology II	7	16	3	21	Research Methodology I	0.129
HLCM101	Health care management I	7	8	3	22	N	0.0645
RSJS101	Restorative Justice	7	8	3	21	N	0.0645
EMDLI01 ETMH 101	Ethics & Medical Law	7	12	3	22	N	0.096
PPDVI03	Personal and Professional Development III ELECTIVES	7	12	3	22	N	0.096
	Specialisation in						
PTCD101	Cardiology Pathophysiology for Cardiology	7	16	3	21	Pathophysiol ogy II	0.129
PMCD101	Pharmacology for Cardiology	7	8	3	22	All Level 2 subjects	0.0645
CTCA101	Clinical Technology Practice in Cardiology Ia	7	12	3	21	All Level 2 subjects	0.096
CTCB101	Clinical Technology Practice in Cardiology Ib	7	16	3	22	All Level 2 subjects	0.129
ITCA101	Instrumentation and Techniques for Clinical Technology in Cardiology la	7	12	3	21	All Level 2 subjects	0.096
ITCB101	Instrumentation and Techniques for Clinical Technology in Cardiology lb	7	16	3	22	All Level 2 subjects	0.129
	Specialisation in Critical care						
PPCC101	Pathophysiology for Critical Care	7	16	3	21	All Level 2 subjects	0.129
PHCC101	Pharmacology for Critical Care	7	8	3	22	All Level 2 subjects	0.0645
CCCA101	Clinical Technology Practice in Critical Care la	7	12	3	21	All Level 2 subjects	0.096
CCCB101	Clinical Technology Practice in Critical Care Ib	7	16	3	22	All Level 2 subjects	0.129
ICRA101	Instrumentation and Techniques for Clinical Technology in Critical Care la	7	12	3	21	All Level 2 subjects	0.096
ICRB101	Instrumentation and Techniques for Clinical Technology in Critical Care Ib	7	16	3	22	All Level 2 subjects	0.129
	Specialisation in Neurophysiology						
PTNP101	Pathophysiology for Neurophysiology	7	16	3	21	All Level 2 subjects	0.129

	T =-						
PHNP101	Pharmacology for Neurophysiology	7	8	3	22	All Level 2 subjects	0.0645
CTNA101	Clinical Technology Practice in Neurophysiology la	7	12	3	21	All Level 2 subjects	0.096
CTNB101	Clinical Technology Practice in Neurophysiology Ib	7	16	3	22	All Level 2 subjects	0.129
ITNA101	Instrumentation and Techniques for Clinical Technology in Neurophysiology la	7	12	3	21	All Level 2 subjects	0.096
ITNB101	Instrumentation and Techniques for Clinical Technology in Neurophysiology Ib	7	16	3	22	All Level 2 subjects	0.129
	Specialisation in Nephrology						
PTNR101	Pathophysiology for Nephrology	7	16	3	21	All Level 2 subjects	0.129
PHNR101	Pharmacology for Nephrology	7	8	3	22	All Level 2 subjects	0.0645
CTPA101	Clinical Technology Practice in Nephrology Ia	7	12	3	21	All Level 2 subjects	0.096
CTPB101	Clinical Technology Practice in Nephrology Ib	7	16	3	22	All Level 2 subjects	0.129
ITPA101	Instrumentation and Techniques for Clinical Technology in Nephrology Ia	7	12	3	21	All Level 2 subjects	0.096
ITPB101	Instrumentation and Techniques for Clinical Technology in Nephrology Ib	7	16	3	22	All Level 2 subjects	0.129
	Specialisation in Perfusion						
PTPF101	Pathophysiology for Perfusion	7	16	3	21	All Level 2 subjects	0.129
PHPF101	Pharmacology for Perfusion	7	8	3	22	All Level 2 subjects	0.0645
CPPA101	Clinical Technology Practice in Perfusion Ia	7	12	3	21	All Level 2 subjects	0.096
CPPB101	Clinical Technology Practice in Perfusion Ib	7	16	3	22	All Level 2 subjects	0.129
ITFA101	Instrumentation and Techniques for Clinical Technology in Perfusion Ia	7	12	3	21	All Level 2 subjects	0.096
ITFB101	Instrumentation and Techniques for Clinical Technology in Perfusion Ib	7	16	3	22	All Level 2 subjects	0.129
	Specialisation in Pulmonology						
PTPL101	Pathophysiology for Pulmonology	7	16	3	21	All Level 2 subjects	0.129
PHPL101	Pharmacology for Pulmonology	7	8	3	22	All Level 2 subjects	0.0645
CTLAI0I	Clinical Technology Practice in Pulmonology la	7	12	3	21	All Level 2 subjects	0.096

CTLB101	Clinical Technology	7	16	3	22	All Level 2	0.129
	Practice in Pulmonology					subjects	
	lb						
ITLA101	Instrumentation and	7	12	3	21	All Level 2	0.096
	Techniques for Clinical					subjects	
	Technology in						
	Pulmonology Ia						
ITLB101	Instrumentation and	7	16	3	22	All Level 2	0.129
	Techniques for Clinical					subjects	
	Technology in					'	
	Pulmonology Ib						
	Specialisation in						
	Reproductive biology						
PTRB101	Pathophysiology for	7	16	3	21	All Level 2	0.129
	Reproductive Biology	-				subjects	
PHRB101	Pharmacology for	7	8	3	22	All Level 2	0.0645
111110101	Reproductive Biology	,				subjects	0.0015
CTRA101	Clinical Technology	7	12	3	21	All Level 2	0.096
CIRAIUI		/	12	3	Z1		0.076
1	Practice in Reproductive				1	subjects	
CTRRIA	Biology la	 -		3		A II	0.129
CTRBI01	Clinical Technology	7	16	3	22	All Level 2	0.129
	Practice in Reproductive					subjects	
	Biology Ib	<u> </u>				1	
ITBA 101	Instrumentation and	7	12	3	21	All Level 2	0.096
1	Techniques for Clinical				1	subjects	
	Technology in						
L	Reproductive Biology la	1			L_		<u> </u>
ITBB101	Instrumentation and	7	16	3	22	All Level 2	0.129
	Techniques for Clinical					subjects	
	Technology in						
	Reproductive Biology Ib						
HCMPI01	Healthcare Management	8	12	4	22	All Level 3	0.091
	Practice					subjects	
PPDV 104	Personal and Professional	8	12	4	22	Community	0.091
	Development IV					Healthcare	
						and Research	
1					1	III	
RP A101	Research Project a	8	12	4	21	All Level 3	0.091
	<i>'</i>					subjects	
RPJB101	Research Project b	8	16	4	22	All Level 3	0.12
,5.0.	. toodar cir i roject b	`	'	'		subjects	3.12
HLCM201	Health care management	8	16	4	21	All Level 3	0.12
I III III III III III III III III III	Health care management	0	10	7	41	subjects	0.12
CUNUO	"	<u> </u>					0.12
CLIN101	Clinical Instruction	8	16	4	21	All Level 3	0.12
						subjects	
SBSM101	Small Business	8	16	4	21	All Level 3	0.12
L	Management	<u> </u>				subjects	<u> </u>
	Specialisation in						
	Cardiology						
CTCA201	Clinical Technology	8	16	4	21	All Level 3	0.12
	Practice in Cardiology Ila					subjects	
CTCB201	Clinical Technology	8	16	4	22	All Level 3	0.12
	Practice in Cardiology IIb					subjects	
ITCA201	Instrumentation and	8	12	4	21	All Level 3	0.091
11.5,1201	Techniques for Clinical	ľ	1 '2		~'	subjects	0.071
I	Technology in				1	Subjects	1
	I I CCITIOIOEY III	1		1	ı	1	1
	Cardiology IIa						

ITCB201	Instrumentation and	8	16	4	22	All Level 3	0.12
	Techniques for Clinical Technology in					subjects	
	Cardiology IIb Specialisation in						
	Critical care						
CCCA201	Clinical Technology	8	16	4	21	All Level 3	0.12
	Practice in Critical Care					subjects	
CCCB201	Clinical Technology	8	16	4	22	All Level 3	0.12
	Practice in Critical Care lib					subjects	
ICRA201	Instrumentation and	8	12	4	21	All Level 3	0.091
	Techniques for Clinical Technology in Critical					subjects	
	Care Ila						
ICRB201	Instrumentation and	8	16	4	22	All Level 3	0.12
	Techniques for Clinical					subjects	
	Technology in Critical Care IIb						
	Specialisation in						
	Neurophysiology						
CTNA201	Clinical Technology Practice in	8	16	4	21	All Level 3	0.12
	Neurophysiology IIa					subjects	
CTNB201	Clinical Technology	8	16	4	22	All Level 3	0.12
	Practice in					subjects	
ITNA201	Neurophysiology IIb Instrumentation and	8	12	4	21	All Level 3	0.091
IIINAZUI	Techniques for Clinical	0	12	7	21	subjects	0.071
	Technology in					, ,	
ITN IDOOL	Neurophysiology IIa						0.10
ITNB201	Instrumentation and Techniques for Clinical	8	16	4	22	All Level 3 subjects	0.12
	Technology in					Subjects	
	Neurophysiology IIb						
	Specialisation in Nephrology						
CTPA201	Clinical Technology	8	16	4	21	All Level 3	0.12
01171201	Practice in Nephrology IIa		1.0			subjects	0.1.2
CTPB201	Clinical Technology	8	16	4	22	All Level 3	0.12
	Practice in Nephrology					subjects	
ITPA201	Instrumentation and	8	12	4	21	All Level 3	0.091
	Techniques for Clinical					subjects	
	Technology in						
ITPB201	Nephrology IIa Instrumentation and	8	16	4	22	All Level 3	0.12
5201	Techniques for Clinical	•				subjects	
	Technology in						
	Nephrology IIb Specialisation in						
	Perfusion in						
CPPA201	Clinical Technology	8	16	4	21	All Level 3	0.12
CDDCCC	Practice in Perfusion IIa		<u> </u>			subjects	0.10
CPPB201	Clinical Technology Practice in Perfusion IIb	8	16	4	22	All Level 3 subjects	0.12
ITFA201	Instrumentation and	8	12	4	21	All Level 3	0.091
	Techniques for Clinical					subjects	
	Technology in Perfusion IIa						
	i criusion na	l					l

ITFB201	Instrumentation and Techniques for Clinical Technology in Perfusion IIb	8	16	4	22	All Level 3 subjects	0.12
	Specialisation in Pulmonology						
CTLA201	Clinical Technology Practice in Pulmonology Iia	8	16	4	21	All Level 3 subjects	0.12
CTLB201	Clinical Technology Practice in Pulmonology Iib	8	16	4	22	All Level 3 subjects	0.12
ITLA201	Instrumentation and Techniques for Clinical Technology in Pulmonology Ila	8	12	4	21	All Level 3 subjects	0.091
ITLB201	Instrumentation and Techniques for Clinical Technology in Pulmonology Ilb	8	16	4	22	All Level 3 subjects	0.12
	Specialisation in Reproductive Biology						
CTRA201	Clinical Technology Practice in Reproductive Biology IIa	8	16	4	21	All Level 3 subjects	0.12
CTRB201	Clinical Technology Practice in Reproductive Biology IIb	8	16	4	22	All Level 3 subjects	0.12
ITBA201	Instrumentation and Techniques for Clinical Technology in Reproductive Biology IIa	8	12	4	21	All Level 3 subjects	0.091
ITBB201	Instrumentation and Techniques for Clinical Technology in Reproductive Biology IIb	8	16	4	22	All Level 3 subjects	0.12

13.3 PROGRAMME RULES

(Approved by SENATE August 2014)

13.3.1. Minimum admission requirements

In addition to Rule G7*, the minimum entrance requirements for the holder of a valid National Senior Certificate (NSC) or a Senior Certificate or National certificate (Vocational) for entry into a Bachelor's Degree and must include the following subjects at the stated minimum ratings in Table I

Table I: Minimum Admission Requirements

NSC REQUIREMENTS	SENIOR CERTIFICATE		NC (V)	
Compulsory subjects NSC Rating		SC Symbol HG SG		
English (Home language) OR English (Ist additional language)	4	D B		70%
Mathematics	4	D	В	70%
Life Sciences	4	D B		70%
Physical Sciences	4	D	В	70%

And two other 20 credit subjects of	3	Four other subjects, only	70%
which only one may be a language		one of which may be a	
		language	

13.3.2 Minimum Admission Requirements in respect of Work Experience, Age, Maturity, RPL and International Students:

The DUT General Rules G7 (3)* and G7 (8)* respectively will apply. The DUT's Admission Policy for International Students and General Rules G4* and G7 (5)* will apply.

I3.3.3 Selection procedures

All applicants must apply to the Central Applications Office (CAO).

In accordance with Rule G5*, acceptance into the programme is limited. Since more applications are received than can be accommodated, the following selection processes will apply:

- Initial short listing for selection is based on the applicant's academic performance in Grade 11 and/or 12.
- Applicants obtaining more than 25 points increase their chance of selection into the programme.
- The point scores for the **NSC** or the **SC** or the **NC(V)** results is obtained by using the table 2.

Table 2: Point Scores

	NSC	SC		NC(V)
RESULTS		HG	SG	
90 – 99%	8	8	6	4
80 – 89%	7	7	5	4
70 – 79%	6	6	4	4
60 – 69%	5	5	3	3
50 – 59%	4	4	2	
40 – 49%	3	3	I	
30 – 39%	2	2		
0 – 29%	I	I		

NOTE: No points are allocated for ten (10) credit subjects.

- Applicants who meet the minimum departmental admission requirements for the Bachelor of Health Sciences in Clinical Technology will be ranked according to the points scored in Grade 12, and may be invited to participate in the selection process.
- The selection is based on the criteria and weightings in the Table 3:

Table 3: Weighting of assessments

Assessment	Weighting (%)
Results of the Senior certificate/National Senior Certificate	60
Interview scores	40

- Applicants invited to the selection process should have a sound knowledge of the Clinical Technology profession.
- Successful applicants will be placed into either the four-year degree or the fiveyear Extended Curriculum Programme.
- Provisional acceptance is given to selected applicants awaiting National Senior Certificate (NSC) and National Certificate (Vocational) results. If the final Grade 12 NSC/ NC (V) results do not meet the minimum entrance requirements, the provisional acceptance will be automatically withdrawn.
- Applicants whose application has been declined due to poor academic achievement in grade 11 may reapply to the programme should they be able to show improved academic performance in the final grade 12 examinations. Those applicants who wish to reapply should immediately notify the programme of their intention to reapply. In order for the application to be reconsidered, the applicant must submit the final grade 12 results to the Department as soon as these results are available.

13.3.4 Progression rules

In addition to Rules G16*, students must pass all prerequisite modules as per Table I before progressing to a higher level.

13.3.5 Exclusion rule

In addition to the DUT General Rules G17*, a first year student who fails three or more modules with an average of less than 40% in the failed modules during that year, is not permitted to re-register for the Programme. Deregistration from any module is subject to the provisions of rule G6 (2)*.

13.3.6 **Re-registration**

Rule G17* of the General Handbook for Students applies.

13.3.7 Interruption of studies

Should a student interrupt their studies for a period or more than three consecutive years, the student will need to apply to the department for permission to re-register and will need to prove currency of appropriate knowledge prior to being granted permission to continue with registration.

13.3.8 Clinical Technology Practice (CTP)

In addition to Rule G28*, the following should be noted:

- 1. Students must achieve clinical competencies in a Health Professions Council of South Africa (HPCSA)-accredited unit.
- 2. The department is responsible for placement of students in level I-IV. Transportation arrangements to the clinical training sites is the responsibility of individual students from level II, II and IV.
- 3. Students will not be allowed to change specialist categories in the third and the fourth registered level.

4. Disciplinary matters occurring in the unit will, in the first instance, be subject to the disciplinary code of conduct of that specific unit, and then be referred to DUT for student disciplinary action.

13.3.9 Registration with the Health Professions Council of South Africa (professional board of radiography and clinical technology)

Students are required to register as a student Clinical Technologist with the Health Professions Council of South Africa (Board of Radiography and Clinical Technology) in their first year of study. Registration fees and submission of registration documents will be for the responsibility of the student.

14. MASTERS OF HEALTH SCIENCES IN CLINICAL TECHNOLOGY (MHCLTI)

14.1 PROGRAMME INFORMATION

This full research qualification is aligned to Rule G24 and the guidelines in the Post Graduate Student Handbook.

- The Student who successfully completes this qualification will be able to apply advanced problem solving skills and critical, reflective thinking to perform independent research in a chosen field and report their findings in a dissertation that meets the accepted criteria and ethical principles for the profession. In this way they will make a contribution to the existing body of knowledge and initiate change that will help develop and advance the profession of medical technology.
- The qualifying Student will be able to conduct independent research under minimal guidance in a chosen field, and contribute to knowledge production in that field. The research problem, its justification, process and outcome is reported in a dissertation which complies with the generally accepted norms for research at that level.

14.1.1 Assessment and Moderation

In addition to Rule G24 (4), postgraduate assessment of dissertations will be aligned to Postgraduate policies and guidelines. Please refer to the General Student Handbook and the Postgraduate Student Handbook.

14.2 PROGRAMME LEARNING STRUCTURE

Code	Module	Year of Study	Assessment Type	NATED Credits	Pre-requisites	Co-requisites
MHCLTI	Dissertation	2	External Examination	1.0	None	none

14.3. PROGRAMME RULES (Approved by SENATE August 2014)

14.3.1 Minimum Admission Requirements

In addition to the General Handbook for Students Rule G24 (I), candidates must be possession of a Bachelor's Degree in Clinical Technology (NQF Level 8), or must have been granted conferment of status according to Rule G10A. Candidates may also apply for admittance via Recognition of Learning (RPL) in accordance with Rule G7 (8) and / or G10B.

14.3.2 Selection Criteria

In accordance with Rule G5, acceptance into the Masters of Health Sciences programme is limited, and not automatic. Students are selected into the programme once they have completed an intention to study and the department has discussed the viability of the proposed topic for the Masters Qualification. The intention to study/ concept page must include the following: Problem statement or Title of the intended study, Objectives / sub-problems / Research Questions, Rationale/motivation to do the study, Brief literature review, Brief methodology.

Applicants must have an aggregate of 60% overall for the B Tech Degree.

14.3.3 Pass Requirements

Rule G24 and the Postgraduate Student Handbook apply.

Students are encouraged to apply themselves to their research, and strive for the best academic results possible in order to adequately prepare themselves for their future careers.

14.3.4 Re-registration Rules

Rule G24 (2), Rule G26 (5) and the Postgraduate Student Handbook apply.

14.3.5 Exclusion Rules

Rule G24 (1)(d); Rule G24 (2), and the Postgraduate Student Handbook apply.

14.3.6 Minimum and maximum duration

The minimum duration for this programme shall be one (1) year of registered study and the maximum duration shall be three (3) years of registered study.

14.37. Interruption of Studies

Should there be bona fide reasons for the interruption of studies for a period of one (I) year or more once the candidate is formally registered, the student may apply for an interruption of registration. Registration may be interrupted under exceptional circumstances only and is not done retrospectively.

15. DOCTOR OF MEDICAL CLINICAL SCIENCES (DRMCS1)

15.1 PROGRAMME INFORMATION

This full research qualification is aligned to Rule G25 and G26 and the guidelines in the Post Graduate Student Handbook. The purpose of this qualification is to ensure that the student who successfully completes this qualification will be able to apply advanced problem-solving skills and critical, reflective thinking to perform independent research in a chosen field and report their findings in a dissertation that meets the accepted criteria and ethical principles for the profession. In this way they will make a contribution to the existing body of knowledge and initiate change that will help develop and advance the profession of Clinical Technology.

15.1.1 Assessment and Moderation

Post graduate assessment will be aligned to Postgraduate policies and guidelines.

Rule G25 (4) and the Postgraduate Student Handbook apply.

15.2 LEARNING PROGRAMME STRUCTURE

Code	Module	Year o Study	f Assessment Type	NATED Credits	Pre- requisites	Co- requisites
DRMCSI	Dissertation	2	External Examination	2.0	None	none

15.3 PROGRAMME RULES

15.3.1 Minimum Admission Requirements

In addition to the General Handbook for Students Rule G24 (I), candidates must be possession of a Master's Degree in Clinical Technology (NQF Level 9), or must have been granted conferment of status according to Rule G10A. Candidates may also apply for admittance via Recognition of Learning (RPL) in accordance with Rule G7 (8) and / or G10B. Students are selected into the programme once they have completed an intention to study and the department has discussed the viability of the proposed topic for the qualification. A sound knowledge of the fundamental principles and concepts of research and statistical methods is required.

15.3.2 Re-registration Rules

Please refer to Rule G26 (5) and the Postgraduate Student Handbook.

15.3.3 Exclusion Rules

Please refer to Rules G25 (2)(b; c(ii)) in the General Student Handbook; and the Postgraduate Student Handbook.

15.3.4 Minimum and maximum duration

In accordance with Rule G25 (2), the minimum duration for this programme will be two (2) years of registered study and the maximum duration will be four (4) years of registered study.

15.3.5. Interruption of Studies

Should a student interrupt their studies by more than three (3) years, the student will need to apply to the department for permission to reregister and will need to prove currency of appropriate knowledge prior to being given permission to continue with registration. Please refer to the Postgraduate Student Handbook.

15 SUBJECT CONTENT AND ASSESSMENTS

NB:

- The information below might change from time to time to suite national, institutional, faculty and departmental needs as may be approved by the Department of Higher Education, the HPCSA and the DUT relevant committees.
- Students are to read this section in conjunction with the relevant study guide.

16.1 BIOMEDICAL TECHNOLOGY

16.1.1 National Diploma: Biomedical Technology

Module Name	Learning Content	Assessment
	Atomic structure, Periodic table, molecular elements	The CONTINUOUS ASSESSMENT mark shall be made up of
FOUNDATION CHEMISTRY	& compounds, Composition and	Theory tests: 50%
(FCMRI0I)	stoichiometry Amines and amides	Practical tests: 30% Practical reports: 5%
		Assignments: 15%
	B . M	The CONTINUOUS ASSESSMENT
	Basic Mathematics, vectors, Problem solving skills in	mark shall be made up of
(FPHY101)	Physics, Conceptual physics	Theory tests: 60% Practical tests: 40%
		The CONTINUOUS ASSESSMENT
		mark
FOUNDATION	And the state of t	shall be made up of
IMMUNOLOGY	Antibody structure, Complement, HLA,	Theory tests: 50%
(FIMM101)	Structures in general	Practical tests: 30%
		Practical reports: 5%
		Assignment /s: 15%
		The CONTINUOUS ASSESSMENT
		mark
FOUNDATION	Amino acids, Physiological buffers, Structures	shall be made up of
BIOCHEMISTRY	in general, denaturation of proteins/DNA	Theory tests: 50%
(FBIO101)	Ionisation of amino acids	Practical tests: 40%
		Practical reports: 5%
		Assignment /s: 5%
LABORATORY	Solutions, Laboratory Mathematics,	The CONTINUOUS ASSESSMENT

TECHNIQUES	Laboratory ware, Safety, Microscopy	mark	1
(FLBT101)	Laboratory ware, Salety, Microscopy	shall be made up of	
(FEBITOI)			50%
		Theory tests:	
		Practical tests:	40%
		Practical reports:	5%
		Assignment /s:	5%
	Communication strategies, Personal	The CONTINUOUS A	SSESSMENT
	management skills, accessing and processing	mark	
ACADEMIC	information	shall be made up of	
LITERACY*	Language practices and conventions	(a)Tests	
	*This is not a subject on its own but will be	(b) oral presentation	
	incorporated in all the foundation subjects as a tool	(c) individual class exer	cises
	to help the Students.	TI CONTINUINT A	CCECCNENT
INTRODUCTON TO	Medical Technology the profession and the	The CONTINUOUS A	
MEDICAL	professional, Legal and Ethical aspects,	mark shall be ma	ae up or 25%
_	Laboratory safety	Theory Tests:	25%
TECHNOLOGY	Laboratory glassware and plastics, Laboratory	Practical Tests:	
(IMETIOI)	techniques and apparatus, Laboratory	Communication skills:	
	organization	Computer skills:	25%
	General arrangement of the body, The cell	Theomy Tester	2.49/
ANATOMY 6	and tissues, haematology, cardiovascular	Theory Tests:	24%
ANATOMY &	system	Practical Tests:	12%
PHYSIOLOGY I	Lymphatic system, Respiratory system,	Practical reports:	2%
(ANPH104)	Nervous system, Endocrine system,	Project:	2% 60%
	Reproductive system	Examination	60%
	Renal system, Gastrointestinal system		
CALCULATION S	Mathematical calculations: Algebra, Graphs,		
CALCULATION &	Trigonometry	Theory tests:	40%
STATISTICS	Statistical calculations: Descriptive Statistics,	Examination:	60%
(CSTAI0I)	Elementary probability, Probability distributions,		
	Correlation Analysis	T. T	2.40/
	Mechanics, thermal physics, wave	Theory Tests:	24%
PHYSICS I	motion, electricity and magnetism, light	Practical Tests:	12%
(PYSCI05)	and optics,	Practical reports:	2%
· ·	Introduction to atomic and nuclear	Project:	2%
	Physics	Examination	24%
	Matter and Energy, Chemical Equations and	Theory Tests:	12%
CHEMISTRY I	Stoichiometry, solution Chemistry, Rates of	Practical Tests:	2%
(CHMB102)	Reactions and Chemical Equilibrium, Organic	Practical reports:	2%
· ·	Chemistry	Project:	60%
	The normal and the adapted call Call injury and and	Examination	00/6
	The normal and the adapted cell, Cell injury and cell		
	death, Inflammation and repair, Neoplasia, Clinical	Theory Tosts:	32%
PATHOPHYSIOLOGY II	aspects of neoplasia, Genetic disorders, Respiratory	Theory Tests: Project:	8%
(PAPH201)	system disorders, Circulatory system disorders,	Project: Examination	60%
-	Urinary system disorders, Digestive system disorders, Nervous system and sensory organs disorders,	EXAMINIATION	00%
	Endocrine system disorders		
	Bio-elements and biomolecules,	Theory Tests:	24%
	Carbohydrates, Nucleic acids, Proteins	Practical Tests:	12%
BIOCHEMISTRY II	and amino acids	Practical reports:	2%
(BIOA202)	Lipids, Enzymes, ph and buffers, Introduction to	Project:	2%
	metabolism, Metabolism of carbohydrates	Examination	60%
	Introduction to Cytology, Specimen collection &	Examination	50/6
	fixation, Specimen preparation, Staining & mounting		
	Special techniques in Cytology, Biological behaviour	Theory Tests:	24%
IMMUNOLOGY II	of cells and tissues, Evaluation of the cellular sample,	Practical Tests:	12%
(IMMU202)	Histology & cytology of the FGT, Hormonal	Practical reports:	2%
(11 11 10 20 2)	Cytology, Agents of infection, Inflammatory,	Project:	2%
	degenerative and regenerative changes, Premalignant	Examination	60%
	changes, Malignant changes, Rare tumours		
BLOOD TRANSFUSION	Government regulations, General aspects of blood	Theory Tests:	24%
TECHNOLOGY	transfusion, The blood group systems	Practical Tests:	12%
I I ECI II TOLOGI	in ansitusion, The blood group systems	וומכנוכמו ולטנט.	1 4/0

(BLTT201)	Transmission of disease, Pretransfusion testing,	Practical reports:	2%	
(BL11201)	Untoward transfusion reactions, quality Assurance	Project:	2%	
	Ontoward transitision reactions, quality Assurance	•	60%	
	Internal control of the control of t	Examination	60%	
CELLUL AD	Introduction to Histology, Fixation, Tissue	Theory Tests:	24%	
CELLULAR	processing, Dehydration &dealcoholization,	Practical Tests:	13%	
PATHOLOGY I	Impregnation & embedding, Decalcification,	Assignment:	3%	
(CEPAI0I)	Microtomy, Staining, artefacts & pigments, Immunohistochemistry	Examination	60%	
	Illillidionistochemisti y	Theory Tests:		24%
CHEMICAL	Basic principles, Water balance, osmolality,	Practical Tests:		11%
PATHOLOGY I	electrolytes, pH and blood gases, Kidney and tests of	Practical reports:	3%	11/0
(CPATIOI)	renal function, Amino acids and proteins	Project:		2%
(CPATIOI)	renar function, Armino acids and proteins	Examination		60%
	History and development, Survey of Microorganisms			
	and classification, Microscopy and staining, Bacterial	Theory Tests:		24%
MICROBIOLOGY I	structure, reproduction and growth, Bacterial	Practical Tests:		12%
(MCGYI0I)	cultivation, Microbial metabolism, Bacterial genetics,	Practical reports:	2%	
(FICGTIOI)	Host parasite relationships, Control of	Project:		2%
	microorganisms	Examination		60%
		Theory Tests:		24%
CHEMICAL	Enzymes, Liver and tests of hepatic function,	Practical Tests:		11%
PATHOLOGY II	Disorders of carbohydrate metabolism, Lipid	Practical reports:	3%	, 0
(CPAT202)	metabolism	Project:	-,-	2%
(6: 2:202)	Pharmacology,	Examination	60%	270
	Origin and normal development of haematopoietic	Theory Tests:		24%
	elements, the erythrocyte, The leucocytes in the	Practical Tests:		12%
HAEMATOLOGY II	circulation	Practical reports:		2%
(HAEM203)	The platelet/megakaryocytic system, Haemostasis,	Project:		2%
	Basic haematological values	Examination		60%
		Theory Tests:		24%
		Practical Tests:		12%
MICROBIOLOGY II	Parasitology, mycology, virology, introduction to	Practical reports:		2%
(MCGY203)	bacteriology	Project:		2%
		Examination		60%
		Theory Tests:	24%	
CELLULAR	Respiratory tract, Serious effusions, Urinary	Practical Tests:	12%	
PATHOLOGY III	tract, Gastrointestinal tract, Central nervous	Practical reports/Assignment: 2%		
(CEPA301)	system	Project: 2%		
		Examination	60%	
		Theory Tests:	24%	
CHEMICAL	Mineral metabolism, CSF and other body fluids,	Practical Tests:	11%	
PATHOLOGY III	Immunochemical techniques, Endocrinology	Practical reports:	3%	
(CPAT303)	Pharmacology	Project: 2%		
		Examination	60%	
	Red cell morphology; The anaemias; The leucocytes,	Theory Tests:	24%	
HAEMATOLOGY III	The myeloproliferative;syndromes; The acute	Practical Tests:	12%	
(HAEM303)	leukaemias, The myelodysplastic syndromes, The	Practical reports:	2%	
(lymphoproliferative disorders, Platelets, Haemostasis,	Project:	2%	
	Parasites, Quality Assurance	Examination	60%	
	Specimen collection, transport and processing, gram	Theory Tests:	24%	
MICROBIOLOGY III	positive bacteria, gram negative bacteria,	Practical Tests:	12%	
(MCGY301)	mycobacteria, Atypical bacteria, spirochaetes,	Practical reports:	2%	
,	serology, antimicrobial agents, nosocomial infection	Project:	2%	
LABORATORY		Examination	60%	
LABORATORY	Performing, interpretation and integration of			
PRACTICE 3 (WORK	laboratory tests in the following disciplines	Workplace assessment	60%	
INTEGRATED	Medical Microbiology, Virology, Chemical Pathology,	Integrated learning pro		
LEARNING)	Cytology, Histology, Haematology and Blood			
(LABP 301)	Transfusion.			

16.1.2 BTECH: BIOMEDICAL TECHNOLOGY

Module Name	Learning Content	ASSESSMENT
RESEARCH METHOD & TECHNIQUES (RMTQ201)	Biostatistics, Research methods and applications	The CONTINUOUS ASSESSMENT mark shall be made up of Assessment weightings: Article critique: 20% Proposal: 50% Poster: 10% Statistics assignment: 20%
RESEARCH PROJECT (RPBMI01)	Preparation and submission of a research dissertation	Oral presentation 10% Chapter I draft 5% Chapter2 draft 5% Thesis 80%
INTEGRATED PATHOPHYSIOLOGY IV (IPAT401)	Clinical diagnosis and laboratory diagnosis of disorders in Integument, Skeletal, Muscular, nervous, Endocrine, Cardiovascular, lymphatic, Respiratory, Digestive, Urinary, Reproductive	Theory tests: 32% Assignment: 8% Examination 60%
LABORATORY MANAGEMENT (LABM201)	Principles of Management, Laboratory organization, Hunam resourses management, Physical resources management, Financial Management, Quality Assurance and Safety, Entrepreneurship	Theory tests: 24% Project: 16% Examination 60%
MOLECULAR BIOLOGY IV (MOLE401)	DNA structure and gene expression, Bacterial genetics, Regulation of gene function in bacterial and eukaryotic cells, Cancer at genetic level, molecular biology applications	The CONTINUOUS ASSESSMENT mark shall be made up of Theory tests: 60% Practical tests: 40%

16.1.3 BACHELOR OF HEALTH SCIENCES IN MEDICAL LABORATORY SCIENCE

CHEMISTRY	Apply knowledge and principles of general and		
	organic chemistry.		
	Explain with examples the role of chemistry in	Theory tests (average of all):	24%
	everyday life.	Practical tests	10%
	Perform calculations required for solution chemistry.	Practical reports	2%
	Prepare solutions following accurate procedures.	Assignments/oral presentation	: 2%
	Demonstrate understanding of the periodic table of	Tutorials, class/homework	2%
	elements and apply knowledge to general principles	Examination:	60%
	of chemistry.		
	Draw up balanced chemical reaction equations.		
PHYSICS (MODULE I)	MECHANICS		
	Fundamental Units & Dimensional Analysis		
	Vectors and Scalars		
	One Dimension Kinematics		
	Newton's Laws of Motion		
	Work, Energy & Power		
	Impulse and Momentum		
	Rotational Dynamics		
	PROPERTIES OF MATTER	Theory tests (average of all):	26%
	Phases of Matter	Practical tests	14%
	Elasticity	Examination:	60%
	Density and Specific Gravity		
	Pressure in Fluids		
	Atmospheric Pressure and Gauge Pressure		
	Pascal's Principle		
	Buoyancy and Archimedes' Principle		
	Surface Tension		
	Capillary Action		
	Viscosity		

	Poiseuille's Law	
PHYSICS (MODULE 2)	THERMAL PHYSICS	
` ′	Temperature	
	Heat and Temperature Change	
	Thermal Expansion of Solids	
	Heat and Phase Change	
	Calorimetry	
	Heat Transfer Mechanisms	
	WAVES & SOUND	
	Oscillatory Motion	
	Wave Motion & Types of Waves	
	Frequency, Amplitude and Wavelength	
	Speed of Waves on Strings	
	Reflection of Waves	
	Sound Waves	
	Energy and Intensity of Sound Waves	
	Doppler Effect	
	GEOMETRICAL OPTICS	
	Reflection	
	Refraction & Snell's Law	
	Dispersion	
	Critical Angles & Total Internal Reflection	
	Images Formed by Plane Mirrors	
	Images Formed by Spherical Mirrors Images Formed by Refraction: Thin Lenses	Theory tests (average of all): 26%
	ELECTRICITY& MAGNETISM	Practical tests (average of all). 26%
	Electric Charge	Examination: 60%
	Insulators and Conductors	Examination: 00%
	Charging by Friction, Conduction and	
	Induction	
	Coulomb's Law	
	Electric Field & Electric Field Lines	
	Electric Current & Potential Difference	
	Resistance & Ohm's Law	
	Series & Parallel Circuits	
	Fundamentals of Magnetism	
	RADIOACTIVITY & RADIATION	
	Properties of Nuclei	
	Binding Energy	
	Decay Processes (Alpha, Beta & Gamma)	
	Decay Constant & Half-Life	
	Activity	
	Medical Applications of Radioactivity	
	Biological Effects of Ionizing Radiation	
	QUANTUM PHYSICS	
	Blackbody Radiation and Plank's Hypothesis	
	Photoelectric Effect	
	Photons & Electromagnetic Waves	
FUNDAMENTALS OF	Wave Properties of Particles C	
FUNDAMENTALS OF MEDICAL	Pipetting. Use of balances.	
LABORATORY SCIENCE	Units, measurements and calculations related	
LABORATORT SCIENCE	to solution preparation.	
	Operate specified equipment in accordance	CONTINUOUS ASEESSMENT
	with standard operating procedures, using	Theory test: 50%
	different equipment including	Practical Tests: 20%
	spectrophotometers, pH meters, weighing of	Practical Reports: 10%
	chemicals.	Assignment/project: 10%
	Laboratory equipment made of glass and	Lab maths +tuts: 10%
	plastic appropriately	
	Sterilization procedures applicable to	
	different medical laboratory equipment,	
	reagent and surfaces.	
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	Apply and uphold safety procedures and	
	correct disposal of waste in accordance with	
	safety regulations acknowledging occupational	
	health and safety principles.	
	Quality Assurance procedures and principles	
	of maintenance of equipment & test analysis.	
	Role and function of the medical laboratory	
	scientist.	
	Apply ethical, professional, and medico-legal	
	principles and rules in the laboratory as	
	applied when dealing with different laboratory	
	specimen testing	
	Stock control procedures in the laboratory.	
	Communicate within a group using verbal,	
	written and electronic means of	
	communication.	
	Fundamental knowledge of statistical	
	techniques	
	TOPICS	
	HPCSA	
	SMLTSA	
	OHS act	
	Hierarchy	
	Course structure	
	CPD	
	Bathopele principles	
	Specimen types	
	Transportation	
	Anticoagulants	
	Storage	
	Decontamination	
	Disinfection	
	Biological, physical and chemical hazards	
	Evacuation drills	
	General laboratory safety rules	
	Centrifuges and centrifugation	
	Balances and weighing	
	Spectrophotometer and photometry	
	pH meter and pH measurement	
	Laboratory glassware and plastic ware	
	Autoclaving	
	Microscopes	
	Water purification (distillation and	
	deionisation)	
	Refrigeration	
	Use of quality control (QC)	
	Terminology used in QC	
	Record books	
	Filing	
STATISTICS	Introduction to Statistics (The learners will be	
	exposed to the differences between descriptive and	
	inferential statistics and its use in the Applied	
	Sciences and the use of computers in statistics)	
	Collection of Data (The different types data and its	
	method of collection will be discussed)	Theory tests (average of all): 24%
	Presentation of Data (The presentation of data in	Practical tests 10%
	the form of frequency distributions, graphs and	Practical reports 2%
		Assignments/oral presentation: 2%
	charts will be discussed)	Tutorials, class/homework 2%
	Measures of Location and Variation (The learners	Examination: 60%
	will be taught the various calculation methods on	
	the data collected and presented)	
	Correlation and Regression Analysis (An	
	understanding of the relationships between variables	
	will be accomplished through these analyses and its	

	use in the Applied Sciences)	
	Basic Probability and its distributions (The learners	
	will be exposed to the basic probability concepts and	
	its various distributions that exist and its relevance	
	to Applied Sciences)	
ANATOMY AND	The human body. The cell: Fluids and electrolytes,	
PHYSIOLOGY IA	Histology	
	Describe the language relating to anatomy and	
	physiology.	
	[, 8).	
	Describe the organisation of the body, metabolism,	
	and the structure and function of the cell	
	and the structure and ranged on the con-	
	Identify, describe, label & draw tissue types	
	Explain homeostasis at cellular level	
	Explain the importance and role of electrolytes and fluids in cells and tissues.	
	Skeletal system. Joints. Skin. Thermoregulatory	
	system	2 X two hour theory test
	Describe the integumentary system is in terms of	A supplementary test will be made
	structure and function	available.
	Classify & describe the anatomy of the skeleton	Each theory test will carry a weighting of 50%
	Describe the anatomy and physiology of the voluntary muscles.	
	Explain the structure of the skin & its components.	
	Consider the role of the skeletal system muscle &	
	skin as it relates to issues that may occur in the	
	environmental health scenario .e.g. ergonomics	
	Cityli Olimental Health Sechario .e.g. ergonomies	
	Naryous and andocrina systems. Sonsos	
	Nervous and endocrine systems. Senses.	
	Describe the nervous system in terms of	
	organization, structure and function.	
	E I de Company	
	Explain the four special senses and their relationship	
	to each other (taste, smell, hearing and sight)	
	Describe the endocrine system terms of hormones	
	and their effects.	
ANATOMY AND	Heart and circulatory system. Lymphatic system.	
PHYSIOLOGY IB	Respiratory system. Immunology	
	Explain the composition of blood is identified and	
	essential functions are explained.	
	Describe anatomy and physiology of the heart and	
	vascular systems.	2 X two hour theory test
	,	A supplementary test will be made
	Describe anatomy and physiology of the lungs and	available.
	respiratory tree.	
	' '	Each theory test will carry a
	Explain gas exchange in the lungs and body tissues.	weighting of 50%
	Explain mechanism of breathing.	
	Urinary system & reproductive system	
	Describe he anatomy and physiology of the urinary	
	system.	
	Explain the anatomy of the male and female	

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	reproductive systems is described.	
	Discuss the essential functions of the male and	
	female reproductive systems	
	Temale reproductive systems	
	Digestive system & nutrition.	
	Describe the anatomy and physiology of the	
	digestive tract and associated organs.	
	Explain the process of digestion.	
	F F	
	Consider the effects of food and nutrition on the	
	human body as it relates to digestion. E.g. Food	
	poisoning/ chemicals.	
	Describe the role of good nutrition in terms of macro &	
	micro nutrients and the importance of good diet.	
	Discuss the effects of poor nutrition on the human	
	body e.g. malnutrition.	
CELL BIOLOGY	pH and buffers	
	biomolecules and bio elements	TI /
	carbohydrates	Theory tests (average of all): 24%
	nucleic acids	Practical tests 10%
	amino acids	Practical reports 2%
	proteins	Assignments/oral presentation: 2%
	enzymes lipids	Tutorials, class/homework 2% Examination: 60%
	Imetabolism	Examination: 60%
	introduction to Polymerase Chain Reaction (PCR)	
IMMUNOLOGY	Development if immunology as a science; specific	
INTONOLOGI	immune response; non-specific immune response;	
	adaptive and innate immune response; antigen;	
	antibody; self and non self; primary and secondary	
	immune response; lymphoid organs; cells; functions	
	and structure	
	Structure of antigen and antigen receptor; growth	
	factors; relationship between growth factors and	- (() 240
	immune response	Theory tests (average of all): 24%
	Structure of the antibody; functions; induction of	Practical tests 10%
	antibody; effector functions; switch between classes;	Practical reports 2%
	classification and function of classes	Assignments/oral presentation: 2% Tutorials, class/homework 2%
	Humoural immunity; cell mediated immunity; human	Examination: 60%
	lymphocytic antigens;	Examination: 00%
	Histocompatibility	
	Shielding of antigen – recognition as self; disorders of	
	compliment deficiencies; hypersensitivity	
	autoimmune disorders; immune deficiencies; human	
	immunodeficiency virus	
	Properties of complement; nomenclature;	
	complement cascade; amplification loop; tick over;	
	regulation	
CORNERSTONE 101	The module content will be developed around the	
	concept of journeys, across time, across space, and	A weekly blog written by each
	across human relationships; the first use of the	student 20%
	concept will take the journey of the Umgeni River	Tutorial attendance (forfeited if
	(which is close to all DUT campuses) as a metaphor.	student attends less than 80% of
	The module will bring different disciplinary	tutorials) 10%
	perspectives to this content.	Visual artefact 15%
	The module will start with the analysis of a particular	Written report 30%
	issue or metaphor (one critical event or	Oral presentation 15%
	development will be and analysed; the event in focus	Peer assessment 10%
	will be selected on the basis of its connections to the	
	35 35.5ctcd on the basis of its connections to the	

	theme of journeys and its relevance to the issues of ethics, diversity and critical citizenry).		
	ethics, diversity and critical citizentry).		
	The final section of the module will identify and		
	integrate learning from earlier sections, and examine		
	implications for further learning. At each stage of		
	the module, students will be required to engage in		
	activities that involve reflection and build		
	communicative practices. There will be a concluding		
	section in which students will identify their learning		
	and examine the implications for their roles as		
	students and as citizens.		
VALUES IN THE	The module will begin with a reflection on personal		
WORKPLACE	values and move to a discussion on how they		
	intersect with values in the workplace. Small group		
	discussions will be formed around how to build		
	positive values in the workplace and the vital themes		
	of ethics, respect, interconnectedness, honesty,	Assignments 40 %	
	creativity and human diversity will form the basis for	Presentation 20%	
	building "sacred spaces at work." This will set the	Reflection 20%	
	tone to unpack issues around leadership values and	Peer assessment. 20%	
	ethics and ethical decision making. The final section		
	of the module will integrate all these aspects and		
	students will be required to identify the implications		
	of what they have learnt to develop social responsibility and their roles as citizens.		
LAW FOR LIFE	Introduction		
LAW TOKELLE	Civil and criminal law		
	Law of insurance	Assignment	60%
	Road accident fund	Poster	20%
	Law of contract	Will document	20%
	Marriage	· · · · · · · · · · · · · · · · · · ·	2070
	Succession		
WORLD OF WORK	Traditional and Modern CV Writing;		
	Who Am I?; (DISC, MBTI etc)		
	Job Searching;		
	Job Applications;		
	Networking;		
	Interviewing;		
	Body Language;		
	Verbal Communication;		
	Visual/Graphical Presentation;		
	What Is "Business"?		
	Career Path Options;		
	Work Readiness Expectations		
	Business Processes and Goals;		
	Organisational Aspects; Stress:	Tosts (average of all)	60%
	Business Ethics	Tests (average of all) Assignment	30%
	Etiquette - Telephone; Social Media, General	Classwork	10%
	Goal Setting & Time Management;	Classwork	1076
	Personal Finance		
	Numeracy		
	Project Management;		
	Meetings		
	Technical Report Writing;		
	Productivity in the Workplace		
	Quality in the Workplace		
	Health & Safety in the Workplace;		
	Housekeeping;		
	Computer and Technology Applications		
	Problem Identification & Solving;		
	Creativity, innovation and questioning		
1	Interpersonal Skills;		

PERSONAL AND PROFESSIONAL Brief overview of project development, implementation and evaluation Assignment Proposal Presentation	
Teamwork COMMUNITY HEALTH CARE AND RESEARCH I PERSONAL AND PROFESSIONAL Brief overview of problem identification in communities PROFESSIONAL Brief overview of project development, implementation and evaluation Theory Assignment Proposal Presentation	
COMMUNITY HEALTH CARE AND RESEARCH I PERSONAL AND PROFESSIONAL DEVELOPMENT I Brief overview of health systems in South Africa Brief overview of problem identification in communities Brief overview of project development, implementation and evaluation Brief overview of project development, implementation and evaluation Theory Assignment Proposal Presesntation	
CARE AND RESEARCH I PERSONAL AND PROFESSIONAL Brief overview of problem identification in communities Brief overview of project development, implementation and evaluation Brief overview of project development, implementation and evaluation Theory Assignment Proposal Presesntation	
PERSONAL AND PROFESSIONAL Brief overview of project development, implementation and evaluation Assignment Proposal Presentation	2021
PROFESSIONAL Brief overview of project development, Proposal implementation and evaluation Presentation	20%
DEVELOPMENT I implementation and evaluation Presentation	10%
	50%
	20%
Communication	
CLINICAL CHEMISTRY I Anticoagulants and preservatives	
Collection and handling of specimens	
Spectrophotometry	
Quality Assurance	
Reference ranges	
Automation principles and methods	
Amino acids, Plasma protein and albumin Theory tests (average of all):	24%
Principles of electrophoresis Practical tests	10%
Kidney function tests including urinalysis, osmolality,	2%
urine tests, calculi	2%
Tutorials class/homework	2%
Liver metabolites Examination:	60%
Use and maintain lab equipment	
Electrochemical techniques	
Electrolytes.	
Uric acid	
Acid/base balance	
Laboratory mathematics/calculations	
MEDICAL Introduction to medical microbiology	
MICROBIOLOGY I Good laboratory practices in the microbiology	
laboratory	
Instrumentation and its application in the laboratory	
Development of microbiological techniques and Theory tests (average of all):	24%
application Practical tests	10%
Practical reports	2%
Taxonomy and nomenclature of microorganisms Assignments/oral presentation:	2%
Microscopy and staining Tutorials, class/homework	2%
Bacterial cultivation and measurement Examination:	60%
Microbial metabolism (biochemical tests)	
Symbiotic relationship and establishment of disease	
Control of microorganisms	
Microbial genetics and recombinant DNA technology	
MEDICAL BACTERIOLOGY	
MICROBIOLOGY 2A Microbiology terminology and personnel	
responsibilities	
Collection, transport, processing of biological	
specimens	
Storage and disposal of biological specimen	
and waste	
Classification of medically important bacteria	
Laboratory identification of microorganisms	
, ,	24%
and specialised) Practical tests	10%
Practical reports	2%
PARASITOLOGY Assignments/oral presentation:	
Classification of medically important parasites Tutorials, class/homework	2%
, , , , ,	
, , , , , , , , , , , , , , , , , , ,	60%
Parasites pathogenesis	
Epidemiology	
Laboratory identification	
VIROLOGY	
Classification of medically important viruses	
Epidemiology Epidemiology	
Replication cycles Cell culture preparation and identification of	

	medically important viruses	
	MYCOLOGY Classification of medically important fungi	
	Fungal structures and reproduction	
	Classification of mycoses	
HAEMATOLOGY I	Blood formation, Cell development: Red cells, white	
I ALTIA I OLOGI I	cells, platelets	
	Structure and function of the bone marrow, cells,	
	haemoglobin	
	Growth factors and their effects: erythropoietin,	
	thrombopoietin, Interleukins, cytokines, other	
	growth factors	
	Factors affecting release of mature cells from the	
	marrow	
	Nutritional requirements in cell development: iron,	
	vitamin B ₁₂ , folate	
	Metabolic requirements of cells: Hexose	
	monophosphate shunt;	
	Rapaport-Leubering pathway; Glycolytic pathway;	
	Methaemoglobin reduction pathway; Glutathione	
	metabolism pathway	
	Processes leading to red cell destruction, features of	Theory tests (average of all): 24%
	haemolysis	Practical tests 10%
	Structure and function of organs involved in	Practical reports 2%
	haematopoiesis: spleen, thymus, lymph nodes, liver	Assignments/oral presentation: 2%
	The immune system: types of immune mechanisms,	Tutorials, class/homework 2% Examination: 60%
	immune responses The process of haemostasis including the coagulation	Examination: 60%
	cascade and fibrinolysis	
	Properties of a good anticoagulant and their effects	
	on specimens, good quality samples	
	Sites of blood and bone marrow collection,	
	principles and methods of tests and techniques: full	
	blood count, differential count, reticulocyte count,	
	coagulation studies, polymerase chain reaction,	
	diagnostic usefulness of bone marrow specimens	
	Storage protocol and the effects of storage on	
	haematological specimens	
	Protocols on reporting of laboratory results	
	Good laboratory practice including ethics, safety	
	principles	
	Principles of quality control programmes in	
	haematology	
IMMUNOHAEMATOLOGY	Blood donation criteria and testing.	
I	Procedures for the collection, processing and testing.	
	Storage and issuing of blood and blood products. Clinical indications for the use of blood and blood	
	products	
	Haemovigilance and biovigilance	
	Apheresis.	
	Clinical significance of blood group system antigens	Theory tests (average of all): 24%
	and antibodies.	Practical tests 10%
	Basic serological techniques.	Practical reports 2%
	Blood group interpretation	Assignments/oral presentation: 2%
	Causes of false results in laboratory testing	Tutorials, class/homework 2%
	Blood group reaction patterns and interpretation	Examination: 60%
	Compatibility and transfusion testing.	
	Selection of blood for cross-match	
	Risks and benefits associated with blood transfusion.	
	Transfusion transmitted diseases.	
	Haemolytic disease of the foetus and new-born	
	(HDFN)	
	Quality management systems.	

HISTOPATHOLOGY I	Laboratory administration – collection, logging,	
	distribution,	
	data recording, reporting, accession and retrieval of data.	
	Safety in the histopathology laboratory – recognize	
	dangers by fresh,	
	unfixed tissue biopsies. Storage and safe handling of	
	chemical and dyes.	
	Light and electron microscopy – behaviour of light	
	and electrons.	
	Fixation and fixatives – effects of specific fixatives on	Theory tests (average of all): 24%
	tissue and organs.	Practical tests 10%
	Poor fixation and fixation artefacts and corrective	Practical reports 2%
	action.	Assignments/oral presentation: 2%
	Tissue processing – familiar with the handling of the	Tutorials, class/homework 2%
	tissue processor and reagents used. Recognize processing artefacts and	Examination: 60%
	take corrective action.	
	Tissue embedding – embedding techniques of various	
	tissue biopsies.	
	Microtomy – familiar with the safety features and	
	how to use a microtome	
	for sectioning of various tissue types.	
	Staining – preparation and use of reagents used to	
	stain specific tissue	
	components and structures to contribute to	
SVT01 0 SV 1	diagnosis.	
CYTOLOGY I	The origins and role of Cytology as a discipline as	
	well as outline the professional and ethical role of a	
	cytotechnologist functioning in a Cytology laboratory.	
	Quality Assurance programme in a Cytopathology	
	LaboratoryThe role of automation in a cytology	
	laboratory, including Liquid- based Cytology and	
	Automated Screening Systems.	
	Growth and differentiation of cells and tissues.	
	The normal cells and tissues found lining the female	
	genital tract (FGT).	
	Collection and processing of cytological samples	
	specimens from the FGT. Cytological evaluation of specimens of the FGT	
	including normal constituents of the cervical smear,	
	infective agents (bacteria, fungi, parasitic and viral	
	agents), inflammatory, degenerative and regenerative	Theory tests (average of all): 24%
	changes and other non-neoplastic changes, (Acute	Practical tests 10% Practical reports 2%
	inflammation, chronic inflammation, Tissue repair,	Practical reports 2% Assignments/oral presentation: 2%
	follicular cervicitis, atrophic vaginitis, metaplasia,	Tutorials, class/homework 2%
	parakeratosis and hyperkeratosis)	Examination: 60%
	The effects of the reproductive hormones on the	
	cells of the FGT The morphogenesis and cytological presentation of	
	premalignant and malignant conditions of the FGT	
	(Natural history of cervical cancer, Pathogenesis of	
	cervical cancer, LSIL, HSIL, Squamous carcinoma,	
	,Adenocarcinoma, Rare Tumours (Clear cell	
	carcinoma, Hydatidiform mole; Choriocarcinoma;	
	Adenosquamous carcinoma, Lymphomas; Melanoma;	
	Sarcomas/ Mixed Mesodermal Tumours,	
	Extrauterine malignancies (ovary/ vulva); Metastatic	
	tumours)	
	Treatment of pre-malignant lesions, cytologic effects	
	of radiation and chemotherapy. General diagnostic application of	
	immunocytochemical techniques and molecular	
	minute Cytochemical techniques and molecular	

	biology to cytological samples including PCR of HPV	
	and genotyping.	
MOLECULAR BIOLOGY	Basic overview of DNA and RNA, the history and	
	their structure	
	Prokaryotic and Eukaryotic Genomes and DNA	
	replication DNA extraction; PCR Working with RNA; RNA extraction; Reverse Transcription and	Theory tests (average of all): 24%
	RT-PCR	Practical tests (average of all): 24%
	Gel Electrophoresis	Practical reports 2%
	DNA Sequencing	Assignments/oral presentation: 2%
	Restriction enzymes, Restriction mapping	Tutorials, class/homework 2%
	Cloning Vectors: plasmids, bacteriophages, cosmids	Examination: 60%
	Cloning: Ligation, transformation; construction of	Examinación.
	Gene (genomic)	
	libraries Cloning of cDNA libraries; Screening for	
	recombinant DNA	
FUNDAMENTALS OF	Medical terminology and internationally recognised	
PATHOLOGY	acronyms	Theory tests (average of all): 32%
	Cell adaptation and injury	Assignments/oral presentation: 5%
	Inflammation and healing.	Tutorials, class/homework 3%
	Classification, types and nomenclature of neoplasia	Examination: 60%
	Body fluid regulation and disturbances	
SYSTEMIC	Classification of body organs and systems	
PATHOPHYSIOLOGY	Disorders and diseases in the following systems:	
	 Cardiovascular system 	
	 Respiratory system 	Theory tests (average of all): 32%
	- Lymphatic system	Assignments/oral presentation: 5%
	- Digestive system	Tutorials, class/homework 3%
	- Endocrine system	Examination: 60%
	- Renal system	
	- Skeletal system	
	The physiological effects of each disorder.	
	The effects of the disorders on other body systems	
COMMUNITY HEALTH CARE AND RESEARCH II	Health systems in South Africa in comparison with	
CARE AND RESEARCH II	other successful third world countries like Brazil Brief overview of problem identification in	
	communities and identification of sector in which	Theory 20%
	primary problem is embedded	Assignment 10%
	Brief overview of project development,	Proposal 50%
	implementation and evaluation	Presentation 20%
	Communication and consultation to academic	20/0
	community	
	Communication to receivers of care	
THE ENTREPRENEURIAL	BECOMING AN ENTREPRENEUR	
EDGE	Understanding yourself	
	What kind of business will suite me best	
	A vision for the business	
	Why become an entrepreneur	
	Who are entrepreneurs	
	Entrepreneurial Resources	
	Entrepreneurial myths	
	Entrepreneurial transition	two tests and one assignment. The
	ADDRESSING RISK	weighting of all three
		assessments are equal. These
	Risks the banks are concerned with	three marks need to exceed
	From the perspective of the bank Risks and interest rates	50% for a pass.
	Researching to reduce my risks	
	Understanding my risks and prospects	
	Problem solving	
	Competitive advantage	
	Business successes and failures	
	Eddiness successes and land es	

UNDERSTANDING MY MARKET

What does my market look like

Sharing the market

Competitors

Suppliers

Customer Relations Management

PI ANNING

The environment

Strategic planning

Operation al planning

Types of plans

Setting the business vision

Determining the business mission

Setting business objectives

Finding and evaluating suppliers

FINANCIAL OBJECTIVES

Costing a product / service Funding the business

MARKETING

What you should now about products and

services

Considering the price

Finding the proper location

What to consider when advertising and doing promotions

ETHICS AND SOCIAL RESPONSIBILITY

Considering ethical issues to address

Drawing up an ethics standard

Being held ethically responsible

Being responsible to your stakeholders

THE GLOBAL ENVIRONMENT

The module content will include the following themes:

Environmental Pollution (Air, water and soil)

Differences between air, water and soil pollution in

terms of cause and effect.

Social, economic and personal impact on environmental pollution.

Pollution control strategies.

Local case studies.

Population growth vs. natural resources

Population growth trends in developed vs developing

Social, economic and environmental impacts of human population growth in the global context. Strategies to curb population growth

Climate change and global warming

Causes of increased global mean temperatures. Impact of climate change on extreme weather

Consequences of climate change on human health, natural resources and biodiversity.

Sustainable development

Concept of sustainable development within the

South African and global context

Inter-relationships between sustainable development, social responsibility, economic development and environmental protection.

FOLIALITY ASS	To		
EQUALITY AND	Concepts and terminology – e.g. diversity, equality,		
DIVERSITY	inclusion, power, oppression		
	Parameters of diversity as listed in section 9 of the	Theory	33%
	SA Constitution	Reflective writing assignment	17%
	Prejudice, discrimination and inequality	Group presentation	17%
	The diversity competence continuum	Diversity festival	33%
	Steps to develop competence/sensitivity in relation	Diversity lestival	JJ /6
	to diverse others		
	Selected topics		
CLINICAL CHEMISTRY 2	Endocrinology		
	Secretion and regulation, hormones of hypothalamus, pituitary, pineal, thyroid, adrenal, gonads, pancreas, GIT		
	Carbohydrate metabolism Intermediary carbohydrate metabolism, hormonal regulation, disorders [glucose, lactate], ketogenesis, glycosylated Hb,		
	fructosamine, xylose		
	Lipid metabolism Lipid constituents, lipoproteins and		
	disorders, serum lipid and lipoprotein analyses, total fecal fat/steatocrit/oral fat loading test		
	Body fluid analysis CSF [glucose, proteins], amniotic fluid [congenital disease, neural tube defects,	Theory tests (average of all): Practical tests	24% 10%
	hemolytic disease, gestational age, fetal pulmonary development], sweat [inc	Practical reports Assignments/oral presentation Tutorials, class/homework	2% n: 2% 2%
	sweat analysis], synovial fluid, serous fluid [pleural, pericardial, peritoneal], transudates and exudates	Examination:	60%
	Tumour markers Properties, classification, markers: PSA, AFP, CEA, CA 125, 153, 199		
	Pharmacology Introduction [classification, routes of administration,		
	terminology], receptor theory, elementary pharmakokinetics, drugs subjected to TDM [Digoxin,		
	Phenytoin, Phenobarbitol, Carbamazapine,		
	Theophylline, Valproic acid, Lithium, Paracetamol,		
	Salicylates, Tricyclic Antidepressants, Cyclosporin,		
	Amikacin, Gentamycin and Vancomycin], techniques		
	of drug analysis [EMIT, ELISA, EI, HPLC, GLC, TLC], toxicology [ethanol, salicylates, paracetamol, barbiturates]		
MEDICAL	Laboratory administration – collection, logging,		
MICROBIOLOGY 2B	distribution,		
MICROBIOLOGI 2B	data recording, reporting, accession and retrieval of		
	data.		
	Safety in the histopathology laboratory – recognize	Theory tests (average of all):	24%
	dangers by fresh,	Practical tests	10%
	unfixed tissue biopsies. Storage and safe handling of	Practical reports	2%
	chemical and dyes.	Assignments/oral presentation	ո։ 2%
	Light and electron microscopy – behaviour of light	Tutorials, class/homework	2%
	and electrons.	Examination:	60%
	Fixation and fixatives - effects of specific fixatives on		
	tissue and organs.		
	Poor fixation and fixation artefacts and corrective		
	action.		
		1	

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	Tissue processing – familiar with the handling of the	
	tissue processor and	
	reagents used. Recognize processing artefacts and	
	take corrective action.	
	Tissue embedding – embedding techniques of various	
	tissue biopsies.	
	Microtomy – familiar with the safety features and how to use a microtome	
	for sectioning of various tissue types.	
	Staining – preparation and use of reagents used to	
	stain specific tissue	
	components and structures to contribute to	
	diagnosis.	
PERSONAL AND	Revision of the basic elements of Writing.	
PROFESSIONAL	Intermediate elements of Writing.	
DEVELOPMENT II	Effective communication and self-expression.	
DEVELOTTIENT II	Community: Experience other communities; a	Write critically reflective pieces on
	variety of social contexts, identify the problems and	each experience, guided by a series
	see if they can play a role in addressing them.	of questions (e.g. a SWOT
	8 Experiences: E.g. Soup kitchen, Children's home,	analysis), identifying the role
	Playhouse (4 disadvantaged settings, I western	players in the community and
	traditional experience, I Indian cultural experience,	seeing their roles.
	I African traditional experience, I outdoor	seeing their roles.
	experience)	
	The student would be required to choose to attend	
	at least 4 of these	
HAEMATOLOGY 2	Classification and clinical features, causes, laboratory	
	features and management of anaemias, leukaemias,	
	malignancies, platelet and haemostatic disorders and	
	disorders associated with systemic non-	
	haematological disorders	Theory tests (average of all): 24%
	Principles of quality control and quality assurance	Practical tests 10%
	and troubleshooting	Practical reports 2%
	Assessment of specimen suitability	Assignments/oral presentation: 2%
	Correct terminology when reporting results	Tutorials, class/homework 2%
	The clinical significance of laboratory results,	Examination: 60%
	including reticulocyte counts,	
	full blood counts, coagulation tests, screening tests,	
	confirmatory tests	
CYTOLOGY 2	Collection and preparation of cytological specimens	
	and the normal cells and tissues found lining the	
	following sites in the body:	
	- respiratory tract	
	-serous effusions	
	-urinary tract	
	-central nervous system	
	-gastro intestinal tract.	
	Cytological evaluation of specimens including normal	
	constituents , infective agents (bacteria, fungi,	Theory tests (average of all): 24%
	parasitic and viral agents), inflammatory,	Practical tests 10%
	degenerative and regenerative changes and other	Practical reports 2%
	non-neoplastic changes of the respiratory tract,	Assignments/oral presentation: 2%
	serous effusions, urinary tract, central nervous	Tutorials, class/homework 2% Examination: 60%
	system and gastro intestinal tract.	Examination: 60%
	The morphogenesis and cytological presentation of premalignant and malignant conditions of the	
	respiratory tract, serous effusions, urinary tract, central nervous system and gastro intestinal tract.	
	General diagnostic application of	
	immunocytochemical techniques and molecular	
	biology to cytological samples including PCR as	
	applicable.	
	Respiratory Tract: collection and microscopic	
	mespiratory fract, collection and microscopic	

features in sputa and bronchial brushings/ lavages and FNAB

Inflammation: Non-specific inflammation,

Tuberculosis, Eosinophilia

Common infective agents and characteristic cytopathic effect for each agent, including Entamoeba sp, Actinomyces sp, Candida sp, Blastomyces sp, Cryptococcus sp, Aspergillus sp, Histoplasmosis sp, Coccidioides sp, Cryptococcus sp, Pneumocystis sp, Echinococcus sp, Entamoeba sp.

Other elements: Ferruginous bodies, Curshmann's spirals, Vegetable cells, Charcot-Leyden crystals. Benign reactive: Bronchial hyperplasia and bronchial metaplasia. without/ with atvoia.

Lung cancer and its pathogenesis, including known carcinogens

Malignant: Squamous carcinoma, Bronchogenic adenocarcinoma, and Bronchoalveolar carcinoma, Sm (neuroendocrine) carcinoma, Large cell undifferentiat carcinoma, Outline other primary/ metastatic tumour The effects of radiation and chemotherapeutic agents

on benign and malignant cells

Urinary tract: Collection techniques, Cytological changes that occur with different inflammatory processes, including those associated with pathogens (esp. Schistosoma haematobium) Casts (e.g. hyaline, granular, cellular) and pathologically significant crystals. Potential sources of diagnostic error in evaluating urinary tract specimens including ileal bladder urine, lithiasis, malakoplakia, etc Malignancies of kidney and urinary tract: (urine/ FNAB): Epithelial tumours of renal pelvis, ureter and urinary bladder: Transitional cell carcinoma, Adenocarcinoma, Squamous carcinoma, Renal cell carcinoma, Wilms' tumour, Other, Metastases.

Effects of radiation and chemotherapeutic agents on benign/ malignant cells, transplant rejection. atypia and its causes, including lithiasis and malakoplakia. latrogenic changes (incl. ileal conduits) and potential pitfalls. Transplant rejection changes.

Central nervous system: Anatomy of brain and spinal cord Macroscopic presentation and significance, fixation, preparatory techniques. "Normal" cells (shunt picture). Meningitis: Bacterial, Viral, TB, Cryptococcal; Parasites. Primary tumours of the CNS; Neural crest tumours; Lymphoma/ leukaemia, midline tumours and miscellaneous 1º tumours, metastatic malignancy.

Gastro intestinal tract

Anatomy of brain and spinal cord. Macroscopic presentation and significance, fixation, preparatory techniques. "Normal" cells (shunt picture). Meningitis: Bacterial, Viral, TB, Cryptococcal; Parasites

Primary tumours of the CNS; Neural crest tumours; Lymphoma/ leukaemia. Miscellaneous 1º tumours. Metastatic malignancy

CLINICAL LABORATORY

PRACTICE I

Clinical Chemistry

Specimen / chemical safety procedures.

Quality control and workflow.

Laboratory calculations and preparation of solutions.

Description of the automated instrument.

Compulsory analytes: Sodium, potassium, chloride, total C02, urea, creatinine and glucose.

All laboratory tests / profiles in chemical pathology.

Selection of the following topics: -

Atomic absorption

Blood gases

Chromatography

Drugs

Electrophoresis

Endocrinology

Nephelometry

Urinalysis

Medical Microbiology

Biosafety protocols applicable to the Microbiology laboratory.

Explain the principles of automated instruments used in the laboratory (where applicable).

Process the following specimens in the laboratory: -

Faeces

Swabs and Pus

CSF

Sputum

Ürine

(Range Statement: Includes staining, microscopy, culture, antibiotic susceptibility and identification of organism/s).

Culture media preparation

(Range Statement: Basic principles of selective, enriched and differential media including antibiotic containing media).

Quality assurance systems.

Virology

Safety

Processing of viral specimens:

Culture and identify viruses in specimens

Media preparation and cell cultures

Serology (HIV, Hepatitis other)

PCR

Blood Transfusion discipline

Donor selection

ABO and Rh Crossmatching

ABO and Rh blood typing

Cytology

Set up microscope incl. Köhler illumination

Female genital tract

Inflammation; Benign proliferative reactions

Reactive cellular changes; Microorganisms/ agents of infection

Squamous abnormalities: ASCUS, LSIL, HSIL, SCC Glandular abnormalities: AGUS (outline),

adenocarcinomas

Urinary tract

Normal, Agents of infection (esp Schistosoma)

Average mark obtained from discipline based assessments 60% Portfolio 30% Learning logs 10%

Malignancy: transitional cell carcinoma, squamous ca, adenocarcinoma

Respiratory tract

Normal; Non-cellular findings (incl. ferruginous bodies); Agents of infection

Inflammation (incl. asthma); Bronchial metaplasia and hyperplasia;

Malignancy: adenocarcinoma, squamous carcinoma, undifferentiated

Serous effusion

Normal; Inflammatory/ non-malignant disease states; Malignancy 10 / 20 tumours, incl. carcinoma, lymphoma, melanoma

Serous effusion: prepare and stain two samples (Pap; MGG stain)

Complete assignment on filter preparations independent

Histopathology

Embedding; Microtomy; Routine H&E staining and mounting

Trim blocks and cut 8 sections of kidney tissue biopsies for special staining techniques.

Special staining techniques:

PAS; PAS/D; Alcian blue; Verhoeff's; Methanamine silver, Toluidine blue; Reticulin, Masson's Trichrome Special techniques: Transmission electron microscope; Immunohistochemistry Frozen sections Stain two sections: one by rapid H&E method and the other for fat.

Electron Microscopy. Molecular laboratory.

Haematology

Specimen processing, handling, safety procedures and ethics.

Quality control principles.

Perform tests and techniques, following standard operating procedures.

Interpretation of laboratory results, correlation of FBC with the findings of

the peripheral blood film.

Professional conduct, principles of good laboratory practice including ward visits for BM, finger-prick and/or blood collection

PRINCIPLES OF	Management Principles (Planning, leading	T	
MANAGEMENT	organizing and control, problem identification &		
	solving, decision making, communication,		220/
	negotiation, conflict resolution, leadership,		32%
	motivation)	Assignments/oral presentation	
	Organisational Development	Tutorials, class/homework	3%
	Change Management	Examination:	60%
	Resource Management		
	Industrial Relations		
	Quality Assurance and Safety including Legislation		
RESTORATIVE JUSTICE	Relevance of a restorative approach in the SA		
,	context.		
	Aspects of legislation and policy.		
	Restorative philosophy and practice in indigenous		
	communities.		20%
	Factors in crime, violence and conflict in modern	Group work	10%
	societies.		10%
	The social control window.	Independent study	40%
		Presentations	10%
	Restoration versus retribution.		
	Shaming, integration, healing and forgiveness.		
	The restorative practices continuum.		
	Informal and informal restorative conferencing.		
COMMUNITY HEALTH	Transformation of Health systems in South Africa in		
CARE AND RESEARCH III	comparison with other successful third world		
	countries like Brazil		
	Brief overview of project evaluation in communities		
	and identification of and evaluation of performance	Theory	20%
	of sector in which primary problem is embedded	Assignment	10%
	Continue project development, implementation and		50%
	evaluation		20%
	Communication and consultation to academic		
	community		
	Communication to receivers of care		
	Communication to high level stakeholders		
PERSONAL AND	In groups of four students, Identify a sustainable		
PROFESSIONAL	community upliftment project		
DEVELOPMENT III	Term I – Formulate a proposal for the project,		
DEVELOPMENT III	including funding proposals, project plan and business		
	plan		
	Writing a proposal, a project plan, and a business		
	plan.		
	Terms 2 and 3 – Implement the project and submit	Portfolio of evidence: Proposal	
	monthly progress report	monthly progress reports and	Final
	Gimme 5 Units: Environmental Awareness and	report	
	Professionalism & Work Ethics.		
	Responsibilities and effects of change from each		
	stage of development: social adjustments.		
	Term 4 – Write a full report on the project,		
	including outcomes and plans to ensure its		
	sustainability		
	Writing a report		
PRINCIPLES OF	The use of the library		
RESEARCH	Referencing		15%
	Plagiarism	Journal article	10%
	Writing up of research findings; posters, publication,	Poster	10%
	dissertation thesis	Research Proposal	10%
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DESEADOU DROIECT			
RESEARCH PROJECT	Statistics reinforce	This module will remain incom	plete
RESEARCH PROJECT MODULE A	Statistics reinforce Literature review	This module will remain incomin Semester I of the fourth year	
	Statistics reinforce Literature review Research methods	in Semester I of the fourth yea	r of
	Statistics reinforce Literature review Research methods Research ethics	in Semester I of the fourth year study. The module is linked to	r of the
	Statistics reinforce Literature review Research methods Research ethics Plagiarism	in Semester I of the fourth year study. The module is linked to Research Project Module B off	r of the
	Statistics reinforce Literature review Research methods Research ethics	in Semester I of the fourth year study. The module is linked to	r of the

DECEMBELL DROVE OF	In	1
RESEARCH PROJECT	Research methods	Research project Mod A mark 30%
MODULE B	Literature review	Draft chapters 20%
	Writing up of research findings: posters, publication,	Complete light bound dissertation50%
	dissertation thesis	F 6
	General aspects of disease	
	Chromosomal disorders	
	Pathophysiology of the following systems and	
	integrating these with other systems and laboratory	
	results	No exam, mark contributes to course
	Central nervous system	mark calculation in Module B
	Endocrine system	
	Cardiovascular	
	Respiratory	
	· · · ·	
1117500 4 750	Immunology	
INTEGRATED	Pathophysiology of the following systems and	TI (
PATHOPHYSIOLOGY	integrating these with other systems and laboratory	Theory test (average of all) 24%
MODULE B	results	Assignemnt/oral presentation 8%
	Gastrointestinal	Case studies (tuts) 4%
	Renal	Online tuts 4%
	Blood and bone marrow Reproductive systems	Examination 60%
	Integumentary	
LABORATORY	Legal and social aspects of Healthcare	
MANAGEMENT	Resource management in healthcare settings	Theory tests 24%
	Budgeting and financial management in Healthcare	Oral Presentation 8%
	Leadership in Healthcare settings	Reflective journal 8%
	Relevant legislation pertaining to private practice	Examination 60%
		Examination 60%
CLUMICAL LABORATORY	Laboratory accreditation	
CLINICAL LABORATORY		
PRACTICE 2: INCLUDES		
THE FOLLOWING		
SPECIALISATION		
OPTIONS FROM I = 10		
BELOW (THE STUDENT		As somethis absence also etimo balance
WILL HAVE TO SELECT		As per the chosen elective below
ONE OF THESE		
ADVANCED		
SPECIALIZATION		
MODULES AT 52		
CREDITS):		
CLINICAL PATHOLOGY	Statutory regulations and ethics	
MODULE A	Specimen requirements and suitability including	
MODULE A	storage for all laboratory analysis	
	Laboratory equipment (all types of equipment	
	Laboratory reagents	
	Total Quality management ; Quality control	
	Personnel (personnel documents and records)	
	Stock control (storage, receipt, procurement, expiry	
	date)	
	Documentation	
	Laboratory safety	
	Laboratory related mathematics	
	Molecular biology techniques	
	Special tests and specimens related to the following	
	specific disciplines:	
	Clinical Chemistry	
	Safety and GLP	
	Workflow, collection and processing of routine	
	samples in a Chemical Pathology laboratory.	
	Knowledge of quantitative, semi-qualitative and	
	qualitative tests (automated or manual) for analytes	
	on either blood, serum, plasma, urine (timed and	
	random), CSF, aspirates/ fluids with particular	
	reference to:	

Reagent, controls and calibrators preparation; Calibration and Q.C procedure;

Operation of instrument/ method procedure

Medical Microbiology

Specimen collection, transport, processing and disposal of specimen with pathogenic microorganisms

Identification of pathogenic microorganisms from clinical specimens.

Quality assurance system

TB/HIV management system

Haematology:

The full blood count including all calculations and interpretation of scatter grams; manual and automated cell counts

Preparation of all types of smears and the calculation of absolute counts:

Collection and handling of blood samples pathogenesis,

laboratory diagnosis and interpretation of morphology of peripheral blood and bone marrow smears of normal red cell and red cell disorders Tests used in the diagnosis and monitoring of red cell disorders haemolytic anaemias the pathogenesis, the interpretation and correlation of the tests with the clinical presentation.

Basic blood transfusion techniques including blood grouping and direct antiglobulin test (Coombs test).

CLINICAL PATHOLOGY MODULE B

Clinical Chemistry

Workflow, transportation and processing of specialised tests in a Chemical Pathology laboratory. Knowledge of quantitative, semi-qualitative and qualitative tests (automated or manual) for analytes on faeces and amniotic fluid with particular reference to: Operation of instrument/ method procedure Safety and GLP.

Medical Microbiology

Infection control

Laboratory accreditation and administration

Water examination Milk examination

Haematology:

The full blood count including all calculations and interpretation of scatter grams; manual and automated cell counts

Collection and handling of blood samples pathogenesis, laboratory diagnosis and interpretation of morphology of peripheral blood and bone marrow smears of normal white cell and haematological malignancies

Tests used in the diagnosis and monitoring of white cell disorders, the interpretation and correlation of the tests with the clinical presentation. Understanding the current classifications including both WHO and FAB.

CD4 counting with all gating strategies Cytochemistry, immunophenotyping (principles, application and interpretation of flow cytochemistry) Practical tests + workbook Assignment 5% Examination: 50%

Theory tests (average of all): 15%

30%

CLINICAL CHEMISTRY 3A	Knowledge of quantitative, semi-qualitative and		
	qualitative tests (automated or manual) for the		
	following analytes on either blood, serum, plasma,		
	urine (timed and random), CSF, aspirates/ fluids,		
	faeces and amniotic fluid with particular reference		
	to:		
	Reagent, controls and calibrators preparation; Calibration and Q.C procedure;		
	Operation of instrument/ method procedure;		
	Sodium, Potassium, Chloride, Bicarbonate (TCO ₂),		
	Urea, Creatinine, Cystatin C, Uric Acid, Calcium,		
	Ionized Calcium, Magnesium and Inorganic		
	Phosphorous.		
	Glucose, Ketones, Hb A1c (Glycated Haemoglobin),		
	Fructosamine and MAU (Microalbumin).		
	Cholesterol, High Density Lipoprotein (HDL), Low		
	Density Lipoprotein (LDL), Triglyceride,		
	Lipoprotein (a) and Apolipoprotein A&B.		
	Total Protein, Albumin, Globulin, Total Bilirubin,		
	Conjugated and Unconjugated Bilirubin, ALP, GGT,		
	AST, ALT and LDH.		
	Amylase, Lipase & Cholinesterase (serum & red cell).		
	CK, CKMB (mass/Activity), Troponin (T/I),		
	Myoglobin, Pro-BNP/ BNP and Homocysteine.		
	Iron Studies: Ferritin, Iron and Transferrin	No exam, assessment marks	
	Lactate, Ammonia.	contribute to course mark.	
	Digoxin, Phenytoin, Phenobarbitol, Carbamazapine,		
	Theophylline, Valproic acid, Lithium, Paracetamol,		
	Salicylates, Tricyclic Antidepressants,		
	Cyclosporin, Amikacin, Gentamycin and		
	Vancomycin, Benzodiazepine, Cannabis,		
	Amphetamine, Barbiturate, Cocaine, Methadone,		
	Methaqualone, Opiate and PCP		
	TSH, T3, T4 (Free and Total), Qualitative and		
	Quantitative bHCG, FSH, LH, Estradiol (E2),		
	Growth Hormone, Testosterone, Progesterone,		
	Prolactin, Aldosterone, Cortisol, Gastrin,		
	Histamine, Insulin, Renin, Vitamin		
	B12, Folate, PTH and ACTH		
	PSA, AFP, CEA, CA markers 125, 153 & 199.		
	CRP, Ultra-sensitive CRP, PCT (procalcitonin).		
	IgE, IgM, IgG, IgA, b2 Microglobulin, C3 and C4,		
	Haptoglobins, SACE,		
	Caeruloplasmin.		
	Xylose, Phenylalanine, Ascorbic acid		
	Osmolality		
	Blood Gases and Co-oximetry Neonatal bilirubin		
	Catecholamines, 5HIAA, 17 Hydroxycorticosteroids.		
	Total Faecal Fat/ Steotocrit/ Oral Fat Loading Test.		
CLINICAL CHEMISTRY 3B	Knowledge of quantitative, semi-qualitative and		
CENTICAL CHEMISTRY 3B	qualitative tests (automated or manual) for the		
	following analytes on either blood, serum, plasma,		
	urine (timed and random), CSF, aspirates/ fluids,		
	faeces and amniotic fluid with particular reference		
	to:	Theory tests (average of all):	5%
	Reagent, controls and calibrators preparation;		0%
	Calibration and Q.C procedure;	Assignment	5%
	Operation of instrument/ method procedure;		50%
	Serum and urine Protein Electrophoresis, IFE /		
	Kappa and Lambda free light chains.		
	Urine bHCG and Dry Chemistry (dipstick and		
	ketostix).		
	Faecal & urine reducing substances, Porphobilinogen,		

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	Porphyrin. Occult Blood/ Faecal Haemoglobin/ Colon Albumin. Calculus analysis Knowledge of the following laboratory function tests or profiles with reference to: Association/ relevanc to the specific organ, Association/ correlation between the tests, The significance and interpretation of abnormal results, Procedure when results do not concur with clinical picture Renal: Sodium, Potassium, Urea and Creatinine including Creatinine Clearance, pH and Base Excess. Liver: ALT, AST, GGT, ALP, LDH, Total Protein, Total and Conjugated Bilirubin. Cardiac: CK, CKMB, Troponin and Myoglobin. Lungs: pH, PCO2, PO2, TCO2 and O2 Saturation; Actual and Standard Bicarbonate, and Base excess. Thyroid: TSH, Free T3 & T4. Pancreas: Amylase (Total and Pancreatic), Lipase. Toxicology: Organophosphate and Salicylate poisoning. Menopausal Screen: LH, FSH and E2 (Estradial)		
MEDICAL			
MICROBIOLOGY 3A	Specimen collection, transport, processing and disposal of specimen with rare / unusual microorganisms Identification of rare / unusual microorganisms from clinical specimens. TB/HIV management system Genotyping characterisation of microorganisms		
MEDICAL	7. 3	The same tracks (s. 10)	1.07
MEDICAL	Infection control and epidemiology		15%
MICROBIOLOGY 3B	Laboratory accreditation and administration		20%
	Quality management system	Assignment	5%
	Public Health	Examination:	50%
CYTOLOGY 3A	Anatomy, histology, cytology, applications and techniques, benign lesions and malignant lesions from the following sites: breast and nipple secretions, thyroid, lymph nodes, salivary glands, liver, pancreas, testes, ovaries, prostate. Principles of specialised sample collection techniques from the sites of the organs listed above including fine needle aspiration biopsies (FNAB). Tests and techniques for the interpretation and distinction between normal and abnormal cytology results. Correlation of results with clinical information. Safety, ethics and quality control principles General diagnostic application of immunocytochemical techniques and molecular biology to cytological samples including PCR as applicable including PCR of HPV and genotyping.		
CTTOLOGT 3B	Anatomy, nistology, cytology, applications and techniques, benign lesions and malignant lesions from the following sites: Rare Tumours of the female genital tract (Clear cell carcinoma, Hydatidiform mole; Choriocarcinoma; Adenosquamous carcinoma, Lymphomas; Melanoma; Sarcomas/ Mixed Mesodermal Tumours, Extrauterine malignancies	Theory tests (average of all): Practical tests Practical reports Assignments/oral presentation: Tutorials, class/homework	24% 20% 2% 2% 2% 50%
	(ovary/ vulva); Metastatic tumours). Principles of specialised sample collection techniques from the sites of the organs listed above including	Examination:	30%

	fine needle aspiration biopsies (FNAB).	
	Tests and techniques for the interpretation and	
	distinction between normal and abnormal cytology	
	results.	
	Correlation of results with clinical information.	
	Safety, ethics and quality control principles.	
	Treatment of pre-malignant gynaecologic lesions and	
	cytologic effects of radiation and chemotherapy.	
	General diagnostic application of	
	immunocytochemical techniques and molecular	
	biology to cytological samples including PCR as	
	applicable including PCR of HPV and genotyping.	
HAEMATOLOGY 3A	Routine and specialised haematology investigations:	
	the full blood count including all calculations and	
	interpretation of scatter grams; manual and	
	automated reticulocyte counts; differential counts	
	including the preparation of all types of smears and	
	the calculation of absolute counts; erythrocyte	
	sedimentation rate; collection and handling of blood	
	samples; CD4 counting with all gating strategies.	
	Pathogenesis, laboratory diagnosis and interpretation	
	of morphology of smears of peripheral blood and	
	bone marrow of normal; all anaemias; inclusion	
	bodies in red cells; blood parasites; haemolysis and	
	haemolytic anaemias.	
	Basic blood transfusion techniques including blood	
	grouping and direct antiglobulin test (Coombs test).	
	Good laboratory practice including laboratory safety	
	and ethics	
HAEMATOLOGY 3B	Routine and specialised haematology investigations:	
	the full blood count including all calculations and	
	interpretation of scatter grams; differential counts	
	and the calculation of absolute counts; CD4 counting	
	with all gating strategies.	
	Pathogenesis, laboratory diagnosis and interpretation	
	of morphology of smears of peripheral blood and	
	bone marrow of normal; benign white cell disorders;	
	myeloproliferative disorders; myelodysplasia;	
	lymphoproliferative disorders; acute leukaemias;	
	platelet disorders; inclusion bodies in white cells; the	
	effects of HIV on blood smears and the theoretical	
	knowledge of bone marrow features of disorders;	
	tests used in the diagnosis and monitoring of	Theory tests (average of all): 24%
	haemostatic disorders including thrombosis and	Practical tests (average of all). 24%
	anticoagulant therapy; vascular disorders; factor	Practical reports 2%
	inhibitors; theoretical knowledge of haemophilia	Assignments/oral presentation: 2%
	factor V Leiden and other inherited thrombophilia	Tutorials, class/homework 2%
	disorders and PK assay.	Examination: 50%
	The pathogenesis and laboratory diagnosis of all	Examination. 50%
	haematological malignancies, the interpretation and	
	correlation of the tests with the clinical presentation,	
	understanding the current classifications including	
	both WHO and FAB including cytochemistry,	
	immunophenotyping (principles, application and	
	interpretation of flow cytochemistry), principle of	
	ISHAGE gating strategy of the enumeration of	
	CD34+ stem cells, cytogenetic techniques, FISH and	
	molecular diagnostic techniques in	
	haematopathology.	
	Good laboratory practice including laboratory safety	
	and ethics	
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HISTOPATHOLOGY 2A	Embedding of various tissue biopsies according to	
	their structural features.	
	Microtomy – thorough knowledge of microtomes	
	and microtome knives.	
	Able to section various tissue biopsies and recognise	
	cutting artefacts and	
	employ corrective measures.	
	Frozen sections – assist in the diagnosis of urgent	
	biopsies that require	
	the use of a cryostat to produce frozen sections.	
	Staining of specific elements – deduce which	
	stain to use for a specific	
	component / structure. Recognise staining artefacts	
	and use corrective	
	measures. 'Trouble-shoot' out of the ordinary	
	staining reactions.	
	Histology of tissues - Identify and describe the	
	tissue types as well as the	
	structure of each organ system. Identify the	
	structures specific to each	
	organ or system.	
HISTOPATHOLOGY 2B	Molecular Biology – have a thorough knowledge	
	of the tests required in	
	Molecular biology to diagnose tumours and bacteria.	
	Knowledge of in situ hybridisation (DISH)	
	Enzyme histochemistry – Simultaneous capture,	
	post-incubation coupling,	
	self coloured substrate and intramolecular	
	rearrangement.	Theory tests (average of all): 24%
	Metal precipitation for enzyme detection.	Practical tests (average of all). 24%
	Immunocytochemistry – able to distinguish between the various	
		Assignments/oral presentation: 2%
	0 1	Tutorials, class/homework 2%
	cases that cannot be	Examination: 50%
	assessed with special staining procedures.	
	Electron microscopy – fixation and processing of	
	specimens for analyses	
	under an electron microscope. Recognise	
	ultrastructural organelles and	
	components of the cells using an electron	
	microscope.	
IMMUNOHAEMATOLOGY	Ethics	
2 A	Health and Safety	
	Transfusion transmitted diseases	
	Blood donation	
	Blood Processing and component therapy	
	Donation testing	
	Storage and issue of blood and blood products	
	Blood cold chain	
	Clinical indications for the use of blood and blood	
	products	
	Introduction to risks and benefits associated with	
	transfusion.	
	Introduction to the haemolytic disease of the foetus	
	and new-born (HDFN)	
	Haemovigilance and biovigilance	
ĺ		
i	Apheresis.	
	Apheresis.	
	Apheresis. Clinical significance of blood group system antigens	
	Apheresis. Clinical significance of blood group system antigens and antibodies.	
	Apheresis. Clinical significance of blood group system antigens and antibodies. Basic serological techniques	
	Apheresis. Clinical significance of blood group system antigens and antibodies. Basic serological techniques Causes of false results in laboratory testing	
	Apheresis. Clinical significance of blood group system antigens and antibodies. Basic serological techniques Causes of false results in laboratory testing Antigen antibody reactions in transfusion testing	

	Compatibility and transfusion testing.		
	Quality management systems.		
IMMUNOHAEMATOLOGY	Risks and benefits associated with transfusion.		
2B	Haemolytic disease of the foetus and new-born (HDFN)		
	Reagent preparation and standardization		
	Paternity testing	Theory tests (average of all):	24%
	HLA testing	Practical tests	20%
	Transfusion reaction investigations	Practical reports	2%
	Antenatal Investigations	Assignments/oral presentation	n: 2%
	Postnatal (Cord and Maternal) Cases	Tutorials, class/homework	2%
	Transfusion reaction investigations	Examination:	50%
	Antenatal Investigations		
	Postnatal (Cord and Maternal) Cases		
	Quality management systems.		

16.2 SUBJECT CONTENT: CLINICAL TECHNOLOGY

16.2.1 ND: CLINICAL TECHNOLOGY

NB: Students to read this section in conjunction with the relevant Student guides

Module Name	Learning Content	ASSESSMENT	
	-	The CONTINUOUS ASSE	SSMENT
FOUNDATION PHYSICS	Basic Mathematics, vectors, Problem solving	mark shall be made up of	
(FPYCI0I)	skills in Physics, Conceptual physics	Theory tests:	60%
		Practical tests:	40%
	Introduction to biomedical instrumentation,	The CONTINUOUS ASSES	SMENT
FOUNDATION BIOMEDICAL	Medical terminology and physiological	mark shall be made up of	
APPARATUS (FBAPI0I)	measurements, Bio-signals and noise, Bio-	Theory tests	60%
ATTAKATOS (I BATTOT)	medical electronics – Analog and digital, and SI	Practical tests	30%
	metric units and equivalencies.	Assignment	10%
	Introduction to specialist categories, Infection		
	control, Sterilisation and disinfection techniques,		
INTRODUCTION TO	Medical and surgical asepsis, Communicable	Theory tests	50%
CLINICAL TECHNOLOGY	disease patient control, Laboratory techniques	Practical tests	30%
(ICLT101)	(microscopes, incubators, refrigerators and	Assignments	20%
	autoclaves), Safety, and Language practices and		
	conventions		
	Introduction to inflammation, Diseases caused	The CONTINUIOUS ACCES	CMENIT
FOUNDATION ORGANS &	by inflammation and associated changes to tissue architecture, Introduction to genetics and	The CONTINUOUS ASSES mark shall be made up	
SYSTEMS	diseases, Introduction to genetics and	Theory tests	p of 70%
PATHOPHYSIOLOGY (FOI0I)	mechanisms related to pathogenesis, and	Assignments	30%
	Introduction to cell injury and cell death	V2218111161172	30/6
	Introduction, Nervous system, Endocrine		
	system, Cardiovascular system, Immunology	Theory Tests	30%
PHYSIOLOGY I (PSII02)	Respiratory system, Gastrointestinal system,	Practical Tests	10%
	Renal system, Reproductive system	Examination Mark	60%
	renai system, reproductive system	Theory Tests 20)%
		Practical Work	16%
	Introduction to Anatomy, Thorax, Abdomen	Attendance	4%
ANATOMY I (ANAYI0I)	and Pelvis, Limbs, Neuroanatomy, Head and	Examination Mark	60%
` ,	Neck	PAPER I: Theory (75% of Ex	kam Mark)
		and	,
		PAPER II: Spotter (25% of Exa	am Mark).
	Atomic structure, Periodic table, Molecular	Assessment Plan	
CHEMISTRY (CHEMI01)	elements and compounds	Theory tests 20	
C.I.I. I.STR. (CIILIIIOI)	Composition and stoichiometry, Amines and	Practical tests 20	
	amides	Examination 60%	
		The CONTINUOUS ASSES	SMENT
COMPUTER APPLICATIONS	Introduction to computing, Hardware, software,	mark shall be made up of	10/
I(CAPPIOI)	communication Microsoft Word, Excel &	Theory tests 20	
` '	PowerPoint (Beginner to intermediate)	Practical tests 70	1%
		Assignment 10%	
	Introduction & Mathematical Concepts,		
	Kinematics in One Dimension, Forces and		
	Newton's Laws of Motion	Theomy Tests 340	,
	Dynamics of Uniform Circular Motion, Work	Theory Tests 26%	-
PHYSICS I (PYSCI05)	and Energy, Rotational Dynamics, Fluids Heat	Practical test 10%	
` ′	and the transfer of heat, Simple Harmonic	Practical book 4% Examination 60%	-
	Motion and Elasticity, Waves and Sound, Electric Circuits	Examination 60%	0
	The Reflection of Light: Mirrors, Lenses and		
	Optical Instruments		

CALCULATIONS & STATISTICS (CSTAI0)	Quadratics, Exponents, Logarithms, Graphs, Equations of a straight line, Conversion of experimental data to linear form, Linear programming, Collection & presentation of data, Sampling techniques, Measures of tendency / dispersion for raw & grouped data, The normal curve	Theory tests Examination	40% 60%
ANATOMY AND PHYSIOLOGY 2 (ANAPH202)	The Nervous System inclusive of the Central & Peripheral Nervous System and Sensory Physiology The Cardiovascular System including Blood Vessels Hemodynamics The Respiratory System including Physical Aspects and Mechanics of Ventilation and Acid-Base Balance The Urinary System inclusive of Urine Production and Renal Control of Electrolyte and Acid-Base Balance The Reproductive System inclusive of the endocrine regulation of both the male and females systems as well as fertilization, pregnancy and parturition	Theory Tests Practical Tests Examination Mark	30% 10% 60%
ORGAN AND SYSTEM PATHOPHYSIOLOGY 2 (OSPP201)	Diseases of Immunity, Fluid and haemodynamic derangements, Nutritional disorders, Systemic diseases, and Infectious diseases Introductory Concepts with reference to the following systems: Respiratory system, Circulatory system, Urinary system, Digestive system, Nervous system and sense organs, Endocrine system, Reproductive system	Theory Tests - 40% Examination Mark	20% 60%
PHARMACOLOGY II (PHAR201)	General Aspects of Drug Therapy, Pharmacokinetics and Pharmacodynamics, Administration of drugs to patients, Adverse effects of drugs, Drugs affecting the autonomic, somatic and sensory nervous system, Drugs affecting the central nervous system, Analgesics and anti-inflammatory drugs, Antihistamines, Hormones and hormone antagonists, Antimicrobial and other anti-infective drugs, Cardiovascular drugs, Drugs affecting the haemopoietic system, Drugs that affect the respiratory system, Drugs that affect the digestive tract, and Poisoning and drug treatment in emergencies	Theory Tests Examination Mark	40% 60%
BIOMEDICAL APPARATUS AND PROCEDURES II (BAPO201)	Introduction to Biomedical Instrumentation Systems Biometrics, Introduction to the Man-Instrument System and Problems Encountered in Measuring a Living System Basic Transducer Principle The Transducer and Transducer Principle, Active Transducers, Passive Transducers and Transducer for Biomedical Applications Electrodes Electrodes Theory, Bio-potential Electrodes, Biochemical Transducers and Blood gas analyser Overview Of Biomedical Instrumentation Systems for the following: Cardiology, Respiratory System, Cardiovascular Perfusion, Neurophysiology, Renal System and Reproductive Biology	Theory tests - 30% Practical tests - 10% Examination - 60%	26% 14% 60%
PSYCHODYNAMICS II	Personality, learning, memory and	Theory tests	24%
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(PYDNI0I)	adjustivebehaviour Basic Principles of human development and the	Assignments 16% Examination 60%
	biological basis of behaviour Attachment theory and psychoanalytic concepts of development Psychological, cognitive and social learning theories of development Psychological, cognitive and social learning theories of development. Emotions, motivation and perception Legal and ethical responsibilities, patient's right charter, Batho Pele principle, National Health Act and Health Professions Act, 1974.	
CARDIOLOGY: BIOMEDICAL APPARATUS 3 (CPA301)	Electrocardiography, Exercise stress testing, Arrhythmia monitoring, Cardiac catheterization, Pacemakers, Echocardiography, Intra-aortic balloon pump, Intra vascular ultrasound system, Defibrillator, Blood gas analyzer, Electrical Safety	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
CARDIOLOGY: CLINICAL PRACTICE 3 (CACP310)	Electrocardiography, Exercise stress testing, Arrhythmia monitoring, Cardiac catheterization, Pacemakers, Echocardiography, Intra-aortic balloon pump, Intra vascular ultrasound system, Defibrillator Blood gas analyzer, Electrical Safety	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
CARDIOLOGY: CLINICAL TECHNOLOGY PRACTICE 3 (CCTP310)	Left and right heart catheterization; Coronary angiography; Percutaneous coronary intervention; Pacemakers Intra-aortic balloon pump; Intravascular ultrasound; Defibrillation; Exercise stress testing; Holter monitoring; Head-up tilt test; Pacemaker check-ups; Programming of pacemakers; Echocardiography;	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20%
CARDIOVASCULAR PERFUSION: BIOMEDICAL APPARATUS 3 (CCBA301)	Embryology of cardiovascular system, Anatomy and physiology of the heart, Anatomy and physiology of the lungs Oxygenators, Gas exchange, Heat exchangers, Blood gas analyser, Arterial and venous cannulae, Coagulation Anatomy and physiology of the kidney, Ultrasonic scanning, Blood pressure monitoring equipments, Pumps Cardiotomy reservoir, Cell saver, Filters, Cardioplegia, Thermoregulators, Ultrafiltration, Electrocardiography Transesophageal echocardiography, Pacemakers, Pulse oximeter	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
CARDIOVASCULAR PERFUSION: CLINICAL PRACTICE 3 (CCC301)	Pulmonary diseases, blood disorders, Coagulation disorders, Effects of oxygenatorsConstrains on rate of heat transfer, Functions of CPB, Renal Failure, Cannulation Blood pressure measurements, Pumps, Heat exchangers, Venting, Ultrafiltration Cardiovascular disorders, Myocardial injuries, Anticoagulation, Electrocardiography Hemodynamic monitoring, Thermoregulation, Cardioplegia, Neurological monitoring Blood gas analyses, Diuretics, benzodiazepine, antiarrhythmics and inotropes	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%

CARDIOVASCULAR PERFUSION: CLINICAL TECHNOLOGY PRACTICE 3 (CTPR301)	Calculation of blood flow rate, selection of bypass circuitry and cannulae Aseptic setting-up of bypass circuitry, priming, and debubbling Calibration and zeroing of pressure transducers and troubleshooting Placement of reliable and rapidly sensing safety devices and monitors Monitoring of urinary output Analysis of blood gas and electrolytes Monitoring of anticoagulation Supervised conduct of cardiopulmonary bypass procedure Monitoring of electrocardiography and hemodynamic parameters	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30% Proficiency based practical tests 80% Process portfolio 20%
CRITICAL CARE: BIOMEDICAL APPARATUS 3 (NEAP301)	Cardiovascular anatomy & physiology Blood Pressure monitoring equipments, Pulse oximeter& co-oximeter, Venous flow measurement Electrocardiography, Cardioversion and defibrillation, Blood flow meters Respiratory system anatomy and physiology, Respiratory therapy equipments Gastrointestinal tract anatomy and physiology History of anaesthesia ,Anaesthetic equipment, Drugs used in anaesthesia Oxygen sensors, Medical gas cylinders and their associated components Thermo-regulatory device, Neurological disorders Hematological measurements including activated clotting time [ACT], Infections	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30% Theory test 70% Assignments 30%
CRITICAL CARE: CLINICAL PRACTICE 3 (NCLI301)	Topics covered: Blood Pressure monitoring equipments, Pulse oximeter& co-oximeter, Venous flow measurement Cardiovascular disorders, Acute renal failure, Electrocardiography, Cardioversion and defibrillation Blood flow meters, Respiratory therapy equipments, Respiratory disorders, GIT disorders, Endocrine disorders History of anaesthesia ,Anaesthetic equipment, Drugs used in anaesthesia Oxygen sensors, Medical gas cylinders and their associated components Thermo-regulatory device, Neurological disorders Hematological measurements including activated clotting time [ACT], Infections	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
CRITICAL CARE: CLINICAL TECHNOLOGY PRACTICE 3 (NCTP301)	Table 12 Lead ECG; Measurement of hemodynamic parameters i.e. BP, Pulse, and RR; Blood gas analysis; Patient care before, during and after the procedure; Thermoregulation Patient transport, oxygen therapy, pulse oximetry and capnography. Prepare anaesthetic and ventilation equipment Effectively assist with bronchoscopy, performance of CPR and during anaesthesia. Intubation and intravenous cannulation. Measure an interpret ACT, glucose, Hct, ESR and SG; Maintenance of the prescribed theatre and ICU equipments. History of Dialysis, Principles of Dialysis,	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20% The CONTINUOUS ASSESSMENT
TELL LINGLOGI. BIOFIEDICAL	r iistor y or Diarysis, i rinciples or Diarysis,	THE CONTINUOUS ASSESSMENT

APPARATUS 3 (NBAMA301)	Sterility and safety, Dialysis Apparatus, Dialysis	mark shall be made up of Theory
	Reprocessing	test 70%
	Water Treatment, Dialysis Facility Design	Assignments 30%
NEPHROLOGY: CLINICAL PRACTICE 3 (NCLP301)	Anatomy & Physiology of the Excretory system Pathophysiology of Renal Disease Blood result analysis & Clinical Invasive and Non-invasive investigation Initiation of Dialysis, Patient observation and Cardio-Pulmonary Resuscitation Anticoagulation, Vascular Access, Peritoneal Dialysis, Hypertension, Diabetis Mellitus Complications during dialysis Drugs used in Dialysis and Transplantation Blood Transfusions and Universal Precautions, Haemoperfusion, Plasmapheresis Continuous Renal Replacement Therapies, Acute and Chronic Dialysis Prescription Nutrition, Pediatric Dialysis	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
NEPHROLOGY: CLINICAL TECHNOLOGY PRACTICE 3 (NTPR301)	Observe patient's vital signs [i.e. heart rate, blood pressure, temperature]; physical appearance of a patient and interpretation of blood results. Apply aseptic techniques and follow safety procedures. Set up disposables / equipment for following procedures:- Chronic Hemodialysis, Acute Hemodialysis, Continuous therapies, Apheresis, Haemoperfusion Paediatric procedures.	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20%
NEUROPHYSIOLOGY: BIOMEDICAL APPARATUS 3 (FBAP301)	Electroencephalography Modes of Operation of an EEG Components: Selection of recording systems, Pre and main amplifiers, Simulators, Electrode Terminals, Ohmeter Types of Electrode, Sensors and Cables, Control Functions effect and Calibrations. Preparation, use and maintenance Electromyography and Nerve Conduction Studies Principle utilised in EMG/ENG Recordings. Modes of Operation of EMG/ENG components: Composition, Accessories, Power supply, Earth; Display and Recording Systems, Control functions, effect and Calibration. Audio Monitor, Signal Delay and Storage unit, Theory of a Strain Gauge Amplifier. Evoked Potential Systems Modes of operation of Evoked Potential Recording systems component: Pre and main Amplifiers, Recording and Display systems, Stimulators, Electrode Terminals Earth (Patient as well as equipment), Control Functions effect and Calibration Averager and other Computer facilities, Memory Storage Facilities, Cursors. TranscranialDopplers Mode of operation, Recording and Display systems, Probes, Hydrocephalus and SAH Polysomnography Instrumentation	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%

NEUROPHYSIOLOGY: CLINICAL PRACTICE 3	Recording Systems, Pre and Main Amplifiers. Electrode Terminals, Earth (Patient as well as equipment) Electrodes, Sensors and Cables, Modules for Recording of Additional Parameters. Epilepsy Monitoring Principles of Epilepsy monitoring; Recording Electroencephalography, Electromyography And Nerve Conduction Studies, Evoked Potential Systems, Transcranial Dopplers,	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70%
(PCTP301)	Polysomnography Instrumentation and Epilepsy Monitoring	Assignments 30%
NEUROPHYSIOLOGY: CLINICAL TECHNOLOGY PRACTICE 3 (PCTP301)	Perform Electroencephalography Perform Nerve Conduction Studies Perform Evoked Potential Testing Perform Trans-cranial Dopplers Assist in Sleep studies and In Long Term Epilepsy Monitoring Perform Polysomnography Practice electrical and laboratory safety	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20%
PULMONOLOGY: BIOMEDICAL APPARATUS 3 (PBAP301)	Anatomy and physiology of the airways Heart and lung circulation Basic lung function equipment Spirometer, Flow measuring devices, Transcutaneous monitoring devices, Gas chromatography Mass spectrometer, Oxygen analysers, Nitrogen analysers, Blood gas analysers, Lung mechanics Pulmonary gas exchange Transport of respiratory gases Control of respiration Systems for the determination of lung function Spirometry and flow-volume systems, Computerised lung function systems, Whole body plethysmograph Diffusion capacity systems, Exercise study equipment, Bronchoscopy	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
PULMONOLOGY: CLINICAL PRACTICE 3 (PCLP301)	Lung injury, Respiratory diseases, Infectious diseases, Immunological disorders, Cardiovascular disorders, Pulmonary function laboratory safety, Pulmonary function measurement, Lung volume evaluation Ventilation tests and artificial ventilation, Basic flow-volume curves, Gas distribution evaluations Diffusion tests, Bronchial provocation, Bronchodilators, Diagnostic bronchoscopy, Allergy investigations	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
PULMONOLOGY: CLINICAL TECHNOLOGY PRACTICE 3 (PTPR301)	Spirometry tests, Plethysmography and a diffusion measurement; Histamine challenge; Pulse oximetry& blood gas analysis; MIP and MEP; Vital signs monitoring; Assist with bronchoscopy.	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20%

REPRODUCTIVE BIOLOGY: BIOMEDICAL APPARATUS 3 (RBAP301)	Applied Embryology, Pituitary and Hypothalamus, Anatomy& Physiology of Male and Female Reproductive Organs & System, Spermatogenesis, Oogenesis, Physiology of Cervical mucus Apparatus for semen analysis, Preparation of media, ART Laboratory Equipment, Aspiration, Identification, Evaluation and Manipulation of Ova, Fertilization and transfer of ova, Embryo transfer and artificial insemination, Cryopreservation of semen, ova, and embryos Reproductive Imaging (Hysterosalphingography) and Contraception	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
REPRODUCTIVE BIOLOGY: CLINICAL PRACTICE 3 (RCPR301	Congenital Anomalies of Male and Female Reproductive tract. Pathophysiology of Male and Female Reproductive organs & Systems Semen analysis, Cervical mucus Examinations, Semen (Spermatosoa) - Cervical mucus-interaction tests Extended antispermatosoa antibody tests in semen, cervical mucus and blood serum Sexual transmitted infections and blood borne viruses in ART Identification, judgement and manipulation of ova, Fertilization and transfer of ova and embryos Cryopreservation of semen, ova and embryos, Embryo scoring for transfer/cryopreservation, Infertility and Persistent Pregnancy Failure, Quality Assurance, Risk management and Laboratory organization, and Patient-Technologist-Relationship	The CONTINUOUS ASSESSMENT mark shall be made up of Theory test 70% Assignments 30%
REPRODUCTIVE BIOLOGY: CLINICAL TECHNOLOGY PRACTICE (RTPR301)	Sterility and Washing Procedures, Sperm counts, Preparation of culture media and dishes, Blood/Serum concentration and processing, Diagnostic semen analyses, Oocyte retrieval: Screening and Grading TSE/MSA/PESA aspiration, Testicular Biopsy processing, Removal of granulosa cells, Fertilization evaluation Embryo transfer in sterile room and at patient, Cryopreservation, Sperm processing for corrective procedures and Insemination procedures	The CONTINUOUS ASSESSMENT mark shall be made up of Proficiency based practical tests 80% Process portfolio 20%

16.2.2 BTECH CLINICAL TECHNOLOGY

Module Name	Learning Content	Assessment
	A. Biostatistics	The CONTINUOUS ASSESSMENT
	Statistics: general introduction, Measures of	mark shall be made up of
	location and dispersion, Ordering of multi-	
	variable data, Probability theory, Probability	Assignments 50%
	distributions, Confidence intervals	
RESEARCH METHODOLOGY	Hypothesis testing, Correlation, The chi-	
(RMDYI0I)	square statistic, Analysis of variance	
()	B. Research Methodology	
	The aim of research, Steps in the research	
	process, Measurements of incidence, Study	
	structures in research, Causality; Risk; Bias;	
	Measurement, The research protocol C. Application	
	The same of	The CONTINUOUS ASSESSMENT
	Foundations of management, Management theory	
PRINCIPLES OF	and perspectives, The complete organisational	test 24%
MANAGEMENT (PRMG10	environment, Social responsibility and ethics, Plan, Organise, Lead & Control, Quality,	Assignments 16%
	productivity and consumer satisfaction	Exams 60%
	,	The CONTINUOUS ASSESSMENT
ADVANCED	Electroencephalography, Polysomnography, Evoked potentials, and	mark shall be made up of Clinical
NEUROPHYSIOLOGIC	Electromyography/neurography	competency – 50%
(ANPT401)	Liecti omyograpnymeur ograpny	Assignment – 10%
(3.11.1.1)		Portfolio 40%
	Micro-manipulation, Cell culturing, Bio-assays,	
	Sperm function tests, Computer assisted sperm	
ADVANCED REPRODUCTIVE	motility, Fluorescence micxroscopy, Electron	
TRECHNOLOGY (ARPT401)	microscopy, Biochemical separation techniques,	
	Sperm quality controls	Portfolio 40%
_	Physiology calculations of flow rates and cannulas,	The CONTINUOUS ASSESSMENT
	Physiological fluids, Effects of temperature changes,	
	Monitoring pre- intra- post, Cardiac drugs —	Clinical competency –
ADVANCED PERFUSION	anaesthetic, Cardioplegia, Perfusion organs, Tissue	50% 10%
TECHNOLOGY (APFT401)	changes, Blood physiology, Pathology of cardio-	Assignment – 10%
	pulmonary bypass on different organs, Flow	Portfolio 40%
	dynamics, Blood conservations, Differential	
	perfusion, and paediatric perfusion	
	A. Specialised Echocardiography	The CONTINUOUS ASSESSMENT
	Current technological advances, Specialised	
	procedures, Doppler estimation of volume	
	flow, Complex congenital defects, Foetal	
	echocardiography, Extensive ventricular	
	assessment, Pericardial disease, Cardiac	
	tumours and masses, Prosthetic heart valves,	
	and Cardiac transplantation B. Mechanisms Of Arrythmogenesis	
	B. Mechanisms Of Arrythmogenesis Disorders of impulse formation, Disorders of	
ADVANCED CARDIAC	impulse conduction, and Combined disorders	
TECHNOLOGY (ACDT401	C. Advanced Electrophysiological Studies	
TECHNOLOGT (ACDT401	Aberrant conduction, Newer approaches in	
	the investigation of sinus-node disorders,	
	Atrioventricular conduction delays and	
	blocks, Investigation of tachycardias,	
	Mechanisms of tachycardias, and Drug studies	
	D. Interventional Management Of	
	Arrythmias	
	E. Cardiac Pharmacology	
	Arrhythmias, Cardiac Failure, and Ischaemic	
	Heart Failure	
ADVANCED RENAL	Anatomy of the Renal System, Functions of the	The CONTINUOUS ASSESSMENT
TECHNOLOGY (ARNT401)	Kidney, The Three Basic Mechanisms Underlying	
		some of made up of

	the Excretory Function Of The Kidney, Renal Processing Of Individual Substances, Water Balance,	50% 10%
	Micturition and Renal Function Tests and Abnormalities.	Assignment – 10% Portfolio 40%
ADVANCED RESPIRATORY TECHNOLOGY (ARST401)	All sections to include detail studies on: Equipment, Techniques and procedures, Patient evaluation Evaluation of results obtained: Exercise Studies - Cardiopulmonary evaluation, Athletes, Metabolic studies Sleep Studies - Sleep Apnoea, Diagnostics, CPAP titrations, other respiratory abnormalities during sleep Advanced Body Plethismographic Studies —RAW, ITGV, IMP's, MEP's, Compliance Control of Ventillation (CO2 Response) Studies Industrial Respiratory Disease Allergies - Skin testing, Bronchial and other provocation techniques, IgE mediated reactions Clinical trials and procedures Bronchoscopic procedures including laser techniques Nebulisation, and pharmacology of nebulised medications Pulmonary related procedures, with diagnostic radiology, cat scanning Ventilation/perfusion studies with radioactive materials	The CONTINUOUS ASSESSMENT mark shall be made up of Clinical competency – 50% 10% Assignment – 10% Portfolio 40%
ADVANCED CRITICAL CARE TECHNOLOGY (ACRT401)		The CONTINUOUS ASSESSMENT mark shall be made up of Clinical competency – 50% 10% Assignment – 10% Portfolio 40%
CLINICAL TECHNOLOGY RESEARCH PROJECT (CLRP101	Preparation and submission of a research thesis	The CONTINUOUS ASSESSMENT mark shall be made up of Thesis 50% Presentation – 30% Poster – 20%

16.2.2. Bachelor of Health Sciences in Clinical Technology (BHCLTI)

Module		Content Assessment plan			
			Assessment plan		
Clinical	to	Introduction and overview of the seven specialist categories in Clinical Technology	Continuous assessmentOral presentations		
Technology		 Role of the Clinical technologist in each category Laboratory techniques (microscopes, incubators, refrigerators and autoclaves Health care system (clinical health governance structure and Health legislative acts & policy). 	(20%) • Reflective journal (20%) • Written theory assessment (60%)		
		5. Organizational structure of the hospital (human resource and sectors) 6. Basic principles of health-care ethics (applied ethics, biomedical ethics, Batho Pele principles) National Health Act, Basic conditions of Employment, Health Professions Act			
Chemistry			THEORY TESTS		

	the state of the s	Total Caranal
	 introduction to chemistry 	Two Tests on General
	• measurements	Inorganic and Physical
	 energy and matter 	Chemistry and Two Tests
	 atoms and elements 	on Organic Chemistry).
	 compounds and their bonds 	PRACTICAL
	 chemical reactions and quantities 	ASSESSMENT_
	 gases 	FINAL EXAM MARK
	 solutions 	$= CM \times 0.4 + EM \times 0.6$
	 acids & bases 	
	nuclear radiation	
	alkanes and cycloalkanes	
	unsaturated hydrocarbons	
	organic compounds with oxygen and	
	sulphur	
	carboxylic acid and esters	
DI : 101	amines and amides	
Physics 101	MECHANICS PROPERTIES OF MATTER	Continuous Assessment
	PROPERTIES OF MATTER	70 % -f -h
		70 % of the average of the 2 Theory Tests
		Theory Tests 30 % of the Practical Mark,
		where
		L. raesiea.
		practical book + 65% practical test]
Dhysias 201	- 4b - b	Continuous Assessment
Physics 201	• thermal physics	Continuous Assessment
	waves & sound	70 % of the average of the 2
	waves & soulid	Theory Tests
	geometrical optics	30 % of the Practical Mark,
	geometrical optics	where
	electricity& magnetism	[Practical Mark = 35%
	electricity & magnetism	practical book + 65%
	radioactivity & radiation	practical test]
	quantum physics	-
Anatomy I	wave properties of particles	Continuous assessment
Anatomy I	Unit I	unit I- theory (20%) and
	o Introduction	practical (15%)
	 Respiratory Anatomy 	practical (13/6)
	 Cardiovascular anatomy 	unit 2- theory (20%) and
	 Genitourinary Anatomy 	practical (15%)
	• Unit 2	p. acaca. (1570)
	 Neuroanatomy 	unit 3- practical (15%) and
	 Head and neck 	assignment (15%)
		3 (,
	• Unit 3	Internally moderated
	 Limbs 	<u> </u>
Physiology I	 Anatomy and physiology are defined. 	Continous Assessement
	• The relationships between anatomy and	Each of the three units will
	physiology are explained.	be assessed as follows:
	UNIT I	 A two hour theory
	 Cells and tissues, 	test at the end of the
	 Integumentary system, 	
	- , ,	

	Muscular system	unit (Minimum of
	Skeletal system	I 20 marks)
	,	One practical test at
	UNIT 2	the end of the course
	Nervous system	
	Endocrine system,	
	Cardiovascular system,	
	Immunity and the Lymphatic system,	
	Blood	
	UNIT 3	
	Respiratory system,	
	Reproductive system	
Pathophysiology I	Basic Immunology: introductory concepts	Semester mark calculations:
1 7 57	Cells of the immune system	- Two written theory
	Innate and adaptive immune responses	assessment (20% each)
	(humoural and cellular)	- Assignments (Essay 15%;
	Antigen-antibody interactions	Presentation 30%)
	Immunological tolerance and memory	- Reflective journaling: (15%) exam=60%; semester mark
	Autoimmunity	= 40%]
	Basic microbiology	10/0]
	 Introduction to Medical microbiology (micobacterium bacilli, streptococcus, 	
	staphylococcus, HI virus)	
	Infection control, medical and surgical	
	asepsis	
	·	
	Communicable disease patient control	
Instrumentation	Introduction to Man-instrumentation	Semester mark calculations:
for Clinical Technology I	systems; O Biometrics	- Two written theory assessment (20% each)
i eciliology i	BiometricsIntroduction to the Man-	- Assignments (Essay 15%;
	Instrument System	Presentation 15%)
	o Problems Encountered in	- Practical assessment (30%)
	Measuring a Living System	- Moderation: Internally
	 Basic physiological parameters; 	moderated.
	2.1. Heart rate / pulse rate	Final manks
	• 2.2. Blood pressure	Final marks: Course mark 40%
	2.3. Stroke volume / Cardiac output	Exam mark 60%
	2.4. Respiratory rate 2.5. Tidely selected (spiratory rate)	
	 2.5. Tidal volume / minute volume Basic Physiological transducers; 	
	Basic Physiological transducers; The Transducer and	
	Transducer Principle	
	Active Transducers	
	 Passive Transducers 	
	Electrodes	
	Electrode theory	
	Biopotential electrodes	
	Biochemical electrodes Modical terminology	
	 Medical terminology Electrical safety. 	
	Liecti icai saiety.	
Second level		

Applied Anatomy and Physiology	Unit 1: The Cardiovascular System Blood & Heart Unit 2: The Respiratory Physiology Functions of the Respiratory System Pulmonary Diseases Unit 3: Nervous system Unit 4: Endocrine System Unit 5: Reproductive systems	Continuous assessment: A two and half hour test at the end of a unit (including theory and applied practical components). Minimum of 150 marks of which a minimum of 10% will comprise the practical component.
Clinical Technology Practice Instrumentation	 Setting-up of equipment: Basic haemodynamic monitoring Basic Electrophysiological procedures: Other basic diagnostic and therapeutic procedures: Spirometry measurement. Anthropometric measurement. Activating clotting time testing. Oral and axillary temperature measurement. Non- provocative nebulisers. Oxygen therapy (mask and nasal cannula). BIOMEDICAL INSTRUMENTATION 	Continuous assessment as follows: Proficiency assessment (60%) Hospital Visit Reports (20%) Presentations (20%)
for Clinical Technology II	SYSTEMS FOR CARDIOLOGY BIOMEDICAL INSTRUMENTATION SYSTEM FOR RESPIRATORY SYSTEM BIOMEDICAL INSTRUMENTATION SYSTEM FOR CRITICAL CARE BIOMEDICAL INSTRUMENTATION FOR CARDIOVASCULAR PERFUSION BIOMEDICAL INSTRUMENTATION SYSTEM FOR NEUROPHYSIOLOGY. BIOMEDICAL INSTRUMENTATION FOR RENAL SYSTEM BIOMEDICAL INSTRUMENTATION FOR RENAL SYSTEM BIOMEDICAL INSTRUMENTATION SYSTEM FOR REPRODUCTIVE BIOLOGY	Semester mark 40%; exam mark 60 %; Semester mark calculations: 3 theory tests (60%) Assignments and presentations (40%)
Clinical Pathophysiology I	 Epidemiology and related medical terminology Overview of Blood disorders Selected Infectious diseases Neoplasia Cardiovascular system Neurological system Respiratory system Pathophysiology of selected disorders of Calcium Metabolism Pathophysiology of selected Hypothalamic and pituitary diseases and overview of Thyroid disease Diabetes Mellitus 	Examination Semester 40%; exam mark 60 % semester mark calculation: 3 written theory tests (60%) 2 x assignments [presentation and written] (40%) Moderation: Internal according to DUT policies

_	T	
	Liver Disease	
	Selected Pancreatic disorders	
	Digestive system and Skin disorders	
	Selected disorders of the Renal system	
	Selected disorders of the male and female	
	Reproductive system	
Basic Pharmacology	This module is divided into 3 Units:	Assessment will be
	UNIT I	continuous.
	General aspects of drug therapy	A two hour theory
	 Pharmacokinetics 	test at the end of each
	 Pharmacodynamics 	unit.
	 Administration of drugs to patients 	 Each theory test will
	Adverse effects of drugs	be weighted as follows
	Autonomic, Somatic and Sensory	_
	Nervous systems	● Theory test I – 30%
		• Theory test 2 – 35%
	UNIT 2	• Theory test 3 – 35%
	 Antimicrobials and other anti-infectives 	
	 Drugs affecting the CNS 	
	 Drugs affecting the CVS 	
	Haemopoetic drugs	
	Analgesics and anti-inflammatories	
	UNIT 3	
	Hormones and Hormone antagonists	
	Antihistamines	
	Respiratory Drugs	
	GIT Drugs	
	Poisoning and emergency drug treatment	
Research	Research Paradigms	Continuous assessment
Methodology I	- The 3 basic research paradigms	Each assessment has a
-	(positivism, interprets and critical theory)	specific weighting i.e. counts
	Research study design (Longitudinal,	a certain % towards the final
	cross-sectional, bi-directional;	mark:
	Quantitative, qualitative, mixed-	Article critique (20%)
	method; reliability, validity and ethics)	• 2 x assignments (80%)
	Research methods and methodology	
	Sampling methods (observations,	
	questionnaire, interviews, surveys, case	
	studies, laboratory experiments)	
	 Data analysis techniques (descriptive 	
	statistics)	
	 Introduction to the review of the 	
	Literature	
	 Referencing styles and plagiarism 	
Research	The steps and stages in the research	Continuous assessment
Methodology II	process.	The final marks:
	The research purpose based on a problem.	• Submission of a
	The literature review	research proposal
	Selecting an appropriate research design	(70%)
	Developing an appropriate sampling plan	• I x assignment (30%)
	for a hypothetical study in terms of	
	feasibility, representativeness and available	Moderation will be
	,, 1	conducted in accordance

		SI DUT I
	resources.	with DUT rules.
	Developing an appropriate data collection	
	plan	
	 Statistical analysis for the data analysis 	
	process.	
	 Ethical issues relating to the conduct of 	
	research	
Health Care	Basic concepts of Healthcare management	Continuous assessment
Management I	(managers and management)	the final mark:
Tranagement 1	`	I written theory test (60%)
	Basic principles of Healthcare management	written theory test (60%)
	(organizational culture, quality	
	management, time management,	I x assignment
	Teamwork)	[presentation and written]
	 Basic Healthcare information systems 	(40%)
	CARDIOLOGY	
Pathophysiology	 Congenital Heart disease 	Continuous assessment
for Cardiology	 Arrhythmias 	The final mark:
	Valvular Heart disease	2 written theory tests (60%)
	 Coronary artery disease 	2 x assignments
	Pericardial disease	[presentation and written]
	Hypertension	(40%)
	Heart Failure	
	Oedema	
	Peripheral vascular disease	
Pharmacology for	 Understand the application for the 	Examination
Cardiology	following therapeutic classes: Anti-	
	arrhythmia therapy, Anti-anginals,	Final mark = 40% course
	Antihypertensives, Diuretic, Pressins,	mark + 60% exam mark
	cardiostimulatories and inhibitors,	
	thrombolytics, vasoconstrictors and	Course mark calculated as
	vasodilators	follows:
	 Understand the pharmacological 	2 written theory tests (60%)
	applications for the following	I x assignment
	cardiovascular disorders:	[presentation and written]
		(40%)
	Angina	(1070)
	Arrhythmia	
	Oedema	
	Heart failure	
	 Systemic and pulmonary hypertension 	
	 Hypotension 	
	Myocardial infarction	
Clinical	Perform the following procedures and explain	Continuous assessment
Technology	the indications, contra-indications, advantages	The final mark:
Practice in	and disadvantages or limitations and	Continuous Proficiency
Cardiology Ia	complications of the following procedures:	Assessment based on the
	Exercise stress testing	application and performance
	Arrhythmia monitoring (Holter)	of the procedures or
	,	techniques as outlined in
	Cardiac cacricterization fere and right freat	module content (80%)
	procedures	
	Intra-aortic balloon pumping	Compilation of a logbook of
	 Single and dual chamber pacing 	procedures (20%)
	 Basic electrophysiology studies 	procedures (20%)
	 Echocardiography 	

Clinical Technology Practice in Cardiology Ib	Describe the haemodynamics related to angiography and echocardiography for the following conditions: • pericardial disease • Congestive heart failure • Coronary artery disease • Valvular heart disease • Cangenital heart disease • Cardiac resynchronization therapy Describe the underlying pathophysiology of symptom production in the conditions in (2) above.	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
	Infection control Cardio-version. Defibrillation. General equipment management. Assist with ICU/Trauma/Theatre clinical procedures. • Physiological data management.	
Instrumentations and Techniques for Clinical Technology in Cardiology I	Electrocardiography Telemetry Basic terminology relating to Biomedical instrumentation and transduction Instrumentation used and procedures for arrhythmia monitoring or termination(non-invasive): Exercise stress testing laboratory equipment Holter Internal and external defibrillation	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Techniques for Clinical Technology in Cardiology Ib	 Invasive monitoring and diagnostic instrumentation and procedures: Monitoring and blood gas equipment in the cardiac catheterization laboratory Catheters used and procedures in the cardiac catheterization laboratory on adult patients (diagnostic angiography and intervention, cardiac output, IVUS, IABP, pericardiocentesis, electrophysiology and pacing) Resonance and damping; Cardiac output measurements Blood gas machine Coagulation instrumentation; Equipment bench testing, diagnostics and quality control; Simulators; Left ventricular assist devices 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Pathophysiology for Critical Care	Myocardial infarction;Heart failure (left & right);	Continuous assessment The final mark: 2 written theory tests (60%)
	 Compensatory mechanisms for a falling CO; Shock; Abdominal compartment syndrome; 	2 x assignments [presentation and written] (40%)

		1 : f-:1	
	•	Liver failure;	
	•	Pancreatic failure;	
	•	Coagulopathies, DIC;	
	•	Endocrine disorders;	
	•	COPD, Asthma, Pneumonia and	
		Aspiration;	
	•	Pulmonary embolism,	
		pneumothorax;	
	•	Respiratory failure;	
	•	Gaseous exchange abnormalities;	
	•	ARDS;	
	•	Neurological assessment for altered	
		levels of consciousness	
Dhawaa aala fan			Formation at an
Pharmacology for	•	Understand the application for the	Examination
Critical Care		following:	Final mark = 40% course
	•	Drugs used in Hypertension and	mark + 60% exam mark
		Angina	mark + 60% exam mark
	•	Drugs used in Heart failure.	Course mark calculated as
	•	Resuscitation drugs	follows:
	•	Local Anaesthetics, Anesthetic	2 written theory tests (60%)
		agents (Inhalational and	I x assignment
		intravenous),	[presentation and written]
	•	Drugs acting at Neuromuscular	(40%)
		Junction and Autonomic Nervous	(1070)
		System.	
	•	Antibiotics, Antimicrobial,	
	•	Antifungal and Antiviral Drugs.	
	•	Understand the pharmacological	
		applications for the following	
		disorders:	
	•	Myocardial infarction;	
	•	Heart failure (left & right);	
	•	Compensatory mechanisms for a	
		falling CO;	
	•	Shock;	
	•	Abdominal compartment syndrome;	
	•	Liver failure;	
	•	Pancreatic failure;	
	•	Coagulopathies, DIC;	
	•	Endocrine disorders;	
	•	COPD, Asthma, Pneumonia and	
		Aspiration;	
	•	Pulmonary embolism,	
		pneumothorax;	
	•	Respiratory failure;	
	•	Gaseous exchange abnormalities;	
	•	ARDS;	
Clinical	• Ir	fection control	Continuous assessment
Technology	• (Quality Control of life Support equipment.	The final mark:
Practice in	• S	catistical analysis and patient scoring.	Continuous Proficiency
Critical Care la			Assessment based on the

	 Blood gas sampling, measurement and interpretation Invasive heamodynamic monitoring procedures. Set up equipment for Intra-hospital transportation of critically ill patients, non-invasive heamodynamic monitoring, monitoring of an anesthetized patient. Preparation of ICU drugs. Handling of Infusion devices and drugs. Capnography. 	application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Clinical Technology Practice in Critical Care Ib	 Assists with bronchoscopy and right heart catheterization. Advanced Cardiac Life Support (ACLS). CPR. Intubation, intravenous cannulation, emergency drug therapy. Ventilation therapy: monitoring and resuscitation. Determine blood flow (Doppler). Cardio-version. Defibrillation. Electrolyte determination. General equipment management. Assist with ICU/Trauma/Theatre clinical procedures. Physiological data management. 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Instrumentations and Techniques for Clinical Technology in Critical Care la	Electrocardiography Telemetry Invasive pressure monitoring equipment; Resonance and damping; Cardiac output measurements Blood gas machine Ventilators and ventilation modes Anesthetic machine and accessories Hemofiltration Thermoregulatory devices Coagulation instrumentation; Arterio- venous flow measurements Infusion devices Gas and vapour analysers Transcutanous gas measurements Autologous cell recovery Thromboelastograms Point of care analysers (Glucose, Hb, Bilirubin)	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Techniques for Clinical Technology in Critical Care Ib	 Endoscopes; Equipment bench testing, diagnostics and quality control; Simulators; Left ventricular assist devices Therapeutic gas delivery systems 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)

	Peripheral nerve stimulators;	T
	reliplieral herve sumulators;	
	Level of consciousness monitors	
	NEUROPHYSIOLOGY	
Dath anhysiology		Continuous assessment
Pathophysiology for	Abnormalities of Consciousness Abnormalities of the brain	The final mark:
Neurophysiology	Epilepsy	2 written theory tests (60%)
· · · · · · · · · · · · · · · · · · ·	Stroke	2 x assignments
	Dementia	[presentation and written]
	Parkinson	(40%)
	Multiple Sclerosis	
	Encephalopathies	
	Meningitis	
	Headaches	
	Hydrocephalus	
	Haemorrhage	
	Aneurysm	
	• Coma	
	Brain death	
	 Abnormalities of Hearing and Vision 	
	Myasthenia gravis	
	Peripheral nerve disorders	
	Entrapment neuropathies	
	Guillain Barre syndrome/CIDP	
	Diabetic and HIV neuropathy	
	Brachial plexopathies	
	 Drug related neuropathies 	
	Critical illness neuropathy	
	 Abnormalities of sleep 	
	General neurological abnormalities	
Dhawaa aa la sa fa a		Examination
Pharmacology for Neurophysiology	 Understand the pharmacological application for the following: 	Examination
Treat opinysiology	Neurotransmitters	Final mark = 40% course
	Blood-brain barrier	mark + 60% exam mark
	Cholinergic pharmacology	
	Adrenergic Pharmacology	Course mark calculated as
	Local anaesthetic pharmacology	follows:
	Understand the pharmacological	2 written theory tests (60%) I x assignment
	applications for the following disorders:	[presentation and written]
	Abnormalities of consciousness	(40%)
	Abnormalities of Hearing and Vision	
	Myasthenia gravis	
	Peripheral nerve disorders	
	Abnormalities of sleep General pour place of sheep	
	General neurological abnormalities	

Clinical Technology Practice in Neurophysiology la	 Brain mapping Assist in Electromyography Nerve conduction studies 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%) Continuous assessment
Technology Practice in Neurophysiology Ib	 Polysomnography Long-term epilepsy monitoring video studies Memory testing and WADA testing 	The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Instrumentation and Techniques for Clinical Technology in Neurophysiology la	 ELECTROENCEPHALOGRAPHY ELECTROMYOGRAPHY AND NERVE CONDUCTION STUDIES Principle utilised in EMG/ENG Recordings. MEDICAL TERMINOLOGY ELECTRICAL SAFETY 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentation and Techniques for Clinical Technology in Neurophysiology Ib	EVOKED POTENTIAL SYSTEMS TRANSCRANIAL DOPPLERs POLYSOMNOGRAPHY INSTRUMENTATION	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Pathophysiology for Nephrology	Nephrology Clinical Manifestations of Renal Diseases Major Clinical Renal Syndromes (renal failure, tubular defects, urinary tract infections, calculi) Diagnosis of Renal Disease (biopsy, microscopy) Congenital abnormalities of the kidney Glomerular disease Nephrotic syndrome Diabetes mellitus Renal hypertension Anaemia	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Pharmacology for Nephrology	 Understand the application for the following: Drug dosing methods and influencing factors 	Examination Final mark = 40% course mark + 60% exam mark

	Anti hyportansiyas	
	 Anti-hypertensives ACE-Inhibitors, Angiotensin-receptor blockers, Diuretics Beta Adrenergic Blocking Drugs Calcium Channel Blockers Dyslipidaemia management Anaemia management Understand the pharmacological applications for the following disorders: Major Clinical Renal Syndromes (renal failure, tubular defects, urinary tract infections, calculi) Diagnosis of Renal Disease (biopsy, microscopy) Congenital abnormalities of the kidney Glomerular disease Nephrotic syndrome Diabetes mellitus Renal hypertension 	Course mark calculated as follows: 2 written theory tests (60%) 1 x assignment [presentation and written] (40%)
Clinical	Handwashing technique and infection	Continuous assessment
Technology Practice in Nephrology Ia	 Handwashing technique and infection control; Setting up of equipments for HD and PD therapies; Organise equipments for emergencies; Priming and disinfection; Preparation of access sites (PD & HD); Subcutanous administration; Intravenous administration; Water sampling testing; Preassement of patient Monitoring of hemodynamics of HD and PD; Phlebotomy; Commencement and discontinuation techniques of HD and PD. Post hemodynamic monitoring of HD and PD 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Clinical Technology Practice in Nephrology Ib	 Post hemodynamic monitoring of HD and PD Cannulation using sterile techniques of arteriovenous fistula; Sterile techniques for connection of catheters; Perform chronic hemodialysis therapy; Perform chronic peritoneal dialysis therapy; Hemodynamic monitoring of both above procedures; Management of acute complications during HD and PD; Management of chronic complications of HD and PD; Setting up of equipments for acute HD/PD and CRRT; Hemodynamic monitoring acute HD/PD. 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)

Instrumentation and Techniques for Clinical Technology in Nephrology la	 Development of dialysis equipment Theory of haemo-dialysis and PD. Method of solute transport and ultrafiltration. Types Dialyzers Blood and dialysate compartments Monitoring devices Calibration, servicing and disinfection of equipments Design, operation and SOP of Hemodialysis equipments; Design, operation and SOP of Peritoneal equipments 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentation and Techniques for Clinical Technology in Nephrology Ib	 Optimization of dialysis with regards to acute- and chronic dialysis therapy. Dialysate used in haemodialysis, peritoneal dialysis and continuous therapies. Water treatment for haemodialysis Emergency equipment; General and health and safety in the renal unit. Design, operation and SOP of acute dialysis and CRRT equipments; Blood gas analysis 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Pathophysiology	PERFUSION • Ischemic Heart Disease	Continuous assessment
	Ischemic Heart Disease	Continuous assessment
for Perfusion Pharmacology for	 Myocardial Infarction Valvular Heart Disease (Acquired and Congenital), Congestive Heart Failure Diseases of the Great Arteries (Dissection, Aneurysm, Pulmonary Embolism) Pulmonary Hypertension Bacterial Endicarditis and Rheumatic Fever Cardiomyopathy and Heart & Lung Transplant Congenital Heart Disease. Understand the application for the 	The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)

	Understand the pharmacological applications for the following cardiovascular disorders: Angina Arrhythmia Oedema Heart failure Systemic and pulmonary hypertension Hypotension Myocardial infarction	
Clinical Technology Practice in Perfusion la	Assessing the Physiological Health of Patient; Use Various Cardioulmonary Components; Electrocardiography (ECG) Measurement; Perform Advanced Cardiac Life Support; Spirometry Measurement, Anthropometric Measurement; Anticoagulation Testing (ACT), Blood Pressure Measurement, Temperature Monitoring, Pulse Measurement; Perform Bloodgas Analysis; Oximetry Measurement; Blenders, Vaporizers, Perform Capnography; Use of Non-provocative Nebulizers; Administer Oxygen Therapy, Calibrate the Transducers; Use of Infusion Devices; Perform Phlebotomy; Utilize Intra-Aortic Balloon Pumps; Perform Autologous Blood Salvage; Monitor Haemodynamic Parameters; Operate Flowmeters;	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Clinical Technology Practice in Perfusion Ib	 Perform Cardiopulmonary Resuscitation (CPR); Utilize the Left Ventricular Assist Devices (LVAD); Administer Drugs; Perform Basic Echocardiography (ECHO); Perform Vascular Sonography; Interpretation and Analysis of Diagnostic Data; Perform External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG) Measurement, Perform Stress Test, Monitor the Basic Electroencephalography (EEG); Application of Defibrillator and Cardioversion; 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)

1	1	1	
	•	Integrate Hemodialyzer;	
	•	Interpret Magnetic Resonance Imaging	
		(MRI);	
	•	Perform Extracorporeal Membrane	
		Oxygenation (ECMO);	
Instrumentations	•	Electrocardiography (ECG);	Continuous assessment
and Techniques	•	Advanced Cardiac Life Support;	The final mark:
for Clinical	•	Measurement of Spirometry,	2 written theory tests (60%)
Technology in	•	Anthropometric,	2 x assignments
Perfusion Ia	•	Anti Coagulation Testing (ACT),	[presentation and written]
	•	Blood Pressure,	(40%)
	•	Temperature, Pulse;	
	•	Bloodgas Analysis;	
	•	Blenders; Vaporizers;	
		Oximetry;	
		Capnography;	
		Non-provocative Nebulizers;	
		Oxygen Therapy,	
		Calibration of Transducers;	
	_	ŕ	
	•	Ventilators;	
	•	Infusion Devices,	
	•	Phlebotomy,	
Instrumentations	•	Intra-Aortic Balloon Pumps;	Continuous assessment
and Techniques	•	Autologous Blood Salvage;	The final mark:
for Clinical	•	Cardiovascular Monitoring;	2 written theory tests (60%)
Technology in	•	Cardiopulmonary Resuscitation (CPR);	2 x assignments
Perfusion Ib	•	Left Ventricular Assist Devices (LVAD);	[presentation and written]
1	_		(400/)
	•	Drug Administration, Echocardiography	(40%)
		Drug Administration, Echocardiography (ECHO);	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography;	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP),	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG),	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test,	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG);	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators,	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters,	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers,	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers;	(40%)
	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters;	(40%)
Pathophysiology	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY	
Pathophysiology for Pulmonology	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury	Continuous assessment
Pathophysiology for Pulmonology	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases	Continuous assessment The final mark:
. , .,	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases	Continuous assessment The final mark: 2 written theory tests (60%)
. , .,		Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Immunological disorders	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments
. , .,	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases	Continuous assessment The final mark: 2 written theory tests (60%)
for Pulmonology		Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Immunological disorders Cardiovascular disorders	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written]
. , .,		Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Immunological disorders Cardiovascular disorders	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
for Pulmonology Pharmacology for		Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Immunological disorders Cardiovascular disorders	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
for Pulmonology Pharmacology for	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Inmunological disorders Cardiovascular disorders Understand the pharmacological application for the following classes: Pressins	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%) Examination
for Pulmonology Pharmacology for	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Inmunological disorders Cardiovascular disorders Understand the pharmacological application for the following classes: Pressins cardiostimulatories and inhibitors	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%) Examination Final mark = 40% course
for Pulmonology Pharmacology for	•	Drug Administration, Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters; PULMONOLOGY Lung injury Respiratory diseases Infectious diseases Inmunological disorders Cardiovascular disorders Understand the pharmacological application for the following classes: Pressins	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%) Examination Final mark = 40% course

	Understand the pharmacological applications for the following disorders:	follows: 2 written theory tests (60%) 1 x assignment [presentation and written] (40%)
Clinical Technology Practice in Pulmonology IA	 Pulmonary function laboratory safety Pulmonary function measurement Lung volume evaluation Ventilation tests and artificial ventilation Basic flow-volume curves Gas distribution evaluations 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of
Clinical Technology Practice in Pulmonology IB	 Diffusion tests Bronchial provocation Bronchodilators Diagnostic bronchoscopy Allergy investigations 	procedures (20%) Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Instrumentations and Procedures for Clinical Technology in Pulmonology Ia	Basic lung function equipment i. Spirometer ii. Flow measuring devices iii. Transcutaneous monitoring devices iv. Gas chromatography v. Mass spectrometer vi. Oxygen analysers vii. Nitrogen analysers viii. Blood gas analysers ix. Lung mechanics	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Procedures for Clinical Technology in Pulmonology Ib	Systems for the determination of lung function i. Spirometry and flow-volume systems ii. Computerised lung function systems iii. Whole body plethysmograph iv. Diffusion capacity systems v. Exercise study equipment Bronchoscopy	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)

	REPRODUCTIVE BIOLOGY	
Pathophysiology	Congenital Anomalies of Male and	Continuous assessment
for Reproductive	Female Reproductive tract.	The final mark:
Biology	 Pathophysiology of Male and Female 	2 written theory tests (60%)
	Reproductive organs & Systems	2 x assignments
	 Infertility and Persistent Pregnancy 	[presentation and written]
	Failure	(40%)
	 Microbiology 	
	 Ectopic pregnancy , placenta previa , 	
	sacrococcygeal teratoma	
	• Genetic disorders (eg Klinefelter	
	syndrome, Turner's syndrome,	
Dhawaa aa la sa fa a	Down's syndrome)	Examination
Pharmacology for Reproductive	 Understand the pharmacological application for the following classes: 	Examination
Biology	Ovulation induction drugs	Final mark = 40% course
Biology	Contraception	mark + 60% exam mark
	Understand the pharmacological	
	applications for the following disorders:	Course mark calculated as
	 Congenital Anomalies of Male and Female 	follows:
	Reproductive tract.	2 written theory tests (60%)
	 Infertility and Persistent Pregnancy Failure 	I x assignment
	Microbiology	[presentation and written]
	 Ectopic pregnancy , placenta previa , 	(40%)
	sacrococcygeal teratoma	
	 Genetic disorders (eg Klinefelter 	
	syndrome, Turner's syndrome, Down's	
	syndrome)Cardiovascular disorders	
Clinical	Fundamentals of Clinical Embryology	Continuous assessment
Technology	 Introduction to In Vitro Fertilisation 	The final mark:
Practice in	and Embryo Culture	Continuous Proficiency
Reproductive	 Congenital Anomalies of Male and 	Assessment based on the
Biology Ia	Female Reproductive tract.	application and performance
	 Pathophysiology of Male and Female 	of the procedures or techniques as outlined in
	Reproductive organs & Systems	module content (80%)
	Semen analysis	module content (60%)
	Cervical mucus Examinations	Compilation of a logbook of
	 Semen (Spermatosoa) - Cervical mucus-interaction tests 	procedures (20%)
		. , ,
	 Extended antispermatosoa antibody tests in semen, cervical mucus and 	
	blood serum	
Clinical	 Sexual transmitted infections and blood 	Continuous assessment
Technology	borne viruses in ART	The final mark:
Practice in	o Identification, judgement and manipulation	Continuous Proficiency
Reproductive	of ova.	Assessment based on the
Biology Ib	 Fertilization of ova and embryos 	application and performance
	o Cryopreservation of semen, ova and	of the procedures or
	embryos	techniques as outlined in
	o Infertility and Persistent Pregnancy Failure	module content (80%)
	(a). Fertility Preservation in	Commission of the best for
	Cancer Patients	Compilation of a logbook of
	(b). Infections and Infertility	procedures (20%)

	(c). Male and Female	
Instrumentations and Techniques for Clinical Technology in Reproductive Biology la	(c). Male and Female Infertility (d). Artificial Insemination (e). Induction of Ovulation Ouality Assurance, Risk management and Laboratory organisation Patient-Technologist-Relationship Apparatus for the following procedures: Semen analysis Preparation of media ART Laboratory Equipment Maintenance of Apparatus Quality control	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Techniques for Clinical Technology in Reproductive Biology Ib	 Reproductive Imaging (Hysterosalphingography, Laparoscopy) Contraception Hormonal Contraception Modern Concepts in Intrauterine Devices Surgical Sterilization 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
	Fourth level	
Health Care Management II	 Legal and social aspects of Healthcare Human resource management in healthcare settings Budgeting and financial management in Healthcare Leadership in Healthcare settings Community relations in Healthcare settings 	Continuous assessment The final mark: 2 x written theory tests (60%) 1 x assignment [presentation and written] (40%)
Research Methodology III	 Conduct a research project and collect data using appropriate research methodology. Perform data analysis using appropriate statistical tests and packages. Interpret findings and present these according to set criteria and formatting requirements in the form of a dissertation. Demonstrate an ability to act professionally and ethically when conducting research 	Continuous assessment The final mark: Research project =70% Presentation of research = 30% Externally moderated
Clinical Instruction (Elective I)	 Learning Process and Models of Instruction Teaching and Learning Styles Teaching, Learning, Assessment, and Study Skills Strategies Curriculum Development and Classroom Management Academic Writing and Presentation Mentorship 	Continuous assessment with external moderation : Theory tests (60%) Assignments (40%)

Small business management (Elective 2)	 Introduction to Entrepreneurship Theory Self-awareness and development of personal attributes Industry and business classification Business Plan development Marketing for Entrepreneurs Finance, business calculations and financial record keeping for Entrepreneurs Operations Management for Entrepreneurs Human Resources for Entrepreneurs Presentation Skills 	Continuous assessment with external moderation: - Theory Tests – Open or closed Book 70% - Individual Participation/Graduate Attributes 10% Business Plan (group work) 20%
	CARDIOLOGY	
Clinical Technology Practice in Cardiology IIa	Setting up and monitoring of the following invasive procedures: Intra-aortic balloon pumping Intravascular ultrasound and fractional flow reserve Right and left heart catheterisation on paediatrics Electrophysiology and ablation Bi-ventricular pacing Implantable cardiac defibrillators Setting up and monitoring of the following invasive procedures:	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
Clinical Technology Practice in Cardiology IIb Instrumentations and Techniques for Clinical Technology in Cardiology IIa	Head-up tilt testing External synchronised cardiac defibrillation Advanced cardiopulmonary resuscitation Perform echocardiography and correctly report on the following: adult and paediatric congenital heart disease valvular heart disease Infective endocarditis Pericardial disease Dobutamine stress echocardiography Intra-Aortic Balloon Pump. Intra-aortic balloon pump Intravascular ultrasound and fractional flow reserve equipment Right and left heart catheterisation on paediatrics: wires, catheters	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%) Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Techniques for Clinical Technology in	 Electrophysiology and ablation equipment and catheters Bi-ventricular pacing: leads, wires and generators Implantable cardiac defibrillators: leads, wires, defibrillator 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments

Condialogy III	F. I. B. J.	[succentation and vunitary]
Cardiology IIb	Echocardiography: transoesophageal	[presentation and written]
	echocardiography and	(40%)
	Dobutamine stress	
	echocardiography;	
	pericardiocentesis	
	Drug Administration and	
	management of side effects.	
	CRITICAL CARE	
Clinical	 Intubation. 	Continuous assessment
Technology	 Assist with acute haemodialysis and 	The final mark:
Practice in	continuous renal replacement	Continuous Proficiency
Critical Care IIa	therapy (CRRT).	Assessment based on the
	 Autologous blood recovery. 	application and performance
	 Cell saving. 	of the procedures or
	 Monitor Intra-Aortic Balloon Pump 	techniques as outlined in
	 Metabolic studies. 	module content (80%)
	 Left ventricle assist therapy. 	
	 Coagulation studies. 	Compilation of a logbook of
	Endoscopy.	procedures (20%)
Clinical	Ultrasonography.	Continuous assessment
Technology	Drug Administration and	The final mark:
Practice in	management of side effects.	Continuous Proficiency
Critical Care IIb	Advanced patient transport (inter-	Assessment based on the
	hospital and international transport).	application and performance
	General equipment management.	of the procedures or
	Physiological data management.	techniques as outlined in
	Neonatal:	module content (80%)
	Set up, apply and maintain the following	, ,
	equipment:	Compilation of a logbook of
	• Incubators;	procedures (20%)
	Humidifiers;	
	Phototherapy;	
	1	
	 Neonatal therapeutic gas administration; 	
	Respiratory support devices.	
	Invasive and non-invasive	
	monitoring	
Instrumentations	Intra-Aortic Balloon Pump.	Continuous assessment
and Techniques	haemodialysis machine	The final mark:
for Clinical	,	2 written theory tests (60%)
Technology in	 Continuous renal replacement therapy equipments (CRRT). 	2 x assignments
Critical care IIa	Autologous blood recovery.	[presentation and written]
	,	(40%)
	Cell saving. Literary pages by	
	Ultrasonography.	
	Neonatal: Incubators; Humidifiers And Physical Action 1988 The state of the	
	and Phototherapy;	
	Acute renal failure;	
	Chronic renal failure;	
	Hepatic failure;	
	Gullian-Barre syndrome, status	
	epilepticus, meningitis, and	
	myasthenia gravis;	

	- During Law 132 - 13	
	Brain herniation, intracranial prossure changes:	
	pressure changes; Drug Administration and	
	management of side effects.	
Instrumentations	Intra-Aortic Balloon Pump.	Continuous assessment
and Techniques	haemodialysis machine	The final mark:
for Clinical	Continuous renal replacement	2 written theory tests (60%)
Technology in	therapy equipments (CRRT).	2 x assignments
Critical care IIb	 Autologous blood recovery. 	[presentation and written]
	 Cell saving. 	(40%)
	 Ultrasonography. 	
	Neonatal: Incubators; Humidifiers	
	and Phototherapy;	
	Acute renal failure;	
	Chronic renal failure;	
	Hepatic failure; Cullian Pages augustus attatus	
	 Gullian-Barre syndrome, status epilepticus, meningitis, and 	
	myasthenia gravis;	
	Brain herniation, intracranial	
	pressure changes;	
	Drug Administration and	
	management of side effects.	
	NEUROPHYSIOLOGY	
Clinical	Paediatric electroencephalography	Continuous assessment
Technology	(EEG)	The final mark:
Practice in Neurophysiology	The electroencephalogram in the	Continuous Proficiency Assessment based on the
lia	unconscious patient in the intensive care	application and performance
	 Sleep and long term 	of the procedures or
	electroencephalography	techniques as outlined in
	Multiple sleep latency testing	module content (80%)
		6 10 611 16
		Compilation of a logbook of procedures (20%)
Clinical	Intra-operative monitoring	Continuous assessment
Technology	Trans-cranial Doppler's	The final mark:
Practice in	Sub-dural monitoring	Continuous Proficiency
Neurophysiology	Drug administration and	Assessment based on the
lib	management of side-effects	application and performance
	Ü	of the procedures or
		techniques as outlined in
		module content (80%)
		Compilation of a logbook of
		procedures (20%)
Instrumentation	Calibration procedures on	Continuous assessment
and Techniques	neurophysiological equipment	The final mark:
for Clinical	• Design, operation and trouble-shooting	2 written theory tests (60%)
Technology in	skills on the equipment for the following	2 x assignments
Neurophysiology IIa	procedures:	[presentation and written] (40%)
114	Paediatric electroencephalography (EEG) The selectroencephalography (EEG) The selectroencephalography (EEG)	(70/0)
	The electroencephalogram in the unconscious patient in the intensive care.	
1	unconscious patient in the intensive care	

	a Cloop and lane town	
	Sleep and long term electroencephalography	
	electroencephalography Multiple sleep latency testing	
	Finduple sleep latency testing	
Instrumentation	Intra-operative monitoring	Continuous assessment
and Techniques	Sub-dural monitoring	The final mark:
for Clinical	Selection of clinical instrumentation and	2 written theory tests (60%)
Technology in	stock control	2 x assignments
Neurophysiology		[presentation and written]
lib	NIEDI IDAI	(40%)
Clinian	NEPHROLOGY	6 .:
Clinical Technology	Acute Hemodialysis;	Continuous assessment The final mark:
Practice in	Acute peritoneal dialysis;	Continuous Proficiency
Nephrology IIa	Paediatric dialysis; Management of transplant actions (and	Assessment based on the
repinology na	 Management of transplant patients (pre and post); 	application and performance
	CRRT therapies:	of the procedures or
	 Plasma exchange; 	techniques as outlined in
	CVVHD;	module content (80%)
	Hemoperfusion	
	'	Compilation of a logbook of
Clinian	CDDT	procedures (20%)
Clinical	CRRT therapies:	Continuous assessment The final mark:
Technology Practice in	CVVH;CAVVH:	Continuous Proficiency
Nephrology lib	CAVVH;SCUF, CVVHD, CVVHDF	Assessment based on the
rtepin ology no	0 3601, 644110, 6441101	application and performance
	Cell saver;	of the procedures or
	,	techniques as outlined in
		module content (80%)
		Compilation of a logbook of procedures (20%)
Instrumentation	Equipments for Acute Hemodialysis;	Continuous assessment
and Techniques	Acute peritoneal dialysis;	The final mark:
for Clinical	 Paediatric dialysis; 	2 written theory tests (60%)
Technology in	Management of transplant patients (pre	2 x assignments
Nephrology IIa	and post);	[presentation and written]
	Equipments for CRRT therapies:	(40%)
	 Plasma exchange; 	
	o CVVHD;	
	 Hemoperfusion 	
Instrumentation	F (CDDT II	Continuous assessment
Instrumentation and Techniques	Equipments for CRRT therapies: (20/4)	Continuous assessment The final mark:
for Clinical	CVVH;CAVVH;	2 written theory tests (60%)
Technology in	SCUF, CVVHD, CVVHDF	2 x assignments
Nephrology IIb	0	[presentation and written]
,	Cell saver;	(40%)
	PERFUSION	
Clinical	Assessing the Physiological Health of Patient;	Continuous assessment
Technology	Use Various Cardioulmonary Components;	The final mark:
Practice in	Electrocardiography (ECG) Measurement;	Continuous Proficiency
Perfusion IIa	Perform Advanced Cardiac Life Support;	Assessment based on the
	Spirometry Measurement, Anthropometric	application and performance

	Measurement; Anticoagulation Testing (ACT), Blood Pressure Measurement, Temperature Monitoring, Pulse Measurement; Perform Bloodgas Analysis; Oximetry Measurement; Blenders, Vaporizers, Perform Capnography; Use of Non-provocative Nebulizers; Administer Oxygen Therapy, Calibrate the Transducers; Use of Ventilators; Use of Infusion Devices; Perform Phlebotomy; Utilize Intra-Aortic Balloon Pumps; Perform Autologous Blood Salvage; Monitor Haemodynamic Parameters; Operate Flowmeters; Perform Cardiopulmonary Resuscitation (CPR); Utilize the Left Ventricular Assist Devices (LVAD);	of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of procedures (20%)
	Administer Drugs	
Clinical Technology Practice in Perfusion lib	Perform Basic Echocardiography (ECHO); Perform Vascular Sonography; Interpretation and Analysis of Diagnostic Data; Perform External Counterpulsation (ECP), 3- Dimensional Cardiography (3DVG) Measurement, Perform Stress Test, Monitor the Basic Electroencephalography (EEG);	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in
Instrumentations	Application of Defibrillator and Cardioversion; Integrate Hemodialyzer; Interpret Magnetic Resonance Imaging (MRI); Perform Extracorporeal Membrane Oxygenation (ECMO)	module content (80%) Compilation of a logbook of procedures (20%)
Instrumentations and Techniques for Clinical Technology in Perfusion II	12 Lead Electrocardiography (ECG); Advanced Cardiac Life Support; Lung Dynamics and Measurement, Ventilation/Perfusion Monitoring, Haemodynamic Monitoring, Blood Gas Analysis; Blenders; Vaporizers; Capnography; Provocative Nebulizers; Ventilators; Infusion Devices, Phlebotomy, Intra-Aortic Balloon Pumps; Autologous Blood Salvage; Cardiovascular Monitoring; Cardiopulmonary Resuscitation (CPR); Left Ventricular Assist Devices (LVAD); Drug Administration,	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Techniques for Clinical Technology in Perfusion II	Echocardiography (ECHO); Vascular Sonography; Interpretation and Analysis of Diagnostic Data. External Counterpulsation (ECP), 3-Dimensional Cardiography (3DVG), Stress Test, Basic Electroencephalography (EEG); Defibrillators, Cardioverters, Transducers, Cell Savers; Flowmeters;	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
	PULMONOLOGY	
Clinical Technology Practice in Pulmonology IIa	Assessing the Physiological Health of Patient; Use Various Cardioulmonary Components; Electrocardiography (ECG) Measurement; Perform Advanced Cardiac Life Support; Anthropometric Measurement; Anticoagulation Testing (ACT), Blood Pressure Measurement, Oximetry Measurement; Blenders, Vaporizers, Perform Capnography;	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of

	Use of Non-provocative Nebulizers; Administer Oxygen Therapy, Calibrate the Transducers;	procedures (20%)
Clinical Technology Practice in Pulmonology IIb	 CEPT (cardio pulmonary exercise testing) Skin allergy investigations using skin prick tests Provocation tests Sleep studies Nitric oxide testing) 	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%)
Instrumentations and Procedures for Clinical Technology in Pulmonology IIa	Exercise study equipment Sleep study equipment	Compilation of a logbook of procedures (20%) Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Instrumentations and Procedures for Clinical Technology in Pulmonology IIb	 Provocation testing equipment Nitric oxide machine (NiOx) 	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written] (40%)
Clinical Technology Practice in Reproductive Biology lia	REPRODUCTIVE BIOLOGY Embryo scoring for transfer/cryopreservation IVF and Embryo Culture Micromanipulation Cryobiology and Cryopreservation	Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%)
Clinical Technology Practice in Reproductive Biology lib	Quality Assurance, Risk management and Laboratory organisation Pre-implantation genetic disease Fluorescence in-situ hybridization Ethics and Law for Embryologists	Compilation of a logbook of procedures (20%) Continuous assessment The final mark: Continuous Proficiency Assessment based on the application and performance of the procedures or techniques as outlined in module content (80%) Compilation of a logbook of
Instrumentations and Techniques for Clinical Technology in Reproductive	Equipment/APPARATUS for the following procedures: Aspiration, Identification, Evaluation and Manipulation of Ova. Fertilization and transfer of ova	Continuous assessment The final mark: 2 written theory tests (60%) 2 x assignments [presentation and written]

Biology lia	Embryo transfer and artificial insemination	(40%)
	•	
Instrumentations and Techniques	 Cryopreservation of semen, ova, and embryos 	Continuous assessment The final mark:
for Clinical Technology in Reproductive Biology lib	Testicular biopsyGenetic screening and analysisQuality control procedures	2 written theory tests (60%) 2 x assignments [presentation and written] (40%)