BOARD NOTICES

BOARD NOTICE 152 OF 2014

THE SOUTH AFRICAN PHARMACY COUNCIL

SCOPES OF PRACTICE AND QUALIFICATIONS FOR SPECIALIST PHARMACISTS

The South African Pharmacy Council (Council) intends to request the Minister of Health to:

- (a) publish amendments to the *Regulations relating to the registration of persons* and the maintenance of registers to make provision for specific categories for existing specialist pharmacists and new categories of specialist pharmacists:
 - (i) Radiopharmacist (existing);
 - (ii) Pharmacokineticist (existing);
 - (iii) Clinical Pharmacist (new);
 - (iv) Public Health Pharmacy and Management (new).
- (b) publish amendments to the *Regulations relating to the practice of pharmacy* to make provision for the scopes of practice of the abovementioned specialist pharmacists; and
- (c) publish regulations in terms of Sections 33 and 49(mA) to provide the required qualifications for the specialist pharmacists.

The qualifications and the proposed scopes of practice are published herewith for public comment prior to the said request to the Minister of Health.

SCHEDULE

- Radiopharmacy:
 - (a) Scope of practice for the specialist pharmacist in Radiopharmacy; and
 - (b) Qualification for the specialist pharmacist in Radiopharmacy.
- 2. Clinical Pharmacy:
 - (a) Scope of practice for the specialist pharmacist in Clinical Pharmacy; and
 - (b) Qualification for the specialist pharmacist in Clinical Pharmacy.
- 3. Public Health Pharmacy and Management:
 - (a) Scope of practice for the specialist pharmacist in Public Health Pharmacy and Management; and
 - (b) Qualification for the specialist pharmacist in Public Health Pharmacy and Management.

4. Pharmacokineticist:

(a) Scope of practice for the Pharmacokineticist.

In this notice "the Act" shall mean the Pharmacy Act, 53 of 1974 (as amended), and any expression to which a meaning has been assigned in the Act shall bear such meaning.

Interested persons are invited to submit within 60 days of publication of this notice, substantiated comments or representations on the qualifications and scopes of practice to the Registrar, The South African Pharmacy Council, Private Bag X40040, Arcadia, 0007, or Fax 0865063010 or email: BN@sapc.za.org.(for the attention of the Senior Manager: Legal Services and Professional Conduct).

TA MASANGO REGISTRAR

Address:

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SPECIALITIES FOR PHARMACISTS

AIM AND GOALS

To enable pharmacists to specialise and to meet advanced pharmaceutical care and the service needs of the country.

The goals for creating specialist pharmacists are to:

- (a) recognise expertise in pharmacy;
- (b) create a career framework, being career progression and job satisfaction;
- (c) move the profession forward;
- (d) achieve better outcomes for patients;
- (e) establish a referral system within the pharmacy profession;
- (f) manage risk and public safety; and
- (g) support the training of academics (teaching staff).

PRINCIPLES

- (a) The creation of specialist pharmacists must be needs driven;
- (b) The speciality in pharmacy must be based on advanced knowledge in the field of specialisation;
- (c) The speciality in pharmacy must be based on advance practical experience in the field of specialisation;
- (d) The speciality will be recognised if the postgraduate degree is pharmacy related; and
- (e) Broad specialist pharmacist would be created with an allowance to create subspecialities within the broad category when that sub-speciality has been well established in practice.

RADIOPHARMACISTS

SCOPE OF PRACTICE

- (a) Perform acts and services specially pertaining to the profession of a pharmacist;
- (b) Take a leading pharmaceutical role in protocol and guideline development in radiopharmacy and nuclear medicine;
- (c) Take a leading pharmaceutical role in compounding and/or manufacturing radiopharmaceuticals;
- (d) Act as a leading pharmaceutical partner within a multi-professional healthcare team in nuclear medicine departments and in industry;
- (e) Develop, implement, evaluate and provide strategic leadership for radiopharmacy services;
- (f) Appraise information, make informed decisions regarding supply and use of radiopharmaceuticals with the evidence available and be able to justify/defend the decisions;
- (g) Develop policies and procedures specifically for the specialty area;
- (h) Develop a quality and an evaluative culture within radiopharmaceutical services;
- (i) Perform pharmaceutical risk management;
- (j) Provide education and training related to radiopharmacy; and
- (k) Research, reach and publish in the field of radiopharmacy;

QUALIFICATION - PROFESSIONAL MASTER'S DEGREE IN RADIOPHARMACY

SYNOPSIS:

To provide a curriculum for a professional Master's Degree in Radiopharmacy, to enable students to register with the South African Pharmacy Council (hereafter referred to as Council) as specialists with a post-qualification that complies with Council's requirements.

Table 1: Summary of the proposed qualification

	Professional Master's Degree in Radiopharmacy
Duration:	Two years
Entry criteria:	Bachelor's Degree in Pharmacy
HEQF-level:	Level 9
Field (CESM):	09 Health Sciences and Social Services
Sub-field:	Curative Health
SAQA-credits:	360 credits
Qualification type:	Professional, exit-level, career-orientated, whole qualification
Final assessment and evaluation:	 Final, exit-level examination(s) will need to be passed in accordance with the relevant Higher Education provider's rules and regulations. In addition, a comprehensive portfolio of evidence will need to be submitted and successfully passed by the accredited provider. Requirements for registration as a specialist after obtaining the professional Master's Degree are presented in Appendix A to this document.
CPD requirements for annual reregistration:	As required by Council
Professional status:	Registration with Council as a practising Radiopharmacist
Articulation:	DPharm / Doctoral Degree

QUALIFICATION OUTLINE:

1. QUALIFICATION TITLE:

Master of Pharmacy in Radiopharmacy

☐ Abbreviation: MPharm (Radiopharmacy)

2. QUALIFICATION TYPE:

Professional Master's Degree

3. FIELD AND SUB-FIELD:

	Field: [09] Health Sciences and Social Services
П	Sub-field: Curative Health

4. LEVEL:

NQF/HEQF Level 9 (Master's Degree)

5. CREDITS:

Total credits: 360

6. RATIONALE FOR THE QUALIFICATION:

A shortage of radiopharmacists has been identified in South Africa and in Africa as a whole. Currently there are only two Council-registered specialist radiopharmacists in South Africa.

Radiopharmaceuticals are used in the diagnosis and treatment of many endstate organ diseases and life-threatening conditions such as major cardiac, renal, endocrine and cerebral disorders, as well as cancers and obscure infections. Their use is growing as they are key agents in the newer diagnostic modalities such as SPECT-CT and PET scintigraphy. Radiopharmaceuticals must be handled with care for both safety and efficacy. Their dosage form design, production and manipulation are often highly technical and sensitive to poor handling techniques, which render them ineffective or dangerous. Hence Radiopharmacy is a specialised area which is key to the diagnostic and treatment services offered in Nuclear Medicine.

There is a need for a qualified Radiopharmacist in every academic hospital Nuclear Medicine department, as well as in many private hospitals. Currently there are no posts for these professionals in the public sector, which presents a major obstacle. In addition, South Africa has major production centres for radiopharmaceuticals, which are sold and used throughout Africa, yet not one of these facilities has a qualified radiopharmacist. Inappropriate rolesubstitution therefore occurs in most facilities which radiopharmaceuticals. In hospitals, some of the tasks that should be performed radiopharmacists are performed by radiographers, whilst other radiopharmacy tasks are simply not performed at all. In production facilities there is role-substitution by radiochemists, medical physicists and pharmacists who have been trained in the workplace.

The existence of this speciality does not preclude the current practice of pharmacists already dispensing radiopharmaceuticals. Pharmacists should continue to perform the acts pertaining to the scope of practice of a pharmacist. Radiopharmacists should perform a leading pharmaceutical role in all activities which relate to radiopharmaceuticals. The role includes:

- (a) Procurement: Order, receipt, storage and inventory control of radiopharmaceuticals, ancillary drugs, supplies and related materials.
- (b) Compounding: Generator elution, kit reconstitution, preparation of products not commercially available and other radiolabelling procedures.
- (c) Manufacture: Radionuclide production and quality control of radiopharmaceuticals according to Good Manufacturing Practice in an industrial setting.
- (d) Quality assurance: Functional checks of instruments, equipment and devices and determination of radiopharmaceutical quality and purity (e.g. radionuclidic purity, radiochemical purity, chemical purity, particle size, sterility, apyrogenicity).
- (e) Dispensing: Preparation of bulk vials or individual patient doses for delivery to the user.
- (f) Distribution: Packaging, labelling and transport of radiopharmaceuticals to the user.
- (g) Health and safety: Radiation protection practices and proper handling of hazardous chemicals and biological specimens.
- (h) Provision of information and consultation: Communication of radiopharmaceutical-related information to others, i.e. general applicability (e.g. teaching), organisational (e.g. policies and procedures), or information concerning the care of specific patients.
- (i) Monitoring patient outcomes: Activities to assure optimal outcomes for individual patients, which includes patient preparation before radiopharmaceutical administration; prevention, recognition, investigation and rectification of clinical problems, such as drug interactions.
- Research and development: Laboratory testing (i) of radiopharmaceuticals, new compounding procedures, or new quality control methods. participation clinical trials and in radiopharmaceuticals.

The rationale for the Radiopharmacy postgraduate qualification is to train radiopharmacists who are able to register with Council as specialists in order to ensure safe and effective production and use of radiopharmaceuticals.

7. PURPOSE:

The purpose of this professional **Master's Degree** is to provide pharmacists who meet the minimum requirements for entry (Bachelor's Degree in Pharmacy) with the opportunity of becoming **specialists in the field of radiopharmacy** by expanding their basic knowledge, skills, values and attitudes, and therefore enabling them to meet the minimum requirements of Council. The degree is inherently a practice-based degree with a large component of work-integrated learning.

Council's Scope of Practice for Radiopharmacists

(a) Perform acts and services specially pertaining to the profession of a pharmacist.

- (b) Take a leading pharmaceutical role in protocol and guideline development in radiopharmacy and nuclear medicine.
- (c) Take a leading pharmaceutical role in compounding and/or manufacturing radiopharmaceuticals
- (d) Act as a leading pharmaceutical partner within a multi-professional healthcare team in nuclear medicine departments and in industry.
- (e) Develop, implement, evaluate and provide strategic leadership for radiopharmacy services.
- (f) Appraise information, make informed decisions regarding supply and use of radiopharmaceuticals with the evidence available and be able to justify/defend the decisions.
- (g) Develop policies and procedures specifically for the specialty area.
- (h) Develop a quality and an evaluative culture within radiopharmaceutical services.
- (i) Perform pharmaceutical risk management.
- (j) Provide education and training related to radiopharmacy.
- (k) Research, reach and publish in the field of radiopharmacy.

8. RULES OF COMBINATION:

Fundamental credits:	108
Core credits:	236
Elective credits:	16
Total:	360

9. ACCESS TO THE QUALIFICATION:

The minimum admission requirement is a four-year Degree in Pharmacy (NQF level 8) or equivalent; registration with the SA Pharmacy Council as an academic intern or as a pharmacist and placement in the area of specialisation.

10. LEARNING ASSUMED TO BE IN PLACE:

ir-year Degree in Pharmacy (NQF level 8) assuming the following is in
Professional and ethical practice
Communication (collaboration with members of the healthcare team) and self-management
Optimal use of medicines (therapeutic decision-making) and medication management
Anatomy and physiology
Pharmaceutics
Pharmacy practice (including aseptic experience, standard operating procedures, GMP and quality assurance)
Pharmacology
Research methodology

Candidates have to comply with all of the theoretical requirements set by Council for registration as a specialist pharmacist as well as annual registration, and must have a thorough understanding and working knowledge of South African *Good Pharmacy Practice* rules.

11. EXIT LEVEL OUTCOMES AND THEIR ASSOCIATED ASSESSMENT CRITERIA:

See Table 2

Table 2: Curriculum Outline

Learning Area	Exit Level Outcome	Credits	Notion and a second
Fundamental	Exit Level Outcome 1: Apply scientific knowledge in radionharmacy services	64	640
Fundamental	Exit Level Outcome 2: Promote safe handling of radiation sources and radiopharmaceuticals in compliance with relevant South African legislation	20	200
Fundamental	Exit Level Outcome 3: Institute quality management in radiopharmacy according to current Good Radiopharmacy Practice (cGRPP 1) and in compliance with GMP ² in radiopharmaceutical production	24	240
Core	Exit Level Outcome 4: Produce, procure, distribute and dispose of radiopharmaceuticals according to cGRPP and in compliance with GMP in radiopharmaceutical production	16	160
Core	Exit Level Outcome 5: Compound and dispense radiopharmaceuticals and radiolabelled blood elements according to GPP, cGRPP and recognised international standards and applicable legislation	28	280
Core	Exit Level Outcome 6: Conduct and management for radiopharmaceuticals and instrumentation in the radiopharmacy	20	200
Core	Exit Level Outcome 7: Monitor and promote diagnostic accuracy and successful treatment outcomes as an active member of the nuclear medicine team	40	360
Core	Exit Level Outcome 8: Act as part of a multidisciplinary team to provide information and consultation on radiopharmaceuticals and good radiopharmacy practice in clinical trials	12	160
Core	Exit Level Outcome 9: Conduct research and prepare for publication in the field of radiopharmacy	120	1200
Elective	Exit Level Outcome 10: Choose an elective topic	16	160
MPharm (Radiopharmacy)	TOTAL	360	3600

¹ Guidelines on current Good Radiopharmacy Practice (cGRPP) in the preparation of radiopharmaceuticals (most current version). EANM Radiopharmacy Committee Republic of South Africa. [Department of Health] (most current version). Medicines Control Council: South African Guide to GMP. Pretoria: Government Printers.

Notional Hours	640						
Associated Assessment Criteria	Criteria for Exit Level Outcome the role of Radiopharmac	Nuclear Medicine in diagnosis and therapy. 2. Medical physics: Explain atomic theory, decay processes, mathematics of	radioactivity decay, interaction of radiation with matter, types of radioactivity and radiation detection (instrumentation and cameras at basic level only).	3. Radiochemistry: Describe and explain production of radionuclides (natural, reactor, cyclotron, generators). Explain	properties of commonly-used diagnostic and therapeutic radionuclides, their chemistry and the principles of the use of ligands and chelating agents.	4. Radiopharmacology: Explain the localisation and mode of action of common radionuclides and radiopharmaceuticals, physical and biological half-life and docimetry.	5. Radiopharmaceutics: Explain and demonstrate aseptic radiolabelling techniques and quality control for radiopharmaceuticals.
Exit Level Outcomes	wledge in	radiopharmacy services Range statement: The range of scientific knowledge will include, but is not limited to:	 Radiation theory and medical physics instrumentation Production and properties of radionuclides 	 Radiopharmaceutical localisation, mode of action, half-life and dosimetry 	Aseptic preparation and quality control or radiopharmaceuticals	[64 credits]	
Learning Area	Fundamental						
Specific Exit Level	Master's Degree in Radiopharmacy (MPharm)						

200	
Assessment Criteria for Exit Level Outcome 2:	 Explain and apply legislation relevant to radiopharmacy services in the South African context³. Discuss and apply local and international guidelines relevant to the production, distribution, use and disposal of radionuclides and radiopharmaceutical products. Describe and demonstrate the principles of the "as low as reasonably achievable" (ALARA) concept and the importance of distance, shielding and time in radiation protection and radiation exposure limits. Demonstrate the practical implementation of radiation protection principles.
Exit Level Outcome 2:	Promote safe handling of radiation sources and radiopharmaceuticals in compliance with relevant South African legislation. [20 credits]
Fundamental	
Master's Degree in Fundamental Radiopharmacy	(MPharm)

³ Department of Minerals and Energy (most current version). Radioactive Waste Management Policy and Strategy for the Republic of South Africa, Pretoria, South Africa AND Department of Health (most current version). Directorate Radiation Control. Code of Practice for the Management and Disposal of Non-Nuclear Radioactive Waste.WSCP91-1, Pretoria, South Africa AND Republic of South Africa. [Department of Health]. 1965. Medicines and Related Substances Control Act (Act 101 of 1965). Pretoria.

240													-						
Assessment Criteria for Exit Level Outcome 3: 2	.	Design and implement environmental requirements for a radiopharmacy.	including choice, operation and	maintenance requirements of laminar flow hoods and isolators.	3. Undertake facility inspections and audits.	2	operating procedures (SOPs) for	radiopharmacy processes.	5. Assure radiopharmacy equipment	calibration and implement maintenance and	cleaning programmes.	6. Complete documents and maintain and	review records in accordance with	applicable legislation and SOPs.	7. Discuss the role of international	organisations in training and standards.	8. Describe the GMP approach for	radiopharmaceuticals and explain validation	processes.
Exit Level Outcome 3:	Institute quality management in radiopharmacy according to current	Good Radiopharmacy Practice (cGRPP) and in compliance with	GMP in radiopharmaceutical	production.		[24 credits]													
Fundamental																			
Master's Degree in Fundamental Radiopharmacv	(MPharm)																		

Master's Degree in Core	Core	Exit Level Outcome 4:	Assessment Criteria for Exit Level Outcome 4: 160	09
Radiopharmacy				
(MPharm)		Produce, procure, distribute and	Produce, procure, distribute and 1. Describe the legislative status of key	
		dispose of radiopharmaceuticals	radiopharmaceuticals and radionuclides.	-
		according to cGRPP and in	2. Explain and apply the production principles	
		compliance with GMP in		
		radiopharmaceutical production.	cyclotrons and generators.	And American
			3. Order, receive, store and maintain the	
			inventory of radiopharmaceuticals, ancillary	
			drugs, supplies and related materials	
		16 credits	according to cGRPP.	
			4. Distribute radiopharmaceuticals to the user	
			according to cGRPP (packaging, labelling	
			and transport).	
			5. Conduct radionuclide and	
			radiopharmaceutical waste management	
			according to current South African	**************************************
			legislation ⁴ and cGRPP.	
			6. Manage the ordering of and record-keeping	
			for Section 21 radiopharmaceuticals.	

Department of Minerals and Energy. (most current version). Radioactive Waste Management Policy and Strategy for the Republic of South Africa. Pretoria. South Africa. AND Department of Health. (most current version). Directorate Radiation Control. Code of Practice for the Management and Disposal of Non-Nuclear Radioactive Waste. WSCP91-1, Pretoria, South Africa.

280

Assessment Criteria for Exit Level Outcome 5:

Compound radiopharmaceuticals according to GPP and cGRPP. Perform generator

dispense

and

Compound

Exit Level Outcome 5:

Core

Master's Degree in Radiopharmacy (MPharm)

radiopharmaceuticals, radiolabelled

													-										
olition bit roconstitution proparation of			2. Dispense radiopharmaceuticals according	to GPP and cGRPP, including evaluation of	the prescription, preparation of bulk vials or	individual patient doses for delivery to the	user and prepare and reconstitute cold kits.	3. Blood products: Prepare radiolabelled red	and white cells and other blood elements	according to local or ISORBE protocols.	4. Compound, manipulate and prepare sterile	admixtures according to SOPs, in	accordance with aseptic techniques and	principles of GMP and/or GPP.	5. Appraise sterilisation methods for	commonly used radiopharmaceuticals.	6. Manage radiopharmacy cleaning	programmes so that sources and risks of	microorganism contamination are reduced.	7. Manage record systems for	radiopharmaceutical preparations produced	in accordance with legal requirements and	organisational policies and procedures.
blood olomonto biologicolo ond	other novel radiopharmaceutical	dosage forms according to GPP,	cGRPP and recognised	international standards and	applicable legislation ⁵ .	•			[28 credits]	1													

National Health Act (61 of 2003). Chapter 8. Control of use of blood, blood products, tissue and gametes in humans. Sections 53-68 and all relevant Regulations thereunder. ⁵ Medicines Control Council (Most current version). Guidelines for similar biological medicines (biosimilar medicines). Non-clinical and clinical requirements. AND. The

Master's Degree in Core	Core	Exit Level Outcome 6:		Assessment Criteria for Exit Level Outcome 6: 200
(MPharm)		Conduct and monitor	quality	quality 1. Describe in detail the principles of
		management	for	radiopharmacy quality management in
		radiopharmaceuticals	and	hospitals and in production facilities.
		instrumentation in	the	2. Conduct functional checks of instruments,
		radiopharmacy.		equipment and devices.
				3. Determine radiopharmaceutical quality and
				purity requirements for radionuclidic,
		[20 credits]		radiochemical and chemical purity.
				4. Evaluate and ensure particle size, sterility
				and apyrogenicity of radiopharmaceuticals.
				5. Ensure completion and filing of appropriate
				records in accordance with cGRPP.

400	
Lesessment Criteria for Exit Level Outcome 7: 1. Describe the pathophysiology of key disease states seen in nuclear medicine. 2. Apply the principles of pharmaceutical care and patient monitoring. 3. Interpret clinical laboratory results. 4. Interpret laboratory tests associated with the identification and quantification of pathogens. 5. Explain the mode of action of common radionuclides and radiopharmaceuticals. 6. Analyse the rationale for the choice of specific radiopharmaceuticals in common conditions (disease or suspected diagnosis, age and gender of the patient, contraindications, 7. Evaluate patient preparation with regard to prevention or recognition of drug or food interactions before radiopharmaceutical administration. 8. Appraise the administration and clinical use of commonly used radionuclides and radio-	pharmaceuticals. 9. Demonstrate active participation in decision-making in the nuclear medicine team.
Monitor and promote diagnostic accuracy and successful treatment outcomes as an active member of the nuclear medicine team. Range statement: The range of conditions includes but is not limited to disorders and diseases, commonly seen in nuclear medicine, of the following systems: Cardiovascular Cardiovascular Cardiovascular Cardiovascular Hepatobiliary Lymphatic Pulmonary Skeletal Skeletal	
Master's Degree in Core Radiopharmacy (MPharm)	

Act as part of a multidisciplinary team to provide information and consultation radiopharmaceuticals and <i>Good Radiopharmacy Practice</i> and in clinical trials. Act as part of a multidisciplinary 1. Communicate radiopharmacy information (e.g. teaching, policies and procedures for the healthcare team. Assessment Criteria for Exit Level Outcome 8: (e.g. teaching, policies and procedures for the healthcare team. Assessment Criteria for adiopharmacy information the care of specific patients) to members of the healthcare team. Badiopharmacy Practice and in 2. Record, identify and address radiopharmaceutical causes of scintigraphic anomalies.	120 1. Communicate radiopharmacy information (e.g. teaching, policies and procedures for the care of specific patients) to members of the healthcare team. 2. Record, identify and address radiopharmaceutical causes of scintigraphic anomalies.
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1200														-					
Assessment Criteria for Exit Level Outcome 9:	1. Critically evaluate information sources, literature and research on medicines and			αi	methodology in the development of a	research protocol. Obtain ethical clearance	if necessary.	3. Conduct research in accordance with	established research methodology and	ethics, as well as Good Clinical Practice	where necessary.	4.	results and formulate conclusions and	recommendations.	5. Write and submit a technical report,	manuscript for publication or minor	dissertation and obtain approval.		
Exit Level Outcome 9:	Conduct research and prepare for publication in the field of	radiopharmacy. Range statement:	Research may include, but is not	limited to, the following areas:	Development of new	radiopharmaceuticals,	Laboratory testing of	radiopharmaceuticals,	Compounding procedures,	Quality assurance or quality	control methods,	Clinical use of	radiopharmaceuticals,	Radiopharmaceuticals	management.			[120 credits]	
Core																			
Master's Degree in Core	(MPharm)																		

Outcome 160	ansition to	
Assessment Criteria for Exit Level Outcome	Demonstrate a deep knowledge of the chosen elective field of radiopharmacy, for transition to independent practice.	
Assessment C	Demonstrate a deep lective field of radiopindependent practice.	
Exit Level Outcome 10:	Choose an elective topic. Topics for electives may include but are not limited to: • Hospital radiopharmacy • Radiopharmaceutical manufacture, production or compounding • Radiopharmaceutical clinical trials • Regulation of radiopharmaceuticals	[16 credits]
Elective		
Master's Degree in	(MPharm)	

12. CRITICAL CROSS-FIELD OUTCOMES:

The following critical cross-field outcomes will form an integral part of the exit level outcomes of this programme:

- Identify and solve problems in which responses display that responsible decisions using critical and creative thinking have been made; Work effectively with others as a member of a team, group, organisation and community: Organise and manage oneself and one's activities responsibly and effectively: Collect, analyse, organise and critically evaluate information: Communicate effectively using visual, mathematical and/or language skills in the modes of oral and/or written persuasion; Use science and technology effectively and critically, show responsibility towards the environment and health of others; Demonstrate an understanding of the world as a set of related systems by recognising that problem-solving contexts do not exist in isolation; Promote the personal and professional development of each learner in the programme, and the social and economic development of the society
 - reflecting on and exploring a variety of strategies to learn more effectively;
 - participating as responsible citizens in the life of local, national and global communities;
 - being culturally and aesthetically sensitive across a range of social contexts;
 - exploring education and career opportunities; and

at large, by creating an awareness of the importance of:

developing entrepreneurial opportunities.

13. INTERNATIONAL COMPARABILITY:

The following examples are provided to illustrate the proposed curriculum's competitiveness and comparability among both developed and developing countries.

Radiopharmaceuticals fall into two major groups — those used for scintigraphy and single photon emission computed tomography (SPECT) and those used for positron emission tomography (PET). PET radiopharmaceuticals are often produced in cyclotrons. Cyclotron operation necessitates specialised training. In Sub-Saharan Africa, there are very few cyclotrons. In other parts of the world, some radiopharmaceutics degrees deal only with cyclotron-produced radiopharmaceuticals. South Africa has four cyclotrons (two in Pretoria and two in Cape Town). In the Southern African context, a degree which deals with cyclotron produced-radiopharmaceuticals as well as SPECT radiopharmaceuticals is required.

In addition, South Africa has a need for radiopharmacists in the clinical setting, hence the clinical use of diagnostic and therapeutic radiopharmaceuticals is an essential area for postgraduate study.

Few Radiopharmacy / nuclear pharmacy postgraduate degrees are listed internationally. Some qualifications for nuclear medicine are stated to lead to radiopharmacy careers.

Radiopharmacy/Nuclear Pharmacy Degrees

The following degree courses have been identified and are summarised below. More details follow.

United Kingdom (Kings College MSc Radiopharmaceutics and PET Radiochemistry)

Co	re programme content:
	Module 1 – Introduction to Medical Imaging Sciences
	Module 2 – Radiopharmacology Formulation and Manufacture
	Module 3a – Radiopharmaceutical Chemistry
	or
	Module 3b - Radiopharmaceutical Chemistry and Radiopharmaceutical
	Design
	Module 4a – Cyclotron Engineering and Nuclear Chemistry
	or
	Module 4b – Radiopharmaceuticals in Practice
	Module 5 – Research Project

FORMAT AND ASSESSMENT

Written examinations (modules 1, 2, 3a, 3b and 4a); practical laboratory work and reports (modules 1, 2, 3a, 3b, 4a and 5); case studies and oral presentation (module 4b); workshops (all modules); audio-visual presentations (all modules); laboratory or library-based research project (module 5).

Iran (Tehran University of Medical Sciences)

The course includes the following topics:

- Health physics and radiobiology
- Radiochemistry
- Instrumental and analytical methods
- Synthesis of radiolabelled compounds
- Pharmacology
- Medical statistics

Macedonia (University of Goce Delcev - Stip)

- Basic applied pharmacy
- Radiopharmaceutical chemistry
- Radiopharmaceutical preparation
- Quality control of radiopharmaceuticals
- Nuclear physics, radiation safety and regulations
- Nuclear medicine aspects of clinical practice
- Radiopharmaceutical preparation SPECT, PET and therapeutic
- Operation of a GMP facility
- Quality control of radiopharmaceuticals
- Clinical application of radiopharmaceuticals in nuclear medicine
- Master's thesis

United States of America (USA)

Radiopharmacy (nuclear pharmacy) services in the USA are often centralised.

A radiopharmacist must possess an active pharmacist licence and have received didactic instruction (200 hours) and/or supervised professional experience in the practice of nuclear pharmacy (500 hours). (APhA-APPM Section on Nuclear Pharmacy: Nuclear Pharmacy Practice Guidelines).

- University of Purdue 200 hours clerkships in industry, centralised radiopharmacy or nuclear medicine. The coursework covers: radiation physics, radiation safety, regulatory issue, proper use of equipment, and radiation biology. The advanced clinical clerkship includes information resources pertaining to nuclear medicine and nuclear pharmacy practice, information services, centralised unit dose radiopharmacy service and nuclear medicine department-based hot labs, the receipt of orders, preparation of prescriptions, compounding of radiopharmaceuticals, performance of quality control and quality assurance tests of compounded radiopharmaceuticals and the compounding environment, and the packaging and delivery of nuclear pharmacy products. Also knowledge of risks associated with administered the radiopharmaceuticals and radiation exposure.
- University of New Mexico. The certificate course has 200 hours of didactic learning and 500 hours of experiential training. It includes an introduction to radiopharmacy, nuclear pharmacy instrumentation, radiopharmaceutical chemistry, chemistry, radiopharmacy health and radiation biology, and radiopharmacology. Experiential training is in clinical and institutional radiopharmacy.
- Nuclear Education Online (NEO) offers an online course for certification purposes. The course covers: nuclear physics, instrumentation, radiation safety and regulations, radiation biology and radiochemistry.

European specialisation certificate in radiopharmacy

The Radiopharmacy Committee of the European Association of Nuclear Medicine (EANM) has established a European postgraduate specialisation certificate in radiopharmacy. A certificate after successful attendance may be awarded to participants, who, in the view of the EANM Radiopharmacy Board, are suitably qualified, in that they have:

- acquired a university postgraduate diploma through attendance at appropriate courses teaching the theoretical components of the radiopharmacy syllabus;
- completed a two-year period of experience in a radiopharmacy department during which they have completed the practical components of the syllabus; and
- completed a nationally acceptable course on radiation safety.

14. INTEGRATED ASSESSMENT:

A combination of integrated assessment strategies, which will combine both
formative and summative assessment and evaluation, will be used to ensure
that the purpose of the qualification is achieved. Assessments may include, but
are not limited to, the following strategies:

Ш	Portiolios of evidence
	Practical experience work-place assessments

	Written and oral assessments and examinations Written assignments OSPEs Case studies Journal clubs Self-assessment strategies, peer-group assessment and preceptor evaluation
CRE	EDIT ACCUMULATION AND TRANSFER:
	didates may apply for recognition of credits obtained as part of an mplete qualification at the same or a different institution, depending on

individual institutional policies. 16. ARTICULATION (PROGRESSION):

Completion of a Master's Degree meets the minimum entry requirement for admission to a Doctoral Degree, usually in the area of specialisation in the Master's Degree.

Articulation may also be horizontal to entries into other Master's Degrees in a similar or related field, or area of specialisation.

17. MODERATION OPTIONS:

Suitable moderating options should be included in each application for accreditation to provide this qualification in accordance with the stipulations of the Council on Higher Education (CHE), as well as the relevant ETQA (i.e. Council). Both internal and external moderation should form an integral part of the provision of this qualification.

18. CRITERIA FOR THE APPOINTMENT OF ASSESSORS:

Assessors in the field of radiopharmacy must have a suitable background with a proven track record and relevant experience to enable them to make sound judgements through their expert application of the assessment criteria specified for this qualification.

19. NOTES:

15.

All candidates must, in addition to their current registration as academic interns or pharmacists, be registered with Council for study towards the specialisation for the duration of the period of learning as specified in current relevant legislation.
The range of elective learning areas offered will be dependent on the approval of the provider and ETQA.
Credit values reflected for each exit level outcome in Table 2 should be regarded only as a guideline.
The respective assessment criteria aim to test the achievement of the specific learning outcomes. Some of these criteria are practice-based, thus providers are required to include periods in their curricula for this purpose.
After attaining the Master's Degree, the candidate may commence with the process for registration as a specialist pharmacist in radiopharmacy

with Council. Requirements for this registration process will be determined by Council.

(See Appendix A)

APPENDIX A

Requirements for registration as a specialist after obtaining the Professional Master of Pharmacy in Radiopharmacy

The prospective candidate should be a registered pharmacist with Council

Training Site

A site registered with the Council as a training institution, pharmacy, health or manufacturing facility where radiopharmaceuticals are routinely handled.

Tutor or supervisor

A postgraduate pharmacist or specialist medical practitioner in nuclear medicine, with at least two years' experience in the field.

Practical training

As stipulated by Council

Evaluation and panel

As stipulated by Council.

CLINICAL PHARMACIST

SCOPE OF PRACTICE - CLINICAL PHARMACIST

- (a) Perform acts and services pertaining to the profession of a pharmacist;
- (b) Provide advanced clinical pharmacy services to a variety of specialities;
- (c) Act as a leading pharmaceutical partner within a multi-professional healthcare team:
- (d) Develop, implement, evaluate and provide strategic leadership for clinical pharmacy services;
- (e) Appraise clinical pharmacy information, make informed decisions with the evidence available and be able to justify/defend the decisions;
- (f) Take a pharmaceutical leadership role in clinical protocol and guideline development;
- (g) Lead clinical audits of medicine use;
- (h) Develop policies and procedures specifically for clinical pharmacy;
- (i) Provide education and training related to clinical pharmacy;
- (j) Perform research, teach and publish in clinical pharmacy; and
- (k) Initiate and participate in pharmacovigilance related to clinical practice.

QUALIFICATION - PROFESSIONAL MASTER'S DEGREE IN CLINICAL PHARMACY

SYNOPSIS:

The aim is to provide a curriculum for a Professional Master's Degree in Clinical Pharmacy, to enable students to register with the South African Pharmacy Council (hereafter referred to as "Council") as specialists.

Table 1: Summary of the proposed qualification

	Professional Master's Degree in Clinical Pharmacy
Duration:	Two years
Entry criteria:	Bachelor's Degree in Pharmacy (Level 8)
HEQF-level:	Level 9
Field:	09 Health Sciences and Social Services
Sub-field:	Curative Health
SAQA-credits:	360 credits
Qualification type:	Professional, exit level, career-orientated, whole qualification
Final assessment and evaluation:	 Final, exit level examination(s) will need to be passed in accordance with the relevant Higher Education provider's rules and regulations. In addition, a comprehensive portfolio of evidence will need to be submitted and successfully passed by the accredited provider. Requirements for registration as a specialist after obtaining the professional Master's Degree are presented in Appendix A to this document.
CPD requirements for annual reregistration:	As required by Council
Professional status:	Registration with Council as a practising clinical pharmacist
Articulation:	DPharm / Doctoral Degree

QUALIFICATION OUTLINE:

QUALIFICATION TITLE:

Master of Pharmacy in Clinical Pharmacy Abbreviation: MPharm (Clinical Pharmacy) 2. **QUALIFICATION TYPE:** Professional Master's Degree

3. FIELD AND SUB-FIELD:

Field: [09] Health Sciences and Social Services
Sub-field: Curative Health

4. LEVEL:

1.

NQF/HEQF Level 9 (Master's Degree)

5. **CREDITS:**

Total credits: 360

6. **RATIONALE FOR THE QUALIFICATION:**

The rationale for the qualification is to train advanced level clinical pharmacists who are able to register with Council as specialists who contribute to capacity building in the field of clinical pharmacy, and to create specialists in the field of pharmacy for the advancement of healthcare in South Africa.

According to Van Mil (2004): "If we want to try to prove that the structured provision of pharmaceutical care has an effect on outcomes, we must first of all make sure that the care provided matches the needs of the patients in that specific health system".6

Historically, the role of the pharmacist, regardless of the health setting, was to ensure prompt and efficient medication supply, adequate stock, accurate dispensing, compounding, storage and transport, and to ensure that medicines were easily accessible to patients who needed them. The pharmacist was also responsible for the selection of medicines, dosage forms, and monitoring of patient compliance.7

Clinical pharmacy is aimed at the development and promotion of rational and appropriate use of medicines and pharmaceutical care, in the interest of the patient and the community.8

⁶ Van Mil F. 2004. Proving the benefits of pharmaceutical care. *Pharmacy World and Science*, 26:123.

⁷ Hepler CD, and Strand LM.1990. Opportunities and responsibilities in pharmaceutical care. American Journal of Hospital Pharmacy, 47:533-43.

⁸ The South African Society of Clinical Pharmacy (SASOCP). 2011. Constitution of the South African Society of Clinical Pharmacy. Available from: http://www.sasocp.co.za. (Accessed: 01/08/2014).

Patients with advanced, untreatable diseases have multiple symptoms and treatment becomes complicated⁹. This makes it difficult for carers to manage their patients' medication, which leads to patients' symptoms being inadequately controlled and a low level of compliance¹⁰. Pharmacists have the responsibility to identify, resolve, and prevent each patient's medicine therapy problems. These responsibilities are met by using the caring paradigm in a patient-centred manner.

Pharmaceutical care mandates that pharmacists should not only dispense medication, but also assume the responsibility of improving the quality of life of patients and improving therapy outcomes¹¹.

Pharmaceutical care involves the implementation of the following steps¹²:

- The assessment of patient health and formulation of a treatment plan to treat disease and symptoms
- Monitoring of patient response to therapy to ensure optimum therapeutic effects
- Performing medication reviews to detect and resolve medication-related problems
- Documentation of the care provided and provision of advice to patients in a way that patients understand.

In South Africa, clinical pharmacists are currently not part of the traditional ward staff, as seen in the United States (US) or the United Kingdom (UK)¹. This situation may be due to lack of human resources and inadequate training and the occupational levels of pharmacists. There is a need to develop and accredit formal qualifications which will enable qualifying pharmacists to render professional services within a recommended scope of practice, and under the auspices of the statutory body, namely Council.

7. PURPOSE:

The primary purpose of a professional Master's Degree is to educate and train graduates who can contribute to the development of knowledge at an advanced level so they are prepared for specialised professional employment.

In some cases, a professional Master's Degree may be designed in consultation with a professional body, or fulfil all or part of the requirements for professional registration or recognition, and may include appropriate forms of work-integrated learning.

⁹ MacRobbie A, Addie S, & Grant E. 2009. The pharmacist in palliative care. Scottish Palliative Care Pharmacists Association (SPCPA). Available from:

http://www.nes.scot.nhs.uk/media/347752/14551_20nes_20pall_care_20dl_20v9_20final.pdf(Accessed: 01/08/2014).

¹⁰ MacRobbie A, Addie S, & Grant E. 2009. The pharmacist in palliative care. Scottish Palliative Care Pharmacists Association (SPCPA). Available from:

http://www.nes.scot.nhs.uk/media/347752/14551 20nes 20pall care 20dl 20v9 20final.pdf(Accessed: 01/08/2014).

11 Hughes CM, Hawwa AF, Scullin C, Anderson C, Bernsten CB, Bjo rnsdo ttir I, Cordina MA, Alves da Costa M, De Wulf I, Eichenberger P, Foulon V, Henman MC, Hersberger KE, Schaefer MA, Sondergaard M, Tully MP, Westerlund T & McElnay JC. 2010. Provision of pharmaceutical care by community pharmacists: a comparison across Europe. Springer science & business media. Available from: http://upload.sitesystem.ch/B2DBB48B7E/EE929BDFA5/4D7608D543.pdf. (Accessed: 01/08/2014).

12 Minnesota Senate. 2005. Medication management care. 8th Legislative session, No. 973,1st Engrossment. Available from: https://www.revisor.leg.state.mn.us/bin/bldbill.php?bill=SO973.1&session=Is84. (Accessed: 01/08/2014).

Successful completion of a programme requires a high level of theoretical engagement and intellectual independence as well as a demonstration of the ability to relate knowledge to the resolution of complex problems in appropriate areas of professional practice. In addition, a professional Master's Degree must include an independent study component that comprises at least a quarter of the credits at NQF level 9, consisting of either a single research or technical project or a series of smaller projects demonstrating innovation or professional expertise.

Master's graduates must be able to deal with complex issues both systematically and creatively, design and critically appraise analytical writing, make sound judgements using data and information at their disposal and communicate their conclusions clearly to specialist and non-specialist audiences, demonstrate self-direction and originality in tackling and solving problems, act autonomously in planning and implementing tasks with a professional orientation, and continue to advance their knowledge, understanding and skills relevant to a particular profession.

The purpose of this professional **Master's Degree** is to provide pharmacists, who meet the minimum requirements for entry (Bachelor's Degree in Pharmacy) with the opportunity of becoming **specialists in the field of clinical pharmacy** by expanding their basic knowledge, skills, values and attitudes, and therefore enabling them to meet the minimum requirements of Council. The degree is inherently a practice-based degree with a large component of work-integrated learning.

Council's Scope of Practice for Clinical Pharmacists

- 1. Perform acts and services pertaining to the profession of a pharmacist.
- 2. Provide advanced clinical pharmacy services to a variety of specialities.
- 3. Act as a leading pharmaceutical partner within a multi-professional healthcare team.
- 4. Develop, implement, evaluate and provide strategic leadership for clinical pharmacy services.
- 5. Appraise clinical pharmacy information, make informed decisions with the evidence available and be able to justify/defend the decisions.
- 6. Take a pharmaceutical leadership role in clinical protocol and guideline development.
- 7. Lead clinical audits of medicine use.
- 8. Develop policies and procedures specifically for clinical pharmacy.
- 9. Provide education and training related to clinical pharmacy.
- 10. Perform research, teach and publish in clinical pharmacy.
- 11. Initiate and participate in pharmacovigilance related to clinical practice.

8. RULES OF COMBINATION:

Fundamental credits:	60
Core credits:	284
Elective credits:	16
<u>Total:</u>	360

9. ACCESS TO THE QUALIFICATION:

The minimum admission requirement is a four-year Degree in Pharmacy (NQF level 8) or equivalent; registration with Council as an academic intern or as a pharmacist, and placement in the area of specialisation.

10. LEARNING ASSUMED TO BE IN PLACE:

of South African Good Pharmacy Practice rules.

	ur-year Degree in Pharmacy (NQF level 8), or equivalent, assuming the
follov	ving is in place:
	Professional and ethical practice
	Communication (collaboration with members of the healthcare team) and self-management
	Optimal use of medicines (therapeutic decision-making) and medication management
	Pharmacology
	Research methodology.
Cour	didates have to comply with all of the theoretical requirements set by not for registration as a specialist pharmacist as well as annual tration, and must have a thorough understanding and working knowledge

11. EXIT LEVEL OUTCOMES AND THEIR ASSOCIATED ASSESSMENT CRITERIA:

See Table 2

Table 2: Curriculum Outline

Learning Area	Exit Level Outcome	Credits	Notional Hours
Fundamental	Exit Level Outcome 1: Formulate pharmaceutical care plans and interpret prescriptions to guide treatment for individual patients and counsel patients to improve treatment outcomes	20	200
Fundamental	Exit Level Outcome 2: Use clinical information, laboratory and diagnostic tests and results to assist with, or support, therapeutic assessments and decisions, including medicine therapy	20	200
Fundamental	Exit Level Outcome 3: Apply basic and clinical pharmacokinetics, pharmacogenomics and pharmacodynamics in medicine therapy for individualised patient care	20	200
Core	Exit Level Outcome 4: Optimise therapy for infectious diseases	28	280
Core	Exit Level Outcome 5: Optimise therapy for disorders related to the endocrine system, including obstetric, gynaecological and urological conditions	20	200
Core	Exit Level Outcome 6: Optimise therapy for disorders related to the gastrointestinal system	20	200
Core	Exit Level Outcome 7 Optimise therapy for disorders related to the cardiovascular system	28	280
Core	Exit Level Outcome 8: Optimise therapy for disorders related to the renal system	20	200
Core	Exit Level Outcome 9: Optimise therapy for neurological and psychiatric disorders	28	280
Core	Exit Level Outcome 10: Optimise therapy for disorders related to the respiratory system	20	200
Core	Exit Level Outcome 11: Conduct research and prepare for publication in the field of clinical pharmacy	120	1200
Elective	Exit Level Outcome 12: Optimise therapy for disorders and/or optimise clinical pharmacy practice related to any one (1) chosen elective topic	16	160
MPharm (Clinical Pharmacy)		360	3600

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Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's	Fundamental	Exit Level Outcome 2:	Assessment Criteria for Exit Level Outcome	200
Degree In Clinical		I se clinical information Jahoratory and	·ii	
Pharmacv			1. Describe, analyse, review and apply	
(MPharm)		support, therapeutic assessments and	normal/reference ranges for commonly	
		decisions, including medicine therapy.	used tests.	
		Range statement: The range of topics will	2. Appraise and explain the possible	
		include, but is not limited to, the following	aetiology of, and pathology related to,	
		points:	clinical laboratory results which are	
		 Vital signs and clinical condition 	outside these ranges.	
		 Urea and electrolytes 	3. Interpret and apply the impact of the	
		 Medical microbiology, immunology 	aetiology of, or pathology related to,	
		• Genetics	clinical laboratory test results on	
		Full blood count	medicine therapy of individual patients.	
		 Organ function tests 		
		 Pathology and pathophysiology as 		
		related to these tests		
		[20 credits]		

Apply basic and clinical pharmacokinetics, Pharmacy In medicine therapy for individualised pharmacokynamics and pharmacodynamics and pharmacognomics and pharmacodynamics Pharmacy and pharmacodynamics and pharmacodynamic definitions and pharmacodynamics and pharmacodynamics Principles of pharmacokinetics, pharmacodynamics and disease associated genetic variations that pharmacodynamics Patient disease state and the individualised dosing calculation of laboratory values and harmacodynamics its influence on medicine therapy in interpreted (e.g., clearance, volume of distribution and half-life). Secretary of topics will a concepts and nonenclature. Individualised dosing calculations interpreted dosing calculations in interpreted (e.g., clearance, volume of distribution and half-life). Patient disease state and the harmacodynamic endering medicine therapy and papropriate policy and interpreted of graphory and papropriate policy and papropriate paginations. Page pharmacodynamics or page pharmacodynamic decisions unage and papropriate page pharmacodynamic policy and papropriate page pharmacodynamic papropriate page pharmacodynamic page pharmacodynamic papropriate page pharmacodynamic papropriate page pharmacodynamic papropriate page pharmacodynamic papropriate page pharmacodynamic pharmacodynamic page pharmacodynamic pharmac	Specific Exit Level	Learning Area	Exit Level Outcomes		Notional Hours
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using alterations in pharmacokinetic and pharm				make appropriate therapeutic decisions	
pharmacodynamic dosing alterations.				using alterations in pharmacokinetic and	
				pharmacodynamic dosing alterations.	

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's	Core	Exit Level Outcome 4:	Assessment Criteria for Exit Level Outcome 4:	280
Clinical		Optimise therapy for infectious diseases.	1. Classify common pathogens and describe mechanisms related to the development of	
Priarriacy (MPharm)		include, but is not limited to, the following	acquiring resistance.	
		points:		
		 Pathogens and laboratory tests 	9	
		 Pathophysiology of the conditions 	antimicrobials.	
		 Medication-related problems 	d implem	
		Evidence-based, patient-specific	antimicrobial stewardship principles as appulicable to clinical practice.	
		medication treatment plans	4. Define. discuss and appraise	
		 I reatment plans, including assisting the 	pathophysiology of the diseases as	
		patient	by microorganisms.	
		 Patient response to and modification of 	5. Use pharmacodynamic principles to guide	
		pharmacotherapy	and ensure effective antimicrobial therapy.	
		 Patient interventions and antimicrobial 	6. Define, discuss and apply infectious disease	
		stewardship	principles to the various infective conditions.	
		-	7. Appraise, organise and evaluate patient	
			information.	
			8. Recognise, categorise and interpret	
		2000	ē	
			appropriate interventions.	
			9. Formulate patient-specific, evidence-based	
			medication treatment plans.	
			10. Formulate and implement treatment plans	
			(including non-medicine treatment) and	
			assist the patient with implementation.	
			11. Monitor and evaluate pharmacotherapy to	
			assess patient response.	
			12. Document patient interventions in	
			accordance with professional and legal	
			requirements.	

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's Degree in	Core	Exit Level Outcome 6:	Assessment Criteria for Exit Level Outcome	200
Clinical		Optimise therapy for disorders related to the	Ši.	
Pharmacy		gastrointestinal system.	1. Define, discuss and appraise the	
(MPharm)		Range statement: The range of topics will	pathophysiology of disorders related to	
		Include, but is not limited to, the following matters:	ine gastrointestinal system. 2. Appraise, organise and evaluate patient	
		 Anatomy and physiology of the 		
			3. Identify, categorise and interpret	
		 Gastro-oesophageal reflux (GORD) 	medication-related problems and make	
		 Peptic ulcer disease 	appropriate interventions.	
		 Inflammatory bowel disease 	4. Formulate and apply patient-specific,	
		 Treatment of nausea and vomiting 	evidence-based medication treatment	
		 Irritable bowel syndrome 		
		 Treatment of constipation and diarrhoea 	5. Monitor and evaluate clinical nutrition	
		 Hepatic medicine metabolism 	when required according to patient	
		 Alcoholic liver disease 	Specific disease states.	
		 Drug-induced liver disease 		
		Pancreatitis	(including	
		 Hepatitis (viral, acute and chronic) 	implementation.	
		 Identify and manage diseases related to 	7. Monitor and evaluate pharmacotherapy	
		nutritional disorders (including a basic		
		understanding of clinical nutrition)	8. Document patient interventions in	
			accordance with professional and legal requirements	
		[20 credits]		

Master's Core Exit Let Degree in Clinical Optim Pharmacy (MPharm)	Exit Level Outcome 7: Optimise therapy for disorders related to the cardiovascular system.		
.⊑ δ̂ €	imise therapy for disorders related to the diovascular system.	Assessment Criteria for Exit Level Outcome	280
ò ̂ €	imise therapy for disorders related to the diovascular system.		
δ̂ c	diovascular system.		
	ii	1. Define, discuss and appraise	
includ condition which was the condition of the condition	Range statement: The range of topics will	pathophysiology of various cardiac	
condition of the condit	include, but is not limited to, the following	disorders.	
······································	conditions:	2. Appraise, organise and evaluate patient	
	Hypertension	information including laboratory tests	
N A A A A A A A A A A A A A A A A A A A	Heart failure	specific to the cardiac system.	
S A A A A A A A A A A A A A A A A A A A	Ischaemic heart disease	3. Recognise, categorise and interpret	
Ary Popularian	Myocardial infarction (MI)	medication-related problems and make	
Dy Ac	Arrhythmias	appropriate interventions.	
- AC	Dvslipidaemia	4. Formulate and apply patient-specific,	
Ac Ac	Thromboembolic disease	evidence-based medication treatment	
Ac Ac		plans.	
e	coronary syndrome	5. Formulate and implement treatment	
S	lests used to evaluate the	plans (including non-medicine	
	cardiovascular system	nt) and assist the	
		implementation.	
		6. Monitor and evaluate pharmacotherapy	
50 02	בס כופחוס]	to assess patient response.	
		7. Document patient interventions in	
		accordance with professional and legal	
		requirements.	

				Hours
Sinical Pharmacy	Core	Exit Level Outcome 8:	Assessment Criteria for Exit Level Outcome	200
harmacy		Optimise therapy for disorders related to the	i.	
		renal system.	1. Define, discuss and appraise	
(MPnarm)		Range statement: The range of topics will	of disorders r	
		include, but is not limited to, the following	•	
		points:	2. Appraise, organise and evaluate patient	
		 Assessment and quantification of renal 	information.	
		function	3. Recognise, categorise and interpret	
		 Acute renal failure 	medication-related problems and make	
		 Chronic renal failure and end stage 	appropriate interventions.	
		renal failure	4. Formulate and apply patient-specific,	
		 Drug-induced renal disease 	evidence-based medication treatment	
		 Appropriate calculations in adjustment 		
		of medicine therapy in renal failure	5. Formulate and implement treatment	
		Assessment and management of the	plans (including non-medicine	
		hydration status of a hospitalised patient	treatment) and assist the patient with	
			implementation.	
			6. Monitor and evaluate pharmacotherapy	
		Stinary CCI	to assess patient response.	
			7. Document patient interventions in	
			accordance with professional and legal	
			requirements.	,,

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's	Core	Exit Level Outcome 9:	Assessment Criteria for Exit Level Outcome	280
Degree in				
Clinical		Uptimise therapy for neurological and	000000000000000000000000000000000000000	
///Indiliacy		psychiatric disorders.	i. Dellile, discuss and applaise	
(MFnarm)		include but is not limited to the following	the control notices ever (including	
		module, but is not infilled to, the following	lile central nervous system (including	
		conditions:		
		Psychiatric Disorders	2. Appraise, organise and evaluate patient	
		 Assessment of psychiatric illness 	information.	
		Schizophrenic disorders	3. Recognise, categorise and interpret	
		* Depression	medication-related problems and make	
		Bipolar disorders	appropriate interventions.	
		Anxietv	4. Formulate and apply patient-specific,	
		Obsessive compulsive disorders	evidence-based medication treatment	
		Pharmacological involvement in	plans.	
		ilities	late and impleme	
			plans (including non-medicine	
		Neurological Disorders	treatment) and assist the patient with	
		Enilopey eyadromos	implementation.	
		Epilepsy syridronies	6. Monitor and evaluate pharmacotherapy	
		Parkinson's disease		
		Alzheimer's disease	7 Document nations interventions in	
		* Stroke	ď	
		Multiple sclerosis	requirements	
		 Attention deficit hyperactivity disorder 		
		[28 credits]		

Master's Core Degree in Clinical Pharmacy (MPharm)	Exit Level Outcome 10: Optimise therapy for disorders related to the respiratory system. Range statement: The range of topics will include, but is not limited to, the following points: The assessment of pulmonary function Asthma	Assessment Criteria for Exit Level Outcome 10:	
inical larmacy IPharm)	Optimise therapy for disorders related to the respiratory system. Range statement: The range of topics will include, but is not limited to, the following points: The assessment of pulmonary function Asthma		200
IPharm)	respiratory system. Range statement: The range of topics will include, but is not limited to, the following points: The assessment of pulmonary function Asthma		
Pharm)	Range statement: The range of topics will include, but is not limited to, the following points: The assessment of pulmonary function Asthma	1. Define, discuss and appraise	
	include, but is not limited to, the following points: The assessment of pulmonary function Asthma	pathophysiology of the disorders related	
	points: The assessment of pulmonary function Asthma		
	I ne assessment or pulmonary functionAsthma	 Appraise, organise and evaluate patient information. 	
	* Asthma		
		3. Recognise, categorise and interpret	-
	 Chronic obstructive pulmonary disease 	medication-related problems and make	
	Drug-induced lung disease	appropriate interventions.	
	Pulmonary hypertension	4. Formulate and apply patient-specific,	
	Occupational pulmonary diseases	evidence-based medication treatment	
		plans.	
		5. Formulate and implement treatment	
	[stipera CC]	plans (including non-medicine	
		treatment) and assist the patient with	
		implementation.	
		6. Monitor and evaluate pharmacotherapy	
		to assess patient response.	
		7. Document patient interventions in	
		accordance with professional and legal	
		requirements.	

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's Degree in	Core	Exit Level Outcome 11:	Assessment Criteria for Exit Level Outcome	1200
Clinical Pharmacy		Conduct research and prepare for publication in the field of clinical pharmacy.	1. Critically evaluate information sources,	
(MPharm)		Range statement: The range of topics will encompass any suitable postgraduate	literature and research on medicines and practices in terms of evidence for	
		tudy İr	decision-making and implementation in	
		pharmacy.		
			z. Apply the principles of research methodology in the development of a	
		[120 credits]	research protocol. Obtain ethical	
			clearance if necessary.	
			3. Conduct research in accordance with	
			established research methodology and	
			ethics, as well as Good Clinical Practice	
			where necessary.	
			4. Analyse data, interpret findings and/or	
			results and formulate conclusions and	
			recommendations.	
			5. Write and submit a technical report,	
			manuscript for publication or minor	
			dissertation based on the research	
			outcomes and obtain approval.	

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's Degree in	Elective	Exit Level Outcome 12:	Assessment Criterion for Exit Level Outcome 12:	160
Clinical		Optimise therapy for disorders and/or		
Pharmacy		optimise clinical pharmacy practice related	Demonstrate extensive knowledge of the	
(MPharm)		to any one (1) chosen elective topic.	chosen elective field of clinical pharmacy,	
		natige statement. The range of topics may include, but is not limited to, the following	of transition to independent practice.	
		selected examples:		
		* Paediatrics		
		Clinical drug development		
		Critical care in adults		
		Oncological pharmacy		
		 Pharmacovigilance 		
		Dermatology		
		Geriatrics		
		Pharmacogenomics		
		 Pharmacoeconomics 		
		[16 credits]		

12. CRITICAL CROSS-FIELD OUTCOMES:

The following critical cross-field outcomes will form an integral part of the exit level outcomes of this programme:

- П Identify and solve problems in which responses display that responsible decisions using critical and creative thinking have been made; Work effectively with others as a member of a team, group, organisation and community; П Organise and manage oneself and one's activities responsibly and effectively; Collect, analyse, organise and critically evaluate information; Communicate effectively using visual, mathematical and/or language skills in the modes of oral and/or written persuasion; Use science and technology effectively and critically, show responsibility towards the environment and health of others; Demonstrate an understanding of the world as a set of related systems by recognising that problem-solving contexts do not exist in isolation; П Promote the personal and professional development of each learner in the programme, and the social and economic development of the society at large, by creating an awareness of the importance of:
 - reflecting on and exploring a variety of strategies to learn more effectively;
 - participating as responsible citizens in the life of local, national and global communities;
 - being culturally and aesthetically sensitive across a range of social contexts;
 - exploring education and career opportunities; and
 - developing entrepreneurial opportunities.

13. INTERNATIONAL COMPARABILITY:

The following examples are provided to illustrate the proposed curriculum's competitiveness and comparability among both developed and developing countries.

Africa: Example Ethiopia

An Eastern African country classified as a low income country with a high prevalence of infectious diseases. Ethiopia has similar health-related issues to South Africa and is also experiencing an epidemiological transition with diabetes, hypertension and other cardiovascular diseases. ¹³

As in South Africa, Ethiopia is experiencing a shortage of pharmacists and pharmacy support staff, and has also identified the need for pharmaceutical care services to meet and respond to healthcare needs. In 2009 the School of Pharmacy of Jimma University launched its first postgraduate programme, a Master of Science in Clinical Pharmacy. The initiative was started as part of a partnership with the Ethiopian Pharmaceutical Association, Strengthening Pharmaceutical Systems Programme of Management Sciences for Health, and the University of Washington.

The following points pertain to Ethiopia's training and education:

- Undergraduate and postgraduate students and faculty members are trained in South Africa, an emphasis on introducing clinical pharmacy in undergraduate programmes to train true specialists has been made, and postgraduate programmes similar to this programme exist.
- The objectives and priority areas of their Master's Degree include a strong emphasis on pharmaceutical care and pharmacovigilance in South Africa pharmaceutical care is

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¹³ Odegard PS, Tadeg H, Downing D, Suleman S, Bedada W, Paulos, G, Mekonnen H, Negussu M, Barlein R and Stergachis A. Strengthening Pharmaceutical Care Education in Ethiopa through collaboration. American Journal of Pharmaceutical Education. 2011;75 (7)134

a core component of the curriculum. However, due to the disease burden in South Africa strong emphasis is placed on a disease-driven Master's Degree. Pharmacovigilance is included as an elective. Pharmacovigilance is part of the Master's Degree in Medicines Management. Other courses are available.

Europe: Example Germany

In Europe, general specialisation towards clinical pharmacy follows the same route as that proposed in South Africa – a basic BPharm degree (in some countries it is three years, in others four years), followed by an internship plus a final examination by a competent entity. Formal education in a university, with a practical component, allows entrance to a specialisation. Special degrees for pharmacists are possible after advanced training of at least three years in specialities such as clinical pharmacy (as proposed in this document). Clinical pharmacy, as part of the basic undergraduate programme in Germany, has also been included in recent years – as in South Africa.¹⁴

United States of America (USA)

Pharmacy practice has shifted to include more clinical services. This has been supported in the schools of pharmacy by education and training. The USA has moved from a Bachelor of Science in Pharmacy to a Doctor of Pharmacy, with additional years of training – from four years of training (Bachelor of Science in Pharmacy) to a minimum of six years of training.

The core curriculum is comparable with the subjects presented in a Master's Degree in Clinical Pharmacy, as the basic pharmacy degree presented in South Africa is still very reliant on natural sciences. The Doctor of Pharmacy core curriculum includes:

□ Pathophysiology
□ Pharmacology

☐ Clinical problem solving☐ Laboratory monitoring

Therapeutics

☐ Physical assessment skills for many diseases.

This curriculum is supported by practical clinical rounds with medical students accompanying physicians. The latest curricular guidelines from the Accreditation Council for Pharmacy Education (ACPE) mandates early pharmacy practice experience through training/shadowing in a physician's office and clinical hospital setting, exposing the pharmacist student to collaborative practice environments. Master's and Doctoral level training are still being undertaken in clinical pharmacy, with additional residencies to encourage advanced level practice and specialisation, similar to the proposal in this document.

Clinical speciality certifications, endorsed by the Board of Pharmacy Specialties (BPS), are available for pharmacists in the following areas:

 · · · · · · · · · · · · · · · · · · ·
Pharmacotherapy Specialist (BCPS)
Nuclear Pharmacist (BCNP)
Nutrition Support Pharmacist (BCNSP)
Oncology Pharmacist (BCOP)
Psychiatric Pharmacist (BCPP)
Ambulatory Care Pharmacist (BCACP)

The board certification is NOT required for pharmacists, and is different from that required for specialist physicians – this is different from the proposed South African degree outlined in this document.¹⁵

¹⁴ Buschauer A. Pharmacy Education in Germany. Presentation. 2011.

¹⁵ Office of the Chief Pharmacist. Improving Patient and Health System Outcomes through Advanced Pharmacy Practice: A Report to the U.S. Surgeon General. 2011.

Australia

The principal pharmacy degree in Australia remains the four year Bachelor of Pharmacy, but some universities have started offering an entry level Master's Degree in Pharmacy. The curricula for the two degrees remain the same and the one has no advantage/disadvantage over the other. As in South Africa, no pharmacy school currently offers a Doctor of Pharmacy (PharmD) as the entry point for registration as a pharmacist. However, clinical pharmacy is being offered as part of postgraduate degrees such as the Master of Clinical Pharmacy, Doctor of Clinical Pharmacy, graduate diploma or graduate certificate awards, in addition to research degrees such as research master's and PhD degrees. The emphasis in the curricula for these postgraduate programmes in clinical pharmacy differs from institution to institution. What is similar to the proposal in this document is that they generally include components of therapeutics and, at the master's level and beyond, completion of a practice-based research project.¹⁶

India

The Master of Pharmacy in Clinical Pharmacy is offered as a two-year degree programme by seventeen institutions across India. Students who have passed their Bachelor of Pharmacy are eligible to enrol. In India, clinical pharmacy is the branch of pharmacy in which pharmacists provide patient care that optimises the use of medicines and promotes health, wellness and disease prevention. The degree is designed to prepare pharmacists for expanded roles as providers of direct patient care with emphasis on physiology, applied therapeutics and pharmacy practice skills.

The subjects in the first year include clinical pharmacy practice, clinical pharmacokinetics, pharmacotherapeutics, biostatistics and research methods. Second year studies include a research project, general medicine clerkship and biotechnology. Specialisation areas include clinical trials, new medicine discovery and hospital pharmacy.

The role of a clinical pharmacist in India is to support and provide the best quality medicine therapy for patients. This role may include:

- Prescription monitoring to maximise medicine efficiency, minimise medicine toxicity and promote cost effectiveness;
- Therapeutic medicine monitoring of medicines with narrow therapeutic index;
- Medicine information services:
- Patient services and counselling;
- Improving patient compliance through collecting past medical history; and
- Offering recommendations to the physician for an optimised medical treatment that is completely patient oriented.¹⁷

14. INTEGRATED ASSESSMENT:

A combination of integrated assessment strategies, which will combine both formative and summative assessment and evaluation, will be used to ensure that the purpose of the qualification is achieved. Assessments may include, but are not limited to, the following strategies:

og.co.
Portfolios of evidence
Simulations, role play and workplace assessments
Written and oral assessments and examinations
Written assignments

¹⁶ Marriot JL, Nation RL, Roller L, Costelloe M, Galbraith K, Stewart P & Charman WN. Pharmacy Education in the Context of Australian Practice. American Journal of Pharmaceutical Education. 2008;72 (6):131

^{&#}x27;Viswanad V, Prabhakar V. 2011. The emergence of the clinical pharmacist and the Indian scenario. Inventi Rapid: Pharmacy Practice 2, (1). Published on Web 21/02/2011, www.inventi.in

		Case studies Journal clubs Self-assessment strategies, peer-group assessment and preceptor evaluation Objective structured clinical examination (OSCEs).
15.	CRE	DIT ACCUMULATION AND TRANSFER:
		didates may apply for recognition of credits obtained as part of an incomplete fication at the same or a different institution, depending on individual institutional ies.
16.	ART	ICULATION (PROGRESSION):
		pletion of a Professional Master's Degree meets the minimum entry requirement for ission to a Doctoral Degree, usually in the area of specialisation in the Master's Degree.
		ulation may also be horizontal to entries into other Master's Degrees in a similar or ed field or area of specialisation.
17.	MOE	DERATION OPTIONS:
	provi Educ	able moderating options should be included in each application for accreditation to ide this qualification in accordance with the stipulations of the Council on Higher cation (CHE), as well as the relevant ETQA (i.e. Council). Both internal and external eration should form an integral part of the provision of this qualification.
18.	CRIT	TERIA FOR THE APPOINTMENT OF ASSESSORS:
	track	essors in the field of clinical pharmacy must have a suitable background with a proven record and relevant experience to enable them to make sound judgements through expert application of the assessment criteria specified for this qualification.
19.	NOT	ES:
		All candidates must, in addition to their current registration as academic interns or pharmacists, be registered with Council for study towards the specialisation for the duration of the period of learning as specified in current relevant legislation.
		The range of elective learning areas offered will be dependent on the approval of the provider and ETQA.
		Credit values reflected for each exit level outcome in Table 2 should be regarded only as a guideline.
		The respective assessment criteria aim to test the achievement of the specific learning outcomes. Some of these criteria are practice-based, thus providers are required to
		include periods in their curricula for this purpose. After attaining the professional Master's Degree, the candidate may commence with the process for registration as a specialist pharmacist in clinical pharmacy with Council.

(See Appendix A)

Requirements for this registration process will be determined by Council.

APPENDIX A

Requirements for registration as a specialist after obtaining the professional Master of Pharmacy in Clinical Pharmacy

The prospective candidate should be a registered pharmacist with Council.

Training Site

A site registered with Council as a training institution, pharmacy or health facility where clinical pharmacy is routinely practised.

Tutor or supervisor

A registered clinical pharmacist or postgraduate pharmacist tutor/trainer with at least two years' experience in clinical pharmacy.

Practical training

As stipulated by Council.

Evaluation and panel

As stipulated by Council

PUBLIC HEALTH PHARMACY AND MANAGEMENT

SCOPE OF PRACTICE - PUBLIC HEALTH PHARMACY AND MANAGEMENT

- (a) Perform acts and services specially pertaining to the profession of a pharmacist;
- (b) Lead and manage surveillance and assessment of the pharmaceutical services;
- (c) Lead projects to protect and promote health and wellbeing, including communicable disease control and environmental health;
- (d) Manage, analyse and interpret information and statistics;
- (e) Develop and analyse pharmaceutical public health policy for the better use of existing and new medicines/technologies and rational use of all medicines, to improve health services;
- (f) Provide strategic leadership for medicine supply management;
- (g) Provide education and training related to public health and management;
- (h) Manage knowledge and transfer research evidence into practice;
- (i) Develop policies and procedures for public health and management;
- (j) Manage, analyse, interpret and advise on pharmacoeconomic information for rational use of medicines; and
- (k) Perform research, teach and publish in the field of public health and management.

QUALIFICATION: PROFESSIONAL MASTER'S DEGREE IN PUBLIC HEALTH PHARMACY AND MANAGEMENT

SYNOPSIS:

To provide a curriculum for a professional Master's Degree in Public Health Pharmacy and Management to enable students to register with the South African Pharmacy Council (hereafter referred to as Council) as specialists with a post-qualification that complies with Council's requirements.

Table 1: Summary of the proposed qualification

	Professional Master of Pharmacy in Public Health Pharmacy and Management
Duration:	Two years
Entry criteria:	Bachelor's Degree in Pharmacy (NQF Level 8)
NQF-level:	Level 9
Field:	09 Health Sciences and Social Services
Sub-field:	Curative Health
SAQA-credits:	360 credits
Qualification type:	Professional, exit level, career-orientated, whole qualification
Final assessment and evaluation:	 Final, exit level examination(s) will need to be passed in accordance with the relevant Higher Education provider's rules and regulations. In addition, a comprehensive portfolio of evidence will need to be submitted and successfully passed by the accredited provider. Requirements for registration as a specialist after obtaining the Professional Master's Degree will be determined by Council.
CPD requirements for annual reregistration:	As required by Council
Professional status:	Registration with Council as a Pharmacist Specialist in Public Health Pharmacy and Management
Articulation	DPharm /Doctoral Degree

QUALIFICATION OUTLINE:

1. Qualification Title:

Master of Pharmacy in Public Health Pharmacy and Management

Abbreviation: MPharm (Public Health Pharmacy and Management)

2. Qualification Type:

Professional Master's Degree

3. Field and Subfield:

Field: [09] Health Sciences and Social Services
Subfield: Curative Health

4. Level:

NQF/HEQF Level 9 (Master's Degree)

5. Credits:

Total credits: 360

6. Rationale for the Qualification:

There is a need in South Africa, especially with the implementation of the National Health Insurance (NHI) and the re-engineering of primary healthcare, for pharmacists to have the necessary skills and expertise to implement public health standards and management principles in the delivery of pharmaceutical services to the population. This need is in line with current local and international efforts to stop the increase of chronic non-communicable diseases. The commitment of pharmacists to combat non-communicable diseases (including cardiovascular diseases, diabetes, chronic respiratory diseases and cancers) was noticeably evident in 2011 when the Durban Declaration was issued at a conference jointly hosted by the Commonwealth Pharmacists' Association, the Pharmaceutical Society of South Africa and the South African Pharmacy Council (SAPC).¹⁸

Internationally, the professional practice role of the pharmacist in public health, and the increasing contribution pharmacists should be making to the provision of public health, has been highlighted.

Postgraduate specialisation for pharmacists in pharmaceutical public health was advocated at the 9th International Conference on Life Long Learning in Pharmacy in New Zealand in 2011.¹⁹ Having specialist public health pharmacists within the profession would be in line with the progressive move away from the main role of the pharmacist as a dispenser or distributor of medicine towards a more patient-oriented focus.

In March 2014, the Royal Pharmaceutical Society in the United Kingdom published *Professional Standards for Public Health Practice for Pharmacy*,

 $^{^{\}rm 18}$ Commonwealth Pharmacists' Association. The Durban Declaration. S Afr Pharm J. 2011; 78:10.

¹⁹ Shaw.J. 20:20 Vision Focusing on the Future of Pharmacy Practice and Education. School of Pharmacy. Life Long Learning in Pharmacy Conference, Rotorua, New Zealand, 29 June 2011.

which sets out a best practice framework for the delivery of public health services across all pharmacy settings in England and Wales.²⁰ These standards emphasise the pharmacy profession's integral part in public health and the public health workforce aimed at delivering a service that will improve the health and wellbeing of the public health community.

The professional Master's Degree in Public Health Pharmacy and Management, NQF Level 9, was developed to meet the requirements of Council, the statutory body for the pharmacy profession, for specialists in pharmacy with specific reference to public health pharmacy and management. With this qualification, specialist pharmacist training is aligned to the needs of the health system and will contribute to capacity building for better management of pharmaceutical services, provide for professional recognition of the pharmacist's role in public health activities and preserve pharmacists, as a scarce resource, in South Africa.

'Public health' is defined as the science and art of promoting and protecting health and wellbeing, preventing ill-health and prolonging life through the organised efforts of society.²¹ The World Health Organisation (WHO)²² further states explicitly that public health refers to all organised measures, whether **public or private**, to prevent disease, promote health, and prolong life among the population as a whole. Public health activities are therefore aimed at improving health for entire populations and not only individual patients or a particular disease. The WHO⁵ and the Royal Pharmaceutical Society³ identified three main public health functions or domains. The pharmacy profession has a role to play across all three:

- Health protection which entails the assessment and monitoring of the health of communities and populations at risk to identify health problems and priorities. This includes infectious diseases, environmental hazards and emergency preparedness.
- Health service delivery and quality, including service planning, efficiency, audit, evaluation and the formulation of public policies designed to solve identified local and national health problems and priorities.
- Health improvement, which includes health promotion and disease prevention services, to ensure that all populations have access to appropriate and cost-effective care.

'Public health pharmacy' and **'pharmaceutical public health'** are terminology used commonly to describe the role or involvement of the pharmacist in public health. Pharmaceutical public health has been defined as the application of pharmaceutical knowledge, skills and resources to the science and art of preventing disease, prolonging life, promoting, protecting and improving health for all through organised efforts of society.²³

The focus of pharmaceutical public health is on the development of pharmacy services and expertise to enhance the health and wellbeing of a whole population. This definition does not however cover all the key aspects and potential roles of pharmacists in public health, categorised previously as micro-

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²⁰ Royal Pharmaceutical Society. Professional Standards for Public Health Practice for Pharmacy. March 2014.

²¹ Adapted from the original definition in the Public Health in England report by Sir Donald Acheson, 1988. In: Royal Pharmaceutical Society, Professional Standards for Public Health Practice for Pharmacy, March 2014.

²² World Health Organisation (WHO). Public health. Trade, foreign policy, diplomacy and health. Available from: http://www.who.int/trade/glossary/story076/en/.

²³ Walker R. Pharmaceutical public health: the end of pharmaceutical care? Pharmaceutical Journal. 2000; 264:340-341.

and macro-level activities.²⁴ **Micro-level activities** focus on individual health promotion and disease prevention services, while **macro-level activities** comprise population-wide approaches, including policy formulation, planning and management functions.²⁵ The specialist qualification in public health pharmacy and management will predominantly be appropriate for pharmacists involved in macro-level activities in the public and private sectors.

This professional Master's Degree is designed to meet the needs of pharmacists who have completed the BPharm degree and who wish to further their competencies in the field of pharmaceutical services and develop their careers in the public health pharmacy and management practice area. An increase in the number of pharmacists with specialised knowledge in public health pharmacy and management will contribute to capacity building in this field, for the overall development of healthcare in South Africa. Pharmacists with this qualification will practise at a higher level and be senior in the health system. Positions of practice would include, for example, pharmaceutical management in the public sector (facility pharmacy manager, sub-district or district pharmacist, provincial head office, policy or human resources, medical depot, national level positions), academia (especially pharmacy practice or public health), private sector (medical aid, community pharmacy), non-profit organisation (USAID-funded, MSH, MSF), general management positions in health systems or hospital management (public, private, non-profit) and public private partnerships (part of non-governmental organisations (NGOs) and NHI).

The development and introduction of this qualification and curriculum outline will assist higher education institutions in the training of these specialist pharmacists who can register with Council as specialists in public health pharmacy and management. Although the sub-field for this qualification at present is listed as curative health, it also includes preventative health, health promotion, health education, environmental health and occupational health.

7. Purpose:

The purpose of this professional **Master's Degree** is to extend the public health and pharmaceutical management competencies of pharmacists to become **specialists in the field of public health pharmacy and management**, apply their expertise in this field and add value to the provision of pharmaceutical services within the health system. Successful completion of this qualification will enable specialist pharmacists to contribute to public health outcomes and pharmaceutical services management. The degree is inherently a practice-based degree with a large component of work-integrated learning.

Council's Scope of Practice for Public Health Pharmacy and Management Pharmacists

- 1. Perform acts and services specially pertaining to the profession of a pharmacist.
- 2. Lead and manage surveillance and assessment of the pharmaceutical services.
- 3. Lead projects to protect and promote health and wellbeing, including communicable disease control and environmental health.

²⁴ Rappaport H, et al. Assessment of realistic public health roles for pharmacists. Journal of Social and Administrative Pharmacy. 1984; 2(2):57-66.

²⁵ Bradley H, Sanders D, Bheekie A. Public health: every pharmacist's business! S Afr Pharm J. 2011; 78(10):34-36.

- 4. Manage, analyse and interpret information and statistics.
- 5. Develop and analyse pharmaceutical public health policy for the better use of existing and new medicines/technologies and rational use of all medicines, to improve health services.
- 6. Provide strategic leadership for medicine supply management.
- 7. Provide education and training related to public health and management.
- 8. Manage knowledge and transfer research evidence into practice.
- 9. Develop policies and procedures for public health and management.
- Manage, analyse, interpret and advise on pharmacoeconomic information for rational use of medicines.
- 11. Perform research, teach and publish in the field of public health and management.

8. RULES OF COMBINATION:

Fundamental credits:	64
Core credits:	264
Elective credits:	32
Total:	360

9. ACCESS TO THE QUALIFICATION:

The minimum admission requirement is a four-year Degree in Pharmacy (NQF level 8) or equivalent, registration with Council as a pharmacist post-community service, and placement in the area of specialisation.

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10. LEARNING ASSUMED TO BE IN PLACE:

A TO	ur-year Degree in Pnarmacy (NQF level 8) assuming the following is in
place	9:
	Professional and ethical practice
	Communication (collaboration with members of the healthcare team) and self-management
	Optimal use of medicines (therapeutic decision-making) and medication management
	Basic knowledge of healthcare and pharmaceuticals management
	Pharmacology
	Research methodology.

Candidates have to comply with all of the theoretical requirements set by Council for registration as a specialist pharmacist as well as annual registration, and must have a thorough understanding and working knowledge of South African *Good Pharmacy Practice* rules.

11. EXIT LEVEL OUTCOMES AND THEIR ASSOCIATED ASSESSMENT CRITERIA:

See Table 2

Table 2: Curriculum Outline

Learning Area	Exit Level Outcome	Credits	Notional Hours
Fundamental	Exit Level Outcome 1: Practise as a specialist pharmacist within the regulatory and policy framework of public health pharmacy	28	280
Fundamental	Exit Level Outcome 2: Apply basic epidemiology and biostatistics in disease prevention, health promotion, healthcare delivery and policy development	36	360
Core	Exit Level Outcome 3: Apply strategic management and leadership to ensure an effective and efficient health system	36	360
Core	Exit Level Outcome 4: Implement the concepts and principles of public health to protect and promote general health and wellbeing	98	360
Core	Exit Level Outcome 5: Provide strategic leadership for pharmaceuticals management in the health system	36	360
Core	Exit Level Outcome 6: Design and implement strategies for the rational use of pharmaceuticals to improve health services	36	360
Core	Exit Level Outcome 7: Conduct research and prepare for publication in one of the specialisation fields of public health pharmacy and management	120	1200
Elective	Exit Level Outcome 8: Deepen knowledge of work in research interest area for transition to independent work in public health pharmacy and management	32	320
MPharm (Public Health Pharmacy and Management)	TOTAL	360	3600

Notional Hours	280 49
Associated Assessment Criteria	1. Interpret and explain the sources of South African law, how it is developed and the interrelationship between the constitution, legislation and the functioning of the courts. 2. Identify, analyse and interpret relevant legislation and policy in the delivery and management of public health pharmacy services. 3. Examine and evaluate the implementation of National Health Insurance within the regulatory and policy framework of public health. 4. Analyse the Bill of Rights (equity), Patient Rights Charter and Batho Pele Principles and appraise their application to the health sector. 5. Appraise and apply the process of development and amendment of legislation and policies. 6. Identify, appraise and obligations within an ethical framework in providing optimal care to communities.
Exit Level Outcomes	Exit Level Outcome 1: Practise as a specialist pharmacist 1. within the regulatory and policy framework of public health pharmacy. [28 credits] 3. 5.
Learning Area	Fundamental
Specific Exit Level	Master's Degree in Public Health Pharmacy and Management (MPharm)

Notional Hours	996
Associated Assessment Criteria	Assessment Criteria for Exit Level Outcome 2: 1. Critically evaluate and use the common causes of death, disease and disability in a particular community in the planning and design of health programmes. 2. Identify, analyse and evaluate the main determinants of health for potential implementation into health policy and health services. 3. Conduct and interpret a community health needs assessment to plan healthcare and public health programmes. 4. Apply the principles and methods of epidemiology in public health. 5. Use epidemiological data to appraise the effectiveness and efficiency of healthcare delivery. 6. Design appropriate studies to determine causes of disease, prognosis, prevention and the evaluation of therapy. 7. Design and demonstrate the ability to implement in practice interventions to prevent and control disease. 8. Apply key biostatistical concepts and methods to summarise, display, evaluate and interpret medical and healthcare data.
Exit Level Outcomes	Apply basic epidemiology and biostatistics in disease prevention, health promotion, healthcare delivery and policy development. [36 credits]
Learning Area	Fundamental
Specific Exit Level	Master's Degree in Public Health Pharmacy and Management (MPharm)

Specific Exit Level	Learning Area	Exit Level Outcomes	Associated Assessment Criteria	Notional Hours
Master's	Core	Exit Level Outcome 4:	Assessment Criteria for Exit Level Outcome 4:	360
Degree In Public Health		Implement the concepts and principles	1. Outline and appraise the context of the public	
Pharmacy and		of public health to protect and promote	health environment.	
Management		general health and wellbeing.	 Critically explore and all apply the design, 	
(MPharm)			implementation, evaluation and review of public	
			health policies and procedures. 4 Explain and evaluate the application of the	
			_	
			levels of pharmaceutical service delivery.	
			5. Demonstrate the ability to develop public health	
			policies for the management and rational use of	
			medicines to improve health services.	
			6. Analyse policy instruments for the delivery of	
			pharmaceutical services.	
			7. Compile a policy and procedure manual for the	
			healthcare organisation.	
			8. Demonstrate the ability to implement policy	
			instruments and a policy and procedure manual.	
			9. Design and implement screening services for	
			health promotion.	
			10. Apply social, psychological and behavioural	
			aspects in health promotion, education and the	
			design of interventions for the health and	
			wellbeing of the community.	
			11. Apply and appraise the principles of cold chain	
			management and immunisation according to	
			required standards.	
			12. Design and use surveillance tools to collect	
			information on community health.	

Assessment Criteria for Exit Level Outcome 5:
strategic leadership for the 1. Evaluate and critically appraise access to
management of pharmaceuticals in the health system.
concept in the selection of medicines for essential medicines lists.
3. Analyse and implement the framework and
components of
4. Utilise a health management information
system for decision-making and to improve access to pharmaceuticals.
5. Appraise
management principles
6. Demonstrate the ability to manage and
develop human resources
7. Demonstrate the ability to implement
quality and risk management programme for effective pharmaceutical supply and
nse.
8. Design tools to monitor and evaluate the
supply chain system and provide feedback

Notional Hours	360
Associated Assessment Criteria	Assessment Criteria for Exit Level Outcome 6: 1. Evaluate the use of pharmaceuticals within the medicines management cycle and health system. 2. Appraise and enhance rational drug use through implementation of and participation in all the activities of the pharmacy and therapeutics committee by all stakeholders. 3. Distinguish and apply the different types of costs in pharmacoeconomic analysis. 4. Appraise and correctly apply the appropriate pharmacoeconomic tools to conduct analyses for the rational use of pharmacoeconomic analyses and decision-making. 5. Critique pharmacoeconomic analyses and decision-making. 6. Construct a simple model for pharmacoeconomic evaluation and decision-making. 7. Identify and analyse priorities for rational drug use interventions and design strategies for interventions. 8. Demonstrate the ability to implement and monitor drug use interventions. 9. Design, apply and evaluate programmes for quality assurance of medicines use (e.g. adherence, medicine safety, medication errors). 10. Design pharmacovigilance and surveillance programmes for patient safety. 11. Demonstrate the ability to implement pharmacovigilance, surveillance and quality assurance programmes.
Exit Level Outcomes	Exit Level Outcome 6: Design and implement strategies for the rational use of pharmaceuticals to improve health services.
Learning Area	Core
Specific Exit Level	Master's Degree in Public Health Pharmacy and Management (MPharm)

Notional Hours	1200
Associated Assessment Criteria	1. Critically evaluate information sources, literature and research on medicines and practices in terms of evidence for decision-making and implementation in practice. 2. Apply the principles of research methodology in the development of a research protocol and obtain ethical clearance. 3. Conduct research in accordance with established research methodology and ethics, as well as <i>Good Clinical Practice</i> where necessary. 4. Analyse data, interpret findings and/or results and formulate conclusions and recommendations. 5. Write and submit a technical report, manuscript for publication or minor dissertation and obtain approval.
Exit Level Outcomes	Exit Level Outcome 7: Conduct research and prepare for publication in one of the specialisation fields of public health pharmacy and management. [120 credits]
Learning Area	Core
Specific Exit Level	Master's Degree in Public Health Pharmacy and Management (MPharm)

O Company O
<u>Exit Level Outcome 8:</u>
Deepen knowledge of work in an appropriate interest area from the
options in the range statement.
for electives may include, but is not
limited to, the following examples:
Pharmaceutical policy
macoeconomi
stics managem
<u>~</u>
macovigilance
Health promotion
entative health
Programme on Immunisation (EPI)
[32 credits]

12. CRITICAL CROSS-FIELD OUTCOMES:

The following critical cross-field outcomes will form an integral part of the exit level outcomes of this programme:

- П Identify and solve problems in which responses display that responsible decisions using critical and creative thinking have been made; Work effectively with others as a member of a team, group, organisation and community; Organise and manage oneself and one's activities responsibly and effectively; Collect, analyse, organise and critically evaluate information; Communicate effectively using visual, mathematical and/or language skills in the modes of oral and/or written presentation; Use science and technology effectively and critically, showing responsibility towards П the environment and health of others: Demonstrate an understanding of the world as a set of related systems by recognising that problem-solving contexts do not exist in isolation; П Promote the personal and professional development of each learner in the programme, and the social and economic development of the society at large, by creating an awareness of the importance of:
 - reflecting on and exploring a variety of strategies to learn more effectively;
 - participating as responsible citizens in the life of local, national and global communities;
 - being culturally and aesthetically sensitive across a range of social contexts;
 - exploring education and career opportunities; and
 - developing entrepreneurial opportunities.

13. INTERNATIONAL COMPARABILITY:

The professional Master's Degree in Public Health Pharmacy and Management has been designed and generated with the standards and guidelines as displayed in the qualifications being offered by institutions in South Africa, Tanzania, Australia, the United States of America and the United Kingdom.

Although these countries offer training in pharmacy administration, public health and management, the training is not identical to the qualification proposed in this document. Certain courses or modules offered by the programmes are comparable and were therefore used for benchmarking. The following examples are provided to illustrate the proposed curriculum's competitiveness and comparability among both developed and developing countries.

Africa: South Africa

The Master of Public Health (MPH) is offered by a number of institutions in South Africa – University of Pretoria, University of the Witwatersrand, University of Limpopo (Medunsa Campus), University of the Western Cape, University of Cape Town and the University of KwaZulu-Natal. The general focus of the MPH programme is to prepare professionals for leadership roles in the evaluation of health, health interventions, management and the healthcare system. Similar to the qualification proposed in this document, "public health" refers to the health of entire populations and is not limited to public sector health. Another similarity is that the MPH is a practice-oriented degree and not a research degree.

The MPH exposes students to different disciplines. However, the field of health systems and public health is very wide and students cannot become a specialist in all its aspects, which is also the situation with the proposed qualification. The MPH programmes are structured in a way that students will acquire a good understanding of the entire field of health systems and

public health, but they select one particular track or focus area in which they will develop detailed and sufficient competence.

The course content of the various MPH programmes examined are different, however there are certain topics covered by most MPH programmes that are also covered, to a certain extent, by the qualification proposed in this document.

Com	parable modules or courses include the following topics:
	Epidemiology and Biostatistics
	Health Policy and Management
	Environmental and Occupational Health
	Disease Control
	Health Research Ethics
	Health Promotion
	Financial Management in the Public Sector
	Project Management
Africa: Tanzania	

The Muhimbili University of Health and Allied Sciences (MUHAS), Dar-es-Salaam, Tanzania, offers two degrees, of which some of the components are comparable with the proposed qualification.

The School of Pharmacy offers a Master of Science in Pharmaceutical Management, which is a four-semester degree programme, each semester consisting of 24 weeks.

The degree contains a dissertation comprising 45% of the total credits for the degree. This is similar to the coursework of the South African general Master's Degree, which contains a research project comprising a minimum of 60 credits at NOF Level 9, and culminates in a mini-dissertation, technical report, one or more creative performances or works, or a series of peer-reviewed articles or other research-equivalent outputs.

The o	degree programme contains the following courses:
	Bioethics
	Epidemiology & Research
	Healthcare Delivery and Pharmaceutical Regulatory Framework
	General Management
	Financial Management
	Educational Principles and Practices for the Health Sciences Professionals
	Pharmaceutical Supply Chain Management
	Managing Rational Use of Medicines
	Drug and Commodity Management in Health Facilities
	Pharmaceutical Marketing
	Fieldwork in Pharmaceutical Management
П	Dissertation

The School of Public Health and Social Sciences offers a Master of Public Health (MPH) Executive Track, which is a modular programme. The aim of the programme is to train candidates to become public health specialists in government and non-governmental organisations (NGOs) as well as national and international organisations. The Research Methods module constitutes 8.5% and the Dissertation 20% of the total credits of the MPH, which is in line with the Higher Education Qualifications Sub-Framework recommendation for professional Master's Degrees in South Africa.

The MPH programme contains the following courses, which illustrates the similarity with the qualification outlined in this document: Principles of Public Health Epidemiology and Biostatistics Implementing Change Special Public Issues Health Policy, Planning and Management Health Economics Financing and Evaluation Research Methods Dissertation
Australia: Queensland
The James Cook University School of Pharmacy offers a Master of Pharmaceutical Public Health over a period of two years. They define pharmaceutical public health as the development of pharmacy services and expertise to enhance the health and wellbeing of a whole population. The programme is designed to enable pharmacists to focus beyond the specific needs of individual patients and meet health goals for the whole community. The course is structured for pharmacists who want to learn the principles of public health and develop services in their different fields of practice. These principles are in line with the scope of practice for this qualification.
The course consists of the following three core subjects: □ Epidemiology for public health □ Management of pharmaceutical services □ An option between public health management and public health leadership and crisis management. For the degree to be awarded, students should complete a dissertation, the three core subjects (above) and additional elective subjects.
United States: Boston University
Boston University School of Public Health offers a Public Health Pharmaceuticals Programme for students to gain knowledge and expertise to address pharmaceutical issues from a public health perspective. The programme is offered at a master's level for students considering careers in the pharmaceutical industry, service delivery programmes, or pharmaceutical policy-making agencies. The pharmaceuticals programme prepares students for positions in both the public and private sectors, including positions in federal and state government agencies, the pharmaceutical industry, contracting research organisations and international agencies. Students are given a solid foundation in pharmaceuticals while providing flexibility to tailor their coursework towards a specific career path in policy, industry, or health programmes and non-governmental organisations.
The following three main tracks are offered, with a mandatory 'Pharmaceuticals in Public Health' course for all tracks: Policy track Health policy Health services research methods Pharmacovigilance Clinical trials Patent law Insurance systems Qualitative research methods

Indus	stry track Project management Good clinical practice Discovery and development Clinical trials Regulatory affairs Intellectual property	
Serv	ice delivery track Project management Infectious diseases Rational drug management and medication adherence Vaccines Corruption Qualitative research	
Unit	ed States: Virginia	
The School of Business, Virginia Commonwealth University, offers a combined Doctor of Pharmacy (PharmD) and Master of Business Administration (MBA). The programme is designed to take advantage of efficiencies and electives in both the PharmD and MBA programmes and seeks to prepare pharmacists for careers that encompass pharmacy and business theories and principles. Students in the combined programme can earn both degrees and save a year or more over the time required for enrolling in the programmes separately.		
found Man	btain both degrees, students need to take all the pharmacy courses, the seven business dation courses, the nine MBA core courses and three elective courses (see below). y of the topics covered in the MBA programme compare well with the topics covered in proposed qualification.	
Busiii	ness Foundation courses: Fundamentals of Accounting Concepts in Economics Financial Concepts of Management Statistical Elements of Quantitative Management Fundamentals of the Legal Environment of Business Management Theory and Practice Concepts and Issues in Marketing	
MBA	Managerial Economics Organisation Leadership and Project Team Management Financial Management Remainder of the Advanced Programme Managerial Accounting Information Systems for Managers Information Systems for Business Intelligence Business Policy Operations Management Marketing Management	

United Kingdom: Professional Standards for Public Health Practice for Pharmacy²⁶

The Professional Standards for Public Health Practice for Pharmacy, published by the Royal

Pharmaceutical Society in the UK, set out a best practice framework for the delivery of public health services. The standards are intended to provide a framework to help pharmacy teams, commissioners and those contracting services to design, implement, deliver and monitor high quality public health practice through pharmacy, regardless of the pharmacy settings from which services are delivered. The following nine key areas are covered by the standards: Surveillance and assessment of the population's health and wellbeing: Data are collected from a variety of sources to support a better understanding of the health and wellbeing needs of a population or community. Public health intelligence: Information and analysis of the health and wellbeing needs of the population or community are used to inform the development of pharmacy public health practice. Assessing the evidence of effectiveness of health and healthcare interventions, programmes and services: Population health is improved by the assessment and application of evidence-based public health interventions, programmes and public health services. Health improvement: Pharmacists and their teams improve the health and wellbeing of the population and help to reduce health inequalities by: proactively promoting health and wellbeing messages; supporting and enabling people to adopt healthier lifestyles and to take responsibility for their own and their family's health; and supporting the concept of self-care. Health protection: The population's health and wellbeing are protected by supporting the prevention and transmission of communicable and other infectious diseases, screening for risk factors and disease, ensuring prudent use of antibiotics in helping to mitigate the risks of antimicrobial resistance, protecting against pharmaceutical hazards, and supporting the pharmacy response to an emergency. Health and social service quality (also known as Healthcare Public Health): Innovative, high quality pharmacy public health services improve health outcomes and ensure fair and effective targeting of available resources. Policy and strategy development and implementation: Local and national policies and strategies are developed and implemented in accordance with local and national needs to improve and protect the health of the community or population. Strategic leadership and collaborative working for health: Pharmacists and their teams take the lead in ensuring pharmacy's contribution to public health is recognised strategically and collaboratively in partnership with other practitioners and agencies to improve and protect the health and wellbeing of populations, helping to reduce health inequalities. Academic public health: Everyone working in pharmacy has a role in contributing to the evidence base for the contribution of pharmacy in improving and protecting the health of the population. This is strengthened by academic research and pharmacy practice research across the profession.

Conclusion:

Although the Master of Pharmacy in Public Health Pharmacy and Management is a unique qualification and is geared towards meeting the specific needs of South Africa, it is evident that it compares favourably with modules or courses offered by postgraduate programmes internationally, as well as the Royal Pharmaceutical Society of the UK's recently published *Professional Standards for Public Health Practice for Pharmacy*.

²⁶ Royal Pharmaceutical Society. Professional Standards for Public Health Practice for Pharmacy. March 2014.

14. INTEGRATED ASSESSMENT:

A con	nbination of integrated assessment strategies, which will combine both formative and
summ	native assessment and evaluation, will be used to ensure that the purpose of the
qualifi	ication is achieved. Assessments may include, but are not limited to, the following
strate	gies:
	Portfolios of evidence

□ Portfolios of evidence
□ Simulations, role play and work-place assessments
□ Written and oral assessments and examinations
□ Written assignments
□ Case studies
□ Journal clubs

Self-assessment strategies, peer-group assessment and preceptor evaluation.

15. CREDIT ACCUMULATION AND TRANSFER:

Candidates may apply for recognition of credits obtained as part of an incomplete qualification at the same or a different institution, depending on individual institutional policies.

16. ARTICULATION (PROGRESSION):

Completion of a Professional Master's Degree meets the minimum entry requirement for admission to a Doctoral Degree, usually in the area of specialisation in the Master's Degree.

Articulation may also be horizontal to entries into other Master's Degrees in a similar or related field or area of specialisation. Horizontal articulation possibilities with this qualification include a Master of Public Health.

17. MODERATION OPTIONS:

Suitable moderating options should be included in each application for accreditation to provide this qualification in accordance with the stipulations of the Council on Higher Education (CHE), as well as the relevant ETQA (i.e. Council). Both internal and external moderation should form an integral part of the provision of this qualification.

18. CRITERIA FOR THE APPOINTMENT OF ASSESSORS:

Assessors in the field of public health pharmacy and management must have a suitable background, with a proven track record and relevant experience, to enable them to make sound judgements through their expert application of the assessment criteria specified for this qualification.

Assessors should be in possession of suitably related postgraduate qualification (i.e. Master's Degree and/or Doctoral Degree level) in public health pharmacy and Management, and/or other related fields of study, and have a good working knowledge of the higher education environment in South Africa.

19. NOTES:

All candidates must, in addition to their current registration as academic interns or
pharmacists, be registered with Council for study towards the specialisation for the
duration of the period of learning as specified in current relevant legislation.
The range of elective learning areas offered will be dependent on the approval of the
provider and ETQA.
Credit values reflected for each exit level outcome in Table 2 should be regarded only
as a guideline

The respective assessment criteria aim to test the achievement of the specific learning
outcomes. Some of these criteria are practice based, thus providers are required to
include periods in their curricula for this purpose.
After attaining the Master's Degree, the candidate may commence with the process for
registration as a specialist pharmacist in public health and management with Council.

(See Appendix A)

APPENDIX A

Requirements for registration as a specialist after obtaining the professional Master's degree in Public Health Pharmacy and Management

Requirements for this registration process will be determined by Council.

The prospective candidate should be a registered pharmacist with Council.

Training Site

A site recognised by or registered with Council as having the necessary scope to train specialist pharmacists in public health pharmacy and management at any of the levels of pharmaceutical services and public health (operational level, middle management or strategic management). Different rotation sites must be available for the candidate to gain experience in various fields of public health pharmacy and management.

Tutor or supervisor

An appropriately trained and qualified person, with extensive experience in the fields of public health pharmacy and management, jointly approved by the training institution and SAPC.

Practical training

As stipulated by Council.

Evaluation and panel

As stipulated by Council.

PHARMACOKINETICIST

SCOPE OF PRACTICE - PHARMACOKINETICIST

- (a) Perform acts and services specially pertaining to the profession of a pharmacist Provide pharmacokinetic consultations to a variety of specialities;
- (b) Take a leading role in identifying drug groups that require pharmacokinetic monitoring due to their increased risk of toxicity, narrow therapeutic index and challenges with adherence or efficacy;
- (c) Take a leading role in identifying patients that need a pharmacokinetic consultation due to a high risk of non-adherence, experiencing adverse drug reactions or drug toxicity due to their specific condition or diagnosis;
- (d) Take a leading role in requesting and interpreting drug concentrations in patients at risk;
- (e) Act as a leading pharmaceutical partner within a multi professional health care team;
- (f) Appraise information, make informed decisions with the drug concentrations available and be able to justify/defend the decisions;
- (g) Advise on dosing adjustment and patient monitoring based on drug concentrations;
- (h) Take a leading role in pharmacokinetic input for clinical protocol and guideline development;
- (i) Provide education and training related to pharmacokinetics; and
- (j) Perform research and publish in pharmacokinetics.