

Ulster University

Report of the outcomes of the adaptation to the GOC education & training requirements

Postgraduate Certificate in Independent Prescribing for Optometrists

ULS-IP1-ETR

Report confirmed by GOC 05 December 2025

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SECTION ONE - ABOUT THIS DOCUMENT

1.1 ABOUT THIS DOCUMENT

This report outlines the outcomes of the review of Ulster University's (provider) adapted Postgraduate Certificate in Independent Prescribing for Optometrists qualification (qualification) against the Requirements for Approved Qualifications in Additional Supply (AS), Supplementary Prescribing (SP) and/or Independent Prescribing (IP) (January 2022).

It includes:

- Feedback against each relevant standard (as listed in the Adaptation Form).
- The status of all the standards reviewed as part of the adaptation process (which includes the formal response process).
- Any action Ulster University is required to take.

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SECTION TWO - PROVIDER DETAILS

2.1 TYPE OF PROVIDER	
Provider	\boxtimes
Sole responsibility for the entire route to registration.	
Awarding Organisation (AO)	_
Sole responsibility for the entire route to registration with centres delivering the	
qualification(s).	

2.2 CENTRE DETAILS	
Centre name(s)	Not applicable.

2.3 EXTERNAL PARTNERS DELIVERING AND/OR MANAGING AREAS OF THE QUALIFICATION	
Not applicable.	

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SECTION THREE - QUALIFICATION DETAILS

3.1 QUALIFICATION DE	TAILS
Qualification title	Postgraduate Certificate in Independent Prescribing for Optometrists
Qualification level	Level 7 (Regulated Qualifications Framework [RQF])
Duration of qualification	One academic year
Number of cohorts per academic year	One
Month(s) of student intake	September
Delivery method(s)	Blended learning
Alternative exit award(s)	No alternative exit awards
Total number of students per cohort	15

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SECTION FOUR – SUMMARY OF THE OUTCOMES OF THE ADAPTATION PROCESS

4.1 QUALITY ASSURANCE ACTIVITY		
Type of activity	Review of the provider's adapted Postgraduate Certificate in	
	Independent Prescribing for Optometrists qualification against	
	the Requirements for Approved Qualifications in Additional	
	Supply (AS), Supplementary Prescribing (SP) and/or	
	Independent Prescribing (IP) (January 2022).	

4.2 GOC REVIEW TEAM	
Officer	Georgia Smith – Education Development Officer
Manager	Lisa Venables – Education Development Manager
Decision maker	Samara Morgan – Head of Education & CPD
Education Visitor Panel (panel) members	 Professor Andy Husband – Lay Chair Dr David Hill – Optometrist / Independent Prescribing Optometrist member Kiki Soteri – Optometrist / Independent Prescribing Optometrist member Pam McClean – Optometrist / Independent Prescribing Optometrist member

4.3 SUMMARY OF CONDITIONS AND RECOMMENDATIONS		
Conditions	The qualification has been set one condition against the	
	following standards:	
	• S3.6	
Recommendations	The qualification has been set two recommendations against	
	the following standards:	
	• S3.5	
	• S3.12	

Commentary against all of the standards reviewed are set out in section 4.4.

The qualification will remain subject to the GOC's quality assurance and enhancement methods (QAEM) on an ongoing basis.

4.4 STANDARDS OVERVIEW

The standards reviewed as part of the adaptation process for approved qualifications (as outlined in the Adaptation Form*) are listed below along with the outcomes, statuses, actions, and any relevant deadlines. Actions may include the following:

- A **condition** is set when the information submitted did not provide the necessary evidence and assurance that a standard is met; further action is required.
- A recommendation is set when the information submitted currently provides the
 necessary evidence and assurance that a standard is met. However, the GOC has
 identified this may be an area that could be enhanced or that will need to be reviewed to
 ensure the standard continues to be met.

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• **No further action** is required – the information submitted provides the necessary assurance that a standard is met.

*The following standards listed were **not** reviewed as part of the adaptation process but are monitored as part of the GOC's Quality Assurance and Enhancement Methods (QAEM):

- Standard one public and patient safety: S1.1, S1.2, S1.3, S1.4
- Standard two admissions of students: S2.2, S2.3, S2.4, S2.6
- Standard three assessment of outcomes and curriculum design: S3.7, S3.8, S3.9, S3.10, S3.16, S3.17
- Standard four management, monitoring and review of approved qualifications: S4.1, S4.4, S4.5, S4.6, S4.7, S4.8, S4.9, S4.10, S4.12
- Standard five leadership, resources and capacity: S5.3, S5.4, S5.5

Further details on the evidence that the provider was required to complete or submit as part of the education and training requirements (ETR) adaptation process can be found on our <u>qualifications in additional supply (AS)</u>, <u>supplementary prescribing (SP) and/or independent prescribing (IP)</u> webpage.

Standard no.	S2.1
Standard	Selection and admission criteria must be appropriate for entry to an
description	approved qualification for specialist entry to the GOC register (AS, SP and/or IP categories) including relevant health, character and fitness to practise checks. For overseas trainees, this should include evidence of proficiency in the English language of at least level 7 overall (with no individual section lower than 6.5) on the International English Language Testing System (IELTS) scale or equivalent.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	 The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: A completed 'Template 2 – criteria narrative'. The provider's 'Course Document' which outlines confirmation that:

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The provider has appropriate, clear, and comprehensive entry and IELTS requirements.

Standard no.	S2.5				
Standard	Recognition of prior learning must be supported by effective and robust				
description	policies and systems. These must ensure that trainees admitted at a point				
	other than the start of a programme have the potential to meet the				
	outcomes for the award of the approved qualification. Prior learning must				
	be recognised in accordance with guidance issued by The Quality Assurance Agency for Higher Education (QAA) and/or The Office of Qualifications and Examinations Regulation (Ofgual) / Scottish				
	Assurance Agency for Higher Education (QAA) and/or The Office of				
	Qualifications and Examinations Regulation (Ofqual) / Scottish				
	Qualifications Authority (SQA) / Qualifications Wales / Department for the				
	Economy in Northern Ireland and must not exempt trainees from				
	summative assessments leading to the award of the approved				
	qualification. (If necessary, separate arrangements will be made for the				
	safe transition of trainees who have not yet completed GOC-approved				
	therapeutic prescribing qualifications programmes prior to the introduction				
	of the new outcomes and standards.)				
Status	MET – no further action required at this stage				
Deadline	Not applicable.				
Rationale	The evidence reviewed provided the necessary assurance that this				
	standard is MET.				
	Supporting evidence reviewed included, but was not limited to:				
	A completed 'Template 2 – criteria narrative'.				
	The provider's 'Recognition of Prior Learning Policy'.				
	The provider's 'Course Document' which outlines confirmation that:				
	Trainees who are successful in their RPL application will not be				
	exempt from undertaking all summative assessments.				
	short and and and an administrative account to the				
	The information reviewed evidenced, amongst other elements, that:				
	The provider's recognition of prior learning (RPL) criteria and process				
	is fairly and consistently applied.				
	is rain, and sensitivity applied.				

Standard no.	S3.1
Standard	There must be a clear assessment strategy for the award of an approved
description	qualification. The strategy must describe how the outcomes will be
	assessed, how assessment will measure trainees' achievement of
	outcomes at the required level (Miller's Pyramid) and how this leads to an
	award of an approved qualification.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this
	standard is MET.
	Supporting evidence reviewed included, but was not limited to:
	A completed 'Template 2 – criteria narrative'.

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A completed 'Template 4 – assessment strategy'.
A completed 'Template 5 – module outcome map'.
The provider's 'Course Document' which outlines:
 The proposed assessment strategy.
 The assessment handbook.
 Qualification regulations.
The provider's 'Ulster University Assessment Code of Practice'
document.
The provider's 'mapping of GOC learning outcomes' document which
outlines the:
 Module each learning outcome will be assessed and;
 which assessment methods will be used.
The information reviewed evidenced, amongst other elements, that:
The provider has a clear assessment strategy mapped against learning
outcomes.

Standard no.	S3.2				
Standard	The approved qualification must be taught and assessed (diagnostically,				
description	formatively and summatively) in a progressive and integrated manner. The				
•	component parts should be linked into a cohesive programme (for				
	example, Harden's spiral curriculum), introducing, progressing and				
	assessing knowledge, skills and behaviour until the outcomes are				
	achieved.				
Status	MET – no further action required at this stage				
Deadline	Not applicable.				
Rationale	The evidence reviewed provided the necessary assurance that this				
rationalo	standard is MET.				
	Standard IS MET.				
	Supporting evidence reviewed included, but was not limited to:				
	Supporting evidence reviewed included, but was not limited to:				
	A completed 'Template 2 – criteria narrative'. A completed 'Template 4 – conseguent atratagu'				
	A completed 'Template 4 – assessment strategy'. A completed 'Template 5 – module outcome man'.				
	A completed 'Template 5 – module outcome map'.				
	The provider's 'Course Document' which outlines confirmation of the				
	course structure.				
	The provider's 'Mapping of GOC Learning Outcomes to Adapted IP				
	Programme' document.				
	The provider's assessment rubrics.				
	Narrative provided in support of the formal response process including:				
	 Confirmation of the in-person activities conducted as part of the 				
	qualification e.g., the placement module.				
	quantitation orgi, the placement mediale.				
	The information reviewed evidenced, amongst other elements, that:				
	 The information reviewed evidenced, amongst other elements, triat. The provider has an appropriate and consistent assessment strategy 				
	mapped against the learning outcomes.				
	 The provider has appropriate assessment methods for the qualification. 				
	The provider has appropriate assessment methods for the qualification.				

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Standard no.	S3.3
Standard	Curriculum design and the assessment of outcomes must involve and be
description	informed by feedback from a range of stakeholders such as patients,
	employers, trainees, commissioners, placement providers, members of the
	eye-care team and other healthcare professionals.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET.
	 Supporting evidence reviewed included, but was not limited to: A completed 'Template 2 – criteria narrative'. A completed 'Template 5 – module outcome map'. The provider's 'Course Document'. The provider's 'Employers Liaison Minutes'. The provider's 'Patient Liaison Board Minutes June 2023'. The provider's 'Patient Liaison Board Minutes June 2024'. The provider's 'Placement Handbook for Independent Prescribing'.
	 The information reviewed evidenced, amongst other elements, that: There was an appropriate level of engagement with a variety of stakeholders. It is clear where feedback has been used in the design and delivery of the qualification.

Standard no.	S3.4			
Standard	The outcomes must be assessed using a range of methods and all final,			
description	summative assessments must be passed. This means that compensation,			
•	trailing and extended re-sit opportunities within and between modules			
	where outcomes are assessed is not permitted.			
Status	MET – no further action required at this stage			
Deadline	Not applicable.			
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Template 2 – criteria narrative'. • A completed 'Template 4 – assessment strategy'. • A completed 'Template 5 – module outcome map'. • The provider's 'Course Document'.			
	 The information reviewed evidenced, amongst other elements, that: The provider utilises a clear range of assessment methods. All final summative assessments must be passed. Trailing and extended resit opportunities are not permitted. 			

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Standard no.	S3.5
Standard	Assessment (including lowest pass) criteria, choice and design of
description	assessment items (diagnostic, formative and summative) leading to the
	award of an approved qualification must ensure safe and effective practice
	and be appropriate for a qualification for specialist entry to the GOC
	register (AS, SP and/or IP).
Status	MET – a recommendation is set
Deadline	Response to the recommendation to be submitted by Monday 28 September 2026.
Rationale	The evidence reviewed provided the necessary assurance that this
	standard is MET.
	Supporting evidence reviewed included but was not limited to:
	A completed 'Template 2 – criteria narrative'.
	A completed 'Template 4 – assessment strategy'.
	A completed 'Template 5 – module outcome map'.
	The provider's 'Course Document'.
	The provider's 'Mapping of GOC Learning Outcomes to Adapted IP
	Programme' document.
	The provider's assessment rubrics.
	Narrative provided in support of the formal response process including:
	 Further information on the placement module.
	The information reviewed evidenced, amongst other elements, that:
	The provider's choice and design of assessment items and
	assessment criteria is appropriate.
	Whilst teaching, learning and assessment methods for trainees with
	specific needs may be modified, the outcomes cannot be modified and
	must be met in full.
	Although the information reviewed provided sufficient assurance that this
	standard is met, a recommendation has been set in relation to this
	standard as the GOC considers that it can be enhanced.
	Possible areas of evidence that can be submitted, are (this list is non-
	exhaustive):
	Evidence that the OSCE placement module assessment has been
	developed around specific station examples, demonstrating how the
	range of stations will assess specific learning outcomes.
	 Evidence that the module examiners have been provided with
	appropriate pass/fail criteria for the OSCE placement module.
	appropriate passitali criteria for the OOCE placement module.

Standard no.	S3.6
Standard	Assessment (including lowest pass) criteria must be explicit and set using
description	an appropriate and tested standard-setting process. This includes
	assessments which occur during learning and experience in practice.
Status	NOT MET – a condition is set

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Deadline	Monday 28 September 2026
Rationale	The evidence did not provide the necessary assurance and therefore this standard is NOT MET.
	 Supporting evidence reviewed included but was not limited to: A completed 'Template 2 – criteria narrative'. A completed 'Template 4 – assessment strategy'. A completed 'Template 5 – module outcome map'. The provider's 'Course Document'. The provider's 'Mapping to GOC Learning Outcomes to Adapted IP Programme' document. Narrative provided in support of the formal response process including: An assessment rubric for the portfolio assessment.
	The evidence did not provide the necessary assurance that this standard is met. There was insufficient evidence in the following areas:
	The standard setting process used to develop the assessment criteria, including the lowest pass.
	Possible areas of evidence that can be submitted, are (this list is non-exhaustive): • Evidence demonstrating the standard setting process used to develop the assessment criteria (including lowest pass) for the placement module within the qualification.
	Although a condition has been set, the GOC notes the progress the provider has made towards meeting this standard through providing clarity surrounding its placement module, including the assessment rubric for the coursework assessment. Further assurance is required regarding the standard setting process used to develop this module.

Standard no.	S3.11
Standard	A range of teaching and learning methods must be used to deliver the
description	outcomes.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET.
	Supporting evidence reviewed included, but was not limited to:
	A completed 'Template 2 – criteria narrative'.
	A completed 'Template 4 – assessment strategy'.
	The provider's 'Course Document' which outlines the:

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 Module descriptors detailing assessments and assessment
methods.
The provider's 'mapping of GOC learning outcomes' document which
outlines the:
 Module where each learning outcome will be assessed and;
 Which assessment methods will be used.
The information reviewed evidenced, amongst other elements, that:
The assessment and teaching methods appear to be appropriate to
deliver the learning outcomes at the required level.

Standard no.	S3.12
Standard description	To enable the development of trainees' clinical, diagnostic and prescribing skills to meet the outcomes, the approved qualification must integrate
description	learning and experience in practice (as a guide, approximately 90 hours).
	The supervision of a trainee's learning and experience in practice must be
	co-ordinated by an appropriately trained and qualified registered
Otativa	healthcare professional (DPP) with independent prescribing rights.
Status	MET – a recommendation is set
Deadline	Response to the recommendation to be submitted by Monday 28 September 2026.
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET.
	Supporting evidence reviewed included but was not limited to:
	A completed 'Template 2 – criteria narrative'.
	The provider's 'Course Document'.
	 The provider's 'Placement Handbook Independent Prescribing' document.
	 Narrative provided in support of the formal response process including: Details on how Designated Prescribing Practitioners (DPPs) are assessed for their suitability, training offered to DPPs, their roles, responsibilities and expectations and confirmation of trainee placement period(s).
	The information reviewed evidenced, amongst other elements, that: There are an appropriate number of hours in practice for the trainee
	under the co-ordinated supervision of a DPP.
	Although the information reviewed provided sufficient assurance that this standard is met, a recommendation has been set in relation to this
	standard is the GOC considers that it can be enhanced.
	Possible areas of evidence that can be submitted, are (this list is non-exhaustive):

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•	Evidence that the DPP training materials are explicit when referring to
	how the placement and caseload variation enables relevant experience
	for trainees to meet the learning outcomes.

S3.13
Outcomes delivered and assessed during learning and experience
in practice must be clearly identified, included within the assessment
strategy and fully integrated within the programme leading to the
award of an approved qualification.
MET – no further action required at this stage
Not applicable.
The evidence reviewed provided the necessary assurance that this standard is MET.
 Supporting evidence reviewed included, but was not limited to: A completed 'Template 2 – criteria narrative'. A completed 'Template 4 – assessment strategy'. A completed 'Template 5 – module outcome map'. The provider's 'Course Document'. The provider's 'Mapping of GOC Learning Outcomes to Adapted IP Programme' document. Narrative provided in support of the formal response process including: Confirmation of trainees' placement dates. The information reviewed evidenced, amongst other elements, that:
Learning and experience in practice is integrated into the qualification.

Standard no.	S3.14
Standard	The selection of outcomes to be taught and assessed during periods of
description	learning and experience in practice and the choice and design of assessment items must be informed by feedback from a variety of
	sources, such as patients, employers, trainees, DPPs, members of the eye-care team and other healthcare professionals.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET.
	 Supporting evidence reviewed included but was not limited to: A completed 'Template 2 – criteria narrative'. The provider's 'Course Document'.
	The provider's 'Employers Liaison Board Minutes June 2023'.
	The provider's 'Patient Liaison Board Minutes June 2023'.
	The provider's 'Patient Liaison Board Minutes June 2024'.
	Narrative provided in support of the formal response process including:

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 Confirmation of the date(s) trainees can commence their placement.
 The information reviewed evidenced, amongst other elements, that: The provider has considered stakeholder feedback in the outcomes taught and assessed during learning and experience.

Standard no.	S3.15
Standard	Equality and diversity data and its analysis must inform curriculum design,
description	delivery and assessment of the approved qualification. This analysis must
•	include trainees' progression by protected characteristic. In addition, the
	principles of equality, diversity and inclusion must be embedded in
	curriculum design and assessment and used to enhance trainees'
	experience of studying on a programme leading to an approved
	qualification.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this
	standard is MET.
	Supporting evidence reviewed included, but was not limited to:
	A completed 'Template 2 – criteria narrative'.
	The provider's 'Equality, Diversity and Inclusion (EDI) 2025 training
	schedule'.
	Narrative that outlines:
	How trainees are encouraged to bring their own experiences to
	course and project work.
	 Examples of where the curriculum design is accommodating of
	different needs.
	 How EDI and cultural awareness have been integrated into the
	curriculum.
	The information reviewed evidenced, amongst other elements, that:
	The provider has clearly considered EDI within its qualification design
	and delivery.
	It is clear how EDI has informed qualification design.
	There are clear examples of how EDI training is undertaken by staff
	and assessors etc.
	1 414 40000010 010.

Standard no.	S4.2
Standard	The organisation responsible for the award of the approved qualification
description	must be legally incorporated (i.e. not be an unincorporated association)
	and have the authority and capability to award the approved qualification.
Status	MET – no further action required at this stage
Deadline	Not applicable.

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Rationale	The evidence reviewed provided the necessary assurance that this standard is MET.
	Supporting evidence reviewed included, but was not limited to: • A completed 'Template 2 – criteria narrative'.
	 The information reviewed evidenced, amongst other elements, that: The provider has clear awarding powers and is a legally incorporated higher education institution.

Standard no.	S4.3
Standard	The provider must have a named point of contact for the approved
description	qualification.
Status	MET – no further action required at this stage
Deadline	Not applicable.
Rationale	The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. The information reviewed evidenced, amongst other elements, that:
	 The provider has a suitably qualified and experienced named individual for the qualification.

There must be an effective mechanism to identify risks to the quality of delivery and assessment of the approved qualification and to identify an requiring attention or development. Status MET – no further action required at this stage Deadline Not applicable. The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: A completed 'Form 2B – notification of adaptation'. A completed 'Template 2 – criteria narrative'. The provider's 'Clinical Learning in Practice Handbook'. The provider's 'School of Biomedical Sciences Risk Register'. The provider's 'Course Document' which outlines staff and students		
delivery and assessment of the approved qualification and to identify an requiring attention or development. Status MET – no further action required at this stage Deadline Not applicable. The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. • The provider's 'Clinical Learning in Practice Handbook'. • The provider's 'School of Biomedical Sciences Risk Register'. • The provider's 'Course Document' which outlines staff and students'	Standard no.	S4.11
requiring attention or development. Status MET – no further action required at this stage Deadline Not applicable. The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. • The provider's 'Clinical Learning in Practice Handbook'. • The provider's 'School of Biomedical Sciences Risk Register'. • The provider's 'Course Document' which outlines staff and students	Standard	There must be an effective mechanism to identify risks to the quality of the
Status Deadline Not applicable. The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. • The provider's 'Clinical Learning in Practice Handbook'. • The provider's 'School of Biomedical Sciences Risk Register'. • The provider's 'Course Document' which outlines staff and students	description	delivery and assessment of the approved qualification and to identify areas
 Not applicable. Rationale The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: A completed 'Form 2B – notification of adaptation'. A completed 'Template 2 – criteria narrative'. The provider's 'Clinical Learning in Practice Handbook'. The provider's 'School of Biomedical Sciences Risk Register'. The provider's 'Course Document' which outlines staff and students 		requiring attention or development.
The evidence reviewed provided the necessary assurance that this standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. • The provider's 'Clinical Learning in Practice Handbook'. • The provider's 'School of Biomedical Sciences Risk Register'. • The provider's 'Course Document' which outlines staff and students	Status	MET – no further action required at this stage
standard is MET. Supporting evidence reviewed included, but was not limited to: • A completed 'Form 2B – notification of adaptation'. • A completed 'Template 2 – criteria narrative'. • The provider's 'Clinical Learning in Practice Handbook'. • The provider's 'School of Biomedical Sciences Risk Register'. • The provider's 'Course Document' which outlines staff and students	Deadline	Not applicable.
 A completed 'Form 2B – notification of adaptation'. A completed 'Template 2 – criteria narrative'. The provider's 'Clinical Learning in Practice Handbook'. The provider's 'School of Biomedical Sciences Risk Register'. The provider's 'Course Document' which outlines staff and students 	Rationale	· · · · · · · · · · · · · · · · · · ·
		 A completed 'Form 2B – notification of adaptation'. A completed 'Template 2 – criteria narrative'. The provider's 'Clinical Learning in Practice Handbook'. The provider's 'School of Biomedical Sciences Risk Register'. The provider's 'Course Document' which outlines staff and students' responsibilities to raise concerns. Narrative provided in support of the formal response process including: Details of how programme-specific risks are appropriately managed once identified.

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The provider has appropriate processes in place to raise, assess and
mitigate risks to the qualification.

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Standard no.	S5.1	
Standard	There must be robust and transparent mechanisms for identifying,	
description	securing and maintaining a sufficient and appropriate level of ongoing resources to deliver the outcomes to meet these standards, including	
	human and physical resources that are fit for purpose and clearly	
	integrated into strategic and business plans. Evaluations of resources and	
	capacity must be evidenced together with evidence of recommendations	
	, ,	
Status	considered and implemented. MET – no further action required at this stage.	
Deadline	MET – no further action required at this stage	
	Not applicable.	
Rationale	The evidence reviewed provided the necessary assurance that this	
	standard is MET.	
	Supporting evidence reviewed included, but was not limited to:	
	A completed 'Template 2 – criteria narrative'.	
	The provider's 'Course Document' which outlines:	
	The resource allocation and planning for the next five years,	
	including details of the staff, facilities and accommodation.	
	The provider's 'School of Biomedical Sciences Risk Register'.	
	The provider's Ochool of Biomedical Ociences Nisk Negister.	
	The information reviewed evidenced, amongst other elements, that:	
	The provider has appropriately assessed and mitigated risks.	
	The provider's teaching and learning environments are suitable and	
	have sufficient capacity for the planned trainee numbers.	
	The provider has sufficient resource (human and physical) to deliver	
	the qualification.	

Standard no.	S5.2		
Standard	There must be a sufficient and appropriately qualified and experienced		
description	staff team. This must include:		
	an appropriately qualified and experienced programme leader,		
	supported to succeed in their role; and		
	sufficient staff responsible for the teaching and assessment of the		
	outcomes, including GOC registrants and other suitably qualified		
	healthcare professionals.		
Status	MET – no further action required at this stage		
Deadline	Not applicable.		
Rationale	The evidence reviewed provided the necessary assurance that this		
	standard is MET.		
	Supporting evidence reviewed included, but was not limited to:		
	 A completed 'Form 2B – notification of adaptation'. 		
	A completed 'Template 2 – criteria narrative'.		
	The provider's 'Course Document' which outlines:		

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- The key members of staff supporting with the qualification delivery.
- The experience and professional backgrounds of the staff members.
- The resource allocation and planning for the next five years, including details of the staff, facilities and accommodation.

The information reviewed evidenced, amongst other elements, that:

- The qualification has appropriate leadership.
- The provider has appropriately experienced and qualified staff members to deliver the qualification.

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