

Schedule of Accreditation

issued by

United Kingdom Accreditation Service

2 Pine Trees, Chertsey Lane, Staines-upon-Thames, TW18 3HR, UK

 <p>Accredited to ISO/IEC 17065:2012 to provide product conformity certification</p>	LRQA Apave Limited	
	Issue No: 002 Issue date: 28 August 2026	
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DETAIL OF ACCREDITATION

Accreditation for the purpose of UK Approved Body Activity in accordance with UKCA Requirements and UKAS Publication GEN 5			
Directive/Regulation	Conformity Assessment procedure/Module/Article	Category of products or individual products	Methods / Procedures Essential requirements: Product specification/ Properties / Standards
Pressure Equipment (Safety) Regulations 2016, SI 2016 No 1105 as amended	Conformity assessment procedures in accordance with Regulation 42 of the SI Schedule 1A Part 6 – Module D1 Quality assurance of the production process Schedule 1A Part 8 – Module E1 Quality assurance of final pressure equipment inspection and testing	Category II Equipment (or below)	Schedule 2, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
	Schedule 1A Part 3 – Module B Design Type Examination Schedule 1A Part 4 – Module C2 Conformity to type based on internal production control plus supervised pressure equipment checks at random intervals Schedule 1A Part 5 – Module D Conformity to type based on quality assurance in the production process Schedule 1A Part 7 – Module E Conformity to type based on pressure equipment quality assurance	Category III Equipment (or below)	Schedule 2, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.



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Directive/Regulation	Conformity Assessment procedure/Module/Article	Category of products or individual products	Methods / Procedures Essential requirements: Product specification/ Properties / Standards
Pressure Equipment (Safety) Regulations 2016, SI 2016 No 1105 as amended (cont'd)	Schedule 1A Part 9 – Module F Conformity to type based on pressure equipment verification	Category III Equipment (or below) (cont'd)	Schedule 2, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
	Part 3 – Module B Production Type Examination Part 5 – Module D Conformity to type based on quality assurance in the production process Part 9 – Module F Conformity to type based on pressure equipment verification Part 10 – Module G Conformity based on unit verification Part 12 – Module H1 Conformity based on full quality assurance plus design examination	Category IV Equipment (or below)	Schedule 2, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
END			