Schedule of Accreditation

issued by

United Kingdom Accreditation Service

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

6906 **Accredited to ISO/IEC 17065:2012 to provide product conformity certification**	**LRQA Apave Limited** **Issue No:** 001 **Issue date:** 28 November 2023	
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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	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	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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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 (cont'd)	Schedule 1A Part 9 – Module F Conformity to type based on pressure equipment verification	Category III Equipment (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	Schedule 2, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.

Accreditation for the purpose of Notified Body Activity relating to the Northern Ireland market (CE + UKNI) taking into account EA-2/17

Directive/Regulation	Conformity Assessment procedure/Module/Article	Category of products or individual products	Methods / Procedures Essential requirements: Product specification/ Properties / Standards
Pressure Equipment (Directive 2014/68/EU) implemented in Northern Ireland by the Pressure Equipment (Safety) Regulation 2016	**Conformity assessment procedures in accordance with Article 14 of the Directive** Annex III.6 Module D1 Quality assurance of the production process Annex III.1 Module E1 Quality assurance of final pressure equipment inspection and testing	Category II Equipment	Annex 1, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
	Annex III.3 Module B Design Type Examination Annex III.4 Module C2 Conformity to type based on internal production control plus supervised pressure equipment checks at random intervals Annex III.5 Module D Conformity to type based on quality assurance in the production process Annex III.7 Part 7 – Module E Conformity to type based on pressure equipment quality assurance	Category III Equipment	Annex 1, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.

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Accreditation for the purpose of Notified Body Activity relating to the Northern Ireland market (CE + UKNI) taking into account EA-2/17

Directive/Regulation	**Conformity Assessment procedure/Module/Article**	**Category of products or individual products**	**Methods / Procedures** **Essential requirements:** **Product specification/ Properties / Standards**
Pressure Equipment (Directive 2014/68/EU) as implemented in Northern Ireland by the Pressure Equipment (Safety) Regulation 2016, SI 2016 No 1105 as amended (cont'd)	Annex III.9 Module F Conformity to type based on pressure equipment verification	Category III Equipment (cont'd)	Annex 1, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
	Annex III.3 Module B Production Type Examination Annex III.5 Module D Conformity to type based on quality assurance in the production process Annex III.9 Module F Conformity to type based on pressure equipment verification Annex III.10 Module G Conformity based on unit verification Annex III.12 Module H1 Conformity based on full quality assurance plus design examination	Category IV Equipment	Annex 1, Essential safety requirements and assessment of technical documentation, quality system and/or products in accordance with the relevant Module.
END			