Title	Describe laboratory quality systems		
Level	5	Credits	4

Purpose People credited with this unit standard are able to describe: the aims of laboratory quality systems; the management of laboratory quality systems; and the technical requirements of laboratory quality systems in accredited laboratories.

Classification	Science > Science - Core	
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Available grade	Achieved	76,
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Guidance Information

- All work must be carried out in accordance with the quality management system, documented protocol system or Standard Operating Procedures acceptable in a commercial or research laboratory.
- 2 Health and Safety practices must conform to Australian/New Zealand Standard AS/NZS 2243 *Safety in Laboratories* Parts 1, 2, 3, 7 and 10 available at http://www.standards.co.nz.
- 3 Legislation applicable to this unit standard includes: Health and Safety at Work Act 2015; Hazardous Substances and New Organisms Act 1996.
- 4 Glossary

Accredited laboratory refers to a laboratory accredited according to the Australian/New Zealand Standard NZS ISO/IEC 17025:2005 General requirements for the competence of testing and calibration laboratories.

Outcomes and performance criteria

Outcome 1

Describe the aims of laboratory quality systems in accredited laboratories.

Performance criteria

- 1.1 Concepts of quality, fitness for purpose, and continuous improvement are described in laboratory quality systems.
- 1.2 Client expectations are described and related to the reliability of laboratory results.

Outcome 2

Describe the management of laboratory quality systems in accredited laboratories.

Range section 4 of New Zealand Standard 17025.

Performance criteria

2.1 The management requirements of the quality system are described in relation to the accredited laboratory.

Outcome 3

Describe the technical requirements of laboratory quality systems in accredited laboratories.

Range section 5 of New Zealand Standard 17025.

Performance criteria

3.1 The technical requirements of the quality system are described in relation to the accredited laboratory.

This unit standard is expiring. Assessment against the standard must take place by the last date for assessment set out below.

Status information and last date for assessment for superseded versions

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Process	Version	Date	Last Date for Assessment	
Registration	1	22 December 1996	31 December 2014	
Revision	2	19 February 1998	31 December 2014	
Review	3	23 November 1999	31 December 2014	
Review	4	26 February 2003	31 December 2014	
Review	5	21 May 2010	31 December 2025	
Rollover	6	27 January 2015	31 December 2025	
Review	7	27 September 2018	31 December 2025	
Review	8	30 November 2023	31 December 2025	

Consent and Moderation Requirements (CMR) reference	0113
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This CMR can be accessed at http://www.nzqa.govt.nz/framework/search/index.do.