Medical Laboratories —Particular Requirements for Quality and Competence

(A New Zealand adoption of ISO 15189:2007)

COMMITTEE REPRESENTATION

Committee P 15189, Medical laboratories – Quality and competence was responsible for the adoption of this International Standard. The P 15189 Committee was established by the Standards Council under the Standards Act 1988.

The committee consisted of representatives of the following:

Nominating Organisations

Canterbury District Health Board
National Cervical Screening Programme, Ministry of Health
New Zealand Blood Service
Southern Community Laboratories
Waitemata District Health Board

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NOTES

CON	ITEN	TS	Page	
Comi	mittee	Representation	IFC	
Ackn	owled	gement	IFC	
Сору	right		IFC	
Fore	word		4	
Introd	ductio	n	5	
Secti	ion			
1		oe	7	
2		native references		
3		Terms and definitions		
4	Management requirements			
7	4.1			
	4.2	Quality management system		
	4.3	Document control		
	4.4	Review of contracts		
	4.5	Examination by referral laboratories		
	4.6	External services and supplies		
		• •		
	4.7	Advisory services		
	4.8	Resolution of complaints		
	4.9	Identification and control of nonconformities		
		Corrective action		
		Preventive action		
		Continual improvement		
		Quality and technical records		
		Internal audits		
		Management review		
5	Tech	19		
	5.1	Personnel		
	5.2	Accommodation and environmental conditions	21	
	5.3	Laboratory equipment	22	
	5.4	Pre-examination procedures	24	
	5.5	Examination procedures	26	
	5.6	Assuring quality of examination procedures	28	
	5.7	Post-examination procedures	29	
	5.8	Reporting of results	29	
Biblio	graph	ıy	43	
Anne	ex			
Α	Corre	elation with ISO 9001:2000 and ISO/IEC 17025:2005 (Informative)	32	
В	Reco			
_		36		
С	-	es in laboratory medicine (Informative)		
Table	<u> </u>			
A.1			32	
A.2				

Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

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The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

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ISO 15189 was prepared by Technical Committee ISO/TC 212, Clinical laboratory testing and in vitro diagnostic test systems.

This second edition cancels and replaces the first edition (ISO 15189:2003) which has been technically revised in order to align it more closely with the second edition of ISO/IEC 17025.