CE Technical Documentation Review Report

Applicant:	Fine Care Corporation 3th Floor,Sun House,B/h old high court,Ashram road,Ahmedabad-380009
Examination intent:	Examination of the completeness of the Technical Documentation according to the requirements of the In Vitro Diagnostic Medical Devices Directive 98/79/EC Annex III
Product(s):	Micropipettes
Type(s)/Model(s):	Fixed Volumes, Variable Volumes
Classification:	Other IVD products (according to manufacturers declaration)
Examination period:	2005 Jan.20, 2005 Jan.29-30
Review result:	During the examination of the provided Technical Documentation (dated 2002-04) no Non-compliance according to the requirements of the In Vitro Diagnostic Medical Devices Directive 98/79/EC Annex III was detected.

TÜV Rheinland (Shanghai) Co., Ltd.

Xiaoli REN
Manager (Asia Region)
Medical Services

Product Safety · Quality Assurance
TÜV Rheinland
Approved

Rev.02, 2005-02-25

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