PROMOSENI 08 MODE

BECKMAN COULTER

DECLARATION OF CONFORMITY

Beckman Coulter Inc. hereby ensures and declares that the product(s) listed below comply with the requirement of the European Union In Vitro Diagnostic Medical Device Directive 98/79/EC.

Beckman Coulter, Inc. assure et déclare par la présente que le(s) produit(s) listé(s) ci-dessous sont conformes aux exigences de la directive européenne 98/79/CE relative aux dispositifs médicaux de diagnostic in vitro.

Beckman Coulter, Inc. dichiara ed assicura che i prodotti qui elencati sono conformi ai requisiti della direttiva comunitaria 98/79/CE relativa ai dispositivi medico-diagnostici in vitro.

Beckman Coulter, Inc. versichert und erklärt hiermit, daß die im Folgenden aufgeführten Produkte den Auflagen der IVD-Richtlinie für In-vitro-Diagnostika der Europäischen Union (98/79/EC) entsprechen.

Beckman Coulter, Inc. asegura y declara que los productos listados a continuación cumplen con los requisitos establecidos en la directiva 98/79/EC de la Comunidad Europea para dispositivos médicos de diagnóstico in vitro.

Product(s) / Produkt(e) / Prodotto(i) / Produit(s) / Producto(s):

LH750 Hematology Analyzer
LH750 SM (Slide Maker)
LH750 SS (Slide Stainer)

Gabriel Compton, Manager
Product Compliance Engineering

Date: November 15, 2004

[AR] Authorized Representative
Beckman Coulter Ireland Inc.
Mervue Business Park
Mervue Galway, Ireland

Quality Systems Registration 404

IQNet
I.S. EN ISO 9001

Conformity Assessment Procedure
Annex III, Self-Declared

Document Control

Issue Date:	December 2, 2003	
Revision Lev:	1.2	
Revision Date:	November 15, 2004	
Starting Serial #:	LH750	AG45676
	LH750 SM	AG47189
	LH750 SS	AG39115
Filename:	2004-913-029I	

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