

UNIPATH

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EC DECLARATION OF CONFORMITY

We, Unipath Limited, declare that the following in-vitro diagnostic medical device meets the essential requirements of Council Directive 98/79/EC, concerning in-vitro diagnostic medical devices (published 27 October 1998). The procedure for examination of Unipath's quality assurance system for compliance with Annex IV (with the exception of sections 4 and 6) of the directive followed. A copy of the current EC Full Quality Assurance System Approval Certificate No. 275 is held in the Product Technical File.

CLEARVIEW CHLAMYDIA MF

Signed for and on behalf of Unipath Limited :-



Mark Gittins
Quality Manager

2nd February 2004

Date