



EC Declaration of Conformity

Manufacturer: QUIDEL Corporation
10165 McKellar Court
San Diego, California 92121
USA

Product Names: QuickVue® Influenza A+B Test (Catalog #20183 and #20185)

Description: The QuickVue Influenza Test is intended for the qualitative detection of influenza type A and type B antigens as an aid in the rapid differential diagnosis of acute influenza type A and type B virus infections.

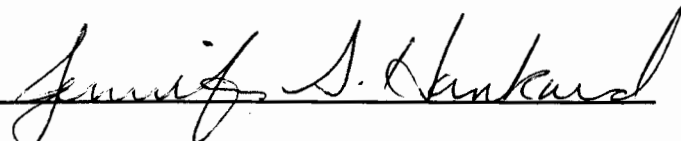
Classification: General In Vitro Diagnostic Device

Conformity Assessment Route: Annex III of Council Directive 98/79/EC Concerning In Vitro diagnostic devices

Estimated Start of CE-Marking: June 21, 2004

QUIDEL Corporation, being the manufacturer / distributor within the European Economic Area hereby declares that the products covered by the declaration conform to the Essential Requirements of EC Directive 98/79/EC. All supporting documentation is retained under the premises of the manufacturer.

This declaration is valid for all devices described in this document.

Signed: 

Date: 6/11/2004

Jennifer S. Hankard
Regulatory Affairs Manager
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