



EC Declaration of Conformity

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

Product Names: QuickVue® Influenza Test (Catalog #00317)

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 diagnosis of acute influenza 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: November 19, 2003

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: _____

11/19/2003

Jennifer S. Hankard
Regulatory Affairs Manager
QUIDEL Corporation
10165 McKellar Court
San Diego, California 92121
USA