

## **EC DECLARATION OF CONFORMITY**

We,

**Thermo Fisher Scientific Oy, Clinical Diagnostics Finland**

hereby declare that the reagent kit with

Trade name: **IMMUNOGLOBULIN M (IgM)**  
Code: **981670 and 981938**

comply with the In Vitro Diagnostic Medical Devices Directive 98/79/EC.

This Declaration is valid for all batches which are placed on the market by ourselves on or after February 15, 2002 and which bear the CE marking.

Vantaa, September 17, 2010

Thermo Fisher Scientific Oy



Silja Halme  
Quality Manager  
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