

## **EC DECLARATION OF CONFORMITY**

We,

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

hereby declare that the reagent kit with

Trade name: **CRP High Sensitivity**  
Code: **981798**

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 25<sup>th</sup> October, 2004 and which bear the CE marking.

Vantaa, July 28, 2009

Thermo Fisher Scientific Oy



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