

## **EC DECLARATION OF CONFORMITY**

We,

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

hereby declare that the reagent kit with

Trade name: **SPECITROL**  
Code: **981250**

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 June 26, 2003 and which bear the CE marking.

Vantaa, July 28, 2009

Thermo Fisher Scientific Oy



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Quality Manager  
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