

## **EC DECLARATION OF CONFORMITY**

We,

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

Hereby declare that the Selective Clinical Chemistry Analyzers with

Trade name: **Konelab PRIME 60 ISE KUSTI**  
Type: **982**

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

This Declaration is valid for all devices which are placed on the market by ourselves on or after December 28, 2006 and which bear the CE marking.

Vantaa, February 11, 2010

Thermo Fisher Scientific Oy



Silja Halme  
Quality Manager  
Quality, Regulatory and Compliance  
Clinical Diagnostics Finland