

## **EC DECLARATION OF CONFORMITY**

We,

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

Hereby declare that the Selective Clinical Chemistry Analyzers with

Trade name:	<b>Konelab 30i</b>	<b>Konelab 30</b>
Type:	<b>964</b>	<b>963</b>

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 December 8, 2003 and which bear the CE marking.

Vantaa, August 06, 2009

Thermo Fisher Scientific Oy



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