

EC DECLARATION OF CONFORMITY

We,

Thermo Fisher Scientific Oy, Clinical Diagnostics Finland

hereby declare that the accessory of Konelab family of clinical chemistry analyzers (in vitro diagnostics devices) with

Trade name: **Multicell Cuvettes**
Code: **984000**

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 1st, 2007 and which bear the CE marking.

Vantaa, October 4th, 2007

Thermo Fisher Scientific Oy



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