

## **EC DECLARATION OF CONFORMITY**

We,

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

hereby declare that the reagent kit with

Trade name:	<b>Lipotrol HDL/LDL Abnormal</b>
Code:	<b>981907</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 September 24, 2009 and which bear the CE marking.

Vantaa, September 28<sup>th</sup>, 2009

Thermo Fisher Scientific Oy



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