

EC DECLARATION OF CONFORMITY

We,

Thermo Fisher Scientific Oy, Clinical Diagnostics Finland

Hereby declare that the *TCAutomation Modules* with

Trade name	Type
Multitube Carrier	1001
Conveyor Module 600 mm	1002
Conveyor Module 900 mm	1003
Conveyor Module 1200 mm	1004
Conveyor Module 1500 mm	1005
Conveyor Module 1800 mm	1006
Conveyor Module 2400 mm	1007
U-Turn Module	1008
L-Turn Module	1009
Parallel Bypass Module	1010
Perpendicular Bypass Module	1011
Entry Module	1012
Exit Module	1013
Rack Entry Module	1014
Rack Exit Module	1015
Combined Rack Entry/Exit Module	1016
Sample Check Module	1017
Buffer Module	1018
Decapper Module	1019
Centrifuge Module	1020
Aliquoter and Labeller Module	1021
Conveyor Module 2100 mm	1022
T-Turn Module	1024
Internal U-Turn	1025
Z-Turn Right Module	1028
Z-Turn Left Module	1029
U-Turn Mek	1034
Recapper	1027
Modular Interface	1023
Decapper Flex	1031
Parallel Bypass Module	1041
Stago STA-R Interface	1042
ES Flex	1038

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 8, 2003 and which bear the CE marking.

Vantaa, December 02nd, 2009

Thermo Fisher Scientific Oy



Silja Halme
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Quality, Regulatory and Compliance
Clinical Diagnostics Finland