ROSCO DIAGNOSTICA A/S
Taastrupgaardsvej 30
DK-2630 Taastrup, Denmark

EU DECLARATION OF CONFORMITY for KITS FOR BACTERIAL IDENTIFICATION

ROSCO DIAGNOSTICA A/S hereby declares that the following products:

Neo-Sensitabs™ Screen ID Pack

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

According to the directive the above kits are classified in the group of 'Other Devices', i.e. devices not listed in List A or B in Annex II and devices not intended for Performance Evaluation. Conformity route: Annex III.

The Declaration covers the Kits (DVL0004) distributed from ROSCO DIAGNOSTICA A/S, which have been supplied with a CE-mark for compliance.

Date of Validity: 22.02.2010

Authorization:

Mikkel Pandrup Duer-Jensen
Manager Quality Assurance

ROSCO DIAGNOSTICA's List of Kits for bacterial identification = DVL0004 is available on request, and shall continuously be updated.