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

EU DECLARATION OF CONFORMITY for NEO-SENSITABS™

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

NEO-SENSITABS™

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

According to the directive NEO-SENSITABS™ 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 all Neo-Sensitabs™ (DVL0001) distributed from ROSCO DIAGNOSTICA A/S, which have been supplied with a CE-mark for compliance.

Date of Validity:

01.01.2011

Authorization:

Mikkel Pandrup Duer-Jensen
QA Manager

ROSCO DIAGNOSTICA's List of Neo-Sensitabs = DVL0001 is available on request, and shall continuously be updated.