

EU DECLARATION OF CONFORMITY for NEO-SENSITABS DISPENSERS

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

NEO-SENSITABS DISPENSERS

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

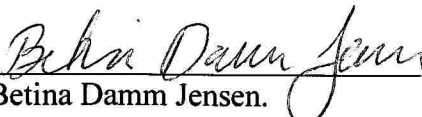
According to the directive the above dispensers are accessories to Neo-Sensitabs and classified in the group of 'Other Devices', i.e. devices not listed in List A or B in Annex II and devices not intended for Performance Evaluation. Conformity route: Annex III.

The Declaration covers the above dispensers (DVL0003) distributed from ROSCO DIAGNOSTICA A/S, which have been supplied with a CE-mark for compliance.

Date of Validity:

04.02.2009

Authorization:


Betina Damm Jensen.
QA Manager

ROSCO DIAGNOSTICA's List of Dispensers, mobile = DVL0003 is available on request, and shall continuously be updated.