

EC DECLARATION OF CONFORMITY

According to Annex III of the IVD Directive 98/79/EC

We, the manufacturers: BioPorto Diagnostics A/S
Tuborg Havnevej 15, st.
DK-2900 Hellerup
Denmark,

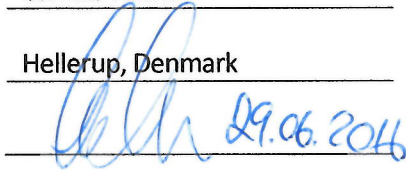
certify that the product identified below conforms to the provisions of the IVD Directive 98/79/EC as transposed into Danish national law as the Danish Act No. 1269 of December 12, 2005, to consolidate the law on medical equipment for in vitro diagnostics.

Product name	Catalogue number
The NGAL Test™ Reagent Kit	ST001CA
The NGAL Test™ Calibrator Kit	ST002CA
The NGAL Test™ Control Kit	ST003CA

Name: Camilla Recke

Position: QA & RA Manager

Location: Hellerup, Denmark

Signature:  29.06.2016

Date: 29.06.2016