SIEMENS

EU Declaration of Conformity



We hereby declare that the product described below conforms to all applicable requirements of Council Directive 98/79/EC for in vitro diagnostic medical devices.

Legal Manufacturer: Siemens Healthcare Diagnostics Products Ltd.

Glyn Rhonwy

Llanberis, Gwynedd, LL55 4EL, UK

Place of Manufacture: Siemens Healthcare Diagnostics Products Ltd.

Glyn Rhonwy

Llanberis, Gwynedd, LL55 4EL, UK

Product Name: IMMULITE/IMMULITE 2000 Free PSA

Catalogue Number (REF): L2KPF2

Siemens Material Number (SMN): 10380984

Classification: ANNEX II, List B

Conformity Assessment Route: ANNEX IV

Notified Body: Lloyd's Register Quality Assurance Ltd.

1 Trinity Park, Bickenhill Lane Solihull, B37 7ES, UK Identification No. 0088

Document Identifier: EC DEC_IMM 2000 Free PSA L2KPF

Version: 01

This declaration of conformity is issued under the sole responsibility of Siemens Healthcare Diagnostics Products Ltd.

This declaration supersedes any declaration issued previously for the same product.

Souta Wample

Signature:

Anita Wample

Head of Quality Management

Siemens Healthcare Diagnostics Products Ltd.

LLanberis Gwynedd LL55 4EL, UK

2016 08-72 Date

[YYYY-MM-DD]

Document No. EC DEC_IMM 2000 Free PSA L2KPF Ver. 01

Page 1 of 1