# Declaration of Conformity

EUROIMMUN Medizinische Labordiagnostika AG
Seekamp 31, D-23560 Lübeck, Germany

declare under our sole responsibility that the products

| | |
|---|---|
| *IF Sprinter (model IFA with ReSa-Trax)* | *YG 0032-0101* |
| *IF Sprinter (model IFA)* | *YG 0032-0101-1* |
| *IF Sprinter (model IFA/ELISA)* | *YG 0032-0101-2* |
| *IF Sprinter (model IFA/ELISA with ReSa-Trax)* | *YG 0032-0101-3* |

(product name, order no)

meet the demands of

*Directive 98/79/EC on in vitro diagnostic medical devices of 27 October 1998*

The following standards are applied:
- EN 61010-1
- EN 61010-2-81
- EN 61326-1
- EN 14971

Lübeck, 25.08.2011

(Place and date of issue)

Dr. Wolfgang Schlumberger
- Member of the Board -

Susanne Aleksandrowicz
- Member of the Board -