

## EU Declaration of Conformity



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

<b>Legal Manufacturer:</b>	Siemens Healthcare Diagnostics Products Ltd. Glyn Rhonwy Llanberis, Gwynedd, LL55 4EL, UK
<b>Place of Manufacture:</b>	Siemens Healthcare Diagnostics Products Ltd. Glyn Rhonwy Llanberis, Gwynedd, LL55 4EL, UK
<b>EU Authorized Representative:</b>	Siemens Healthcare Diagnostics Manufacturing Ltd. Chapel Lane Swords, Co. Dublin, Ireland
<b>Product Name:</b>	IMMULITE 2000 Anti-TPO Ab
<b>Catalogue Number (REF):</b>	L2KTO2 L2KTO6
<b>Siemens Material Number (SMN):</b>	10381650 10381649
<b>Classification:</b>	General IVD
<b>Conformity Assessment Route:</b>	ANNEX III
<b>Document Identifier:</b>	EC DEC_IMM 2000 Anti-TPO Ab L2KTO
<b>Version:</b>	02

*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.*

<b>Signature:</b>	2019-02-04
<b>Malgorzata Robak</b> Regulatory Affairs Supervisor Siemens Healthcare Diagnostics Products Ltd. Llanberis, Gwynedd LL55 4EL, UK	<b>Date</b> [YYYY-MM-DD]