

EC – DECLARATION OF CONFORMITY

The product named hereinafter was developed, designed, and manufactured in compliance with the relevant, fundamental safety and health requirements of the listed EC directives and norms.

In the event of modifications that were not authorised by us or if the product is used in a manner that is not in line with the intended purpose, this declaration will be rendered void.

<i>Product designation:</i>	Laboratory centrifuge
<i>Product name:</i>	Sigma 2-7 Sigma 2-7 Cyto
<i>Order number:</i>	10226 10228
<i>Directives:</i>	2006/42/EC Machinery Directive 2014/35/EU Low Voltage Directive 2014/30/EU EMC Directive (EU) 2015/863 RoHS Directive
<i>Normes:</i>	EN 61010-2-020:2017 EN IEC 61326-1:2021

Sigma Laborzentrifugen GmbH

An der Unteren Söse 50
37520 Osterode
Germany

Authorised representative
for CE matters:
Eckhard Tödteberg

Osterode, 2024-12-04



Managing Director

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<i>Product designation:</i>	Laboratory centrifuge
<i>Product name:</i>	Sigma 2-7 IVD
<i>Part number:</i>	10227
<i>Basic UDI as referred to in Part C of Annex VI:</i>	426073439IVD01001JQCJ4
<i>Manufacturer:</i>	Sigma Laborzentrifugen GmbH An der Unteren Söse 50 37520 Osterode Germany
<i>Single Registration Number (SRN):</i>	DE-MF-000009414

As the manufacturer of the unit(s), we assume full responsibility and hereby declare that the product(s) mentioned hereinabove comply with the requirements as set out in the following regulation(s)/directive(s).

<i>Regulations:</i>	(EU) 2017/746 Regulation on in vitro diagnostics
<i>Directives:</i>	(EU) 2015/863 RoHS directive
<i>Risk class in accordance with Annex VIII</i>	Class A

Osterode, 2024-04-25



Managing Director