CE

Declaration of Conformity

technoclone

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Manufacturer	Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH Brunner Str. 67 1230 Vienna Austria
SRN	**AT-MF-000024115**
Product(s)	Ceveron 100 series System Solution
Basic UDI-DI	912003686TC990000003UE

UDI-DI (GTIN)	**Product name**	**REF**
09120036864122	Ceveron 100 series System Solution	9820201

Classification	Class A according to Annex VIII rule 5(a)
Common Standards	N/A
Notified Body Involvement	N/A
Conformity assessment route	Following article 48(10), the EU declaration of conformity is issued after drawing up the technical documentation set out in Annexes II and III (IVDR)
Initial date of first declaration of conformity	26-May-2022
Valid until	25-May-2025

This declaration of conformity is issued under the sole responsibility of Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH. We hereby declare that the device(s) specified above meet the provision of the Regulation for *in vitro* diagnostic medical devices (EU) 2017/746 Annex I.

Name: Veronika Binder **Position:** Chief Executive Officer

Signed: **Date:** 01-Feb-2023

This declaration of conformity is issued in Vienna, Austria on behalf of Technoclone Herstellung von Diagnostika und Arzneimitteln GmbH.