Marienfeld * Am Wöllerspfad 4 * 97922 Lauda-Königshofen * Germany

EC Declaration of Conformity

In accordance with the requirements of Directive 98/79/EC for in vitro diagnostic medical devices, Annex I, we declare under our sole responsibility the conformity of the products.

We, the Paul Marienfeld GmbH & Co. KG, are registered as manufacturer respectively marketing authorisation holder of in-vitro diagnostic devices at DIMDI (Deutsches Institut für Medizinische Dokumentation und Information).

We hereby certify that the following in-vitro diagnostic devices comply with the essential requirements of Annex I to the Directive 98/79/EC and are suitable for use in accordance with these regulations.

Product specification	**Registration no.**
Cover glasses for blood and tissue sections	DE/CA 37/IVD/8/12

This registration number covers the products with the following Art.No.

0100032	0101040	0101172	0102042	0103172	0111600	0895222
0100042	0101050	0101173	0102052	0103192	0111620	0895242
0100052	0101052	0101182	0102062	0103222	0111640	
0100062	0101053	0101192	0102112	0103242	0111650	
0100112	0101060	0101193	0102122	0107032	0111700	
0100122	0101062	0101202	0102142	0107052	0117500	
0100142	0101092	0101212	0102152	0107222	0117520	
0100172	0101102	0101222	0102172	0107242	0117530	
0100192	0101103	0101224	0102192	0111500	0117580	
0100222	0101112	0101232	0102222	0111520	0117640	
0100242	0101122	0101233	0102242	0111530	0117650	
0101000	0101123	0101242	0103032	0111540	0895002	
0101010	0101142	0101243	0103042	0111550	0895012	
0101020	0101143	0101244	0103052	0111560	0895022	
0101030	0101152	0102032	0103062	0111580	0895202	

The conformity assessment procedure has been carried out in accordance with Annex III of Directive 98/78/EC.
The above mentioned in-vitro diagnostic devices are neither covered by Annex II nor intended for self-application.

This declaration is valid until 31st December 2019.

Best regards,

Harry Marienfeld
Managing Director
Lauda-Königshofen, 27th January 2017