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Certificate of Conformity (CoC)

According to DIN EN ISO 10993-5:2009-10

Certificate No.: A2 20231025 003
Test Report Ref. No.: 23AA2292 Version 1

Certificate Holder

Manufacturer Address	Notter GmbH In den Erlen 10 D-75248 Ölbronn
Contact Person ^{#1}	Mr. Fabian Seifried
Order number	23AA2292
Issuing date of the test report	16.10.2023
Issuing date of the CoC	25.10.2023

Product

Product	TiCN (redish colour)								
Date of receipt of the product	22.09.2023								
Condition	The test samples arrived undamaged and individually packed.								
Identification	<table border="0"> <tr> <td>Description</td> <td>TiCN (redish colour) from 18.09.2023</td> </tr> <tr> <td>Process:</td> <td>TiCN300_Verd_GHI-DEF_Planet_Locator_violett</td> </tr> <tr> <td>Lot No.</td> <td>C, TiCN_23-013</td> </tr> <tr> <td>Layer thickness</td> <td>1.7 ± 0.1 µm</td> </tr> </table>	Description	TiCN (redish colour) from 18.09.2023	Process:	TiCN300_Verd_GHI-DEF_Planet_Locator_violett	Lot No.	C, TiCN_23-013	Layer thickness	1.7 ± 0.1 µm
Description	TiCN (redish colour) from 18.09.2023								
Process:	TiCN300_Verd_GHI-DEF_Planet_Locator_violett								
Lot No.	C, TiCN_23-013								
Layer thickness	1.7 ± 0.1 µm								

Tested according to

Standard / Guideline	DIN EN ISO 10993-5:2009-10 21 CFR Part 58 (Gesetz zum Schutz vor gefährlichen Stoffen [Chemikaliengesetz – ChemG] § 19b Abs.1 Chemikaliengesetz)
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This certificate refers to the above mentioned product. This is to certify that the test sample is in conformity with the requirements of DIN EN ISO 10993-5:2009-10 **to confirm biocompatibility regarding cytotoxicity**. The holder of the certificate is entitled to use this certificate for the declaration of conformity according to DIN EN ISO 10993-5:2009-10 and 21 CFR Part 58. This certificate does not imply assessment of the production of the product. Detailed specification of the tested product are shown in the above mentioned test report.

This certificate is valid provided that the test specifications and construction of the product remain unchanged and refers only to the product tested as received by the laboratory and cannot be transferred to other products.

Ansbach, 25.10.2023



Dr. Katrin Sulzer
Head of BioLabs

Dokumentnummer	Version	Autor	freigegeben	Dokumentenquellname
FB-BL-106	V2	KS	PH 19.10.2021	FB-BL-106_CoC_23AA2292_2023-10-25_V1