ARTZT
quality sports and health equipment

KONFORMITÄTSERKLÄRUNG
DECLARATION OF CONFORMITY

1) **Manufacturer** *(Name, department)*:

Name: Ludwig Artzt GmbH
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2) **Products** *(name, type or model/batch number, etc.)*:

Product Category according to (EU) 2017/2185:
MDN1205 (Non-active, non-implantable, orthopedic and rehabilitation devices)

Risk Class: 1
Rule: 1
CND Code: Z120602
Basic UDI-DI: 426007163LA6001N3

Device Family	Generic Device Group	Product Code	Device Name	Variant
Massage Tools	Instrument Assisted Massage	LA-7010	ARTZT thepro Fazer	Set 1-4
		LA-7011	ARTZT thepro Fazer	1
		LA-7012	ARTZT thepro Fazer	2
		LA-7013	ARTZT thepro Fazer	3
		LA-7014	ARTZT thepro Fazer	4
		LA-7015	ARTZT thepro Fazer	5

3) The products described above are in conformity with:

Document No.	Title	Edition/Date of Issue
2017/745	Medical Device Regulation	Apr 5th 2017

Dornburg, Germany, June 21st 2024

Felix Artzt, CEO, Ludwig Artzt GmbH