



AFINA s.r.o.  
Consulting Agency

# Certificate of CE-registration

This is to certify that, in accordance with either medical device Regulation MDR 2017/745,

**AFINA s.r.o.**

agree to perform all duties and responsibilities as the Authorized Representative for

**"ELASTIKUM" Company Limited Liability**

**Address: Yangikurgon-2 28, Toshkent viloiati, 111305, Uzbekistan**

**SRN: UZ-MF-000036753**

as stipulated and demanded by the afore-mentioned Regulation. The Czech competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

Registration number	Name of medical device	Registration number of manufacturer	Manufacturer
01184105	Medical bleached cloth gauze non-sterile in pieces.	082747	"ELASTIKUM" Company Limited Liability
	Identification code variants	Name add-on	Catalogue number
	001	84 cm x 1 m	
	002	84 cm x 2 m	
	003	84 cm x 3 m	
	004	84 cm x 5 m	
	005	84 cm x 10 m	
	006	84 cm x 25 m	
	007	90 cm x 1 m	
	008	90 cm x 2 m	
	009	90 cm x 3 m	
	0010	90 cm x 5 m	
	0011	90 cm x 10 m	
	0012	90 cm x 25 m	



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The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfil the essential

requirements of either Regalement MDR 2017/745. A safety officer has been appointed for Czechia and therefore is in full compliance with § 31 MPG.

Signed on 10. 10. 2023



Ing. Frejdlin Arkadij  
Director