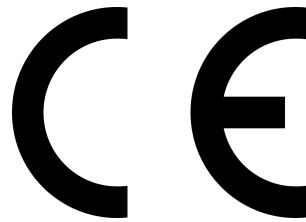




Certificate of CE-registration



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

AFINA s.r.o.

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

DGNCT LLC

Southeast 2nd Avenue, 333, Miami, FL, 33131, USA

as stipulated and demanded by the afore-mentioned Regalement. 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
01174942	Diagnocat AI	082229	DGNCT LLC

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 1. 9. 2023



Ing. Frejdlin Arkadij
Director