



CERTIFICATE

This certifies that the Quality management system for medical devices of company

Dr. Müller Pharma s.r.o.

U Mostku 182, 503 41 Hradec Králové, Czech Republic

**Dr. Müller
PHARMA**

has been assessed by 3EC International and found to be in conformance with the following standard:

EN ISO 13485:2016

for the following scope:

DESIGN AND DEVELOPMENT, MANUFACTURING AND DISTRIBUTION OF NON-ACTIVE MEDICAL DEVICES: LUBRICANT GELS, VAGINAL GELS, ULTRASOUND GELS, NASAL SPRAYS, SUPPOSITORIES AND LICE REMEDY

Certificate No.: M-0388/19

Date of issuance: November 20th, 2019

Original date of approval: December 1st, 2011

This certificate is valid from **November 30th, 2019** to **November 29th, 2022** on condition that organization will maintain effective Quality management system for medical devices. To verify the validity of this certificate please contact our office at: +421 (0)2 5831 8343.

Issuing office: 3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovak Republic




Dr. Katarína Tomin Srdošová
Head of Certification Body 3EC International a.s.

Certification body 3EC International a.s. is accredited by SNAS under registration number 305/Q-054 with accreditation certificate No. Q-054 for certification of Quality management systems for medical devices.