

CERTIFICATE

NO: AQS/Q/175512026

SI CERTIFICA CHE IL SISTEMA DI GESTIONE PER QUALITÀ DI:
WE HEREBY CERTIFY THAT THE QUALITY MANAGEMENT SYSTEM OPERATED BY:

MYLAV S.r.l.

P.IVA / VAT: IT04197610753

SEDE LEGALE / LEGAL HEADQUARTERS:
VIA VECCHIA MONTESARDO, 21, ALESSANO 73031 - LE - ITALY.

SEDE OPERATIVA / OPERATIONAL HEADQUARTERS:
VIA SIRTORI, 9 FRAZ. PASSIRANA, RHO 20017 - MI - ITALY.

E' CONFORME ALLA NORMA / IS IN COMPLIANCE WITH THE STANDARD

ISO 9001:2015

PER LE SEGUENTI ATTIVITÀ / FOR THE FOLLOWING ACTIVITIES

IT: Laboratorio di analisi veterinarie con settori di specializzazione in: chimica clinica, ematologia, endocrinologia, microbiologia, biologia molecolare, genetica e genetica oncologica, immunologia e immunologia oncologica, istologia, citologia e sierologia.

ENG: Veterinary analysis laboratory with fields of specialization in: clinical chemistry, hematology, endocrinology, microbiology, molecular biology, genetics and oncological genetics, immunology and oncological immunology, histology, cytology and serology.

IAF Code: 38.

VALIDITÀ CERTIFICATO / VALIDITY OF THE CERTIFICATE

THE USE AND THE VALIDITY OF THE CERTIFICATE SHALL SATISFY THE REQUIREMENTS OF THE RULES FOR CERTIFICATION OF MANAGEMENT SYSTEMS

FIRST ISSUE DATE: 05/06/2020

LATEST ISSUANCE DATE: 05/06/2026

EXPIRY DATE: 04/06/2029

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For Certification Body

Eriola Curo

CERTIFICATE

NO: AQS/DM/175522026

SI CERTIFICA CHE IL SISTEMA DI GESTIONE DELLA QUALITÀ PER DISPOSITIVI MEDICI DI/
WE HEREBY CERTIFY THAT THE QUALITY MANAGEMENT SYSTEM FOR MEDICAL DEVICES OPERATED BY:

MYLAV S.r.l.

P.IVA / VAT: IT04197610753

SEDE LEGALE / LEGAL HEADQUARTERS:
VIA VECCHIA MONTESARDO, 21, ALESSANO 73031 - LE - ITALY.

SEDE OPERATIVA / OPERATIONAL HEADQUARTERS:
VIA SIRTORI, 9 FRAZ. PASSIRANA, RHO 20017 - MI - ITALY.

E' CONFORME ALLA NORMA / IS IN COMPLIANCE WITH THE STANDARD

ISO 13485:2016

PER LE SEGUENTI ATTIVITA' / FOR THE FOLLOWING ACTIVITIES

IT: Progettazione, sviluppo e immissione in commercio di dispositivi medico diagnostici in vitro consumabili di uso generale in plastica riciclata (materiale plastico green) per analisi di campioni. Progettazione, produzione e immissione in commercio di dispositivi medici su misura non sterili fisici e virtuali per supporto alla pianificazione chirurgica, alla diagnosi e relativi servizi di assistenza con la fabbricazione additiva per la visualizzazione di modelli 3D specifici per il paziente da utilizzare in fase pre ed intra operatoria e le relative attività di assistenza e post-vendita.

ENG: Design, development, and marketing of general-purpose in vitro diagnostic consumable medical devices made of recycled plastic (green plastic) for sample analysis. Design, production, and marketing of custom, non-sterile physical and virtual medical devices to support surgical planning, diagnosis, and related support services using additive manufacturing for the visualization of patient-specific 3D models for pre- and intra-operative use, as well as related support and post-sales services.

Categorie / Categories: A.1.1.B; A.1.2; A, A.1.4, A.1.7

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For Certification Body

Eriola Cuni