

Compliance statement regarding MDR 2017/745 and IVDR 2017/746

INTEGRA Biosciences is committed to meeting all applicable legal and regulatory requirements in the markets where we operate. We place great emphasis on the safety and quality of our products and ensure that all relevant regulations pertaining to our product categories and services are fully adhered to.

Although INTEGRA is not subject to the requirements of EU Regulations MDR 2017/745 and IVDR 2017/746, we still maintain the highest standards in terms of product safety, quality, and regulatory compliance. Our processes align with best practices and international standards to meet the requirements of the respective markets and the products we offer.

We are committed to continuously monitoring the quality and safety of our products and ensuring that they consistently meet the applicable requirements and standards.

INTEGRA Instruments	
ASSIST	MINI 96
ASSIST PLUS	MIRO CANVAS
COLDPLATE and BIOSHAKE	PIPETBOY Acu 2 / PRO / GENIUS
D-ONE Pipettes	VACUBOY
DOSE IT	VACUSAFE
EVOLVE Pipettes	VACUSIP
FIREBOY	VIAFLO 96 / 384
FLAMEBOY	VIAFLO Pipettes
MAG and HEATMAG	VOYAGER Pipettes
MEDIACLAVE 10 / 30	WELLJET
MEDIAJET (VARIO)	SWITCH

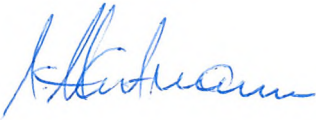
INTEGRA Consumables	
12 well reagent reservoirs	INTEGRA serological pipets
Automation GRIPTIPS®	Multichannel reagent reservoirs
Automation friendly reagent reservoirs	Divided reagent reservoirs
ECO rack GRIPTIPS® for handheld pipettes	

INTEGRA Magnetic Beads	
MAGFLO™ NGS	MAGFLO™ PCR

These above-mentioned INTEGRA Products are not classified as medical device or in vitro diagnostic medical devices in the country of origin and the European Union, and they do not fall under MDR 2017/745 or IVDR 2017/746.

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Zizers, August 2025



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If you require further information or a more detailed explanation on a specific aspect, please do not hesitate to contact us!

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