Mphi MDR family
is available!

Multiwave LockedSystem

Discover
MLS® Laser Therapy

ALWAYS BY YOUR SIDE.

Mphi MDR devices are reliable and easy to use. Now more than ever.

Expanded application modalities: all programs (preset and personal) can be performed not only in Point-by-Point and Scan mode, but also in fixed point modality using the Charlie multi-diode applicator on the special arm.

Extreme versatility: the Trigger Point mode is always available through the handpiece for all protocols, while the laserpuncture mode with conical lightguide is only available for Mphi 75 MDR.

High level of professionalism: through constant dosage control (J/cm^2), the user can **objectify and replicate successful therapies**.

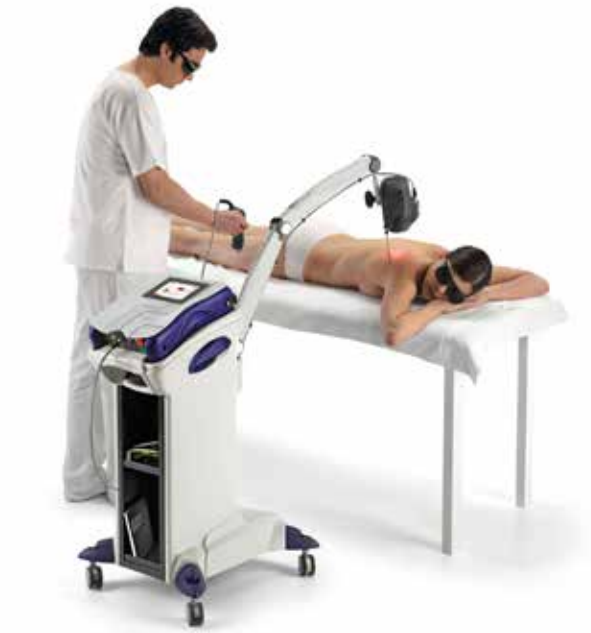
Intuitive approach: thanks to the EU MDR protocols, MLS® treatment modalities are based on an extremely intuitive therapeutic approach centred on the **Target Tissues.**

Completely renewed user experience
New capacitive touch screen, increased sensitivity and improved usability of the SW interface.

New SW and new media library
The software is completely renewed, is available in 26 languages and provides the user with +400 new images and videos of specific treatments according to the chosen protocol, application mode and optical terminal.

Increased reliability
The new air cooling system is silenced and thermostat-controlled to keep the laser sources in optimal working condition over time.

New electronics and sensors
Mphi MDR family offers improved performance, speed and memory, advanced control sensors and warning lights for increased reliability and safety.

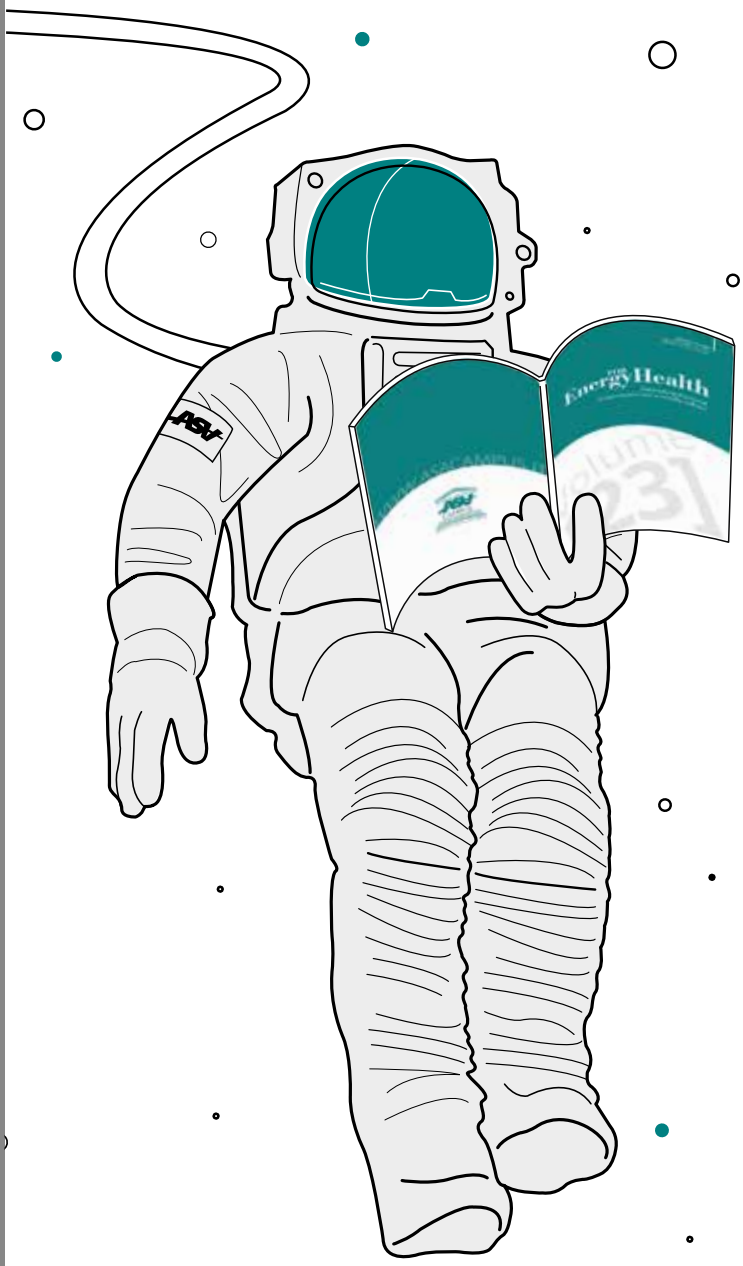


10/2024 – EN

Choose EU MDR devices!

European Union Medical Device Regulation (EU MDR) certified medical devices ensure **quality**, **safety** and **effectiveness** of the systems and treatments offered to patients.

Here's why investing in **ASA EU MDR medical devices** is the best choice.



PATIENT SAFETY AND PROTECTION

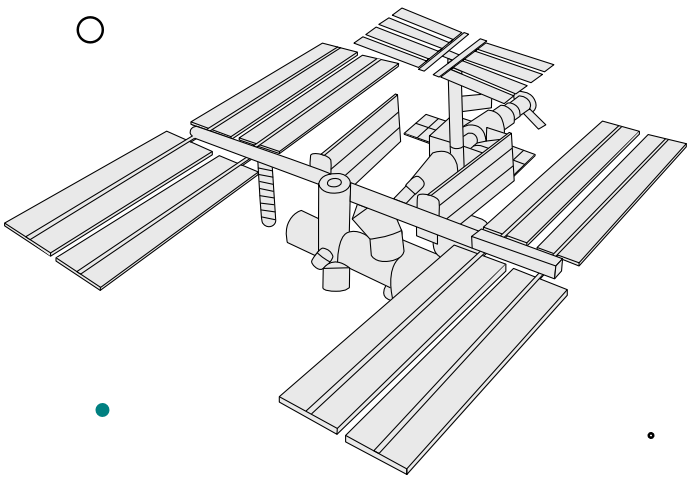
EU MDR ensures that only tested and safe devices are on the market: with more than **30,000 devices** worldwide and **over 200 publications**, **ASA** stands for **efficacy and safety**. ASA systems are scientifically and clinically validated through extensive **research** work at **ASAcampus**.

AMONG THE TOP OF THE CLASS

ASA **Quality System** is certified by **TÜV SÜD** Product Service GmbH. ASA received **EU MDR** certification in **April 2023**. ASA EU MDR systems undergo **rigorous compliance checks and audits**, thus ensuring durable, reliable and effective Therapeutic Solutions.

RIGOROUS CLINICAL EVALUATION

To demonstrate efficacy and safety, ASA EU MDR devices undergo extensive evaluations based on **robust clinical data.** EU MDR requires maximum **transparency from the manufacturer to the patient**: ASA has long adopted **regular monitoring** of its devices worldwide.



Did you know that some experiments conducted by ASAcampus were performed on the International Space Station?