**Eurofins Medical Device Services Europe acquires Italian Studio Ambiente**

Milan, 28 January 2026 – Eurofins Medical Device Services Europe, a leading provider of comprehensive testing, packaging, sterilisation and consulting services for medical devices is proud to announce the acquisition of Studio Ambiente, a leading testing and consultancy partner for medical devices in Dossobuono di Villafranca, Italy.

This strategic acquisition reinforces Eurofins Medical Device Services Europe's commitment to offering the best possible medical device safety and performance testing to clients across Europe.

With this acquisition, Eurofins Medical Device Services Europe continues to build on its legacy of scientific excellence and customer-centric service, offering clients a broader portfolio of high-quality and bespoke consulting and testing solutions and sterile packaging services, supporting faster access to market and strengthening support for medical device manufacturers across the entire product lifecycle. Studio Ambiente, with over 25 years of experience, offers certified microbiological testing and integrated regulatory consulting for medical devices, pharmaceuticals, and cosmetics. Combining ISO 9001/13485 quality systems with global market access expertise, it supports manufacturers from product development through CE marking, FDA 510(k), and MDSAP certification.

**About Studio Ambiente S.r.l.**

Founded in 1996 and headquartered in Dossobuono di Villafranca, Italy, Studio Ambiente is a leading testing and consultancy partner for medical devices, pharmaceuticals, industrials, and cosmetics.

The company offers a wide range of testing services with an in-house microbiological laboratory accredited ISO 17025 and certified according to ISO 9001, ISO 13485 and GLP, for routine monitoring of products and validation of special processes, as well as analysis of production sites, in compliance with international standards.

Studio Ambiente also supports manufacturers from product development through global market access, offering services such as quality systems implementation, CE marking and FDA 510(k).

Its multidisciplinary team of biologists and biomedical engineers ensures comprehensive solutions for non-active, implantable, and active medical devices, backed by decades of technical and regulatory experience.

**About Eurofins Medical Device Services Europe**

Eurofins Medical Device Services Europe, part of the global network of Eurofins laboratories, has over 40 years of experience in medical device testing, consulting, and sterile packaging. With more than 20 laboratories and 2 sterile packaging sites across Europe, North America, and Asia Pacific, Eurofins provides end-to-end support for Class I, II, and III medical devices. Its services cover biocompatibility, microbiology, mechanical testing, regulatory consulting, and packaging validation, all compliant with cGMP, GLP, and ISO 17025 standards. Eurofins is dedicated to helping manufacturers bring safe, effective, and compliant products to market quickly and efficiently. https://www.eurofins.com/medical-device

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