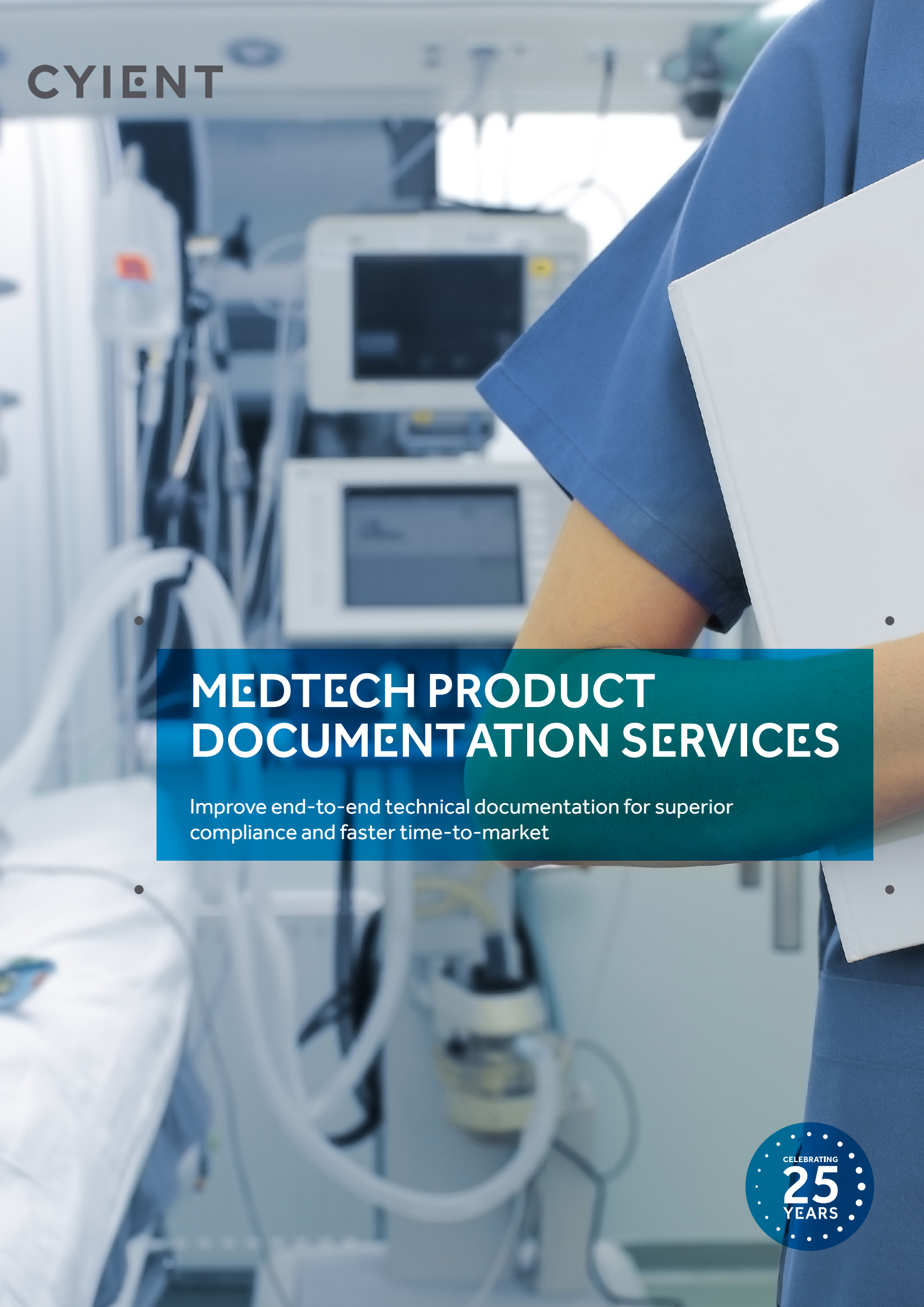
CYIENT



MEDTECH PRODUCT DOCUMENTATION SERVICES

Improve end-to-end technical documentation for superior compliance and faster time-to-market

CELEBRATING 25 YEARS

The stringent regulatory and industry standards such as FDA, CE, and ISO 13485 put enormous pressure on medical device original equipment manufacturers (OEMs) to produce accurate and concise documentation that demonstrates quality, design, risk management, and marketing compliance. This is a major challenge given the increasing product complexity and multiple design changes that typically lead to several versions of the product. The need to convert documents from paper-to-digital, create illustrations, and employ resources with a grasp of various software and tools further compounds the challenge.

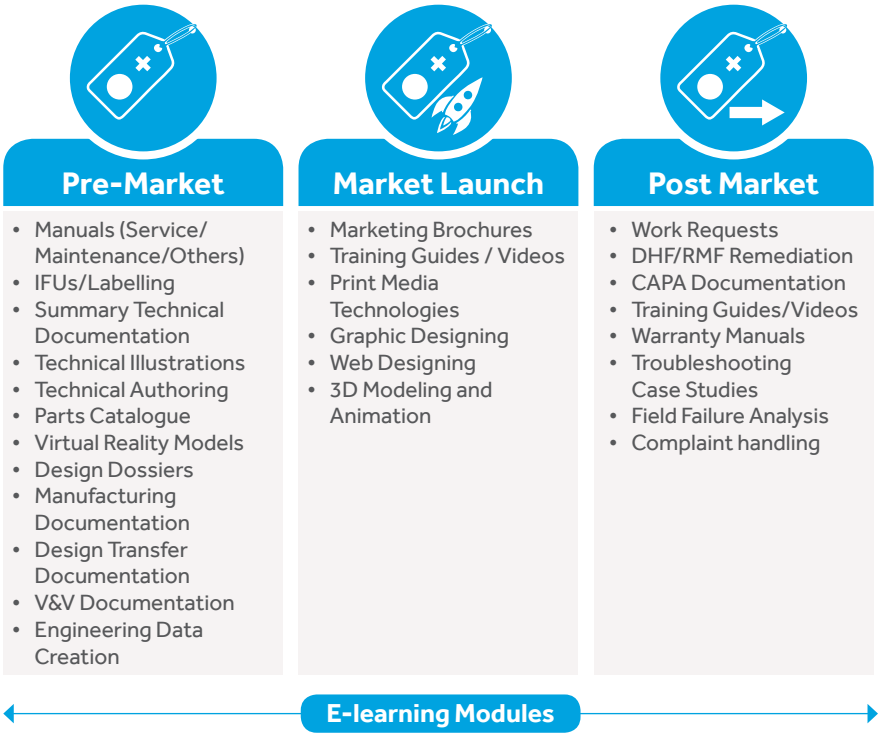
Cyient Portfolio:

Comprehensive Product Documentation Services

Cyient provides end-to-end product documentation support across the entire medical device product life cycle—from pre-market to market launch and post-market. Our streamlined, easy-to-use documentation services enable efficient content creation and management, resulting in enhanced compliance, faster time-to-market for products, and cost optimization. We closely work with the product development team to completely align the documentation services leading to shorter development cycles. Our sound yet simplified documentation practices help you deliver a superior product experience to your customers, and reduce warranty/support costs as well as the risk of liability.

Offerings in Medical Technology and Healthcare Space

We deliver end-to-end technical documentation throughout the product life cycle in the MedTech space



What Sets Us Apart

We provide leading innovation and quality while remaining affordable. Our team of domain experts ensure superior service delivery aligned with your business needs through our ready-to-use infrastructure, methods, and processes. By supporting product documentation that spans the entire product life cycle, we enable you to simultaneously drive compliance and innovation, thereby facilitating sustained growth and success. Our one-stop solution is underpinned by deep medical device domain expertise and more than 5.4 million man-hours of cross-industry experience. We house a dedicated Product Documentation Center of Excellence (CoE) for one of the leading in-vitro diagnostics products companies.

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