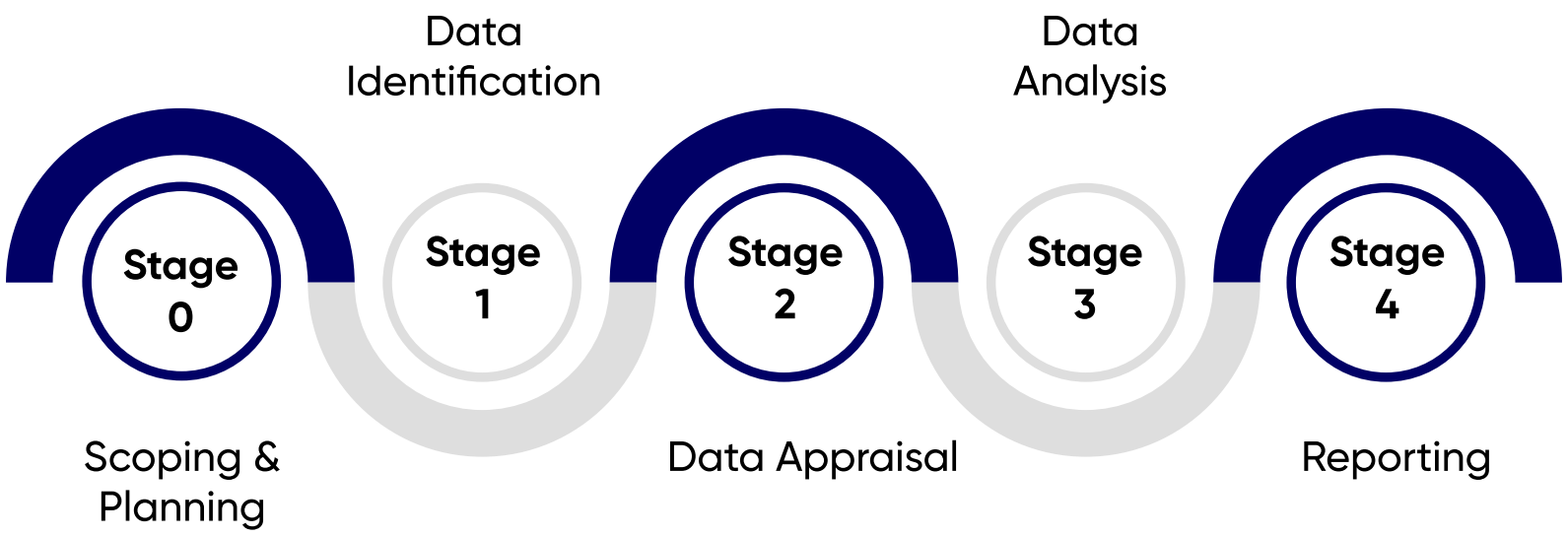
Clinical Evaluation Support

Your Path to CE Marking, Made Easy

At COD Research, we provide a comprehensive suite of services for Medical Devices, ensuring compliance with applicable laws and regulations while supporting manufacturers in meeting the highest quality and safety standards.

With our expertise, you gain a reliable partner to ensure your CERs are compliant, evidence-based, and aligned with the latest regulatory standards.



From planning and literature review to CER writing, gap analysis, and PMS integration, we help your team create compliant, regulator-ready Clinical Evaluation Reports. Our services ensure seamless updates, collaborative workflows— all aligned with MEDDEV 2.7/1 Rev. 4 and the EUMDR.

We offer a full spectrum of services tailored to meet the unique needs of medical device safety. Our expertise includes:

- Drug-Device Combinations: Drug-Eluting Stents, Inhalers, Transdermal Patches etc.
- Invasive Devices: Surgical Instruments, Implants etc.
- Non-Invasive Devices: Diagnostic Imaging Equipment, Therapeutic Devices etc.

CLINICAL EVALUATION SUPPORT

Our Service

Clinical evaluation plan

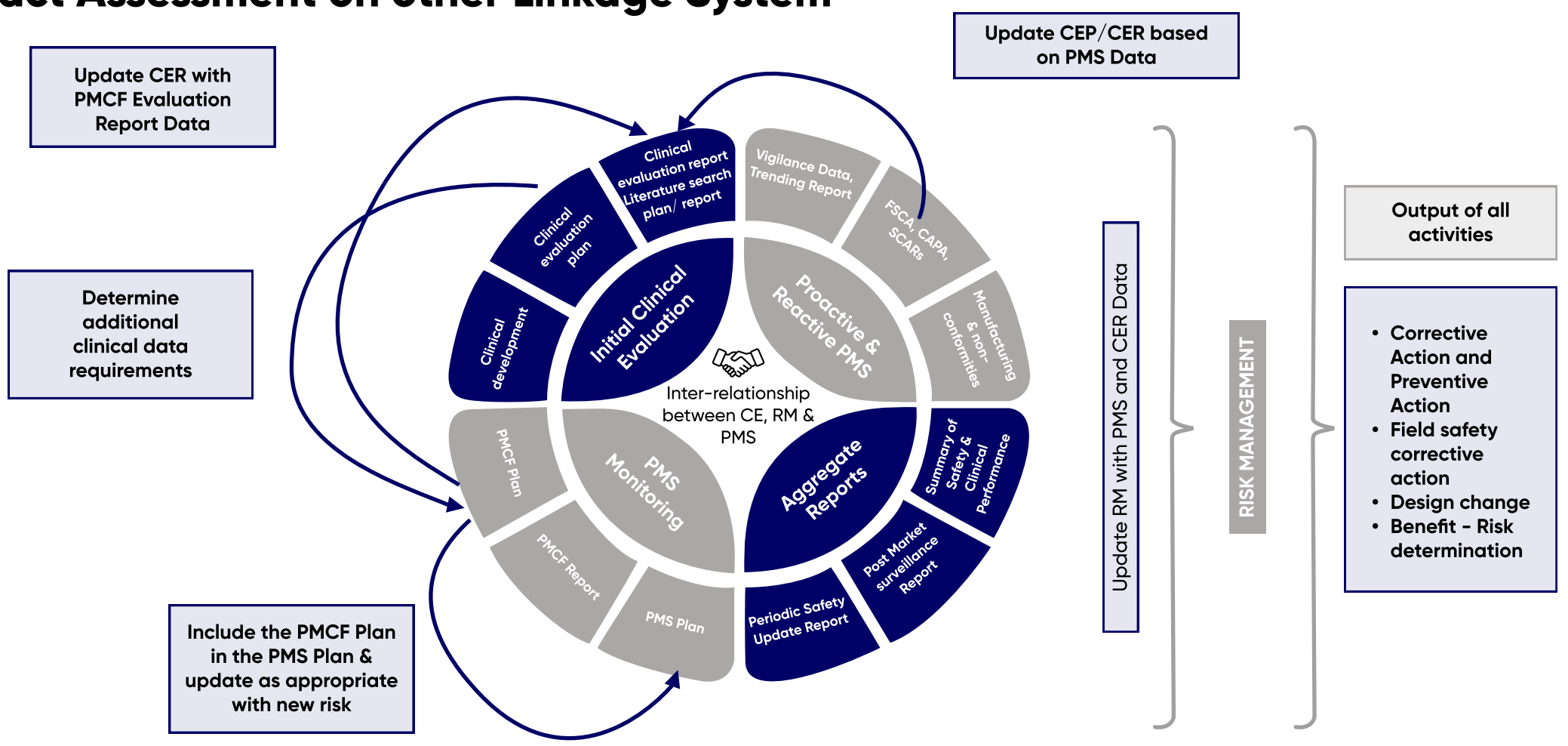
Clinical evaluation report

Literature search plan & report

- Systematic analysis
- State-of-the-art (SOTA) analysis
- With quantitative and qualitative analysis (PRISMA & Cochrane)

Regulatory database search plan & report

Impact Assessment on other Linkage System



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