

WHY PROFESSIONAL REGULATORY WRITING MATTERS

Regulatory submissions require precision, compliance, and clarity. Professional regulatory writing ensures adherence to global guidelines, accurate documentation, and higher success rates in approvals from regulatory authorities such as FDA, EMA, and ICH.

WHAT WE OFFER



Regulatory Document Preparation (CTD, eCTD, IND, NDA)



Clinical Study Reports (CSR) Writing



Investigator's Brochure (IB) Development



Protocol & Synopsis Writing



Regulatory Submission Support & Publishing



Medical & Scientific Writing Compliance

PUBRICA'S PROCESS



Requirement Analysis & Regulatory Strategy



Document Planning & Structuring



Regulatory Writing & Draft Development



Quality Review & Compliance Check



Final Submission & Publishing Support

WHY CHOOSE PUBRICA?



Experienced Regulatory Writing Experts



Compliance with FDA, EMA, ICH Guidelines



High-Quality Scientific Documentation



Confidential & Reliable Service



Timely Delivery with Accuracy

CONTACT



sales@pubrica.com



+44 161 394 0786(UK)
+1-972-502-9262(USA)
+91 98843 50006(IN)



www.pubrica.com