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This guidance document is intended to assist pharmaceutical companies with the submission of regulatory information in electronic Common Technical Document format (eCTD) to the National Competent Authorities

Harmonised Technical Guidance for eCTD Submissions in the EU

The electronic Common Technical Document (eCTD) allows for the electronic submission of the Common Technical Document (CTD) from applicant to regulator. While the table of content is consistent with the harmonised CTD, the eCTD also provides a harmonised technical solution to implementing the CTD electronically.

eCTD v3.2 - eSubmission: Projects

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10 mistakes to avoid for eCTD success

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