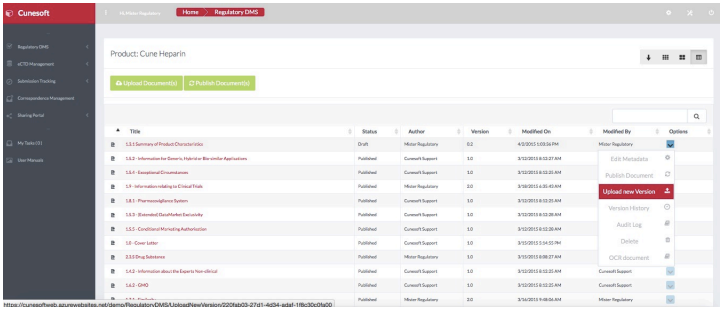
cune-RDMS

cunesoft
compliance delivered

- Expedites the creation of new documents and the uploading of existing documents including large content libraries efficiently from outside the system.
- Configures review and approval workflows to support paperless, efficient document creation, editing and approval processes.
- Documents electronic/digital signatures according to FDA requirements published in 21 CFR part 11.
- Tags documents with ICH and/or xEVMPD specific metadata as required.
- Navigates through large content libraries with ease using full text search, filtering mechanisms and personalized views.
- Converts documents into eCTD compliant PDF versions automatically with OCR option once the submission stage is reached.
- Controls document versions and allows only "submission ready" documents to be used in the cune-eCTD module.
- Recognizes and addresses the need for a more collaborative workspace where both internal and external users (even those working in various regions or remotely) can work more effectively.

Regulatory Software Online

cune-RDMS	cune-eCTD	cune-Track	cune-SOP



Cut submission time

Save 30-50% in license costs

Cunesoft Advantages

Reduce maintenance costs

Cut audit preparation time

About Cunesoft: We offer a ready to use, pre-validated and highly secure cloud based regulatory compliance suite. Our software has been built to fulfill regulatory needs holistically and end to end. Our vision is to keep it as simple as possible while delivering compliance and fulfilling FDA 21CFR11, EU Annex 11, ICH, and GxP requirements.

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