**Revised Script Option 5: 3010 characters, 329 words**

Many content authors in the Life Sciences are struggling to keep up.

They rely on multiple systems and methods to create and update their content, and this can produce several issues when it comes to the organization and management of their writing. Furthermore, the old way of copying and pasting is prone to error, making it impossible to stay on schedule. There is simply no way of knowing when source content changes, leading to errors and delays when updates need to be made. The dizzying headache that comes from this mish-mash approach to content management is only made worse for many content authors because they still rely on email for document reviews and approvals.

When we look at the situation, we see that the process of authoring content has become stuck in the dark ages, whereby inefficient and antiquated methods are the norm. What if you could simply reuse your approved content while saving massive amounts of time authoring and updating your documents?

Meet Docuvera.

Built specifically for the Life Sciences industry, Docuvera provides a single platform to author, review, approve, translate, and publish regulated documents by reusing approved, component-based content. It combines an intuitive interface with the latest web technologies to provide a powerful, yet easy-to-use service that alleviates the pain of traditional document creation.

Docuvera allows authors to reuse approved blocks of content to assemble documents without the need for technical expertise.

And, when reused content changes, Docuvera alerts authors so they can seamlessly incorporate updates into their documents.

Collaboration between departments is enhanced via an online interface for reviews, discussions and resolutions. No more manual merging of Word track changes.

With our omni-channel approach authors can create Word, PDF, HTML and other documents from the same content by applying different styles during the publishing process. No more manual creation needed.

Docuvera provides detailed reporting and audit trails enabling your team with the most advance ways of storing a document history for compliance and other reasons.

Your document creation efforts will be reduced while your output is dramatically increased. All without having to add additional staff. Docuvera’s end-to-end solution reduces content production times, ensures alignment for compliant content, dramatically reduce human errors and creates faster time-to-market for regulatory submissions.

Ready to see what Docuvera can do for you? Get a demo today.