

STRUCTURED COMPONENT AUTHORIZING WILL TRANSFORM HOW YOU CREATE AND MANAGE YOUR DOCUMENTATION.

Antiquated systems lead to massive inefficiencies in creating Life Sciences documentation.

Cut-paste thousands of pages—\$325 MM in documentation costs per new drug launch

Email draft documents for review—Increased compliance violations, work stoppages

Final documents isolated on hard drives—20 hours/week lost due to searching, recreating content

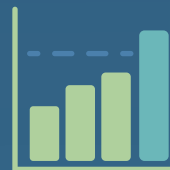


What if you could change all this by implementing structured component authoring? SCA can achieve:



82% CONTENT REUSE

With SCA, companies achieve high levels of reuse of existing, approved content – thus eliminating the need to create/review/approve/translate redundant content.



110% PRODUCTIVITY GAINS

Improved efficiencies allow existing teams to take on more initiatives without adding staff.



12-MONTH BREAK EVEN

With an SCA solution that generates broad productivity gains while achieving widespread adoption, companies reach rapid break even of initial investment in implementation and services costs.



145% ROI

A typical customer achieved a NPV of \$3.2MM and a 145% ROI, as users were able to devote time saved to new projects – and financial benefits will only increase with wider adoption and new use cases.

**Docuvera is SCA designed specifically for Life Sciences.
Let us show you what Docuvera can do for you.**



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