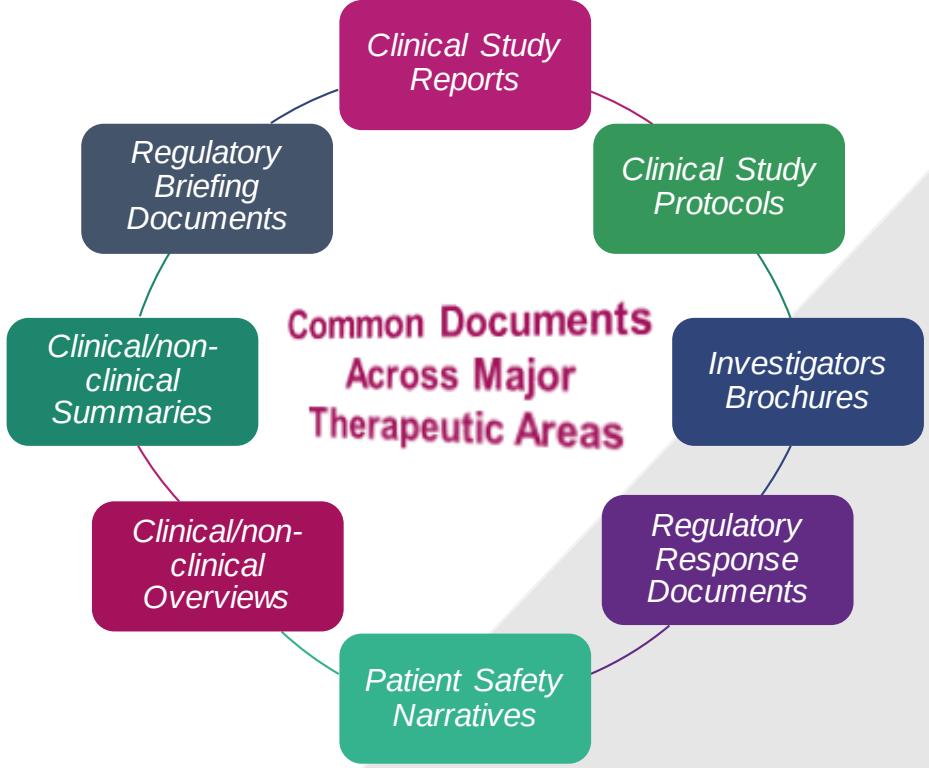
Kinapse Document QC Service

Poor quality documents pose a risk to patient safety, cause delays in regulatory review and result in wasted effort and budget.

"Consistently achieving Right first time" quality is becoming ever more challenging due to increasing:

1. *Number of clinical trials*
2. *Amount of regulatory documentation per trial*
3. *Authoring via contractors or vendors outside of the direct control of the sponsor*



Many sponsors recognize that independent Quality Control (QC) of documents improves the quality and efficiency of the regulatory document process.

8
Years delivering QC services

100%
Success rate in quality and timelines

>3900
Documents QC reviewed

4
of top 10 pharma companies are clients

Value of Independent QC

- *Analysis of QC findings by client to identify specific training and coaching needs for authors*
- *Allows a client's internal medical writers to focus on document authoring and submission.*
- *Objective perspective of independent QC*

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