Accelerating Clinical Submissions via Text Analytics AI

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Regulatory submissions are a critical hurdle that every biopharmaceutical company must overcome to conduct clinical trials, commercialize products and maintain products in the market. The majority of all submission types carry human made errors which are costing Pharma, and CROs valuable time and significant cost to amend.

BeaconCure offers a unique Text Analytics solution to address submission documentation quality control needs. With BeaconCure, Pharma and CRO companies can quickly and effectively validate the summary data against the sources, correct inconsistencies in clinical data tables and text, and produce error free submissions ready for regulatory approval.

Direct Key benefits to Pharma and CROs:

1. **Accelerated Turnaround:** dramatically reduce months of Quality Control team work to a few days
2. **Eliminate Inconsistences:** Confirm summary documentation is clean of human errors and fully backed up by the source data collected during the trial