We can design and conduct your clinical studies and help you submit them to FDA according to all regulatory requirements.

The Study Data Technical Conformance Guide posted by FDA on November 2016, identifies specific rules for using CDISC standards on FDA submissions. These requirements are mandatory to all studies initiated after December 16, 2016.

Blueclinical has submitted several clinical studies with SAS datasets prepared in accordance with the latest FDA CDISC specifications. To sum up, we are happy to let you know that **Blueclinical is ready for your next FDA application process!**

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