EXPEDITE YOUR TRIAL DATABASE BUILDS AND INCREASE TRIAL EFFICIENCY WITH FIRMA CLINICAL DATA SERVICES

Clinical trial database build times affect downstream data management tasks.

Data entry cannot be performed if the database is not available at first patient first visit (FPFV). Quicker, high-quality database builds allow for faster launch and entry of clinical data.



Database build and release take an average 68 days for most companies¹



Retrospective data entry delays database lock



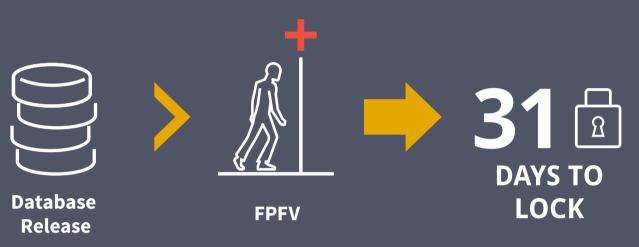




Database Release



83% of life science companies experience an average of 54 days to database lock when database release takes place *after* FPFV.



17%

83%

17% of life science companies experience an average of 31 days to database lock when database release takes place *before* FPFV.

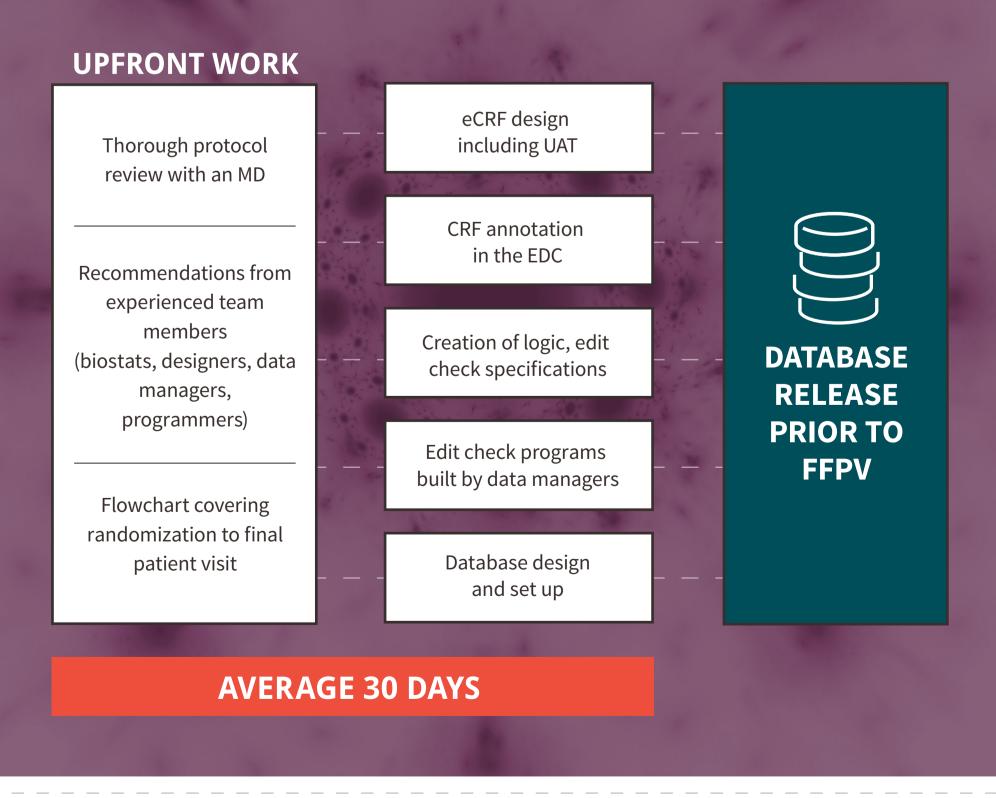
Firma's efficient process drives fasterdatabase builds

We've taken the typical sequential build process and **pre-loaded it with work and process enhancements**, expanding efficiencies and resulting in a **30-day build and release timeline**.

30 DAYS with Firma's efficient process



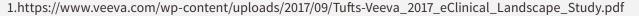
Communication and collaboration are key to expedite database build times. At Firma, we believe in communicating directly with the data management team throughout the entire build process.



DATA SERVICES

Leverage our expertise and training to shorten your database build. Schedule a call with our Data Services Leadership team today: email us at **info@firmaclinical.com** or visit **firmaclinical.com**.





2.https://www.fda.gov/forpatients/approvals/fast/ucm20041766.htm