The client needed to run 3 healthy-normal trials at one Celerion clinic in addition to using two hospitals to run a hepatic and a renal study.

Utilized Celerion’s data management services for the 5 study program.

Experienced data managers designed Case Report Forms (CRFs) for the hospital data collection that was compatible with the electronic CRFs generated using Celerion’s proprietary ClinQuick® system.

Data programmers were able to utilize a consistent format for all 5 studies that followed the Clinical Data Interchange Standards Consortium (CDISC) Study Data Tabulation Model (SDTM) format.

Celerion was able to deliver a quality, consistent database for all 5 studies, achieving the clients’s needed database lock timelines.