

Pharmaceutical and Device-Multi Center Clinical Trial Algorithm

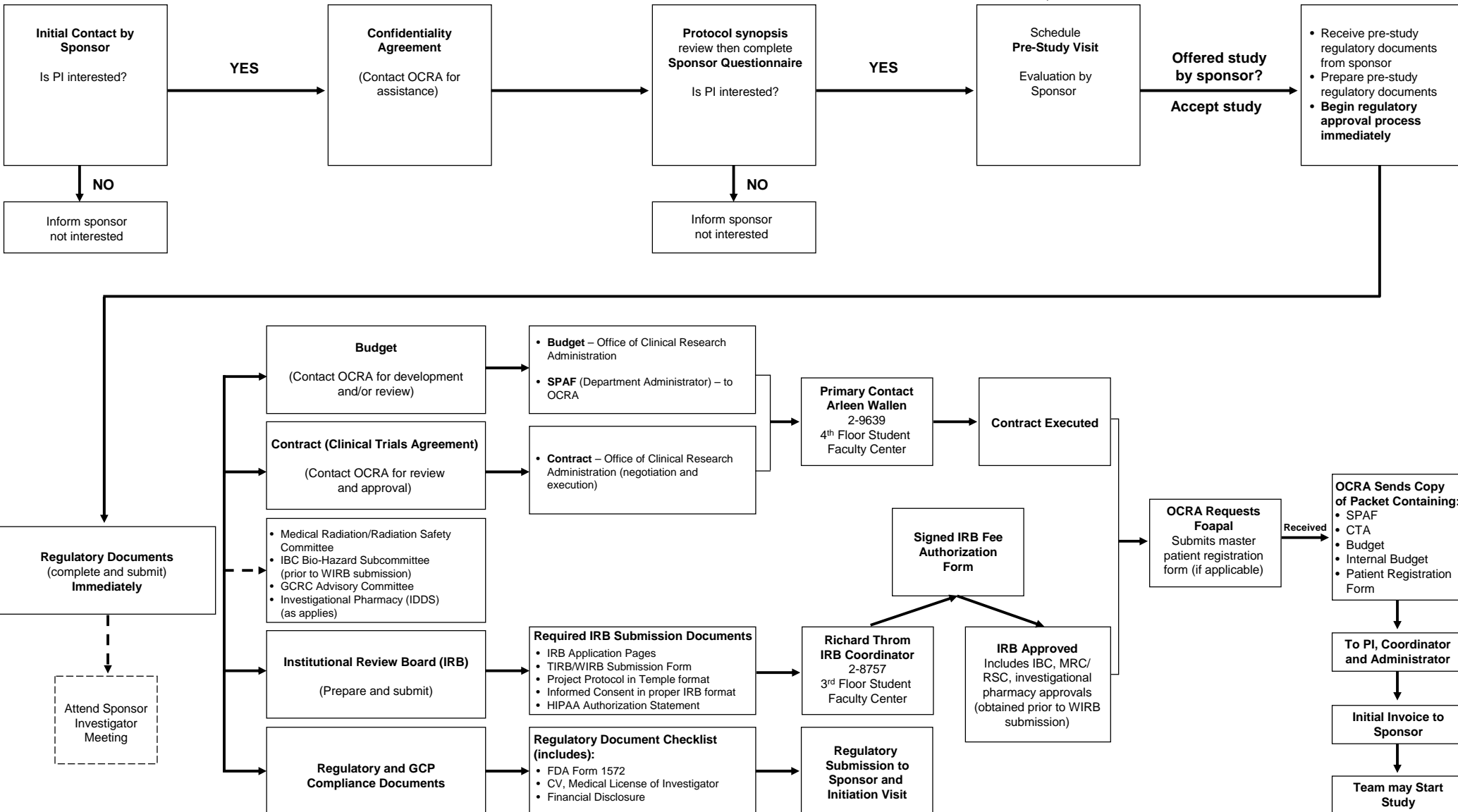
A Temple Guide for Study Start and Approval

Office of Clinical Research Administration (OCRA)

Reviews, negotiates and processes all industry clinical trial agreements, provides guidance and GCP training regarding study start up process and conduct of clinical trial.

Contact Us:

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IRB submission process can occur concurrently with budget and contract review provided OCRA receives a signed IRB fee authorization form