Investigator-Initiated Study (IIS) Concept Sheet

Complete the form below and attach specified documentation.
Return to IIS Program Coordinator at iis-grants@lexpharma.com

Date Request Submitted
Study Type Preclinical, Clinical, or Preclinical & Clinical
Request Type Drug Only, Funding Only, or Drug & Funding
Primary Investigator (PI) Include CV with the submission
Name and title:
Institution:
Mailing address:
Phone:
Fax:
Email:
Co-PI(s)
Study Site(s) Institutions and sites where the study will be conducted
Study Title
Background & Rationale Describe the background and rationale for the study, including a brief review of the literature. Attach references with your submission as needed

Study Objective(s) & Scientific Hypothesis Provide basic description of the study objectives and/or hypothesis
Study Design Describe the study design, including information regarding endpoints, inclusion/exclusion criteria, control groups, and sample size/power calculations as appropriate
Study Timeline Estimated study duration. For clinical studies include estimated duration from initiation (FPI) to completion (LPO), including follow-up and completion of study report
Budget Include summary of total budget request and estimated budget for each year of the study. Enclose detailed itemized budget with the submission
Publication Plan Include anticipated submission of abstracts and full manuscripts, with lead author, preliminary list of authors, timing of submission, and target congress and/or journal