IIS Submission Guidance

Daiichi Sankyo accepts unsolicited IIS proposals for clinical or non-clinical studies. The following minimum information must be submitted to Daiichi Sankyo at IIS@dsi.com in order for us to conduct a review:

- Principal Investigator
- Institution Name
- Type of Study (clinical, non-clinical)
- Type of Request (Drug Product, Financial, or both)
- Budget and Drug Product Request details
- Study Rationale
- Study Design (e.g. control, number of treatment arms, product doses, treatment duration, numbers of animals)
- Primary Objective(s)
- Planned timelines

The following are additional information requirements for clinical IIS proposals:

- Inclusion/Exclusion Criteria
- Study Population/Indication
- Study Endpoints
- Planned Sample Size and Statistical Powering Justification
- Planned number of subjects and sites/site locations (country only)
- Study Visit Schedule of Events

If you would like to request a template to assist you, or prefer to submit a full application, please contact us at IIS@dsi.com.