

## REQUEST SUITABILITY OF THE FACILITIES

**In order to start a study** at Germans Trias i Pujol University Hospital, **it is mandatory for the study to be registered** at the site by the HUGTiP REC.

- **Clinical trials with medicinal products or medical devices** will be registered by **requesting the suitability of the facilities document** to the secretariat of the HUGTiP Research Ethics Committee.

To **request the signature** of the suitability of the facilities, you must send to **avaluacionsceic.germanstrias@gencat.cat** the following requirements:

- **Submit the full** protocol (in Spanish or English)
  - State **which is the assessing REC**.
  - Report on **the status of the assessment application** (not submitted, under evaluation, approved)
  - Indicate the name of principal investigator at HUGTiP
  - **Submit [Suitability of the site's facilities](#)** along with study data and the site's involved departments.
- **Other kind of studies** will be registered by **submitting the initial assessment or feasibility review application** to the HUGTiP REC.

To **request the initial assessment or feasibility review** from a study you must send the following documents to: **avaluacionsceic.germanstrias@gencat.cat** the stipulated requirements depending on the type of study, available at the following link: **[ASSESSMENT REQUIREMENTS/FEASIBILITY REVIEW](#)**