STUDY NUMBER: 2600180

Dear pioneer,

A study to **treat a hormonal disorder** will soon start at our research centre in Edegem.
Are you interested in participating and giving science a boost?
Hereby we would like to give you an overview of the study: duration of **± 7 weeks.**

In this cohort, the study medication will be administered **orally** (by mouth). In addition, a hormone stimulation test will be administered **intravenously** (via the vein).

INCLUSION CRITERIA

- **Last participation in other clinical study (final study visit*):**
  - Cohort **A1.1**: no later than **02SEP2026** (3 months prior to dosing)

  **Be careful: Final study visit of the previous study needs to be performed to be able to participate at screening.*
- Healthy men women between age of 18 and 55 years (inclusive)
- BMI: between 18.5 and 30.0 kg/m² (inclusive)
  - Minimum body weight of 50.0 kg for males and 45.0 kg for females
- Non-smoker or willing to stop smoking from screening until the final study visit, including nicotine-containing products such as nicotine patches or electronic cigarettes
- Not using any medication, vitamins, or homeopathic substances
- Contraception:
  - Infertile women: **Hormone replacement therapy is not permitted**
    - postmenopausal (= no menstruation for at least 12 months)
    - OR removal of uterus
    - OR removal of both ovaries
    - OR removal of both fallopian tubes
    - other medically documented causes of infertility
  - Men: (From administration of study medication until 90 days after the administration)
    - No sperm donation (from screening until 90 days after the administration of study medication)
    - Use of a condom
    - Azoospermia (due to vasectomy or another medical cause)
- You will be asked to follow the restrictions regarding physical activity, consumption of poppy seeds, grapefruit (juice), citrus fruits, quinine-containing beverages (e.g. tonic water), caffeine- and alcoholic products.
- No participation in any of the previous cohorts of this study
- You are not eligible to participate if you:
  - are hypersensitive to sunlight or UV radiation
  - live in a different time zone than Belgium
  - work night shifts or have a disrupted sleep pattern
  - have donated and/or lost approximately 500 mL of blood within the 3 months prior to the screening visit. For plasma donations, a washout period of 30 days before administration of the study medication applies

INTEREST IN PARTICIPATING

Registration for this study only indicates your interest and does not obligate you to participate, nor us to include you as a participant.

Thank you in advance for your interest and we hope to welcome you soon for study 2600180

SGS CPU