

A study to evaluate the effect of various degrees of reduced liver function on the processing of fenebrutinib in the body

Submission date 05/10/2023	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 10/10/2023	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 13/05/2024	Condition category Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

A disease of the brain and spinal cord (central nervous system) called multiple sclerosis (MS), is a long-lasting (chronic) disease in which the body attacks the protective covering around nerves and damages the nerves. Fenebrutinib is an experimental drug being developed for the treatment of MS. Health authorities have not yet approved fenebrutinib for the treatment of MS.

The main aim of the study is to find out how mild or moderate liver damage (mild or moderate hepatic impairment) will affect the breakdown and removal of the study drug (fenebrutinib) from the body (this is called pharmacokinetics [PK]).

Who can participate?

People between 18 and 75 years of age with normal liver function or with mild or moderate liver damage (mild or moderate hepatic impairment) can participate in this study.

What does the study involve?

Participants will need to be a part of the study for about 5 weeks including the screening period. The study will include the following parts:

1. A screening period of up to 28 days to check the eligibility of participants to take part in the study.
2. A dosing/treatment period: Participants with normal liver function and mild or moderate liver damage will receive a single dose of fenebrutinib, by mouth (orally) on Day 1. Participants will have to get admitted to the clinic 1 day before receiving the treatment (Day -1) and stay in the clinic until Day 5.
3. A follow-up phone call to check on the participant will be made 7 days after the study drug administration to check on their well-being.

What are the possible benefits and risks of participating?

Fenebrutinib is an experimental drug and is being given purely for research purposes, it is not

intended that participants will receive any benefit from this study, but the information learned from this study may be useful to treat future patients of multiple sclerosis. Participants may receive monetary compensation for taking part in the study.

Participants may have side effects due to fenebrutinib or procedures used in this study. Side effects can be mild to severe and even life-threatening or fatal, and they can vary from person to person. Full information on risks associated with fenebrutinib is provided to volunteers in the Informed Consent Form. When investigating new medicines there is also a risk of unexpected side effects and occasionally allergic reactions. All volunteers will be closely monitored during the study and safety assessments will be performed at regular intervals. Risks are further mitigated by ensuring that only volunteers who meet all inclusion/exclusion criteria are included and that if the safety of any volunteer represents a concern they will be withdrawn. Blood samples will be collected during the study. Collection of these samples can cause pain, bruising, or infection where the needle is inserted. Electrocardiograms (ECGs) will be taken in this study. ECG patches may cause a skin reaction such as redness or itching, or localized skin discomforts and/or hair loss associated with the placement of ECG leads.

There may be a risk in exposing an unborn child to the study drug, and all risks are not known at this time. Women and men must take precautions to avoid exposing an unborn child to the study drug. Participants who are pregnant, become pregnant or are currently breastfeeding cannot take part in this study.

Where is the study run from?
Genentech Inc. (Switzerland)

When is the study starting and how long is it expected to run for?
July 2023 to January 2025

Who is funding the study?
Genentech Inc. (Switzerland)

Who is the main contact?
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Contact information

Type(s)
Public, Scientific, Principal Investigator

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number**ClinicalTrials.gov number**

Nil known

Secondary identifying numbers

GP44943

Study information

Scientific Title

A phase I, open-Label, single-Dose study to evaluate the effect of mild or moderate hepatic impairment on the pharmacokinetics of fenebrutinib

Study objectives

The main aim of this study is to determine the pharmacokinetics (PK) of a single oral dose of fenebrutinib in participants with mild or moderate hepatic impairment compared to demographically matched healthy participants with normal hepatic function.

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 16/08/2023, Salus IRB (2111 West Braker Lane, Suite 100, Austin, Texas, 78758, United States of America; +1 512-380-1244; salus@salusirb.com), ref: None provided

Study design

Phase I multicenter non-randomized open-label parallel-group single-dose study

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Other

Study type(s)

Other

Participant information sheet

No participant information sheet available

Health condition(s) or problem(s) studied

Multiple sclerosis (MS) (study conducted in volunteers with normal hepatic function or mild/moderate hepatic impairment)

Interventions

Cohort 1: Participants with normal hepatic function will receive a single dose of fenebrutinib, 200 milligrams (mg), orally, on Day 1.

Cohort 2: Participants with mild hepatic impairment will receive a single dose of fenebrutinib, 200 mg, orally, on Day 1.

Cohort 3: Participants with moderate hepatic impairment will receive a single dose of fenebrutinib, 200 mg, orally, on Day 1.

Intervention Type

Drug

Pharmaceutical study type(s)

Pharmacokinetic

Phase

Phase I

Drug/device/biological/vaccine name(s)

Fenebrutinib

Primary outcome measure

1. Area under the concentration-time curve from hour 0 to the last measurable concentration (AUC_{0-t}) of fenebrutinib measured using a validated liquid chromatography with tandem mass spectrometry (LC-MS/MS) assay from the blood samples collected at multiple time points from Day 1 to Day 5
2. AUC extrapolated to infinity (AUC_{0-∞}) of fenebrutinib measured using a validated LC-MS/MS assay from the blood samples collected at multiple time points from Day 1 to Day 5
3. Maximum observed concentration (C_{max}) of fenebrutinib measured using a validated LC-MS/MS assay from the blood samples collected at multiple time points from Day 1 to Day 5

Secondary outcome measures

1. Number of participants with adverse events (AEs) and severity of AEs graded as per National Cancer Institute Common Terminology Criteria for Adverse Events version 5.0 (NCI CTCAE; v5.0) from screening up to 14 days after the dose of study drug (approximately 6 weeks)

Overall study start date

20/07/2023

Completion date

14/01/2025

Eligibility

Key inclusion criteria

1. Male and female participants between 18 to 75 years of age, inclusive, at screening
2. Body weight ≥ 45 kilograms (kg) and within body mass index (BMI) range 18.0 to 42.0 kilograms per square meter (kg/m²), inclusive

Additional inclusion criteria for participants with normal hepatic function (Cohort 1) only:

1. In reasonably good health for their age
2. Matched to participants with mild or moderate hepatic impairment in sex, age (± 10 years), and body weight ($\pm 15\%$)
3. Negative hepatitis panel
4. Normal hepatic function and no history of clinically significant hepatic dysfunction
5. Estimated glomerular filtration rate ≥ 80 millilitre per minute (mL/min) at screening, as calculated by the Chronic Kidney Disease Epidemiology Collaboration equation.

Additional inclusion criteria for participants with hepatic impairment (Cohorts 2 and 3) only:

1. Chronic (>6 months), stable (no acute episodes of illness within the previous 1 month prior to Screening due to deterioration in hepatic function) hepatic insufficiency with features of cirrhosis due to any etiology. Participants must also remain stable throughout the Screening period.
2. Considered to have mild or moderate hepatic impairment and has been clinically stable for at least 1 month prior to Screening. To be classified as having hepatic impairment, participants must have a Child-Pugh score of 5 to 6 (inclusive; mild) or 7 to 9 (inclusive; moderate) with known medical history of liver disease and unambiguous medical history.
3. Negative hepatitis B virus core antibody and HbsAg screens and negative hepatitis C viral load.
4. Estimated glomerular filtration rate ≥ 60 mL/min at screening, as calculated by the Chronic Kidney Disease Epidemiology Collaboration equation.

Other protocol defined inclusion criteria could apply.

Participant type(s)

Healthy volunteer, Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

75 Years

Sex

Both

Target number of participants

32

Key exclusion criteria

1. Participants who are pregnant or breastfeeding or intending to become pregnant during the study or within 28 days after the dose of study drug.
2. History of surgical or artificial shunts (i.e., transjugular intrahepatic portosystemic procedure).
3. History of malignancy within 5 years prior to screening, except for completely excised basal

cell carcinoma or squamous cell carcinoma of the skin or cervical carcinoma in situ.

4. Significant illness, including infections, surgery, or hospitalization within the 2 weeks prior to dosing.

5. History of stomach or intestinal surgery or resection that would potentially alter absorption and/or excretion of orally administered drugs, except that uncomplicated hernia repair, appendectomy, and/or cholecystectomy will be allowed.

6. Malabsorption syndrome or other condition that would interfere with enteral absorption.

Additional exclusion criteria for participants with hepatic impairment (Cohorts 2 and 3) only:

1. Evidence of progressive liver disease that has worsened or is worsening within 1 month prior to Screening.

2. Evidence of hepatorenal syndrome

3. Ascites requiring paracentesis within 3 months prior to Check-in (Day -1) or other intervention, with the exception of diuretics.

4. Required treatment for gastrointestinal bleeding within 12 months prior to Check-in (Day -1)

5. Receipt of a liver transplant

6. Required additional medication for hepatic encephalopathy within 12 months prior to check-in (Day -1), unless deemed acceptable by the investigator.

7. One or more of the following laboratory results:

- Alkaline phosphatase $>10 \times$ upper limit of normal (ULN).

- Alanine aminotransferase or aspartate aminotransferase $>5 \times$ ULN.

- Bilirubin $>3 \times$ ULN in the absence of Gilbert's Syndrome or hemolysis.

Other protocol defined exclusion criteria could apply.

Date of first enrolment

18/10/2023

Date of final enrolment

07/01/2025

Locations

Countries of recruitment

Canada

New Zealand

Study participating centre

Altasciences Montreal

1200 Beaumont Avenue

Montréal

Canada

H3P 3P1

Study participating centre

NZCR - Auckland
Ferncroft St
Grafton, Auckland
New Zealand
1010

Sponsor information

Organisation

Genentech Inc.

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Sponsor type

Industry

Website

<https://www.roche.com/about/>

Funder(s)

Funder type

Industry

Funder Name

Genentech Inc.

Results and Publications

Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal

Intention to publish date

14/01/2026

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are not expected to be made available due to participant-level data not being a regulatory requirement.

IPD sharing plan summary

Not expected to be made available