

Comparing the effects of infliximab and alpha interferon in the treatment of Behçet's Disease

Submission date 09/03/2016	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 09/03/2016	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 24/10/2024	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Behçet's disease (BD) is a rare and poorly understood condition that causes inflammation (swelling) throughout the body. The inflammation associated with BD can cause a wide range of symptoms, including genital and mouth ulcers, eye problems (which can lead to blindness), skin rashes and inflammation of the blood vessels, which increases a person's risk of developing blood clots and stroke. The exact cause of BD is unknown however many believe it is an autoimmune disorder (in which the body's immune system mistakenly attacks the body's own, healthy tissue). Traditional treatments include corticosteroids (anti-inflammatory medicine) and immunosuppressant drugs (medications that reduce the activity of the immune system), but they can have unpleasant side effects. Biologic therapies are a newer type of medication, which target the biological processes involved in the process of inflammation more selectively. Recently, the biologic drugs infliximab and interferon alpha (aIFN) have been used successfully in treating BD however they are expensive (£16,000/year and £4,000/year, respectively), not always effective and there have been very few studies looking at their effectiveness. The Department of Health has recently commissioned three national Centres of Excellence for BD in London, Birmingham and Liverpool, with the remit to deliver care for the approximate 500 UK patients currently with this condition. The aim of this study is to compare the effectiveness of infliximab with aIFN at treating patients with BD.

Who can participate?

Adults with BD who are suitable to receive biological therapy.

What does the study involve?

Participants are randomly allocated to one of two groups. Those in the first group are treated with infliximab. This is given through a drip (intravenous infusion) at the standard dose of 5mg/kg at the start of the study, week 2, week 6 and then every 8 weeks until the end of the study. Those in the second group are treated with aIFN. This is given as an injection under the skin (subcutaneous injection) starting at a dose of 3 million units one a day for 3 days, then 6 million units (for men who weigh more than 80kg) or 4.5 million units (for women or men weighing less than 80kg) every day. This dose can be reduced down every 2-4 weeks if necessary to 3 million

units twice a week over the course of the study. After 3 and 6 months of treatment, participants in both groups are assessed in order to find out if there has been any improvement to their condition.

What are the possible benefits and risks of participating?
Not provided at time of registration.

Where is the study run from?
University of Liverpool (UK)

When is the study starting and how long is it expected to run for?
April 2016 to June 2021 (updated 10/11/2020, previously: November 2020) (updated 09/01/2020, previously: December 2019)

Who is funding the study?
Medical Research Council (UK)

Who is the main contact?
Miss Elizabeth Blennerhassett

Contact information

Type(s)
Scientific

Contact name
Dr Diane Carlton

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Additional identifiers

EudraCT/CTIS number
2014-005390-36

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
20233

Study information

Scientific Title

BIO BEHÇET'S: Optimal utilisation of biologic drugs in Behçet's Disease: a randomised controlled trial of infliximab vs alpha interferon, with genotyping and metabolomic profiling, towards a stratified medicines approach to treatment

Acronym

BIO BEHÇET'S

Study objectives

The aim of the study is to create the evidence base to underpin clinically effective prescribing of the biologic drugs infliximab and alpha interferon for Behçet's Disease.

The objectives of the study are to:

1. Undertake a randomised controlled trial to compare IFX versus aIFN in patients with BD who are unresponsive to standard oral therapy
2. Examine whether IFNL3 and IFNL4 SNPs can predict response to aIFN and/or IFX in BD
3. Examine the potential for urine metabolomics to act as biomarker for drug response to IFX and /or aIFN in BD

Ethics approval required

Old ethics approval format

Ethics approval(s)

North West - Liverpool Central Research Ethics Committee, 17/02/2015, 15/NW/0008

Study design

Randomised two-arm parallel open-label design

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Behçet disease

Interventions

Participants are randomly allocated to one of two groups.

Group 1: Participants will receive infliximab at a standard dose of 5mg/kg at week 0, week 2 and week 6 as loading then every 8 weeks for the length of the trial.

Group 2: Interferon alpha (Roferon) subcutaneous injection in tapering dose starting at 3 million units once daily for three days, then 6** million units once daily (male with body weight more than 80 kg) or 4.5 million units once daily (female, or male with weight less than 80 kg). Dose tapered down every 2-4 weeks according to defined clinical criteria to 3 million units twice a week, over the period of the trial.

** The six million units dose will only be administered to males of over 80 kg with major organ threatening disease (e.g., severe eye involvement). Males of less than 80kg or females will start with 4.5 million.

Intervention Type

Other

Primary outcome measure

Disease severity is measured using the Behcet's disease activity index (BDAI) after 3 months of treatment.

Secondary outcome measures

1. Disease severity is measured using the Behcet's disease activity index (BDAI) after six months of treatment,
2. Significant improvement in organ involvement within the primary organ system that resulted in the decision to start a biologic agent and other organ systems assessed by:
 - 2.1. Ocular: reduction in vitreous haze using the SUN consensus group grading scale and visual acuity change from baseline
 - 2.2. Oral ulcer activity: change in ulcer severity score (USS)
 - 2.3. Change in number of genital ulcers
 - 2.4. Musculoskeletal: Likert pain score
3. Adverse events in each group
4. Reduction in dose of prednisolone (or equivalent glucocorticoid) at three months: a clinically meaningful reduction is considered to be 50% of baseline or dose of <15mg/day prednisolone
5. Reduction in dose of prednisolone (or equivalent glucocorticoid) at six months: a clinically meaningful reduction is considered to be 50% of baseline or dose of <7.5mg/day prednisolone
6. Quality of life is measured using EQ-5D and BD-QoL at baseline, 3 and 6 months
7. Disease activity is measured using the Physician's Global Assessment of disease activity (a 7 point Likert Scale completed as part of (but assessed independently of) the BDAI) at 3 and 6 months

Overall study start date

01/04/2016

Completion date

30/06/2021

Eligibility

Key inclusion criteria

1. Diagnosed to have BD by International Study Group (ISG) criteria or International Criteria for BD (ICBD)
2. Have refractory disease as defined by the UK Centres of Excellence criteria (failure to respond to steroid and/or immunosuppressive therapy with significant or major organ-threatening disease) and therefore qualify for biologic therapy with either IFX or aIFN
3. Able to give informed consent
4. Have not previously received a biologic agent
5. Aged over 18 years

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 100; UK Sample Size: 100

Total final enrolment

79

Key exclusion criteria

1. Have a contraindication to either IFX or aIFN (eg active infection, severe liver disease, neutropenia, previous malignancy)
2. Are likely to not comply: for example cannot attend assessments because of excessive travel requirements
3. Are already, or likely to become, pregnant during the study
4. Express a strong preference for one of the two potential therapies
5. Have heart disease or severe heart failure
6. Have been diagnosed with Multiple sclerosis
7. Have evidence of infection with HIV

Date of first enrolment

01/04/2016

Date of final enrolment

28/02/2020

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre**University of Liverpool**

Department of Health Services Research
1-3 Brownlow Street
Liverpool
United Kingdom
L69 3GL

Sponsor information**Organisation**

University of Liverpool

Sponsor details

Whelan Building
Quadrangle
Brownlow Hill
Liverpool
England
United Kingdom
L69 3GB

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/04xs57h96>

Funder(s)**Funder type**

Government

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

30/06/2021

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
HRA research summary			28/06/2023	No	No
Results article		21/10/2024	24/10/2024	Yes	No