

Pilot trial of omeprazole in idiopathic pulmonary fibrosis (PPIPF Study)

Submission date 10/02/2014	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 11/02/2014	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 21/06/2019	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Idiopathic pulmonary fibrosis (IPF) is a disease of unknown cause in which areas of normal lung tissue are replaced by scars. As a result it becomes harder for the lungs to inhale oxygen from the air. IPF is commonly progressive, and around 50% of patients diagnosed with the disease die after about 3 years. The most common troublesome symptoms of IPF are breathlessness on exertion and cough. No drug treatments have been clearly shown to improve the death rate or to significantly improve symptoms in IPF.

In recent years it has been recognised that cough can be caused by small amounts of liquid coming up from the stomach and going down the wrong way into the lungs, a process commonly known as reflux. As liquid in the stomach is usually acidic, patients lungs may repeatedly be exposed to small amounts of acid. Reflux is unusually common in IPF and could potentially contribute to the unbearable cough found with the disease. However, there are many potential causes for cough in IPF.

Stomach acid can be efficiently switched off by drugs called proton pump inhibitors, one of which is called omeprazole. If reflux of stomach acid does contribute to cough in IPF, omeprazole might be expected to reduce cough. The purpose of this study is therefore to test whether omeprazole does reduce cough in patients with IPF.

Who can participate?

Patients with idiopathic pulmonary fibrosis aged between 40 and 85 years.

What does study involve?

Participants will be randomly allocated to one of two groups. One group will take 20 mg of Omeprazole twice daily and the other group will take a matching placebo (dummy). Both groups will take their medications orally before food for 90 days.

Before starting the study and towards the end of the study participants will undergo tests which will be distributed over six visits.

What are the possible benefits and potential risks for participants?

There is no direct benefit to participants. If the results are encouraging, this would pave the way for larger studies to comprehensively assess the potential role of omeprazole as a new effective treatment for IPF.

The risks from taking part are considered to be low. Cough monitoring is a research tool, but is not associated with any known risks. All of the other tests are routinely carried out in clinical practice, and all are considered to be low-risk procedures. Omeprazole has been used in clinical practice by millions of patients and is regarded as having an extremely low rate of serious side effects.

Where is the study run from?

Interstitial Lung Disease Clinic in Royal Victoria Infirmary, Newcastle Upon Tyne, UK.

When is the study starting and how long is it expected to run for?

The recruitment is expected to start in February - March 2014. Participants will be enrolled for period of 3 months.

Who is funding the study?

British Lung Foundation, UK.

Who is contact for study?

Professor John Simpson

Contact information

Type(s)

Scientific

Contact name

Dr Prosenjit Dutta

Contact details

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Additional identifiers

EudraCT/CTIS number

2013-003301-26

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

15732

Study information

Scientific Title

A randomised double blinded placebo-controlled pilot trial of omeprazole in idiopathic pulmonary fibrosis (IPF)

Acronym

PPIPF

Study objectives

Omeprazole reduces cough in IPF.

More details can be found at: <http://public.ukcrn.org.uk/search/StudyDetail.aspx?StudyID=15732>

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES committee of Yorkshire & The Humber - Leeds West, 10/01/2014, REC ref: 13/YH/0284

Study design

Randomised; Interventional; Design type: Treatment

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

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

Idiopathic Pulmonary Fibrosis - Respiratory (Chest) Disease

Interventions

Omeprazole or matched placebo

Experimental Arm: Omeprazole 20 mg twice daily

Comparator Arm: Matched placebo twice daily

Study Entry: Single Randomisation only

Patients will be randomised 1:1 to Omeprazole 20 mg twice daily of matching placebo to be taken orally before food for 90 days. Prior to starting and towards the end of the study participants will undergo the following tests which will be distributed over six visits.

Visit 1: breathing tests, walk test and 24-hour cough monitor test

Visit 2: reflux study

Visit 3: start trial medication for 3 months

Visit 4, 5, and 6: ideally during the last 3 days (but can be within 2 weeks) of completion of study medication

Visit 4: duplicate of Visit 1

Visit 5: duplicate of Visit 2

Visit 6: telescopic test for lungs (bronchoscopy)

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Omeprazole

Primary outcome measure

1. Change in objectively measured cough frequency from beginning of study (baseline) to end of treatment (within last 2 weeks of completion of treatment). This will be compared in two groups.
2. Range of other measurements will be assessed such as the numbers of patients eligible to take part, agreeing to randomization and providing outcome data.

Secondary outcome measures

1. Changes in symptoms of cough and reflux at the end of treatment (as measured by validated questionnaires)
2. Changes in amount of acid and nonacid reflux at the end of treatment
3. Changes in lung function tests and distance walked in 6 minutes after treatment
4. Amount of inflammation and rate of infection in lungs
5. Any adverse events related to medication
6. Rates of eligibility, recruitment, randomization and study completion: feasibility and acceptability; Timepoint(s): end of study period.

Overall study start date

01/02/2014

Completion date

30/09/2016

Eligibility

Key inclusion criteria

A pragmatic clinical definition of IPF will be used, in which recruited patients must fulfill all of the following criteria:

1. IPF is considered the most likely diagnosis by the regional interstitial lung disease multidisciplinary team meeting (ILD-MDT)
2. History of cough, with or without exertional dyspnoea
3. High resolution computed tomography (HRCT) scan features of honeycombing in a predominantly basal and subpleural distribution

4. Bibasal crackles on auscultation
5. Features of a restrictive ventilatory defect (vital capacity (VC) <90% predicted and/or diffusion factor for carbon monoxide (Tco) <90% predicted)
6. Aged 40-85 years

Patients with radiological emphysema will be eligible so long as the diagnosis of IPF is secure, i.e., all the features above are satisfied.

If the regional ILD-MDT cannot reach a clear consensus as to the diagnosis, the case will be referred to two experts in ILD from outside the region, and the patient will be eligible if both consider IPF to be the most likely diagnosis.

Patients taking a PPI during screening will potentially be eligible. In these cases the indication for ongoing treatment will be reviewed.

1. Patients taking short courses (e.g., 2 months) of PPI will be eligible once the treatment has been discontinued for a minimum of 1 month.
2. There are few licensed indications for long-term omeprazole other than Zollinger-Ellison syndrome. Therefore, unless there is a known diagnosis of Zollinger-Ellison or a history of significant dyspepsia or gastrointestinal bleeding during a previous discontinuation of PPI, patients on long-term PPI will be asked to consider a trial of supervised discontinuation. If he/she agrees and provides written consent to a trial of discontinuation, the GP will be contacted. If both the patient and the GP agree to a trial of discontinuation, the patient will be eligible for the study if he/she provides written consent after at least 2 weeks without PPI.
3. Patients taking antacids, prokinetics or raft alginates at the time of screening will be eligible if they have been off these treatments for a period of at least 2 weeks.
4. Target Gender: Male & Female; Upper Age Limit 85 years ; Lower Age Limit 40 years

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

Total 60 patients - 30 in each arm

Total final enrolment

45

Key exclusion criteria

1. Known allergy to omeprazole or other PPI
2. Concomitant use of warfarin, diazepam, phenytoin or ketoconazole
3. Concomitant use of a regular PPI, antacid, prokinetic or raft alginate during the trial period.
4. History of upper respiratory tract infection, lower respiratory tract infection or exacerbation of IPF in the 4 weeks before starting study drugs
5. Active trial of treatment for IPF (e.g., prednisolone, pirfenidone, N-acetylcysteine) started in

the 4 weeks before starting study drugs

6. Documented history of hepatic cirrhosis

7. Pregnancy or lactation

8. ILD-MDT considers the most likely cause of the patients ILD to be a condition other than IPF, for example rheumatoid lung, systemic sclerosis ILD, asbestosis, chronic hypersensitivity pneumonitis, sarcoidosis, etc.

9. Concurrent enrolment in a trial of a Clinical Trials of Investigational Medicinal Product (CTIMP) for IPF

Date of first enrolment

01/02/2014

Date of final enrolment

30/09/2016

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

4th Floor William Leech Building

Newcastle Upon Tyne

United Kingdom

NE2 4HH

Sponsor information

Organisation

The Newcastle upon Tyne Hospitals NHS Foundation Trust (UK)

Sponsor details

c/o Jillian Peacock

Joint Research Office

Level 6 Leazes Wing

Royal Victoria Infirmary

Queen Victoria Road

Newcastle Upon Tyne

England

United Kingdom

NE1 4LP

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/05p40t847>

Funder(s)

Funder type

Charity

Funder Name

British Lung Foundation (UK) Grant Codes: IPFPSG12-7

Alternative Name(s)

BLF

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

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?
Basic results			21/06/2019	No	No
HRA research summary			28/06/2023	No	No