

Trial of vitamin D supplementation in chronic obstructive pulmonary disease (COPD)

Submission date 12/05/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/05/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 03/02/2016	Condition category Respiratory	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Vitamin D - the sunshine vitamin - is best known for its effects on bone health. Profound deficiency causes rickets, a condition that causes the bones on children to become soft and weak, which, in turn, can lead to bone deformities. More moderate deficiency, commonly seen in the UK during winter and spring, can make people more susceptible to respiratory infections. Respiratory infections cause 20% of GP consultations, 300,000 hospital admissions and 30,000 deaths per year. Patients with chronic obstructive pulmonary disease (COPD) are at high risk of such infections. Studies have shown that vitamin D 'switches on' the production of natural antibiotic substances that can kill viruses and bacteria in cells that fight infection. One small study, originally designed to look at the effects of vitamin D on bone health has shown that patients receiving high-dose vitamin D were 3 times less likely to have cold and 'flu symptoms than those who received placebo (dummy pill). The primary aim of the study is to determine whether vitamin D supplementation is a cost-effective and acceptable way to reduce acute respiratory illness in patients with COPD.

Who can participate?

Patients aged 40 years and over and diagnosed with COPD

What does the study involve?

Participants are randomly allocated to one of two groups. Those in group 1 are given Vigantol (a form of vitamin D). Those in group 2 are given a placebo. All participants attend five study visits over the course of a year and are also contacted by telephone on five occasions at intervals between scheduled visits. Participants are asked to complete a daily diary of chest symptoms, give blood samples and perform breathing and muscle strength tests at the beginning, the middle and the end of the study.

What are the possible benefits and risks of participating?

Not provided at time of registration

Where is the study run from?

A number of GP surgeries in London and Norfolk.

When is the study starting and how long is it expected to run for?
September 2009 to August 2013

Who is funding the study?
National Institute for Health Research (UK)

Who is the main contact?
Ms Wai Yee James

Contact information

Type(s)
Scientific

Contact name
Ms Wai Yee James

Contact details
Centre for Health Sciences
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E1 2AT

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number
NCT00977873

Secondary identifying numbers
7831

Study information

Scientific Title
Randomised, multicentre, double-blind, placebo-controlled trial of vitamin D supplementation in patients with chronic obstructive pulmonary disease (COPD)

Study objectives
The primary aim of the study is to determine whether vitamin D supplementation is a cost-effective and acceptable strategy to reduce acute respiratory illness in patients with COPD.

Ethics approval required
Old ethics approval format

Ethics approval(s)

Study design

Multicentre randomised interventional prevention trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Prevention

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

Topic: Inflammatory and Immune System; Subtopic: Inflammatory and Immune System (all Subtopics); Disease: Immunology and inflammation

Interventions

Vigantol, 3 mg or miglyol oil (placebo), to be administered every two months over a twelve months period, in total of six doses. Patients will be followed up at these post-dose time points: 2 months, 6 months and 12 months.

Intervention Type

Supplement

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Vitamin D

Primary outcome measure

Time from randomisation to first moderate or severe COPD exacerbation

Secondary outcome measures

Respiratory morbidity. Measured at screen visit, randomisation visit, 2 months post-dose, 6 months post-dose and 12 months post-dose.

Overall study start date

11/09/2009

Completion date

Eligibility

Key inclusion criteria

1. Medical record diagnosis of COPD, emphysema or bronchitis
2. Post-bronchodilator forced expiratory volume in one second (FEV1)/forced vital capacity (FVC) less than 70% or post-bronchodilator FEV1/slow vital capacity (VC) less than 70%
3. Post-bronchodilator FEV1 less than 80% predicted
4. Age 40 years on day of first dose of IMP, either sex
5. Smoking history 15 pack-years
6. Exacerbation of COPD requiring treatment with antibiotics and/or systemic corticosteroids within 12 months of screening visit
7. Contactable by telephone and able to attend face-to-face review at 2, 6 and 12 months post-enrolment
8. If a woman of child-bearing potential, is sexually abstinent or has negative pregnancy test within 7 days of recruitment and agrees to use reliable form of contraception until she has completed the study
9. Able to give written informed consent to participate

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Planned sample size: 240

Key exclusion criteria

1. Current diagnosis of asthma
2. Known clinically significant bronchiectasis
3. Known sarcoidosis, hyperparathyroidism, nephrolithiasis, active tuberculosis, vitamin D intolerance, liver failure, renal failure, terminal illness, lymphoma or other malignancy not in remission for 3 years
4. Any other condition that, in an investigator's judgement, might compromise patient safety or compliance, interfere with evaluation or preclude completion of the study
5. COPD requiring long-term oxygen therapy 12 hours per day
6. Taking benzothiadiazine derivative, cardiac glycoside, carbamazepine, phenobarbital, phenytoin or primidone
7. Taking dietary supplement containing vitamin D up to 2 months before first dose of IMP
8. Treatment with any investigational medical product or device up to 4 months before first dose of IMP
9. Breastfeeding, pregnant or planning a pregnancy
10. Baseline corrected serum calcium greater than 2.65 mmol/L
11. Baseline serum creatinine greater than 125 micromol/L
12. Upper respiratory tract infection (URTI) or COPD exacerbation up to 28 days before first dose of IMP

- 13. Inability to use spirometer
- 14. Inability to complete symptom diary

Date of first enrolment

11/09/2009

Date of final enrolment

30/08/2013

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Centre for Health Sciences

London

United Kingdom

E1 2AT

Sponsor information

Organisation

Barts and The London School of Medicine and Dentistry

Sponsor details

Blizard Institute of Cell and Molecular Science (ICMS)

The Blizard Building

4 Newark Street

London

England

United Kingdom

E1 2AT

Sponsor type

University/education

Website

<http://www.icms.qmul.ac.uk/>

ROR

<https://ror.org/00b31g692>

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

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?
Results article	results	01/12/2014		Yes	No
HRA research summary			28/06/2023	No	No