ISRCTN18247441 https://doi.org/10.1186/ISRCTN18247441

An evaluation of systemic metronidazole and amoxycillin as an adjunct to routine therapy in the treatment of severe periodontal disease

Submission date 23/01/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 23/01/2004	**Overall study status** Completed	
Last Edited 06/01/2010	**Condition category** Oral Health	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mrs Jane Rooney

Contact details
University of Bristol
Dental Hospital and School
Lower Maudlin Street
Bristol
United Kingdom
BS1 2LY
+44 (0)117 9294472
J.C.Rooney@bristol.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

R/ROONEY/291

Study information

Scientific Title

Study objectives
The aim of this study is to evaluate systematic metronidazole and amoxycillin, alone and combined, as adjuncts to root planning in the treatment of patients with severe adult periodontis. In addition, microbiological monitoring will be performed in order to determine of pre-treatment microbiological status influences treatment outcome.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Oral health/stomatognathic diseases: Severe periodontal disease

Interventions
Not provided at time of registration

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
metronidazole and amoxycillin

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/1994

Completion date
01/06/1997

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
Added 06/01/10: 66

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/1994

Date of final enrolment
01/06/1997

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

University of Bristol
Bristol
United Kingdom
BS1 2LY

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive South West (UK)

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	01/04/2002		Yes	No