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

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

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

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# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers

# 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