

# A comparison of resorbable versus metal plates for the management of fractured mandibles

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 14/03/2014	<b>Condition category</b> Oral Health	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

**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**  
N0190133723

## Study information

## **Scientific Title**

### **Study objectives**

To establish the efficacy and complication rate of the use of resorbable plates in fractured mandibles compared to the established treatment of metal plates.

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Not provided at time of registration

### **Study design**

Blinded 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: Oral medicine

### **Interventions**

Resorbable versus metal plates

### **Intervention Type**

Other

### **Phase**

Not Specified

### **Primary outcome measure**

Not provided at time of registration

### **Secondary outcome measures**

Not provided at time of registration

### **Overall study start date**

01/03/2004

**Completion date**

30/09/2007

## Eligibility

**Key inclusion criteria**

Not provided at time of registration

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

41 patients

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/03/2004

**Date of final enrolment**

30/09/2007

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

The Queen Victoria Hospital NHS Trust

East Grinstead

United Kingdom

RH19 3DZ

## Sponsor information

**Organisation**

Department of Health

**Sponsor details**

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

**Sponsor type**

Government

**Website**

<http://www.dh.gov.uk/Home/fs/en>

**Funder(s)**

**Funder type**

Government

**Funder Name**

Queen Victoria Hospital NHS Trust (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