# Impact of treatment of chronic periodontitis on the serum levels of prohepcidin in patients with chronic kidney disease

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
07/11/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
18/11/2010	Completed	[X] Results
Last Edited	Condition category	[] Individual participant data
03/01/2012	Oral Health	

## Plain English summary of protocol

Not provided at time of registration

## Contact information

## Type(s)

Scientific

#### Contact name

Prof Eduardo Vilela

#### Contact details

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## Additional identifiers

**EudraCT/CTIS** number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers # 942.248.2006/report # 327/2006

## Study information

#### Scientific Title

Impact of treatment of chronic periodontitis on the serum levels of prohepcidin in patients with chronic kidney disease: Interventional controlled clinical assay

#### Acronym

**UFJF** 

#### **Study objectives**

We hypothesized that part of the chronic inflammatory response seen in chronic kidney disease (CKD) patients stems from chronic periodontitis (CP), which, through the increase in the expression of inflammatory markers such as interleukin-6 (IL-6), stimulates hepcidin synthesis.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethics Committee in Research on Human Beings from UFJF approved on the 6th of December 2006 (ref: 942.248.2006 / report: 327/2006)

#### Study design

Interventional controlled clinical assay

#### 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 contact details below to request a patient information sheet.

#### Health condition(s) or problem(s) studied

Chronic periodontitis (CP); chronic kidney disease (CKD); Prohepcidin

#### **Interventions**

Patients with CP were divided into two groups: CKD patients and controls Both groups received instructions on oral hygiene techniques, including the manual techniques of tooth and interproximal brushing, and on how to use dental floss and perform supragingival prophylaxis. The nonsurgical periodontal therapy (PT) consisted of radicular scraping and subgingival curettage, using standard instrumentation with Gracey curettes and ultrasound devices performed in 1 hour sessions over an average period of 4 weeks. Local anesthesia was used when necessary.

Upon concluding the periodontal treatment, participants were followed up after 15, 30, 60, and 90 days. At each return visit, instructions on oral hygiene and supragingival phrophylaxis were provided.

Blood samples were collected for biochemical analysis at baseline and 3 months after PT. Venous blood after 12 hours fasting was collected in vacuum tubes between 7:00am and 9: 00am. Plasma samples with EDTA/heparin and serum samples were immediately placed in ice, aliquoted within 1 hour and stored at -80C until use.

#### Results:

The efficacy of PT was proven by the statistically significant decrease in the levels of inflammatory markers and the improvement of clinical parameters of CP observed 3 months after completion of the PT. Prohepcidin, IL-6 and us-PCR levels diminished significantly after PT in both groups. In the control group, besides the decrease in the inflammatory markers, a significant increase could also be observed in the levels of haemoglobin and ferritin associated with PT.

#### Intervention Type

Other

#### **Phase**

Not Applicable

#### Primary outcome measure

Inflammatory markers, assessed at baseline and 3 months post-PT

- 1. C-reactive protein (us-CRP)
- 2. Interleukin-6 (IL-6)
- 3. Prohepcidin

#### Secondary outcome measures

Biochemical analyses from blood sample in EDTA, at baseline and 3 months post-PT

- 1. Complete haemogram (automated Coulter STKS)
- 2. Serum iron (ferrozine)
- 3. Ferritin (electrochemiluminescence)
- 4. Transferrine saturation index (labtest ferrozine)

#### Overall study start date

10/08/2008

#### Completion date

10/08/2010

## **Eligibility**

#### Key inclusion criteria

1. Patients with CP, allocated to one of two groups depending on CKD status

1.1. CKD group:

- 1.1.1. Patients with CKD at stages 3 to 5 and undergoing conservative treatment
- 1.1.2. Recruited from the PREVENRIM, a CKD prevention clinic at the Interdisciplinary Nucleus of Studies, Research and Treatment in Nephrology (NIEPEN) of the Universidade Federal de Juiz de Fora (UFJF)
- 1.2. Control patients:
- 1.2.1. Patients with no systemic disease from the Periodontology Clinic of the School of Dentistry at UFJF
- 2. Over 18 years of age, either sex
- 2. Minimum of 20 natural teeth and without periapical lesions
- 3. Have received no periodontal, antimicrobial, or anti-inflammatory treatment within the last 6 months
- 4. Have not used steroids or immunosuppressant drugs within the last 6 months

#### Participant type(s)

Patient

#### Age group

Adult

#### Lower age limit

18 Years

#### Sex

Both

#### Target number of participants

56

#### Key exclusion criteria

- 1. Pregnant or breast feeding women
- 2. Smokers or ex-smokers who had quit smoking within the last 10 years

#### Date of first enrolment

10/08/2008

#### Date of final enrolment

10/08/2010

## Locations

#### Countries of recruitment

Brazil

### Study participating centre Rua Dr Geraldo Moutinho 55

Juiz de Fora Brazil 36036348

## Sponsor information

### Organisation

Federal University of Juiz de Fora (Universidade Federal de Juiz de Fora [UFJF]) (Brazil)

#### Sponsor details

Rua Dr Geraldo Moutinho 55 Jardins Imperiais Juiz de Fora Brazil 36036348

#### Sponsor type

University/education

#### **ROR**

https://ror.org/04yqw9c44

## Funder(s)

## Funder type

University/education

#### **Funder Name**

Federal University of Juiz de Fora (Universidade Federal de Juiz de Fora [UFJF]) (Brazil)

## **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	01/07/2011		Yes	No