Lipid-A project

Submission date	Recruitment status No longer recruiting	Prospectively registered		
08/01/2015		☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
08/01/2015	Completed	[X] Results		
Last Edited 23/01/2019	Condition category Oral Health	[] Individual participant data		
23/01/2013	Oraciricateri			

Plain English summary of protocol

Background and study aims

Periodontitis is a chronic inflammatory disease of the tissues supporting the teeth and one of the main causes of tooth loss. It is caused by bacteria present in a shallow crevice between the gums and the teeth which triggers the host inflammatory response. This exacerbated host inflammatory response is responsible for tissue damage and the loss of the bone that supports teeth. Despite being a very common oral disease, diagnostic procedure and treatment option for periodontitis have not changed significantly for more than a century. It is usually diagnosed too late, once bone less has already occurred, and treated, not very successfully, by mechanical removal of subgingival deposites. Therefore, there is a great need for new diagnostic approaches and treatment options for this widespread disease. Bacterial species that are responsible for development of periodontal diseases are able to produce a very potent substance, lipopolysaccharide (LPS), that triggers the host inflammatory response. If this type of bacterial antigens are detected in the mouth before the development or in early stages of the disease, a successful preventive regimen could be employed and harmful consequences avoided. The aim of this study is to see if there are any differences in the chemical composition of lipopolysaccharides isolated from subgingival dental plaque, saliva from patients with gum diseases and patients with healthy gums.

Who can participate?
Adults with or without chronic periodontitis

What does the study involve?

Saliva and subgingival deposit samples are collected from both patients with periodontitis and healthy volunteers treated at the Peninsula School of Dentistry clinics in Plymouth and Truro. LPS is extracted from these samples and its chemical composition (with a focus on Lipid-A, its most potent part) examined in the research laboratories of the Plymouth University. The chemical composition of LPS is compared between healthy participants and patients with periodontitis as well as between samples from the same periodontitis patient taken before and after routine periodontal treatment.

What are the possible benefits and risks of participating? Not provided at time of registration Where is the study run from?
The Peninsula School of Dentistry clinics in Plymouth and Truro (UK)

When is the study starting and how long is it expected to run for? August 2014 to June 2015

Who is funding the study?
GSK and Oral and Dental Research Trust (UK)

Who is the main contact? Dr Svetislav Zaric

Contact information

Type(s)

Scientific

Contact name

Dr Svetislav Zaric

ORCID ID

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 18081

Study information

Scientific Title

Subgingival plaque lipid-A profile as a bacterially-derived biomarker for chronic periodontitis

Study objectives

Are there differences in the chemical composition of lipopolysaccharides isolated from subgingival dental plaque and saliva from patients with gum diseases and patients with healthy gums?

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES Committee South West - Cornwall and Plymouth, 03/03/2014, ref:14/SW/0020

Study design

Non-randomised study

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Oral & Dental; Subtopic: Oral & Dental Public Health; Disease: All Oral & Dental

Interventions

- 1. Full mouth periodontal therapy
- 2. Full mouth periodontal debridement

Intervention Type

Procedure/Surgery

Primary outcome measure

Change in subginigval lipid-A profile; Timepoint(s): 3 months after completion of periodontal therapy

Secondary outcome measures

N/A

Overall study start date

20/08/2014

Completion date

Eligibility

Key inclusion criteria

- 1. Healthy (PD <4 mm, no evidence of attachment and bone loss, and <10% of sites with BOP)
- 2. Chronic periodontitis (presence of 1/3 bone loss or more and presence of 6 mm pocket or more, with evidence of attachment loss at 2 or more teeth per quadrant and evidence of generalized bleeding)

Participant type(s)

Mixed

Age group

Adult

Sex

Both

Target number of participants

Planned Sample Size: 60; UK Sample Size: 60; Description: 30 patients with chronic periodontitis and 30 individuals with healthy periodontium

Key exclusion criteria

- 1. Systemic chronic diseases
- 2. Acute medical interventions or diseases 4 weeks before baseline
- 3. Pregnancy
- 4. Antibiotics 6 months before intake and or repeated use of NSAID's medications 4 weeks before intake

Date of first enrolment

20/08/2014

Date of final enrolment

30/06/2015

Locations

Countries of recruitment

England

United Kingdom

Study participating centre University of Plymouth

Peninsula Dental School Plymouth University Portland Square C406 Plymouth United Kingdom PL4 8AA

Sponsor information

Organisation

Peninsula Dental Social Enterprise CIC

Sponsor details

Peninsula Dental School Plymouth University Portland Square C406 Plymouth England United Kingdom PL4 8AA

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/04dtfyh05

Funder(s)

Funder type

Government

Funder Name

GSK and Oral and Dental Research Trust (UK)

Results and Publications

Publication and dissemination plan

Results of the study will be reported and disseminated by conference presentations and peer reviewed scientific journal publications by the end of 2017.

Intention to publish date

31/12/2017

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are not expected to be made available. The participant level data is confidential and is stored in the patients' record system SoelHealth.

IPD sharing plan summary

Not expected to be made available

Study outputs

Output type	Details results	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		01/09/2019	22/01/2019	Yes	No
HRA research summary			28/06/2023	No	No