# The effect of the natural mineral supplement Aquamin™ together with the short chain fructooligosaccaride Nutraflora® on bone health in post-menopausal women

Submission date	Recruitment status  No longer recruiting	<ul><li>Prospectively registered</li></ul>			
12/08/2009		☐ Protocol			
Registration date	Overall study status Completed	Statistical analysis plan			
02/10/2009		[X] Results			
Last Edited	Condition category	Individual participant data			
01/08/2022	Musculoskeletal Diseases				

## Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Dr Emeir Duffy

#### Contact details

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# Additional identifiers

**Protocol serial number** N/A

# Study information

#### Scientific Title

The effect of the mineral supplement Aquamin™ together with the short chain fructooligosaccaride Nutraflora® on bone mineral density and bone turnover markers in postmenopausal women: a double-blind, randomised, placebo-controlled trial

#### Study objectives

Osteoporosis and low bone mass is becoming a major public health problem with postmenopausal women being at the highest risk. Osteoporosis is a disease characterised by low bone mineral density (BMD), where the structure of bone deteriorates, making it more susceptible to fractures, especially in the spine, hip and wrist. A recent animal study has shown the mineral supplement Aquamin™ to enhance bone mineral density. Furthermore, there is also evidence that short-chain fructooligosaccharide (Nutraflora®) supplementation enhances bone mineral density in rats. This study sets out to investigate the effect of supplementation of Aquamin™ together with the short chain fructooligosaccaride Nutraflora® on bone health using bone mineral density and bone turnover markers as indicators of bone health in postmenopausal women.

### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Research Ethics Committee of the University of Ulster approved on the 21st July 2008 (ref: REC /08/0083)

#### Study design

Double-blind randomised placebo-controlled trial

## Primary study design

Interventional

## Study type(s)

Treatment

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

Osteoporosis

#### **Interventions**

Each treatment is incorporated into chocolate chews and will be supplemented with 2 x chocolate flavoured chews. Individuals are supplemented per day for 2 years.

Treatment 1: Placebo (maltrodextrin)

Treatment 2: Aquamin<sup>™</sup> (1800 mg/d) alone

Treatment 3: Aquamin™ (1800 mg/d) with Nutraflora® (3.6 g/d)

## Intervention Type

Drug

#### Phase

Not Applicable

# Drug/device/biological/vaccine name(s)

Aquamin™, Nutraflora®

#### Primary outcome(s)

Bone mineral density: dual energy X-ray absorptiometry

#### Key secondary outcome(s))

- 1. Bone resorption and formation markers:
- 1.1. Serum CTx: ELISA
- 1.2. Urinary deoxypryridinoline cross-links: ELISA
- 1.3. Urinary CTx: ELISA
- 2. Serum intact osteocalcin: ELISA
- 3. Serum N-terminal midfragment OC: ELISA
- 4. Vitamin D: ELISA
- 5. Free calcium: ELISA
- 6. Parathyroid hormone: ELISA
- 7. Salivary Cortisol: ELISACalcium: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 8. Phosphate: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 9. Full Blood Profile: Sysmex full blood analyser
- 10. Electrolytes: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 11. Liver function:
- 11.1. Albumin: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, United)
- 11.2. Alanine transaminase: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 11.3. Gamma-glutamyltransferase: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, United)
- 11.4. Total bilirubin: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, United)
- 11.5. Alkaline Phosphatase: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 12. Total protein: Automated instrument (Instrumentation Laboratories UK Ltd, Warrington, UK)
- 13. Anthropometric measurements: height, weight, waist circumference, measured using standardised procedures
- 14. Blood pressure
- 15. Mean daily dietary intake (4-day diet diary)
- 16. Habitual dietary intake: Food Frequency Questionnaire
- 17. Exercise: Physical Activtiy Questionnaire
- 18. Sun Exposure: Questionnaire

#### Completion date

31/07/2011

# Eligibility

#### Key inclusion criteria

- 1. Apparently healthy post-menopausal female volunteers (defined as having no menstrual period, bleeding or spotting during 1 year prior to enrolment)
- 2. Aged 48 75 years old
- 3. Weight less than 136 kg (DEXA [dual energy x-ray absorptiometry] limit 136 kg)
- 4. Community dwelling and fully mobile, with hormone implants (if used) removed at least one year prior to randomisation

#### Participant type(s)

Patient

#### Healthy volunteers allowed

No

#### Age group

Adult

#### Sex

**Female** 

#### Total final enrolment

300

#### Key exclusion criteria

- 1. Osteoporotic bone density (T-score less than -2.5)
- 2. Corticosteroid medications during the previous 6 months
- 3. History/presence of chronic renal, hepatic, gastrointestinal disease or traumatic lumbar compression fracture
- 4. Evidence of collapsed or focal vertebral sclerosis
- 5. Menopause before the age of 40 years
- 6. Bone diseases or other condition known to affect bone status
- 7. Treatment with specific therapy for osteoporosis
- 8. Uncontrolled hypertension or heart failure, renal calculi
- 9. Volunteers should not have used any prescribed medication known to affect bone status
- 10. Use of dietary supplements containing calcium and vitamin D three months prior to the study

#### Date of first enrolment

01/01/2009

#### Date of final enrolment

31/07/2011

# Locations

#### Countries of recruitment

**United Kingdom** 

Northern Ireland

Ireland

Study participating centre University of Ulster

Coleraine United Kingdom BT52 1SA

# Sponsor information

# Organisation

Marigot Ltd (Ireland)

#### **ROR**

https://ror.org/05kmpn815

# Funder(s)

# Funder type

Industry

#### Funder Name

Marigot Ltd (Ireland)

# **Results and Publications**

# Individual participant data (IPD) sharing plan

Not provided at time of registration

# IPD sharing plan summary

Not provided at time of registration

# **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient- Pfacing?
Results article	results	01/03 /2014		Yes	No
Other publications	Associations of long chain polyunsaturated fatty acids with bone mineral density and bone turnover	30/07 /2022	01/08 /2022	Yes	No
Participant information sheet	Participant information sheet	11/11 /2025	11/11 /2025	No	Yes