



Abstract **Protective Effects of Green Shelled Mussels in Osteoarthritis** ⁺

Parkpoom. Siriarchavatana ¹, Marlena C. Kruger ^{2,3,4}, Matt Miller ⁵, Hong (Sabrina) Tian ⁶ and Fran M. Wolber ⁷*

- ¹ School of Food and Advanced Technology, Massey University, Palmerston North, New Zealand; P.Siriarchavatana@massey.ac.nz
- ² College of Health, Massey University, Palmerston North, New Zealand; M.C.Kruger@massey.ac.nz
- ³ School of Food and Advanced Technology, College of Sciences, Massey University, Palmerston North, New Zealand
- ⁴ Riddet Institute, Massey University, Palmerston North, New Zealand
- ⁵ Cawthron Institute, Nelson, New Zealand; Matt.Miller@cawthron.org.nz
- ⁶ Sanford Ltd., Auckland, New Zealand; STian@sanford.co.nz
- ⁷ Centre for Metabolic Health Research, Massey University, Palmerston North, New Zealand
- * correspondence: F.M.Wolber@massey.ac.nz
- + Presented at the 2018 Nutrition Society of New Zealand Annual Conference, Auckland, New Zealand, 28–30 November 2018.

Published: 13 March 2019

Background: Obesity-induced chronic inflammation is associated with metabolic syndrome, and often leads to the development of osteoarthritis. In osteoarthritis, inflammatory cells act on both bone cells and cartilage cells to cause destruction of the joint. Green shelled mussels (GSM), a seafood native to New Zealand, have been shown to inhibit inflammation and reduce pain in the joints of animals with arthritis.

Methods: Female Sprague-Dawley rats (N \ge 10 per group) were fed a normal control (CON) or high-fat/high-sugar (HFHS) diet with or without the inclusion of freeze-dried whole GSM from age 3 months to 7 months. Rats were assessed for serum levels of CTX-2, a biomarker for cartilage degradation. In addition, GSM extracts were used in vitro to treat undifferentiated bone cells (RAW 264.7 cells) and assessed for their ability to prevent the cells from differentiating into bone-destroying osteoclasts and producing tartrate-resistant acid phosphatase (TRAP).

Results: Unsurprisingly, rats fed a HSHS diet gained more weight and produced more CTX-2 compared to those fed a CON diet. However, the inclusion of GSM in both diets resulted in a reduction of cartilage degradation: CTX-2 levels in rats fed CON vs. CON + GSM were 194 ± 26 vs. 161 ± 25 pg/mL, while CTX-2 levels in rats fed HFHS vs HFHS + GSM were 241 ± 31 vs. 151 ± 19 pg/mL (p = 0.02). In vitro, a non-polar lipid extract of GSM significantly reduced osteoclast differentiation in a dose-dependent manner, with treatment at 20 µg/mL reducing differentiation by 80% and TRAP production by 85%, whereas a polar lipid extract had no effect.

Conclusions: Dietary GSM significantly reduces the development of joint osteoarthritis caused by diet-induced metabolic syndrome in obese rats. Non-polar lipids in GSM in vitro significantly reduces the development of bone-resorbing osteoclast cells. Inclusion of GSM in the human diet is likely to protect both joint and bone health.

Supplementary Material: The poster is available online at www.mdpi.com/2504-3900/8/1/50/s1.



© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).