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Increased bioavailability of phenolic acids and enhanced vascular function following intake of feruloyl esterase-processed high fibre bread: a randomized, controlled, single blind, crossover human intervention trial.

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Short Running Head: Free FA bread and vascular function

Clinical Trial Registry: No: NCT03946293; **Website:** www.clinicaltrials.gov

2 ABSTRACT

3 **Background and Aims:** Clinical trial data have indicated an association between wholegrain
4 consumption and a reduction in surrogate markers of cardiovascular disease. Phenolics present in
5 wholegrain bound to arabinoxylan fibre may contribute these effects, particularly when released
6 enzymatically from the fiber prior to ingestion. The aim of the present study was therefore to
7 determine whether the intake of high fibre bread containing higher free ferulic acid (FA) levels
8 (enzymatically released during processing) enhances human endothelium-dependent vascular
9 function.

10 **Methods:** A randomized, single masked, controlled, crossover, human intervention study was
11 conducted on 19 healthy men. Individuals consumed either a high fibre flatbread with enzymatically
12 released free FA (14.22 mg), an equivalent standard high fibre bread (2.34 mg), or a white bread
13 control (0.48 mg) and markers of vascular function and plasma phenolic acid concentrations were
14 measured at baseline, 2, 5 and 7 hours post consumption.

15 **Results:** Significantly increased brachial arterial dilation was observed following consumption of
16 the high free FA ('enzyme-treated') high fibre bread verses both a white flatbread (2h: $p<0.05$; 5h:
17 $p<0.01$) and a standard high fibre flatbread (5h: $p<0.05$). Concurrently, significant increases in
18 plasma FA levels were observed, at 2 h ($p<0.01$) after consumption of the enzyme-treated flatbread,
19 relative to control treatments. Blood pressure, heart rate, DVP-SI and DVP-RI were not
20 significantly altered following intake of any of the breads ($p>0.05$).

21 **Conclusion:** Dietary intake of bread, processed enzymatically to release FA from arabinoxylan
22 fiber during production increases the bioavailability of FA, and induces acute endothelium-
23 dependent vasodilation.

24 **Key Words:** bioavailability, ferulic acid, vascular function, high fibre bread, human intervention,
25 feruloyl esterase.

26

Abbreviations: Cardio-Vascular Disease (CVD), Flow Mediated Dilatation (FMD), Laser Doppler Imaging (LDI), Digital Pulse Wave (DVP), Blood Pressure (BP), acetylcholine (Ach), 3,5-Dichloro-4-hydroxybenzoic acid (3,5DHBA)

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28 INTRODUCTION

29 Clinical data suggest that the regular intake of foods/beverages rich in phenolic compounds, such as
30 cocoa, green tea, and berries, improves human vascular function and maintains a healthy circulatory
31 function [1-4]. However, such foods/beverages tend to be expensive, have limited availability and
32 are not consumed by large proportions of the population. Thus, to increase the intake of phenolic
33 compounds at the population level, in a sustainable, economically viable way, requires their
34 delivery through foods derived from staple crops such as rice, maize and wheat. Wholegrain wheat
35 is a rich source of phenolic acids, in particular ferulic acid (FA) [5,6], in addition to vitamins,
36 minerals, phytosterols, unsaturated fatty acids and lignans [7,8], and is consumed as both white and
37 wholemeal breads and other cereal products. There is robust scientific evidence that diets rich in
38 wholegrain have a beneficial effect on cardiovascular health [9-12] and current dietary guidelines
39 emphasize the importance of wholegrain in the diet [13].

40 A number of components of wheat have been suggested to contribute to the cardiovascular health
41 benefits associated with wholegrain consumption, most notably the fiber component [6,11,14,15].
42 However, phenolic compounds that are bound to arabinoxylan fibre, particularly the
43 hydroxycinnamate FA, may also contribute, in part, to vascular health effects [16-18]. FA
44 comprises up to 90 % of the phenolic acids present in wholegrain wheat and is concentrated in the
45 outer and aleurone layers of the grain (the bran) where it is present in three forms: as soluble free
46 acid, soluble conjugates, and insoluble bound forms [6,19]. The latter constitute the majority of FA,
47 and is covalently linked to arabinoxylan polysaccharides via stable ester bonds. Bound FA may also
48 act as structural cross-links in cell walls, by oxidative dimerization [20]. Although cleavage of the
49 ester linkage to release free FA is possible in the colon via fecal microbial xylanases and esterases,
50 it is estimated that only about 25% of bound FA is released during this phase of digestion [21-24].
51 Consequently, its liberation and absorption in the gastrointestinal tract and potential to induce
52 beneficial vascular changes in humans is limited, although slow release and absorption of phenolic
53 acids in the gut may contribute to the long-term benefits of regular wholegrain consumption [25].

54 In the present study, we produced high fibre bread with high amounts of free FA, released
55 enzymatically during dough development, in order to determine whether dietary-relevant amounts
56 of free FA delivered in high fibre bread are capable of inducing acute beneficial effects upon
57 endothelial function.

58 **SUBJECTS AND METHODS**

59 **Production and composition of breads**

60 *Triticum aestivum* cv Hereward was grown at Rothamsted Research (Harpenden, UK) and milled by
61 Campden BRI (Chipping Campden, UK) using a Buhler MLU-202 mill (Urzwil, Switzerland) to
62 give 10 fractions; Reduction 1 (R1), Reduction 2 (R2), Reduction 3 (R3), Break 1 (B1), Break 2
63 (B2), Break 3 (B3), Bran Flour (BF), Offal Flour (OF), Bran Off-tails (BO) and Offal Off-tails
64 (OOT). Selected fractions were combined to produce experimental unleavened breads: white flour
65 was prepared by combining breaks 1, 2 and 3 and reductions 1, 2 and 3 with an extraction rate of
66 72.0% of total flour. 'High fibre' flour was prepared by combining 9 fractions: breaks 1, 2 and 3,
67 reductions 1, 2 and 3, bran flour, offal flour, and offal off tails with an extraction rate of 85% of
68 total flour. The bran off-tails were not included in the 'high fibre' flour as their coarse texture
69 reduced the palatability of the breads. The FA content of the milled wheat fractions is shown in
70 **Supplemental data, Table S1**. Intervention breads (unleavened, flatbreads) were produced by
71 combining flour, water, enzyme (Ultraflo L[®]) and salt (**Table 1**) for 5 min using a Buhler mixer.
72 Ultraflo L[®], a commercial β -glucanase, which also possesses xylanase and feruloyl esterase
73 activities, was used to release FA bound to arabinoxylan. For the breads used in the intervention
74 study this enzyme mixture was diluted in water at a ratio of 95:5 (water:Ultraflo L[®], (v/v)). Dough
75 was made using flour, water (containing the 5% enzyme solution) and salt in the following
76 proportions 60.5% flour; 38.7% water; 0.7% salt. The dough was left to prove for 4 hours in a
77 temperature-controlled environment (20°C), before being divided into 50 g portions and rolled to 13
78 cm x 2mm disks. The unleavened flatbreads were baked for 7 minutes at 230°C and left to cool.

79 The control breads (white and high fibre) contained deactivated Ultraflo L® (95°C for 30 minutes,
80 then cooled on ice).

81

82 The white intervention bread differed in appearance to the high fiber breads due to the flours used
83 in each, although there were no observable differences between the standard and enzyme-treated
84 high fibre breads. The optimum conditions for enzymatic release of FA from arabinoxylan during
85 dough proving were established prior to the production of breads for intervention (Supplementary
86 Figures 2A and 2B). All breads were made at the same time and stored at -20°C until use. No
87 significant degradation of FA was observed within breads following 2 or 4 weeks of storage at -
88 20°C ($p>0.05$) (data not shown).

89

90 Phenolic acids present in the original milling fractions ($n=3$) (Supplementary Table S1) and in the
91 final intervention breads ($n=3$) (Table 1) were quantified as described by [6]. Briefly, after addition
92 of an internal standard free phenolic acids were extracted from finely ground material in 80% (v/v)
93 ethanol. Bound phenolics were determined in samples after the removal of free and conjugated
94 phenolic acids. Internal standard was added to the remaining residue and 2M NaOH added to
95 release the esterified phenolic acids. Following centrifugation, the supernatant was acidified with
96 12M HCl, mixed thoroughly, ethyl acetate added, mixed and centrifuged again. The upper phase
97 was removed, and the ethyl acetate extraction repeated a further two times. The combined
98 supernatants are evaporated to dryness in a Speedvac. The dry sample was then re-suspended in 2%
99 (v/v) acetic acid, centrifuged and FA levels quantified using an Agilent 1100 HPLC equipped with a
100 diode array detector (as described by [6], using authentic standards and an internal standard (3,5-
101 Dichloro-4-hydroxybenzoic acid). All fractions were dried prior to re-suspension in 2% (v/v) acetic
102 acid. 50 μ l of each extract was separated using a Discovery RP-Amide C16 column (250 x4.6mm,
103 5 μ m) maintained at 30°C. The mobile phases consisted of a mixture of acetonitrile (solution A)
104 and 2% (v/v) acetic acid in water (solution B) with a flow rate of 1.0 mL/min. The following

105 gradient system was used (min/% B): 0/100, 30/85, 50/50, 70/30, with 10 min post-run for column
106 re-equilibration. The wavelength used for quantification of FA was 280 nm and spectral
107 characteristics were scanned over the range 200–600 nm. All data were analyzed using
108 ChemStation software.

109 **Ethics**

110 The clinical trial was registered at clinicaltrials.gov (NCT03946293) and conducted according to
111 the Declaration of Helsinki following Good Clinical Practice (GCP) and was approved for conduct
112 by the University of Reading's Research Ethics Committee (ethics reference number 12/06). All
113 volunteers signed an informed consent form before commencing the study.

114 **Intervention study population**

115 Healthy male volunteers ($n = 19$) (**Figure 1**) were recruited from the University of Reading and
116 surrounding areas by use of the Hugh Sinclair Unit volunteer database, poster advertisement within
117 the university and local community via local websites. Volunteers were screened and selected
118 according to the following inclusion criteria: 1) fasting lipids in the upper half of the normal range
119 (triacylglycerol 0.8-3.2 mmol/l and total cholesterol 6.0-8.0 mmol/l); 2) BMI 25-32 kg/m²; 3) non-
120 smoker; 4) not diabetic (diagnosed or fasting glucose < 7 mmol/l) or suffer from endocrine
121 disorders; 5) hemoglobin and liver enzymes levels within the normal range [Alanine Transaminase
122 (ALT): 0-55 IU/L; Alkaline Phosphatase (ALP): 38-126 U/L; Aspartate Transaminase (AST): 0-45
123 IU/L; Gamma Glutamyl Transferase (GGT): 12-58 IU/L]); 6) not having suffered a myocardial
124 infarction/stroke in the past 12 months; 7) not suffering from renal or bowel disease or have a
125 history of cholestatic liver or pancreatitis; 8) not on drug treatment for hyperlipidemia,
126 hypertension, inflammation or hyper-coagulation; 9) not taking any fish oil, fatty acid or vitamin
127 and mineral supplements; 10) no history of alcohol misuse; 11) not planning, or on a weight
128 reduction regime; 12) not having taken antibiotics in the 6 months prior to the study; and 13) being
129 able to consume the study interventions. The 19 individuals selected for the study were instructed

130 not to alter their usual dietary or fluid intake, with respect to amount consumed. Volunteers were
131 requested to refrain from the following, 24 h prior to, and during, the study visits: 1) consumption
132 of polyphenol-rich foods including fruits, vegetables, wholegrain bread/pasta/rice, cocoa, coffee, tea
133 and wine; 2) participating in vigorous exercise (> 3 x 20 min/week); and 3) consuming more than
134 168 g of alcohol (any form) per week. Volunteers were further asked to fast for 12 hours before
135 each study visit and during that period only consume low-nitrate water provided. The Baseline
136 characteristics of the study population are presented in Table 1.

137 **Study design**

138 The study was a randomized, single blind, crossover, controlled intervention trial in which subjects
139 attended the Hugh Sinclair Unit of Human Nutrition on 3 separate occasions. We considered the
140 trial to be single blinded (researcher) as interventions differed slightly in appearance, although no
141 information regarding the interventions could be implied from this. Visits were separated by at
142 least 2 weeks, and treatments to subjects were allocated randomly using a restricted Williams
143 design (as patients were recruited sequentially) by an independent researcher, who implemented the
144 allocation sequence on the first visit. On arrival, subjects were rested in the supine position for at
145 least 30 minutes in a quiet, air-conditioned environment (22-24°C), during which time they were
146 cannulated for venous blood collection and baseline vascular measurements were taken, including
147 FMD of the brachial artery (primary outcome), laser Doppler imaging (LDI) with iontophoresis,
148 and digital pulse wave (DVP), and systolic and diastolic blood pressure (Omron MX2 automatic
149 digital upper arm blood pressure monitor). Following collection of baseline measurements,
150 volunteers were requested to consume three 30 g flatbreads (white, standard high fibre, or enzyme-
151 treated high fibre) within a 5 min period and further blood was collected at 1, 2, 3, 4, 5, 6 and 7 h
152 and vascular measurements were conducted at 2, 5 and 7 h. Volunteers were provided with a low-
153 fat, low-polyphenol lunch (consisting of 2 slices of white bread, low fat cream cheese containing
154 3% fat, lightly salted crisps and a low-fat vanilla yoghurt) at 3 h and had free access to bottled water
155 containing low nitrate/nitrite levels) throughout the study day. Following completion of each

156 experimental arm, participants followed a washout period of 14 days prior, where they were asked
157 to follow their normal habitual diet, before switching to an alternate arm of the trial. A qualified
158 nurse was responsible for cannulation and blood collection. All study personnel, including research
159 nurses and research staff were blinded to the interventions. The study followed ethical standards,
160 which were in accordance with the University of Reading Ethics Committee. No changes were
161 made to any of the trial methods or outcomes following the commencement of the trial.
162 Recruitment for the trial took place between Oct 2012 and February 2013, whilst the trial was
163 conducted between January 2013 and January 2014.

164 **Flow Mediated Dilatation (FMD)**

165 Subjects were positioned into the supine position with the right arm rested on a custom-made arm
166 support at a 90° angle with their hand facing towards the ceiling. Electrocardiogram (ECG) pads
167 were attached to time each image frame with respect to the cardiac cycle. A sphygmomanometric
168 cuff was placed on the forearm and an image of the longitudinal brachial artery was obtained using
169 a 3.5 x 1.0 cm hockey-stick probe, attached to a flexible holding device with locking action to allow
170 for accurate measurements, and ATL Ultrasound HDI5000 broadband ultrasound system. The live
171 image was captured using image-grabbing software (MIA-11c) and collected images at 0.25
172 frames/second. A baseline measurement was recorded for 40 seconds using the flow. After this the
173 main FMD measurement was started by recording the image for 60 seconds before arterial
174 occlusion was created by inflating the cuff to 220mmHg causing ischemia of the arterial vessel.
175 After 5 minutes the cuff was deflated, rapidly causing a decrease in pressure leading to reactive
176 hyperemia to accommodate the dilated resistance levels. Image collection continued for 5 minutes
177 post release and analysis of the arterial diameter was performed to assess the flexibility of the artery
178 calculated as the relative diastolic diameter change from baseline compared to the peak diastolic
179 diameter. Analysis was performed using a semi-automated computerized analysis system (Brachial
180 Analyzer; Medical Imaging Applications-11c). In order to avoid potential inter-individual, and
181 intra-individual variability of measurements within our experimental group, the same researcher

182 conducted all FMD measurements. Additionally, care was taken to conduct brachial artery flow-
183 mediated dilatation measurements consistently, with probe placements and cuff positioning
184 carefully recorded and replicated between and across individuals.

185

186 **Laser Doppler Imaging with iontophoresis (LDI)**

187 The assessment of peripheral microvascular function [26,27] was conducted via the administration
188 of an endothelial-dependent vasodilator, acetylcholine ACh (1%), and an endothelium in-dependent
189 vasodilator, sodium nitroprusside (SNP) (1%), to the skin, which were delivered transdermally
190 using iontophoresis. Subjects were rested in the supine position, with their right arm supported by
191 an armrest. Two ION 6 chambers were applied to the forearm, within 1cm of each other, using
192 double sided adhesive disks. 2.5ml of freshly prepared ACh (1% w/v in 0.5% (w/v) NaCl solution;
193 Sigma Aldrich, Poole, Dorset, UK) was introduced to the anodal chamber and 2.5ml of SNP (1%
194 w/v in 0.5% (w/v) NaCl solution; Sigma Aldrich, Poole, Dorset, UK) into the cathodal chamber. A
195 coverslip was placed over each chamber to prevent any liquid from escaping. After baseline of skin
196 perfusion was measured, an incremental electrical current was delivered simultaneously across the
197 two chambers using the laser Doppler imager (LD12-VR) and MIC2 iontophoresis controller (Moor
198 instruments Ltd) (increased in 5 μ A steps; 5, 10, 15 and 20 μ A to yield a total charge of 8
199 milliCoulombs within 12 minutes). A total of 20 scans were performed. An indicator of
200 microvascular response due to ACh and SNP was calculated by the area under the flux versus time
201 curve during the 20 scans.

202 **Digital Volume Pulse (DVP)**

203 The DVP stiffness index (DVP-SI) and DVP reflexion index (DVP-RI) was calculated by placing a
204 PulseTrace PCA 2 with a photoplethysmograph transducer (MircoMedical, Kent) on the index
205 finger of the left hand, transmitting an infrared light at 940nm. DVP was conducted as previously
206 described [28-30]. The infrared light transmission through the finger records the systolic and
207 diastolic waveforms of the pulse, and also records the stiffness index (DVP-SI) and reflexion index

208 (DVP-RI). The DVP-SI (in m/s) is related to large arterial stiffness and is measured by dividing the
209 height of the subject by the time between the first and the second wave peaks of the DVP. It is
210 related to CVD risk factors such as blood pressure, age and waist to hip ratio. The DVP-RI is
211 related to smaller artery stiffness and is defined as the relative height of the second peak compared
212 with the first.

213 **Plasma FA analysis**

214 Blood was collected at baseline and 1, 2, 3, 4, 5,6 and 7h after ingestion and processed as described
215 in [31] and analyzed as follows. LC-MS analysis, as described by [32], with minor modifications
216 was also used to assess FA levels and other phenolic acids in human plasma samples. Measurements
217 were performed on an ABSciex 4000QTRAP MS system linked to an Agilent 1200 HPLC, column
218 oven set to 30 °C, flow rate 200 µl/min. The column eluent was introduced into the mass spectrometer
219 by a TurboIonSpray probe operating at 475 °C, with ion spray voltage set in negative mode to -4500
220 V. Both the nebulizer gas pressure (GS1) and turbo heater gas (GS2) were set to 60 psi. The curtain
221 gas flow was set to 25 l/min. The MS data were collected in MRM scan mode with compound-
222 dependent parameters.

223 **Data handling, power calculations and statistical analysis**

224 The power calculation was based on the primary clinical outcome measure (FMD of the brachial
225 artery) in order to determine the minimum number of participants required for the study. The
226 minimal physiologically significant improvement on FMD measurable is an absolute change of
227 between 1 to 2 % in FMD, considering a baseline vasodilatation of 4-6 %. The sample size was
228 calculated based on the variance of repeated measurement in the control group and on control data.
229 Consequently, with a standard deviation within patients of 2.3% (based on previous studies
230 performed in our group), a significance level of 0.05 and a power of 80%, 18 subjects were required
231 in order to determine a significant within-subject difference between treatments of at least 2.1%.
232 Two-way repeated measured ANOVA was used to analyze all data using GraphPad Prism version 4
233 (GraphPad Software Inc., San Diego, CA, US). Post hoc analysis was conducted using a Bonferroni

234 multiple comparisons test. Significance was defined as $P < 0.05$, with P values represented in the
235 figures as * $P = 0.01-0.05$, ** $P = 0.001-0.01$, *** $P = < 0.001$, **** $P = < 0.0001$. The incremental
236 area under the LDI flux versus time curve (IAUC) was calculated using the trapezoidal method.
237 Prism (GraphPad software, USA) was used to analyze the data. Differences by treatment were
238 identified using a two-way ANOVA with repeated measures. Post hoc analysis was subjected to
239 Bonferroni correction. P values less than 0.05 were treated as significant.

240

241 **RESULTS**

242 **Baseline measurements**

243 All baseline measurements of the subjects were within the normal range (**Table 2**) including blood
244 pressure and FMD. All treatments were well tolerated, with only one participant withdrawing due to
245 an adverse event (gastrointestinal discomfort on both high fibre interventions).

246

247 **Release of free FA from flatbreads**

248 The levels of free, conjugated and bound FA in the white and high fibre flatbreads are presented in
249 Table 1. Enzyme treatment of high fibre bread resulted in the release of more than 60 % of bound
250 FA, with 30g of enzyme-treated high fibre flatbread providing 4.74 mg of *free* FA, compared to
251 0.78 mg in standard high fibre and 0.16 mg in white bread. Three flatbreads were provided per
252 treatment, delivering a 22.21mg of *total* FA after consumption of the enzyme-treated high fibre
253 flatbread, compared to 18.87mg from the non-enzyme treated high fibre flatbread and 1.38mg from
254 the white flatbread control.

255

256 Vascular function

257 A time-dependent increase in FMD was observed following consumption of the enzyme-treated
258 high fibre bread with significantly increased brachial arterial dilation at 2 h ($p < 0.05$), 5 h ($p < 0.01$)
259 and 7 h ($p < 0.05$), relative to the white bread and at 5 h ($p < 0.01$), relative to non-treated high fibre
260 control (n = 19) (**Figure 2**). FMD increased by $0.9 \pm 0.5\%$ at 2 h, $1.5 \pm 1.1\%$ at 5 h and $1.2 \pm 0.9\%$
261 at 7 h for the enzyme-treated high fibre flatbread, relative to baseline. No significant differences in
262 FMD were observed at baseline between groups ($p > 0.05$) or following intake of either the white
263 bread or the non-treated high fibre bread, relative to baseline ($p > 0.05$), although FMD levels were
264 elevated following consumption of non-treated high fibre flatbread at 7 h ($p = 0.098$, n = 19).
265 Blood pressure, heart rate, DVP-SI and DVP-RI were not significantly altered following the
266 consumption of any of the breads (Table 2) or between baseline and post intervention ($p > 0.05$; 2-
267 way ANOVA with Bonferroni post-hoc analysis). Following consumption of the enzyme-treated
268 high fibre bread, there was a rise in endothelium-dependent vasodilation (as indicated by increases
269 in skin erythrocyte flux in the presence of acetylcholine chloride) at 5h and a return to baseline at 7h
270 (**Figure 3 (A)**), although this was not statistically significant. Similarly, no changes were
271 observed in response to SNP (endothelium independent) for any of the treatments (**Figure 3 (B)**).

272

273 Plasma ferulic acid analysis

274 No significant increases in total plasma phenolic acids were observed over the 7 h period (**Figure**
275 **4(A)**). However, a significant increase in plasma ferulic acid was observed following intake of the
276 enzyme-treated high fibre flatbread at 2 h ($p < 0.001$) and 5 h ($p < 0.01$), versus white bread and at 2 h
277 ($p < 0.001$) and 5 h ($p < 0.05$), versus non-treated high fibre control (**Figure 4 (B)**). None of the other
278 individual phenolic acids measured in the study differed between treatments ($p < 0.05$;

279 Supplementary Figure S2).

280

281 **DISCUSSION**

282 In the present study, we show improved acute vascular function following the consumption of the
283 high fibre flatbread which had been treated with an esterase enzyme to release FA bound to
284 arabinoxylan fibre. The release of FA during bread making resulted in a significantly higher level
285 of ferulic acid reaching the circulation (relative to control breads) and resulted in a significant
286 increase in FMD at 2 ($p<0.05$) and 5 h ($p<0.01$) compared to both the high fibre and white flatbread
287 controls (Figure 2). These results are in agreement with a study which showed that intake of wine
288 containing a similar amount of phenolic acids improves endothelial-independent vasodilation
289 acutely in healthy volunteers [33]. Furthermore, with respect to FMD effect size, our data are also
290 in agreement with previous data sets showing increases in brachial artery dilation at 1-2 and 6 h
291 following ingestion of, for example the equivalent of 240g fresh weight of blueberries (containing
292 766-1791 mg of polyphenols) [34], cocoa [35, 36], red wine [37] and decaffeinated coffee [38].

293

294 The increase in FMD observed in response to the enzyme-treated high fibre flatbread was paralleled
295 by an increase in plasma FA. Previous studies have reported free FA concentrations in plasma of
296 between 150-210 nM at 1 to 3 hours post consumption of a high bran cereal [39] and
297 640 μ mol/min/L after consumption of bread [40], with these foods delivering phenolics primarily in
298 the bound-form with no attempt to release FA prior to consumption. There is evidence that FA can
299 be released from arabinoxylan via the action of microbiota-derived esterases and xylanases [21,41-
300 43], although this appears to be of limited efficiency, with only about 25% released by fecal
301 esterases [21,44,45]. Previous work has also shown that processing (fermentation and enzyme
302 treatment) of wheat bran can increase free FA in bread by up to 8-fold [46] and lead to increased
303 release of FA in the colon, and consequently increases in derived metabolites such as 3,4-
304 dihydroxyphenylpropionic acid, 3-hydroxyphenylpropionic acid and 3-phenylpropionic acid. Such
305 bioprocessed bread has been shown to increase the bioavailability of phenolic acids and lead to
306 subsequent immunomodulatory effects in humans [40].

307

308 With respect to mechanism of action, there are very few studies regarding the physiological actions
309 of FA and its metabolites on endothelial function. Animal studies indicate the potential for FA to
310 improve endothelium dysfunction in aortas of rats with spontaneous hypersensitivity [47], and in
311 conjunction with astragaloside IV in streptozotocin-induced diabetic rats [48]. Both studies
312 concluded that FA increased NO bioavailability via a decrease in NADPH-dependent superoxide
313 anion levels [47] and by regulating eNOS activity [48]. In humans, the intake of oats, which is also
314 rich in FA and other phenolics, has been shown to enhance the FMD response in overweight,
315 dyslipidemic adults, particularly in post-menopausal women [49], providing further support for an
316 NO-mediated mechanism. In addition to such a pathway of activity, it has also been suggested that
317 the intake of any cereal food rich in aleurone cells (which have high contents of fibre, phenolics and
318 other potentially beneficial components) may also improve markers of immune function status, such
319 as C-reactive protein and adhesion molecules [50]. Additional support for the causal activity of
320 phenolic acids in vascular benefits, comes from the late improvement in FMD following
321 consumption of the untreated high fibre bread, in the current study, where the excretion of total
322 phenolic acids was highest. Such metabolites include isoferulic acid, hippuric acid and
323 protocatechuic acid, metabolites known to be products of hydroxycinnamic acids, after bacterial
324 metabolism in the large intestine [51-53]. Similarly, small phenolic metabolites derived from
325 anthocyanin and chlorogenic acid were proposed to mediate improvements in FMD at 6 h post-
326 consumption of blueberry flavonoids [33].

327

328 Another potential mechanism may involve an increase in steady-state NO levels, via inhibition of
329 NADPH oxidase. Ferulic acid may inhibit NADPH oxidase, in a similar manner to the structurally
330 related drug apocynin, thus reducing superoxide levels. This reduction in superoxide formation
331 may maintain nitric oxide levels by limiting the reaction of nitric oxide with superoxide (to form
332 peroxynitrite) and consequently sparing nitric oxide bioavailability to the vasculature. Inhibition of

333 NADPH oxidase activity by phenolic metabolites, such as FA and isoferulic acid, has been
334 observed in clinical trials following intake of blueberry polyphenols, which occurred concurrently
335 with an increase in FMD [33] and has been noted in cell studies with flavanols [54,55]. FA, and its
336 metabolites, have structural homology to apocynin, a known NADPH oxidase inhibitor [56], with
337 ferulic, vanillic, homovanillic, and hippuric acids, as well as tyrosol and hydroxytyrosol, all
338 possessing significant inhibitory activity [55, 56] . Further investigation is required to define the
339 detailed mechanism by which wholegrain phenolics exert their activity in humans, however, it
340 appears likely that the interactions of these small phenolics with the endothelium and its local
341 environment may underpin such physiological events.

342 In conclusion, this study has shown that enzymatic processing of high fibre bread can increase the
343 bioavailability of FA and lead to improvements in human vascular function. Our data also suggest
344 that the global influence of wholegrain intake on human cardiovascular health may also relate to the
345 combined influence of both fibre and phenolic acids present within the wholegrain. Further
346 development of innovative baking and bread processing techniques, such as enzymatic treatment of
347 dough to release potential bioactives, may represent a promising approach to improve the bio-
348 accessibility of these beneficial components of wholegrain in a cost-effective way to a wide section
349 of the population.

350

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356 **Conflict of Interest (COI) Statement**

357 The authors have declared no conflict of interest.

358

359 **Authors' Contributions**

360 Study design (PRS, JPES), phenolic analysis (AA), human study implementation (AA), solid phase
361 extraction (AA), mass spectrometry (LVM, AL), statistical analysis (AA, JPES), data interpretation
362 (AA, AL, PRS, JPES), manuscript preparation (AA, AL, PRS, JPES), manuscript approval (AA,
363 LVM, AL, PRS, JPES).

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Table 1.
Baseline fasting characteristics of study population.¹

Characteristic	Value
Age (yrs)	26.6 ± 6.4
Height (m)	1.8 ± 0.1
Weight (kg)	76.6 ± 11.7
BMI (kg/m ²)	24.2 ± 2.4
Waist circumference (cm)	81.9 ± 5.7
BP (mmHg)	
Systolic	121.2 ± 10.8
Diastolic	70.3 ± 8.0
Total cholesterol (mmol/l)	4.4 ± 0.8
Glucose (mmol/l)	5.3 ± 0.4
TAG (mmol/l)	0.8 ± 0.3
Brachial artery diameter (mm)	4.1 ± 0.5
FMD %	4.8 ± 1.6

¹ Values are means ± SD. (n=19).

Table 2.Compositional analysis of white, non-treated high fibre and enzyme-treated high fibre flatbreads.¹

Compounds	Composition		
	White	Non-Treated High Fibre	Enzyme-treated High Fibre
White flour (%)	60.5	0.0	0
High Fibre flour (%)	0.0	60.5	60.5
Water (%)	34.8	34.8	34.8
Ultraflo L ® activated (%)	0.0	0.0	3.9
Ultraflo L ® de-activated (%)	3.9	3.9	0
Salt (%)	0.8	0.8	0.8
Energy (k joules) (kj/100g)	1169.0	1193.0	1247.0
Energy (k calories) (kcal/100g)	276.0	282.0	294.0
Protein (g/100g)	7.70	8.90	9.20
Total Carbohydrate (g/100g)	60.60	61.10	63.00
Total sugars (g/100g)	3.30	4.00	3.70
Fat (g/100g)	0.80	1.20	1.30
Saturates (g/100g)	0.14	0.24	0.26
Monounsaturates (g/100g)	1.23	0.24	0.18
Polyunsaturates (g/100g)	0.39	0.67	0.80
Trans fatty acids (g/100g)	<0.1	<0.1	<0.1
Total fibre (g/100g)	2.40	4.60	3.20
Sodium (mg/100g)	315.00	320.00	315.00
Moisture content (g/100g)	29.70	27.30	25.00
Ash content (g/100g)	1.18	1.48	1.54
Free FA (mg/100g)	0.53	2.60	15.80
Conjugated FA (mg/100g)	0.01	1.47	0.33
Bound FA (mg/100g)	1.00	16.90	9.43
Total FA (mg/100g)	1.54	20.97	25.56

364 ¹ Values based on freeze dried flatbread.

365

Table 3. Acute effects of interventions on blood pressure, heart rate and DVP.¹

	Baseline	2h	5h	7h
Systolic blood pressure				
(mm Hg)				
White	124 ± 2	121 ± 2	120 ± 3	121 ± 2
Non-Treated High Fibre	121 ± 2	120 ± 2	120 ± 2	121 ± 2
Enzyme-Treated High Fibre	121 ± 2	121 ± 2	120 ± 2	120 ± 2
Diastolic blood pressure				
(mm Hg)				
White	71 ± 1	68 ± 1	67 ± 1	72 ± 2
Non-Treated High Fibre	73 ± 2	68 ± 1	68 ± 2	71 ± 2
Enzyme-Treated High Fibre	71 ± 1	69 ± 1	67 ± 1	71 ± 1
Heart rate (beats/min)				
White	56 ± 2	58 ± 2	59 ± 2	57 ± 2
Non-Treated High Fibre	59 ± 2	60 ± 2	62 ± 2	59 ± 2
Enzyme-Treated High Fibre	59 ± 2	61 ± 2	60 ± 2	60 ± 2
DVP-SI (m/s)				
White	5.14 ± 0.8	5.3 ± 0.1	5.2 ± 0.1	5.6 ± 0.4
Non-Treated High Fibre	5.2 ± 0.3	5.5 ± 0.4	5.5 ± 0.4	5.4 ± 0.5
Enzyme-Treated High Fibre	5.5 ± 0.3	5.4 ± 0.4	5.5 ± 0.4	5.4 ± 0.4
DVP-RI (%)				
White	67 ± 12	66.6 ± 13	60.7 ± 13	62.1 ± 10
Non-Treated High Fibre	64.2 ± 15	63.9 ± 10	60.5 ± 11	60.5 ± 7
Enzyme-Treated High Fibre	66.2 ± 12	62.7 ± 14	62.1 ± 14	61 ± 12

¹ Values are means ± SEM. (n=19).

368 **Figure Legends.**

369 **Figure 1.** Consort Diagram. Flow diagram of the progress through the phases of the randomised
370 clinical trial indicating participant numbers at enrolment, intervention allocation, follow-up, and
371 data analysis.

372

373 **Figure 2.** Time dependent changes in FMD (n=19, mean \pm SD) following consumption of enzyme-
374 treated high fibre (\blacktriangle), non-treated high fibre (\blacksquare) and white bread (\bullet). Data were analyzed using 2-
375 way ANOVA with post-hoc analysis conducted by using a Bonferroni multiple-comparisons test.
376 Levels of significance between samples are indicated as follows: enzyme-treated high fibre
377 flatbread versus white flatbread; 2 h (* $p < 0.05$), 5 h (** $p < 0.01$) and 7 h (* $p < 0.05$) and enzyme-
378 treated high fibre flatbread vs non-treated high fibre flatbread 5 h (b: $p < 0.01$).

379

380 **Figure 3.** Response of forearm skin erythrocyte flux following the iontophoresis of (A)
381 acetylcholine chloride and (B) sodium nitroprusside (n=19) after consumption of white (\bullet), non-
382 treated high fibre (\blacksquare) and enzyme-treated high fibre (\blacktriangle) flatbread. Values are means \pm SEM.

383

384 **Figure 4.** (A). Total phenolic acids, in plasma at baseline, 2, 5 and 7 h following consumption of
385 white (\bullet), non-treated high fibre (\blacksquare) and enzyme-treated high fibre (\blacktriangle) flatbread (n=18; mean \pm
386 SEM). (B). Concentration of free FA in plasma at baseline, 2, 5 and 7 h following consumption of
387 white (\bullet), non-treated high fibre (\blacksquare) and enzyme-treated high fibre (\blacktriangle) flatbread (n=18; mean \pm
388 SEM). Data were measured using 2-factor repeated measures ANOVA with time and treatment as
389 the 2 factors. Post hoc analysis was conducted using a Bonferroni multiple comparisons test. a:
390 white vs enzyme-treated high fibre, $p < 0.001$; b: white vs enzyme-treated high fibre, $p < 0.01$; c: non-
391 treated high fibre vs enzyme treated high fibre, $p < 0.001$; d: non-treated high fibre vs enzyme-
392 treated high fibre, $p < 0.5$.