

Title: Identifying chondroprotective diet-derived bioactives and investigating their synergism

¹Rose K Davidson ^{**} Ph.D & ¹Jonathan Green ^{**} Ph.D, ¹Sarah Gardner BSc, ²Yongping Bao Ph.D, ²Aedin Cassidy Ph.D, ¹Ian M Clark ^{*} Ph.D.

Biomedical Research Centre: ¹School of Biological Sciences, ²Norwich Medical School, Department of Nutrition, Norwich Research Park, University of East Anglia, Norfolk, UK, NR4 7TJ.

*Corresponding author

**These authors contributed equally to the work

This work was funded by the BBSRC Diet and Health Research Industry Club grants BB/J500112/1, BB/L025485/1.

*Prof. Ian M Clark,

School of Biological Sciences,

University of East Anglia,

Norwich Research Park,

Norwich,

Norfolk, UK

NR4 7TJ

Email: i.clark@uea.ac.uk.

Tel: +44 (0) 1603 592760

Fax: +44 (0) 1603 591750

Osteoarthritis (OA) is a multifactorial disease and nutrition is a modifiable factor that may contribute to disease onset or progression. A detailed understanding of mechanisms through which diet-derived bioactive molecules function and interact in OA is needed.

We profiled 96 diet-derived, mainly plant-based bioactives using an *in vitro* model in chondrocytes, selecting four candidates for further study. We aimed to determine synergistic interactions between bioactives that affected the expression of key genes in OA.

Selected bioactives, sulforaphane, apigenin, isoliquiritigenin and luteolin, inhibited one or more interleukin-1-induced metalloproteinases implicated in OA (*MMP1*, *MMP13*, *ADAMTS4*, *ADAMTS5*). Isoliquiritigenin and luteolin showed reactive oxygen species scavenging activity in chondrocytes whereas sulforaphane had no effect and apigenin showed only a weak trend. Sulforaphane inhibited the IL-1/NFκB and Wnt3a/TCF/Lef pathways and increased TGFβ/Smad2/3 and BMP6/Smad1/5/8 signalling. Apigenin showed potent inhibition of the IL-1/NFκB and TGFβ/Smad2/3 pathways, whereas luteolin showed only weak inhibition of the IL-1/NFκB pathway. All four bioactives inhibited cytokine-induced aggrecan loss from cartilage tissue explants. The combination of sulforaphane and isoliquiritigenin was synergistic for inhibiting *MMP13* gene expression in chondrocytes.

We conclude that dietary-derived bioactives may be important modulators of cartilage homeostasis and synergistic relationships between bioactives may have an anti-inflammatory and chondroprotective role.

Introduction

Osteoarthritis (OA) is a global problem and a leading cause of disability^{1,2}. OA is the most common type of arthritis affecting 13.2% of males and 19.8% of females in the UK³. There are no effective disease-modifying drugs for OA, with pharmacological treatment restricted to pain relief. Treatment of chronic conditions such as osteoarthritis (OA) is further complicated by co-morbidities as the risk escalates with age⁴. The importance of the health, economic and societal issues surrounding OA was acknowledged by the General Assembly of the United Nations and World Health Organisation^{5,6}

OA is a disease affecting the entire di-artrodial joint where degeneration of the articular cartilage, changes in the underlying (subchondral) bone and synovial inflammation are typically seen. This disease is a major cause of disability, results in a loss of mobility in the affected joint and unmanaged pain symptoms detrimental to quality of life⁷. Metalloproteinases commonly linked to the destruction of articular cartilage are the collagenase MMP-13 and aggrecanases ADAMTS-4 and ADAMTS-5^{8,9}.

A change in the typical approach to chronic disease management may benefit patients¹⁰. The consequence for conditions of slow progression such as osteoarthritis is a late presentation of symptoms, when joint damage has already occurred. This may in part account for the lack of effective disease-modifying interventions for OA. With age as a primary risk factor for OA, the entire population can be considered as 'at risk' for disease. Dietary intervention as a strategy to reduce risk at the population level provides a low risk option with potentially wide-reaching benefits¹¹.

Available epidemiological data suggest that dietary constituents, specifically carotenoids, n3-polyunsaturated fatty acids (PUFA), vitamin K and allium intake are associated with a reduction in the progression of OA in man¹². A number of other plant-derived phytochemicals outside of nutrients themselves, have been proposed to have benefits on joint health and OA e.g.^{13,14} but translation to man has been slow. Detailed molecular studies both *in vitro* and *in vivo* are required in order to establish modes of action, which are different from traditional single target pharmaceuticals, and to optimise clinical trial design in order to gain optimal evidence of efficacy in both disease treatment and prevention.

Previously, our research in this area has focused on sulforaphane, an isothiocyanate derived from the glucosinolate precursor glucoraphanin, which is abundant in cruciferous vegetables, particularly broccoli. It is widely known for its cytoprotective, anti-inflammatory, anti-microbial and anti-tumourigenic activities. There are a number of reports on the potential for SFN in OA describing protection of chondrocytes from cell death, repression of catabolic metalloproteases, intra-articular administration using microspheres to protect cartilage, protection of subchondral bone and cartilage protection in mouse models of OA¹⁵⁻¹⁹.

We sought to extend knowledge in this area by screening a library of diet-derived bioactives for inhibition of either basal or interleukin-1-induced expression of matrix metalloproteinase-13 (MMP13) in two chondrocyte-like cell lines as a surrogate for chondroprotection. The top 'hits' from this screen were further tested in primary human articular chondrocytes which led us to focus on four bioactives for possible synergism (sulforaphane (SFN), apigenin (API), isoliquiritigenin (ISO) and luteolin (LUT)). These compounds were then examined for mechanism of action but also their interaction.

Flavonoids are polyphenolic compounds common to a wide range of plant foods. Flavones are a flavonoid sub-class to which apigenin and luteolin belong. Apigenin is found abundantly in bell peppers, garlic, belimbi fruit, guava, celery, parsley and chamomile. Sources of luteolin are bird's

eye chilli, onions, carrots and olive oil^{20,21}. Luteolin is reported to inhibit aggrecanases ADAMTS4 and ADAMTS5 in ATDC5 cells and murine cartilage explants²² and apigenin is a hyaluronidase inhibitor as demonstrated by *in vitro* turbidity assay and in bovine cartilage explants^{23,24}, and is reported to inhibit metalloproteinase expression in primary rat articular chondrocytes and the human SW1353 chondrosarcoma cell line^{25,26}.

Isoliquiritigenin is also a flavonoid found in liquorice root, though has a chalcone structure. Chalcone derivatives are shown to have wide ranging bioactivities such as anti-inflammatory, anti-oxidant, anti-microbial activity²⁷. Isoliquiritigenin is reported to protect subchondral bone in a mouse model of OA²⁸, repress interleukin-1-induced matrix metalloproteinases in rat primary chondrocytes²⁹, protect cartilage in the anterior cruciate ligament transection (ACLT) mouse model of OA, and inhibit the inflammatory nuclear factor kappa B (NFκB) pathway in a rat model of intracerebral haemorrhage and ATDC5 cells^{30,31}, also affecting the NLRP3 inflammasome and inducing NRF2 signalling³⁰.

Results

Key matrix metalloproteinases in osteoarthritis: diet-derived bioactives inhibit gene expression.

In order to identify compounds which were potentially chondroprotective, ninety six diet-derived bioactives were screened at 10 μM against basal and IL-1-induced *MMP13* expression (measured by RT-qPCR) in SW1353 human chondrosarcoma and C28/I2 immortalised human costal chondrocytes. Each screen was performed once *n*=1, with three replicates of each condition, with the 10 μM dose chosen pragmatically in order to reveal activity.

The heat-map (Supplementary data S1) shows distinct areas of high and low *MMP13* expression for IL-1 treated SW1353 cells. Two clades (highlighted) of low *MMP13* expression across the two cell lines were noted to contain apigenin, isoliquiritigenin and luteolin.

From these initial screens, ten bioactives were selected and with sulforaphane (identified in our previous research¹⁶) they were tested for inhibition of *MMP13* expression, but also other cartilage matrix-degrading proteases *MMP1*, *ADAMTS4* and *ADAMTS5*. Human primary chondrocytes (HACs) isolated from three separate patients *n*=3, were treated with 10 μM genistin, luteolin, polydatin, curcumin, apigenin, myricetin, isoliquiritigenin, ursolic acid, epigallocatechin gallate, naringin or sulforaphane in triplicate for 6 hours, or pre-treated with 10 μM of each bioactive for 30 mins then 5ng/ml IL-1 for 6 hours. Levels of protease gene expression are shown in Table 1 (IL-1-induced) and Table 2 (basal).

IL1-induced *MMP1*, *MMP13*, *ADAMTS4* and *ADAMTS5* mRNA expression was consistently decreased by luteolin but did not reach statistical significance. Apigenin significantly decreased IL-1-induced *MMP13*, *ADAMTS4* and *ADAMTS5* expression (mean fold change and 95% CI: 0.18 (0.09 to 1.55), *p*=0.005, 0.11 (0.20 to 1.58), *p*=0.0001 and 0.22 (-0.06 to 1.62), *p*=0.018 respectively). Isoliquiritigenin significantly decreased the expression of IL-1-induced *MMP13*, *ADAMTS4* and *ADAMTS5* (mean fold change and 95% CI: 0.14 (0.14 to 1.59), *p*=0.0005, 0.08 (0.15 to 1.69), *p*=0.0001, and 0.11 (0.05 to 1.73), *p*=0.001 respectively). Sulforaphane significantly decreased IL-1-induced expression of *MMP13* and *ADAMTS4* (mean fold change and 95% CI: 0.12 (0.15 to 1.61), *p*=0.001, 0.03 (0.28 to 1.66), *p*=0.0001), while showing a trend to decreasing both IL-1-induced *MMP1* and *ADAMTS5* (Table 1). Ursolic acid increased basal *MMP1* expression mean fold change 4.34 (95% CI: -5.87 to -0.82 *p*=0.004) and sulforaphane inhibited basal *ADAMTS5* expression mean fold change 0.25 (95% CI: 0.23 to 1.27, *p*=0.001) (Table 2). Three bioactives apigenin (API),

isoliquiritigenin (ISO) and luteolin (LUT) were taken forwards, along with sulforaphane (SFN), as having activity regulating the expression of four key genes relevant to OA in HACs.

We established a dose response for each bioactive against IL-1-induced *MMP13* expression in HACs using three separate patient cell isolates $n=3$, each in triplicate conditions (Figure 1). SFN reached significance at 12 μM and 24 μM mean fold change and 95% CI: 0.11 (0.53 to 1.26) $p=0.0001$, and 0.05 (0.57 to 1.33) $p=0.0001$ respectively (Figure 1a). API reached significance at 8 μM mean fold change and 95%CI: 0.35 (0.36 to 0.95) $p=0.037$, 16 μM 0.07 (0.65 to 1.22) $p=0.0001$ and 32 μM 0.03 (0.68 to 1.25) $p=0.0001$ (Figure 1b). ISO was significant at 10 μM mean fold change and 95%CI: 0.30 (0.35 to 1.06) $p=0.002$ and 20 μM 0.06 (0.59 to 1.30) $p=0.0001$ (Figure 1c). LUT was significant at 20 μM mean fold change and 95%CI: 0.11 (0.60 to 1.17) $p=0.0001$ and 40 μM 0.04 (0.67 to 1.25) $p=0.0001$ (Figure 1d).

Hence, a wide screen of dietary-derived bioactives led us to focus on a subset of 10 compounds with further analyses highlighting 4 compounds, which dose-dependently inhibit IL-1-induced *MMP13* expression in primary human articular chondrocytes.

Bioactives affect reactive oxygen species (ROS) in primary human chondrocytes

In order to investigate potential mechanism of action, direct antioxidant activity of the bioactives was measured in primary human articular chondrocytes (three separate patient cell isolates $n=3$, each in triplicate). Primary chondrocytes were pre-treated for 30 minutes with the bioactives and then 100 μM tert-butyl hydrogen peroxide (tBHP) was added to induce cellular reactive oxygen species (ROS).

ISO and LUT showed a dose dependent inhibition of tBHP-induced ROS that was statistically significant (Figure 2). ISO inhibited ROS production: (3 μM mean fold change and 95% CI: 0.48 (0.26 to 0.78), $p=0.030$, 6 μM 0.27 (0.47 to 0.99), $p=0.0005$) and 12 μM 0.12 (0.62 to 1.14), $p=0.0001$ (Figure 2c). LUT inhibited ROS production: (1.5 μM mean fold change and 95% CI: 0.33 (0.41 to 0.94), $p=0.0035$, 3 μM 0.10 (0.64 to 1.16), $p=0.0001$, 6 μM 0.04 (0.70 to 1.22), $p=0.0001$ and 12 μM 0.03 (0.71 to 1.24), $p=0.0001$) (Figure 2d). No effect was seen for SFN or API (Figure 2a & 2b).

This showed that both ISO and LUT exert a direct antioxidant effect in chondrocytes, in contrast to SFN and API.

Transcriptional signalling pathway analysis

A number of signalling pathways play a role in cartilage homeostasis and OA. Luciferase reporters for IL-1/nuclear factor-kappa B, TGF- β /Smad2/3, Wnt3A/TCF/Lef, and BMP6/Smad1/5/8 signalling pathways were used to measure transcriptional activity in response to SFN, API, ISO and LUT treatment (Figure 3). Data are from a minimum of three independent experiments ($n=3$) each performed in triplicate.

SFN dose dependently inhibited IL-1-induced NF κ B signalling reaching significance at 12 μM mean fold change and 95% CI: 0.11 (0.8 to 1.7), $p=0.003$ (Figure 3a) and Wnt3A/TCF/Lef signalling (not significant) (Figure 3d). SFN showed trends of increasing TGF- β -induced Smad2/3 and BMP6-induced Smad1/5/8 signalling (not significant) (Figure 3b & 3c).

API strongly inhibited IL-1/NFκB (1.5-12 μM mean fold change and 95%CI: 0.22 (0.30 to 1.26), p=0.009, 3 μM 0.09 (0.43 to 1.40), p=0.0001, 6 μM 0.04 (0.48 to 1.44), p=0.0001, 12 μM 0.03 (0.45 to 1.49), p=0.0001) (Figure 3e) and TGF-β/Smad2/3 signalling at 1.5-12 μM (1.5 μM mean fold change and 95% CI: 0.06 (0.22 to 1.67), p=0.002, 3 μM 0.02 (0.26 to 1.70), p=0.0008, 6 μM 0.01 (0.27 to 1.71) p=0.0001, 12 μM 0.03 (0.19 to 1.76), p=0.001) (Figure 3f). In contrast API had no effect on BMP6/Smad1/5/8 (Figure 3g) and decreased Wnt3A/TCF/Lef activity weakly in a dose dependent manner (not significant) (Figure 3h).

We observed a small decrease in BMP6-induced Smad1/5/8 signalling at 12 μM (not significant) with ISO treatment (Figure 3k) but no effect on the IL-1/NFκB, TGF-β/Smad2/3 or Wnt3A/TCF/Lef reporters (Figure 3i, 3j & 3l).

LUT showed a weak dose dependent inhibition of IL-1/NFκB signalling (not significant) (Figure 3m) and small decreases at the high concentrations for TGF-β/Smad2/3 (Figure 3n), BMP6/Smad1/5/8 (Figure 3o) and Wnt3A/TCF/Lef (Figure 3p) reporters (not significant).

Thus, SFN and API both inhibit NFκB signalling, API also inhibits Smad2/3 signalling. SFN also shows weak induction of Smad2/3 and Smad1/5/8 signalling and repression of canonical Wnt signalling. There was minimal impact of the four compounds on other signalling pathways tested.

Bioactives reduce cartilage tissue degradation *in vitro*

In order to examine the impact of bioactivity on cartilage homeostasis, bovine nasal cartilage explants were induced to degrade using inflammatory cytokines IL-1 and OSM in three independent experiments (n=3) each performed in triplicate. Cartilage was treated with 0-24 μM SFN, API, ISO or LUT ± IL-1 and OSM and the degradation product of aggrecan (glycosaminoglycan) was measured using 1,9-dimethyl-methylene blue assay (Figure 4). All four bioactives showed dose-dependent patterns inhibiting cartilage degradation: IL1/OSM treatment mean % GAG loss and 95% CI: 57.74 (41.1 to 61.11). SFN (24 μM mean % loss and 95% CI: 28.13 (12.06 to 47.16), p=0.0019) and significant linear trend (slope = -0.07, p=0.0009, r² = 0.78) (Figure 4a). API (24 μM mean % loss and 95% CI: 29.45 (5.99 to 50.6), p=0.022) with significant linear trend (slope = -0.08, p=0.0043, r² = 0.90) (Figure 4b). ISO (24 μM mean % loss and 95% CI: 40.37 (1.40 to 33.34), p=0.049) and significant linear trend (slope = -0.04, p=0.013, r² = 0.88) (Figure 4c). Similarly, LUT (24 μM mean % loss and 95% CI: 32.25 (-5.96 to 56.94), p=0.080), and showed a significant linear trend (slope = -0.07, p=0.022, r² = 0.64) (Figure 4d).

All compounds show a statistically significant dose-response (i.e significant decrease across the dose range even though individual lower concentrations are not significant on their own). SFN, API and ISO show statistical significance in preventing IL1/OSM-induced GAG release from cartilage explants at the highest dose tested.

Kinase signalling pathway analysis

It is likely that compounds alter intracellular kinase signalling, therefore we used kinase arrays to identify potential regulation of this in response to IL-1 stimulation (5ng/ml, 10 mins) by SFN or ISO (10 μM) in human primary chondrocytes. SFN inhibited IL-1 phosphorylation of Akt1/2/3 (s473), c-jun, CREB, ERK1/2, PRAS40, WNK1 (all >25% decrease). ISO inhibited Akt1/2/3 (s473), β-catenin,

EGF-R (>25% decrease). SFN reduced basal levels of phosphorylation (>50%) compared to control of c-jun, CREB, ERK1/2, HSP27, JNK1/2/3, p27 and p38 α $n=1$ (Supplementary data S2).

Bioactive synergism for inhibition of *MMP13* expression in primary chondrocytes

Since foods are rarely eaten alone, the synergism between compounds with respect to IL1-induced *MMP13* expression was determined. All synergy experiments used cell isolates from at least three separate patients $n=3$, each performed in duplicate. The dose-effect relationship parameters of each single bioactive SFN, API, ISO and LUT against IL1-induced *MMP13* mRNA expression in HACs were determined. The rank order of potency based on the IC₅₀ values were: ISO>SFN>LUT>API in HACs (Table 3).

The combination index (CI) values of bioactive combinations against IL1-induced *MMP13* mRNA expression in HACs was measured. Combination ratios were based on IC₅₀ values rather than absolute concentrations. Each component had the equipotent combination ratio of: SFN:API:ISO:LUT (S:A:I:L) = 0.75:1:0.625:1.25 μ M. Synergism was only seen in the SFN/ISO (SI) combination (mean CI: 0.724). Nearly additive combinations were SA (mean CI: 1.02) and AI (mean CI: 1.06) where SA \geq AI. All three (and four) bioactive combinations were antagonistic. Rank order for antagonism was SAIL \geq IL \geq LSA \geq ILS \geq AIL \geq SL>SAI>AL (Table 4). Note assigned symbols for the combination indices are taken from Chou³².

The dose reduction index (DRI) values and relationship to CI values in synergistic combinations were determined (Table 5). The two-bioactive combination SFN + ISO show a robust dose reduction from ED₅₀ to ED₉₅. This favourable dose reduction was also seen for SFN +API at ED₅₀ to ED₇₅ and API + ISO at ED₉₀ to ED₉₅ even though the average unweighted CI values suggest a nearly additive effect (Table 4).

A polygonogram summary of two-bioactive combination interactions, for the four bioactives is shown (Supplementary data S3). The combination of SFN and ISO show synergistic activity for reducing IL-1 induced *MMP13* expression in primary chondrocytes at each effect level. The average combination indices for SFN/API and API/ISO were 'nearly additive' however, it is noted that the SFN/API combination shows slight to moderate synergism at lower effect levels.

SFN/ISO combination regulates the expression of other key genes in OA.

In order to investigate the potential of the SFN/ISO combination to interact synergistically to regulate other relevant genes, we used RT-qPCR to measure the expression of proteases *ADAMTS4*, *ADAMTS5*, the Nrf2-induced indirect antioxidant *HO-1*, and the early NF κ B response gene and regulator *A20*. We used the synergism sample set at equipotent IC₅₀ ratios of 0.75:0.625 μ M where $n=3$ (figure 5).

IL-1-induced *ADAMTS4* was inhibited by SFN and ISO singly (not significant) and in combination (SI (1.5:1.25 μ M) mean fold change 1.27 (95% CI: -1.39 to 0.84), $p=0.99$), (SI (3:2.5 μ M) mean fold change 0.16 (95% CI: -0.27 to 1.96), $p=0.014$), (SI (6:5 μ M) mean fold change 0.005 (95% CI -0.12 to 2.11), $p=0.034$) (Figure 5a).

IL-1-induced *ADAMTS5* was inhibited by SFN and ISO singly (not significant) and in combination (SI (1.5:1.25 μ M) mean fold change 0.57 (95% CI: -0.26 to 1.40), $p=0.73$), (SI (3:2.5 μ M) mean fold change 0.71 (95% CI: -0.12 to 1.54), $p=0.007$), (SI (6:5 μ M) mean fold change 0.09 (95% CI: 0.08 to 1.74), $p=0.0001$) (Figure 5b).

HO-1 was induced by SFN singly and in combination with ISO compared to control (SFN 1.5 μ M mean fold change 6.59 (95% CI: -16.31 to 7.58), $p=0.025$), (3 μ M mean fold change 7.44 (95% CI: -17.17 to 6.72), $p=0.022$), (6 μ M mean fold change 8.96 (95% CI: -18.69 to 5.20), $p=0.012$). ISO alone did not induce *HO-1* expression. In combination, (SI (1.5:1.25 μ M) mean fold change 13.74 (95% CI -23.47 to 0.42), $p=0.0018$), (SI (3:2.5 μ M) mean fold change 22.5 (95% CI -32.22 to -8.33), $p=0.0001$), (SI (6:5 μ M) mean fold change 23.86 (95% CI: -33.59 to -9.70), $p=0.0001$) (Figure 5c).

IL-1-induced *A20* was inhibited by SFN singly (not significant) and in combination with ISO (SI (1.5:2.5 μ M) mean fold change 1.02 (95% CI: -1.68 to 1.63), $p=0.997$), (SI (3:2.5 μ M) mean fold change 0.23 (95% CI: -0.89 to 2.42), $p=0.16$), (SI (6:5 μ M) mean fold change 0.04 (95% CI: -0.69 to 2.62), $p=0.004$) (Figure 5d).

Whilst mathematical synergism for these genes wasn't determined as for *MMP13* above, the combination of SFN and ISO together was clearly more effective than either compound alone, confirming the utility of the approach.

Discussion

A strategy to prevent or treat OA using dietary modification, supplementation or the development of functional foods is clearly attractive. It also has wide public health implications particularly in a chronic disease which is increasing in prevalence with no disease-modifying treatments currently available. There is a need to understand the mechanism by which diet-derived bioactives can impact upon the joint, both in vitro and in vivo. This is vital for the design of appropriate intervention trials and therefore the translation to treatment in man.

Our strategy was to screen diet-derived compounds for their ability to inhibit either basal or IL1-induced *MMP13* expression since this identified SFN, which was then shown to be chondroprotective in animal models of OA¹⁶. Further we hypothesised that if bioactives regulate gene expression through differing mechanisms, then they are more likely to be synergistic. Of the 96 bioactives in our initial screen for expression of key OA metalloproteinase genes, the four most efficacious compounds were SFN, API, ISO and LUT.

The effect of SFN on metalloproteinase expression and cartilage degradation was in line with earlier work^{16,33,34} and unsurprisingly, had no direct effect on ROS production since it is well documented by others that SFN is an indirect antioxidant³⁵. SFN affected each of the four cytokine-stimulated pathways; IL-1, TGF β , BMP6 and Wnt3A with differing efficacy. Of the four pathways analysed, SFN interacted most strongly with the IL-1/NF κ B pathway, likely in part as a function of SFN's known indirect antioxidant and anti-inflammatory activities, which we have reported previously¹⁶. SFN also inhibited Wnt3a canonical signalling, an observation seen previously in breast cancer stem cells³⁶ but diverged to enhance both TGF β /Smad signalling, (which is reported similarly in colorectal cancer cells though not in a murine model of muscle fibrosis^{37,38}), and BMP6 signalling. Moreover, BMP6/Smad1/5/8 pathway is reported to induce *HO-1* expression^{39,40} and so SFN could be viewed overall as a dietary modulator of cartilage homeostasis. This is reinforced by its efficacy in animal models of OA^{16,17,19}. Since SFN is a well-known activator of the NRF2 signalling pathway⁴¹ and though not measured here, this may be a significant contributing mechanism of action.

Our data support previous studies where API inhibited IL-1-induced metalloproteinase expression^{25,26}, and is reflected by the decrease in glycosaminoglycan loss also seen by Lim et al²⁶ in rabbit articular cartilage. API showed remarkably strong and targeted activity for inhibiting IL-1/NF κ B and

TGF- β /Smad2/3 signalling pathways while having no effect on BMP6/Smad1/5/8 signalling or only a weak inhibitory effect on the Wnt3A/TCF/Lef pathway. Inhibition of NF κ B signalling by API agrees with some earlier studies^{42,43} (though not Lim et al²⁶), as does the inhibition of TGF β /Smad signalling in prostate cancer cell lines⁴⁴. Conversely, Zhang et al demonstrate apigenin activation of the TGF β /Smad pathway in skin fibroblasts to induce collagen synthesis, albeit at a later time point (12 hours)⁴⁵. Similarly for SFN, we did not observe direct antioxidant activity for API in primary chondrocytes though the literature describes varied outcomes and mechanisms for API including antioxidant activity that may be cell type dependent. Yokomizo & Moriwaki describe a lack of radical scavenging activity for API in Caco-2 cells⁴⁶, and API is described as a driver of ROS production⁴⁷, which we did not observe in chondrocytes (data not shown). Conversely, Crasci et al describe free radical scavenging activity for both API and LUT in chondrocytes⁴⁸.

Our data on both metalloproteinase expression and explant cartilage degradation support a recent study demonstrating *in vivo*, the potential for ISO (administered by intraperitoneal injection) to protect articular cartilage, where the authors also show protection of subchondral bone tissue in the joint of mice²⁸. This followed another recent report of ISO inhibition of IL-1-induced metalloproteinase expression in rats²⁹. Interestingly and in contrast to SFN and API, ISO showed a consistent dose dependent inhibition of tBHP-induced ROS production indicating some direct antioxidant activity. We did not find evidence to support an interaction of ISO with any of the reporters tested and suggest an interaction with Nrf2 to be more likely^{30,49,50}.

LUT showed a similar activity profile to ISO overall despite being structurally more similar to API (one hydroxyl group difference in the B ring). ISO inhibition of tBHP-induced ROS production was striking in relation to other activity measures and supports an earlier report in Caco-2 cells⁴⁶, where no effect by API was also seen. There are a number of reports of LUT that describe reactive oxygen and nitrogen species scavenging, transition metal chelation and antioxidant enzyme induction and is reviewed in detail elsewhere²¹. Nonetheless, we observed no marked interaction with the pathway reporters excepting the weak interaction with the IL-1/NF κ B reporter. Despite the repression of metalloproteinase expression by LUT (also seen by others^{22,51}) that was borne out in our cartilage degradation assay, it was surprising to observe an antagonistic relationship for reducing *MMP13* expression between LUT and each of the remaining bioactives.

It is possible that the anti-inflammatory/cytoprotective activities of SFN and antioxidant activities of ISO culminate in the synergistic interaction to reverse IL-1-induced *MMP13* expression that we have identified. The nearly additive relationship of SFN and API could be explained by their similar activity profiles, which may saturate signalling pathways, whilst neither having direct antioxidant activities in our hands. However, this may be simplistic since the interaction profile of SFN/API with *MMP13* expression differs substantially at different effect levels, being mild to moderately synergistic at lower effect levels. A high effect level, though clinically desirable, is not a necessity in terms of OA prevention or treatment.

Our data suggest antagonistic interactions (i.e. the effect of both compounds is less than the additive effect of the individual compounds) between SFN/LUT and ISO/LUT, which are more difficult to explain. Interestingly all combinations with 3 or more bioactives were antagonistic, except for the SFN/API/ISO combination, which was only moderately antagonistic, possibly negated by the SFN/ISO interaction. More broadly speaking, food matrices or food-drug interactions present a far more complicated milieu.

We also assayed the Nrf2-induced HO-1 gene and the NFκB-induced A20 gene, both readouts of key pathways in OA, where treatment with both SFN and ISO showed a greater effect than the additive effects of the single compounds.

We have clearly identified dietary-bioactives with the strong potential to be chondroprotective *in vivo* and with SFN and ISO displaying synergy. The concentrations of bioactives in our *in vitro* assays are either equivalent or higher than reported plasma concentrations after consumption (e.g. ⁵²⁻⁵⁴). However, compounds or their metabolites can accumulate with repeated consumption, or be concentrated intracellularly⁵⁵. The potential for functional foods or supplementation also increases the achievable concentration *in vivo*. It is worth noting that dietary compounds undergo extensive metabolism following ingestion and even when metabolism is well established, the mix of parent compound and metabolites that act upon the articular joint is unknown. It is also likely that the metabolites produced upon ingestion may be more active than the parent compound ⁵⁶. The only way to understand this is to perform feeding experiments in animal models of disease using chemically-defined foods as we have done for SFN alone¹⁶. *In vivo* experiments also enable a comparison of mechanism-of-action with the *in vitro* and *ex vivo* analyses. This is the direction of our future research in this area.

We conclude that diet-derived bioactives may be important modulators of cartilage homeostasis and synergistic relationships between bioactives (potentially SFN/ISO) in primary chondrocytes may have a role in chondroprotection and counterbalancing inflammation.

Materials and Methods

Materials

SFN was from Toronto Research chemicals, CAN. All other bioactives were obtained from Stratech, UK. All cytokines were obtained from Firstlink Ltd (UK) except Wingless-Type MMTV Integration Site Family 3A (WNT3A), which was obtained from R&D Systems. The Proteome Profiler Array (cat: ARY003B) was obtained from R&D Systems, UK.

Methods

A flow diagram to summarise the experimental process is shown in Supplementary Information.

Cell culture and treatments

The human chondrosarcoma SW1353 cell line was purchased from ATCC and the C28/I2 cell line was a kind gift from Mary Goldring, HSS Research Institute (USA). Primary chondrocytes were obtained as described previously¹⁶. All experimental protocols were approved by NRES Committee East of England (ref: 08/h0304/85+5). All tissue donors gave informed written consent. All methods were performed in accordance with relevant laboratory guidelines and institutional regulations. All treatments included 30 min pre-incubation with bioactive compound followed by cytokine treatment. Cells were propagated in Dulbecco's modified Eagles medium (DMEM, GlutaMAX) with 10% v/v foetal calf serum and 1000IU/penicillin, 100µg/ml streptomycin, at 37°C, 5% CO₂.

Complementary DNA (cDNA) synthesis and quantitative reverse transcription–polymerase chain reaction (qRT-PCR).

Whole cell lysates were harvested with Cells-to-cDNA II Cell Lysis Buffer (Ambion). Lysates were treated with DNase I (Ambion) and reverse transcribed in a total volume of 20 µl, using 200 ng

random primers and 100 units Moloney murine leukaemia virus reverse transcriptase (Invitrogen), according to the manufacturer's instructions, in the presence of 40 units RNasin (Promega). Relative quantification of genes was performed using an ABI Prism 7500 Sequence Detection System (Applied Biosystems). PCRs used 5 μ l of reverse-transcribed RNA (a 10-fold dilution of cDNA was used for 18S analyses). The MMP and ADAMTS primers and probes were previously described^{57,58}. The primers and probes for *HO-1*, *A20* were designed using the Universal Probe Library (Roche). Gene expression was normalised to ribosomal *18S* and relative quantification was calculated using $2^{-\Delta\Delta Ct}$.

Gene promoter reporter assays.

SW1353 cells were plated at 6×10^3 cells/well in 96-well plates and left to adhere overnight. At least three independent experiments were carried out each performed in triplicate ($n=3$). Transfections were carried out using 150 ng reporter plasmid DNA and 50ng renilla control plasmid and Lipofectamine 2000 (Fisher Scientific) for 24 hours. Cells were synchronised by serum starvation overnight, prior to treatments. Optimal time and cytokine concentrations were determined for each reporter individually and used in all downstream experiments. Cytokine treatments were: interleukin-1 (IL1) (5ng/ml) 2 hours; transforming growth factor beta (TGF β) (1ng/ml), 6 hours; bone morphogenic protein 6 (BMP6) (200ng/ml), 24 hours; wingless-type MMTV integration site family 3A (WNT3a) (100ng/ml), 24 hours. Luminescence was measured using the Dual-Glo[®] Luciferase Assay System (Promega) and EnVision Multilabel Plate Reader (PerkinElmer). pGL3 BRE Luciferase was a gift from Martine Roussel & Peter ten Dijke (Addgene plasmid # 45126)⁵⁹, pGL4.32[*luc2P*/NF κ B-RE/Hygro] (Promega), pGL3[CAGA-*luc*] was a kind gift from Jean-Michel Gauthier⁶⁰, M50 Super 8x TOPFlash was a gift from Randall Moon (Addgene plasmid # 12456)⁶¹.

Reactive Oxygen Species (ROS) Analysis

ROS were detected using 2',7'-dichlorofluorescein diacetate (DCFDA Cellular ROS Detection Assay Kit from Abcam (cat: ab113851)) according to the manufacturer's instructions. Briefly, primary chondrocytes were plated at 8×10^3 cells per well in 96-well plates. The experiment was repeated three times using three different patient cell isolates ($n=3$), each performed in triplicate. Treatments were 1.5-12 μ M SFN, API, ISO or LUT pre-treatment for 30 mins \pm (100 μ M) tert-Butyl hydroperoxide (tBHP) for 4 hours. ROS levels were measured at Ex/Em 492/530nm using EnVision Multilabel Plate Reader (PerkinElmer).

In vitro cartilage degradation assays

Cartilage explants were pre-treated with 0-24 μ M SFN, API, ISO or LUT. Cytokines IL-1 and OSM (0.5 ng/ml and 5 ng/ml respectively) were added to induce cartilage breakdown. All treatments were performed in triplicate and the experiment performed three times ($n=3$). Remaining cartilage was papain-digested overnight at 65°C. Glycosaminoglycan (GAG) was measured in the media and digested cartilage as described previously^{62,63}.

Kinase signalling pathway analysis

Primary chondrocytes were plated at 3×10^6 cells per well in 150 mm plates and allowed to adhere overnight. Chondrocytes were treated with IL-1 β (5ng/ml), for 10 mins with or without SFN, API or ISO (10 μ M) pre-treatment (30 mins). Cell lysates were prepared and applied to Proteome Profiler Array (R & D Systems) according to the manufacturer's instructions. Phosphorylation levels were semi-quantified using ImageJ software.

Cytotoxicity in HACs

HACs were treated with 10 μ M of each bioactive for 24 hours in three independent cell isolates ($n=3$). Lactate dehydrogenase (LDH) was measured in the culture media using the CytoTox 96[®] NonRadioactive Cytotoxicity Assay (Promega) according to the manufacturer's instructions. Total LDH was measured by lysing cells. Absorbance was measured at 492nm using an EnVision Multilabel Plate Reader (PerkinElmer) (Supplementary data S4).

Data availability

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

Statistical testing

Fold change values were log transformed and one-way analysis of variance (ANOVA) with Dunnett's post-test, were performed using GraphPad Prism version 7.03 for Windows to obtain adjusted p-values. Mean fold change and confidence intervals are reported for fold change values. $p<0.05$ was considered significant. Synergism was analysed using the Chou & Talalay method⁶⁴ and CompuSyn v1.0 software. <http://www.combosyn.com/index.html>

Acknowledgements

We give thanks to all patients who volunteered to donate their cartilage tissue and the staff at Spire Norwich Hospital (Norfolk, UK) for their help in collecting these samples. We would like to thank H.G Blake Ltd (Norfolk, UK) for their generous gift of bovine cartilage tissue.

Additional Information

Author Contributions

Contributors I.M.C., R.K.D., J.G., S.G., Y.P.B., and A.C. were involved in the conception and design of the study, acquisition of data, analysis and interpretation of data; drafting and revising of manuscript.

Competing Interests

The authors declare no competing interests.

References

- 1 Cross, M. *et al.* The global burden of hip and knee osteoarthritis: estimates from the Global Burden of Disease 2010 study. *Annals of the Rheumatic Diseases*, doi:10.1136/annrheumdis-2013-204763 (2014).
- 2 Brennan-Olsen, S. L. *et al.* Prevalence of arthritis according to age, sex and socioeconomic status in six low and middle income countries: analysis of data from the World Health Organization study on global AGEing and adult health (SAGE) Wave 1. *BMC Musculoskeletal Disord* **18**, 271, doi:10.1186/s12891-017-1624-z (2017).
- 3 ArthritisResearchUK. Osteoarthritis in general practice. Data and perspectives. *Keel Data Report* (2013).
- 4 Barnett, K. *et al.* Epidemiology of multimorbidity and implications for health care, research, and medical education: a cross-sectional study. *Lancet* **380**, 37-43, doi:10.1016/S0140-6736(12)60240-2 (2012).
- 5 Woolf, A. D. The Bone and Joint Decade 2000–2010. *Annals of the Rheumatic Diseases* **59**, 81-82, doi:10.1136/ard.59.2.81 (2000).

- 6 Woolf, A. D. & Pfleger, B. Burden of major musculoskeletal conditions. *Bulletin of the World Health Organisation* **81**, 646-656 (2003).
- 7 Kiadaliri, A. A. *et al.* Association of knee pain and different definitions of knee osteoarthritis with health-related quality of life: a population-based cohort study in southern Sweden. *Health Qual Life Outcomes* **14**, 121, doi:10.1186/s12955-016-0525-4 (2016).
- 8 Heathfield, T. F., Onnerfjord, P., Dahlberg, L. & Heinegard, D. Cleavage of fibromodulin in cartilage explants involves removal of the N-terminal tyrosine sulfate-rich region by proteolysis at a site that is sensitive to matrix metalloproteinase-13. *J Biol Chem* **279**, 6286-6295, doi:10.1074/jbc.M307765200 (2004).
- 9 Bondeson, J., Wainwright, S., Hughes, C. & Caterson, B. The regulation of the ADAMTS4 and ADAMTS5 aggrecanases in osteoarthritis: a review. *Clin Exp Rheumatol* **26**, 139-145 (2008).
- 10 Allam, M. F. & Arjona, O. Health promotion or pharmacological treatment for chronic diseases? *J Prev Med Hyg* **54**, 11-13 (2013).
- 11 Organization, G. W. H. Global status report on noncommunicable diseases 2010. 176 (World Health Organization, Italy, 2011).
- 12 Green, J. A. *et al.* The potential for dietary factors to prevent or treat osteoarthritis. *Proc Nutr Soc* **73**, 278-288, doi:10.1017/s0029665113003935 (2014).
- 13 Mobasheri, A., Henrotin, Y., Biesalski, H. K. & Shakibaei, M. Scientific evidence and rationale for the development of curcumin and resveratrol as nutraceuticals for joint health. *International journal of molecular sciences* **13**, 4202-4232, doi:10.3390/ijms13044202 (2012).
- 14 Williams, F. M. *et al.* Dietary garlic and hip osteoarthritis: evidence of a protective effect and putative mechanism of action. *BMC musculoskeletal disorders* **11**, 280, doi:10.1186/1471-2474-11-280 (2010).
- 15 Facchini, A. *et al.* Sulforaphane protects human chondrocytes against cell death induced by various stimuli. *J Cell Physiol* **226**, 1771-1779, doi:10.1002/jcp.22506 (2011).
- 16 Davidson, R. K. *et al.* Sulforaphane represses matrix-degrading proteases and protects cartilage from destruction in vitro and in vivo. *Arthritis Rheum* **65**, 3130-3140, doi:10.1002/art.38133 (2013).
- 17 Ko, J. Y., Choi, Y. J., Jeong, G. J. & Im, G. I. Sulforaphane-PLGA microspheres for the intra-articular treatment of osteoarthritis. *Biomaterials* **34**, 5359-5368, doi:10.1016/j.biomaterials.2013.03.066 (2013).
- 18 Berenbaum, F. Does broccoli protect from osteoarthritis? *Joint Bone Spine* **81**, 284-286, doi:10.1016/j.jbspin.2014.04.001 (2014).
- 19 Javaheri, B. *et al.* Stable sulforaphane protects against gait anomalies and modifies bone microarchitecture in the spontaneous STR/Ort model of osteoarthritis. *Bone* **103**, 308-317, doi:10.1016/j.bone.2017.07.028 (2017).
- 20 Miesan, K. H. & Mohamed, S. Flavonoid (myricetin, quercetin, kaempferol, luteolin, and apigenin) content of edible tropical plants. *J Agric Food Chem* **49**, 3106-3112 (2001).
- 21 Lopez-Lazaro, M. Distribution and biological activities of the flavonoid luteolin. *Mini Rev Med Chem* **9**, 31-59 (2009).
- 22 Moncada-Pazos, A., Obaya, A. J., Vilorio, C. G., Lopez-Otin, C. & Cal, S. The nutraceutical flavonoid luteolin inhibits ADAMTS-4 and ADAMTS-5 aggrecanase activities. *J Mol Med (Berl)* **89**, 611-619, doi:10.1007/s00109-011-0741-7 (2011).
- 23 Kuppusamy, U. R. & Das, N. P. Inhibitory effects of flavonoids on several venom hyaluronidases. *Experientia* **47**, 1196-1200 (1991).
- 24 Durigova, M. *et al.* Characterization of an ADAMTS-5-mediated cleavage site in aggrecan in OSM-stimulated bovine cartilage. *Osteoarthritis Cartilage* **16**, 1245-1252, doi:10.1016/j.joca.2008.02.013 (2008).
- 25 Park, J. S. *et al.* Apigenin Regulates Interleukin-1beta-Induced Production of Matrix Metalloproteinase Both in the Knee Joint of Rat and in Primary Cultured Articular

- Chondrocytes. *Biomolecules & therapeutics* **24**, 163-170, doi:10.4062/biomolther.2015.217 (2016).
- 26 Lim, H., Park, H. & Kim, H. P. Effects of flavonoids on matrix metalloproteinase-13 expression of interleukin-1beta-treated articular chondrocytes and their cellular mechanisms: inhibition of c-Fos/AP-1 and JAK/STAT signaling pathways. *J Pharmacol Sci* **116**, 221-231 (2011).
- 27 Zhuang, C. *et al.* Chalcone: A Privileged Structure in Medicinal Chemistry. *Chem Rev* **117**, 7762-7810, doi:10.1021/acs.chemrev.7b00020 (2017).
- 28 Ji, B. *et al.* Isoliquiritigenin blunts osteoarthritis by inhibition of bone resorption and angiogenesis in subchondral bone. *Scientific Reports* **8**, 1721, doi:10.1038/s41598-018-19162-y (2018).
- 29 Zhang, L., Ma, S., Su, H. & Cheng, J. Isoliquiritigenin Inhibits IL-1beta-Induced Production of Matrix Metalloproteinase in Articular Chondrocytes. *Mol Ther Methods Clin Dev* **9**, 153-159, doi:10.1016/j.omtm.2018.02.006 (2018).
- 30 Zeng, J. *et al.* Isoliquiritigenin alleviates early brain injury after experimental intracerebral hemorrhage via suppressing ROS- and/or NF-κB-mediated NLRP3 inflammasome activation by promoting Nrf2 antioxidant pathway. *Journal of Neuroinflammation* **14**, 119, doi:10.1186/s12974-017-0895-5 (2017).
- 31 Ji, B. *et al.* Isoliquiritigenin suppresses IL-1beta induced apoptosis and inflammation in chondrocyte-like ATDC5 cells by inhibiting NF-kappaB and exerts chondroprotective effects on a mouse model of anterior cruciate ligament transection. *Int J Mol Med* **40**, 1709-1718, doi:10.3892/ijmm.2017.3177 (2017).
- 32 Chou, T.-C. Theoretical Basis, Experimental Design, and Computerized Simulation of Synergism and Antagonism in Drug Combination Studies. *Pharmacological Reviews* **58**, 621-681, doi:10.1124/pr.58.3.10 (2006).
- 33 Heiss, E., Herhaus, C., Klimo, K., Bartsch, H. & GerhÄuser, C. Nuclear Factor κB Is a Molecular Target for Sulforaphane-mediated Anti-inflammatory Mechanisms. *Journal of Biological Chemistry* **276**, 32008-32015, doi:10.1074/jbc.M104794200 (2001).
- 34 Choi, Y. J., Lee, W.-S., Lee, E.-G., Sung, M.-S. & Yoo, W.-H. Sulforaphane Inhibits IL-1β-Induced Proliferation of Rheumatoid Arthritis Synovial Fibroblasts and the Production of MMPs, COX-2, and PGE2. *Inflammation* **37**, 1496-1503, doi:10.1007/s10753-014-9875-4 (2014).
- 35 Dinkova-Kostova, A. T. & Talalay, P. Direct and indirect antioxidant properties of inducers of cytoprotective proteins. *Mol Nutr Food Res* **52 Suppl 1**, S128-138, doi:10.1002/mnfr.200700195 (2008).
- 36 Li, Y. *et al.* Sulforaphane, a dietary component of broccoli/broccoli sprouts, inhibits breast cancer stem cells. *Clin Cancer Res* **16**, 2580-2590, doi:10.1158/1078-0432.ccr-09-2937 (2010).
- 37 Kaminski, B. M. *et al.* Isothiocyanate sulforaphane inhibits protooncogenic ornithine decarboxylase activity in colorectal cancer cells via induction of the TGF-beta/Smad signaling pathway. *Mol Nutr Food Res* **54**, 1486-1496, doi:10.1002/mnfr.201000105 (2010).
- 38 Sun, C., Li, S. & Li, D. Sulforaphane mitigates muscle fibrosis in mdx mice via Nrf2-mediated inhibition of TGF-beta/Smad signaling. *Journal of applied physiology (Bethesda, Md. : 1985)* **120**, 377-390, doi:10.1152/jappphysiol.00721.2015 (2016).
- 39 Wang, C. *et al.* BMP-6 inhibits MMP-9 expression by regulating heme oxygenase-1 in MCF-7 breast cancer cells. *Journal of Cancer Research and Clinical Oncology* **137**, 985-995, doi:10.1007/s00432-010-0963-z (2011).
- 40 Yan, J., Yang, S., Zhang, J., Zhai, C. & Zhu, T. BMP6 attenuates oxidant injury in HK-2 cells via Smad-dependent HO-1 induction. *Free Radic Biol Med* **46**, 1275-1282, doi:10.1016/j.freeradbiomed.2009.02.007 (2009).

- 41 Dinkova-Kostova, A. T., Fahey, J. W., Kostov, R. V. & Kensler, T. W. KEAP1 and Done? Targeting the NRF2 Pathway with Sulforaphane. *Trends Food Sci Technol* **69**, 257-269, doi:10.1016/j.tifs.2017.02.002 (2017).
- 42 Balez, R. *et al.* Neuroprotective effects of apigenin against inflammation, neuronal excitability and apoptosis in an induced pluripotent stem cell model of Alzheimer's disease. *Scientific Reports* **6**, 31450, doi:10.1038/srep31450 (2016).
- 43 Wang, J. *et al.* Anti-Inflammatory Effects of Apigenin in Lipopolysaccharide-Induced Inflammatory in Acute Lung Injury by Suppressing COX-2 and NF-kB Pathway. *Inflammation* **37**, 2085-2090, doi:10.1007/s10753-014-9942-x (2014).
- 44 Mirzoeva, S., Franzen, C. A. & Pelling, J. C. Apigenin inhibits TGF-beta-induced VEGF expression in human prostate carcinoma cells via a Smad2/3- and Src-dependent mechanism. *Mol Carcinog* **53**, 598-609, doi:10.1002/mc.22005 (2014).
- 45 Zhang, Y. *et al.* Apigenin induces dermal collagen synthesis via smad2/3 signaling pathway. *European journal of histochemistry : EJH* **59**, 2467, doi:10.4081/ejh.2015.2467 (2015).
- 46 Yokomizo, A. & Moriwaki, M. Effects of uptake of flavonoids on oxidative stress induced by hydrogen peroxide in human intestinal Caco-2 cells. *Biosci Biotechnol Biochem* **70**, 1317-1324 (2006).
- 47 Valdameri, G. *et al.* Involvement of catalase in the apoptotic mechanism induced by apigenin in HepG2 human hepatoma cells. *Chem Biol Interact* **193**, 180-189, doi:10.1016/j.cbi.2011.06.009 (2011).
- 48 Crasci, L., Cardile, V., Longhitano, G., Nanfita, F. & Panico, A. Anti-degenerative effect of Apigenin, Luteolin and Quercetin on human keratinocyte and chondrocyte cultures: SAR evaluation. *Drug research* **68**, 132-138, doi:10.1055/s-0043-120662 (2018).
- 49 Liu, Q., Lv, H., Wen, Z., Ci, X. & Peng, L. Isoliquiritigenin Activates Nuclear Factor Erythroid-2 Related Factor 2 to Suppress the NOD-Like Receptor Protein 3 Inflammasome and Inhibits the NF-kappaB Pathway in Macrophages and in Acute Lung Injury. *Frontiers in immunology* **8**, 1518, doi:10.3389/fimmu.2017.01518 (2017).
- 50 Chen, H. *et al.* Isoliquiritigenin-induced effects on Nrf2 mediated antioxidant defence in the HL-60 cell monocytic differentiation. *Cell Biol Int* **37**, 1215-1224, doi:10.1002/cbin.10156 (2013).
- 51 Kang, B. J., Ryu, J., Lee, C. J. & Hwang, S. C. Luteolin Inhibits the Activity, Secretion and Gene Expression of MMP-3 in Cultured Articular Chondrocytes and Production of MMP-3 in the Rat Knee. *Biomolecules & therapeutics* **22**, 239-245, doi:10.4062/biomolther.2014.020 (2014).
- 52 Gasper, A. V. *et al.* Glutathione S-transferase M1 polymorphism and metabolism of sulforaphane from standard and high-glucosinolate broccoli. *Am J Clin Nutr* **82**, 1283-1291, doi:10.1093/ajcn/82.6.1283 [pii] (2005).
- 53 Cao, J., Zhang, Y., Chen, W. & Zhao, X. The relationship between fasting plasma concentrations of selected flavonoids and their ordinary dietary intake. *Br J Nutr* **103**, 249-255, doi:10.1017/S000711450999170X (2010).
- 54 Zhang, Y., Cao, J., Wang, Y. & Xiao, S. Simultaneous Determination of Glycyrrhizin and 15 Flavonoids in Licorice and Blood by High Performance Liquid Chromatography with Ultraviolet Detector. *ISRN Analytical Chemistry* **2013**, 7, doi:10.1155/2013/786151 (2013).
- 55 Zhang, Y. Molecular mechanism of rapid cellular accumulation of anticarcinogenic isothiocyanates. *Carcinogenesis* **22**, 425-431 (2001).
- 56 Kay, C. D., Pereira-Caro, G., Ludwig, I. A., Clifford, M. N. & Crozier, A. Anthocyanins and Flavanones Are More Bioavailable than Previously Perceived: A Review of Recent Evidence. *Annu Rev Food Sci Technol* **8**, 155-180, doi:10.1146/annurev-food-030216-025636 (2017).
- 57 Porter, S., Clark, I. M., Kevorkian, L. & Edwards, D. R. The ADAMTS metalloproteinases. *Biochem J* **386**, 15 - 27 (2005).

- 58 Nuttall, R. K. *et al.* Elevated membrane-type matrix metalloproteinases in gliomas revealed by profiling proteases and inhibitors in human cancer cells. *Mol Cancer Res* **1**, 333-345 (2003).
- 59 Korchynskiy, O. & ten Dijke, P. Identification and Functional Characterization of Distinct Critically Important Bone Morphogenetic Protein-specific Response Elements in the Id1 Promoter. *Journal of Biological Chemistry* **277**, 4883-4891, doi:10.1074/jbc.M111023200 (2002).
- 60 Dennler, S. *et al.* Direct binding of Smad3 and Smad4 to critical TGF β -inducible elements in the promoter of human plasminogen activator inhibitor-type 1 gene. Vol. 17 (1998).
- 61 Veeman, M. T., Slusarski, D. C., Kaykas, A., Louie, S. H. & Moon, R. T. Zebrafish prickles, a modulator of noncanonical Wnt/Fz signaling, regulates gastrulation movements. *Curr Biol* **13**, 680-685 (2003).
- 62 Billington, C. J. Cartilage proteoglycan release assay. *Methods Mol Biol* **151**, 451-456 (2001).
- 63 Shingleton, W. D., Ellis, A. J., Rowan, A. D. & Cawston, T. E. Retinoic acid combines with interleukin-1 to promote the degradation of collagen from bovine nasal cartilage: matrix metalloproteinases-1 and -13 are involved in cartilage collagen breakdown. *J Cell Biochem* **79**, 519 - 531 (2000).
- 64 Chou, T. C. & Talalay, P. Quantitative analysis of dose-effect relationships: the combined effects of multiple drugs or enzyme inhibitors. *Adv Enzyme Regul* **22**, 27-55 (1984).

Figure Legends

Figure 1. IL-1-induced *MMP13* gene expression in human primary articular chondrocytes. Four selected bioactives (a) sulforaphane (SFN), (b) apigenin (API), (c) isoliquiritigenin (ISO), (d) luteolin (LUT) were titrated against IL-1 treatment (5ng/ml) in at least three independent isolates of human primary articular chondrocytes ($n=3$) each performed in triplicate, for 6 h to establish a dose response. *MMP13* mRNA expression was measured using RT-qPCR and normalised to *18S*. Data is expressed as mean fold change (FC) \pm SEM. One-way ANOVA, and all treatments were compared to IL-1 treatment with Dunnett's post-test. Statistical significance is shown as $p<0.05$ *, $p<0.01$ **, $p<0.001$ ***, $p=0.0001$ ****

Figure 2. Reactive oxygen species (ROS) measured using the DCFDA assay in human primary articular chondrocytes. Cells were treated with 100 μ M tert-butyl-hydroperoxide (tBHP) alone (4 h) \pm pre-treatment with (a) SFN, (b) API, (c) ISO or (d) LUT at 0.75-12 μ M for 30 mins. Data are from three independent isolates of human primary articular chondrocytes ($n=3$), where each experiment was performed in triplicate. Data are shown as mean fold change (FC) to control \pm SEM. One-way ANOVA, and all treatments were compared to IL-1 treatment with Dunnett's post-test. Statistical significance is shown as: $p<0.01$ **, <0.001 ***, $p\leq 0.0001$ ****

Figure 3. Transcriptional signalling pathway analysis. Luciferase reporter activity for IL-1/nuclear factor-kappa B, TGF- β /Smad2/3, BMP6/Smad1/5/8 and Wnt3A/TCF/Lef (TOPFlash), and using the SW1353 chondrosarcoma cell line in response to (a-d) SFN, (e-h) API, (i-l) ISO or (m-p) LUT treatment. Cells were treated with IL-1 (5ng/ml, 2 h), TGF- β (1ng/ml, 6 h), BMP6 (200ng/ml, 24 h) and Wnt3a (100ng/ml, 24 h). Values given are from at least three independent experiments ($n=3$) where each experiment was performed in triplicate. Data are shown as mean fold change (FC) to control \pm SEM. One-way ANOVA, and all treatments were compared to IL-1 treatment with Dunnett's post-test. Statistical significance is shown as: $p<0.05$ *, $p<0.01$ **, $p<0.001$ ***, $p\leq 0.0001$. RLU – relative light units (normalised to control plasmid).

Figure 4. Cartilage tissue degradation *in vitro*. Aggrecan loss measured using 1,9-dimethyl-methylene blue assay from bovine cartilage explants in response to (a) SFN, (b) API, (c) ISO or (d) LUT treatment (open bars), + IL-1/OSM at 0.5ng/ml and 5ng/ml respectively (black bars). Values given are from three independent experiments ($n=3$) and each performed in triplicate. Data are shown as mean percent loss of total glycosaminoglycan (GAG) \pm SEM. One-way ANOVA, and all treatments with cytokines were compared to IL-1/OSM treatment alone with Dunnett's post-test. Statistical significance is shown as: $p<0.05$ *

Figure 5. Cytokine-induced expression levels of key genes in osteoarthritis with SFN/ISO combination treatment. IL1-induced gene expression of *ADAMTS4*, *ADAMTS5*, *HO-1* and *A20* in human articular primary chondrocytes (HACs). HACs were pre-treated with SFN or ISO individually and in the equipotent SFN:ISO combination ratio: 0.75:0.625 μ M for 30 mins followed by IL1 (5ng/ml) 6 hours. Gene expression was measured in at least three independent isolates of human primary articular chondrocytes ($n=3$) by RT-qPCR and normalised to *18S*. Data is expressed as mean fold change (FC) from IL-1 \pm SEM. One-way ANOVA, and all treatments were compared to IL-1 treatment with Dunnett's post-test. Statistical significance is shown as: $p<0.05$ *, $p<0.01$ **, $p<0.001$ ***, $p\leq 0.0001$ ****.

Tables

Induced Bioactive	MMP1		MMP13		ADAMTS4		ADAMTS5	
	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value
Genistin	0.75 (-0.88 to 1.39)	0.994	0.89 (-0.62 to 0.84)	0.999	0.81 (-0.50 to 0.88)	0.997	0.77 (-0.62 to 1.07)	0.996
Luteolin	0.30 (-0.43 to 1.84)	0.429	0.39 (-0.12 to 1.34)	0.201	0.41 (-0.10 to 1.28)	0.068	0.27 (-0.11 to 1.58)	0.075
Polydatin	0.73 (-0.86 to 1.41)	0.997	0.74 (-0.47 to 0.99)	0.987	0.80 (-0.49 to 0.89)	0.997	0.84 (-0.68 to 1.01)	0.999
Curcumin	0.61 (-0.74 to 1.53)	0.899	0.62 (-0.35 to 1.11)	0.799	0.83 (-0.52 to 0.86)	0.992	0.44 (-0.28 to 1.41)	0.124
Apigenin	0.23 (-0.36 to 1.91)	0.186	0.18 (0.09 to 1.55)	0.005	0.11 (0.20 to 1.58)	0.0001	0.22 (-0.06 to 1.62)	0.018
Myricetin	0.97 (-1.10 to 1.16)	0.999	1.07 (-0.80 to 0.66)	0.999	1.34 (-1.03 to 0.35)	0.994	0.77 (-0.61 to 1.07)	0.991
Isoliqu	0.23 (-0.36 to 1.90)	0.151	0.14 (0.14 to 1.59)	0.0005	0.08 (0.15 to 1.69)	0.0001	0.11 (0.05 to 1.73)	0.001
Ursolic acid	0.82 (-0.95 to 1.32)	0.991	0.31 (-0.03 to 1.43)	0.041	0.69 (-0.38 to 1.00)	0.902	0.33 (-0.17 to 1.52)	0.136
EGCG	1.03 (-1.16 to 1.11)	0.999	1.01 (-0.74 to 0.72)	0.999	1.27 (-0.95 to 0.42)	0.999	0.97 (-0.81 to 0.88)	0.999
Naringin	0.94 (-1.07 to 1.20)	0.999	1.13 (-0.86 to 0.60)	0.999	1.16 (-0.85 to 0.53)	0.999	0.97 (-0.81 to 0.87)	0.999
Sulforaphane	0.17 (-0.31 to 1.96)	0.163	0.12 (0.15 to 1.61)	0.001	0.03 (0.28 to 1.66)	0.0001	0.23 (-0.07 to 1.61)	0.053

Table 1. Inhibition of cytokine-induced levels of key osteoarthritis matrix metalloproteinase mRNA expression in primary chondrocytes. Cytokine-induced matrix metalloproteinase gene expression in human articular primary chondrocytes pre-treated with 10µM of each bioactive 30 mins then 5ng/ml interleukin-1 (IL-1) for 6 hours. Cell isolates from three patients ($n=3$) were used and each independent experiment performed in triplicate. Levels of *MMP1*, *MMP13*, *ADAMTS4* and *ADAMTS5* mRNA, normalised to *18S* were measured by RT-qPCR. Mean fold change (FC) to IL-1, 95% confidence intervals (CI) are shown. One-way ANOVA, and all treatments were compared to IL-1 treatment with Dunnett's post-test corrected p-values shown.

Basal Bioactive	MMP1		MMP13		ADAMTS4		ADAMTS5	
	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value	Mean FC (95% CI)	p- value
Genistin	1.03 (-2.55 to 2.50)	0.999	1.09 (-3.14 to 2.96)	0.999	-	-	0.96 (-0.48 to 0.56)	0.999
Luteolin	0.96 (-2.48 to 2.56)	0.999	1.83 (-3.88 to 2.22)	0.997	-	-	0.71 (-0.23 to 0.81)	0.766
Polydatin	0.90 (-2.42 to 2.62)	0.999	1.88 (-3.93 to 2.16)	0.985	-	-	0.98 (-0.50 to 0.54)	0.999
Curcumin	1.13 (-2.65 to 2.39)	0.999	2.43 (-4.48 to 1.61)	0.822	-	-	0.70 (-0.22 to 0.83)	0.838
Apigenin	0.76 (-2.28 to 2.77)	0.912	2.05 (-4.10 to 2.00)	0.996	-	-	0.59 (-0.11 to 0.93)	0.337
Myricetin	0.74 (-2.26 to 2.78)	0.914	1.89 (-3.94 to 2.16)	0.999	-	-	0.77 (-0.29 to 0.75)	0.965
Isoliqu	0.95 (-2.47 to 2.57)	0.999	2.02 (-4.07 to 2.02)	0.991	-	-	0.49 (-0.03 to 1.01)	0.148
Ursolic acid	4.35 (-5.87 to -0.82)	0.004	0.64 (-3.04 to 3.77)	0.864	-	-	0.69 (-0.21 to 0.83)	0.532
EGCG	1.11 (-2.63 to 2.41)	0.999	1.81 (-3.86 to 2.24)	0.999	-	-	0.62 (-0.14 to 0.90)	0.442
Naringin	0.95 (-2.48 to 2.57)	0.999	1.09 (-3.14 to 2.95)	0.999	-	-	0.92 (-0.44 to 0.60)	0.999
Sulforaphane	0.92 (-2.44 to 2.60)	0.999	1.75 (-3.80 to 2.29)	0.993	-	-	0.25 (0.23 to 1.27)	0.001

Table 2. Inhibition of basal levels of key osteoarthritis matrix metalloproteinase mRNA expression in primary chondrocytes. Basal matrix metalloproteinase gene expression in human articular primary chondrocytes pre-treated with 10µM of each bioactive for 6 hours. Cell isolates from three patients were used ($n=3$) and each independent experiment performed in triplicate. Expression levels of *MMP1*, *MMP13*, *ADAMTS4* and *ADAMTS5* mRNA, normalised to *18S* were measured by RT-qPCR. Mean fold change (FC) to control, 95% confidence intervals (CI) are shown. One-way ANOVA, and all treatments were compared to control with Dunnett's post-test corrected p-values shown.

Bioactive	D_m	m	r
SFN	3.33	1.43	0.75
API	6.76	2.54	0.93
ISO	2.89	1.24	0.73
LUT	3.80	1.32	0.84

Table 3. Dose-effect relationship parameters of SFN, API, ISO and LUT individually against IL1-induced *MMP13* mRNA expression in primary chondrocytes. IC_{50} concentration (D_m) is shown in µM. The slope (m) and linear correlation coefficient (r) of the median-effect plot are given. Values are the mean of three human articular primary chondrocyte isolates ($n=3$) and each experiment contained six concentrations for each bioactive, in duplicate. Gene expression was measured using RT-qPCR and normalised to *18S*.

Bioactive Combo	CI Values					Assigned symbol
	ED ₅₀	ED ₇₅	ED ₉₀	ED ₉₅	mean CI	
SA	0.706	0.879	1.127	1.356	1.017	±
SI	0.872	0.758	0.661	0.604	0.724	++
SL	1.066	1.404	1.851	2.235	1.639	---
AI	1.213	1.072	0.997	0.975	1.064	±
AL	0.887	1.008	1.189	1.361	1.111	-
IL	3.111	2.208	1.569	1.244	2.033	---
SAI	1.066	1.161	1.311	1.454	1.248	--
AIL	1.919	1.754	1.663	1.642	1.744	---
ILS	1.550	1.712	1.896	2.034	1.798	---
LSA	1.753	1.836	1.978	2.119	1.922	---
SAIL	1.368	1.770	2.357	2.917	2.103	---

Table 4. Bioactive combination indices of SFN, API, ISO and LUT for *MMP13* expression in primary chondrocytes. Combination Index (CI) values of bioactive combinations against IL1-induced *MMP13* mRNA expression in human articular primary chondrocytes ($n=3$) measured by RT-qPCR and normalised to *18S*. Combination ratios were based on IC₅₀ values. Each bioactive component had a combination ratio of: S:A:I:L = 0.75:1:0.625:1.25 μ M. Assigned symbols are: nearly additive (\pm), moderate synergism (++), slight antagonism (-), moderate antagonism (--), antagonism (---)³². CI = Combination index.

Combination	DRI Values at:				CI Values at:			
	ED ₅₀	ED ₇₅	ED ₉₀	ED ₉₅	ED ₅₀	ED ₇₅	ED ₉₀	ED ₉₅
S + A (0.75:1)	1.418	3.292	7.641	13.549	0.706	0.879	1.127	1.356
S + I (0.75:0.625)	1.479	2.930	5.808	9.248	0.872	0.758	0.661	0.604
A + I (1:0.625)	3.328	5.781	10.043	14.622	1.213	1.072	0.997	0.975

Table 5. Dose reduction Index (DRI) values for *MMP13* expression in relation to combination index (CI) values for synergistic combinations of SFN/API, SFN/ISO and API/ISO in primary chondrocytes. The DRI describes the fold reduction in bioactive concentration needed in a synergistic combination at each effect level shown (compared with each bioactive alone). Gene expression was measured by RT-qPCR, normalised to *18S*, from three independent experiments, $n=3$. Synergism (CI <1), additivity (CI =1), antagonism (CI >1).









