Spinacia oleracea extract attenuates disease progression and sub-chondral bone changes in monosodium iodoacetate-induced osteoarthritis in rats

Dharmendra Choudhary, Priyanka Kothari, Ashish Kumar Tripathi, Sonu Singh, Sulekha Adhikary, Naseer Ahmad, Sudhir Kumar, Kapil Dev, Vijay Kumar Mishra, Shubha Shukla, Rakesh Maurya, Prabhat R. Mishra and Ritu Trivedi

Abstract

Background: Spinacia oleracea is an important dietary vegetable in India and throughout the world and has many beneficial effects. It is cultivated globally. However, its effect on osteoarthritis that mainly targets the cartilage cells remains unknown. In this study we aimed to evaluate the anti-osteoarthritic and chondro-protective effects of SOE on chemically induced osteoarthritis (OA).

Methods: OA was induced by intra-patellar injection of monosodium iodoacetate (MIA) at the knee joint in rats. SOE was then given orally at 250 and 500 mg.kg^-1 day^-1 doses for 28 days to these rats. Anti-osteoarthritic potential of SOE was evaluated by micro-CT, mRNA and protein expression of pro-inflammatory and chondrogenic genes, clinically relevant biomarker's and behavioural experiments.

Results: In vitro cell free and cell based assays indicated that SOE acts as a strong anti-oxidant and an anti-inflammatory agent. Histological analysis of knee joints at the end of the experiment by safranin-o and toluidine blue staining established its protective effect. Radiological data corroborated the findings with improvement in the joint space and irregularity of the articular and atrophied femoral condyles and tibial plateau. Micro-CT analysis of sub-chondral bone indicated that SOE had the ability to mitigate OA effects by increasing bone volume to tissue volume (BV/TV) which resulted in decrease of trabecular pattern factor (Tb.Pf) by more than 200%. SOE stimulated chondrogenic marker gene expression with reduction in pro-inflammatory markers. Purified compounds isolated from SOE exhibited increased Sox-9 and Col-II protein expression in articular chondrocytes. Serum and urine analysis indicated that SOE had the potential to down-regulate glutathione S-transferase (GST) activity, clinical markers of osteoarthritis like cartilage oligometric matrix protein (COMP) and CTX-II. Overall, this led to a significant improvement in locomotion and balancing activity in rats as assessed by Open-field and Rota rod test.

Conclusion: On the basis of in vitro and in vivo experiments performed with Spinacea oleracea extract we can deduce that SOE has the ability to alleviate the MIA induced deleterious effects.

Keywords: Cartilage, Monosodium iodoacetate (MIA), Osteoarthritis (OA), Spinacia oleracea extract (SOE)
Background
Osteoarthritis (OA) is a high prevalence disease with socio-economic impact. It affects mainly the weight-bearing joints such as hips and knees, and causes physical disabilities. Over 100 million people worldwide suffer from OA [1]. It is a common rheumatologic problem with a prevalence of 22% to 39% in India [2]. Despite, the identified risk factors the exact pathogenesis of osteoarthritis remains unclear. As per reported literature, currently, there is no effective treatment that can cure osteoarthritis leaving joint replacement as the only therapeutic option in patients with severe osteoarthritis [3]. Articular cartilage degeneration is the primary concern in osteoarthritis. Cartilage degradation is associated with structural and metabolic changes in joints such as subchondral bone sclerosis and synovial membrane inflammation [4]. Biomechanical and biochemical interactions with subchondral bone and other joint tissues play important roles in maintaining homeostasis of articular cartilage [5]. Subchondral bone provides the mechanical support for the overlying articular cartilage during the movement of load-bearing joints and experience a constant adaptation in response to changes in the mechanical environment through modelling or remodelling [6]. Clinically, osteophyte formation is the characteristic marker of osteoarthritis [7]. Studies suggest that slow destructive process of joints results from the combination of mechanical stress, inflammation, and biochemical factors that includes mainly reactive oxygen species (ROS) and matrix metalloproteinases (MMPs) [8]. Prophylactic and symptomatic therapies are still available that reduce inflammation. However, therapies that can cure osteoarthritis are still far from reach.

The role of phyto-pharmaceuticals in the pathogenesis of osteoarthritis has drawn a lot of attention in recent years [9]. *Spinacia oleracea* which is cultivated globally is an important dietary food vegetable and a common raw material in the food processing industry [10]. The fresh leaves are cooked and taken as food. *Spinacia oleracea* has been used as dietary supplements or alternative medicinal foods in Indians as well as throughout the world. Spinach has been used as prevention or cure for diabetes [11], heart injury [12], neural disorders [13], hormone replacement therapy [14], anti-cancer [15] and many more. Currently, it is used as a proper health supplement and clinician suggests as healthy food which fight against many disease conditions and health supplements. However, there are no reports concerning the osteoarthritis and chondro-protective effect of leaves of *Spinacia oleracea*. We have previously shown the osteogenic effects of *Spinacia oleracea* extract (SOE) in post-menopausal bone loss and fracture healing activity [16]. In drill hole fracture healing model, SOE not only stimulated osteoblast cells but also mobilized chondrocyte proliferation at the injury site [16]. This prompted us to investigate its role in subchondral bone pathology and articular cartilage degeneration during the progression of osteoarthritis. Studies report that injection of monosodium iodoacetate (MIA) into the knee joints of rats is considered a suitable model for osteoarthritis and used for evaluation of chondroprotective activity of novel agents which also resembles the phenomena observed in human OA [17]. Therefore, the present study was hypothesized to investigate the effect of SOE in cartilage degenerative disease at articular cartilage tissue region with an objective to search and develop novel therapeutic alternatives for the prevention and management of chondrocytes related disorders.

The present study is being carried out to investigate the effect of oral administration of SOE on pathological changes in MIA induced OA in in vivo settings in rats.

Methods
Reagents and chemicals
Cell culture media and supplements such as Dulbecco’s Modified Eagle Medium: Nutrient Mixture F-12 (DMEM/F-12) Media, Fetal bovine serum (FBS), antibiotic cocktail, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), safranin-o and toluidine blue dye, 1, 1-diphenyl-2-picrylhydrazyl radical (DPPH), 2’-azino[3-ethylbenzthiazoline-6-sulfonic acid (ABTS), and Dimethyl methylene blue (DMMB) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Complementary DNA (cDNA) synthesis kit was purchased from Thermo Fisher Scientific (Waltham, Massachusetts, USA) and SYBR Green kit was obtained from Genetix (New Delhi, India). Rat CTX-II (ELISA KIT E-EL-R2554) and Rat COMP (ELISA kit E-EL-R0159) were purchased from Elabsiences (Elabsciences biotechnology co ltd, Hubei province, China).

Preparations of Spinaciaoleracea extract (SOE)
The leaves of *Spinacia oleracea* were collected from district Lucknow, India during the month of November–December, by Dr. Rakesh Maurya and co-workers. It was identified and authenticated by Dr. Lal Babu Choudhary, Principal Scientist at Plant Diversity, Systematics & Herbarium Division CSIR-National Botanical Research Institute, Lucknow India. An institutional Code no. (CDRI-2492) had been deposited in herbarium of investor laboratory, Division of medicinal and process chemistry, CDRI, Lucknow for future reference.

The fresh *Spinacia oleracea* leaves (30 kg) were cut into small pieces and placed in glass percolator with ethanol (20 L) and allowed to stand at room temperature for approx. 24 h. The percolate was collected. This process of extraction was repeated for four times. This extract was further filtered and concentrated under reduced pressure at 45 °C. The extract thus obtained was
further dried in a vacuum desiccator to remove remaining ethanol and moisture and finally, 560 g (1.8% yield) weight of extract was obtained. This dried extract (100% of Spinacia oleracea leaves) was further used for different in-vitro and in-vivo studies as described previously by us [16].

MTT (3-(4, 5-Dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide) assay
Cell viability was assessed using an MTT assay based on the ability of mitochondria of viable cells to convert soluble MTT into an insoluble purple formazan reaction product. First, dissolve SOE in 10 μl DMSO then it’s made up to 1.0 mg/ml by DMEM media; this is used as stock solution. This SOE stock is used for treatment on cultured rat articular chondrocytes cells (RAC) at different concentrations ranging from 1.95 to 1000 μg/ml. RAC was isolated from knee of young pups as per previously published protocols [18]. In brief, chondrocyte cells were isolated from distal femoral and proximal tibial condyles of 1 day-old Sprague-Dawley rats. Condyles tissue was isolated, cleaned from surrounding tissue and digested with trypsin for 30 min then processed in collagenase solution (2 mg/ml) at 37 °C for 4 h followed by passing through a 70 μm filter. Isolated cells were cultured for 48 h then trypsinized and 1 × 10^3 cells/well were seeded in 48 wells plate. After 80–90% confluences of articular chondrocyte cells in monolayer culture, cells were treated with different concentrations of SOE for 24 h in DMEM/F12 medium with 10% fetal bovine serum [19]. After 24 h of SOE treatment, RAC cells were treated with MTT solution (5 mg/ml in Dulbecco’s modified Eagle’s medium (DMEM/F12) without phenol red; Sigma) for 2 h. The MTT solution was then aspirated and replaced with 200 ml/well dimethyl sulfoxide (DMSO). Detection was done at 540 nm OD by spectrophotometer [20].

2, 2′-azinobis [3-ethylbenzthiazoline]-6-sulfonic acid (ABTS) free radical scavenging assay
The antioxidant potential of the Spinach oleracia extracts (SOE) was measured using 2, 2′- azinobis [3-ethylbenzthiazoline]-6-sulfonic acid (ABTS) assay [22, 23]. In brief ABTS (14 mM), and potassium per sulphate (4.88 mM) was mixed, then left overnight (12–16 h) at room temperature in the dark. SOE concentration ranging 15.6 to 1000 μg/ml was measured by adding with ABTS solution to initiate the reaction. The absorbance was read at wavelength 415 nm with incubating for 15 min by a spectrophotometer (molecular devices, USA). The antioxidant capacity of each concentration of the extract was determined based on the reduction of ABTS absorption by calculating percentage of antioxidant activity as given in DPPH assay [26].

Cartilage dissection and explant culture for DMMB assays
Articular cartilage tissues were obtained from knee joints of healthy SD rats [27]. Articular cartilage was disinfected by antibiotics and further washed with PBS before using for explant culture. Cartilage tissues were chopped and placed in DMEM+ F12 media for incubation for 24 h (37 °C, 5% CO2). After incubation, media was removed and replaced with following treatments: untreated culture media (control), IL-1β (10 ng/ml) and IL-1β with most effective doses of SOE at 500, 250, and 125 μg/ml. Explants were incubated (37 °C, 5% CO2) for 6 days and collected the supernatants for evaluation of proteoglycan release., The Lipid peroxidation inhibition activity
Malon-dialdehyde (MDA) assay was used to determine the lipid peroxidation inhibition effect of SOE as described by a previous study [24]. Briefly rat liver tissues (2.0 g) were sliced and homogenized in 10 ml 15 mM KCl-Tris-HCl buffer (pH 7.2). The reaction solution (0.25 ml liver homogenate, 0.1 ml Tris-HCl buffer (pH 7.2), 0.05 ml 1 mM ascorbic acid, 0.05 ml 4 mM FeCl2) and 0.5 ml of plant extract of different concentration of SOE were taken in a tube. The reaction tube was incubated at 37 °C for 1 h. After incubation, 0.5 ml 0.1 N HCl, 0.2 ml 9.8% sodium dodecyl sulphate, 0.9 ml distilled water and 2 ml 0.6% thiobarbituric acid (TBA) were added to each tube and vigorously shaken. Then, the tubes were placed in a boiling water bath at 95 °C for 30 min. After cooling, the flocculent precipitate was removed by adding 5 ml -butanol, mixed well, and centrifuged at 2500 RPM for 10 min. The absorbance (ABS) of the supernatant was measured at 532 nm and percentage of lipid peroxidation by SOE was measured by following equation [25]. Lipid peroxidation inhibition (%) = [Acontrol – Asample / Acontrol] × 100, where Acontrol is absorbance of control and A-sample is absorbance of sample (SOE extract).
metachromatic dye 1,9 dimethyl methylene blue (DMMB) was used to quantify the amount of released proteoglycan especially sulphated glucosamineglycan’s (GAGs). DMMB activity was measured according to the previously developed protocol with little modification. On a 96-well plate, 1:1 sample and DMMB solution were added to each well, before reading the plate, it was kept in dark for 10 min then OD was measured by spectrophotometer at 540 nm. OD of DMMB is directly proportional to the release of proteoglycans [28].

**Western blot analysis**

Rat articular chondrocyte were grown in complete media with and without treatment with 3-Methyl-6,7-(methylenedioxy) quercetagetin and 3-O-Methylpatuletin for 48 h. Cells were washed in chilled PBS and lysed by lysis buffer from Sigma Aldrich (St. Louis, MO, USA) containing a protease inhibitor mixture Sigma Aldrich (St. Louis, MO, USA). 30 μg of protein lysate were loaded in 12% polyacrylamide gel, followed by transfer on PVDF membrane. Membrane blocking was done by 5% BSA (bovine serum albumin) and then incubated with primary antibody at 4 °C overnight. Membrane were then washed and probed with horseradish peroxidase-conjugated secondary antibody (abcam®, Cambridge, USA) and visualized by Chemiluminescence system (LS4000 GE Healthcare Life Sciences, India) [29].

**Induction of osteoarthritis (OA) and experimental design**

The aim of this study was to assess the effect of SOE on MIA induced OA in the biological system of Sprague-dawley rats. For this experiment, 55 adult healthy and pathogen free female rats were used. Rats were divided into four groups (n = 10 in control group and n = 15 in each group of MIA induced rats. Adult healthy and pathogen free female Sprague-dawley rats, 14–16 weeks of age, (180 to 200 g) were obtained from the National Laboratory Animal Centre, Lucknow India. Animals were kept in a 12 h light-dark cycle, five animals in each steel cage with controlled temperature (22–24 °C), humidity (50–60%) and food and drinking water was supplied ad-libitum. This experiment was carried out in the Animal Centre of the central drug research institute for 28 days.

Rats were anesthetized by cocktail of Ketamine and Xylazin (9:1 ratio). After anesthetization all animals except control group received a single injection of MIA through the intra-articular joint of the left knee with 3 mg MIA in 50 μl saline [30]. The control group animals were injected with 50 μl of physiologic saline in their left knee [31]. All animals were closely observed every day and when primary symptoms like swelling in knee and walking disability was observed at the end of 3 days. These animals (N = 10, Total 30) were further selected for treatment of SOE and MIA control. SOE dispersed in 1.0% gum acacia in 250 or 500 mg.kg⁻¹ day⁻¹ doses and was given orally for 28 days once in a day. The control groups received only 1.0% gum acacia orally. After 28 days of treatment animals were euthanatized by CO₂ exposure in the morning at laboratory.

**Histological analysis**

Long bone tissues were isolated from differentially treated rats. Dissected tissues were cleaned by removing muscles and fixed in 4% formaldehyde. Tibia bone was decalcified in EDTA solution [32]. Sections of width 5.0 μm were cut by Leica RM2265 rotatory microtome with the help of sharp blade. Sections were stained with Safranin O and Toudine Blue. Photomicrographs of stained sections were captured using microscope (Evos XL, Life technologies).

**Micro computed tomography (micro-CT) at femoral condyles and tibia plateau region**

The effects of SOE on micro-architectural deterioration in MIA induced osteoarthritis model were evaluated with micro computed tomography (Sky Scan 1076 scanner; Sky Scan, Aartselaar, Belgium). Femoral condyles and tibial plateau regions were scanned. Bones of all animals of each group were scanned at voxel size of 18 μm, at a voltage of 70 kV, a current of 142 mA, field of view of 35 mm, by using filter 1.0 mm aluminium plate, 0.8-degree rotation step with full width [33]. Total tissue volume (TV), trabecular bone volume (BV/TV, %), trabecular number (Tb.N, mm⁻¹), and trabecular separation (Tb.Sp, mm) and trabecular bone pattern factors (Tb.pf) (mm⁻¹), were calculated by BATMAN software provided with the μCT machine [34].

**Quantitative real-time polymerase chain reaction analysis**

The bones were excised during autopsy, removed attached tissue on bone, cleaned and collected in RNA later. For RNA isolation, bones were cut and upper head part containing cartilage and subchondral region were taken for genes expression study. First, isolated bone tissues were cleaned by removing muscles and fixed in 4% formaldehyde. Tibia bone was decalcified in EDTA solution [32]. Sections of width 5.0 μm were cut by Leica RM2265 rotatory microtome with the help of sharp blade. Sections were stained with Safranin O and Toudine Blue. Photomicrographs of stained sections were captured using microscope (Evos XL, Life technologies).

**Open-field activity test**

The locomotor activity was examined on 28th day post-surgery in a sound proof room by Opto-Varimex-5
Table 1 Primer sequences of various rat genes used for qPCR

| Oligo name | Sequence (5’ to 3’) |
|------------|-------------------|
| COL-1      | F: CAT.GTT.CAG.CCT.TGT.GGA.CCT (GGA.CCT) R: GCA.GCT.GAC.TCT.GGA.GAT.GT |
| IL1β       | F: TGT.GAT.GAA.AGA.CGG.CAC.AC R: CTT.CCT.TTG.GTA.TTG.TGT.GG |
| Acan       | F: AAT.GGG.AGG.CAG.CCT.TCA.AC R: TTG.AGA.GCC.AGA.GGG.ACT.TT |
| SOX9       | F: TGA.AGA.GAG.AGA.CGG.AGA.AA R: CAT.GAC.CCT.TCA.GCA.ACT.GT |
| COL10a1    | F: CAC.AGC.CAT.TTC.GAG.CTT TT R: TCT.AAG.TTG.CCC.CAG.GTA.CG |
| MMP13      | F: TGG.ACA.AGC.ATC.TCC.AAA.G R: GTC.CAG.GCC.AGA.GGT.GT |
| MMP9       | F: CCT.CGT.CAT.GAA.GAC.GAT.AA A R: GGT.CAG.CTT.TAG.GGC.CAC.GA |
| MMP1a      | F: GGT.GAT.AAT.GTG.TTC.GCC.TTC R: TCA.GGT.CCA.TCA.AAT.GGT.GTA |
| TNF-α      | F: TGA.ACT.TCG.GGG.TGA.TCG R: GGG.GTG.CTC.ACT.CGA.GTT.TT |
| TIMP-1     | F: CAG.CAA.AAG.CCC.TTC.GTA.AA R: TCG.TGT.AAC.GAG.GGA.ACT |
| COL2a1     | F: CCA.GGT.CCT.GGT.GGA.AAA R: CCT.CCT.TCC.GGC.CCT.TT |
| BMP-2      | F: CCC.CTA.TAT.GCT.GGA.CCT.GT R: AAG.CTT.CCT.CGA.TGG.CCT.TT |
| TIMP-2     | F: CTT.TTT.CCA.ATG.CAG.ACG.TA R: GAT.GGG.GGT.GCC.ATA.GAT.GT |
| B-αCTIN    | F: CCC.GCG.AGT.ACA.ACC.TTC.TT R: CGT.CAT.CCA.TGG.CGA.ACT |

F: Forward Primer, R: Reverse Primer

(Columbus Instruments, USA) which was consist of Plexi glass transparent cages (42.5 cm X 42.5 cm arena size) with a grid of infrared emitters and detectors in order to automatically detect activity measures [37].

Rota Rod test

Rats were individually placed on rotating rod (7.5 cm diameter) and trained through three different sessions on three consecutive days (one session/day), whereby each session included four separate training trials. Rats were trained at 5 rpm on day 1, at 10 rpm on day 2 and at 15 rpm on day 3. At the end of training sessions, animals were tested to ensure that they could maintain themselves on rotating rod for 300 s. On 28th day post-surgery, each group of animals was subjected to Rota rod test to evaluate the latency to fall recorded [37].

GSTs (glutathione S-transferases) activity in serum

GST activity of control, MIA and different dose of SOE was measured in serum by glutathione S-transferase assay kit (Cayman USA, 703,302) according to manufacturer protocols. Briefly, 150 μl assay buffer provided with kit, 20 μl glutathione, 20 μl serum sample and finally add 10 μl 1-chloro-2, 4-dinitrobenzene (CDNB). This reaction mixture was carefully shacked and read at 340 nm by ELISA plate reader (molecular devices, USA) [38].

Elsa

Urinary CTX-II (C-telopeptide of type II collagen) [39] and serum COMP (cartilage oligomeric matrix protein) [40], are biochemical markers for uses in progression of joint destruction. Urinary CTX-II and serum COMP concentrations were determined after 28 days treatment of SOE by enzyme-linked immunosorbent assay kits as per manufacturer’s protocols [41].

Statistical analysis

All data is represented as Mean ± standard error of the mean (Mean ± S.E.M). Group differences were determined using one-way analysis of variance (ANOVA) with Newman–Keuls post hoc tests using Prism version 5.0. software. Probability values of p < 0.05 were considered to be statistically significant. Data for DPPH, ABTS, LPI, DMMB, MTT n = 4 and all assays were replicated three times. Data for qRT-PCR n = 3, serum/urine parameter n = 6 and micro CT analysis n = 10 samples were taken for analysis. Results were obtained from minimum three independent experiments in triplicate and are expressed as Mean ± S.E.M. For significance, P < 0.05 was used in each group, and results were reproducible and there were no disagreements amongst the blinded assessors (RT, PRM and RM).

Results

Cell viability assay of SOE

Cell viability after treatment with different concentrations of SOE was determined by MTT assay on rat articular chondrocytes cells. Data in (Fig. 1a) shows that the SOE concentration from 1.95 to 1000 μg/ml was safe for use and did not alter cells viability at any dose examined in this study. In addition, we evaluated SOE efficacy in acute toxicity rat model as shown in HE staining (see Additional file 1: Figure S1). We also assessed biochemical parameters ALT and AST as liver health indicators (see Additional file 2: Figure S2) and its ‘materials and method’ is given in text file (Additional file 3).

Free radical scavenging activity of SOE

We next assessed the antioxidant activity of SOE at different doses using the DPPH free radical scavenging method. SOE at concentrations between 1.95 to 1000 μg ml−1 showed significant scavenging activity. Maximum scavenging activity by SOE was 17.52%, 23.02% and 23.65% at doses of 250, 500 and 1000 μg.ml−1 respectively (Fig. 1b). Ascorbic acid was taken as positive control as shown in Additional file: 4: Figure S3.
Ex-vivo lipid peroxidation inhibition activity of SOE
Lipid peroxidation inhibition activity was measured ex-vivo by determining the malondialdehyde (MDA) in rat liver homogenate. Inhibition of lipid peroxidation indicated anti-oxidant property of plant extracts. Experiment showed that SOE exhibits MDA inhibition at 250, 500 and 1000 μg.ml⁻¹ by 18.05%, 21.53% and 23.28% respectively (Fig. 1c).

Free radical cation scavenging activity of SOE
ABTS radical scavenging method was adopted for the determination of antiradical and antioxidant activity of SOE and the obtained results are showed in Fig. 1d. We checked anti-oxidant potential of SOE for the concentration range from 15.6 to 1000 μg.ml⁻¹. We observed maximum anti-oxidant potential of SOE at 250, 500 and 1000 μg.ml⁻¹ that was 24.23%, 27.37% and 28.95% respectively (Fig. 1d).

Effect of SOE on GAG release
To assess the preventive effect of SOE we next assessed the glycosamine glycan (GAG) release in conditioned media of IL-1β treated rat articular chondrocytes as IL-1β activates MMPs that cleave the extracellular matrix components of the cartilage. After 6 days of culture in DMEM, we observed 27.75% increase of GAG release in media in the presence of IL-1β as compared to the control. Experiments
in the presence of SOE (IL-1β + SOE) at concentration 125 μg.mL⁻¹, IL-1β + SOE at concentration 250 μg.mL⁻¹ and IL-1β + SOE at concentration 500 μg.mL⁻¹ inhibited GAG release by 8.34%, 11.84% and 16.38% respectively. This experiment suggests that SOE has the ability to prevent the destruction of the cartilage as caused by IL-1β-stimulation and thus GAG release in the media (Fig. 1e).

**Active components of SOE increase chondrogenic proteins**

Treatment with 3-Methyl-6, 7-(methylenedioxy) quercetagetin and 3-O-Methylpatuletin two active compounds isolated from SOE increased protein expression of Sox 9 and Col-2 the chondroprotective markers in articular chondrocytes (Fig. 2) suggesting that the prevention of cartilage damage as observed by SOE administration may have been the contribution of these pure compounds. HPLC data of SOE and its active components are exhibited in Additional file 5: Figure S4.

**Effect of SOE on articular cartilage and subchondral bone**

Figure 3a represents the picture of gross morphological changes at the knee joint after 4 weeks of injection of MIA into the left knee. Grossly, the normal joints revealed smooth and shiny articular surfaces. The knees of the MIA-injected group showed irregular abrasions at the articular cartilage surfaces of the femoral condyle and the tibial plateau. Figure 3 represents the macroscopic photographs of the damaged articular cartilage of the femur condyles (B) and tibial plateau (C) after injection of MIA. OA was characterized by progressive cartilage destruction and alterations in subchondral bone structure. Figure 3d represents radiographies of the left knee joints. Radiographs of the control group showed normal state of the knee characterized with smooth articular surfaces. Due to MIA intraarticular injection, joint represents with complete loss of joint space, irregularity in articular surfaces and atrophied femoral condyles and tibial plateau. Interestingly, the treatments with 250 and 500 mg.kg⁻¹ day⁻¹ doses of SOE for four weeks restored the cartilage morphology. Further, after 4 weeks of MIA injection, μ-CT analysis revealed a significant loss in cartilage and subchondral bone parts in MIA group (Fig. 3e).

In rat model of OA, the micro-CT analyses showed that treatment with 250 and 500 mg.kg⁻¹ day⁻¹ of SOE suppressed osteoarthritis progression.

Further, detailed analysis of tibial bone by microcomputed tomography (μ-CT) indicated that MIA degrades cartilage substantially relative to sham-operated group (Fig. 4a). Histological analysis by safranin O (Fig. 4b) and toluidine blue (Fig. 4c) showed that MIA injected animals undergo maximum loss at articular region as compared to saline treated control animals with notable proteoglycan loss and disappearance of chondrocytes at the deep zone of articular cartilage. Figure 3b and Fig. 3c reflect intact articular cartilage in control group with SOE reducing the MIA-induced cartilage degradation. We observed that the dose of 250 mg.kg⁻¹ day⁻¹ of SOE had less effect on the structural changes in the joints as compared to the 500 mg.kg⁻¹ day⁻¹ dose. The 500 mg.kg⁻¹ day⁻¹ dose resulted in restoration of hyaline cartilage which is revealed by histology of articular cartilage.

Osteoarthritis Research Society International (OARSI) scores revealed degeneration of articular cartilage that was induced after MIA injection and progressed gradually (Fig. 4d).

OA is primarily associated with damage of articular cartilage that leads to increased bone remodelling followed by subchondral bone deterioration with the progression of the disease. Results of μ-CT at the subchondral bone site show that MIA induction increases tissue volume (TV) by 160% and decreased subchondral bone volume/tissue volume (BV/TV) ratio by 58% as compared to sham-operated controls. Increase in subchondral TV after MIA induced OA ultimately led to the reduction of the total BV/TV. Moreover, MIA disrupts the connectivity and microarchitecture of subchondral bone which was indicated by a significantly higher trabecular pattern factor (Tb.Pf) (Fig. 4e-g). Treatment of SOE at 500 mg.kg⁻¹ day⁻¹ treatments decreased the TV by 48% and increased the BV/TV by 79% that resulted in a significant decrease in Tb.Pf by more than 200% after SOE administration. We also assessed μ-CT parameters at femoral condyles (see Additional file 6: Figure S5).

We further assessed the trabecular bone by 3D-μ-CT that clearly showed that SOE has potentially lessened the harmful effects of MIA as indicated in the representative images Fig. 5a. MIA significantly decreased number of trabecules in bone as assessed by Tb.N (Fig. 5b) that leads to disruption of the trabecular connectivity and higher trabecular separation (Tb.Sp) in the MIA rats (Fig. 5c). The
SOE dose of 250 mg kg\(^{-1}\) day\(^{-1}\) reversed the MIA induced effects with increased Tb.N by ~58\% (Fig. 5b) and decreased, Tb.Sp by ~17\% (Fig. 5c) as compared to MIA group. On the other hand, the higher dose of 500 mg kg\(^{-1}\) day\(^{-1}\) maintained the articular cartilage by increasing Tb.N by ~60\% and decreasing Tb.Sp by ~26\% compared to MIA group. Inter-dose comparisons show that SOE at 500 mg kg\(^{-1}\) day\(^{-1}\) was more effective in rescuing articular and sub-chondral bone loss.

**Open-field activity test and Rota rod test after SOE administration**

The protective effect of SOE on the cartilage as observed in the above experiments was concluded on the basis of the
behavioural experiments of locomotion. We observed that MIA treated rats significantly travelled for shorter distance in open-field area experiment (Fig. 5d) and decreased latency to fall as seen in Rota rod test (Fig. 5e), as compared to control group. This could have been because of the articular joint inflammation and destruction as observed by us or due to osteoarthritic pain. Observations showed that 500 mg.kg$^{-1}$ day$^{-1}$ SOE displayed significant improvement in distance travelled and increased in latency to fall in Open-field and Rota rod test respectively, when compared with MIA group. Although we observed positive changes with the lower dose (Fig. 5d and e) of SOE but the overall effect was more pronounced with the 500 mg.kg$^{-1}$ day$^{-1}$ dose.
Effects of SOE on the expression of pro-inflammatory cytokines and chondrogenic genes in articular cartilage region of knee joints

At molecular level expression of pro-inflammatory and chondrogenic genes in isolated cartilage from the knee region of control, MIA and SOE administered animals were investigated. The progressive destruction of articular cartilage is caused by MIA injection that significantly upregulated the expression of pro-inflammatory genes interleukin-1 beta (IL-1β), tumor necrosis factor alpha (TNFα), and matrix-degrading enzymes (Fig. 6a-f). IL-1β and TNFα are considered as the major cytokines, involved in the onset and pathogenesis of OA. MIA increased expression of IL-1β by ~6.0 folds (Fig. 6a) and TNFα by ~7.0 folds (Fig. 6b). This led to an increase in expression of collagen 10 (COL10) (Fig. 6c) that increases hypertrophy like environment in the articular cartilage region. These cytokines further damaged the joints by enhancing the expression of a group of metalloproteinases like interstitial collagenase (MMP-1), stromelysin-1 (MMP-3), and collagenase 3 (MMP-13) (Fig. 6d-f). In addition to these harmful effects, MIA blocked chondrogenic activity by down regulating the synthesis of ECM components, like sex determining region Y-box 9 (SOX9) (Fig. 7a), bone morphogenetic protein 2 (BMP2) (Fig. 7b), collagen type-II (Col2) (Fig. 7c), and aggrecan (Fig. 7d). Further, tissue inhibitors of matrix metalloproteinases (TIMPs) are recognized as endogenous protease inhibitors of MMPs. MIA decreased the expression of TIMP1 (Fig. 7e) and TIMP2 (Fig. 7f).

When we administered SOE at 250 mg.kg⁻¹ day⁻¹ and at 500 mg.kg⁻¹ day⁻¹ we observed reversal of the deleterious effects of MIA. The dose of SOE at 250 mg.kg⁻¹ day⁻¹ significantly decreased the

![Fig. 5 SOE improves micro-architectural and behavioural parameters.](image-url)

- **a** Representative 3D μCT images of tibia plateau region of different groups.
- **b** Trabecular number (Tb.N, 1/mm); (c) Trabecular Separation; (Tb.Sp, mm) (d) Open-field activity and (e) Rota rod test to examine the locomotor function and motor coordination. All values are expressed as Mean ± S.E.M. (n = 6/group); *P < 0.05, **P < 0.01, ***P < 0.001 compared to the MIA group, ###P < 0.001 compared to the 250 mg.kg⁻¹ day⁻¹ dose, NS= non significant.
expression of IL-1β by ~46%, COL10 by ~40%, MMP1 by ~22% and MMP3 by ~40%, and moreover, upregulated the expression of SOX9 by ~103%, aggrecan by ~52%, and TIMP2 by ~56% as compared to MIA group. When a higher dose of SOE was used we observed an inhibition by ~50% of the expression of inflammatory cytokines like IL-1β, TNFα, COL10, MMPs and restoration of the cartilage by upregulation of the extra-cellular matrix components by more than 100% in SOX9, BMP2, COL2, AGGREGAN, TIMP1, TIMP2. Inter-comparison between results associated with two different doses of 250 and 500 mg.kg⁻¹ day⁻¹ of SOE, we observed that the 500 mg.kg⁻¹ day⁻¹ dose was more effective for restoration of the cartilage as a therapeutic treatment.

Glutathione S-transferases (GSTs) activity in serum
An increased serum GST pool of ~20% was found in serum of MIA injected group which indicated that MIA animals produced more GST activity to detoxify itself in response to the harmful effects of MIA. Oral administration of SOE decreased the GST activity by ~34% at 250 mg.kg⁻¹ day⁻¹ and by ~30% at 500 mg.kg⁻¹ day⁻¹ with respect to MIA group. No significance was observed within the SOE administered groups (Fig. 8a).
SOE treatment decreases cartilage turnover marker in biological fluids

MIA increased the levels of serum COMP and urinary CTX-II, which are associated with progression of OA. SOE at 250 and 500 mg.kg$^{-1}$ day$^{-1}$ administration to MIA injected animals lowered the CTX-II and COMP levels as compared to MIA group however significant changes were observed at 500 mg.kg$^{-1}$ day$^{-1}$ (Fig. 8b, c). The clinical markers highlight the potential of SOE in preventing loss of articular cartilage and subchondral bone as potential anti-osteoarthritic agent.

Discussion

In this study, SOE was evaluated at physiological doses for its effect on MIA induced osteoarthritis in rodents. Chondrocytes are the only component cells that are capable of controlling vital activities of the articular cartilage [42]. Therefore, the saving of plausible number of cartilage cells in the joint’s articular structure of SOE treated group was not only interesting but also served as an important finding in OA pathogenesis and progression. Previous studies reported that inflammatory cytokines, chemokines, and...
other inflammatory markers are found higher in osteoarthritis patients [43].

Till date, complete cure for OA remains elusive and the management of OA is largely palliative focusing on the alleviation of symptoms only. Current recommendations for the management of OA include a combination of pharmacological (NSAIDs, DMOADs, paracetamol, etc.) and non-pharmacological interventions (weight loss and exercise). Pharmacological drugs include non-steroidal anti-inflammatory drugs (NSAIDs) that have analgesic and antipyretic actions, with risk of an upper gastrointestinal (GI) event like ulcer, perforations, bleeding and obstruction. Disease modifying OA drugs (DMOADs) are currently in clinical trial phase (like matrix metalloproteinase inhibitors, Fibroblast growth factor 18 (FGF-18), Interleukin 1 inhibitors, Inducible nitric oxide synthase) and long term safety of these drugs is still unclear [44].

Currently, use of herbal products for medicine and dietary supplements or as complementary and alternative medicine have become an important area of research in bone health especially orthopaedics, arthritis, rheumatic diseases, and musculoskeletal [45]. Herbal products have lot of therapeutics and dietary uses with low risk of side effects and toxicity [46–51]. Therefore, in current study, efforts are being made to elucidate the role of natural product like SOE for the prevention and cure of OA. In this regard, this study first evaluated the anti-oxidant and anti-inflammatory effects of SOE. ABTS and DPPH assays indicates that SOE have anti-oxidant and free radicle scavenging activity without using animals showing as a marker of preliminary identification of characterization of novel agent for treatment of OA.

Results taken together by DPPH, ABTS, and LPI experimental data revealed the antioxidant nature of SOE. Furthermore, when cartilage tissue was exposed to IL-1β, it underwent degradation and consequently released GAG, which is a typical clinical symptom of OA [52]. While treating with SOE, the released GAG was significantly reduced as measured by DMMB assay and showed anti-inflammatory potential of SOE.

Further, anti-osteoarthritic activity of SOE was studied under in vivo conditions by making MIA induced osteoarthritic rat model, which resembles the phenomena observed in human OA of knee region and by increasing disease conditions like free radical generation and inflammation [17]. The MIA used in the present study to induced OA condition as MIA suppression of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) that results in a reduction of glycolysis activity in the cartilage chondrocytes which further cause pathological structural/morphologic changes in articular cartilage. The MIA caused histopathologic changes in articular cartilage structure which was confirmed by safranin-o and toluidine blue staining. Histological and ORASI data indicated that SOE treatment prevents the degradation of chondrocytes and extracellular matrix (ECM) components in articular region. Previous studies support that both the cartilage and subchondral bone communicate to each other through biomechanical and biochemical signalling pathways for maintaining homeostasis of the joint environment. Moreover, defects in articular cartilage increase various inflammatory cytokines and decrease the chondrocytes synthesis that further leads to increased bone remodelling and subchondral bone deterioration. Radiography, confirmed the above gross findings of SOE that were corroborated and confirmed quantitatively with micro-CT. It was found that 4 weeks post-treatment with SOE sustained the changes found especially
in the joint spaces. These changes with SOE were brought about by improving the cartilage by increasing total bone volume to tissue volume and trabecular number that provide structural and functional strength to the cartilage. Whereas, in case of those treated with MIA, radiographs showed increased inter condylar fossa, severe joint space loss, atrophied femoral condyles and lack of articular surface smoothness. Impairment of joint space is an important feature in diagnosis and assessment of OA. MIA significantly decreased duration of rotation and balancing on moving rod that are related to inflammation swelling, pain and stiffness in joint whereas, SOE treatment showed increased joint mobility with increased travelling distance and decreased chance of latency to fall, suggesting its role in lessening pain and its associated symptoms.

Pathophysiology of OA, is also associated with increased inflammatory cytokines associated with the disruption of homeostasis of ECM component in OA as it is often subjected to target high mechanical load tissues such as joints by disturbing the catabolic and anabolic processes [53]. The cytokines like IL-1β and TNFα produced by the chondrocytes after defect in articular cartilage, especially IL-1β, play a significant role in the further degradation of cartilage. IL-1β and TNFα are the main cytokine instigators of cartilage degeneration in arthritis, and these induce MMP in chondrocyte cells [54]. Molecular analysis by qRT-PCR revealed that imbalance between the cartilage anabolic and inflammation related genes might account for the advanced condylar destruction and chondrocytic apoptosis following MIA induction that is balanced by SOE. MIA increased the size of articular chondrocytes by an increase in expression of Col10 that leads to upregulated osteogenic environment in the articular region and leads to progression of OA by subchondral bone deterioration. SOE treatment maintained chondrogenic environment of articular chondrocytes by decreasing the expression of Col10 in the joints. As shown by data, MIA affected the joints by enhancing the expression of a group of metalloproteinases, which have a destructive effect on cartilage. These metalloproteinases block chondrocyte activities by down regulating the synthesis of ECM components, interfering with the synthesis of the key structural proteins such as collagen type-II, Sox9 and aggrecan [55]. This was corroborated by the treatment of isolated pure compounds from SOE to rat articular chondrocytes suggesting that the active components present in SOE may be contributing to prevention from the damaging effect of MIA. For better understanding, the schematic diagram is shown (see Additional file 7: Figure S6) in which SOE exhibits chondroprotective effects on subchondral bone and causes the shifting of chondrocytes and cartilage homeostasis towards anabolism.

In addition, SOE treatment remarkably reduced the elevated serum level of GST, COMP and urinary CTX-II associated with MIA in a dose-dependent manner. These findings indicated that SOE treatment decreases the MIA induced inflammation, articular and sub-chondral bone loss. This study indicates that SOE has anti-osteoarthritic and chondro-protective effects in chemically induced osteoarthritis. However, confirmation of the effect of SOE on OA as observed by us further requires validation in higher animals and primates for extrapolation of data to humans.

Conclusions
On the basis of in vitro and in vivo experiments performed with Spinacea oleracea extract and it’s isolated compounds we can deduce that SOE has the ability to alleviate MIA induced deleterious effects.

Additional files

Additional file 1: Figure S1. H&E stained organ sections, isolated from rat after treatment of SOE (acute toxicity study). No noticeable abnormality was observed in major organs including kidney, liver, and spleen. (TIFF 9629 kb)

Additional file 2: Figure S2. Serum ALT and AST level in different groups after 28 days of treatment of SOE. No significant differences in control, MIA, 250 mg/kg, 500 mg/kg. All values are expressed as Mean ± S.E.M (n = 4/group). (TIFF 7976 kb)

Additional file 3: Supplementary information. (DOCX 16.5 kb)

Additional file 4: Figure S3. Ascorbic acid was used as positive control in both DPPH and ABTS assay (a) Ascorbic acid has maximum scavenging activity from 7.81 μg/ml to 1000 μg/ml in constant manner. (b) Minimum scavenging activity was found at 15.62 μg/ml and it was increased in concentration dependent manner. All values are expressed as Mean ± S.E.M (n = 4/group). (TIFF 6.90 mb)

Additional file 5: Figure S4. HPLC data for the SOE and identified compound. (TIFF 1.47 mb)

Additional file 6: Figure S5. 3D images of femoral condyle bone obtained from Micro-CT and their parameters. (TIFF 9.40 mb)

Additional file 7: Figure S6. On the basis of molecular changes, histology, and micro-CT, it is concluded that SOE shows chondro-protective effects on subchondral bone and causes the shifting of chondrocytes and cartilage homeostasis towards anabolism. (TIFF 709 kb)

Abbreviations
ALT: Alanine amino-transferase; AST: Aspartate amino-transferase; BMP2: Bone morphogenetic protein 2 (Fig. 6B); Col10: Collagen type 10; Col2: Collagen type-II; COMP: Cartilage oligomeric matrix protein; CTX-II: C-telopeptide of type II collagen; GSTs: Glutathione S-transferases; IL-1β: Interleukin-1 beta; MMP: Matrix metalloproteinases like interstitial collagenase; MMP-9: Matrix metalloproteinases collagenase 3; MMP-3: Matrix metalloproteinases stromelysin-1; Oxidative stress: ROS: Reactive oxygen species; SOE: Spinacea oleracea extract; SOX9: Sex determining region Y-box 9 (Fig. 6A); TIMPs: Tissue inhibitors of matrix metalloproteinases; TNFα: Tumor necrosis factor alpha

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Availability of data and materials
All the details of data and materials of this study are included in the manuscript.

Authors’ contributions
Conceived and designed the experiments by RT, RM, PRM and DC. RT and CPCSEA), Ministry of Environment, Forest and Climate Change, Animal Control and Supervision of Experiments on Animals (34/GO/Re-SL-Bi-S/99/ CPCSEA), Ministry of Environment, Forest and Climate Change, Animal Welfare Division, Government of India guidelines.

Ethics approval
All animal care and experimental procedures were approved by the Institutional Animal Ethics Committee (IAEC, Central drug research institute) and performed according to the regulations of the Council for the Purpose of Control and Supervision of Experiments on Animals (34/GO/Re-SL-Bi-S/99/ CPCSEA), Ministry of Environment, Forest and Climate Change, Animal Welfare Division, Government of India guidelines.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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Author details
1Division of Endocrinology, CSIR-Central Drug Research Institute, Lucknow 226031, India. 2Division of Medicinal and Process Chemistry, CSIR-Central Drug Research Institute, Lucknow, India. 3Division of Pharmacology, CSIR-Central Drug Research Institute, Lucknow, India. 4Division of Pharmaceutics, CSIR-Central Drug Research Institute, Lucknow, India. 5Academy of Scientific & Innovative Research (AcSIR), CSIR-Central Drug Research Institute, Lucknow 226031, India.

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