Towards prevention of post-traumatic osteoarthritis: report from an international expert working group on considerations for the design and conduct of interventional studies following acute knee injury


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Towards prevention of post-traumatic osteoarthritis: report from an international expert working group on considerations for the design and conduct of interventional studies following acute knee injury


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Osteoarthritis (OA) pathologically represents a continuum from risk exposure, to molecular changes and structural changes with associated pain, which for some people progresses to the need for joint replacement. Detection and treatment of those at high risk of OA could enable effective interventions before any major structural damage has occurred or before pain becomes chronic, that is at a pre-radiographic or even pre-symptomatic stage. Such intervention would be comparable to current early management of diabetes, cardiovascular disease, osteoporosis or pre-rheumatoid arthritis.

Joint injury remains one of the biggest risk factors for OA. In Sweden, approximately 80/100,000 people per year experience anterior cruciate ligament (ACL) rupture; in the U.S. there are 252,000 ACL injuries per year\(^1\). 50% of people with significant knee joint injuries such as ACL rupture and/or acute meniscal tear subsequently develop symptomatic radiographic OA within 10 years, so-called post-traumatic OA (PTOA)\(^1\); at least 33% with acute ACL rupture will have magnetic resonance imaging (MRI)-defined whole joint OA after 5 years\(^1\). PTOA is thought to comprise around 12% of all OA, although its incidence appears to be increasing\(^5,6\).

Whilst preventing joint injury is an intervention to prevent PTOA\(^1\), our focus was on interventions after knee joint trauma. We conducted an evidence review, then consensus process developing considerations and a research agenda. Though the evidence review summarized the use of outcome measures including patient-reported outcome measures (PROMs), no recommendations for specific outcome measures were planned.

Evidence review

An evidence review was conducted to identify experimental, interventional studies following acute knee injury with specific reference to post-traumatic knee OA. Systematic searches were conducted across five databases (Cochrane Library; EMBASE; MEDLINE; CINAHLPlus; AMED) from inception to August 2016. The search strategy was designed in OVID-Medline using text words and subject headings (MeSH), combining terms for knee injury, osteoarthritis and clinical trials or systematic reviews (Supplementary Table 1).

All references were imported into EndNote where duplicates were removed. Screening and study detail extraction was by NC, verified by three others (FW, DM, PC). Study inclusion criteria were as follows: population clearly stated within 6 months of acute knee injury (any setting); interventional study (any intervention, including surgical, pharmacological, non-pharmacological) with any comparator (including active, placebo, sham or no

Methods

The considerations process was facilitated by the Osteoarthritis and Crystal Diseases Clinical Studies Group of Arthritis Research UK (UK’s largest arthritis charity), which was established to develop consensus research priorities and nurture methodologically robust clinical trials.

Conclusions: These considerations represent the first international consensus on the conduct of interventional studies following acute knee joint trauma.

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Consensus group

A group of 32 stakeholders, including physiotherapists, orthopaedic surgeons, rheumatologists, sports and exercise medicine physicians, primary care physicians, radiologists, laboratory scientists, statisticians, clinical trialists, engineers, pharmaceutical company experts and four patient representatives (2 who had a previous knee joint injury) comprised the consensus group. After the evidence review results were circulated, the group convened at a face-to-face meeting. The evidence review, which included a summary of the evidence search or considerations, as they were felt to be prone to bias and not representative of our main focus which related to experimental studies.

Results

Evidence review

The initial search identified 2476 citations (MEDLINE, n = 532; EMBASE, n = 863; CINAHLPlus, n = 489; AMED, n = 60; Cochrane Library, n = 532). 945 duplicates were removed. Screening of the remaining 1531 abstracts yielded 43 eligible studies. Seven systematic reviews identified a further 15 reported trials. From these 58 papers (including 11 conference abstracts), 37 unique studies were included. Details of each study are summarized in Supplementary Table 2. The majority of studies involved ACL injury (n = 20; 54%), patellar dislocation (n = 8; 22%) or tibial plateau fracture.

Table 1

Overview of basic study details, categorized according to type of knee injury

<table>
<thead>
<tr>
<th>Number of studies [papers if different]</th>
<th>ACL</th>
<th>Patellar Dislocation</th>
<th>Tibial plateau fracture</th>
<th>Other</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>RCT/nRCT</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RCT: 20 incl. 1 protocol and 1 pilot</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>nRCT: 1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>RCT: 6 incl. 1 protocol nRCT: 1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RCT: 2 incl. 1 pilot</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RCT: 35 incl. 2 pilots, 2 protocol; nRCT: 2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

| Sample size at randomization           |     |                      |                         |       |       |
| Missing                                | 1   |                      |                         |       | 1     |
| 20-50                                  | 2   |                      |                         |       | 3     |
| >50-100                                | 5   | 5                     | 4                       |       | 14    |
| >100-200                               | 7   | 2                     | 2                       |       | 11    |
| Power calculation                      |     |                      |                         |       |       |
| a priori                               | 5   | 2                     | 2                       |       | 16    |
| post hoc                               | 2   | 1                     | 3                       |       | 3     |
| None                                   | 8   | 3                     | 1                       |       | 14    |
| Unclear                                | 1   | 2                     | 4                       |       | 4     |
| Study adequately powered               |     |                      |                         |       |       |
| (based on sample size)                 |     |                      |                         |       |       |
| Missing                                | 9   | 4                     | 5                       |       | 15    |
| <3 months                              | 1   | 1                     | 2                       |       | 2     |
| 3-6 month                              | 3   | 3                     | 3                       |       | 3     |
| >6 months-1 year                       | 3   | 3                     | 1                       |       | 7     |
| >1-2 years                              | 4   | 3                     | 1                       |       | 7     |
| >2-5 years                              | 3   | 2                     | 5                       |       | 5     |
| >5-10 years                            | 2   | 3                     | 5                       |       | 5     |
| >10 years                              | 4   | 1                     | 6                       |       | 6     |
| Primary outcome measure(s) clearly defined | 9 + 1 used for sample size calculation | 5 | 2 + 1 used for sample size calculation | 1 | 17 + 2 based on sample size calculation |
| Type of interventions                  |     |                      |                         |       |       |
| Surgical vs Surgical                   | 10  | 7                     | 17                      |       |       |
| Surgical vs Other                      | 1   | 8                     | 9                       |       |       |
| Other vs Other                         | 3   | 3                     |                         |       |       |
| Pharmacological vs Pharmacological     | 2 (all placebo) | 1 (placebo) | 3 (all placebo controlled) |       |       |
| Post-op Rehab vs Post-op Rehab         | 2   | 1                     | 3                       |       |       |
| Post-op Pharma vs Post-op Pharma       | 1   | 1                     | 1                       |       |       |
| Post-op Pharma vs No intervention      | 1   | 1                     | 1                       |       |       |

ACL – Anterior cruciate ligament.
RCT – Randomized controlled trial.
nRCT – non-randomized controlled trial.
fracture (n = 7; 19%), with the remaining two studies including any ‘acute knee injury’. Table I summarizes the basic study details grouped according to type of injury. All but two studies were RCTs (n = 35; 95%). Of 16 studies reporting power calculations, 15 met or exceeded the sample size required (Supplementary Table 2). Study duration varied widely, approximately equally distributed across 0–1 years, >1–5 years and >5 years. Most studies (70%) compared a surgical intervention against either another surgical or nonsurgical/non-pharmacological (henceforth referred to as ‘other’) intervention. Comparisons of post-operative rehabilitation interventions, pharmacological studies (the only studies to use a placebo arm) and all other interventions each accounted for ~8% of all studies (Table I).

An overview of inclusion and exclusion criteria for all available full-text papers (n = 32) is shown in Supplementary Table 3. Most studies (88%) had clearly defined eligibility criteria. Sixty percent provided a specific age range, spanning 13–50 years old. Sex was a specified criterion in only three studies, one of which excluded females. Elite professional sports activity and pregnancy were exclusions in 20% of studies.

Pre-existing conditions or other concomitant injuries excluded patients in 80% of studies. For example, previous index (and sometimes contralateral) knee injury and/or surgery were exclusions in >60% of studies and the presence of OA was an exclusion criterion in 25% of studies (Supplementary Tables 3 and 4).

One-hundred-and-forty-seven outcome measures were identified (Table II), including physical examination outcomes (n = 30), patient-reported outcomes (n = 26) of which the Knee Injury and Osteoarthritis Outcome Score (KOOS) was most frequently used; imaging outcomes (n = 43), biomarkers (n = 39) and other (n = 9) (Supplementary Tables 5–9 respectively). Primary outcome measures were identified by only 19 studies (Tables I and II). Ten different OA outcomes included nine imaging structural measures and one surrogate measure, KOOS (Table II, Supplementary Tables 6 and 7). Only five studies (of ACL rupture subjects) used molecular biomarkers as outcome measures (Supplementary Table 8).

Summary of key area discussions

Molecular pathogenesis and biomarkers of the injury response

Recently there has been an increase in our understanding of the molecular pathogenesis of PTOA. Observations from both humans and animal models reveal that diverse signalling pathways (involving inflammation, apoptosis and cell senescence) are activated by injury. This activation is associated with subsequent bone remodelling, cartilage matrix damage and synovial inflammation. Recently there has been an increase in our understanding of the molecular pathogenesis of PTOA. Observations from both humans and animal models reveal that diverse signalling pathways (involving inflammation, apoptosis and cell senescence) are activated by injury. This activation is associated with subsequent bone remodelling, cartilage matrix damage and synovial inflammation. Synovial fluid at the time of joint injury shows marked increases in pro-inflammatory cytokines (e.g., IL-6 is 1000-fold up-regulated) and within 2 weeks shows evidence of matrix catabolism of both aggrecan and type II collagen. The response appears to differ between individuals, and is represented by a tissue inflammatory response, primarily detectable in the synovial fluid . Following injury, a variety of factors may encourage joint homeostasis and resolution (including normal physiological loading), or progression to post-traumatic OA (including excessive loading or further injury). Further injury or surgery would appear to prolong the inflammatory response to trauma. There may be an ‘early therapeutic window’ following joint injury during which inflammatory response genes are up-regulated and matrix degradation is initiated which could be targeted by intervention . The optimal and/or latest times at which degradation could be halted or reversed are currently unknown.

Table II

<table>
<thead>
<tr>
<th>Outcome measure category (n*)</th>
<th>Primary outcomes</th>
<th>Osteoarthritis and surrogate OA outcomes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physical examination n = 30</td>
<td>1. Laxity (n = 4)</td>
<td>1. Knee injury and Osteoarthritis Outcome Score (KOOS) (n = 1)</td>
</tr>
<tr>
<td></td>
<td>2. Patellofemoral stability (n = 2)</td>
<td></td>
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<tr>
<td></td>
<td>3. Limb symmetry indices (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4. Torque (n = 1)</td>
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<td></td>
<td>5. Muscle electrical activity (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6. Functional hop test (n = 1)</td>
<td></td>
</tr>
<tr>
<td>Patient reported n = 26</td>
<td>1. Knee injury and Osteoarthritis Outcome Score (KOOS) (n = 5)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2. Hospital for Special Surgery (HSS) knee score (n = 2)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3. International Knee Documentation Committee (IKDC) Subjective Knee Form (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4. Kujala score (n = 2)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5. The Knee Self-Efficacy Scale (K-SES) (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6. The Physical Activity Scale (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>7. Tegner activity score (n = 1)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>8. Multidimensional Health Locus of Control (n = 1)</td>
<td></td>
</tr>
<tr>
<td>Imaging n = 43</td>
<td>1. Radiographic: Kellgren–Lawrence classification (n = 1)</td>
<td>1. Radiographic: Study specified criteria incl. joint space narrowing, osteophyte grade, subchondral sclerosis and sharpening of subchondral spines (n = 4)</td>
</tr>
<tr>
<td></td>
<td>2. CT: Quality of reduction (n = 1)</td>
<td>2. Radiographic: Kellgren–Lawrence classification (n = 4)</td>
</tr>
<tr>
<td></td>
<td>3. MRI: Morphologic measures of articulating bone curvature (femur, tibia &amp; trochlea) (n = 1)</td>
<td>3. Radiographic: Ahlback classification (n = 2)</td>
</tr>
<tr>
<td></td>
<td>4. MRI: Cartilage thickness of femorotibial medial compartment (n = 1)</td>
<td>4. Radiographic: modified OARSI grading scale for OA (n = 1)</td>
</tr>
<tr>
<td></td>
<td>5. MRI: Anterior Cruciate Ligament Osteoarthritis Score (ACLOS) (n = 1)</td>
<td>5. Radiographic: Medial joint space width (n = 1)</td>
</tr>
<tr>
<td>Biomarkers (n = 39)</td>
<td>1. GAG/proteoglycan marker: ARGS-aggrecan (n = 1)</td>
<td>6. Radiographic: Ahlback &amp; Fairbank composite scale (n = 1)</td>
</tr>
<tr>
<td>Other (n = 9)</td>
<td>2. Safety, tolerability &amp; adverse effects (n = 1)</td>
<td>7. MRI: Whole Organ Magnetic Resonance Imaging Score (WORMS) (n = 1)</td>
</tr>
<tr>
<td>TOTAL n = 147</td>
<td>Total – 21 primary outcomes</td>
<td>Total – 10 measures</td>
</tr>
</tbody>
</table>

*where n = total number of studies with such measures.
Animal models of knee injury

Much work on OA pathogenesis has been accomplished in animal models, which exploit the association between joint injury and OA, using trauma or surgically-induced injury to predictably induce disease: they are therefore particularly suited to testing early interventions in this setting. Findings from murine models such as those involving destabilization of the medial meniscus appear to translate to human studies of ACL rupture or meniscal tear.

The effects of suppressing certain key pathways in these models have been described in knockout mice. Despite this, very few interventions have been tested at the time of injury, in rodents or in man, as opposed to established OA, which could account for some of the failure of translation of OA therapeutics to date.

However, there may be some molecular differences as well as some practical challenges in the testing of intra-articular agents in small animals and in the extrapolation of optimal timing of an intervention from rodent to man.

Examples of potential pharmacological targets

Glutamate concentrations are increased in synovial fluid of arthritic joints in humans and animals, activating glutamate receptors on neurones and synoviocytes to induce pain and cause release of IL-6. Intra-articular inhibition of α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid (AMPA) and kainate glutamate receptors at the time of injury or induction of arthritis in rodent models alleviates pain, inflammation and joint degeneration.

IL-1 causes cartilage degradation in vitro and is upregulated in synovial fluid following joint injury. Blockade of this pathway (with IL1-receptor antagonist (IL1RA)) reduced inflammation and degeneration in a mouse model of arthritis. IL-1 or AMPA/kainate receptors represent potential therapeutic targets for preventing later disease, as their inhibition at the time of injury in models of post-traumatic OA reduced disease. IL1RA is the first therapeutic agent to be tested in human pilot studies at the time of knee injury for this indication. A further example is a small RCT testing steroid injection within 4 days of ACL tear, where the collagen degradation biomarker CTX-II was significantly reduced in synovial fluid in the steroid-treated arms. Since AMPA/kainate receptor antagonists, IL1RA and steroids are already used in man, re-purposing of existing agents is a real possibility.

Imaging in acute injuries

Imaging-based change following knee injury reflects the initial trauma but also the responses to subsequent changed dynamic knee loading after destabilizing injuries. The majority of studies include X-ray and MRI cartilage outcomes, both semi-quantitative and quantitative. Although there are a few high-quality longitudinal imaging studies after ACL rupture, more studies are needed. It is possible to define early OA on either X-ray or MRI, and evidence indicates that MRI changes alone can act as an endpoint. Depending on the target, non-cartilage MR outcomes, either bone-based, such as bone marrow lesions (BMLs) or synovitis-effusion, may be appropriate. Compositional measures using MRI, positron emission tomography (PET) or computed tomography (CT) remain investigational. Composite metric sequences including T1ρ and T2 have been associated with the PROM KOOS, pain after ACL reconstruction and with synovial fluid biomarkers at the time of surgery. Change in these compositional measures may reflect differences in surgical factors after ACL reconstruction and the pre-injury joint structure.

Consistent changes in cartilage thickness occur after ACL rupture: two cartilage regions quickly increase in thickness over time, whilst other areas decrease. Within 3 months of ACL injury, there are marked changes in knee bone curvature. Patellofemoral joint (PFJ) OA appears more prevalent in cohort studies, particularly relating to ACL rupture/reconstruction; however, the PFJ is not always examined by X-ray.

Structural changes generally develop slowly, and traumatic and degenerative changes must be clearly separated, although may appear similar (as in the case of BMLs). Common OA assessment methods are semi-quantitative instruments are only partially applicable in this setting: Whole Organ Magnetic Resonance Imaging Score (WORMS), BLOKS and MOAKS do not differentiate between traumatic and degenerative joint changes, and do not include assessment of post-surgical graft integrity. Anterior Cruciate Ligament Osteoarthritis Score (ACLOAS) is a new tool which addresses some of these issues including clear differentiation of traumatic from degenerative BMLs, extent of baseline traumatic osteochondral damage and assessment of the graft.

Imaging biomarkers predicting OA

A systematic review in this area reported that meniscal lesions, meniscectomy, BMLs, time from injury and altered biomechanics all are associated with cartilage loss over time after ACL rupture. Greater cartilage damage at baseline is associated with worse clinical outcome (although this could represent pre-existing OA). Presence of cortical depression fractures is associated with worse IKDC score at 1 year. MRI-detected inflammation markers (effusion synovitis/Hoffa-synovitis) at 2 years after ACL rupture were associated with OA development at 5 years. Effusion, or presence of BMLs at 1 year, or meniscal tears at any stage were found to be associated with radiological OA at 2 years. Early bone curvature change is predictive of cartilage loss at 5 years and accentuated by the presence of meniscal injury.

Points to consider

These are summarized under overarching considerations and three main areas: eligibility criteria, outcome measures and definition and timing of interventions and comparators in these studies.

Overarching considerations

Key overarching considerations are included in Table III. It was emphasized that a better understanding of disease pathogenesis was important. The appropriate time-window, role and effects of a proposed intervention on underlying processes such as inflammation, mechanical loading and subsequent bone or cartilage change need to be elucidated. Some findings may usefully be translated from animal models; however, it was also noted that there may be important differences between the response to acute knee trauma and a discrete surgically-induced isolated injury to ACL or meniscus. It was agreed that the considerations highlighted in this paper should be reviewed periodically as more data become available, with a maximum of 3 years before the next revision.

Eligibility criteria

Eligibility criteria should be clearly defined and should identify specific groups with a modifiable process following their injury in which to test the intervention (Table IV).

Definition of injury. Examples of well-defined groups based on MRI to be included would be ACL tear combined with other injuries such as traumatic meniscal tear (although different outcomes are probably associated with medial or lateral tears), or chondral damage/cortical depression fracture. Degenerative meniscal lesions should be considered part of early OA and not included in acute post-trauma studies. 55% of patients sustain simultaneous injuries to both ACL and meniscus, the ubiquitous biological
response to joint tissues injury supports broader inclusion of injury sub-types. Combined ligament injuries or fractures should not necessarily be excluded but considered as a separate ‘extreme’ phenotype, as they may be at substantially increased OA risk, which may or may not be reversible.

Time since injury. Some interventions may be most effective if exerting their effect as soon as possible after the early biological changes after injury. The appropriate time window for any intervention after injury needs to be carefully justified, according to its nature.

Age. Those less than 30 years are more likely to have purely traumatic meniscal lesions; those over age 35 could be at risk of pre-existing OA/degenerative meniscal lesions.

Demographics. Elite athletes are more likely to have past/repeated injuries but may have different responses to injury compared to non-elite individuals. As elite athletes are at high risk of OA, they still represent a relevant subgroup for investigation.

Exclusions. Previous substantial knee injury or surgical procedure to the index knee may confound results and should be considered as a possible exclusion. BMI should be documented: excessive obesity has independent effects on disease risk, joint loading and inflammation.

Outcome measures
Key considerations are shown in Table V.

PROMs. In addition to the collection of longer-term PROMs, repeated, multiple early measures will allow examination of potential earlier surrogate endpoints in the future.

Imaging. Baseline and longitudinal evaluation should differentiate pre-existing degenerative from acute traumatic structural joint damage. The contralateral knee may subsequently be affected,

Table III
Recommendations for points to consider: overarching considerations

<table>
<thead>
<tr>
<th>Consideration</th>
<th>Recommendation</th>
</tr>
</thead>
</table>
| 1. General    | - Considerations should be relevant to the design of all forms of interventional study following joint injury, unless otherwise stated  
                - CONSORT or STROBE criteria should be adopted in the design and reporting of any interventional or cohort study in this area  
                - Patients and the public should be involved throughout the process of study design and delivery |
| 2. Regulatory | - Current and future regulatory considerations and requirements in this area should be considered in design of future studies  
                - The community should work closely with regulatory bodies to establish evidence and precedent for outcomes and design of interventional trials  
                - Responder criteria, the number needed to treat for benefit (NNT), and cost-effectiveness should be measured |
| 3. Feasibility| - Feasibility, patient burden and cost considerations of, for example, type of imaging, or intervening near to the injury should be carefully weighed against the scientific/therapeutic benefits of the proposed approach  
                - For any given study, a balance should be found between scientific rigour in design and pragmatic considerations regarding recruitment and generalizability to clinical practice |
| 4. Specific targets | - Some of the considerations around study design (including eligibility criteria, outcomes and time-window of intervention) may be different, depending on the nature of the intervention  
                        - There may be particular biomarker(s) which are specific and sensitive for a particular intervention |
| 5. Stratification| - The assessment of personal or individualized risk was noted to be important  
                        - Novel molecular or imaging biomarkers might be used in the future as stratifiers at the point of entry to the study, or as intermediate (surrogate) outcomes, but none are validated for these purposes currently  
                        - Effective stratification of an individual's personal risk of post-traumatic OA is not yet possible based on current knowledge |

Table IV
Recommendations for points to consider: eligibility criteria

<table>
<thead>
<tr>
<th>Consideration</th>
<th>Recommendation</th>
</tr>
</thead>
</table>
| 1. Definition of acute knee injury | - The extent and characteristics of acute structural joint damage should be fully classified by magnetic resonance imaging  
                                        - Subgroups/types of injury for inclusion such as ACL and/or meniscal tear should be carefully defined  
                                        - Different types of injury may be associated with different biomechanical outcomes and responsiveness to any given intervention, so the target population needs to be carefully defined  
                                        - In the case of meniscal tears, the individual's age, history of a clear injurious episode, plus MR appearances are all important in identifying traumatic tears (and excluding degenerative lesions from these studies)  
                                        - Caution should be exercised in the inclusion of extreme phenotypes, for example those with isolated ACL tears or very extensive injuries |
| 2. Time since injury | - Establishing an appropriate therapeutic time-window will be relevant for each new target/intervention  
                        - Certain interventions targeting the early response to injury may benefit from being tested within days of injury, or up to a maximum of 4–6 weeks from injury |
| 3. Age | - Upper age limit should be carefully considered; an upper age limit of 35 was proposed  
        - Challenges were highlighted around intervening in paediatric populations who lack capacity to give informed consent or who have immature growth plates |
| 4. Demographics | - People of both sexes should be included  
                        - Studies may include, but should not be restricted, to professional athletes |
| 5. Proposed exclusions | - Other existing causes of joint pathology  
                                - inflammatory arthritis or pre-existing established osteoarthritis  
                                - other disorders of bone, current or past  
                                - previous substantial injury or surgery of index knee (particularly where there would be an associated markedly increased risk of PTOA)  
                                - other concomitant body injury or surgery (in some circumstances as may confound biomarkers)  
                                - Pregnancy or breast-feeding  
                                - Heavy use of alcohol, or recreational drug use  
                                - Morbid obesity |
therefore differentiating index from control knee is important. Considerations around type of imaging and its frequency include evidence of specific outcome performance metrics, feasibility and cost. Where trials are multi-center, MRI protocols need to be carefully designed (for example, compositional imaging may be challenging in a multi-center setting, and magnet strength should be considered in the context of ACL reconstruction and metal artefact).

Selection of imaging biomarker (semi-quantitative or quantitative) requires understanding of the validity, reliability and responsiveness of each measure. MRI techniques that assess early cartilage changes may be useful. Measures of synovial or fat pad inflammation may be important for anti-inflammatory therapeutics and MRI techniques that quantify synovitis may be considered. Early changes in 3D bone shape seen after injury which predict subsequent OA warrant further study as a potential surrogate endpoint.

Molecular biomarkers. These were noted to be under development as stratifiers and as outcome measures: none were yet sufficiently evidence-based to act as independent surrogate measures as either an early OA diagnostic, prognostic or patient selection aid for interventional studies. Irrespective of target, to accelerate therapeutic advances, it is important that bio-samples be collected in all cohorts and clinical trials where possible. DNA storage would allow the international community to work collaboratively to identify novel genetic predictors of outcome.

Synovial fluid was highlighted as a potentially important bio-sample, showing biologically important molecular changes after injury and after intervention; synovial fluid molecular changes are likely to have increased utility compared to serum13-15. Contra-lateral aspiration of synovial fluid was controversial, as the contralateral knee is not always a good control and it is difficult to aspirate normal joints. It is important that non-surgical studies access synovial fluid to avoid bias towards surgical intervention studies. In some cohorts, serum/plasma/urine may be available prior to the injury (e.g., participants in a biobank or military cohorts): measuring change within an individual was noted as analytically powerful. Regarding biomarker choice, the most qualified biomarkers to date, e.g., CTX-II, could be included if cartilage matrix catabolism is a target; synovial inflammation or bone biomarkers, or specific cytokine measurements may be relevant depending on target27.

Functional outcomes. Symptoms of instability could be more reliable than any examination-based measures. However, their sensitivity to change compared with existing measures such as pain should be evaluated further27.

**Definition and timing of intervention and comparator**

The choice of timing of the intervention will depend on the nature and mode of its action and intended effects, as well as the measured outcome. An optimal 'therapeutic window' should be carefully defined for any intervention (Table VI), see also Eligibility Criteria: 'Time Since Injury'. It may be that identification of high risk phenotypes is possible by imaging or molecular biomarkers at defined times after the injury.

Types of intervention are highly varied; where multi-modality interventions are used, these should be carefully defined, and controlled. Drugs could be given systemically or intra-articularly, as single or multiple doses, dependent on agent and duration of treatment, safety considerations and acceptability.

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**Table V**

Recommendations for points to consider: outcome measures

<table>
<thead>
<tr>
<th>Consideration</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. General</td>
<td></td>
</tr>
<tr>
<td>2. Patient reported outcome measures (PROMs)</td>
<td></td>
</tr>
<tr>
<td>3. Imaging</td>
<td></td>
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<tr>
<td>4. Molecular biomarkers</td>
<td></td>
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<tr>
<td>5. Functional outcomes</td>
<td></td>
</tr>
</tbody>
</table>

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Research recommendations

The particular challenges and questions highlighted as needing further research are included in Table VII.

Table VII
Recommendations for points to consider: definition and timing of intervention and comparators

<table>
<thead>
<tr>
<th>Consideration</th>
<th>Recommendation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. General</td>
<td>□ Optimal time-window for administration of any given intervention should be validated and clearly defined</td>
</tr>
<tr>
<td></td>
<td>□ Assumptions should be avoided; different proposed time-windows for intervention should be tested head to head in feasibility studies if necessary, to ensure patient acceptability, recruitment and likely translation into clinical care</td>
</tr>
<tr>
<td>2. Comparators</td>
<td>□ A comparator and/or placebo or sham arm should always be used where possible</td>
</tr>
<tr>
<td></td>
<td>○ Choice will depend on whether study is efficacy or pragmatic</td>
</tr>
<tr>
<td></td>
<td>○ Patients should be randomized to intervention or comparator arms</td>
</tr>
<tr>
<td></td>
<td>○ Assessment of acceptability of sham treatments, particularly when invasive, is paramount when considering design and feasibility</td>
</tr>
<tr>
<td></td>
<td>□ Double blind protocols should be used where possible</td>
</tr>
<tr>
<td></td>
<td>□ While double-blinding is not always possible, blinded observer/assessor almost always is</td>
</tr>
<tr>
<td>3. Multimodality intervention</td>
<td>□ Multimodality interventions may be particularly suited to this area</td>
</tr>
<tr>
<td></td>
<td>○ Such studies are very challenging to design and deliver and require expert input</td>
</tr>
<tr>
<td></td>
<td>○ Choice of each component ideally requires a priori evidence of effect</td>
</tr>
<tr>
<td></td>
<td>□ The interaction of different interventions is an important consideration in this area, given that multi-modal intervention is common in clinical practice.</td>
</tr>
</tbody>
</table>

A comparator and/or placebo or sham arm should be used, because of the known substantial placebo effect in OA studies48. The comparator will often be standard or usual care, rather than no treatment and requires careful definition. Randomization and placebo control are important principles not only for pharmacological interventions, but also for device and surgical studies, where a large placebo effect would be anticipated and which is not otherwise controlled49.

There are a number of practical considerations for successful recruitment, randomisation strategies, the standardisation of the intervention (particularly if surgical) and allocation concealment in these types of studies, particularly when they are multi-site18. This should be carefully considered during study design and a number of existing OARSI recommendations in trials of prevention of joint injury and of established OA are highly relevant here7,8,50,51.

Patient representatives highlighted concerns for the potential for over-diagnosis or overtreatment in the absence of risk stratification, and further Patient and Public Involvement is encouraged in this area now, and as the field develops.

Further evidence is needed for which outcomes should be used in this setting, and what measurement(s) (whether a molecular or imaging biomarker or PROM) might act as an acceptable surrogate short term outcome for future OA (given that 5–10 year interventional trials are not feasible). Although these current considerations address interventional studies, the consensus group acknowledged that ancillary/cohort studies which establish associations between PROMs, biomarkers and imaging outcomes could address key knowledge gaps to provide evidence for future trials. The design of these studies should be carefully considered and outcomes appropriately powered, but they may include more exploratory outcomes. Sensitive, specific early measures which might shorten studies should be sought.

The Consensus group noted that animal studies can inform human studies, and such programs were justifiable to facilitate early translation of targets to humans.
Discussion

Our review of the literature has highlighted a lack of conformity in design of interventional studies in this area. Evidence from the review and expert consensus has been synthesised in producing these first international considerations on the design and conduct of interventional studies aiming at prevention of OA following acute knee injury. Critical knowledge gaps limiting such trials have been highlighted, and summarised as research recommendations. These considerations are intended to underpin future guidelines as this field evolves. Collaborative working on cohort and feasibility studies is needed to provide better evidence for interventional study design.

Studies need to include those patients who are at the highest risk, but whose risk is modifiable by the proposed intervention. There was an awareness of the identity of extreme phenotypes, such as combined ligament injuries, which may fall outside these criteria. As in OA, predictive risk modelling is needed for knee trauma. A better understanding of underlying disease mechanisms from both animal and human studies is needed. Understanding how related mechanisms such as inflammation and mechanical loading of the joint after trauma contribute to either resolution or progression to OA was deemed essential for the development of new interventions.

The feasibility and acceptability of testing interventions in an acute setting can be challenging. Informed consent for sham or placebo treatments at the time of knee injury needs careful review by patients, healthcare providers and trialists. Sham-controlled trials including surgical trials are often needed to provide the best possible level of evidence. Recent consensus in classification of early knee OA will facilitate such trials. Alternative surrogate outcome measures need to be developed to shorten trial duration and improve the likelihood of drugs being developed by industry. MRI costs are relatively high, but may be justified by allowing researchers to examine earlier outcomes. Whilst X-ray follow-up may appear more feasible, it’s use as a lone imaging modality must be adequately powered.

There are some limitations to the approach used. The literature review was performed to provide evidence for discussions, rather than as a stand-alone piece of work; it was clear after the initial search that areas of interest, such as pharmacological interventions, were not well represented in the current literature, and limitations of generalizability to all types of interventions should therefore be borne in mind. A critical appraisal of the studies was not performed as it was not felt necessary for the requirements of this review, which was pragmatic in nature. Given the relatively low number of RCTs identified in this area, non-randomized controlled trials as well as RCTs were included where identified. Not all opinions might be equally represented from this type of approach. However, a wide range of stakeholders and groups were involved, including patients. Effort was made to ensure diversity; pre-appointed facilitators and reporters with note-keeping and voice recording of sessions ensured a transparent and consistent process. More detailed discussions on considerations of recruitment/randomization/allocation concealment strategies were beyond our scope.

In summary, these initial considerations provide a starting point for further work in this area. These points are intended to be complimentary to, and should be considered alongside, OARSI Clinical Trials Recommendations on prevention of joint injury, the design, analysis and reporting of OA RCTs and clinical requirements for development of therapeutics in OA. The regulatory considerations for a new indication of preventing symptoms or OA structural change following joint injury are unique. Engagement with both regulators and the pharmaceutical industry is essential if the area is to progress and overcome current hurdles. Although such trial designs may be challenging, in order to develop new therapeutics with the aim of patient benefit, the consensus was that progress in this area is both possible and urgently required.

Author contributions
All authors made substantial contributions to all three of sections (1), (2) and (3) below:

(1) acquisition of data, or analysis and interpretation of data.
(2) drafting the article or revising it critically for important intellectual content.
(3) final approval of the version to be submitted.

DJM, FEW and PGC in addition conceived and designed the work, and take collective responsibility for data integrity as a whole.

Competing interest statement
FEW – none.
NC – none.
SK – none.
RF – none.
ME – none.
DF – none.
ML – employee of Abbvie pharmaceutical company.
SM – none.
CW – none.
DB – none.
SL – none.
VBK – none.
FWR – Shareholder Boston Imaging Core Lab. (BICL), LLC., a company providing radiologic image assessment services to academia and the pharmaceutical industry.

DJM – co-inventor on a patent related to the use of glutamate receptor antagonists to prevent osteoarthritis (WO2015001349).

PGC – none.

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Supplementary data

Supplementary data related to this article can be found at https://doi.org/10.1016/j.joca.2018.08.001.

References


