Adipose-derived mesenchymal stem cell therapy in the treatment of knee osteoarthritis: a randomized controlled trial

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Aim: To evaluate the efficacy of autologous adipose-derived mesenchymal stem cell (ADMSC) therapy on pain, function and disease modification in knee osteoarthritis. Methods: 30 participants with symptomatic knee osteoarthritis were randomized into three groups. Two treatment groups received intra-articular ADMSC therapy consisting of either a single injection (100 × 10^6 ADMSCs) or two injections (100 × 10^6 ADMSCs at baseline and 6 months). The third group served as control and continued conservative management. Results: No serious adverse events were observed. Both treatment groups receiving ADMSCs showed clinically significant pain and functional improvement at completion of follow-up at 12 months. Radiological analysis using the Magnetic Resonance Imaging Osteoarthritis Knee Score indicated modification of disease progression. Conclusion: Autologous ADMSC therapy appears to be a safe and effective therapy for knee osteoarthritis and may have the potential to prevent disease progression.

Trial registration number: ACTRN12614000814673

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Osteoarthritis (OA) is an increasingly prevalent and progressive condition causing chronic pain and loss of function. Worldwide it is acknowledged as the fourth leading cause of disability and an important factor affecting disability-adjusted life years [1,2]. Symptomatic arthritis has a prevalence of 10% of males and 18% of females over the age of 45 years with radiological prevalence in up to 80% of the population at 65 years of age [3–6]. With an aging population this prevalence is expected to increase [3,4].

Conventional conservative medical treatments for knee OA are aimed at symptomatic control rather than disease modification. These pharmaceutical treatments demonstrate only modest and limited benefits and may have unwanted side effects [7–10]. Surgical total knee replacement, which remains the chosen treatment for symptomatic knee OA that is uncontrolled by conservative therapies, is not without significant complication with as many as 20% of patients having persistent pain or loss of function at 12 months [11–13].

Whilst historically primary OA has been considered an age-related degenerative condition, we now have a much greater understanding of the pathophysiology of this condition, with an acceptance that OA results due to an imbalance between catabolic/pro-inflammatory and anabolic pathways [14]. Release of reactive oxygen species (free radicals) as a result of mechanical or biological stressors has been shown to lead to chondrocyte cell senescence [15]. This is accompanied by an observed increase in inflammatory cytokine expression including IL-1, TNF-α and matrix metallopeptidase-13 [14–16]. These cytokines further lead to production of other pro-inflammatory factors, including IL-8, IL-6, leukotriene inhibiting factor, proteases and prostaglandin E2 (PGE2). This host of inflammatory compounds leads to a pro-catabolic state favoring matrix degradation and development of OA.
The ability of mesenchymal stem cells (MSCs) to differentiate along a mesodermal lineage, including osteoblasts and chondrocytes, has suggested their intrinsic potential in tissue repair and regeneration [17–19]. It is, however, likely that it is via paracrine signaling, rather than direct differentiation that MSCs manipulate the local environment seen in OA, which leads to the disease modification [20].

MSCs are known to have immunomodulatory and anti-inflammatory benefits through suppression of inflammatory T-cell proliferation, inhibition of monocyte maturation and expression of anti-inflammatory cytokines [21–24]. Expression of anabolic cytokines may lead to upregulation of previously senescent but metabolically active and responsive chondrocytes [25].

Preclinical research has successfully shown benefit of intra-articular MSC therapy in observed pain and functional improvement [26–29]. While disease modification has been observed in some trials, this has not been consistent. Initial clinical trials have encouragingly observed similar pain and functional improvement in symptomatic knee OA following MSC therapy [30–33]. The use of expanded autologous adipose derived MSCs (ADMSCs) in an early Phase I/II trial resulted in radiological evidence of improved cartilage volume and confirmed hyaline-like cartilage on histology [32]. Similarly, the use of allogeneic isolated and expanded bone marrow-derived MSCs had observed improvements in pain and function with improvement in cartilage quality and disease stabilization in comparison to the control group at 12 months [33].

Based upon published clinical trial outcomes, MSC therapy appears safe. In a formal systematic review and meta-analysis of trials involving intravascular administration of either autologous or allogeneic expanded MSC therapies (totaling over 1000 participants) no significant adverse events except transient fever were identified [34]. A systematic review of clinical trials using intra-articular autologous expanded MSC therapies with a mean follow-up of 21 months and assessing 844 procedures similarly showed no association with adverse events including infection, malignancy or death [35].

We present a randomized controlled trial (RCT) to assess the efficacy of autologous ADMSCs in the treatment of symptomatic knee OA. As past studies involving only a single intra-articular MSC injection have postulated potential benefits of multiple injections we have chosen to assess the efficacy of different dosing regimes. While the primary objective is to assess safety, pain and functional changes, a secondary objective is to determine the potential of MSC therapy to achieve disease modification, as detected through radiological examination using magnetic resonance imaging (MRI) techniques.

Materials & methods

Trial design

This RCT was approved by the Human Research Ethics Committee (HREC) of Monash University and registered on the Australian and New Zealand Clinic Trial Registry (Trial ID: ACTRN12614000814673). All participants completed written informed consent. The study was a single centred trial.

The trial design consisted of a total of 30 participants randomly and equally allocated to a control group and two treatment groups. Participants were not blinded to their treatment allocation.

Treatment groups were to receive different treatment dosing regimes of autologous ADMSCs:

- Control: Ongoing conventional conservative management only
- one-injection group: a single intra-articular injection of 100 × 10^6 ADMSCs
- two-injection group: two intra-articular injections of 100 × 10^6 ADMSCs (baseline and 6 months).

A third treatment group receiving five injections of 40 × 10^6 ADMSCs at baseline, 1, 2, 3 and 6 months, respectively, was intended but ceased due to observed and reproducible moderate adverse events in a concurrently run study with the same treatment protocol (documented as increasing self-limiting pain with sequential injections at monthly intervals) [36]. Participants that were already randomly allocated to this treatment group and yet to commence the trial were re-randomized to another trial group. The HREC of Monash University were informed of this trial protocol variation.

Those participants allocated to the control group maintained their pre-existing conventional conservative management program consisting of simple analgesia, weight management and exercise. Participants were not prescribed a trial-specific conservative program. Participants recorded any changes in their conservative management program.
Table 1. Eligibility criteria.

<table>
<thead>
<tr>
<th>Inclusion criteria</th>
<th>Exclusion criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Radiological diagnosis of osteoarthritis using the American College of Rheumatology criteria</td>
<td>1. Pregnancy (accepted contra-indication as no safety data on this population)</td>
</tr>
<tr>
<td>2. Radiological grading of Grade II–III osteoarthritis (OA) of the knee as determined by a qualified radiologist using the Kellgren and Lawrence system</td>
<td>2. Breast feeding (accepted contra-indication as no safety data on this population)</td>
</tr>
<tr>
<td>3. Medial or lateral compartment OA as determined above</td>
<td>3. Have other causes of their knee symptoms suspected to be due to serious pathology such as tumors or referral from the hip or lumbar spine. These conditions are not under investigation within the current project</td>
</tr>
<tr>
<td>4. Conservative OA treatment already undertaken defined as: analgesia/anti-inflammatory medication, supplements approved by the treating clinician (e.g., glucosamine sulphate), an attempted exercise program prescribed by a physiotherapist or medical practitioner for at least 8 weeks, weight loss and nutritional management as prescribed by a dietitian or medical practitioner for at least 8 weeks, and biomechanical management including bracing if appropriate as prescribed by a physiotherapist, podiatrist or medical practitioner. Autologous MSC is an invasive treatment and guidelines recommend trialing conservative measures as the first line of treatment in knee OA</td>
<td>4. Bleeding disorder – i.e., hemophilia (accepted contra-indication as no safety data on this population)</td>
</tr>
<tr>
<td>5. A minimum pain score of 5 on an 11-point numerical rating scale</td>
<td>5. MRI confirmed displaced meniscal tear</td>
</tr>
<tr>
<td>6. Single knee osteoarthritis</td>
<td>6. MRI confirmed Grade IV chondral loss</td>
</tr>
<tr>
<td>7. Less than 5 degrees varus or valgus knee deformity as measured by the long mechanical axis of the knee on x-ray</td>
<td>7. Previous meniscectomy/significant partial meniscectomy or other knee related surgery within the last 12 months</td>
</tr>
<tr>
<td>8. Sufficient English skills to complete the questionnaires required for the study, as well as to understand the instructions given by the study doctors. This is required as no funding is available for translation or interpreters, and the outcome questionnaires to be used have only been validated in English language</td>
<td>8. Previous intra-articular injectable therapies within the last 6 months</td>
</tr>
<tr>
<td></td>
<td>9. History of cancer</td>
</tr>
<tr>
<td></td>
<td>10. History of atypical chronic pain syndrome – i.e., chronic regional pain</td>
</tr>
<tr>
<td></td>
<td>11. History of systemic illness or significant organ impairment/failure (i.e., renal failure)</td>
</tr>
<tr>
<td></td>
<td>12. History of allergy to any substances used within the treatments</td>
</tr>
<tr>
<td></td>
<td>13. Plans at the time of enrollment to undergo surgery in the following 12 months. This criterion is aimed at avoiding co-interventions that may confound the results of the study. While involvement in the project will not strictly prevent participants from undertaking such interventions if required, we will exclude volunteers who already have such procedures scheduled</td>
</tr>
</tbody>
</table>

Participants

Eligible participants 18 years or older with unilateral moderate symptomatic knee pain and Grade II–III medial or lateral compartment knee OA according to Kellgren Lawrence criteria were recruited. All participants had previously attempted primary conservative management of OA including simple analgesia, an exercise and weight management program and biomechanical adjustment with bracing or orthotics if relevant. Formal eligibility criteria is outlined in Table 1.

Baseline assessment involving a subjective and abbreviated musculoskeletal physical examination was performed with diagnostic x-ray and MRI assessed to determine eligibility criteria as indicated in Table 1. Participants received formal written information about the requirement of involvement in the trial and were invited to provide written informed consent.

Intervention

*Autologous ADMSC preparation*

Adipose tissue was chosen as the source of autologous MSCs due to the relative ease of harvest, abundance of MSCs and shared chondrogenic potential in comparison to other source sites such as bone marrow [37–39].

The harvest, isolation and expansion of ADMSCs has been previously published [36,40,41]. Participants randomly allocated to a treatment group underwent an abdominal lipoharvest procedure. Using a lateral abdominal approach, the subcutaneous fat was infiltrated with up to 300 ml of tumescent fluid (comprising of 30 ml of 2% lidocaine, 1 ml of 1:1000 adrenaline and 1 ml of 8.4% bicarbonate suspended in a normal saline solution to a total 1000 ml). Following this, up to 60 ml of adipose tissue and tumescent fluid was aspirated through a 4 mm lipoaspirate cannula.
Table 2. Fluorescence-activated cell sorting surface marker analysis. Mean data (n = 20) is presented with standard deviation in brackets.

<table>
<thead>
<tr>
<th>Histograms - positive markers</th>
<th>Histograms - negative markers</th>
</tr>
</thead>
<tbody>
<tr>
<td>CD90+ve</td>
<td>CD14+ve</td>
</tr>
<tr>
<td>CD73+ve</td>
<td>CD19+ve</td>
</tr>
<tr>
<td>CD105+ve</td>
<td>CD34+ve</td>
</tr>
<tr>
<td>CD14+ve</td>
<td>CD45+ve</td>
</tr>
<tr>
<td>Percentage</td>
<td>Percentage</td>
</tr>
<tr>
<td>98.17 (2.4)</td>
<td>0.89 (1.0)</td>
</tr>
<tr>
<td>97.75 (3.1)</td>
<td>0.31 (0.4)</td>
</tr>
<tr>
<td>93.78 (6.6)</td>
<td>0.95 (1.1)</td>
</tr>
<tr>
<td>0.95 (1.1)</td>
<td>0.63 (0.7)</td>
</tr>
</tbody>
</table>

Table 3. Average cell count and viability confirmed using a muse cell analyzer.

<table>
<thead>
<tr>
<th>Cell count (mean ± standard deviation)</th>
<th>Viability (mean ± standard deviation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>One-injection group</td>
<td></td>
</tr>
<tr>
<td>Baseline injection</td>
<td>103.9 million (7.7)</td>
</tr>
<tr>
<td></td>
<td>95.4% (2.3)</td>
</tr>
<tr>
<td>Two-injection group</td>
<td></td>
</tr>
<tr>
<td>Baseline injection</td>
<td>95.1 million (11.1)</td>
</tr>
<tr>
<td></td>
<td>93% (4.2)</td>
</tr>
<tr>
<td>6 month injection</td>
<td>102.6 million (8.3)</td>
</tr>
<tr>
<td></td>
<td>92.9% (3.6)</td>
</tr>
</tbody>
</table>

Characterization of isolated ADMSCs was confirmed using international established standards as published by the International Society of Cellular Therapy [44]. Flow Cytometry Fluorescent Activated Cell Sorting (FACS) analysis was completed independently on all isolated ADMSC samples at Monash University. Samples were tested for the presence of MSC surface markers CD90, CD73 and CD105 and absence of hematopoietic surface markers CD14, CD19, CD34 and CD45 (Table 2 and Supplementary Material). All isolated ADMSC samples underwent additional independent sterility testing for microbial growth/contamination.

**Injection method**

On the day of injection stored cells were thawed using a sterile water bath with cryoprotectant removed under centrifugation and washed in chilled phosphate buffered saline (PBS). The resultant cell pellet was resuspended in injectable sterile isotonic (0.9%) normal saline to a total of 3 ml. Cell count and viability was confirmed using a Muse Cell Analyzer (Merck Millipore, USA; Table 3 and Supplementary Material). For a single participant in the two-injection group the muse cell analyzer was not available for the baseline injection and a manual hemocytometer count was performed.

At the time of injection, the knee of the participants allocated to an active treatment group was prepped using standard sterile protocols. All injections were performed under direct visualization using ultrasound guidance to ensure accuracy. Local anesthetic (2 ml of 1% lidocaine) was infiltrated superficial to the knee joint capsule. Under aseptic conditions, and using a superolateral patella approach, the re-suspended autologous ADMSCs were injected into the intra-articular knee space.

**Post-operative rehabilitation**

Participants in the treatment groups were provided with post injection analgesia as required. They were advised to remain nonweight bearing with the use of crutches for 4 weeks. Education regarding range of motion and quadriceps activation exercises was provided. Participants in the two injection group were not required to be non-weight bearing after the second injection at 6 months.

**Outcome measures**

Primary outcome aims of this trial were pain and functional changes as measured by the Numeric Pain Rating Scale (NPRS), the Knee Injury and Osteoarthritis Outcome Score and Western Ontario and McMaster Universities...
Osteoarthritis Index (WOMAC). Secondary outcomes included structural assessment through use of MRI and additional outcome questionnaires (see Table 4).

**Questionnaires**

Outcome questionnaires were recorded using the software program Clinical Intelligence (Clinical Intelligence, Melbourne, Australia) and were recorded at baseline, 1, 3, 6 and 12 months.

- **Numeric Pain Rating Scale (NPRS).** Participants were asked to rate their average pain intensity over the past week on a scale of 0–10. The NPRS has been validated for use in people with knee OA [45].

- **The Knee Injury and Osteoarthritis Outcome Score (KOOS).** The KOOS is a validated score that consists of 5 subscales pain improvement, symptom improvement, activities of daily living, function in sport and recreation and knee-related quality of life. A normalized score is calculated for each subscale with 100 indicating no symptoms and 0 indicating maximal symptoms [46].

- **The Western Ontario and McMaster Universities Arthritis Index (WOMAC Index 3.0).** A validated quality of life score, the WOMAC score quantitatively assesses pain, stiffness and physical function in patients with symptomatic OA [47]. In this trial the WOMAC score is presented as an inverse percentage to be more easily compared with the KOOS subscales where 100 indicates no symptoms.

**Magnetic resonance imaging**

Magnetic resonance imaging (MRI) offers the ability to perform a non-invasive evaluation of structural changes associated with OA. The MRI Osteoarthritis Knee Score (MOAKS) is a validated semi-quantitative tool for the evaluation of knee OA [48]. The knee is divided into 14 sub-regional divisions (patella – medial and lateral; femoral – medial and lateral trochlea/central/posterior; tibia – medial and lateral, anterior/central/posterior) with each region achieving a score for features of OA. Features assessed for the purpose of this study were:

- **Bone marrow lesions (BMLs) and cysts.** Each region is graded for the percentage of volume of each BML and whether there is an associated cyst.

- **Articular cartilage.** Each region is graded on the percentage area of cartilage with partial to full thickness loss and also separately on area with full thickness loss.

- **Osteophytes.** Size of osteophytes in 14 sub-regions are graded as Grade 0 (none), Grade 1 (small), Grade 2 (medium) and Grade 3 (large).

- **Synovitis.** Volume of effusion-synovitis is graded as Grade 0 (none), Grade 1 (small), Grade 2 (medium) and Grade 3 (large).

- **Meniscus.** Medial and lateral meniscus are assessed for signal and morphology of the tear including vertical, radial, horizontal, complex, root tear, partial or complete maceration.

- **Peri-articular features.** Popliteal cyst – scored as present or not present.

MOAKs scores were used to describe OA progression over time as previously published [49]. Subregional scores were collated into the broader regions of medial, lateral and patella-femoral compartments.

MRI analysis was performed on a 1.5 T or greater MRI system with a dedicated peripheral knee coil. Proton density (PD) and proton density fat saturated (PDFS) images were completed in coronal, sagittal and axial planes.

MRI assessment was conducted by a musculoskeletal physician trained in MRI analysis using a single blinded approach with no knowledge of the treatment group.

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Table 4. Outcome measures.

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Measurement point (months)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Primary Outcome Measure</strong></td>
<td></td>
</tr>
<tr>
<td>1. Numeric Pain Rating Scale (NPRS)</td>
<td>0, 1, 3, 6, 12</td>
</tr>
<tr>
<td>2. Knee Injury and Osteoarthritis Outcome Score (KOOS)</td>
<td>0, 1, 3, 6, 12</td>
</tr>
<tr>
<td>3. Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC)</td>
<td>0, 1, 3, 6, 12</td>
</tr>
<tr>
<td><strong>Secondary Outcome Measure</strong></td>
<td></td>
</tr>
<tr>
<td>1. MRI analysis</td>
<td>0, 12</td>
</tr>
<tr>
<td>– MRI Osteoarthritis Knee Score (MOAKS)</td>
<td></td>
</tr>
</tbody>
</table>
Sample size
To date there is insufficient published data on the effects of autologous MSC therapy in OA. As such, sample size calculations are not feasible. This study represents a pilot RCT to assess treatment protocols for safety and efficacy prior to development of a larger RCT and is consistent with previously published study guidelines [50].

Randomization & allocation
Participant group allocation was prepared in advance using a web-based automated random number generator (www.randomizer.org). This was prepared by a designated researcher not involved in the recruitment, screening, assessment, enrolment or treatment process. All researchers with direct involvement in participant recruitment and treatment were blinded to this process.

Adverse events
Adverse events (AEs) were defined as an undesirable clinical development in a participant which was not present at baseline or which increased in severity after treatment. AEs were assessed and graded in regards to severity:

- Mild: awareness of a sign or symptom which does not interfere with the participants’ usual daily activity and/or is transient, resolving with use of simple interventions including simple analgesia.
- Moderate: interferes with the participants’ usual daily activity and/or requires symptomatic treatment including regular analgesia (i.e., opiate analgesia).
- Severe: symptom(s) causing severe discomfort with significant impact of the participants’ usual daily activity.
- Serious: any unexpected medical incident which requires hospitalization, results in long term disability, is life-threatening or results in death.

Statistical analysis
SigmaPlot 10 graphical and statistical software (Systat Software, Inc., CA, USA) was used to conduct the analysis. Two Way Repeated Measures Anova (One Factor Repetition, General Linear Model) was utilized to detect mean differences in the independent variables of Treatment (two injections, one injection or control) and Time (baseline, 1, 3, 6 and 12 months) for the dependent variables of NPRS, WOMAC and KOOS scores.

The objectives of the data analysis were first to determine if differences in means occurred between treatments at each of the 5 time points, and second, to determine if mean differences occurred within each of the three groups when comparing 1, 3, 6 and 12 months data to baseline measurements.

When statistically significant (p < 0.05) interactions between the factors of Treatment and Time were apparent an All Pairwise Multiple Comparison Procedure (Holm-Sidak method) was employed to determine the specific mean differences for both Time and Treatment factors with alpha again set at 0.05.

For the AEs and MOAKs categorical data analysis, Chi-square tests were performed. For the MOAKs data analysis the percentage data was first converted to categories (yes or no) so as to perform the Chi-square tests. Note that due to low sample sizes and low numbers in each category these tests were typically below a power level of 5.

Patient & public involvement
Patients and/or the public were not involved in the development of the research question, choice of outcome measures, design of the trial, recruitment of participants and conduct of the trial. Results of the trial will be disseminated to study participants through direct consultation with a trial clinician at completion of the trial and on publication of results.

Results
Intergroup comparison & demographics
As indicated in the Figure 1 flow chart, two participants (one in the Control Group and also another in the two-injection group) were excluded from radiological follow-up analysis as there was significant delay in follow-up MRI (>3 months). One participant was excluded from questionnaire analysis due to incomplete questionnaire data collection (though completed radiological follow-up).

Demographics of the study participants are presented in Table 5. Despite randomization there was a difference in body mass index (BMI) between groups with the control group being, on average, overweight, while both treatment groups were within the obese range.
Adipose-derived mesenchymal stem cell therapy in the treatment of knee osteoarthritis

**Figure 1. Flow chart.**
ADMSC: Adipose-derived mesenchymal stem cells.

**Table 5. Demographic characteristics for the Control and Intervention Groups.** Mean data are presented with standard deviations in brackets. Note the significant difference ($p < 0.05$) for body mass index with the one-injection group having a significantly higher body mass index than the Control group.

<table>
<thead>
<tr>
<th>Demographic Characteristics</th>
<th>Control</th>
<th>One injection</th>
<th>Two injections</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total number (n)</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Gender (% male)</td>
<td>50</td>
<td>70</td>
<td>40</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>51.5 (6.1)</td>
<td>54.6 (6.3)</td>
<td>54.7 (10.2)</td>
<td>0.58</td>
</tr>
<tr>
<td>Height (m)</td>
<td>1.73 (0.1)</td>
<td>1.75 (0.1)</td>
<td>1.7 (0.1)</td>
<td>0.59</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>76 (14.5)</td>
<td>97.1 (20.2)</td>
<td>88.7 (23.5)</td>
<td>0.07</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>25.2 (3.4)</td>
<td>31.6 (5.9)</td>
<td>30.4 (5.6)</td>
<td>0.023</td>
</tr>
</tbody>
</table>

**Pain & functional outcome measures**

Pain, as measured by the NPRS, in the one-injection group and two-injection group improved from a mean (standard deviation) of 6.7 (1.7) and 6.5 (1.4) to 2.6 (1.8) and 2.3 (2) respectively at completion of the study. Within group improvement when compared with baseline was statistically significant ($< 0.05$) throughout all time points in both the single and two-injection protocol treatment groups (Figure 2). Similarly, between group differences revealed that both the one-injection and two-injection groups had significantly less pain at 12 months compared with the Control group (Figure 2). No significant differences were found between the two treatment groups.

KOOS subscale analysis showed consistent improvement in all subscales during follow-up and this was maintained until completion of follow-up at 12 months. There was no significant difference at 12 months in all subscales.
Figure 2. Numeric pain rating scale.
* indicates a significant difference between the one-injection group and the Control group at that time point.
# represents a significant difference between the two-injection group and the Control group at that time point. Error bars represent 95% confidence intervals. Within-group differences (not displayed) are detailed in preceding text and are presented in Supplementary Table data.

Figure 3. Knee injury and osteoarthritis outcome score pain improvement subscale.
* indicates a significant difference between the one-injection group and the Control group at that time point.
# represents a significant difference between the two-injection group and the Control group at that time point. Error bars represent 95% confidence intervals. Within group differences presented in Supplementary Table data.
Figure 4. Knee injury and osteoarthritis outcome score pain improvement subscale.
* indicates a significant difference between the one injection group and the Control group at that time point.
# represents a significant difference between the two-injection group and the Control group at that time point. ∧ indicates a significant difference between treatment groups. Error bars represent 95% confidence intervals. Within group differences presented in Supplementary Table data.

Figure 5. Knee injury and osteoarthritis outcome score Activities of Daily Living subscale.
* indicates a significant difference between the one-injection group and the Control group at that time point.
# represents a significant difference between the two-injection group and the Control group at that time point. Error bars represent 95% confidence intervals. Within-group differences presented in Supplementary Table data.

ADL: Activities of daily living; KOOS: Knee injury and osteoarthritis outcome score.
Figure 6. Knee injury and osteoarthritis outcome score sport and recreation subscale.
*indicates a significant difference between the one-injection group and the Control group at that time point.
* indicates a significant difference between treatment groups at that time point. Error bars represent 95% confidence intervals. Within-group differences presented in Supplementary Table data.

Figure 7. Knee injury and osteoarthritis outcome score Quality of Life subscale.
*indicates a significant difference between the one-injection group and the Control group at that time point.
* indicates a significant difference between the two-injection group and the Control group at that time point. Error bars represent 95% confidence intervals. Within-group differences presented in supplementary table data.
KOOS: Knee Injury and Osteoarthritis Outcome Score.
between the two treatment groups. All subscales displayed statistically significant improvement against control at 12 months of follow-up. The one-injection group had more consistent improvement against Control throughout the follow-up period. See Figures 3–7.

Global WOMAC score expressed as an inverse percentage improved throughout follow-up in both the one-injection group and two-injection group with a mean (standard deviation) of 59.6 (17.9) and 54.4 (18.2) at baseline improving to 84 (9.4) and 87.3 (8) at 12 months. Statistically significant improvement versus baseline measures was observed as early as 1 month and until final follow-up at 12 months. Both treatment groups had statistically significant improvement at 12 months in comparison to control (Figure 8).

The mean power for all statistical tests performed for pain and functional outcome measures (n = 21) was 0.877 ± 15 SD.

Importantly, the percentage of participants achieving a minimal clinically important difference, when combining all pain and functional outcome measures, was 84.1 and 87.1% in the 1 injection and two-injection groups, respectively (Table 6). Age and gender were not found to influence response to therapy.

See Supplementary Material for tabulated NPRS, KOOS and WOMAC results.

Structural analysis

MRI Osteoarthritis Knee Scores (MOAKS) were completed at baseline and at 12 months follow-up. One participant in the control group and one participant in the two-injection group were disregarded from MRI analysis due to considerable delay in 12 month imaging. Table 7 indicates the baseline prevalence and change in Global MOAKS at 12 months follow-up of the control and treatment groups (see Supplementary Material for regional results). A total of 67% of participants within the control group had progression of cartilage loss with a further 56% having extension of osteophyte formation. By comparison, in the one-injection group only 30% of participants had further cartilage loss although 50% had progression of osteophyte formation at 12 months. In the two-injection group 89% of participants had improvement in cartilage or no progression in cartilage loss with stabilization of OA also indicated by 89% having no progression in osteophyte formation. The two-injection group was the only group to
Table 6. Minimal Clinically Important Difference at 12 months in comparison to Baseline data for the three groups.

<table>
<thead>
<tr>
<th>Outcome measures</th>
<th>Control Group</th>
<th>One-injection group</th>
<th>Two-injection group</th>
</tr>
</thead>
<tbody>
<tr>
<td>NPRS</td>
<td>40</td>
<td>87.5</td>
<td>100</td>
</tr>
<tr>
<td>WOMAC</td>
<td>20</td>
<td>100</td>
<td>90</td>
</tr>
<tr>
<td>KOOS (pain improvement)</td>
<td>10</td>
<td>90</td>
<td>80</td>
</tr>
<tr>
<td>KOOS (symptom improvement)</td>
<td>30</td>
<td>66.7</td>
<td>70</td>
</tr>
<tr>
<td>KOOS (ADL)</td>
<td>30</td>
<td>77.8</td>
<td>90</td>
</tr>
<tr>
<td>KOOS (Sport &amp; Rec)</td>
<td>30</td>
<td>77.8</td>
<td>100</td>
</tr>
<tr>
<td>KOOS (QoL)</td>
<td>20</td>
<td>88.9</td>
<td>80</td>
</tr>
<tr>
<td>Averages</td>
<td>25.7%</td>
<td>84.1%</td>
<td>87.1%</td>
</tr>
</tbody>
</table>

All data are in percentages and expressed as the percentage of subjects who registered a MCID at 12 months when compared with their baseline value. Note that an MCID equates to a decrease of 1 point or more on a 10 point Numeric Pain Rating Scale and an increase of 8 points or more on a 100 point Knee Injury and Osteoarthritis Outcome Score or Western Ontario and McMaster Universities Arthritis Index score.

ADL: Activities of daily living; KOOS: Knee Injury and Osteoarthritis Outcome Score; NPRS: Numeric Pain Rating Scale; WOMAC: Western Ontario and McMaster Universities Arthritis Index.

Table 7. Baseline prevalence and change in Global MOAKS at 12 months follow-up.

<table>
<thead>
<tr>
<th>MOAKS features</th>
<th>Control group</th>
<th>One injection</th>
<th>Two injection</th>
<th>Chi-square p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bone marrow lesion</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>9/9(100%)</td>
<td>10/10(100%)</td>
<td>9/9(100%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>3/9 (33%)</td>
<td>3/10 (30%)</td>
<td>5/9 (56%)</td>
<td>0.474</td>
</tr>
<tr>
<td>– No change</td>
<td>4/9 (44%)</td>
<td>3/10 (30%)</td>
<td>3/9 (33%)</td>
<td>0.577</td>
</tr>
<tr>
<td>– Improvement</td>
<td>2/9 (22%)</td>
<td>4/10 (40%)</td>
<td>1/9 (11%)</td>
<td>0.122</td>
</tr>
<tr>
<td>Articular cartilage pathology</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>9/9 (100%)</td>
<td>10/10 (100%)</td>
<td>9/9 (100%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>6/9 (67%)</td>
<td>3/10 (30%)</td>
<td>1/9 (11%)</td>
<td>0.043</td>
</tr>
<tr>
<td>– No change</td>
<td>3/9 (33%)</td>
<td>7/10 (70%)</td>
<td>7/9 (78%)</td>
<td>0.117</td>
</tr>
<tr>
<td>– Improvement</td>
<td>0/9 (0%)</td>
<td>0/10 (0%)</td>
<td>1/9 (11%)</td>
<td>0.335</td>
</tr>
<tr>
<td>Osteophytes</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>9/9 (100%)</td>
<td>9/10 (90%)</td>
<td>9/9 (100%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>5/9 (56%)</td>
<td>5/10 (50%)</td>
<td>1/9 (11%)</td>
<td>0.107</td>
</tr>
<tr>
<td>– No change</td>
<td>4/9 (44%)</td>
<td>5/10 (50%)</td>
<td>8/9 (89%)</td>
<td>0.107</td>
</tr>
<tr>
<td>– Improvement</td>
<td>0/9 (0%)</td>
<td>0/10 (0%)</td>
<td>0/9 (0%)</td>
<td>0.996</td>
</tr>
<tr>
<td>Synovitis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>8/9 (89%)</td>
<td>7/10 (70%)</td>
<td>9/9 (100%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>2/9 (22%)</td>
<td>1/10 (10%)</td>
<td>0/9 (0%)</td>
<td>0.312</td>
</tr>
<tr>
<td>– No change</td>
<td>4/9 (44%)</td>
<td>7/10 (70%)</td>
<td>6/9 (67%)</td>
<td>0.474</td>
</tr>
<tr>
<td>– Improvement</td>
<td>3/9 (33%)</td>
<td>2/10 (20%)</td>
<td>3/9 (33%)</td>
<td>0.756</td>
</tr>
<tr>
<td>Meniscus pathology</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>9/9 (100%)</td>
<td>10/10 (100%)</td>
<td>9/9 (100%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>1/9 (11%)</td>
<td>1/10 (10%)</td>
<td>0/9 (0%)</td>
<td>0.598</td>
</tr>
<tr>
<td>– No Change</td>
<td>8/9 (89%)</td>
<td>9/10 (90%)</td>
<td>8/9 (89%)</td>
<td>0.996</td>
</tr>
<tr>
<td>– Improvement</td>
<td>0/9 (0%)</td>
<td>0/10 (0%)</td>
<td>1/9 (11%)</td>
<td>0.335</td>
</tr>
<tr>
<td>Popliteal cyst</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>– Baseline prevalence</td>
<td>3/9 (33%)</td>
<td>5/10 (50%)</td>
<td>3/9 (33%)</td>
<td></td>
</tr>
<tr>
<td>– Progression</td>
<td>0/9 (0%)</td>
<td>1/10 (10%)</td>
<td>1/9 (11%)</td>
<td>0.598</td>
</tr>
<tr>
<td>– No change</td>
<td>8/9 (89%)</td>
<td>8/10 (80%)</td>
<td>8/9 (89%)</td>
<td>0.813</td>
</tr>
<tr>
<td>– Improvement</td>
<td>1/9 (11%)</td>
<td>1/10 (10%)</td>
<td>0/9 (0%)</td>
<td>0.598</td>
</tr>
</tbody>
</table>

Note the significant difference in proportions (Chi-square analysis) with articular cartilage pathology progression but with an average power of the performed tests being only 25.
Table 8. Adverse events for the one- and two-injection treatment groups. Note that no significant difference in the proportions (Chi-square analysis) of adverse events between groups were apparent (p = 0.303) with the power of the performed test being 0.47.

<table>
<thead>
<tr>
<th>Adverse events</th>
<th>One injection (baseline)</th>
<th>Two injections (baseline injection)</th>
<th>Two injections (6 month injection)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nil</td>
<td>20%</td>
<td>10%</td>
<td>0%</td>
</tr>
<tr>
<td>Mild</td>
<td>60%</td>
<td>50%</td>
<td>40%</td>
</tr>
<tr>
<td>Moderate</td>
<td>10%</td>
<td>30%</td>
<td>60%</td>
</tr>
<tr>
<td>Severe</td>
<td>10%</td>
<td>10%</td>
<td>0%</td>
</tr>
<tr>
<td>Serious</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
</tr>
</tbody>
</table>

show improvement in cartilage loss based on MOAKS analysis (a single participant). Bone marrow lesions (BMLs), synovitis, meniscus pathology or popliteal cysts showed no appreciable difference between groups.

Complications & adverse events

Minor discomfort and bruising was commonly noted in both treatment groups after their lipoharvest procedure. This resolved without further intervention indicating a mild expected AE. Table 8 indicates AEs associated with the intra-articular injections of autologous ADMSC therapy at baseline and also at 6 months in the two-injection group. Many participants in the treatment groups experienced discomfort and/or swelling post ADMSC therapy. These AEs were self-limiting, requiring a period of unloading, analgesia and/or oral anti-inflammatory use. Two participants reported pain and swelling for 4 weeks following ADMSC therapy and due to observed impact on their usual daily activity this was categorised as a severe AE. The second injection of MSC therapy at 6 months in the two-injection group was associated with a modest increase in reported moderate AEs in comparison to the initial injection. No related serious AEs were observed.

Discussion

Previous studies have indicated the potential benefit of MSC therapy in symptomatic knee OA. It is important to differentiate the results of this trial with past trials using more heterogeneous cell based products including stromal vascular fraction [51,52]. This study is the first to assess and compare the safety and efficacy of different treatment protocols with single versus multiple injections and using autologous isolated and expanded ADMSCs. The results indicate that autologous MSC therapy is a safe and effective treatment modality for moderate knee OA with the ability to modify disease progression.

Pain and functional improvement was comparable in both treatment groups. Both statistical and clinically significant change (MCID) was observed in comparison to both baseline and the control group using the NPRS, WOMAC and KOOS subscale scores. The two injection group had an observed mild reduction in improvement in KOOS subscale scores at 6 months although this was as a direct consequence of outcome questionnaires being completed by some participants in the period directly after their second injection (which was associated with a self-limiting period of pain and swelling). Despite some studies indicating stem cell age may be associated with reduced function, this trial did not observe any age-related difference in pain and functional outcome [53].

Pain and functional improvements observed in this pilot RCT compared favorably against conventional treatments for OA. NPRS improved by a percentage of 69% from baseline to final follow-up at 12 months in both treatment groups. This is in comparison to a <14% improvement in pain scores at 12 months following arthroscopic debridement or up to 12% improvement with a prescribed exercise program [54,55].

Past clinical trials had indicated the ability of intra-articular MSC therapy to modify the progression of OA. Jo and colleagues observed increased cartilage volume at 6 months following MSC therapy, with a later study using allogeneic bone marrow derived MSCs successfully stabilizing OA and improving observed cartilage quality [33,35]. Using the semi-quantitative and validated MOAKS assessment to describe structural change over 12 months, the two injection protocol was observed to achieve a trend of greater disease stabilization in comparison to the single injection group. Observed stabilization rather than cartilage regrowth suggests that the modality of action of MSCs is via paracrine and local supportive pathways rather than direct differentiation along a chondrocyte lineage. This is consistent with preclinical trials which have failed to show significant incorporation of labeled MSCs into regenerative cartilage tissue [28].
Surprisingly, features including BMLs, synovitis and the presence of a popliteal cyst were not associated with pain and functional outcome. Several participants in both treatment groups exhibited clinically significant pain improvement despite no change or in fact progression in BMLs. The lack of correlation of improvement with BMLs and synovitis/effusion is in contrast to previous studies indicating these features to be strongly associated with pain in OA \([56,57]\). This perhaps reflects the observation of lack of improvement in knee pain and function following treatment with bisphosphonates (residronate) in previous RCTs \([58,59]\). Furthermore, MOAKs analysis indicating successful stabilization despite persistent BMLs is also in contrast to past research which has shown an association between BMLs and progression in OA \([60]\).

Both treatment protocols were well tolerated with no serious AE documented from either the stem cell harvest technique (liposuction) or ADMSC therapy. Whilst two participants experienced pain and swelling, which was classed as a severe AE, this was managed appropriately with a period of unloading and oral analgesia/anti-inflammatories and resolved without further intervention required. All AEs were expected and similar to past adverse outcomes reported in previous trials using intra-articular MSC therapy. A modest increase in reported moderate AEs at the second injection in the two-injection group suggests that timing of repeat MSC therapy may influence therapy tolerability.

The main limitation in this RCT is lack of a formal placebo with all groups unblinded to treatment. With the requirement of a lipoharvest procedure to harvest autologous ADMSCs it was deemed unethical to perform an unnecessary surgical procedure on the control group if they were later only to receive a sham placebo injection. This trial represents a pilot study and further research with greater participant numbers is warranted. The MRI analysis is limited in that MOAKS is a semi-quantitative measure. In future research the authors would consider the use of more precise MRI quantitative measures including cartilage volume as described by Cicuttini and colleagues and also the novel and recently validated method of cartilage quality assessment using T2 mapping techniques \([61,62]\).

**Conclusion**

OA is a cause of significant morbidity and with an aging population it promises to remain an enormous community and economic burden. Current conservative management strategies fail to alter disease progression and surgical management in the form of joint replacement is associated with considerable risk. In this RCT, autologous ADMSC therapy was associated with clinically significant improvement in pain and function in symptomatic knee OA. These improvements were well beyond that achieved by conventional methods. Furthermore, ADMSCs were associated with successful disease modification with comparison of treatment protocols indicating that two intra-articular injections of ADMSCs separated by 6 months achieved more consistent OA stabilization than a single injection. Intra-articular autologous ADMSC therapy was shown to be safe with no serious AEs.

**Translational perspective**

Regenerative cellular therapies rather than being unique and novel are now well-established and form part of accepted medical practice in areas including bone marrow and tissue transplantation, blood transfusion and reproductive in vitro fertilization. First described by Dr Alexander Friedenstein over 40 years ago, MSCs, with their observed multipotency and presence throughout the body, have been postulated as a future biological cell therapy for tissue repair and regeneration. Recent years have seen a significant increase in the number of clinical trials involving MSCs with international groups including the International Society for Stem Cell Research publishing guidelines for appropriate stem cell research and clinical translation \([63]\). An acknowledgment of the various different cellular therapies being both studied and used clinically in the area of musculoskeletal medicine, and yet often described in broad inclusive terminology, has seen the development of a minimum information for studies evaluating biologics in orthopedics (MIBO) \([64]\).

This trial represents the first published RCT on the use of autologous isolated and expanded ADMSCs in OA. Significant pain and functional improvement and observed disease modification with structural stabilization indicate that ADMSC therapy may represent an exciting advancement in the treatment of OA. Additional research with direct comparison against accepted conventional treatment options needs to be conducted.

**Supplementary data**

To view the supplementary data that accompany this paper please visit the journal website at: www.futuremedicine.com/doi/full/10.2217/rme-2018-0161
Summary points

- Osteoarthritis (OA) is the fourth leading cause of disability worldwide.
- Symptomatic arthritis affects 10% of males and 18% of females over the age of 45 years.
- 80% of patients over the age of 65 have radiological evidence of OA.
- Current conservative management strategies are aimed at symptom control and do not modify the progression of OA. Pharmaceutical treatments are limited and may have serious side effects.
- Surgical total joint replacement remains the accepted treatment for knee OA uncontrolled by conservative therapies and is not without complication with up to 20% of patients experiencing persistent pain and loss of function at 12 months.
- Initial preclinical and clinical trials have indicated the ability of intra-articular MSC therapies to improve pain and function.
- This is the first randomized controlled trial to assess the safety and efficacy of autologous isolated and expanded adipose derived mesenchymal stem cell (ADMSC) therapy in symptomatic knee OA.
- Over 12 months of follow-up statistical and importantly clinically significant pain and functional improvements were observed in both treatment groups against control and baseline values as measured by validated outcome scores including Numeric Pain Rating Scale, Western Ontario and McMaster Universities Arthritis Index and Knee Injury and Osteoarthritis Outcome Score.
- Semi-quantitative radiological analysis using the MRI based Osteoarthritis Knee Score assessment indicated modification of disease progression over 12 months of follow-up with greatest stabilization achieved by the treatment group receiving two injections of ADMSCs.
- No serious adverse events were observed.
- Improvement in pain scores compared favorably to previous studies on the effectiveness of current conventional therapies.
- Autologous ADMSC therapy represents a potential exciting advancement in the active management of OA.

Financial & competing interests disclosure

J Freitag, D Bates, L Huguenin, A Tenen are clinic partners within Melbourne Stem Cell Centre.

J Freitag, D Bates, R Boyd, K Shah, L Huguenin, A Tenen are associated with Magellan Stem Cells and are part of its Medical and Scientific Advisory Committee.

This proposed study was funded jointly by both Magellan Stem Cells and Melbourne Stem Cell Centre. Members of their Medical and Scientific Advisory board have been involved in the study conception and design and are listed as co-authors of this paper. The authors received no payment for their involvement in the study. Interpretation of results, and subsequent submission and publication decisions have been made independent of the sponsors.

The authors have no other relevant affiliations or financial involvement with any organization or entity with a financial interest in or financial conflict with the subject matter or materials discussed in the manuscript apart from those disclosed.

Author contribution

J Freitag, D Bates, R Boyd, K Shah, L Huguenin, A Tenen, K Paterson were involved in conception of the study. J Freitag, D Bates, J Wickham, K Shah, R Boyd, L Huguenin, A Tenen, K Paterson designed the study protocol. J Wickham was involved in developing the statistical analysis strategy.

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- R Castelluccio – Nurse Project Manager.
- M Grogan and E Picken – Study Co-ordinators.
- N Shah, I Wickramasinghe and M Lopez – Laboratory Staff.

Ethical conduct of research

Ethics approval was given for the study by the Human Research Ethics Committee of Monash University and registered on the Australian and New Zealand Clinic Trial Registry (Trial ID: ACTRN12614000814673). In addition, informed consent has been obtained from the participants involved.
Data sharing statement

Individual de-identified participant data which underlie results reported in this article will be available upon publication. The study protocol will also be available. Material will be accessible to investigators whose proposed use of data has been approved by an independent review committee and for data meta-analysis. Requests are to be directed to the corresponding author of the article.

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