The use of hip injection (HI) in the treatment of osteoarthritis (OA) has gained wide popularity. The relatively low cost, fast and simple method of pain relief are amongst its many advantages. Over time, the content of the injection has also evolved from local anesthetic (LA) agents to corticosteroids (CSs), hyaluronic acid (HA) and platelet-rich plasma (PRP).[1] The scope of use of injections in the hip region has grown from traditional aspiration to therapeutic injections. The two main substances used in recent times for pain relief are CSs and HA gel. For decades, low doses of CS were given to surgically unfit patients and to those who are not keen on joint replacement surgery.[2]
The recent surge in the use of high-molecular-weight HA for knee OA has been expanded as a treatment option for hip OA. The popularity of the administration of HA has been mounting with very little outcome data to support its use. Administration of HA injections has shown some promise in a selected subset of patients suffering from early OA of the hip.\textsuperscript{[3,4]} Most papers report insufficient sample size and had a varied follow-up period which results in difficulty formulating and implementing national guidelines and clinical recommendations. Current literature advocates the safe use of CS injections for early hip OA.\textsuperscript{[5]} Although there is no concrete evidence supporting HA injections, this has not dissuaded researchers from injecting PRP, mesenchymal stem cells (MSCs), LA agents, NSAIDS and many different combinations into the hip. The true extent of their benefits is still being debated.\textsuperscript{[6]} In this review, we outline recent trends, discuss the role of HIs, and summarize complications of the technique.

**SEARCH STRATEGY AND SELECTION CRITERIA**

We conducted a review of literature on intraarticular injections for OA of the hip. The PubMed, Cochrane Libra and DOAJ were accessed, and articles written in the English language with keywords “hip osteoarthritis injection”, and those that published relevant literature on humans were included in our search. We focused on publications from the past 11 years (2010 to 2021). Meta-analyses, systematic reviews, randomized and non-randomized clinical trials on hip OA that were published in the English language were included. All other types of articles were excluded from this review. On typing the keywords “hip osteoarthritis and hip injection”, a total of 785 articles were identified in the search, out of which only 232 articles were found to be relevant and hence selected. Out of these 232, 141 articles were excluded due to the unsuitable nature of the article, and lack of well-defined inclusion and
The efficacy of intraarticular hip injection for patients with Hip OA

CONTEMPORARY TRENDS AND CLINICAL RECOMMENDATIONS

Intraarticular HIs have been administered for decades. As early as 1947, Crowe reported satisfactory results with intraarticular acid phosphate injections for the treatment of hip OA. He also recommended the anterior approach for administration of the injection as the easiest, least painful, and most accurate. In 1956, Leveaux and Quin published their results on “Local injection of hydrocortisone and procaine in osteoarthritis of the hip joint” and concluded that the combination of these two substances was of valuable palliative management for the painful osteoarthritic hip joint. On the other hand, the American Academy of Orthopaedic Surgeons (AAOS) guidelines on the management of OA of the hip published in 2017 considered only intraarticular CSs and HA worthy of any recommendation. AAOS supported the use of intraarticular CSs to improve short-term function and pain for patients with symptomatic OA of the hip. However, they did not support the use of intraarticular HA, citing equal efficacy to placebo for function, stiffness, and pain in patients with symptomatic OA of the hip. In a meta-analysis conducted by Gazendam et al., only minimal clinically important differences were observed from the baseline to six months after all HIs, and the results were similar in the intervention and placebo groups.

ROLE OF INTRAARTICULAR CORTICOSTEROIDS FOR HIP OSTEOARTHRITIS

Synovitis is a major cause of pain in hip OA. Local anti-inflammatory treatment such as intraarticular CS is effective in ameliorating pain in OA of the hip. Downregulating genetic expression of several proinflammatory proteins and limiting the interaction between white blood cells involved in immune response appears to be the mechanism of action of the therapy. These injections are frequently given in combination with an LA agent. Methylprednisolone or triamcinolone is combined with 1% lidocaine or 0.5% bupivacaine. The dose administered depends on patient-specific factors and surgeon experience. The dose of methylprednisolone ranges from 40 to 120 mg, while the dose of triamcinolone ranges from 20 to 80 mg. These agents have been selected by the virtue of them being less soluble in water (particulate CSs). The more the solubility of a particular steroid injection, the less the duration of effect of the injection. Only preservative-free anesthetics must be used as a solvent for these steroids to prevent particulate precipitation. Mixing steroids with an LA agent has distinct advantages of reducing infiltration discomfort, increasing the volume of the injection and better distribution of the solution throughout the joint.

McCabe et al. published their systematic review on the efficacy of intraarticular steroids in hip OA in 2016. They recommended methodologically rigorous trials to verify whether intraarticular CSs were beneficial, and the duration of efficacy. They included randomized-controlled trials (RCTs) assessing the efficacy of hip intraarticular steroid injection on pain. A total of five RCTs were included though all were marred by a limited sample size. They determined that these injections were well tolerated and were effective in reducing some amount of pain for up to four weeks.

In 2018, Lai et al. published a retrospective analysis of all intraarticular hip steroid injections performed for hip OA between January 2010 and December 2012. Around 20% showed no response, less than 50% showed an immediate response (≤2 weeks of pain relief), and the remaining showed a continued response (>2 weeks of pain relief). Age, obesity, duration of symptoms and radiological grading of hip OA were not found to have a significant correlation with intraarticular CS injection. Total hip replacement within two years was required in almost 50% of the patients, which ultimately led to the authors’ recommendation of considering hip arthroplasty early in the disease.

Multiple authors have reported that the effect of intraarticular steroid administration on pain and function disappears rapidly, but Deshmukh et al. reported that steroids could provide long-term relief, unlike others. Additionally, it was reported that there was a relationship between the reduction of pain and the severity of the disease, contrary to Lai et al.

Moreover, since the coronavirus pandemic has drastically reduced the occurrence of elective arthroplasty surgery, intraarticular steroids are an effective intervention for patients awaiting joint replacement surgery. However, these injections are not without risks such as the risk of infection and progression of cartilage damage. Therefore, we should remember to use these injections judiciously.
ROLE OF INTRAARTICULAR VISCOSUPPLEMENTATION FOR HIP OSTEOARTHRITIS

Hyaluronic acid has been said to improve the rheological properties of synovial fluid with some chondroprotective and anti-inflammatory effects. Although the positive clinical effects of HA have been demonstrated in the knee joint, guidelines and the latest literature recommend otherwise for its use in the hip. In 2017, Eymard et al., on behalf of the Osteoarthritis Group of the French Society of Rheumatology and the French Research Group in Interventional Rheumatology, published their results from a multi-center, open-label, prospective, trial. They elaborated on the subset of hip OA patients that benefited the most by a single intraarticular injection of a cross-linked HA combined with mannitol on Day 90. Patients with moderate pain, moderate disability, moderate joint space narrowing, superomedial and axial femoral head migration, with femoral-acetabular impingement or coxa profunda displayed better results. This drug combination was able to decrease pain by 50%, in half of their patients at day 90.

Benefits in an almost similar subset of patients have been reported by other authors, as well. Pogliacomi et al. in their original study published in 2018 reported the efficacy of intraarticular HI of a single dose of high-weight HA in patients under a follow-up of 12 months. They concluded that patients with a moderate grade of OA are the ones that benefit the most from the said injection.

Extravasation of HA has been shown to cause inflammation of periarticular tissues and must be avoided, as it can sometimes mimic a septic reaction. Acute local reactions have been associated with multiple injections, leading to the conclusion that a single injection is more beneficial. In a meta-analysis focused on adverse reactions after HA HIs; Wu et al. could not find an increased adverse reaction rate with HA compared to controls.

ROLE OF INTRAARTICULAR PLATELET-RICH PLASMA FOR HIP OSTEOARTHRITIS

We did not find high-quality studies comparing PRP with placebo up to April 15th, 2016. In 2018, the Royal Australian College of General Practitioners released their guideline for the management of knee and hip OA, and due to a lack of high-quality evidence, they were unable to make any recommendation for PRP injections.

Platelet-rich plasma is a biological treatment with great perspective, but standardization of treatment is lacking, and this has led to conflicting evidence on the effect of intraarticular PRP injections for the management of hip OA. Dong et al. published a meta-analysis of high-powered RCTs conducted on the effect of intraarticular PRP on OA, up to June 2019. They included three RCTs on hip OA and concluded that large scale double-blinded RCTs are required to evaluate the effect of PRP injection in hip OA. An RCT comparing the efficacy of ultrasound-guided intraarticular injections of PRP versus HA for hip OA, published by Battaglia et al., recognized the efficacy of PRP in terms of functional enhancement and decrease in pain, but concluded that PRP was not superior to HA up to 12-month follow-up.

Di Sante et al. published an RCT comparing the efficacy of PRP vs HA injection in hip OA and established that intraarticular PRP had a pain-relieving effect on hip OA that lasted up to four weeks. The PRP is being marketed as a promising new product of regenerative medicine that is superior to other current therapies. However, unfortunately, it still lacks robust evidence to support its use in clinical practice. Orthopedic surgeons should be aware of the ongoing uncertainty about the evidence behind PRP therapies and inform patients about this fact. Overall, there is insufficient evidence to support the use of PRP for hip OA in the current literature.

ROLE OF INTRAARTICULAR MESENCHYMAL STEM CELLS FOR HIP OSTEOARTHRITIS

A study from Iran by Emadedin et al. published their results on injections of autologous bone marrow-derived MSCs for patients with OA of the hip, knee, and ankle. A total of 17 patients were evaluated clinically, as well as with magnetic resonance imaging (MRI) scans for a period of up to 30 months post-injection. They did not observe any serious adverse effects (systemic effects, tumors, pulmonary embolism, and death) and found the injection to be safe. Additionally, the study group had decreased pain, improved walking ability and functional scores. Furthermore, an increase in cartilage thickness and a decrease in subchondral oedema was observed on MRI. A major drawback of this study was the small sample size, particularly while evaluating effects on hip OA.

Researchers from Chile published their well-designed study of 10 patients who were more than 60 years old and were suffering from hip OA (up to moderate grade). These patients were given an intraarticular injection of ex vivo expanded autologous bone marrow-derived MSCs. Patients without pain or mild pain were not included in the study. Pain,
stiffness, functionality, and range of motion were evaluated. Patients were followed up to a maximum of 40 weeks. Improvement in all the clinical parameters was noted in all, but one patient. The radiographic progression of OA was also arrested and these occurred without any major side effects.\[30\]

A systematic review published in 2018 included 28 studies for critical review. Although HIs were a minuscule part of the whole study group; the general trend was favorable to MSC injections without major complications.\[31\] Considering the potential of stem cell therapy, there is still a need for high-quality research on this topic.

**NOVEL INTRAARTICULAR INJECTIONS FOR HIP OSTEOARTHRITIS**

A combination of MSCs and constituents of PRP is known as bone marrow aspirate concentrate (BMAC). The BMAC is one of the few United States Food and Drug Administration (FDA) approved methods for providing stem cells.\[32\]

A study by Rodriguez-Fontan et al.\[33\] included 25 joints (10 knees, 15 hips), that were injected with intraarticular BMAC. Only patients with Kellgren-Lawrence Grade I-II/ Tönnis Grade I-II were included in the study. Maximum follow-up was up to 24 months. A total of 63.2% of patients were satisfied with the procedure and this injection was found to be safe. Darrow et al.\[34\] published their study of repeated BMAC injections in the hip for OA. All patients reported decreased pain and improved function. They hypothesized that multiple injections in a short period were responsible for this significant improvement from baseline. The BMAC has great potential, and large-scale, placebo-controlled RCTs can pave the way for a futuristic regenerative method of treatment for early to moderate hip OA.

Another treatment modality that is worth mentioning due to its low cost is prolotherapy; a technique that has seen more waxing and waning than any other therapy in orthopedic medicine. It has recently gained popularity for hip OA. However, placebo-controlled trials are still lacking.

A treatment method that is gaining popularity is the intraarticular injection of NSAIDs. They are less potent anti-inflammatory agents (as compared to CSs). Park et al.\[35\] published the results of their retrospective comparative study where 50 patients received intraarticular CS injection and 48 received intraarticular ketorolac injection. Ketorolac HI was found to be as effective as CS HI.

**COMPLICATIONS OF INTRAARTICULAR HIP INJECTIONS**

Procedural complications such as pain and bleeding at the injection site are easily manageable. The fear of septic arthritis although real is exceedingly rare, if proper aseptic precautions have been followed.\[36\] In studies on a substance other than steroids and LAs, complications related to injection have been reported with local adverse effects.\[27,29\]

Patients planned for intraarticular steroid injection must be warned of the side effects such as mild headaches, slight exaggeration in pain (steroid flare), lack of sleep and facial flushing (particularly common in females).\[37\] Diabetics often enquire regarding the influence of these injections on their blood glucose levels. Clinicians must put forward the available evidence that a transient spike in blood glucose levels lasting less than a week can occur in selected individuals after CS injections.\[37\] There is also an increased risk of infection in diabetic patients undergoing intraarticular steroid injections.

The list of complications of intraarticular steroids would be incomplete without mentioning the so-called “corticosteroid arthropathy”.\[38\] Having said that, a study with human subjects did not display radiological evidence of cartilage damage, despite repeated injections of intraarticular CSs in their knees for over two years.\[39\] Inversely, an annual MRI-based study comparing intraarticular triamcinolone and intraarticular saline every 12 weeks for two years found that triamcinolone resulted in significantly greater cartilage volume loss.\[40\]

In the practice of orthopedics, there is one common fear in the usage of intraarticular injections, which is the risk of postoperative infection in the hip that has been previously injected with a CS. Steroids are well known to dampen the intraarticular immune response. Extensive studies have been done on this topic. Werner et al.\[41\] proved beyond doubt that the interval between HI and hip arthroplasty must be maintained for more than three months to mitigate the risk of prosthetic joint infection. These studies would probably be quoted more frequently in the coming years as more patients are being treated with intraarticular HIs in recent times due to the coronavirus pandemic putting a halt to elective arthroplasty surgeries.

Another popular opinion is the chondrotoxic effect of LA agents used either alone or along with particulate CSs for intraarticular injection. Jayaram et al.\[42\] recognized that there was variability in the chondrotoxic effect of different LA agents and there was no consensus on an ideal LA for intraarticular...
They carried out a systematic review on the effect of LA agents on knee cartilage. Literature from 1990 to 2018 was utilized with 16 studies included using chondrocyte viability, morphology, and histology as the outcome measures. Data related to commonly used LA agents such as lidocaine, bupivacaine, ropivacaine, levobupivacaine, and mepivacaine were extracted. Each drug had different spectrums of chondrocyte damage culminating in increased apoptosis, extracellular matrix damage and mitochondrial dysfunction. These effects were dose- and duration-dependent and concomitant CS administration was found to exacerbate these effects. The toxicity spectrum was found to be maximum for bupivacaine and the least for ropivacaine (in concentrations less than 0.75%). However, we should keep in mind that these observations were largely based on in vitro studies.

**IMAGING-GUIDED INTRAARTICULAR INJECTION**

The intraarticular injection procedure can be technically challenging due to the deep placement of the hip joint. Unsuccessful injection potentially exposes the patient to complications from the procedure (pain, infection, bleeding) and may contribute to diagnosis and treatment delays. Intraarticular HI with the help of anatomic landmarks is frequently performed over the signs defined for hip arthroscopy by Wettstein and Dienst[42] for accessing the peripheral compartment of the hip joint. However, the success rate of HIs without imaging guidance is not satisfactory. A meta-analysis about success rates of injection techniques reported that operators using ultrasound had an injection success of 100%, while operators using anatomical landmark-guided injections were 72% accurate.[43]

The most used imaging method for intraarticular injections of the hip is the fluoroscopic technique. Radiopaque contrast material can be used for confirming that the needle is in the joint. However, access to the fluoroscopy device can sometimes be not possible. Due to the difficulty, ultrasound guidance, which can be easily applied in office conditions, is recommended for HIs by several authors.[44,45] In an animal experiment evaluating the success rate, the accuracy was 90% for the ultrasound-guided procedure and 75% for the fluoroscopy-guided intraarticular injection. There was a statistically significant difference between the two groups.[46] Computed tomography (CT) can theoretically provide joint injection with high success, but it is not preferred due to the high radiation exposure of the patient and operator. There is a rare need to CT guidance for magnetic resonance arthrography, when qualified personnel are not available, with reduced radiation dose can be used for contrast agent injection (Table 1).[47]

In conclusion, intraarticular CSs have established themselves as a pain-alleviating agent for hip OA; however, they must not be administered frequently, as a gap of three to six months must be maintained after the injection and before hip arthroplasty surgery. Additionally, the choice of LA that is combined with CS must be evidence-based and the clinician must be aware of the chondrotoxic effects of LA agents. The HA has not been able to replicate the results obtained in knee OA. At best, it has shown a promising effect in moderate grade hip OA. Biological agents including PRP/ MSC/ BMAC have great potential, but unrestricted usage cannot be recommended, as high-quality evidence is still lacking in modern literature.[48]

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