

Cigna Medical Coverage Policies – Musculoskeletal Knee Surgery: Arthroscopic and Open Procedures

Effective May 31, 2023



Instructions for use

The following coverage policy applies to health benefit plans administered by Cigna. Coverage policies are intended to provide guidance in interpreting certain standard Cigna benefit plans and are used by medical directors and other health care professionals in making medical necessity and other coverage determinations. Please note the terms of a customer's particular benefit plan document may differ significantly from the standard benefit plans upon which these coverage policies are based. For example, a customer's benefit plan document may contain a specific exclusion related to a topic addressed in a coverage policy.

In the event of a conflict, a customer's benefit plan document always supersedes the information in the coverage policy. In the absence of federal or state coverage mandates, benefits are ultimately determined by the terms of the applicable benefit plan document. Coverage determinations in each specific instance require consideration of:

1. The terms of the applicable benefit plan document in effect on the date of service
2. Any applicable laws and regulations
3. Any relevant collateral source materials including coverage policies
4. The specific facts of the particular situation

Coverage policies relate exclusively to the administration of health benefit plans. Coverage policies are not recommendations for treatment and should never be used as treatment guidelines.

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CMM-312:Knee Surgery - Arthroscopic and Open Procedures

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Definitions

- **Arthrofibrosis:** a condition of the appendicular skeletal system that has resulted from disease, injury, or surgery, and results in pain and restricted range of motion due to internal scarring of the joint with consequent stiffness.
- **Autologous Chondrocyte Implantation (ACI) or Autologous Chondrocyte Transplantation (ACT):** a cell-based cartilage repair surgical technique which utilizes an individual's own cells in an effort to repair damage to articular cartilage with the goal of improving joint function and reducing pain. The procedure involves the collection and culture of articular cartilage cells (i.e., chondrocytes) that are then implanted into the cartilage defect with the intent that the cultured cells will contribute to the regeneration and repair of the articular surface.
 - ◆ **Hybrid autologous Chondrocyte Implantation (ACI):** ACI is combined with other surgical repair techniques of cartilage defects (e.g., osteochondral autograft transfer).
- **Kellgren-Lawrence Grading System:** a radiographic grading system that has been developed for describing osteoarthritic changes to the tibial-femoral joint of the knee. When used, the radiographic findings on plain x-rays are typically reported within one of the following categories:
 - ◆ **Grade 0** – No radiographic features of osteoarthritis are present
 - ◆ **Grade I** – Doubtful narrowing of joint space and possible osteophytic lipping
 - ◆ **Grade II** – Definite osteophytes and possible narrowing of joint space
 - ◆ **Grade III** – Moderate multiple osteophytes, definite narrowing of joint space, some sclerosis, and possible deformity of bone contour
 - ◆ **Grade IV** – Large osteophytes, marked narrowing of joint space, severe sclerosis, and definite deformity of bone contour
- **Kissing Lesion:** an articular cartilage defect on opposing joint surfaces of the knee and that are in contact either between the patella and distal femur or the distal femur and tibia (e.g., bipolar lesion).
- **MACI[®] Implant (Vericel Corporation, Cambridge, MA [formerly Genzyme Biosurgery]):** Until recently, Carticel[®] (Vericel Corporation, Cambridge, MA [formerly Genzyme Biosurgery]) was the only technology that received FDA approval for the culturing of chondrocytes. MACI[®] Implant (Matrix Induced Autologous Chondrocyte Implant) received approval from the U.S. Food and Drug Administration December 2016 as an autologous cellularized scaffold indicated for repair of single or multiple symptomatic, full-thickness cartilage defects of the knee with or without bone involvement in adults. MACI[®] Implant is utilized as part of an ACI procedure in which cartilage cells are removed during arthroscopy, and shipped to a laboratory, where the cells are cultured over a period of several weeks. The cells are seeded on a porcine collagen membrane, and once the culturing process is complete, the cells seeded on the membrane are returned to the surgeon for implantation during the procedure. The membrane is placed into the defect, and over several months the cells create a matrix that is intended to cover the articular surface of the knee. The safety and effectiveness of MACI[®] Implant in joints other than the knee has not been established.
- **Modified Outerbridge Classification:** a system that has been developed for judging articular cartilage injury to the knee. This system allows delineation of

varying areas of chondral pathology, based on the qualitative appearance of the cartilage surface as viewed on MRI and can assist in identifying those injuries that are suitable for repair techniques. The characterization of cartilage in this system is as follows:

- ◆ **Grade I** – Softening with swelling
 - ◆ **Grade II** – Fragmentation and fissuring less than one square centimeter (1 cm²)
 - ◆ **Grade III** – Fragmentation and fissuring greater than one square centimeter (1 cm²)
 - ◆ **Grade IV** – Subchondral bone exposed
- **Mosaicplasty** (or osteochondral cylinder transplantation): a surgical technique which consists of harvesting cylindrical bone-cartilage grafts and transplanting them into focal chondral or osteochondral defects in the knee. After excision of the chondral lesion, an abrasion arthroplasty is performed to refresh the base of the defect. The grafting procedure involves collecting grafts from the posterior aspect of the distal femoral articular surfaces (medial condyle, lateral condyle or trochlea) and implanting the grafts in a mosaic-like pattern that will contribute to regeneration and repair the articular surface. A recipient tunnel is created and sized with a drill bit slightly larger than the length of the graft. The harvested graft is placed in the tunnel by a press-fit method. All subsequent grafts are inserted in a similar pattern.
- **Non-surgical management** (with regard to the treatment of knee pain): any provider-directed non-surgical treatment which has been demonstrated in the scientific literature as efficacious and/or is considered reasonable care in the treatment of knee pain. The types of treatment involved can include, but are not limited to: ice, relative rest/activity modification, acupuncture, weight loss, supervised physiotherapy modalities, and therapeutic exercises, prescription and non-prescription medications, assistive devices (e.g., brace, cane, crutches, walker, wheelchair), and/or intra-articular injections (e.g., steroid and/or viscosupplementation).
- **Osteochondral Allograft Transplantation (OATS Procedure)**: a procedure that is similar to mosaicplasty, involving the use of a larger, single plug that usually fills an entire defect. It is often performed to graft chondral defects that are also associated with anterior cruciate ligament (ACL) tears. This method allows arthroscopic access to both the ACL and the chondral defect for the performance of a repair and the grafting procedure.
- **Outerbridge Classification**: a system that has been developed for judging articular cartilage injury to the knee. This system allows delineation of varying areas of chondral pathology, based on the qualitative appearance of the cartilage surface as viewed by direct visualization intraoperatively, and can assist in identifying those injuries that are suitable for repair techniques. The characterization of cartilage in this system is as follows:
- ◆ **Grade I** - Softening with swelling
 - ◆ **Grade II** - Fragmentation and fissuring less than one square centimeter (1 cm²)
 - ◆ **Grade III** - Fragmentation and fissuring greater than one square centimeter (1 cm²)
 - ◆ **Grade IV** - Subchondral bone exposed

- **Subchondral Drilling or Microfracturing:** a surgical procedure which is performed after the calcified cartilage is debrided and the surgeon creates tiny fractures in the adjacent bones (through the use of an awl). Blood and bone marrow (which contains stem cells) seep out of the fractures, creating a blood clot that releases cartilage-building cells. The microfractures are treated as an injury by the body, which is why the surgery results in new, replacement cartilage. Studies have shown that microfracturing techniques do not fill the chondral defect fully and the repair material that forms is fibrocartilage. Fibrocartilage is not as mechanically sound as the original hyaline cartilage; it is much denser and isn't able to withstand the demands of everyday activities as well as hyaline cartilage and is, therefore, at a higher risk of breaking down. The procedure is less effective in treating older individuals, overweight individuals, or in larger cartilage lesions. Furthermore, chances are high that after only one or two years, symptoms start to return as the fibrocartilage wears away, forcing the individual to reengage in articular cartilage repair.

General Guidelines

Application of Guideline

- The determination of medical necessity for the performance of knee surgery is always made on a case-by-case basis.
- Manipulation of a knee joint under general anesthesia is included in all arthroscopic knee procedures and is therefore considered incidental to the base procedure requiring medical necessity review.
- For advanced imaging indications for conditions about the knee, refer to **MS-25: Knee**
- For coverage indications for articular cartilage allograft materials, please reference the **Cigna Medical Coverage Policy: 0118 Bone, Cartilage and Ligament Graft Substitutes**

Arthroscopic or Open Procedure for Fracture, Tumor, Infection, or Foreign Body

Arthroscopic or open knee surgery may be considered **medically necessary** for **ANY** of the following:

- When surgery is being performed for fracture, tumor, infection or foreign body that has led to or will likely lead to progressive destruction.

Diagnostic Arthroscopy

Diagnostic Arthroscopy Indications

- Diagnostic arthroscopy is considered **medically necessary** as a stand-alone procedure when **ALL** of the following criteria have been met:

- ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment) for at least six (6) months in duration
- ◆ Any **ONE** of the following physical examination findings:
 - Limited range of motion
 - Evidence of joint swelling/effusion
 - Joint line tenderness
- ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
- ◆ Absence of Kellgren-Lawrence Grade II or greater findings on plain radiographs
- ◆ MRI or CT arthrogram is inconclusive for internal derangement/pathology

Diagnostic Arthroscopy Non-Indications

Not Medically Necessary

- Diagnostic Arthroscopy is considered **not medically necessary** for **ANY** other indication or condition.

Experimental Investigational, or Unproven (EIU)

- Based on lack of scientific evidence of efficacy and safety, the following is considered **experimental, investigational, or unproven (EIU)**:
 - ◆ “In-office” diagnostic arthroscopy (e.g., Mi-Eye™, VisionScope®)

Arthroscopic Debridement (Chondroplasty)/Loose Body/Foreign Body Removal

Arthroscopic Debridement (Chondroplasty) and Loose Body/Foreign Body Removal Indications

- Arthroscopic debridement (chondroplasty), loose body removal, and foreign body removal are considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Individual reports pain and any **ONE** of the following mechanical symptoms:
 - Knee range of motion is “blocked” due to pain
 - Giving way, subjective weakness, or buckling of the knee
 - Painful locking, clicking, catching, or popping during weight-bearing activities
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - **Note:** In the presence of painful locking, clicking, catching, or popping during weight-bearing activities secondary to an intra-articular loose body or foreign body, three (3) months of provider-directed non-surgical management is not required.
 - ◆ Concurrent findings on **EITHER** of the following:
 - MRI or CT arthrogram demonstrates articular cartilage degeneration and any **ONE** of the following:

- Loose body or foreign body within the joint
- Unstable flaps of articular cartilage
- Meniscal tear that extends to the articular surface (not simply degenerative changes, i.e., fraying) in conjunction with articular cartilage degeneration within the same compartment
- Impinging osteophytes, which would be reasonably expected to result in mechanical symptoms and loss of knee joint function
- Orthogonal radiographs demonstrate a loose body within the tibiofemoral or patellofemoral joint space

Arthroscopic Debridement (Chondroplasty) and Loose Body/Foreign Body Removal Non-Indications

- Arthroscopic debridement (chondroplasty) and loose body removal is considered **not medically necessary** in the presence of Kellgren-Lawrence Grade II or greater findings on plain radiographs except for loose body removal in the presence of an acutely locked knee on physical examination.
- Arthroscopic debridement (chondroplasty), loose body removal, and foreign body removal are considered **not medically necessary** for **ANY** other indication or condition.

Synovectomy

Synovectomy Indications

- Synovectomy (limited [e.g., plica or shelf resection], as a stand-alone procedure, or a major procedure with 2 or more compartments [e.g., medial and lateral]) is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Any **ONE** of the following physical examination findings:
 - Limited range of motion
 - Evidence of joint swelling/effusion
 - Joint line or plica tenderness
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - ◆ MRI or CT arthrogram demonstrates evidence of synovitis or plica
 - **Note:** Advanced imaging is not required for the clinical diagnosis of patellar clunk syndrome following knee replacement surgery
 - ◆ Absence of Kellgren-Lawrence Grade IV findings on plain radiographs
 - ◆ Presence of any **ONE** of the following:
 - Plica syndrome
 - Inflammatory arthritis (i.e., rheumatoid arthritis, gout, pseudogout, psoriatic arthritis)
 - Pigmented villonodular synovitis (PVNS)
 - Synovial chondromatosis
 - Lyme synovitis

- Hemophilia
- Hemochromatosis
- Non-specific synovitis (including proliferative synovitis, post-operative synovitis as a sequela from a knee replacement, patellar clunk syndrome, cyclops lesion, etc.)
- Recurrent hemarthrosis (i.e., secondary to sickle cell anemia, bleeding diathesis, etc.)

Synovectomy Non-Indications

- Synovectomy is considered **not medically necessary** for **ANY** other indication or condition.

Meniscectomy or Meniscal Repair

Meniscectomy or Meniscal Repair Indications

- Meniscectomy (partial or total) **or** meniscal repair is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ **TWO OR MORE** of the following physical examination findings:
 - Limited range of motion
 - Evidence of joint swelling/effusion
 - Joint line tenderness
 - Positive McMurray's Test, Thessaly Test, or Apley's Compression Test
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - **Note:** Meniscal tear with a locked knee on physical examination does not require three (3) months of provider-directed non-surgical management.
 - ◆ MRI or CT arthrogram demonstrates a meniscal tear that extends to the articular surface (not simply degenerative changes, i.e., fraying) and correlates with the individual's reported symptoms and physical examination findings
 - ◆ Absence of Kellgren-Lawrence Grade II or greater findings on plain radiographs
 - **Note:** Acute meniscal tear with associated function-limiting pain or locked knee on physical examination does not require absence of Kellgren-Lawrence Grade 2 or greater findings on plain radiographs
- Meniscectomy/saucerization for discoid lateral meniscus is considered **medically necessary** when MRI confirms the presence of a discoid meniscus and **ALL** of the above criteria are met (other than demonstration of a meniscal tear).
- Meniscal debridement is considered **medically necessary** when performed in conjunction with other medically necessary arthroscopic procedures on the knee (e.g., anterior cruciate reconstruction).

Meniscectomy or Meniscal Repair Non-Indications

- Meniscectomy (partial or total) or meniscal repair is considered **not medically necessary** for **ANY** other indication or condition.

Meniscal Allograft Transplantation

Meniscal Allograft Transplantation Indications

- Meniscal allograft transplantation is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands or employment)
 - ◆ Prior significant trauma resulting in an irreparable meniscal tear or has undergone a meniscectomy where at least 50% of the meniscus has been removed
 - ◆ Any **ONE** of the following physical examination findings:
 - Limited range of motion
 - Evidence of joint swelling/effusion
 - Joint line tenderness
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - ◆ Individual is not considered an appropriate candidate for total knee arthroplasty
 - ◆ Body Mass Index (BMI) 35 or less
 - ◆ Age 49 years or younger

Meniscal Allograft Transplantation Non-Indications

- Meniscal allograft transplantation is considered **not medically necessary** for **ANY** other indication or condition including, when **EITHER** of the following are present:
 - ◆ Upon standing radiographs, individual demonstrates osteoarthritic change in the knee including joint space narrowing and osteophytes which is classified by the Kellgren-Lawrence Scale as Grade III or IV
 - ◆ Upon MRI, individual demonstrates articular degeneration in affected compartment which is classified by the Modified Outerbridge Classification as Grade III or IV

Anterior Cruciate Ligament (ACL) Reconstruction

Anterior Cruciate Ligament (ACL) Reconstruction Indications

- Anterior cruciate ligament (ACL) reconstruction with allograft or autograft is considered **medically necessary** when **ALL** the following criteria have been met:
 - ◆ Function-limiting pain and/or a of knee function during the course of preoperative treatment which interferes with **ANY** of the following:
 - Ability to carry out age appropriate activities of daily living
 - Demands of employment
 - Need to return to activities that require cutting, pivoting, and/or agility in which ACL insufficiency may predispose to further instability episodes that may result in new articular or meniscal cartilage injuries
 - ◆ Individual reports knee instability which is noted as giving way, subjective weakness, or “buckling” during the course of preoperative treatment
 - ◆ Any **ONE** of the following physical examination findings:

- Positive Lachman's Test
- Positive Anterior Drawer Test
- Positive Pivot Shift Test
- ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration, except in an acute injury setting where joint instability has been documented and **ANY** of the following are present:
 - Need to return to activities that require cutting, pivoting, and/or agility activities in which ACL insufficiency may predispose to further instability episodes that may result in new articular or meniscal cartilage injuries
 - A confirmed ACL tear and a repairable meniscus tear
 - Concomitant ligament injuries (i.e., multi-ligamentous knee injury) that require reconstruction to provide stability
- ◆ MRI, CT arthrogram, or arthroscopy demonstrates a tear/disruption or significant laxity of the anterior cruciate ligament (ACL)

Anterior Cruciate Ligament (ACL) Reconstruction Non-Indications

- Anterior cruciate ligament (ACL) reconstruction is considered **not medically necessary** for **ANY** other indication or condition.

Anterior Cruciate Ligament (ACL) Repair

- Anterior cruciate ligament (ACL) repair is considered **experimental, investigational, or unproven (EIU)**.

Anterolateral Ligament (ALL) Reconstruction

Anterolateral Ligament (ALL) Reconstruction Indications

- Anterolateral ligament reconstruction is considered **medically necessary** when **ALL** of the following criteria have been met:
- Anterolateral ligament (ALL) reconstruction is required to augment the anterior cruciate ligament (ACL) reconstruction
 - ◆ Function-limiting pain and/or a documented loss of knee function during the course of preoperative treatment which interferes with **ANY** of the following:
 - Ability to carry out age appropriate activities of daily living
 - Demands of employment
 - Need to return to activities that require cutting, pivoting, and/or agility in which ACL insufficiency may predispose to further instability episodes that may result in new articular or meniscal cartilage injuries
 - ◆ Individual reports knee instability which is noted as giving way, subjective weakness, or "buckling" during the course of preoperative treatment
 - ◆ Any **ONE** of the following physical examination findings:
 - Positive Lachman's Test
 - Positive Anterior Drawer Test
 - Positive Pivot Shift Test
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration, (except in an acute injury setting where joint instability has been documented), and **ANY** of the following are present:

- Need to return to high-demand sports that require cutting, pivoting, and/or agility activities in which ACL insufficiency may predispose to further instability episodes that may result in new articular or meniscal cartilage injuries
- A confirmed ACL tear and a repairable meniscus tear
- Concomitant ligament injuries (i.e., multi-ligamentous knee injury) that require reconstruction to provide stability
- ◆ MRI, CT arthrogram, or arthroscopy demonstrates a tear/disruption or significant laxity of the anterior cruciate ligament (ACL)

Anterolateral Ligament (ALL) Reconstruction Non-Indications

- Anterolateral Ligament (ALL) reconstruction is considered **not medically necessary** for **ANY** other indication or condition.

Posterior Cruciate Ligament (PCL) Reconstruction

Posterior Cruciate Ligament (PCL) Reconstruction Indications

- Posterior cruciate ligament (PCL) reconstruction with allograft or autograft is considered **medically necessary** when **ALL** the following criteria have been met:
 - ◆ Function-limiting pain and a documented loss of knee function which interferes with the ability to carry out the age appropriate activities of daily living and/or demands of employment
 - ◆ Any **ONE** of the following physical examination/radiographic imaging findings:
 - Positive Posterior Drawer Sign
 - Positive Posterior Sag Sign or Tibial Drop Back Test
 - Positive Quadriceps Active Test
 - Eight (8) millimeters or more of increased posterior translation on stress radiographs
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration, except in an acute injury setting where hemarthrosis, effusion and joint instability have been documented and **EITHER** of the following are present:
 - Need to return to activities that require cutting, pivoting, and/or agility activities in which PCL insufficiency may predispose to further instability episodes that may result in new articular or meniscal cartilage injuries
 - Concomitant ligament injuries (i.e., multi-ligamentous knee) that require reconstruction to provide stability
 - ◆ MRI, CT Arthrogram, or arthroscopy demonstrates a tear/disruption or significant laxity of the posterior cruciate ligament (PCL)

Posterior Cruciate Ligament (PCL) Reconstruction Non-Indications

- Posterior cruciate ligament (PCL) reconstruction is considered **not medically necessary** for **ANY** other indication or condition.

Medial/Lateral Collateral Ligament (MCL/LCL) Repair/ Reconstruction

Medial/Lateral Collateral Ligament (MCL/LCL) Repair/Reconstruction Indications

- Medial/lateral collateral ligament (MCL/LCL) repair/reconstruction with allograft or autograft is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain and/or loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment
 - ◆ Individual reports knee instability which is noted as giving way, subjective weakness, or buckling
 - ◆ **EITHER** of the following physical examination findings:
 - Positive Valgus Stress Test (Medial)
 - Positive Varus Stress Test (Lateral)
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration, except in an acute injury setting of the lateral collateral ligament (LCL) (including the posterolateral corner) where total disruption of the ligament documented on MRI or CT arthrogram and effusion and joint instability have been documented on physical examination
 - ◆ MRI or CT arthrogram demonstrates a tear/disruption of the medial or lateral collateral ligament (MCL/LCL)

Medial/Lateral Collateral Ligament (MCL/LCL) Repair/Reconstruction Non-Indications

- Medial collateral ligament (MCL) repair/reconstruction is considered **not medically necessary** in an acute injury setting, including an isolated MCL repair.
- Medial/lateral collateral ligament (MCL/LCL) repair/reconstruction is considered **not medically necessary** for any other indication or condition.

Autologous Chondrocyte Implantation (ACI) or Autologous Chondrocyte Transplantation (ACT)

Autologous Chondrocyte Implantation (ACI) or Autologous Chondrocyte Transplantation (ACT) Indications

- Autologous chondrocyte implantation (ACI) or autologous chondrocyte transplantation (ACT) (using the MACI[®] implant) is considered **medically necessary** for the treatment of symptomatic single or multiple full-thickness cartilage defects of the distal femoral articular surface (i.e., medial condyle, lateral condyle, or trochlea) and/or patella caused by acute or repetitive trauma when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Presence of **BOTH** of the following findings:

- A stable knee with intact or reconstructed ligaments (ACL or PCL) and menisci.
 - A concurrent ligament stabilization or meniscal procedure at the time of ACI would be acceptable.
- Normal tibial-femoral and/or patella-femoral alignment
- ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
- ◆ A full-thickness distal femoral articular surface (i.e., medial condyle, lateral condyle or trochlea) and/or patellar chondral defect of 1-10cm² in size has been identified during an MRI or CT arthrogram, or during an arthroscopy and the Modified Outerbridge Classification or Outerbridge Classification is Grade III or Grade IV
- ◆ Absence of osteochondritis dissecans (OCD) lesion that requires bone grafting
- ◆ Absence of inflammatory arthritis or other systemic disease affecting the joints
- ◆ Absence of osteoarthritis or generalized tibial chondromalacia
- ◆ Minimal to absent osteoarthritic changes in the surrounding articular cartilage (e.g., Kellgren-Lawrence Grade II or less)
- ◆ Normal articular cartilage at the lesion border (contained lesion)
- ◆ For femoral and patellar chondral lesions, absence of a corresponding 'kissing lesion' with a Modified Outerbridge Classification of Grade III or IV of the distal femur (trochlea, condyles), patella, or tibia
- ◆ Body Mass Index (BMI) 35 or less
- ◆ Age 15-55 years

Autologous Chondrocyte Implantation (ACI) or Autologous Chondrocyte Transplantation (ACT) Non-Indications

Not Medically Necessary

- Autologous chondrocyte implantation is considered **not medically necessary** for any other indication or condition, including when **ANY** of the following are present:
 - ◆ Any knee joint surgery within six (6) months before screening excluding surgery to procure a biopsy or a concomitant procedure to prepare the knee for a MACI[®] implant
 - ◆ Presence of a corresponding 'kissing lesion' with a Modified Outerbridge Classification Grade III or IV defect(s) on the distal femur (trochlea, condyles), patella, or tibia
 - ◆ Presence of Kellgren-Lawrence Grade III or IV osteoarthritic changes in the surrounding articular cartilage
 - ◆ Total meniscectomy, meniscal allograft, or bucket-handle tear or displaced tear requiring > 50% removal of the meniscus in the target knee
 - ◆ Septic arthritis within one (1) year before screening
 - ◆ Known history of hypersensitivity to gentamicin, other aminoglycosides, or products of porcine or bovine origin
 - ◆ Uncorrected congenital blood coagulation disorders
 - ◆ Cruciate ligament instability

Experimental, Investigational, or Unproven (EIU)

- Autologous chondrocyte implantation using the MACI® implant for treatment of cartilaginous defects other than the distal femur or patella (i.e., proximal tibia) and of cartilaginous defects involving other joints (e.g., ankle, elbow, shoulder) is considered **experimental, investigational, or unproven (EIU)**.
 - ◆ For ACI performed for locations other than the knee, please reference the **Cigna Medical Coverage Policy: 0515 Miscellaneous Musculoskeletal Procedures**
- Hybrid autologous chondrocyte implantation performed with osteochondral autograft transfer system (Hybrid ACI/OATS) technique for the treatment of osteochondral defects is considered **experimental, investigational, or unproven (EIU)**.

Osteochondral Allograft/Autograft Transplantation Systems (OATS)/Mosaicplasty

Osteochondral Allograft/Autograft Transplantation Systems (OATS)/ Mosaicplasty Indications

- Osteochondral allograft/autograft transplantation (OATS)/mosaicplasty is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Presence of **BOTH** of the following findings:
 - A stable knee with intact or reconstructed ligaments (ACL or PCL) and menisci.
 - A concurrent ligament stabilization or meniscal procedure at the time of OATS would be acceptable.
 - Normal tibial-femoral and/or patella-femoral alignment
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - ◆ A full-thickness distal femoral articular surface (i.e., medial condyle, lateral condyle or trochlea) and/or patellar chondral defect that has been identified during an MRI or CT arthrogram, or during an arthroscopy and the Modified Outerbridge Classification or Outerbridge Classification is Grade III or Grade IV
 - ◆ **EITHER** of the following:
 - Osteochondral autograft transplants and mosaicplasty:
 - Small (i.e., ≤ 2.5 cm² total) chondral defects with sharp, definite borders surrounded by normal-appearing hyaline cartilage
 - Osteochondral allograft transplants:
 - Larger (i.e., ≤ 10.0 cm² total) chondral defects with sharp definite borders surrounded by normal appearing hyaline cartilage
 - ◆ Absence of inflammatory arthritis or other systemic disease affecting the joints
 - ◆ Minimal to absent osteoarthritic changes in the surrounding articular cartilage (e.g., Kellgren-Lawrence Grade 2 or less)
 - ◆ Normal articular cartilage at the lesion border (contained lesion)

- ◆ Absence of osteoarthritis or generalized tibial chondromalacia, steroid-induced cartilage or bone disease
- ◆ For femoral and patellar chondral lesions, absence of a corresponding ‘kissing lesion’ with a Modified Outerbridge Classification of Grade III or IV of the distal femur (trochlea, condyles), patella, or tibia
- ◆ Individual is not a candidate for total knee arthroplasty
- ◆ Body Mass Index (BMI) of less than 35
- ◆ Age 49 years or younger

Osteochondral Allograft/Autograft Transplantation Systems (OATS)/ Mosaicplasty Non-Indications

Not Medically Necessary

- Osteochondral allograft/autograft transplantation (OATS)/mosaicplasty of the distal femoral articular or patellar surface is considered **not medically necessary** for **ANY** other indication or condition.

Experimental, Investigational, or Unproven (EIU)

- Osteochondral allograft/autograft transplantation (OATS)/mosaicplasty is considered **experimental, investigational, or unproven (EIU)** for treatment of articular cartilage defects in locations other than the distal femur or patella (i.e., proximal tibia).
 - ◆ For osteochondral grafting performed for locations other than the knee (e.g., ankle, shoulder, elbow), please reference the **Cigna Medical Coverage Policy: 0515 Miscellaneous Musculoskeletal Procedures**
- Hybrid autologous chondrocyte implantation performed with osteochondral autograft transfer system (Hybrid ACI/OATS) technique for the treatment of osteochondral defects is considered **experimental, investigational, or unproven (EIU)**.

Abrasion Arthroplasty/Subchondral Drilling/Microfracturing

Abrasion Arthroplasty/Subchondral Drilling/Microfracturing Indications

- Abrasion arthroplasty, subchondral drilling or microfracturing is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Presence of **BOTH** of the following findings:
 - A stable knee with intact or reconstructed ligaments (ACL or PCL) and menisci.
 - A concurrent ligament stabilization or meniscal procedure at the time of abrasion arthroplasty would be acceptable.
 - Normal tibial-femoral and/or patella-femoral alignment
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration

- ◆ A full-thickness distal femoral articular surface (i.e., medial condyle, lateral condyle or trochlea) and/or patellar chondral defect of $\leq 2.5 \text{ cm}^2$ in size on the weight-bearing surface that has been identified during an MRI or CT arthrogram, or during an arthroscopy and the Modified Outerbridge Classification or Outerbridge Classification is Grade III or IV

Abrasion Arthroplasty/Subchondral Drilling/Microfracturing Non-Indications

- Abrasion arthroplasty, subchondral drilling, or microfracturing is considered **not medically necessary** for **ANY** other indication or condition.

Procedures for Patellofemoral Conditions

- Procedures for anterior knee pain with or without recurrent patellar instability can include both bony and/or soft tissue surgical procedures.
 - ◆ Bony surgical procedures can include tibial tubercle osteotomy/tubercleplasty (e.g., Fulkerson, Maquet) and trochleoplasty.
 - ◆ Soft tissue surgical procedures can include medial patellofemoral ligament (MPFL) reconstruction/repair, extensor realignment and/or muscle advancement or release (e.g. Cambell, Goldthwaite type procedure) and lateral retinacular release.

Procedures for Patellofemoral Conditions Indications

Medial Patellofemoral Ligament (MPFL) Reconstruction/Repair

- Medial patellofemoral ligament (MPFL) reconstruction/repair for anterior knee pain with or without recurrent patellar instability are considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Any **ONE** of the following:
 - Function-limiting anterior knee pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - Recurrent patellar instability interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - ◆ Any **ONE** of the following physical examination findings:
 - Positive J sign
 - Positive moving patellar apprehension test
 - Lateral patellar translation $> \frac{1}{2}$ (one-half) of the patellar width
 - Tenderness of the medial or lateral facets
 - Patellar grind test (Clarke's sign)
- Radiographic or other findings must include a tear of the MPFL identified by **ANY** of the following:
- visualized on MRI, CT, or Ultrasound (US)
 - identified by arthroscopy
 - on physical examination with **EITHER** of the following findings:

- MPFL palpation test findings with the knee in full extension and the patella medially subluxated noting tenderness to palpation of the origin of the MPFL
- Patella glide test findings > 75% lateral subluxation of the patella width at 30 degrees of knee flexion

Trochleoplasty

Trochleoplasty for anterior knee pain with or without recurrent patellar instability is considered **medically necessary** when **ALL** of the following criteria have been met:

- Any **ONE** of the following:
 - ◆ Function-limiting anterior knee pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Recurrent patellar instability interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment
- Failure of provider-directed non-surgical management for at least three (3) months in duration
- Any **ONE** of the following physical examination findings:
 - ◆ Positive J sign
 - ◆ Positive moving patellar apprehension test
 - ◆ Lateral patellar translation > ½ (one-half) of the patellar width
 - ◆ Tenderness of the medial or lateral facets
 - ◆ Patellar grind test (Clarke's sign)
- Include **ANY** of the following radiographic findings:
 - ◆ Supratrochlear spur
 - ◆ Lateral trochlear inclination (LTI) > 11 degrees
 - ◆ Crossing sign
 - ◆ Double-contour sign

Procedures Other Than Medial Patellofemoral Ligament (MPFL) Reconstruction Or Trochleoplasty

Procedures other than medial patellofemoral ligament (mpfl) reconstruction or trochleoplasty performed for anterior knee pain with or without recurrent patellar instability is considered **medically necessary** when **ALL** of the following criteria have been met:

- Any **ONE** of the following:
 - ◆ Function-limiting anterior knee pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Recurrent patellar instability interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment
- Failure of provider-directed non-surgical management for at least three (3) months in duration
- Any **ONE** of the following physical examination findings:
 - ◆ Positive J sign
 - ◆ Positive moving patellar apprehension test

- ◆ Lateral patellar translation > ½ (one-half) of the patellar width
- ◆ Tenderness of the medial or lateral facets
- ◆ Patellar grind test (Clarke's sign)
- Include **ANY** of the following:
 - ◆ Radiographic evidence of patellar tilt > 20 degrees
 - ◆ Patella alta (e.g., Insall-Salvati, Blackburne-Peel, Caton-Deschamps ratios)
 - ◆ Sulcus angle > 145 degrees
 - ◆ Increased tibial tubercle-posterior cruciate distance of > 24 mm
 - ◆ Increased TT-TG (tibial tubercle-trochlear groove) distance of > 20 mm
 - ◆ Concordant osteochondral defect of the patellofemoral joint (e.g., MRI, CT scan, or previous arthroscopic procedure)
 - ◆ Acute patellar dislocation with associated intra-articular fracture

Procedures for Patellofemoral Conditions Non-Indications

- Procedures for patellofemoral conditions are considered **not medically necessary** for any other indication or condition.

High Tibial Osteotomy

High Tibial Osteotomy Indications

- High tibial osteotomy is considered **medically necessary** when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ **ALL** of the following physical examination findings:
 - Less than 15 degrees of fixed varus deformity
 - The individual must be capable of at least 90 degrees of flexion
 - Joint stability in full extension
 - Intact anterior cruciate ligament (ACL)
 - ◆ Failure of provider-directed non-surgical management for at least three (3) months in duration
 - ◆ Unicompartamental osteoarthritis of the knee
 - ◆ Age 60 years or less
 - ◆ Individual is not a candidate for a knee arthroplasty

High Tibial Osteotomy Non-Indications

- High tibial osteotomy is considered **not medically necessary** for **ANY** other indication or condition, including when **ANY** of the following criteria is present:
 - ◆ Inflammatory arthropathy (i.e., rheumatoid arthritis)
 - ◆ Chondrocalcinosis
 - ◆ Anterior cruciate ligament (ACL) tear
 - ◆ Degenerative change affecting more than 1/3 of the femoral condylar surface
 - ◆ Osteochondral defect more than five (5) mm in depth

Lysis of Adhesions

Lysis of Adhesions Indications

- Lysis of adhesions is considered **medically necessary** for arthrofibrosis when **ALL** of the following criteria have been met:
 - ◆ Function-limiting pain (e.g., loss of knee function which interferes with the ability to carry out age appropriate activities of daily living and/or demands of employment)
 - ◆ Individual demonstrates less than 90 degrees of knee flexion by two (2) months after surgery, including knee replacement or trauma
 - ◆ Failure of provider-directed non-surgical management for at least two (2) months in duration, including **BOTH** of the following:
 - Anti-inflammatory medication and/or cortisone injection (unless contraindicated)
 - Physical therapy (i.e., active exercise and manual therapy designed to increase joint mobility and range of motion)

Lysis of Adhesions Non-Indications

- Lysis of adhesions is considered **not medically necessary** for **ANY** other indication or condition.

Procedures Not Addressed Elsewhere

- Based on lack of scientific evidence of efficacy and safety, the following are considered **experimental, investigational, or unproven (EIU)**:
 - ◆ Knee subchondroplasty
 - ◆ Focal resurfacing of a single knee joint defect (e.g., Arthrosurface Knee HemiCAP®, UniCAP®)

Procedure (CPT®) Codes

This guideline relates to the CPT® code set below. Codes are displayed for informational purposes only. Any given code's inclusion on this list does not necessarily indicate prior authorization is required.

CPT®	Code Description/Definition
27331	Arthrotomy, knee; including joint exploration, biopsy, or removal of loose or foreign bodies
27332	Arthrotomy, with excision of semilunar cartilage (meniscectomy) knee; medial OR lateral
27333	Arthrotomy, with excision of semilunar cartilage (meniscectomy) knee; medial AND lateral
27334	Arthrotomy, with synovectomy, knee; anterior OR posterior
27335	Arthrotomy, with synovectomy, knee; anterior AND posterior including popliteal area
27340	Excision, prepatellar bursa
27347	Excision of lesion of meniscus or capsule (eg, cyst, ganglion), knee
27355	Excision or curettage of bone cyst or benign tumor of femur;
27356	Excision or curettage of bone cyst or benign tumor of femur; with allograft
27357	Excision or curettage of bone cyst or benign tumor of femur; with autograft (includes obtaining graft)
27358	Excision or curettage of bone cyst or benign tumor of femur; with internal fixation (List in addition to code for primary procedure)

This guideline relates to the CPT® code set below. Codes are displayed for informational purposes only. Any given code's inclusion on this list does not necessarily indicate prior authorization is required.

CPT®	Code Description/Definition
27360	Partial excision (craterization, saucerization, or diaphysectomy) bone, femur, proximal tibia and/or fibula (eg, osteomyelitis or bone abscess)
27403	Arthrotomy with meniscus repair, knee
27405	Repair, primary, torn ligament and/or capsule, knee; collateral
27407	Repair, primary, torn ligament and/or capsule, knee; cruciate
27409	Repair, primary, torn ligament and or capsule, knee; collateral and cruciate ligaments
27412	Autologous chondrocyte implantation, knee
27415	Osteochondral allograft, knee, open
27416	Osteochondral autograft(s), knee, open (eg, mosaicplasty) (includes harvesting of autograph[s])
27418	Anterior tibial tubercleplasty (eg, Maquet type procedure)
27420	Reconstruction of dislocating patella; (eg, Hauser type procedure)
27422	Reconstruction of dislocating patella; with extensor realignment and/or muscle advancement or release (eg, Campbell, Goldwaite type procedure)
27424	Reconstruction of dislocating patella; with patellectomy
27425	Lateral retinacular release, open
27427	Ligamentous reconstruction (augmentation), knee; extra-articular
27428	Ligamentous reconstruction (augmentation), knee; intra-articular (open)
27429	Ligamentous reconstruction (augmentation), knee; intra-articular (open) and extra-articular
27438	Arthroplasty, patella; with prosthesis
27440	Arthroplasty, knee, tibial plateau;
27442	Arthroplasty, femoral condyles or tibial plateau(s), knee;
29850	Arthroscopically aided treatment of intercondylar spine(s) and/or tuberosity fracture(s) of the knee, with or without manipulation; without internal or external fixation (includes arthroscopy)
29851	Arthroscopically aided treatment of intercondylar spine(s) and/or tuberosity fracture(s) of the knee, with or without manipulation; with internal or external fixation (includes arthroscopy)
29855	Arthroscopically aided treatment of tibial fracture, proximal (plateau); unicondylar, includes internal fixation, when performed (includes arthroscopy)
29856	Arthroscopically aided treatment of tibial fracture, proximal (plateau); bicondylar, includes internal fixation, when performed (includes arthroscopy)
29866	Arthroscopy, knee, surgical; osteochondral autograft(s) (eg, mosaicplasty) (includes harvesting of the autograft[s])
29867	Arthroscopy, knee, surgical; osteochondral allograft (eg, mosaicplasty)
29868	Arthroscopy, knee, surgical; meniscal transplantation (includes arthrotomy for meniscal insertion), medial or lateral
29870	Arthroscopy, knee, diagnostic; with or without synovial biopsy (separate procedure)
29871	Arthroscopy, knee, surgical; for infection, lavage and drainage
29873	Arthroscopy, knee, surgical; with lateral release
29874	Arthroscopy, knee, surgical; for removal of loose body or foreign body (eg, osteochondritis dissecans fragmentation, chondral fragmentation)
29875	Arthroscopy, knee, surgical; synovectomy, limited (eg, plica or shelf resection) (separate procedure)
29876	Arthroscopy, knee, surgical; synovectomy, major, two or more compartments (eg, medial or lateral)
29877	Arthroscopy, knee, surgical; debridement/shaving of articular cartilage (chondroplasty)
29879	Arthroscopy, knee, surgical; abrasion arthroplasty (includes chondroplasty where necessary) or multiple drilling or microfracture
29880	Arthroscopy, knee, surgical; with meniscectomy (medial AND lateral, including any meniscal shaving) including debridement/shaving of articular cartilage (chondroplasty), same or separate compartment(s), when performed

This guideline relates to the CPT® code set below. Codes are displayed for informational purposes only. Any given code's inclusion on this list does not necessarily indicate prior authorization is required.

CPT®	Code Description/Definition
29881	Arthroscopy, knee, surgical; with meniscectomy (medial OR lateral, including any meniscal shaving) including debridement/shaving of articular cartilage (chondroplasty), same or separate compartment(s), when performed
29882	Arthroscopy, knee, surgical; with meniscus repair (medial OR lateral)
29883	Arthroscopy, knee, surgical; with meniscus repair (medial AND lateral)
29884	Arthroscopy, knee, surgical; with lysis of adhesions, with or without manipulation (separate procedure)
29885	Arthroscopy, knee, surgical; drilling for osteochondritis dissecans with bone grafting, with or without internal fixation (including debridement of base of lesion)
29886	Arthroscopy, knee, surgical; drilling for intact osteochondritis dissecans lesion
29887	Arthroscopy, knee, surgical; drilling for intact osteochondritis dissecans lesion with internal fixation
29888	Arthroscopically aided anterior cruciate ligament repair/augmentation or reconstruction
29889	Arthroscopically aided posterior cruciate ligament repair/augmentation or reconstruction

This list may not be all inclusive and is not intended to be used for coding/billing purposes. The final determination of reimbursement for services is the decision of the health plan and is based on the individual's policy or benefit entitlement structure as well as claims processing rules.

Procedure (HCPCS) Codes

This guideline relates to the HCPCS code set below. Codes are displayed for informational purposes only. Any given code's inclusion on this list does not necessarily indicate prior authorization is required.

J7330	Autologous cultured chondrocytes, implant
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This list may not be all inclusive and is not intended to be used for coding/billing purposes. The final determination of reimbursement for services is the decision of the health plan and is based on the individual's policy or benefit entitlement structure as well as claims processing rules.

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