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Appendix:

Using the defined inclusion and exclusion criteria, a detailed search strategy was followed for each joint and location.

Search 1: (osteocondral [All Fields] or osteoarticular [All Fields]) AND (“allografts”[MeSH Terms] OR ‘allografts’[All Fields] OR ‘allograft’[All Fields])) AND (“knee”[MeSH Terms] OR “knee”[All Fields] OR “knee joint”[MeSH Terms] OR (“knee”[All Fields] AND “joint”[All Fields]) OR “knee joint”[All Fields])

Search 2: (osteocondral [All Fields] or osteoarticular [All Fields]) AND (“allografts”[MeSH Terms] OR ‘allografts’[All Fields] OR ‘allograft’[All Fields]) AND “patella”[All Fields]

Search 3: (osteocondral [All Fields] or osteoarticular [All Fields]) AND (“allografts”[MeSH Terms] OR ‘allografts’[All Fields] OR ‘allograft’[All Fields]) AND (“ankle”[All Fields] OR “talus”[All fields] OR “talar”[All Fields])

Search 4: (osteocondral [All Fields] or osteoarticular [All Fields]) AND (“allografts”[MeSH Terms] OR ‘allografts’[All Fields] OR ‘allograft’[All Fields]) AND (‘humeral’[All Terms] OR ‘glenoid’[All Terms] OR ‘shoulder’[All Terms] ‘glenohumeral’[All Terms] AND “Joint”[All Terms])

Search 5: (‘osteocondral’ [All Fields] or ‘osteoarticular’ [All Fields]) AND (“allografts”[MeSH Terms] OR ‘allografts’[All Fields] OR ‘allograft’[All Fields]) AND (‘femoral head’[All Terms] OR ‘acetabulum’[All Terms] OR ‘hip’[All Terms] AND “Joint”[All Terms])

Inclusion/exclusion criteria

The results were filtered to English and full-text articles. References were reviewed to identify additional articles and confirm completeness. Inclusion criteria for clinical studies of each joint

location included original research studies that investigated the use of fresh osteochondral allograft transplantation in humans to treat chondral or osteochondral defects. The inclusion criteria for clinical studies required at least one patient, reported patient outcomes, and minimum follow-up of 12 months. Systematic reviews of clinical studies investigation outcomes of OCA, consensus statements, and health economic studies were included as relevant to this study. Exclusion criteria for all joint locations were as follows: articles not written in English; review or expert opinion articles; instructional courses; surgical technique articles; conference proceedings and presentations; OCAs for tumor; basic science or animal studies, articles that used OCAs that were cryopreserved, frozen, or decellularized; One author performed the literature search (S.M.T) and 2 authors (D.C., S.M.T) independently reviewed the search results. Titles and abstracts were reviewed for all search results. Full-text articles were obtained to determine if the study met the established inclusion and exclusion criteria. In total, 103 articles were identified that reported clinical outcomes of OCA in the knee joint, 32 articles reported outcomes of OCA in the ankle joint, 10 articles report outcomes of OCA in the shoulder joint, and 11 articles report outcomes of OCA in the hip joint.

Appendix Table 1: Statements generated from private payer insurance policies

Statements based on private payer medical criteria	
Defect characteristics	
1	There should be a maximum and minimum defect size for OCA transplantation
2	Defect should be focal, isolated, unipolar
3	Defect should be full thickness and grade III or IV
4	Defect should be located on weight bearing surface of medial or lateral femoral condyles or trochlear region (trochlear groove of femur)
5	Defect can be located on patella
6	The patella region is non-load bearing
7	Weight bearing and non-weight bearing should not be criteria
8	OCA is only for femoral articulation
9	Specific location of the knee should not be criteria
10	The lesion should only involve one side of joint and opposing articular cartilage surface should be generally free of disease or injury including no arthritis on corresponding tibial surface
11	Lesion must/should be “contained”
12	Lesions must/should be adjacent to near normal surrounding articular cartilage with minimal to absent degenerative changes (e.g. Kellgren-Lawrence Grade 2 or less)
13	Lesions must/should be opposed by near normal articulating cartilage (e.g. grade 0, 1, 2)
14	Joint should have absence of a corresponding ‘kissing lesion’ with a Modified Outerbridge Scale of Grade III or IV of the distal femur (trochlea, condyles), patella, or tibia
Joint and health stability	
15	Patient should have normal knee alignment or surgically corrected at time of allograft or prior to grafting
16	The knee should have meniscus that is intact or has stable partial tears
17	Individuals who have had a previous total meniscectomy is a contraindication to OCA transplantation
18	The knee should be stable with intact or reconstructed ligaments
19	Knee should have normal tibial-femoral and/or patella/femoral alignment
20	The joint space should be normal without evidence of inflammation or degenerative changes
21	Presence inflammation or osteoarthritis of the joint is a contraindication to OCA transplantation
22	Joint space should be normal or near-normal with no > 15% joint space narrowing
23	Individuals with a cartilaginous defect associated with osteoarthritis should be a contraindication
24	Individuals with inflammatory joint disease should be a contraindication to OCA

25	Individuals where osteoarthritic or inflammatory process significantly affects the quality of perilesional cartilage should be a contraindication
26	Presence of active infection is a contraindication
27	Presence of synovial disease is a contraindication
28	Patient history of malignancy is a contraindication
29	Presence of a current unresected tumor is a contraindication
Patient selection	
30	The defect should be identified during an MRI or CT arthrogram, or during arthroscopy
31	OCA can treat avascular necrosis lesions of the femoral condyle
32	The chondral defect of the knee should be caused by acute or repetitive trauma
33	OCA transplantation can be used in skeletally immature patients
34	Patients should show skeletal maturity
35	There should be criteria for maximum and minimum age limit.
36	Adult patients should be too young to be considered an appropriate candidate for total or partial knee arthroplasty or other reconstructive knee surgery (e.g. less than or equal to 55 years)
37	There should be criteria for maximum BMI
38	BMI greater than 35kg/m ² is considered severe obesity and a contraindication to OCA
39	Patient selection should include no history of cancer in the bones, cartilage, fat, or muscle of affected limb
40	Patient selection should include absence of localized or systemic infection
41	Patient must be willing and capable to participate in an extensive period of non-weight bearing and supervised post-operative physical rehabilitation program
Patient history	
42	The defect should cause disabling localized knee pain that is unresponsive to conservative treatment
43	Patient should have function-limiting pain (e.g. loss of knee function which interferes with ability to carry out age appropriate activities of daily living and/or demands of employment)
44	Patient should have presence of disabling, localized knee pain of greater than or equal to 6 months duration that is refractory to conservative treatment, as well as, arthroplasty and/or microfracture techniques
45	Patient should have debilitating symptoms that significantly limit ambulation
46	Patient should have at least 6 months of knee pain and failed conservative treatment measures
47	Patient should have failed a provider-directed non-surgical management for at least three (3) months in duration
48	Patient should have disabling symptoms such as locking, swelling, or knee pain that are unresponsive to conservative treatment for a minimum of 2 months
49	Patients should have failed arthroscopic procedure or are not candidates for such procedure because of size, shape, or location of the lesion

50	Patient should have unsatisfactory outcome with previous arthroscopic or other traditional surgical procedure (i.e. microfracture drilling, abrasion, osteochondral graft)
51	If debridement is only prior surgical treatment consideration should be given to marrow-stimulation techniques before osteochondral grafting
52	OCA transplantation should be indicated when other cartilage repair techniques (e.g. microfracture, osteochondral autografting or autologous chondrocyte implantation) would be inadequate due to lesion size, location, or depth
Other locations/joints for OCAs	
53	OCA of all other joints besides the knee is investigational due and does not have scientific evidence of effectiveness improving health outcomes
54	OCA transplantation of the patella unproven/investigational
55	OCA transplantation of the talus unproven/investigational
56	OCA transplantation of the shoulder unproven/investigational
57	OCA transplantation of the elbow unproven/investigational

Appendix Table 2: Level of evidence for statements that reach consensus

MOCA Likert survey results				
Statements	Level of Evidence	% Agree	% Neither agree or disagree	% Disagree
Defect should be full thickness and grade III or IV ^{1,7-100}	II, III, IV	96.7	0.0	3.3
Defect can be located on patella ^{5,19,25,34,38,42,45,60,63,67,83,95,101}	IV	96.7	3.3	0.0
Weight bearing and non-weight bearing should not be criteria ⁸	IV	83.3	10.0	6.7
Specific location of the knee should not be criteria ^{1,3-100}	II, III, IV	80.0	3.3	16.7
The defect should be identified during an MRI or CT arthrogram, or during arthroscopy	NA	93.3	3.3	3.3
Lesions must/should be adjacent to near normal surrounding articular cartilage with minimal to absent degenerative changes (e.g. Kellgren-Lawrence Grade 2 or less) ^{18,30,58,59,95,96}	IV	80.0	13.3	6.7
Patient should have normal knee alignment or surgically corrected at time of allograft or prior to grafting ^{17,25,38,40,41,44,50,52,57-59,80-82,100}	IV	96.7	3.3	0.0
The knee should be stable with intact or reconstructed ligaments ^{13,80,82,87,94}	III, IV	96.7	3.3	0.0
Individuals where osteoarthritic or inflammatory process significantly affects the quality of perilesion cartilage should be a contraindication ^{20,30,58,59,95,96}	IV	80.0	20.0	0.0
Presence of active infection is a contraindication	NA	96.7	0.0	3.3
OCA transplantation can be used in skeletally immature patients ^{64,92}	IV	93.3	3.3	3.3
Patient selection should include absence of localized or systemic infection	NA	100.0	0.0	0.0
Patient must be willing and capable to participate in an extensive period of non-weight bearing and supervised post-operative physical rehabilitation program ⁷³	III	80.0	3.3	16.7
Patient should have function-limiting pain (e.g. loss of knee function which interferes with ability to carry out age appropriate activities of daily living and/or demands of employment)	NA	80.0	10.0	10.0

OCA can treat avascular necrosis lesions of the femoral condyle ^{8,21,59,32,41,86}	II, IV	96.7	0.0	3.3
OCA transplantation should be indicated when other cartilage repair techniques (e.g. microfracture, osteochondral autografting or autologous chondrocyte implantation) would be inadequate due to lesion size, location, or depth ¹⁻¹⁰⁰	II, III, IV	80.0	6.7	13.3
The patella region is non-load bearing	NA	3.3	0.0	96.7
Osteochondral allograft (OCA) is only for femoral articulation ^{5,19,25,34,38,42,45,60,63,67,83,95,101}	IV	6.7	10.0	83.3
Patient history of malignancy is a contraindication	NA	10.0	10.0	80.0
Patients should show skeletal maturity ^{64,92}	IV	16.7	3.3	80.0
There should be criteria for maximum and minimum age limit ^{1,3,14,23,27,51,64,68,92,95}	III, IV	6.7	16.7	76.7
Patient should have presence of disabling, localized knee pain of greater than or equal to 6 months duration that is refractory to conservative treatment, as well as, arthroplasty and/or microfracture techniques ^{33,35,48,58,72,98}	III, IV	10.0	6.7	83.3
Patient should have unsatisfactory outcome with previous arthroscopic or other traditional surgical procedure (i.e. microfracture drilling, abrasion, osteochondral graft) ^{33,35,48,58,72,98}	III, IV	13.3	6.7	80.0
If debridement is only prior surgical treatment consideration should be given to marrow-stimulation techniques before osteochondral grafting ^{33,35,48,58,72,98}	III, IV	3.3	10.0	86.7
OCA of all other joints besides the knee is investigational and does not have scientific evidence of effectiveness improving health outcomes ^{5,19,25,34,38,42,45,60,95,101-152}	II, III, IV	10.0	3.3	86.7
OCA transplantation of patella is unproven/investigational ^{15,25,34,38,42,45,60,63,67,83,95,101}	IV	3.3	0.0	96.7
OCA transplantation of the talus is unproven/investigational ^{102,103,105-107,109-116,118-120,123,125,126,128,134,136,137,141-145,147, 149, 153}	II, III, IV	10.0	0.0	90.0

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