Hyaluronic Acid Derivatives:

Durolane®, Euflexxa®, Gel-One®, GelSyn-3™, GenVisc 850®, Hyalgan®, Hymovis®, Monovisc®, Orthovisc®, Supartz/Supartz FX™, SynoJoynt™, Synvisc®, Synvisc-One®, Triluron™, TriVisc™, VISCO-3™ (Intra-articular)

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I. Length of Authorization

- Initial: Prior authorization validity will be provided initially for 6 months.
- Renewal: Prior authorization validity may be renewed every 12 months thereafter.

II. Dosing Limits

Max Units (per dose and over time) [HCPCS Unit]:*

Drug	HCPCS	1 Billable Unit (BU)	BU per Admin	No. Admins (per knee per 180 days)	Max Units (per 180 days)*
Durolane	J7318	1 mg	60	1	120
Euflexxa	J7323	1 dose	1	3	6
Gel-One	J7326	1 dose	1	1	2
GelSyn-3	J7328	0.1 mg	168	3	1008
GenVisc 850	J7320	1 mg	25	5	250
Hyalgan; Supartz; Supartz FX	J7321	1 dose	1	5	10
Hymovis	J7322	1 mg	24	2	96
Monovisc	J7327	1 dose	1	1	2

Orthovisc	J7324	1 dose	1	4	8
SynoJoynt	J7331	1 mg	20	3	120
Synvisc	J7325	1 mg	16	3	96
Synvisc-One	J7325	1 mg	48	1	96
Triluron	J7332	1 mg	20	3	120
TriVisc	J7329	1 mg	25	3	150
VISCO-3	J7321	1 dose	1	3	10

^{*}Max units are based on administration to both knees

III. Initial Approval Criteria

Coverage is provided in the following conditions:

· Patient must try and have an inadequate response, contraindication, or intolerance to Euflexxa

Universal Criteria 1-15,23-25

- Patient does not have any conditions which would preclude intra-articular injections (e.g., active
 joint infection, unstable joint, bleeding disorders, etc.); AND
- Patient has not received therapy with intra-articular long-acting corticosteroid type drugs (i.e. Zilretta, etc.) within the previous 6 months of therapy; **AND**

Osteoarthritis of the knee † 1-15,23-25,27-29

- Patient has a radiographically* confirmed diagnosis of osteoarthritis of the knee; AND
- Trial and failure of conservative therapy (including physical therapy AND pharmacotherapy [e.g., non-steroidal anti-inflammatory drugs (NSAIDs), acetaminophen (up to 1 g 4 times/day) and/or topical capsaicin cream]) has been attempted and has not resulted in functional improvement after at least 3 months; AND
- The patient has failed to adequately respond to aspiration and injection of intra-articular steroids; **AND**
- The patient reports pain which interferes with functional activities (e.g., ambulation, prolonged standing)

IV. Renewal Criteria 1-15,23-25,27-29

Coverage can be renewed based upon the following criteria:

^{*}Note: Imaging is not required to make the diagnosis in patients with a typical presentation of OA²⁷

[†] FDA Approved Indication(s); ‡ Compendia Recommended Indication(s); ♠ Orphan Drug

- Patient must try and have an inadequate response, contraindication, or intolerance to Euflexxa
- Patient continues to meet the universal and other indication-specific relevant criteria identified in section III; AND
- Disease response with treatment as defined by improvement in signs and symptoms of pain and
 a stabilization or improvement in functional capacity during the 6-month period following the
 previous series of injections as evidenced by objective measures; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: severe joint swelling and pain, severe infections, anaphylactic or anaphylactoid reactions, etc.

V. Dosage/Administration (per knee per 180 days)

Drug	Dose
Durolane	60 mg intra-articularly x 1 administration
Euflexxa	20 mg intra-articularly once weekly x 3 administrations
Gel-One	30 mg intra-articularly x 1 administration
GelSyn-3	16.8 mg intra-articularly once weekly x 3 administrations
GenVisc 850	25 mg intra-articularly once weekly x 5 administrations
Hyalgan	20 mg intra-articularly once weekly x 5 administrations
Hymovis	24 mg intra-articularly once weekly x 2 administrations
Monovisc	88 mg intra-articularly x 1 administration
Orthovisc	30 mg intra-articularly once weekly x 4 administrations
SynoJoynt	20 mg intra-articularly once weekly x 3 administrations
Supartz/Supartz FX	25 mg intra-articularly once weekly x 5 administrations
Synvisc	16 mg intra-articularly once weekly x 3 administrations
Synvisc-One	48 mg intra-articularly x 1 administration
Triluron	20 mg intra-articularly once weekly x 3 administrations
TriVisc	25 mg intra-articularly once weekly x 3 administrations
VISCO-3	25 mg intra-articularly once weekly x 3 administrations

VI. Billing Code/Availability Information

HCPCS Code & NDC:

Drug	HCPCS Code	1 Billable Unit	Dose per Injection	Injections (per knee per 180 days)	NDC
Durolane	J7318	1 mg	60 mg/3 mL	1	89130-2020-xx
Euflexxa	J7323	1 dose	20 mg/2 mL	3	55566-4100-xx

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Gel-One	J7326	1 dose	30 mg/3 mL	1	50016-0957-xx
GelSyn-3	J7328	0.1 mg	16.8 mg/2 mL	3	89130-3111-xx
GenVisc 850	J7320	1 mg	25mg/2.5 ml	5	50653-0006-xx
Hyalgan	J7321	1 dose	20 mg/2 mL	5	89122-0724-xx
Hymovis	J7322	1 mg	24 mg/3 mL	2	89122-0496-xx
Monovisc	J7327	1 dose	88 mg/4 mL	1	59676-0820-xx
Orthovisc	J7324	1 dose	30 mg/2 mL	4	59676-0360-xx
Supartz	J7321	1 dose	25 mg/2.5 mL	5	89130-5555-xx
Supartz FX	J7321	1 dose	25 mg/2.5 mL	5	89130-4444-xx
SynoJoynt	J7331	1 mg	20 mg/2 mL	3	82197-0721-xx
Synvisc	J7325	1 mg	16 mg/2 mL	3	58468-0090-xx
Synvisc-One	J7325	1 mg	48 mg/6 mL	1	58468-0090-xx
Triluron	J7332	1 mg	20 mg/2 mL	3	89122-0879-xx
TriVisc	J7329	1 mg	25 mg/2.5 mL	3	50563-0006-xx
VISCO-3	J7321	1 dose	25mg/2.5 mL	3	50016-0957-xx

VII. References

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Appendix 1 – Covered Diagnosis Codes

ICD-10	ICD-10 Description
M17.0	Bilateral primary osteoarthritis of knee
M17.10	Unilateral primary osteoarthritis, unspecified knee
M17.11	Unilateral primary osteoarthritis, right knee
M17.12	Unilateral primary osteoarthritis, left knee
M17.2	Bilateral post-traumatic osteoarthritis of knee
M17.30	Unilateral post-traumatic osteoarthritis, unspecified knee
M17.31	Unilateral post-traumatic osteoarthritis, right knee
M17.32	Unilateral post-traumatic osteoarthritis, left knee
M17.4	Other bilateral secondary osteoarthritis of knee
M17.5	Other unilateral secondary osteoarthritis of knee
M17.9	Osteoarthritis of knee, unspecified

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

The preceding information is intended for non-Medicare coverage determinations. Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determinations (NCDs) and/or Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. Local Coverage Articles (LCAs) may also exist for claims payment purposes or to clarify benefit eligibility under Part B for drugs which may be self-administered. The following link may be used to search for NCD, LCD, or LCA documents: https://www.cms.gov/medicare-coverage-database/search.aspx. Additional indications, including any preceding information, may be applied at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCA/LCD):

	Medicare Par	t B Covered Diagnosis Codes
Jurisdiction	NCD/LCA/LCD Document (s)	Contractor
J, M	A59030	Palmetto GBA
6, K	A52420	National Government Services, Inc. (NGS)
5, 8	A56157	Wisconsin Physicians Service Insurance Corp (WPS)

Medicare Part B Administrative Contractor (MAC) Jurisdictions			
Jurisdiction	Applicable State/US Territory	Contractor	
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC	
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC	
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)	
6	MN, WI, IL	National Government Services, Inc. (NGS)	
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.	
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)	
N (9)	FL, PR, VI	First Coast Service Options, Inc.	
J (10)	TN, GA, AL	Palmetto GBA	
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA	
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.	
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)	
15	кү, он	CGS Administrators, LLC	