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JOHNS HOPKINS HEALTHCARE		

Johns Hopkins HealthCare LLC	Policy Number	MMDP001
Pharmacy Public Medical Management Drug Policies	Effective Date	01/01/2006
	Review Date	10/19/2022
Subject	Revision Date	10/19/2022
Hyaluronic Acid, Euflexxa, Hyalgan, Orthovisc, Supartz, Synvisc, Monovisc, Gel-one, Gel-Syn 3, Genvisc 850, Supartz FX, Visco 3, Hymovis, Durolane, TriVisc, Synojoynt, Triluron	Page	1 of 4

This document applies to the following Participating Organizations:

US Family Health Plan

<u>Keywords</u>: durolane, euflexxa, gel-one, gel-syn, genvisc 850, hyalgan, hymovis, monovisc, orthovisc, supartz, SynoJoynt, synvisc, Triluron, trivisc, visco 3

Table	e of Contents	Page Number
I.	POLICY	1
II.	POLICY CRITERIA	1
III.	AUTHORIZATION PERIOD/LIMITATIONS	1
IV.	EXCLUSIONS	2
V.	RECOMMENDED DOSAGE	2
VI.	CODES	2
VII.	<u>REFERENCES</u>	3
VIII.	APPROVALS	3

I. POLICY

A. Hyaluronic Acid, Euflexxa, Hyalgan, Orthovisc, Supartz, Synvisc, Monovisc, Gel-one, Gel-Syn 3, Genvisc 850, Supartz FX, Visco 3, Hymovis, Durolane, TriVisc, SynoJoynt and Triluron will require prior-authorization for medical benefit coverage to ensure appropriate use. The process for initiating a prior authorization can be found in policy PHARM 20.

II. POLICY CRITERIA

- A. Hyaluronic acid products may be approved if ALL the following have been met:
 - 1. Patient is 22 years of age or older (18 years of age or older if request is for Euflexxa, Hyalgan, Orthovisc, Supartz, Supartz FX, Gel-Syn 3, or Triluron)
 - 2. Patient has a documented diagnosis of osteoarthritis of the knee
 - 3. Documented trial and inadequate response to at least two non-pharmacological interventions (e.g. physical therapy exercises, a weight loss program (if patient is overweight), tai chi, or a group-based self-management program)
 - 4. Documented trial and inadequate response to at least a one-month trial each of the following:
 - 1. A formulary topical nonsteroidal antiinflammatory drugs (NSAID) [such as diclofenac 1% gel]
 - 2. A dose optimized regimen of a formulary oral NSAID, or acetaminophen (if the use of NSAIDs is contraindicated)
 - 5. Documented trial and inadequate response or intolerance to intra-articular glucocorticoid injection

III. AUTHORIZATION PERIOD/LIMITATIONS

- A. Initial authorization will be limited to three to five weeks depending on the hyaluronic acid product
- B. Authorization for re-treatment may be granted under the following criteria:
 - 1. A minimum of 26 weeks has elapsed since the initial treatment
 - 2. Documentation has been provided showing a beneficial response to initial treatment, evidenced by both of the following:
 - a. Improvement in pain and joint movement
 - b. Reduction in the dosing of NSAIDs, or other analgesics

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IV. EXCLUSIONS

- A. Hyaluronic Acid products will not be approved for the following (not an all-inclusive list):
 - 1. Pediatric patients
 - 2. Patients who are pregnant, or nursing
 - 3. Patients with documented skin diseases, or infection, at the injection site
 - 4. Concurrent use with another hyaluronic acid product
 - 5. Any other indications or uses that are not FDA-approved, or guideline-supported
- B. The use of physician samples, or manufacturer product discounts, does not guarantee coverage under the provisions of the medical and/or pharmacy benefit. All pertinent criteria must be met in order to be eligible for benefit coverage.

V. RECOMMENDED DOSAGE

Please refer to the FDA-approved prescribing information for product-specific dosing details.

VI. CODES

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Note: The following CPT/HCPCS codes are included below for informational purposes. Inclusion or exclusion of a CPT/HCPCS code(s) below does not signify or imply member coverage or provider reimbursement. The member's specific benefit plan determines coverage.

Medication	HCPCS/CPT Code
Hyaluronan or derivative, Euflexxa, for intra- articular injection, per dose (20 mg/2 mL)	J7323
Hyaluronan or derivative, Hyalgan or Supartz or Visco-3, for intra-articular injection, per dose (Hyalgan dose is 20 mg/2 mL, Supartz and Visco-3 dose is 25 mg/2.5 mL)	J7321
Hyaluronan or derivative, Orthovisc, for intra- articular injection, per dose (30 mg/2 mL)	J7324
Hyaluronan or derivative, Synvisc or Synvisc- One, for intra-articular injection, 1 mg	J7325
Hyaluronan or derivative, Monovisc, for intra- articular injection, per dose	J7327
Hyaluronan or derivative, Gel-One, for intra- articular injection, per dose	J7326
Hyaluronan or derivative, GELSYN-3, for intra- articular injection, 0.1 mg	J7328
Hyaluronan or derivative, Genvisc 850, for intra-articular injection, 1 mg	J7320
Hyaluronan or derivative, Hymovis, for intra- articular injection, 1 mg	J7322

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Hyaluronan or derivative, Durolane, for intra- articular injection, 1 mg	J7318
Hyaluronan or derivative, Trivisc, for intra- articular injection, 1 mg	J7329
Hyaluronan or derivative, synojoynt, for intra- articular injection, 1 mg	J7331
Hyaluronan or derivative, triluron, for intra- articular injection, 1 mg	J7332

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VIII. APPROVALS

Signature on file at JHHC

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DATE OF REVISION	SUMMARY OF CHANGE
04/20/2016	Removed background information and added Monovisc/ Gel-One to criteria review, removed process of initiation of request
10/19/2016	Added new formulation Gel-Syn
07/25/2017	Policy section- language update to match operational processes
07/27/2017	Updated Exclusions section regarding physician samples
07/01/2018	Removed EHP Line of Business
12/19/2018	Updated layout, and clarified applicable drugs
01/16/2019	Added Durolane and TriVisc as applicable drugs
10/16/2019	Added SynoJoynt and Triluron as applicable drugs
01/15/2020	No policy changes- Presented policy for USFHP adoption effective 3/1/2020
07/15/2020	Updated policy criteria and limitations based on 2019 OA treatment guidelines, and FDA-approved drug-specific prescribing information
10/21/2020	Removed single treatment cycle limitation for specific drugs
11/09/2021	Removed Priority Partners as an applicable LOB
10/19/2022	Clarified criteria

 $Review\ Dates:\ 04/20/2016,10/19/2016,\ 12/19/2018,\ 01/16/2019,\ 10/16/2019,\ 01/15/2020,\ 07/15/2020,\ 10/21/2020,\ 10/19/2022$

 $Revision\ Dates:\ 1/16/2008,\ 01/13/2009,\ 03/01/2014,\ \ 10/19/2016,\ 07/25/2017,\ 07/27/2017,\ 07/01/2018,\ 12/19/2018,\ 01/16/2019,\ 10/16/2019,\ 06/02/2020,\ 07/15/2020,\ 10/21/2020,\ 11/09/2021,\ 10/19/2022$