

Division: Pharmacy Policy	Subject: State of Florida's Agency for Health Care Administration's
	Prior Authorization Criteria
Original Development Date:	April 22, 2022
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# **Colchicine Agents**

PREFERRED MEDICATIONS	NON-PREFERRED MEDICATIONS
Colchicine 0.6 mg tablet	Colchicine 0.6 mg (Mitigare®) capsule
Probenecid/colchicine tablet	Gloperba® (colchicine solution)
	Colcrys tablet
	Mitigare® (colchicine capsule)

## **LENGTH OF AUTHORIZATION**: 6 months

#### **REVIEW CRITERIA:**

- Familial Mediterranean Fever (FMF) (Colcrys only)
  - O Patient must be  $\geq 4$  years of age.
- Prophylaxis of Gout Flares (Gloperba, Mitigare and Colcrys)
  - o Patient must be  $\geq 18$  years of age.
  - o Patient has tried and failed allopurinol.
  - Patient has tried and failed the preferred medications.
- Treatment of Gout Flares (Colcrys only)
  - Patient must have trial and failure of at least 14 days of NSAID therapy (naproxen, ibuprofen, diclofenac, meloxicam, indomethacin, celecoxib) while on urate lowering therapy (allopurinol, probenecid, febuxostat); OR
  - o Patient must have a history of GI bleeding or comorbidities that would not allow trial of NSAIDs.

### **CONTINUATION OF THERAPY:**

- Patient met initial review criteria; AND
- Documentation of improved clinical response; AND
- Patient has not experienced any treatment-restricting adverse effects; AND
- Dosing is appropriate as per labeling or is supported by compendia.
- For Treatment of Gout Flares:
  - Current history of urate lowering therapy with 100% compliance in the past three months (as per claims history); AND
  - Must have a current history of tophaceous gout (Nodular masses of uric acid crystals [tophi] are deposited in different soft tissue areas of the body); **OR**
  - O Patient has elevated urate level ( $\geq 6$ ) in the past month

#### DOSING AND ADMINISTRATION:

• Refer to product labeling <a href="https://www.accessdata.fda.gov/scripts/cder/daf/">https://www.accessdata.fda.gov/scripts/cder/daf/</a>