





- a) Administer 150 mg subcutaneously at week 0, 1, 2, 3, and 4 and every 4 weeks thereafter. May increase dose to 300 mg every 4 weeks if needed; OR
- b) Administer 150 mg subcutaneously every 4 weeks. May increase dose to 300 mg every 4 weeks if needed; OR
- c) Administer 6 mg/kg by IV infusion at week 0 followed by 1.75 mg/kg every 4 weeks thereafter (max maintenance dose 300 mg per infusion); OR
- d) Administer 1.75 mg/kg by IV infusion every 4 weeks (max maintenance dose 300 mg per infusion). <u>ERA</u>:
- a) Weight-based dosage (see below) is administered by subcutaneous injection at weeks 0, 1, 2, 3%



For reauthorization:

1. Chart notes must show improvement or stabilized signs and symptoms of disease, demonstrated by BSA improvement, etc.

If all the above requirements are met, the medication will be approved for an additional 12 months.

Psoriatic Arthritis (PsA)

For initial authorization:

- 1. Member must be 2 years of age or older; AND
- 2. Medication must be prescribed by or in consultation with a rheumatologist or a dermatologist; AND
- 3. Member has a documented diagnosis of active psoriatic arthritis (PsA); AND
- 4. Member has met a 4-week trial of an NSAID taken at maximally tolerated doses <u>AND</u> a 3-month trial of a conventional DMARD agent (e.g., methotrexate, sulfasalazine, cyclosporine, etc.) <u>unless</u> ONE of the following situations is met:
 - a) Conventional DMARD is **NOT** required for:
 - i) Concomitant axial disease (i.e., involving sacroiliac joint and spine) or enthesitis; OR

b) thus with Brid (the presence with B) [// RE(a) suce TOOD (cequaled do 24:()-7(sym)*nBT/F9(u)-3154.949 rg48] TJ[

i) Severe PsA (defined as having at least **ONE** of the following: erosive disease, active -7()-7(e)



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	the wording of "non-biologic" DMARD to "conventional" DMARD. Clarified reauthorization criteria. Updated references.
11/08/2023	Added HS diagnosis; added/updated references; added IV dosing to applicable dx; simplified TB test requirement wording; added medical benefit option.

References:

- 1. Cosentyx [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; 2023.
- Ward MM, Deodhar A, Gensler LS, et al. 2019 Update of the American College of Rheumatology/Spondylitis Association of America/Spondyloarthritis Research and Treatment Network Recommendations for the Treatment of Ankylosing Spondylitis and Nonradiographic Axial Spondyloarthritis. *Arthritis Rheumatol.* 2019 Oct;71(10):1599-1613. doi: 10.1002/art.41042. Epub 2019 Aug 22.
- 3. Akgul O, Ozgocmen S. Classification criteria for spondyloarthropathies. World J Orthop. 2011;2(12):107