Adalimumab (Humira)

Notes:

- Quantity Limits: Yes
- ^Adequate trial is defined as the following:
 - Phototherapy 8 weeks
 - Systemic non-biologics for psoriasis 6 weeks
 - Topical/oral antibiotics 8 weeks
- * Intolerance excludes adverse drug reactions that are expected, mild in nature, resolve with continued treatment, and do not require medication discontinuation
- **Methotrexate not required if patient has dactylitis (inflammation of finder or toe) and/or enthesitis (inflammation of the entheses)
- Gastroenterology High Risk Classification:
 - Crohn's disease: at least one of the following extensive anatomical involvement, perianal and/or severe rectal disease, deep ulcers, prior surgical resection, stricture and/or penetrating behavior
 - Ulcerative colitis: at least one of the following extensive colitis, deep ulcers, age < 40 years, high CRP and ESR, history of hospitalization, C. difficile infection, CMV infection

<u>Initiation (new start) criteria:</u> Non-formulary **adalimumab (Humira)** will be covered on the prescription drug benefit when the following criteria are met:

- 1. Prescriber is a dermatologist, and patient has a diagnosis of psoriasis
 - Patient has failed an adequate trial[^] of phototherapy (unless documented by prescriber phototherapy not appropriate)
 - Patient has failed an adequate trial, or patient has an allergy or intolerance to at least 1 of the following:
 - Methotrexate
 - Cyclosporine
 - Acitretin
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 2. Prescriber is a dermatologist, and patient has a diagnosis of hidradenitis suppurativa
 - Patient has failed an adequate trial, or patient has an allergy or intolerance to, the following (or contraindication to all):
 - Topical clindamycin 1%
 - Oral antibiotic
 - <u>Brand Biologic with Biosimilar or Unbranded Biologic Available</u> criteria are met

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- 3. Prescriber is a rheumatologist, and patient has a diagnosis of rheumatoid arthritis/inflammatory arthritis
 - Patient has tried and failed/intolerant to as least 1 of the following:
 - Methotrexate
 - o Hydroxychloroquine
 - Sulfasalazine
 - Leflunomide
 - Patient has tried and failed/intolerant to infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 4. Prescriber is a dermatologist or rheumatologist, and patient has a diagnosis of psoriatic arthritis
 - Patient has tried and failed/intolerant to or has contraindication to methotrexate**
 - Patient has tried and failed/intolerant to infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 5. Prescriber is a rheumatologist, and patient has a diagnosis of ankylosing spondylitis/spondyloarthropathy
 - Patient has tried and failed/intolerant to infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 6. Prescriber is a rheumatologist, and patient has a diagnosis of juvenile idiopathic arthritis
 - Patient has tried and failed/intolerant to or has contraindication to methotrexate
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 7. Prescriber is a gastroenterologist, and patient is 17 years of age or younger with a diagnosis of Crohn's disease or ulcerative colitis (if patient is 18 and older, see #8 or #9)
 - Patient has tried and failed/intolerant to infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met

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- 8. Prescriber is a gastroenterologist, and patient is 18 years of age or older with a diagnosis of Crohn's disease
 - If patient is LOW risk:
 - Patient has tried and failed/intolerant to the following:
 - Prednisone or budesonide
 - At least 1 of the following: azathioprine, mercaptopurine, methotrexate
 - Infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
 - If patient is HIGH risk:
 - Patient has tried and failed/intolerant to the following:
 - Infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 9. Prescriber is a gastroenterologist, and patient is 18 years of age or older with a diagnosis of ulcerative colitis
 - If patient is LOW risk:
 - Patient has tried and failed/intolerant to the following:
 - Prednisone
 - At least 1 of the following: mesalamine product (oral or rectal), sulfasalazine
 - At least 1 of the following: azathioprine, mercaptopurine, methotrexate
 - Infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
 - If patient is HIGH risk:
 - Patient has tried and failed/intolerant to the following:
 - Infliximab product (unless documented by prescriber that patient is unable to attend infusion appointments)

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- Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met
- 10. Prescriber is a uveitis specialist, and patient has a diagnosis of iridocyclitis/uveitis
 - Brand Biologic with Biosimilar or Unbranded Biologic Available criteria are met

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<u>Criteria for current Kaiser Permanente members who were previously approved for adalimumab (Humira)</u>: Non-formulary (Humira) will be covered on the prescription drug benefit when the following criteria are met:

- 1. Prescriber is a dermatologist, and patient has a diagnosis of psoriasis or hidradenitis suppurativa
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 2. Prescriber is a rheumatologist, and patient has a diagnosis of rheumatoid arthritis/inflammatory arthritis, psoriatic arthritis, ankylosing spondylitis/spondyloarthropathy, or juvenile idiopathic arthritis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 3. Prescriber is a gastroenterologist, and patient has a diagnosis of Crohn's disease or ulcerative colitis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 4. Prescriber is a uveitis specialist, and patient has a diagnosis of iridocyclitis/uveitis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber

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Adalimumab (Humira)

<u>Criteria</u> for new members entering Kaiser Permanente already taking the medication who have not been reviewed previously: Non-formulary adalimumab (Humira) will be covered on the prescription drug benefit when the following criteria are met:

- 1. Prescriber is a dermatologist, and patient has a diagnosis of psoriasis or hidradenitis suppurativa
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 2. Prescriber is a rheumatologist, and patient has a diagnosis of rheumatoid arthritis/inflammatory arthritis, psoriatic arthritis, ankylosing spondylitis, or juvenile idiopathic arthritis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 3. Prescriber is a gastroenterologist, and patient has a diagnosis of Crohn's disease or ulcerative colitis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) OR
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber
- 4. Prescriber is a uveitis specialist, and patient has a diagnosis of iridocyclitis/uveitis
 - Patient has a documented allergic reaction to adalimumab-atto (does not include injection site reaction) **OR**
 - Patient has tried at least 3 months of adalimumab-atto and experienced persistent new or worsening symptoms of disease as documented with objective findings by the prescriber

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