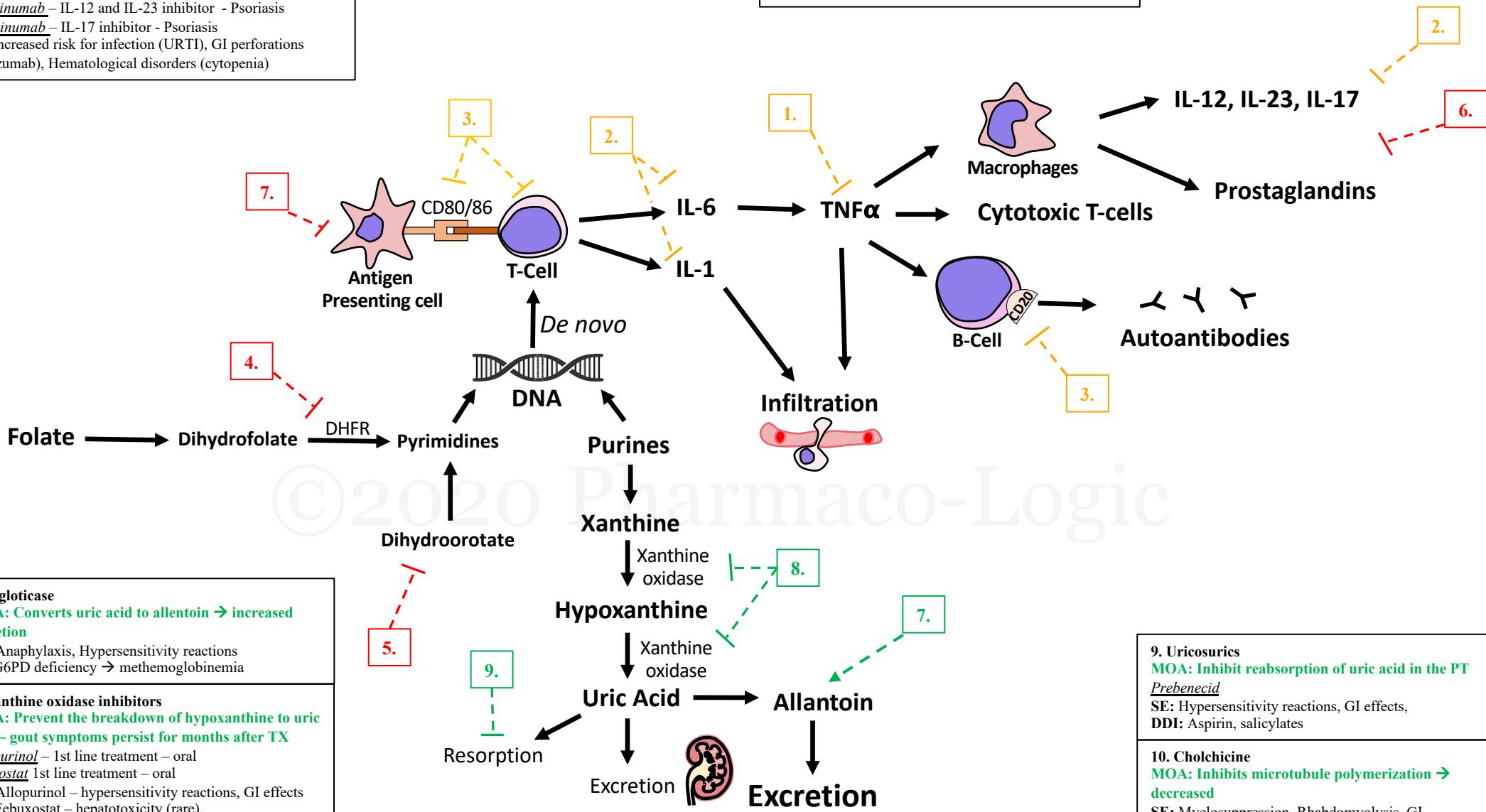


1. TNF alpha antagonists MOA: Inhibit TNF alpha signaling → prevent inflammation and joint destruction <i>Adalimumab</i> <i>Entanercept</i> – soluble TNF receptor <i>Infliximab</i> SE: increased risk for infection (URTI), Hematological disorders (cytopenia), reactivation of TB or HepB	3. Lymphocyte inhibitors MOA: Inhibit TNF alpha signaling → prevent inflammation and joint destruction <i>Rituximab</i> – antibody against CD20 → NK Mediated Death <i>Abatacept</i> – Decreases CD80/86 expression <i>Alefacept</i> - T-cell antagonist – psoriasis SE: Rituximab - Steven Johnson Syndrome, Hep B, PML Abatacept – URTI, Nausea, infections	4. Methotrexate MOA: Inhibitor of dihydrofolate reductase → decreased de novo DNA synthesis USE: RA, psoriasis (low dose), Cancers (high dose) SE: Myelosuppression, hepatotoxicity, nephrotoxicity, stomatitis, Preg Cat X,	6. Sulfasalazine - aminosalicylate MOA: Metabolized to 5-ASA → NSAID – anti-inflammatory USE: RA, Inflammatory bowel disease SE: Myelosuppression, hepatotoxicity, rash, hemolysis in G6PD deficiency
2. Interleukin inhibitors MOA: Inhibit inflammatory interleukin signaling <i>Tocilizumab</i> – IL-6 receptor antagonist <i>Anakinra</i> – IL-1 receptor antagonist <i>Ustekinumab</i> – IL-12 and IL-23 inhibitor - Psoriasis <i>Secukinumab</i> – IL-17 inhibitor - Psoriasis SE: increased risk for infection (URTI), GI perforations (Tolizumab), Hematological disorders (cytopenia)		5. Leflunomide MOA: Metabolized to teriflunomide. Inhibitor of dihydroorotate dehydrogenase → preventing pyrimidine synthesis USE: RA SE: Hepatotoxicity, Preg Cat X,	7. Hydrochloroquine MOA: Inhibit antigen presentation cell function USE: RA, Malaria SE: Rash, bleaching, alopecia, vision problems (regular eye exams)



7. Pegloticase MOA: Converts uric acid to allantoin → increased excretion SE: Anaphylaxis, Hypersensitivity reactions CI: G6PD deficiency → methemoglobinemia
8. Xanthine oxidase inhibitors MOA: Prevent the breakdown of hypoxanthine to uric acid – gout symptoms persist for months after TX <i>Allopurinol</i> – 1st line treatment – oral <i>Febuxostat</i> 1st line treatment – oral SE: Allopurinol – hypersensitivity reactions, GI effects Febuxostat – hepatotoxicity (rare), DDI: Warfarin, Azathioprine, 6-mercaptopurine

9. Uricosurics MOA: Inhibit reabsorption of uric acid in the PT <i>Prebenecid</i> SE: Hypersensitivity reactions, GI effects, DDI: Aspirin, salicylates
10. Cholchicine MOA: Inhibits microtubule polymerization → decreased myelination SE: Myelosuppression, Rhabdomyolysis, GI DDI: Statins, P-Glycoproteins, CYP3A4 inhibitors