

# Systemic Biological Therapies for Psoriasis

If the patient has **moderate to severe disease** - PASI score  $\geq 10$ , DLQI score  $>10$  AND psoriasis has failed to respond to standard systemic therapies including ciclosporin, methotrexate and PUVA, OR the person has a contraindication to, or is intolerant of, these treatments.

**Very severe disease** i.e. PASI score  $\geq 20$ , DLQI score  $>18$  AND psoriasis has failed to respond to standard systemic therapies including ciclosporin, methotrexate and PUVA, OR patient has a contraindication to, or is intolerant of, these treatments.

Choose one of:  
**Apremilast (TA419)**  
**Dimethyl Fumarate (TA475)**

**FIRST LINE BIOLOGICS**  
Choose from the biosimilar agents:  
**Adalimumab (Amgevita) (Anti-TNF) (TA146)**  
**Etanercept (Benepali) (Anti-TNF) (TA146)**  
If pregnant— **Certolizumab (Anti-TNF) (TA574)**

**SECOND LINE BIOLOGICS**  
Choose alternative from:  
**Certolizumab (Anti-TNF) (TA574)**  
**Infliximab (Remsima) (Anti-TNF) (TA134)**  
**Bimekizumab (IL-17A) (TA723)**  
**Brodalumab (IL-17A) (TA350)**  
**Ixekizumab (IL-17A) (TA422)**  
**Secukinumab (IL-17A) (TA350)**  
**Guselkumab (IL-23) (TA521)**  
**Risankizumab (IL-23) (TA596)**  
**Tildrakizumab (IL-23) (TA575)**  
**Ustekinumab (IL-12/23) (TA180)**

**THIRD LINE**  
Choose an alternative mechanism of action from the first and second line choices.

**FOURTH LINE**  
Choose an alternative mechanism of action from the first, second and third line choices.

Choose from biosimilar agents:  
**Infliximab (Remsima) (Anti-TNF) (TA134)**

**If no improvement, stop bDMARD**  
(requests for further bDMARD are subject to IFR process).

An adequate response is defined as: a 75% decrease in PASI score from when treatment started (PASI 75), OR a 50% decrease in PASI score (PASI 50) and a 5 point decrease in DLQI from when treatment started.

If psoriasis has not responded to standard systemic therapies including ciclosporin, methotrexate and PUVA or the person has a contraindication to, or is intolerant of, see flow chart below for treatment options.  
If a person has both psoriasis and psoriatic arthritis, take into account both conditions before initiating or making changes to biological therapy and manage their treatment in consultation with a rheumatologist.

Review if there has been an adequate response to therapy at the following time frames:

- 16 weeks (adalimumab, apremilast, certolizumab, dimethyl fumarate, guselkumab, risankizumab, ustekinumab)
- 12 weeks (brodalumab, etanercept, ixekizumab, secukinumab, tildrakizumab)
- 10 weeks (infliximab)

If secondary non-response or tolerance, move to the next step of the pathway. If positive response, maintain treatment and reassess every 3-6 months.

Choice of Biologic then dependent upon:-

- Cost effectiveness, dosing schedule as per patient needs and patient device/route preference (Adalimumab or Etanercept biosimilar first line when clinically appropriate)
- Proven efficacy and safety profile in long term use
- Patient co-morbidities
- Ease vs complexity of monitoring
- Ustekinumab dose escalation for patients  $<100$  Kg based on patient's response to treatment, local, regional and national experience and close monitoring of effectiveness and side effects profile

**References:** Specialist Pharmacy Service. Updated RMO Advisory Statement: Sequential Use of Biologics. [Internet]. 2020. [Cited March 2022]. Available from <https://www.sps.nhs.uk/articles/rmoc-advisory-statement-sequential-use-of-biologic-medicines/>