

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Ustekinumab

INITIATION – Crohn's disease - adults

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

- ☐ Patient is currently on treatment with ustekinumab commenced prior to 1 February 2023 and met all remaining criteria (criterion 2) below at the time of commencing treatment
- or
- ☐ Patient has active Crohn's disease
- and
- ☐ Patient has had an initial approval for prior biologic therapy for Crohn's disease and has experienced intolerable side effects or insufficient benefit to meet renewal criteria
- or
- ☐ Patient meets the initiation criteria for prior biologic therapies for Crohn's disease
- and
- ☐ Other biologics for Crohn's disease are contraindicated

CONTINUATION – Crohn's disease - adults

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

- ☐ CDAI score has reduced by 100 points, or HBI score has reduced by 3 points, from when the patient was initiated on biologic therapy
- or
- ☐ CDAI score is 150 or less, or HBI is 4 or less
- or
- ☐ The patient has experienced an adequate response to treatment, but CDAI score and/or HBI score cannot be assessed
- and
- ☐ Ustekinumab to be administered at a dose no greater than 90 mg every 8 weeks

INITIATION – Crohn's disease - children*

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

- ☐ Patient is currently on treatment with ustekinumab commenced prior to 1 February 2023 and met all remaining criteria (criterion 2) below at the time of commencing treatment
- or
- ☐ Patient has active Crohn's disease
- and
- ☐ Patient has had an initial approval for prior biologic therapy and has experienced intolerable side effects or insufficient benefit to meet renewal criteria
- or
- ☐ Patient meets the initiation criteria for prior biologic therapies for Crohn's disease
- and
- ☐ Other biologics for Crohn's disease are contraindicated

Note: Indication marked with * is an unapproved indication.

I confirm that the above details are correct:

Signed: Date:

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PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Ustekinumab - continued

CONTINUATION – Crohn's disease - children*

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

- ☐ PCDAI score has reduced by 10 points from when the patient was initiated on biologic therapy
- or
- ☐ PCDAI score is 15 or less
- or
- ☐ The patient has experienced an adequate response to treatment, but CDAI score cannot be assessed

and

- ☐ Ustekinumab to administered at a dose no greater than 90 mg every 8 weeks

Note: Indication marked with * is an unapproved indication.

INITIATION – ulcerative colitis

Re-assessment required after 6 months

Prerequisites (tick boxes where appropriate)

- ☐ Patient is currently on treatment with ustekinumab commenced prior to 1 February 2023 and met all remaining criteria (criterion 2) below at the time of commencing treatment
- or
- ☐ Patient has active ulcerative colitis
- and
- ☐ Patient has had an initial approval for prior biologic therapy for ulcerative colitis and has experienced intolerable side effects or insufficient benefit to meet renewal criteria
- or
- ☐ Patient meets the initiation criteria for prior biologic therapies for ulcerative colitis
- and
- ☐ Other biologics for ulcerative colitis are contraindicated

CONTINUATION – ulcerative colitis

Re-assessment required after 12 months

Prerequisites (tick boxes where appropriate)

- ☐ The SCCAI score has reduced by 2 points or more from the SCCAI score since initiation on biologic therapy
- or
- ☐ PUCAI score has reduced by 10 points or more from the PUCAI score since initiation on biologic therapy*

and

- ☐ Ustekinumab will be used at a dose no greater than 90 mg intravenously every 8 weeks

Note: Criterion marked with * is for an unapproved indication.

I confirm that the above details are correct:

Signed: Date: