## HOSPITAL MEDICINES LIST RESTRICTIONS CHECKLIST

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.

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|-----------------------|---|--|-------------------|--|
| PRESCRI               | IBER  | PATIENT:   | PATIENT:          |  |
| Name:                 |   | Name:  | Name:             |  |
| Ward:                 |   | NHI:   |                   |  |
| Febuxo                | stat  |  |                   |  |
| INITIATIO<br>Prerequi |   | Gout (tick boxes where appropriate)  |                   |  |
| and                   | d   | Patient has been diagnosed with gout   |                   |  |
|                       |   | O The patient has a serum urate level greater than 0.36 mmol/l despite treatment with allopurinol at doses of at lea and addition of probenecid at doses of up to 2 g per day or maximum tolerated dose  | st 600 mg/day     |  |
|                       | or  | The patient has experienced intolerable side effects from allopurinol such that treatment discontinuation is required and serum urate remains greater than 0.36 mmol/l despite use of probenecid at doses of up to 2 g per day or maximum tolerated dose |                   |  |
|                       | or  | O The patient has renal impairment such that probenecid is contraindicated or likely to be ineffective and serum urate remains greater than 0.36 mmol/l despite optimal treatment with allopurinol (see Note)  |                   |  |
|                       | OI OI   | O The patient has previously had an initial Special Authority approval for benzbromarone for treatment of gout.  |                   |  |
| Prerequi              | Re-assessment required after 6 weeks Prerequisites (tick boxes where appropriate)  Prescribed by, or recommended by a haematologist or oncologist, or in accordance with a protocol or guideline that has been endorsed by the Health NZ Hospital.  Patient is scheduled to receive cancer therapy carrying an intermediate or high risk of tumour lysis syndrome |  |                   |  |
| and                   |   | Patient has a documented history of allopurinol intolerance  |                   |  |
| Re-asses              | ssmer<br>isites<br>Preso  | ON – Tumour lysis syndrome It required after 6 weeks (tick box where appropriate)  cribed by, or recommended by a haematologist or oncologist, or in accordance with a protocol or guideline that has been the NZ Hospital.                              | n endorsed by the |  |
| O                     | The treatment remains appropriate and patient is benefitting from treatment   |  |                   |  |
|                       |   |  |                   |  |

| I confirm that the above details are correct: |       |
|---|-------|
| Signed:                                       | Date: |