Technology Guidance

Ustekinumab biosimilar

for treating inflammatory conditions

Technology Guidance from the MOH Drug Advisory Committee

Guidance Recommendations

The Ministry of Health's Drug Advisory Committee has recommended ustekinumab biosimilar (Steqeyma) 45 mg/0.5 mL pre-filled syringe, 90 mg/1 mL pre-filled syringe, and 130 mg/26 mL vial for listing on the MOH Standard Drug List (SDL). The decision was based on the acceptable pricing proposal from the company.

Funding status

SDL subsidy will apply from 1 April 2026 to all registered indications of ustekinumab biosimilar (Steqeyma) 45 mg/0.5 mL pre-filled syringe, 90 mg/1 mL pre-filled syringe, and 130 mg/26 mL vial in Singapore.

SDL subsidy **does not** apply to other brand(s), formulation(s) or strength(s) of ustekinumab.

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Agency for Care Effectiveness - ACE Agency for Care Effectiveness (ACE)

About the Agency

The Agency for Care Effectiveness (ACE) was established by the Ministry of Health (Singapore) to drive better decision-making in healthcare through health technology assessment (HTA), clinical guidance, and education.

As the national HTA agency, ACE conducts evaluations to inform government funding decisions for treatments, diagnostic tests and vaccines, and produces guidance for public hospitals and institutions in Singapore.

The guidance is not, and should not be regarded as, a substitute for professional or medical advice. Please seek the advice of a qualified healthcare professional about any medical condition. The responsibility for making decisions appropriate to the circumstances of the individual patient remains with the healthcare professional.

Find out more about ACE at www.ace-hta.gov.sg/about

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