

Evomela® (melphalan) – Indication withdrawal

- On August 9, 2021, the <u>FDA approved</u> the voluntary removal of Acrotech Biopharma's <u>Evomela</u> (<u>melphalan</u>) indication for palliative treatment of patients with multiple myeloma for whom oral therapy is not appropriate.
- Evomela remains approved for use as a high-dose conditioning treatment prior to hematopoietic progenitor (stem) cell transplantation in patients with multiple myeloma.
- Evomela carries a boxed warning for severe bone marrow suppression, hypersensitivity, and leukemogenicity.



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