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Subcutaneous Injection for UCENTYVIOprefilled syringe andENTYVIOOPEN are intended for subcutaneous use under the guidance and supervision of a healthcare professional

Patients or caregivers may self inject subcutaneousENTYVIOusing either theENTYVIOprefilled syringe orENTYVIOOPEN after training in subcutaneous injection technique Learn more aboutENTYVIO vedolizumab a biologic treatment for moderate to severe ulcerative colitis or Crohn s disease in adults Dec 24 2025 Takeda Pharmaceuticals America Inc

ENTYVIOis indicated in adults for the treatment of moderately to severely active ulcerative colitis UC

moderately to severely active Crohn s disease CD .ENTYVIOis an integrin Entyviois indicated for the treatment of adult patients with moderately to severely active Crohns disease who have had an inadequate response with lost response to or were intolerant to either conventional therapy or a tumour necrosis factor alpha TNF antagonist Single dose prefilled pen This Instructions for Use contains information on how to injectENTYVIO These highlights do not include all the information needed to useENTYVIO safely and effectively

See full prescribing information forENTYVIO Dec 19 2025 Insertthe syringe needle into the vial through the center of the stopper and direct the stream of Sterile Water for Injection to the glass wall of the vial to avoid excessive foaming

Gently swirl the vial for at least 15 seconds to dissolve the lyophilized powder

Do not vigorously shake or invert ReconstituteENTYVIOvial containing lyophilized powder with 4.8 mL of Sterile Water for injection using a syringe with a 21 to 25 gauge needle.Insertthe syringe needle into the vial through the center of the stopper and direct the stream of Sterile Water for injection to the glass wall of the vial to avoid excessive foaming Entyvio vedolizumab for subcutaneous use is an integrin receptor antagonist indicated in adults for the treatment of moderately to severely active ulcerative colitis UC and Crohns disease CD Entyvio300 mg powder for concentrate for solution for infusion is administered as an intravenous infusion over 30 minutes

Patients should be monitored during and after infusion see section 4.4. For instructions on reconstitution and dilution of the medicinal product before administration see section 6.6.

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