



## CSL Behring – Withdrawal of Hizentra® [immune globulin subcutaneous (human)]

- On December 30, 2021 [CSL Behring announced](#) a patient-level withdrawal of one lot of [Hizentra \[immune globulin subcutaneous \(human\)\]](#) due to an increased frequency of reports of injection-site reactions and local hypersensitivity-type of events after administration.
- The withdrawn lot was shipped from CSL Behring beginning on October 5, 2021.

Product Description	NDC	Lot # (Expiration Date)
Hizentra [immune globulin subcutaneous (human)] 20% liquid	44206-0455-10	P100343632 (11/24/2023)

- Hizentra is indicated for the treatment of primary immunodeficiency in adults and pediatric patients 2 years of age and older and for maintenance therapy in adults with chronic inflammatory demyelinating polyneuropathy.
- Injection-site reactions and hypersensitivity are a known risk with subcutaneous immune globulin products.
- Patients should contact their healthcare provider if they have experienced any problems that may be related to using the withdrawn Hizentra.
- Contact CSL Behring Medical Information at **1-800-504-5434** or email [MedinfoNA@cslbehring.com](mailto:MedinfoNA@cslbehring.com) for more information about the withdrawal.



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