Takeda – Withdrawal of Vonvendi® (von Willebrand factor, recombinant)

- On January 26, 2024, Takeda announced a consumer level withdrawal of two lots of Vonvendi (von Willebrand factor, recombinant) because of misprinted product labels with an incorrect expiration date.

- There is no quality issue with Vonvendi. The safety and efficacy profile remains consistent with the labeled information. The expiration date on the outside of the package (June 27, 2025) is six months after the actual expiration date (January 27, 2025).

- The withdrawn lots are listed below:

Product Description	NDC number	Lot number (Exp Date)
Vonvendi (von Willebrand factor, recombinant), 650 IU	0944-7550-01, 0944-7551-02, 0944-7553-02	TVA22004AA (6/27/2025); TVA22004AB (6/27/2025)

- Vonvendi is indicated for use in adults diagnosed with von Willebrand disease (VWD) for on-demand treatment and control of bleeding episodes, perioperative management of bleeding, routine prophylaxis to reduce frequency of bleeding episodes in patients with severe Type 3 VWD receiving on-demand therapy.

- Anyone with the affected products on hand should discontinue use, stop distribution and quarantine product immediately. Patients with impacted product on hand should return to their pharmacy for a replacement.

- Patients should contact their healthcare provider if they have experienced any problems that may be related to using the withdrawn products.

- Consumers may contact Takeda at **1-877-825-3327** for more information.