

Who's in the Community Counts Registry?

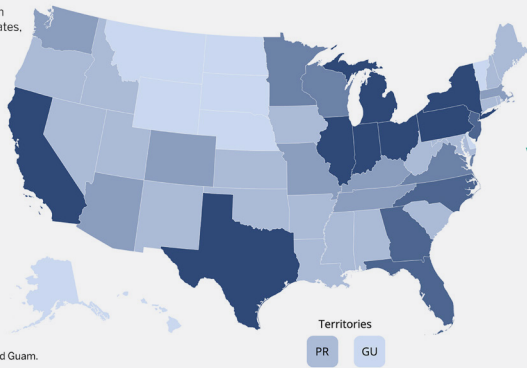
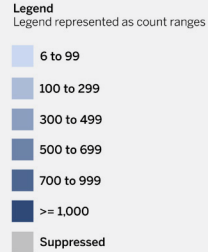
Data from start of the program in 2012 to 11/2022

The Data Visualization Tool Gives Us Answers!

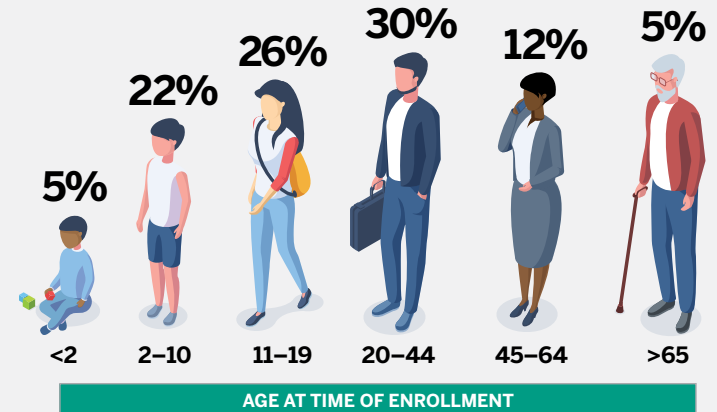
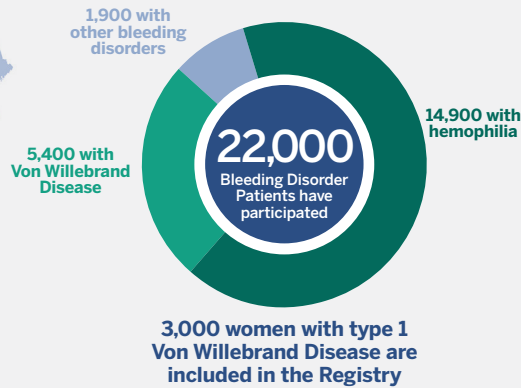
The Registry for Bleeding Disorders Surveillance gathers detailed information on HTC patients with bleeding disorders with the aim to improve the medical care and health outcomes of people with hemophilia and other bleeding disorders.

Patient Geographic Distribution

This map displays the geographic distribution of Registry participants across the United States, Puerto Rico and Guam.



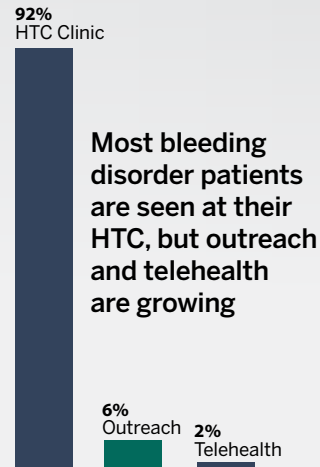
Participants are from all U.S. States, Puerto Rico and Guam.



Things we have learned from the Registry:

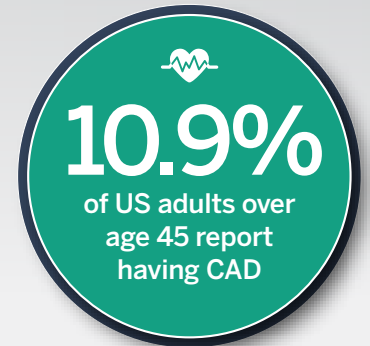
79%

of boys with severe hemophilia under age 10 are on continuous prophylaxis



At enrollment, **5.3%** of participants with hemophilia over age 45 were noted to have Coronary Artery Disease (CAD)

VS.



cdc.gov/aging/publications/coronary-heart-disease-brief.html



Check out the Data Visualization tool yourself! [Click here](#) or scan the QR code with your smart device