

# HANDI HIGHLIGHTS - RARE AND ULTRA RARE BLEEDING DISORDERS

We are celebrating Rare Disease Day by highlighting resources for healthcare providers and people with rare and ultra rare bleeding disorders.



## Educational Resources

[Rare Factor Deficiencies](#)

[Untold Journeys: A Glimpse into Ultra-Rare Bleeding Disorders](#) 

Factor V ([English](#) and [Spanish](#))

Factor VII ([English](#) and [Spanish](#))

- Super Seven: A Story for Rares ([English](#) and [Spanish](#))
- Super Seven: Discussion Guide ([English](#) and [Spanish](#))

Factor X ([English](#) and [Spanish](#))

Factor XIII ([English](#) and [Spanish](#))

Glanzmann's Thrombasthenia ([English](#) and [Spanish](#))

- [Glanzmann's Thrombasthenia and Bernard-Soulier Syndrome](#)

Platelet Storage Pool ([English](#) and [Spanish](#))

[The Power of Connection for People with Rare Bleeding Disorders](#)



## Provider Resources

[Rare Disease Clinical Research Network](#)

[National Organization for Rare Diseases](#)

[Rare Coagulation Disorders Resource Room](#)

[Current Treatments for Ultra-Rare Bleeding Disorders](#) 

[NBDF National Research Blueprint White Paper](#)



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Check your [local chapters](#) for more information on local events

Visit the [HANDI homepage](#) to submit a request for information and to learn the answers to frequently asked questions.

*NBDF recommends that all individuals consult their HTC or primary physician for medical advice and any questions regarding their choice of treatment.*