



Committed to One National Data Source for the Hemostasis and Thrombosis Community

Securely hosted by the Centers for
Disease Control and Prevention (CDC)





Coming soon to your Hemophilia Treatment Center (HTC)... ATHN

Collaborate... *The American Thrombosis and Hemostasis Network (ATHN) is collaborating with Hemophilia Treatment Centers, non-profit organizations and government agencies to collect core data in a standardized electronic format and then use it to advance the science and practice of care for better outcomes, to help preserve access to comprehensive care and to avoid disruption of care in case of an emergency.*

Prepare... *You can have your HTC health data "in your pocket" in case of disaster.*

With ATHN's help, health professionals around the country will define and protect the core meaningful health data needed to support your care as part of a national emergency preparedness plan.

ATHN will be working with HTCs, the CDC, national organizations and local chapters to seek input, test the process and keep you informed.

Ask... *If your HTC is an ATHN Affiliate working to better understand the epidemiology, genetics, natural history and treatments of bleeding and blood clotting disorders.*

How you can consent to have your non-identifiable data pooled to improve outcomes for you, your family and generations to come.

How you can leave a legacy.

our vision. *To advance and improve the care of individuals affected by bleeding and thrombotic disorders.*

our mission. *To provide stewardship of a secure national database, adherent to all privacy guidelines, which will be used to support clinical outcomes analysis, research, advocacy and public health reporting in the hemostasis and thrombosis community.*

our values. *Improving clinical outcomes and care, facilitating continuity of care, fostering collaboration, maintaining confidentiality, conserving resources through a common infrastructure.*



american thrombosis & hemostasis network

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