

INDIAN ASSOCIATION FOR HAEMOPHILIA AND ALLIED DISORDERS

HAEMOPHILIA TREATMENT CENTER REGISTRATION FORM

Please note:

- 1. This is an editable PDF. Please fill this and share back to IAHAD at pe@iahad.org
- 2. You can also print, fill and scan a copy back to us at pe@iahad.org
- 3. Kindly note that only ONE registration will be accepted per HTC. Please discuss with your team before sharing the final version.

Name of the HTC:Address:	
State:	
Email ID of the HTC:	
Name of the Director/HTC In-Charge: Email ID:	
 Total number of patients with a Hereditary Bleedin Total number of patients with a Hereditary Bleedin 	

Bleeding Disorder	Patients Registered	Patients with Annual Follow-Up	Patients with Severe Disease
Haemophilia A		•	
Haemophilia B			
VWD Type 1			
VWD Type 2			
VWD Type 3			
VWD Type Unknown			
Afibrinogenemia (<10mg/dl)			
Hypofibrinogenemia (50-150mg/dl)			
Dysfibrinogenemia (10-50mg/dl)			
Factor II Deficiency			
Factor V Deficiency			
Factor VII Deficiency			
Factor X Deficiency			
Factor XI Deficiency			
Factor XIII Deficiency			
Combined Factor V + VIII			
Deficiency			
Combined Factor II + VII + IX + X			
Deficiency			
Glanzmann's thrombasthenia			
Bernard Soulier Syndrome			
Others			



<u>Haemophilia Treatment Center - Comprehensive Team members details.</u> (Members can be added every year. Member List to be updated every year)

MEMBERS	Name (Full name with initials)	E-mail address
Physician		
Pathologist		
Nurse Coordinator		
Physiotherapist		
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Orthopaedic Surgeon		
Other Surgeons		
Laboratory Scientists		
Data Manager		
Data Manager		
Psycho-Social Team		
Coordinator		
Other		
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^{**} Each Category can have multiple names. ***Leave it blank if not available

Consent: I agree to register our HTC as an Institutional Member of IAHAD.

I hereby declare that the entries made in this form as above is true and correct to the best of my knowledge and belief.

Signature - Head / Physician In-charge of HTC

Name	Signature	Date