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WFH 2026 WORLD CONGRESS

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PP-223	Novel Variants in Hemophilia A Patients: Report From a Referral Hemophilia Treatment Centre in Indonesia	Fitri Primacakti
PP-225	Clinical Profile of Hemophilia Children Admitted to a Limited-Resource Hospital in West Nusa Tenggara, Indonesia: A Cross-Sectional Study	Yudhi Kurniawan
PP-226	Recurrent Hemophilic Pseudotumor in Mild Hemophilia A With Neurofibromatosis Type 1: Experience From a Low- and Middle-Income Country	Purnamasari Natsir Putri
PP-227	Paediatric Hemophilia B in Northern Peninsular Malaysia: A Descriptive Analysis of Patient Profiles and Outcomes	Chee Enn Han
PP-228	Safe Childhood: Helping Children With Hemophilia A Stay Active and Confident	Aamir Aslam Awan

PP-229	Epidural Hematoma on the Left Frontoparietotemporal Region in a Child With Newly Diagnosed Severe Hemophilia A: A Case Report	Salsa Ardhana Makruf
PO-230	Concomitant Bleeding and Thrombosis in a Cohort of Genetically Predisposed Egyptian Children Presenting With Pediatric Thrombosis	Magy Abdelwahab

Physiotherapy and Rehabilitation

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PP-233	Recreative Sport, Joint Health, and Musculoskeletal Power in Children and Young People With Haemophilia	Carlos Cruz-Montecinos
PP-235	Ten Years of Multidisciplinary Rehabilitation Check-Ups for People With Hemophilia in Japan: Long-Term Survivors of HIV Acquired From Historically Contaminated Blood Products	Junko Fujitani
PP-236	A Comparison of Patients Presenting With Musculoskeletal Issues to the Bleeding Disorder Rheumatology Clinic 10 Years Apart	Abigail Polus
PP-237	The Effects of Physical & Mental Health Rehabilitation Program (PMHRP) in Post Operative Periods for Hemophilic Arthritis	Won-Sook Bak
PP-238	A Prospective Ultrasound Framework for Physiotherapists Working in Haemophilia Care	James Cartwright
PP-241	Safety and Efficacy of Physiotherapy in Patients with Severe Hemophilia A Treated with Emicizumab: A Prospective Polish Multicenter Study	Ewa Stefańska-Windyga
PO-243	Role of Physiotherapy to Rehabilitate a Person With Hemophilia Induced Foot Drop: A Case Study	Easmin Ara Doly
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Prophylaxis

POSTER #	TITLE	PRESENTING AUTHOR
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PP-253	Real-World Experience With Emicizumab Prophylaxis From a Single Canadian Tertiary Care Center	Areeb Hassan
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PP-256	The Role of Low Dose Emicizumab Prophylaxis in Improving PedHAL Domains and Clinical Outcomes in Children With Hemophilia: An Insight From a Low-Middle Income Country	Shigy Francis
PP-257	Level of Knowledge of Hemophilia Patient's Families in Implementing Prophylactic Therapy at the Hemophilia Treatment Center Ulin General Hospital , Banjarmasin, Indonesia	Wulandewi Marhaeni
PP-258	Joint Status and Activity Patients With Moderate and Severe Hemophilia in Chiang Mai University Hospital	Rungrote Natesirinilkul
PP-259	Determinants of Adherence to Clotting Factor Prophylaxis in Adolescent and Adult Men With Hemophilia	Alice Sacramento
PP-261	Management for Hemophilia A Pediatric Patients Without Inhibitors Receiving Nationwide Emicizumab Prophylaxis	Sanikarn Nakrong
PP-262	New Confidence Through Modern Prophylaxis: Community Experiences With Emicizumab in Hemophilia A	Aamir Aslam Awan
PP-263	Bleeding Outcomes >3 Years After Discontinuing Emicizumab Among Persons With Hemophilia A in a Low-Resource Setting: Experience From Moi Teaching and Referral Hospital, Kenya	Aliwa Everlyn
PP-264	Outcomes of Emicizumab Prophylaxis in Inhibitor-Positive Severe Hemophilia A: A Nationwide Population-Based Study in Taiwan	Chia-Jen Liu
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Psychosocial Issues

POSTER #	TITLE	PRESENTING AUTHOR
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MP-268	Tea Party Care in the Hemophilia Community: A Psychosocial Healing Program for Parents of Children With Hemophilia	Seunggeun Kim
MP-269	Patient-Reported Outcomes on Artificial Intelligence Use in Hemophilia: A Psychometric Evaluation	Cindy Komar
PP-271	Mental Health Needs Assessment Among People With Hemophilia in Nepal: A Mixed- Methods Study	Toong Youttananukorn
PP-272	Evaluating the World Federation of Hemophilia's Shared Decision-Making Tool: Insights From a Multi-group Impact Study	Donna Coffin
PP-273	Emotional Burden and Marginalization in Individuals With Bleeding Disorders	Midhat Khalid
PP-275	The Importance of the Psychosocial Approach in the Healthcare Team: Current Reality and Future Perspectives	Silvina Grana
PP-276	Advancing Mental Health and Substance Use Care Accessibility for Individuals With Bleeding Disorders: A United States Multi-state Outreach and Champion Facility Identification Model	Shanthi Hegde
PP-278	Navigating Life With Bleeding Disorders: Understanding the Lived Psychosocial Experiences of Youth	Midhat Khalid
PP-279	Parenting Challenges in the Overlap of Autism and Hemophilia: A Phenomenological Case Report From Iran	Fatemeh Maleki
PP-280	Psychosocial Well Being and Exercise Participation of Patients With Haemophilia and the Carers in Hong Kong	Henry Hon-Wai Lam
PP-281	Feasibility and Engagement of an Online CBT-Based Group Intervention for Adolescents With Haemophilia	Anna-Maria Tsilia
PP-283	Neurobiological Perspectives on Hemophilia: A Review of Neurological, Cognitive, and Psychosocial Interactions	Ferose Pallimanhayalil
PP-284	Beyond Carriers: Promoting Recognition, Diagnosis, and Psychosocial Inclusion for Women with Bleeding Disorders in Korea	Eunju Jo
PP-285	Peer-Led Support Groups as a Strategy to Reduce Caregiver Isolation in Kenya's Haemophilia Community	Sylvia Kathurima

PP-286	From Paternalism to Shared Decision Making: A Qualitative Phenomenological Study of Physician–Patient Communication Styles in Pediatric Hemophilia Within the Iranian Cultural Context	Ali Eshghi
PP-287	Comparison of Psycho-Social Difficulties Faced by Haemophilia Patients With and Without HIV/AIDS	Tomie Fujii
PP-288	Creating a Children's Book as a Tool for Education and Support	Darian Smith
PP-290	Innovations in Child Life Services for Hemophilia Care in Kenya: Enhancing Psychosocial Support and Treatment Compliance for Pediatric Patients Living With Hemophilia	Bilha Nyaboke Omari
PP-291	Balancing Hemophilia Care and Cultural Practices: Psychosocial and Clinical Outcomes of Circumcision With Optimized Factor Therapy	Vijayakumar Narayana Pillai
PP-292	Ibero-American Psychosocial Group: A Collaborative Network in Development	Silvina Graña
PP-295	New Educational Strategies for Individuals With Hemophilia and Autism	Tongpil Min
PP-296	Hemophilia and Autism: Where Do Challenges Come From?	Gavin Min
PP-297	A Study on Development of RAG-Based Psychological Counseling Support System via Analysis of Unmet Counseling Needs in Hemophilia Patients	Jihun Chang
PO-298	Bleeding in Silence: A Young Man's Battle for Identity, Dignity, and Hope With Severe Hemophilia A	Midhat Khalid
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MP-303	Psychological Well-Being in Primary Caregivers of People With Coagulopathies	Ezequiel Martinez
PO-304	The Invisible Burden: A Systematic Review of Psychosocial Issues in Patients With Hemophilia	Muhammad Nadeem Iqbal Zahid

Quality of Life / Outcome Research

POSTER #	TITLE	PRESENTING AUTHOR
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MP-306	Hemophilia Life Stages and Changes Global Survey of People With Hemophilia, Caregivers, and Hematologists: Patient Experience According to Disease Severity and Gender	Kim Isenberg
MP-307	Psychological Burden and Quality of Life of Mothers of Hemophiliac Children: A Study of 69 Cases	El Khorassani Mohammed
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PP-310	WBDR RSP Series Part 2/6: Generating Clinical Evidence That Informs Policy – Four Focused Research Areas from 32 RSP Projects (2018–2023)	Toong Youttananukorn
MP-311	WBDR RSP Series Part 3/6: Diagnosis Evidence– Findings From Research Projects Under the WFH WBDR Research Support Program (2018–2023)	Toong Youttananukorn
PP-312	WBDR RSP Series Part 4/6: Quality-of-Life Evidence – Findings From Research Projects Under the WFH WBDR Research Support Program (2018–2023)	Toong Youttananukorn
PP-313	WBDR RSP Series Part 5/6: Prophylaxis Evidence - Findings From Research Projects Under the WFH WBDR Research Support Program (2018–2023)	Toong Youttananukorn
PP-314	WBDR RSP Series Part 6/6: Underserved Populations (WGBD/VWD) Evidence - Findings From Research Projects Under the WFH WBDR Research Support Program (2018–2023)	Toong Youttananukorn
PP-315	Insights Into Physical Function and Treatment Experience With Mim8: Patient Cases From the FRONTIER Trials	Francisco José López-Jaime
PP-316	Patient-Reported Outcomes in Hemophilia in Kosovo: Results From the PROBE Questionnaire	Dawn Rotellini
PP-317	The Quality of Life of Hemophilia A Patients Accessing Treatment Services	Toong Youttananukorn
PP-318	Long-Term Treatment Adherence Decline in Pediatric Hemophilia Patients Receiving Extended Half-Life Therapies: A Cross-Sectional Study in South Korea	Harim Kim
PP-319	Assessing the Change in Bleeding Rates and Related Clinical Outcomes in Haemophilia Patients Before and After Introducing Humanitarian Aid Clotting Factor Concentrates in Malawi	Brian Dimba
PP-320	Real-World Effectiveness and Patient-Reported Burden of Pharmacokinetic-Guided Individualized Prophylaxis in Chinese Hemophilia A Patients	Zeping Zhou

PP-321	Health-Related Quality of Life among Adults with Haemophilia in Southern Nigeria	Toong Youttananukorn
PP-322	Adapting the Patient Reported Outcomes Burdens and Experiences (PROBE) Study to Measure Quality of Life in People With von Willebrand Disease (VWD)	Elizabeth Clearfield
PP-323	Mapping the Journey of Caregivers and Patients With Hemophilia A and B in Brazil	Indianara Galhardo
PP-324	Parental Experience of Hemophilia Diagnosis Disclosure in Morocco: A Mixed-Methods Study on Communicational Quality and Psychosocial Impact	El Khorassani Mohammed
PP-325	Quality of Life Assessment in Parents of Infants With Severe Hemophilia A Without Inhibitors After Emicizumab Prophylaxis	Canan Albayrak
PP-326	Empowering Movement in Hemophilia: A Medically Supervised Trekking Experience	Carlos Kolker
PP-327	Mapping the Journey of Caregivers and Patients Under 6 Years Old With Hemophilia A in Brazil	Mariana Battazza
PP-329	Impact of Hemophilia on Quality of Life and Self-Esteem in Adolescents and Young Adults in South India	Ferose Pallimanhayalil
PP-331	Evaluation of the Quality of Life of Hemophiliac Patients Under Prophylactic Treatment in Cameroon	Toong Youttananukorn
PP-332	Assessment of Joint Arthropathy by FISH and Pain Scoring in Hemophilia Patients: Experience From Pakistan	Munira Borhany
PP-333	Emicizumab Prophylaxis and Its Impact on Bleeding Control, Joint Health, and Psychosocial Well-Being in Hemophilia A	Ayisha Imran
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PO-336	Study on Quality of Life and Health Care in Men With Hemophilia in Uruguay, Comparative Studies	Isabel Sorondo
PO-337	Increasing Patient and Family Knowledge Regarding the Joint Health of Hemophilia Patients: Educational Intervention at Hemophilia Treatment Center Ulin General Hospital, Banjarmasin, Indonesia	Ratih Kumala Sari
PO-339	Impact of Genetic Counseling on Prevention of Factor VII Deficiency in a Family From Pakistan	Lubna Zafar

PO-342	Assessment of Quality of Life in People With Hemophilia on Coagulation Factor Prophylaxis in Brazil	Ricardo Mesquita Camelo
PO-343	Filipino Version of the Haem-a-QOL Questionnaire Among Adult Hemophilia Patients Followed- Up at the Philippine General Hospital: A Validation Study	Ma Angelina L Mirasol

Rare Bleeding Disorders

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PP-346	Hereditary Severe Factor X Deficiency in Childhood: Single Centre Experience at Evelina London Children's Hospital	Jayanthi Alamelu
PP-347	A Sticky Situation: An Illustrated Review of Glanzmann's Thrombasthenia	Eliza VanZweden
PP-348	Use of Emicizumab in Patients With Hemophilia in Low- or Middle-Income Countries	Majid Naderi
PP-349	Development of a Global Database: The Glanzmann Thrombasthenia Real-World Insights Research Study	Amy Owen-Wyard
PP-351	Acquired Hemophilia A in a Resource-Limited Setting: Demographic Profile and Real-World Outcomes From a Multi-Centre Retrospective Study	Rema Ganapathy
PP-352	Susoctocog Alfa for Bleeding Control in High-Titer Acquired Hemophilia A: A Case Series of Four Patients	Jae Joon Han
PP-353	Acquired Deficiency of Vitamin K Dependent Factors: An Alarming Increase in Cases. Experience in Banco Municipal De Sangre DC, Caracas	Apsara Boadas
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Surgical Treatment

POSTER #	TITLE	PRESENTING AUTHOR
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PP-361	Perioperative Management With Lonoctocog Alfa (Afstyla) in Hemophilia A: Real-World Surgical Experience	Olga Benítez Hidalgo
PP-362	Surgical Management in Hemophilia Patients Receiving Efmoroctocog Alfa	Goksel Leblebisatan
PP-363	Expanding Surgical Boundaries in Severe Hemophilia A: Successful Orthognathic Correction Using PK-Guided Extended Half-Life FVIII in a Middle-Income Country	Eileen Viviana Fonseca
PO-364	Surgical Resection and Perioperative Management of Large Haemophilic Pseudotumors: Case Report	Theofanis Adraktas
PO-365	Perioperative Management and Surgical Outcomes in Patients With Inherited Bleeding Disorders in a Public Sector Hospital: The Pakistan Institute of Medical Sciences Experience	Farwa Sijjeel
PO-366	Intussusception Secondary to Meckel's Diverticulitis: A Rare and Life-Threatening Complication in an Infant With Hemophilia A	Serap Karaman
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Trials in Progress

POSTER #	TITLE	PRESENTING AUTHOR
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Von Willebrand Disease

POSTER #	TITLE	PRESENTING AUTHOR
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MP-374	Recombinant von Willebrand Factor, Vonicog Alfa: United Kingdom Experience of Its Use in Children Under 18 Years of Age	Jayanthi Alamelu
MP-375	Clinical Spectrum, Subtype Distribution, And Treatment Outcomes In Von Willebrand Disease: A Prospective Study From A Hemophilia Treatment Center In Pakistan	Munira Borhany
PP-377	Recombinant von Willebrand Factor (rVWF) for Surgical Haemostasis and Bleeding Management: A 5-Year Retrospective Review From a UK Haemophilia Centre	Yu Yan Carmen Lee
PP-378	Congenital Von Willebrand Disease: Management of Pregnancy and Delivery	Vera Geierova
PP-379	Expanding the PIVOT-vWD Programme: An International Patient-Driven Study to Capture the Impact, Voice, and Outcomes of von Willebrand Disease	Robert Sidonio
PP-380	Pregnancy Outcomes and Hemostatic Management in Women With Von Willebrand Disease: A Single-Center Experience	Chin Sheng Ting

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PP-382	Successfully Searching to See Type 1C Von Willebrand Disease	Karen Sims
PP-383	Bridging the Diagnostic Gap: Development and Validation of a Localized Screening Tool for Von Willebrand Disease in a Taiwanese Cohort	Shin-Nan Cheng
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PP-385	Managing Type 3 von Willebrand Disease Complicated by Anti-VWF Alloantibodies: A Therapeutic Challenge	Emna Gouider
MP-386	Global Utilisation and Impact of VWDtest.com: A Multilingual Digital Platform to Improve Awareness and Early Identification of von Willebrand Disease	Fernando F. Corrales-Medina
PO-387	A Single Documented Case of Thalassemia Major With Concomitant Presence of Von Willebrand Disease: Exceptional Co-Occurrence	Farah Hanif

Women and Girls' Health and Research

POSTER #	TITLE	PRESENTING AUTHOR
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MP-391	Strengthening Diagnostics and Patient Care for Women and Girls With Bleeding Disorders Through International Capacity-Building and Twinning Programme	Bitty Kurian
MP-392	Diagnostic Delay and Medical Sexism in Girls and Women With Suspected Bleeding Disorders in the Middle East: An Iranian Cohort and a Proposal for a Gender-Sensitive Algorithm	Shadi Tabibian
MP-394	Bleeding Profile and Health Related Quality of Life Among Females Living With Bleeding Disorders: A Single-Centre Observational Study	Rema G

MP-395	Inherited Bleeding Disorders in Women: A 15 Year Cohort Review From a Single Comprehensive Center in Australia	Kara Cordiner
MP-396	Females Affected by Hemophilia A; Data From the U.S. 8CHECK Program	Barbara Konkle
PP-397	Perceived Health Status in Women with Haemophilia and Carriers	Carla Daffunchio
PP-398	Improving Diagnosis and Treatment of Women With VWD in Pakistan: A Two Phase Analysis Using WBDR Data and Shared Decision Making Tools	Tahira Zafar
PP-399	Advancing Global Data Collection for Women with Bleeding Disorders: Early Insights from the PROBE WBD Module	Dawn Rotellini
PP-400	Unraveling Pathogenic Variants in the F8 Gene in Females With Mild Hemophilia A and Asymptomatic Carriers: Detecting Compound and Double Heterozygous	Yasser Vega
PP-401	Hemostatic Disorders in Adolescents with Severe Menorrhagia: Results of a Stepped Diagnostic-Therapeutic Approach	Yaima Mendez Pérez
PP-403	Cross-Cultural Adaptation and Reliability of the Self-Administered ISTH Bleeding Assessment Tool	Sowmya Nayak
PP-404	The Experiences of Women Surrounding Hemophilia (WSH) in Japan: Carriers, Caregivers, and the Family Context	Mikiko Ito
MP-405	Bleeding Profiles and ISTH-BAT Assessment in Women and Girls With Hemophilia: Insight From an Indonesian Referral Center	Fitri Primacakti
PP-409	Opportunities at Birth: Postpartum Hemorrhage as an Indicator for Early Hemophilia Screening in Low- and Middle-Income Countries	Maureen Mumbua
PP-410	Becoming Stronger in the Bleed: A Phenomenological Case of a Young Pakistani Nurse Living With Severe Von Willebrand Disease Type 3	Wajiha Javaid
PO-413	Gendered Marital Challenges for Girls with von Willebrand Disease in Pakistan	Wajiha Javaid
PO-414	Adolescent Carrier Status Awareness Following Disclosure by a Hemophilia Specialist: A Case Study of a Daughter of a Hemophilia Patient in Japan	Mikiko Ito