The overwhelming majority of people living with inherited bleeding disorders around the world still do not have access to diagnosis, treatment and care.

World Federation of Hemophilia (WFH) is a non-profit organization dedicated to improving and sustaining care for people with inherited bleeding disorders around the world.

We work in partnership with healthcare providers (HCPs), governments, and our global network of national member organizations (NMOs) in 147 countries.
OUR ROADMAP TO 2025

Our vision of **Treatment for All** is for a world where all people with inherited bleeding disorders have access to care, regardless of their type of bleeding disorder, gender, or where they live.

**TO ACHIEVE OUR VISION...**

**WE WILL WORK TOWARDS THESE GOALS...**

More people with inherited bleeding disorders can reliably access safe treatment and care

NMOs and HCPs are equipped and working together effectively to support people living with bleeding disorders

Collaboration is strengthened to better serve the interests of people living with bleeding disorders

**BY FOCUSING ON BUILDING CAPACITY IN THESE AREAS...**

Identify and diagnose people living with bleeding disorders

Provide adequate care and treatment

Collect, interpret, and disseminate data

Advocate for safe and sustainable treatment and care

**INVESTING IN THESE KEY TOOLS...**

WFH treatment guidelines – resources to guide best practices in treatment and care

WFH data collection programs – building capacity and providing infrastructure

WFH Humanitarian Aid Program – to provide predictable and sustainable treatment for those who need it most

**AND PRIORITIZING THESE ACTIVITIES ...**

Train, educate, and fund

Create space for collaboration and exchange

**Influence global health policies to promote access**

Promote standards of excellence

**More people with inherited bleeding disorders can reliably access safe treatment and care**

**NMOs and HCPs are equipped and working together effectively to support people living with bleeding disorders**

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**Identify and diagnose people living with bleeding disorders**

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**WFH treatment guidelines – resources to guide best practices in treatment and care**

**WFH data collection programs – building capacity and providing infrastructure**

**WFH Humanitarian Aid Program – to provide predictable and sustainable treatment for those who need it most**

**Train, educate, and fund**

**Create space for collaboration and exchange**

**Influence global health policies to promote access**

**Promote standards of excellence**
HOW WE’LL KNOW WE ARE MAKING PROGRESS

We will monitor and evaluate our global impact in four key areas. Over the next 5 years, we are aiming to:

- **Identify and Diagnose People Living with Bleeding Disorders**
  - 25% increase in hemophilia diagnoses
    - 2020: 240,000
    - Goal: 300,000
  - 14% increase in VWD diagnoses
    - 2020: 87,500
    - Goal: 100,000

- **Provide Adequate Care and Treatment**
  - 25% increase in number of people <18 yo with severe hemophilia receiving prophylaxis
    - 2020: 14,000
    - Goal: 17,500
  - 20 additional national member organizations reporting complete data in the Annual Global Survey

- **Collect, Interpret, & Disseminate Data**
  - 40 additional hemophilia treatment centres participating in the World Bleeding Disorders Registry

- **Advocate for Safe and Sustainable Treatment and Care**
  - 30 countries demonstrating an increase in government support
  - 20 countries producing or updating national treatment guidelines and policies based on WFH guidelines

**OUR 2021-2025 STRATEGIC PRIORITIES**

Our roadmap shows all that we do and why. To deliver on these aims, we will focus our attention on these priorities over the next five years:

1. Redesign our training and education offerings to maximize impact
2. Set NMO standards of excellence and support their adoption
3. Strengthen our global advocacy work to improve access to treatment and care
4. Increase the percentage of people identified and diagnosed with bleeding disorders through innovation
5. Ensure we have the resources, both financial and human, to help us achieve our goals.

**MAKING AN IMPACT, TOGETHER**

We have charted the way forward, but we cannot do it alone. We hope you will be inspired to join us. Together, we can improve the lives of people living with inherited bleeding disorders around the world.