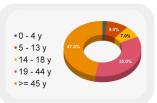
THE HAEMOPHILIA A CARE PATHWAY

in Belgium

April 2025

The overall objective of the white paper is to map the needs and aspirations of patients affected by haemophilia A, in collaboration with key stakeholders in the healthcare ecosystem.





Haemophilia A patients in Belgium DIAGNOSIS

Familial diagnosis or typically at 6-12 months of age when characteristic bleeding symptoms start

REFERENCE CENTRA

Haemophilia A care pathway has been created in collaboration with Belgium's reference centra



Hospital and local pharmacist



Patient

Specialist and general practitioner

> Physiotherapist, osteopath, revalidation specialist and orthopaedist



Patient association

My doctor reminded me that I need to find the right balance between living normally, participating in sports and keeping in mind that my haemophilia isn't cured-it's just silent most of the time.

Patients hope for a life with a haemophilia-free mindset through (ultra) long-acting treatments that sustain higher coagulation activity

Support haemophilia A patients

FUTURE OUTLOOK

CALL TO ACTION



Continue patient education to increase awareness and understanding of haemophilia A



Expand physical, psychological and social support services to address health needs



Encourage the development of new treatments



Establish integrated care coordination between the hospital and home-based care



- ☐ Increased coagulation level and longer bleeding protection
 - ☐ Support by local pharmacists and physiotherapists
 - ☐ Protocols at emergency centres

DAILY LIFE

Wandering joint pain leading to mobility challenges

Needle phobia

Psychological burden



REQUESTS

To improve patient quality-of-life. On a daily basis:

- ☐ Individualised physical activity schemes
- ☐ List of reference centres abroad
- ☐ Easily, understandable information/education
- ☐ Dedicated transport services
- □ Online patient portals
- ☐ Haemophilia health diaries

