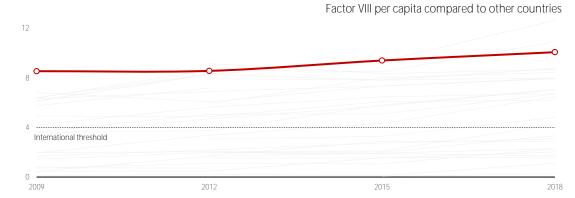


Sweden

This country fact sheet accompanies the EHC heat map and gives you further information on the country's progress over the various EHC surveys.

Here, you will see how the FVIII international units provision has evolved and compares to other countries. You will also find the same parameters as in the heat map.



Change since previous survey: ▲ Improvement ▼ Decline	2009		2012		2015		2018
International units Factor VIII	8.54	A	8.56	A	9.40	A	10.07
per capita Factor IX	1.48	•	0.34	•	0.00	A	2.03
Organisation Comprehensive Care Centres (CCC's)	Yes		Yes		Yes		Yes
of care Haemophilia Treatment Centres	No		No		No		No
National Haemophilia Council or Co-ordinating Group	No		No		No	A	Yes
Number of groups in decision-making on haemophilia care	3	•	1	A	3	•	2
Number of groups choosing haemophilia treatment products	3	•	1	A	3	•	2
National Tender for procurement of factor concentrates	No		No		No		No
Treatment Home Treatment	Yes		Yes		Yes		Yes
regimens % of people with haemophilia using home treatment	76-100%		76-100%		76-100%		76-100%
Treatment delivered to the patient's home	Unknown		Some		Some		Some
Prophylaxis treatment availability	Yes		Yes		Yes		Yes
Children currently on prophy (%)	76-100%		76-100%		76-100%		76-100%
Adults currently on prophy (%)	76-100%		76-100%		76-100%		76-100%
Access to ITI (% of people with inhibitors)	100%	•	76-99%	A	100%		100%
Access to Emergency medicine and acute surgery	Yes		Yes		Yes		Yes
specialist services Paediatrics	Yes		Yes		Yes		Yes
Infectious disease specialists (especially HIV)	Yes		Yes	•	Sometimes	A	Yes
Hepatology	Yes		Yes		Yes		Yes
Rheumatology	Yes		Yes		Yes	•	Sometimes
Orthopaedics	Yes		Yes		Yes		Yes
Physiotherapy	Yes		Yes		Yes		Yes
Dentistry	Yes		Yes		Yes		Yes
Obstetrics and Gynaecology	Yes		Yes		Yes		Yes
Genetics	Sometimes	A	Yes		Yes		Yes
Social and psychological support	Sometimes	A	Yes		Yes		Yes
Pain management	Yes		Yes		Yes		Yes
General surgery	Yes		Yes		Yes		Yes
Urology	Sometimes	A	Yes		Yes		Yes
Share of Expected Haemophilia A	88%	•	86%	A	88%	•	65%
Bleeding Disorder Haemophilia B	99%	•	69%	A	70%	•	57%
Prevalence von Willebrand Disease	16%	A	19%	•	15%	•	2%
Haemophilia Pasma-derived factor concentrate	Rarely		Rarely		Rarely		Rarely
replacement Recombinant factor concentrate	Always		Always		Always		Always
therapy Plasma		•	Rarely		D l	A	Never
Cryoprecipitate	Never	•	Kalely		Rarely	_	INCVCI
VWD replacement Plasma-derived factor concentrate	Never Never	V	Never		Never		Never
VVID replacement riasina derived lactor concentrate		•	,		,	•	
therapy DDAVP	Never	▼	Never	•	Never		Never
	Never Always		Never Always	A	Never Always	A	Never Always