

Together Ahead

Waldenström Finland

WMF - Patient Association:

A national patient organization founded and led by patients. The association brings patients with Waldenström's disease or similar diseases, their relatives and care givers to work together to improve their physical, mental, and social lives. The association aims to raise awareness of Waldenström's disease, safeguard therapeutic and social equality, and improve communication flow.

www.wmfin.fi

More information including detailed contact information can be found on our website.

e-mail: info@wmfin.fi



WALDENSTRÖM FINLAND RY

TOGETHER WE CAN COPE

with Waldenström's macroglobulinemia, or Waldenström disease (WM)

- A rare disease named after its discoverer, Swedish doctor Jan Waldenström (1944)
- Plasmacell lymph node cancer
- In the Cancer Registry since the 50s
- In Finland 678 known cases, 290 women and 397 men
- New diagnoses approx. 15 -35 cases annually
- In Finland 196 patients in 2020
- Treatments have developed tremendously
- Our association was established in 2019
- Online meetings every month
- International network and fact-finding
- Peer support and own Closed Facebook-group

Get in touch & come along!

Chairman Juha Wirekoski +358 50 4316628
Vice Chairman Gerd-Marie Backlund +358 400 850420
Secretary Kaarina Luiro +358 400 916061

www.wmfin.fi