



# Together Ahead!

## Waldenström Finland

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### 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](http://www.wmfin.fi)

More information including detailed contact information can be found on our website.  
e-mail: [info@wmfin.fi](mailto: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 902 known cases, 382 women and 520 men.
- New diagnoses approx. 15 -35 cases annually.
- In Finland 294 patients in 2023.
- 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!

