

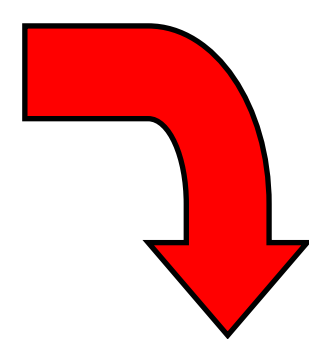
# Together, understanding and living with NMP

For a better quality of life and more effective care



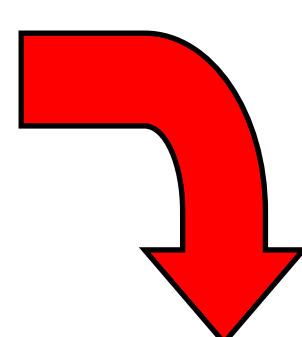
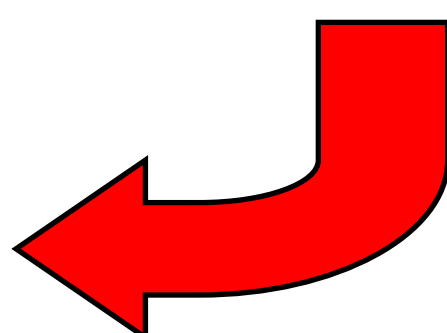
May 2021

Vaquez polycythaemia diagnosis



Discovering the disease via Google and social networks

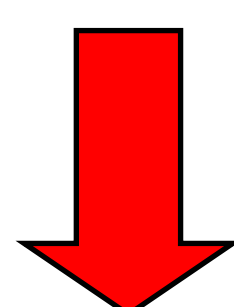
Meeting with other patient associations



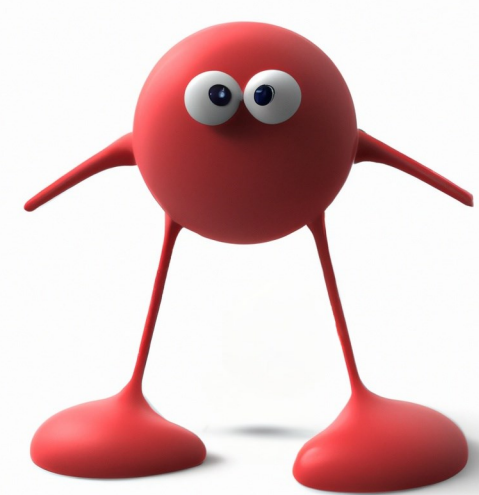
Foundation of NMP Belgium



- Patients, relatives and carers
- General practitioners and specialists
- Pharmaceutical companies
- Public authorities



INFORMATION  
PREVENTION  
COMMUNICATION  
SUPPORT  
LOBBYING



Globulus, our mascot, supports patients and their families



## WHO ?

Cécile Belvaux, Christophe Vanden Broeck (co-founders)

## WHY ?

There wasn't any patient's association for myeloproliferative disorders in French-speaking Belgium. Information, guidance and support were missing .

## WHAT ?

- 1 Website and social networks creation
- 2 Adriano Salaroli join the board as Scientific Advisor
- 3 Information materials for patients, GPs and specialists creation
- 4 Goodies to raise awareness
- 5 Collaboration with various patient's associations and haematology departments

## NEXT STEPS...

- To organise events (theatre piece, symposiums, etc.)
- To gain new members
- To develop sponsorship
- To send a monthly newsletter

## SOME FACTS...

- 🌀 Since our birth, we already reach 130 members...
- 🌀 We take part in various events such as the PEC Expert Patient programme, the BHS annual conference, the WeCan Academy, Septembre rouge, etc.
- 🌀 We work with leading hospitals (Bordet, Brugmann...)
- 🌀 We have joined umbrella associations such as LUSS (association of associations), RadiOrg...

## NMP Belgique

NMP patient association in French-speaking Belgium