

# Pan European Patient Organisation Funding

## Myeloma Patients Europe (MPE)

MPE is an umbrella organisation of multiple myeloma patient groups and associations from across Europe. MPE was formed in 2011 following the merger of the European Myeloma Platform and Myeloma Euronet and is registered as a non-profit organisation (AISBL) under Belgium law. The vision of MPE is to achieve a network of highly efficient, effective and sustainable myeloma patient organisations across Europe. The mission of MPE is to provide education, information and support to member groups and to advocate at European, national and local levels for the best possible research and equal access to the best possible treatment and care.

We have supported this organisation since 2019.

### During 2021:

- Core Programs (Education and Information & Access Atlas) – €81,956
- COVID Programs – €7,030
- Clinical Trial Finder Tool – €22,200

Our support represented 9.52% of their overall income

### During 2020:

- GSK provided
  - € 40,000.00 for MPE Access Program
  - € 25,000.00 for MPE Information Resources
  - € 36,000.00 for Community Advisory Boards (CAB)
  - € 10,000.00 for COVID Support Programs

Our support represented 11.1% of their overall income

### During 2019:

- GSK provided €30,000 for:
  - Supporting the MPE Access Atlas Programme, a platform developed to help build the knowledge and capacity of members through providing the information needed to advocate effectively on national issues
  - Educational Activities

Our support represented ~16% of their overall income

