

Patient Organization Support Payments made between

1st January 2024- 31st December 2024

| Organization | Description / NAF Summary Amount | Amount in GBP |
|---|--|---------------|
| Association for Glycogen Storage Disease (AGSD) | Funding for 2024 activities | 15,000 |
| Beacon: For Rare Diseases | Support for the 'Building Rare Leaders' programme | 5,000 |
| Genetic Alliance UK | Funding for the 'Seeking a Rare Diagnosis' project | 5,000 |
| Genetic Alliance UK | Corporate partnership | 15,000 |
| Genetic Alliance UK | Funding towards Rare Disease Day campaign | 12,800 |
| Kidneys for life | Funding support for 8th Fabry Update conference | 75,000 |
| Medics for Rare Disease | Corporate partnership | 15,000 |
| Medics For Rare Disease | Fee for service for advice | 2,844 |
| Metabolic Support UK | Sponsorship for conference | 10,000 |
| Metabolic Support UK | Funding towards patient education activities | 10,000 |
| MPS Society UK | Corporate roundtable membership | 16,500 |
| MPS Society UK | Fee for service for advice | 970 |
| MPS Society UK | Funding toward advocacy and support services | 20,000 |
| MPS Society UK | Fee for service for speaker | 1,030.30 |
| Pompe Support Network UK | Funding for 2024 activities | 15,000 |
| Pompe Support Network UK | Funding towards disease management guide | 23,000 |
| Pompe Support Network UK | Medical Advisory Board | 472 |
| RDRP (a wholly owned not-for-profit subsidiary of MPS Society UK) | Sponsorship of conference | 23,500 |
| Rareminds | Fee for service for advice | 937.50 |
| Rareminds | Funding for 2024 activities | 5,000 |