

**REGISTERED COMPANY NUMBER: 03253584 (England and Wales)**  
**REGISTERED CHARITY NUMBER: 1058968**

THE PITUITARY FOUNDATION  
(A COMPANY LIMITED BY GUARANTEE)

TRUSTEES' REPORT AND  
UNAUDITED FINANCIAL STATEMENTS  
FOR THE YEAR ENDED 30 JUNE 2025

THE PITUITARY FOUNDATION

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FOR THE YEAR ENDED 30 JUNE 2025

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THE PITUITARY FOUNDATION

REFERENCE AND ADMINISTRATIVE DETAILS  
FOR THE YEAR ENDED 30 JUNE 2025

TRUSTEES	Mr T Sumpster Chair Mr A M Mackintosh Vice Chair Dr I E Macdonald Hon Secretary (resigned 20.5.25) Mr P Rouse Hon Treasurer Dr S E Baldeweg (resigned 20.5.25) Dr A M Brooke Ms D L Cooper Ms A Deleigne Mr A H Matty (resigned 2.10.24) Ms J A Phillips Mrs C E Thatcher (resigned 20.5.25) Ms N Chowdhury (appointed 2.6.25)
COMPANY SECRETARY	Dr I E Macdonald
REGISTERED OFFICE	Brunswick Court Brunswick Square Bristol BS2 8PE
REGISTERED COMPANY NUMBER	03253584 (England and Wales)
REGISTERED CHARITY NUMBER	1058968
SOLICITORS	TLT LLP One Redcliffe Square Bristol BS1 6TP
BANKERS	National Westminster Bank 32 Corn Street Bristol BS1 1IQ  Royal Bank of Scotland 36-38 Baldwin Street Bristol BN1 1NR

**THE PITUITARY FOUNDATION**  
**TRUSTEES' REPORT**  
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The Trustees, who are also directors of the charity for the purposes of the Companies Act 2006, present their report with the financial statements of the charity for the year ended 30 June 2025. The Trustees have adopted the provisions of Accounting and Reporting by Charities: Statement of Recommended Practice applicable to charities preparing their accounts in accordance with the Financial Reporting Standard applicable in the UK and Republic of Ireland (FRS 102) (effective 1 January 2019).

## Objectives and Activities

### **a. The purposes of the charity as set out in our governing document**

The Charity's objects, as described in our Memorandum and Articles of Association, are to promote the relief and treatment of persons suffering from pituitary disorders and related conditions and diseases, and their families, friends and carers, and to provide information and support; and to promote and support research and to disseminate for the public benefit the results of any such research.

The main activities that the Foundation undertakes to deliver on these services are:

- the delivery of general and specialised helplines;
- development and dissemination of resources online and in print;
- presentation of information and holding of community building events online and in person;
- facilitation of patient-patient support services;
- engagement in, and the dissemination of, research;
- support of volunteers; and
- undertaking of fundraising schemes including events.

### **b. Our strategy and core aims**

Our current strategy was developed in 2023 and covers the four years from 2023 to 2027. It states that:

**Our vision** is that all people with pituitary conditions can live well, now and in the future.

**Our mission** (the way in which we achieve our vision) is that:

We work with everyone with a pituitary condition, their support networks and healthcare professionals to raise pituitary awareness, and to reduce time to diagnosis.

We empower patients to navigate the UK healthcare system and obtain the best clinical outcomes possible and provide support to enable everyone with a pituitary condition to live as well as possible.

**Our values** (which inform how we approach achieving our vision) are:

We are welcoming  
We are caring  
We are empowering  
We are trustworthy  
We are collaborative

**Our Objectives** (which break down the practical steps we will take to achieve our vision) are:

#### **1: We will work to reduce the time it takes to be diagnosed with a pituitary condition by increasing awareness**

We have three focus areas to achieve this objective:

- 1.1 Raise awareness amongst Endocrinology specialists
- 1.2 Raise awareness in primary care
- 1.3 Raise the general public's awareness

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**2: We will empower every patient to receive the best care available to them, now and in the future**

We have three areas that we focus on to achieve this objective:

- 2.1 Offer support and education to everyone impacted by pituitary conditions
- 2.2 Advocate for the best medical care possible in the NHS for everyone with pituitary conditions
- 2.3 Champion valuable patient centred investment in the future of pituitary care

**3: We will support all people with pituitary conditions to live well**

We have three areas that we focus on to achieve this objective:

- 3.1 Increase the accessibility and inclusivity of support and services
- 3.2 Provide lifestyle support to people with pituitary conditions
- 3.3 Seek to ensure Psychological Support and broader wellbeing advice is available to everyone impacted by a pituitary condition

**4: We will ensure The Foundation is robust and sustainable**

We have three areas that we focus on to achieve this objective:

- 4.1 Ensure we have the right resources to deliver our strategy
- 4.2 Ensure our financial resources match our ambition
- 4.3 Work sustainably

## Achievements and performance

### a. Overview of activities throughout the year

2024 was a very special year for The Foundation as we celebrated our 30<sup>th</sup> anniversary. It was a fantastic opportunity for us to applaud our community and all the contributions over the decades, and to raise awareness of our activities. As part of our year of celebration we invited nominations for 'pituitary heroes' – people who have made an incredible contribution to our community across support services, clinical care, research, and fundraising. We received over 70 nominations from which 30 amazing individuals were selected. Throughout the year we shone a spotlight on our heroes through our website and social media, culminating in a charity reception held at University College London Hospitals in October 2025.

Volunteers were instrumental in founding the charity 30 years ago and continue to be at the heart of our work. We were pleased to develop our support for volunteers during the year with staff resource enabling us to build relationships, review and update our support materials and services for volunteers, deliver training, and (at the time of writing) start planning for an in-person volunteer weekend. During the year we saw support groups thrive with new groups starting up (in person and online), our volunteer-run helpline open every week day, and our Lived Experience Committee established. We are incredibly grateful to everyone who volunteers with The Foundation. They freely give their time to support the community we are here for, with kindness and understanding.

Our information and support services continue to be highly valued and are the cornerstones of The Foundation's offer for the pituitary community. During the year we saw the effective delivery of our helplines and volunteer run support groups which provide vital local support. We delivered successful in-person 'get together' events to bring the community together in Manchester and Leicester. We also delivered an ambitious programme of online events led by experts in pituitary conditions which were well attended and with high viewing numbers of the recordings.

We have been pleased by the progress we've made in developing our communications and health information. As reported here we have an effective website with continuously updated content, a really strong social media presence, and an engaged community online. We were also delighted to start a project on health information which will see us review, update, and develop all of our health information.

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Providing trusted information about pituitary conditions is a vital part of our role, especially in a changing digital context, so we are pleased to be investing in this area.

Underpinning all our work is our need for income. The financial year we are reporting on saw a shortfall in our income against our costs in the year. We have assessed this within our Finance Committee and with the CEO and Head of Fundraising: a key area of underperformance was income from challenge events, so we have therefore put in place robust plans and support for our fundraisers, and are confident that 2025/26 will deliver the results we are aiming for. As always, we are incredibly grateful to all our supporters for their generosity which enables us to deliver all our services and support.

We started the financial year with many of the elements in place for providing successful delivery: a new CEO and full staff team at the start of 2025, clear strategic framework (2023-2027), excellent infrastructure, an overall sound financial base, and a supportive community. Our annual report 2024-2025 describes a year with many highlights, areas of development, and the effective delivery of our core services.

**b. Delivery against the strategy**

In this section we will look at specific activity relating to each objective, and areas we are planning to develop. We measure our success against qualitative and quantitative measures. For projects, we set objectives at the beginning, whilst for ongoing work including our advocacy we review and respond to feedback from our community. Our Lived Experience Committee assists in reviewing and guiding our work and compliments our Trustees who interrogate the value and impact of our services.

**1: We will work to reduce the time it takes to be diagnosed with a pituitary condition by increasing awareness**

This objective is focused on reducing the time it takes to be diagnosed through increased awareness of pituitary conditions. During the year we achieved the following:

**1 (a) Awareness within Endocrinology**

We are an affiliated patient support group with the Society for Endocrinology and as such contribute to and benefit from being part of a network of specialist organisations liaising directly with the clinical endocrine community.

**Conferences**

Attendance at conferences is an important way in which we spread awareness of the charity and its work amongst clinicians across the country and beyond. This year we attended the Society for Endocrinology's British Endocrinology Society conference in Harrogate in November 2025. We also attended the British Society for Paediatric Endocrinology in Glasgow in October 2025. In October we attended the WAPO world congress (World Alliance of Pituitary Organisations) in Dublin which was a valuable opportunity to connect with patient advocates and patient support organisations from across the globe.

We estimate that we reached more than 4,000 health care professionals through these various conferences, and have been able to talk about our services and share resources with them in the process.

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**Medical Committee**

Our Medical Committee has consistently contributed to our work including ensuring we are connected across the clinical endocrine community. We are incredibly grateful to all members who freely give their time to support our work. A review of membership has led to additional specialist areas being represented, which is important given the complexity of pituitary conditions. At the time of writing there are 23 members of the Medical Committee, including:

- 8 Endocrinologists
- 4 Endocrine nurses
- 1 Paediatric endocrinologist
- 2 Pituitary surgeons
- 1 Pituitary surgery nurse
- 1 Consultant endocrine nurse
- 2 Endocrine pharmacists
- 1 Psychologist
- 1 GP
- 1 Orthoptist
- 1 Radiologist

**1 (b) Awareness in Primary Care**

**Emergency Care**

During the year we collaborated with the Addison's Disease Self Help Group (ADSHG) in our response to the move away from 'red flagging' the home addresses of people with adrenal insufficiency by the AACE (Association of Ambulance Chief Executives). We recognized the importance of sharing information about this shift with our community, the purpose of summary care records (SCR), and other preparations that patients can make in case of an emergency. As part of this collaboration and response we prepared a template letter for patients to download to notify their GP practice to update their SCR.

We continue to provide the steroid emergency card, either by post or print at home versions in our member magazine Pituitary Life. During the year we sent out 455 printed cards and people were able to download from our website, with about 1,700 made available through our Pituitary Life magazine.

**GP Awareness**

Raising awareness of pituitary conditions with GPs is an ongoing priority for us and we are always looking for relevant opportunities to develop this. As well as the patient letter on SCRs described above, we also prepared a similar template letter for patients with AVP deficiency. Again, the aim here was to increase GP awareness of specific pituitary conditions as well as to enable patients to self-advocate and request that their records are accurate in case of emergency.

A new project on prescription length for 'vital medications' was in progress at the time of writing. We became aware of concerns that vital medications were being issued on 28-day prescriptions which presents several issues for our community. Once again, we were delighted to collaborate with Addison's Disease Self Help Group. We carried out a short survey to increase our understanding of the issue and were grateful to the almost 1,400 people who shared their experience with us. At the time of writing, we are working with members of our Medical Committee on next steps.

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**1 (c) Raise the general public's awareness**

**Awareness Month**

Every year in October we run an awareness campaign to mark 'Pituitary Awareness Month'. It is an opportunity for us to set a theme and design activity which delivers against our awareness-raising objectives. In October 2024 our focus was 'back to basics' with events and information to support raising awareness of the importance of the pituitary gland, its life sustaining role, and pituitary conditions.

During the awareness month we held two online events focusing on the basics of pituitary conditions. We also held three in person events including a 'get together' in Manchester, a charity reception to celebrate our Pituitary Heroes in London, and an awareness walk in Bristol.

Across Awareness Month, we saw strong engagement statistics across our social media profiles. The average reach of our Instagram page almost doubled on the previous month. Our audience was particularly interested in content explaining the pituitary gland, and the causes of different pituitary conditions.

**Website**

Our website is one of the main ways for people to access our information and support. Following the launch of our new website in 2023 we have continued to improve its content and visual presentation. Overall, we've seen a 3.3% increase in the number of active users on the site from the previous year. Hopefully this signifies that the website is starting to regain traffic after the previous year's dip, following the launch of the new site.

During the year we published 50 blog posts on our website, including research opportunities, charity updates, expert articles and community stories. Our most popular posts were those about the desmopressin and hydrocortisone medication shortages, highlighting how valuable these updates are for our community.

Overall, our top performing pages are our information pages, specifically our 'Cushing's disease', 'What are hormones', 'What is the pituitary gland', 'Hydrocortisone Sick Day Rules' and 'Adrenal Insufficiency' pages. This suggests that our website is functioning well as an educational resource and supports those seeking out specific condition information and those wanting more general pituitary-related content.

This year, we collaborated with Rareminds (a specialist mental health services provider) to introduce our new Psychological Support Hub; a dedicated page on our website to house all our wellbeing resources and services. Since launching, we've had nearly 800 visitors to the Hub. Many visitors have returned to the Hub multiple times, suggesting that the content is relevant and valuable to them.

**Social Media**

Our social media profiles have continued to see strong growth across this year. In February 2025, we ran a successful Rare Disease Day social media campaign that spotlighted inspiring stories from our community members. The campaign connected well with our audience and led to the highest engagement rate in 6 months, across our Instagram, Facebook and LinkedIn profiles.

Our combined social media following has now surpassed 30,000, which is a huge achievement for a charity of our size. We intend to continue using social media as a tool to empower and inform our audiences and raise awareness of pituitary conditions and our work.

**Press Coverage**

In January, we were pleased to provide expert comments on Cushing's disease for a Telegraph article about the condition. Since then, we've been mentioned in a number of press articles covering Cushing's disease, AVP deficiency and Kallman's syndrome.



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## **Newsletters**

Our monthly members' and general newsletters provide regular touch-points for our community. Combined, our mailing list has grown to over 9,700 subscribers. Our newsletters continue to drive strong engagement with other areas of our work, including promoting attendance at our events, recruiting for volunteer positions, and signposting to our online resources.

## **2: We will empower every patient to receive the best care available to them, now and in the future**

Our second objective focuses on empowering all patients to receive the best care available to them. We play an instrumental role in this by providing support and education to complement that provided by the NHS.

### **2 (a). Offering Support and Education**

#### **Helplines**

Our helplines remain a vital 1-1 support offering. Our objective is to provide information and support in ways that are accessible to our community. Individuals from our community can reach us via our nurse-run helpline, our volunteer-run information and support helpline, and via email.

During the year we responded to over 1,100 enquiries. This was a reduction of 450 based on the previous year. Our analysis shows this was mostly on our volunteer helpline and email helpline. During the year we had gaps on our volunteer helpline which limited the number of hours that it was available. We've focused on recruiting new volunteers and at the time of writing are delighted that the volunteer run helpline is open for more hours through the week.

We are always looking at ways we can develop our helpline service, taking account of the feedback that we receive. We also monitor the helpline service for any common themes that are emerging that we may need to respond to. For example, we were able to identify the issues and concerns about desmopressin spray medication supply which we mentioned elsewhere in this report.

We'd like to take this opportunity to particularly thank all volunteers who deliver the information and support helpline. With 81% of callers rating the service 'extremely valuable' it is clear just how valuable this is to our community, and a reflection of the quality of support that is provided. Some quotes from the year:

*"Still new but very impressed with the information, services and support. thank you"*

*"Satisfied with everything the helpline is providing me with. Thank you."*

*"Keep doing what you're doing! Thank you!"*

#### **Health information resources**

We are proud of having achieved PifTick accreditation from the Patient Information Forum in 2023, and we underwent successful reassessment in 2024. This is a quality mark that reflects that we have applied good practice principles to the preparation of our health information content. We understand the importance of providing trusted information about pituitary conditions and see this as a core part of our service. During the year we were delighted to publish six updated health information booklets for our community which are PifTick accredited:

- Prolactinoma
- Acromegaly
- AVP deficiency (previously known as Diabetes Insipidus)
- Adrenal Insufficiency
- Cushing's
- Surgery and Radiotherapy

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During the latter part of the year, Trustees approved a 'health information project' which will see us carry out a full audit of all our health information resources, start updating existing resources, and develop new resources which meet the needs of our community. As part of this we will work closely with our Lived Experience Committee. We are extremely grateful to our Medical Committee for supporting us with our health information resources.

Also during the year we were delighted to work with SIMBA CoMICS on a new animation on Cushing's disease, including symptoms. We engaged 'patient reviewers' to work with the content creation team, which resulted in an information animation which is now on our website and on YouTube. We would like to develop more health information in different formats such as this animation, and we see it as an important way to make information accessible.

### **Online Events**

Our online events, which started as a test a couple of years ago, have become a core part of our service offer. During the year, we planned and delivered an ambitious programme which represented a doubling of the number of events held in the previous year. They covered many topics, including on specific conditions, employment rights, and nutrition. Our events are delivered 'live' and are recorded for posting on our YouTube channel later.

We are absolutely delighted that over 1,000 people registered for our events. Over 95% of attendees to our online events rated them excellent or good. The main reason why the majority attended was to gain information (92% of attendees), with the second most popular reason being to support The Foundation (34%). Most attendees find out about our online events on social media (41%), with the second most popular being the monthly newsletter (39%). Some quotes:

*"Thank you, very informative. More of them please as I try to attend anything related to my condition"*

*"Thank you for organising these wonderful events. You really do touch lives and help improve people's situations. Thank you"*

*"Really interesting, gave me some insights to my condition, even though I've had it for over 30 years now"*

*"This was one of the best webinars I've attended. I came away from it feeling so much better and less alone. Thank you so, so much."*

We continue to develop this part of our service which is only possible due the generosity of the experts who volunteer their time including members of our Medical Committee.

### **Get Togethers**

Our Get Together events, introduced after the pandemic, have become a regular and valued fixture of our annual calendar. This year we began involving volunteers in the planning and delivery of the Get Together events, and this has been incredibly valuable. During the year we brought our community together in Manchester and Leicester. These events were attended by over 90 people. Once again we were grateful to members of our Medical Committee for their contributions as well as volunteers and Pituitary Ambassadors.

In Manchester, 100% of attendees rated the day as good or excellent, with most attendees looking to gain information, or meet others. This was the first Get Together featuring a patient story presentation, and 100% of attendees found this to be excellent. Here is some of the feedback received:

*"I really enjoyed the day, and I enjoyed talking to new people and hearing their stories. All the information was well presented and extremely useful. Thank you."*

*"An excellent event. Well-structured and very informative. Thank you to all involved in helping to make the day possible. It was so very much appreciated. It's always good to pick up tips from other people's journeys. Thank you"*

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*"A fantastic day. Very well organised. It has given me more strength being among other patients and also learning lots from the speakers. I hope many more of these events are put on around the country helping lots of other people too."*

Over 80% of attendees in Leicester found the day to be excellent or good, with the majority of attendees wanting to improve their understanding of the topics by attending. We had some very positive feedback, including:

*"A most enjoyable day, thank you."*

*"Thank you, the conference ran smoothly with a good opportunity to exchange stories"*

*"I thought [staff] were GREAT ADVOCATES for the foundation, so welcoming, professional and clear in how the day would be. WELL DONE to you both, excellent organisation and on the day attentive. "*

### **Pituitary Life magazine**

Our printed magazine, Pituitary Life, is produced three times a year and was published in Autumn 2024, Spring 2025, and Summer 2025. It is a valued part of our membership offer and is delivered to over 1,700 members. It is also available in a digital format. We are grateful to everyone who contributes content including medical experts and members of our community who generously share their experiences. During the year we established an editorial group to support the involvement of volunteers in the development of the magazine.

Our Summer 2025 was our 60<sup>th</sup> edition of the magazine, reason to celebrate with a special cover image. Feedback from that edition:

*"Avidly read, as always"*

*"I was delighted to see that you have a proforma letter to ask my GP to ensure that my adrenal insufficiency is recorded on my summary care record and have downloaded this"*

### **2 (b). Advocate for the best medical care possible in the NHS for everyone with pituitary conditions**

Advocacy on the issues that matter most to our community is an important part of our role as the leading pituitary charity in the UK. There were a number of areas where our community engagement resulted in advocacy activity.

#### **AVP deficiency name change**

We were part of the international working group supporting the implementation of the AVP deficiency name change from Diabetes Insipidus. We produced materials to raise awareness about the name change for our community including a new updated health information booklet.

#### **Medicines shortages**

We have been proactive in reaching out to the relevant authorities about shortages of medicines which included desmopressin nasal spray and hydrocortisone sodium phosphate injection, reflecting the concerns and challenges of our community. In relation to desmopressin nasal spray, this included surveying our community to find out about their experiences and reflecting these to the relevant bodies.

#### **NHS 10 year plan**

We submitted our response as part of the NHS's engagement on its new 10 year plan, setting out the particular challenges for people with undiagnosed and diagnosed pituitary conditions. We are keen to understand more about changes which are underway including the delegation of the commissioning of specialised services, the dissolution of NHS England, and the implementation of the NHS 10 year plan. We will be monitoring these changes and advocating for the pituitary community within the rare disease space.

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## **Sector Engagement**

We are represented on a number of working groups including: the BSPED working group for AI education standards for children and young people; DI name change working group; NHS England Policy Working group for Acromegaly; Oxytocin Study (Diabetes Insipidus Project | Patient representative); and the newly formed working group for prolactinoma treatments, launched by PROMISE Study Team, Dr Cihan Atila, MD-PhD, University Hospital Basel.

We continue to contribute to the Scottish Medical Consortium Patient Group, the Welsh Patient and Public Interest Group, the NICE Voluntary Sector and Community Sector Forum, and we are members of the ABPI Patient Organisation Forum.

Our Head of Support Services was one of the authors of 'A Core Outcome Set for Pituitary Surgery Research: An International Delphi Consensus Study' which has been accepted for publication by *Pituitary*. (June 2025)

## **2 (c). Champion valuable patient centred investment in the future of pituitary conditions**

### **Networks**

We are an active and affiliated member of the Society for Endocrinology Patient Support Group network, representing a valuable way of ensuring that the patient voice is heard at a foundational level. We are also a member of WAPO (World Alliance of Pituitary Organisation). These offer good opportunities to connect with pituitary patient support organisations in countries around the world.

### **Lived Experience Committee**

We launched our Lived Experience Committee in September 2024. This is chaired by a Trustee and is made up of people with pituitary conditions. It is an important way for us to incorporate the voice of people with pituitary conditions at the heart of The Foundation. We continue to develop the work of the Lived Experience Committee including its connection with our Medical Committee.

### **Research and consultations**

We know that our community is interested in research and we see we have a role to play in awareness of opportunities to get involved in research and to provide information about developments in the world of research. We provide news articles and lay versions of academic research where possible. We also signpost to opportunities to get involved in research where appropriate. At times, staff members will represent our community in research by providing insights based on our extensive knowledge and experience. For example, during the year we:

- contributed to a research piece by the Specialised Healthcare Alliance on the mental health impacts of rare disease;
- supported a research project on hunger in relation to Craniopharyngioma;
- members of our medical committee and a Trustee published an academic paper on AVP-D, name change and developments in testing;
- provided support to a new project on proton beam radiation therapy for people with Acromegaly and Cushing's.

These are examples of the ways in which we contribute directly, signpost and support advocates to become involved, and provide information to our community about research developments in pituitary conditions.

## **3: We will support all people with pituitary conditions to live well**

At the heart of this objective is ensuring that as many people as possible can access, engage with and benefit from our services.

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**3 (a). Increase the accessibility and inclusivity of support and services**

There is a high level of commitment across the charity to providing services and support which are relevant, inclusive, and accessible to everyone living with a pituitary condition. Our use of different communication channels and providing information and services in a variety of formats (for example in person and online) help to ensure that people can access what they need from us when they need it.

**Support Groups**

Our Support Groups have been a valued part of our service offer since the charity's inception and continue to evolve to meet the needs of our community. At the time of writing we are grateful to long standing volunteers for their running of support groups across the UK, as well as being pleased to support the start-up of new groups.

During the year four new Support Groups were started up in Sussex, Kent, Edinburgh, and Leicester. This would not have been possible without the wonderful volunteers now leading those support groups. At the time of writing, this brings the total number of active support groups to 22, both online and in-person. During the year, the staff team visited 11 of the groups.

We have successfully introduced new online, condition specific support groups to complement those that meet in person. New groups meeting during the year include a Prolactinoma group, a young person's group and an Acromegaly group.

At the start of 2025 we clarified our relationship with the Cushing's UK group and are proud that it is now officially a Pituitary Foundation support group. The group has been running successfully for 8 years, coming under the Foundation's umbrella post pandemic, and is now officially a support group. The amazing volunteers running the group currently support almost 3,000 members.

**Membership**

We are continually grateful to those in our community who support our work by becoming members. During the year we continued to put on welcome meetings to members (also non- members welcome) to demonstrate how The Foundation can support people, and the ways in which people can get involved.

At the time of writing we have 2,104 active members, including 284 individuals with 'life membership'. During the year we began sending out renewal reminders by email with the aim of reducing postage costs. We also completed moving around 120 people from the Joint membership (which is no longer available) onto our Concessionary and Individual memberships.

**3 (b). Lifestyle Support**

We recognise that many pituitary conditions are lifelong, and we want to support people throughout their life, not just at the time of diagnosis. At the time of writing we are planning for a Pituitary Awareness Month theme of 'a lifetime of hormone health' which has been designed to cover important life stages and milestones with a pituitary condition.

**3 (c). Psychological Support**

Pituitary conditions can have a significant impact on mental health, so we have continued to prioritise developing our information and services offer in psychological support. This has included the following elements:

**Counselling.** We have been delighted to continue working with specialist mental health service provider, Rareminds. This has resulted in the delivery of over 200 individual counselling sessions. At the time of writing we are also working with Rareminds to deliver group sessions led by a therapist.

**World Mental Health Awareness week (MHAW).** The focus of this year's Mental Health Awareness Week was 'community', and the importance of positive communities in fostering psychological wellbeing. To mark this, we ran a short social media campaign highlighting this focus and thanking our

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community. We also held an online event, delivered by Dr Sue Jackson, on the topic of community and mental health.

**Psychological Hub.** During MHAW, we deployed our new Psychological Support Hub, an online space on our website housing our psychological support resources and services. We hope this hub will make it easier for people to find and access our existing resources, including our videos, longer reads and Rareminds counselling.

**4: We will ensure The Foundation is robust and sustainable**

Our objectives in this area are to ensure we have the right resources (including funding) to deliver our strategy as well be sustainable.

**Governance.** The Board of Trustees approved a new trustee tenure policy in November 2024 which brings The Foundation in line with good practice. This meant that sadly three long standing Trustees retired at our Annual General Meeting in May 2025 with others planning to retire in 2025/26. We are incredibly fortunate to have had many years of committed service from our retiring Trustees and wish them all the best for the future.

We carried out a skills audit earlier in the year in preparation for trustee recruitment. At the time of writing we are in the recruitment process for Medical Trustees and planning to open up recruitment for other trustee roles in the autumn of 2025. All the thinking and work on succession planning has been done to ensure that there is a managed process to support changes to the Board of Trustees.

**Staff.** Our staff team saw some change during the year with our incoming new CEO starting in the same month as our outgoing CEO left (July 2024), a transition managed smoothly by both individuals, for which we are grateful. We also had some other team changes which we've managed with minimum impact to operations and delivery. We are fortunate to have an excellent staff team with everyone demonstrating real commitment to our work. We'd like to acknowledge and thank everyone in the staff team during 2024/25 for their contributions.

Whole staff team training during the year included data protection and DEI (diversity, equality, and inclusion). Individuals in the team have also done training on first aid, volunteer management, gift aid regulations, AI (artificial intelligence), and health information.

**Finance administration.** We continued to roll out our new finance support for local support groups which reduces administration and increases financial controls and visibility. As part of this we produced detailed guidance for volunteers.

**Fundraising.** Please see the full report that follows for more details. During the year we expanded our challenge event offering through partnership with Run4Charity, and carried out a deep dive into our management of the programme including stewardship of fundraisers.

**Policies and procedures.** Our staff handbook includes all relevant policies and is updated when required in partnership with our external HR advisors. During the year Trustees approved a financial Reserves policy, and we updated our Safeguarding policies and processes. We also developed and approved a new 'Working with healthcare companies' policy and published this on our website. This ensures we are in line with good practice guidance published during the year on disclosure of funding from pharmaceutical companies.

**Collaboration.** We continue to strive towards collaboration where it is in the interests of our community. This includes collaborating with other patient support organisations working in the endocrine space as we've reported here.

**Plans for Future Periods**

We reported in our 2023/24 Annual Report that our reserves position provides us with an opportunity to further invest in our services and the sustainability of the charity. During the year we agreed a number of areas for investment including:

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**Health information project:** providing trusted health information is a core part of our services offer. Having embarked on updating our health information, we've invested in a dedicated project to progress this work more fully. This includes a full audit of our health information, updating existing resources, and developing new information to meet the needs of our community.

**Charity sustainability:** Trustees agreed to the funding of a data protection audit which has enabled us to strengthen and develop our policies and procedures in regard to handling data. We have also committed to upgrading IT kit in anticipation of the shift to Windows 11.

**Psychological Support:** as described elsewhere in this report we have further invested in our services and information to support people with the mental health impacts of living with a pituitary condition.

There are two areas we are developing with plans to move these forward in 2025/26:

**Children and young people's project:** This project will engage with younger people in our community to understand their needs and will result in recommendations for specific activity areas.

**Raising awareness:** This piece of development work will see us increasing our awareness raising activity as well as engagement with supporters. We will also develop and deliver more campaigns on the issues that matter most to our community.

#### **Public benefit**

The Trustees have considered the Charity Commission's requirement in respect of Public Benefit. In their view the charity meets, in full, the criteria to satisfy the test. The Trustees' Annual Report describes the activities undertaken to further its charitable purposes for public benefit.

## **Financial review**

### **a. Overview**

After several years in which we successfully generated trading surpluses, in financial terms this was a more challenging year for the charity, and a reflection of the continuing economic pressures faced by everyone.

Our income for the year totalled £447,730; in the previous year we received a very large legacy of £278,000 from the Glayzer family – excluding this our income in that year was £501,565, so on a comparable basis our income this year was about 10% lower. This year we witnessed some pressure on levels of donations generally, received lower contributions from pharmaceutical companies active in the endocrine sector, and in particular saw a substantial reduction in income from Events (organised by the charity or done independently by charity supporters). We did, however, manage to raise record funds from Trusts and Foundations.

Whilst our Events income was disappointing, as ever we are grateful both to the numerous supporters who completed a broad array of challenges on our behalf, some of which are highlighted below, and to all our generous funders during the year.

Once again, our team were effective in managing our costs well, with total costs coming in at £522,778. Of this sum, £28,500 related to planned expenditure on various initiatives from reserves, so excluding that figure our underlying core costs were £494,278, an increase of only about £8,000 over the previous year.

Overall, we report a total net loss for the year of £72,355; excluding the expenditure we funded from reserves, our underlying trading deficit was £43,855.

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Our CEO and Head of Fundraising have been working in recent months to ensure that our Events programme can bounce back this year, and we are pleased to report that early indications have been encouraging.

At the year end, our accumulated reserves had reduced to £706,719, still a very healthy level for the charity, with unrestricted reserves at £589,113 and restricted reserves of £117,606. Our reserves policy, and plans for investing some of these funds, are set out later in the section.

**b. Incoming resources**

In 2024/25 The Foundation made income of £447,730 compared to £779,565 in the previous financial year.

**c. Principal Risks and uncertainties**

The charity is highly reliant on donations from individuals, Trusts and Foundations and companies; these may be affected by broad economic and other circumstances over time. We manage a database of more than two thousand members and supporters and work hard to appeal to them for their support, and we maintain strong relationships with a range of organisations many of which continue to support the charity year on year.

The Foundation receives some of its income from legacies; by its nature this income stream is unpredictable, and therefore the charity seeks to limit its reliance on this element of income, budgeting for a modest amount of legacy income.

**d. Funders**

The Foundation enjoys an extraordinary level of support from its community. The Trustees express their huge gratitude to all our donors: members, donors, fundraisers, companies and Trusts and Foundations. Without this support we simply could not continue. Thank you.

We are fortunate to have a huge number of supporters, many of whom wish to remain anonymous. However, we would like to recognize some of the generosity we have benefited from during this financial year – the next section does not give a complete picture, but hopefully provides a flavour of the support we enjoy.

**e. Events and community fundraising**

As ever, many of our fundraisers went to extreme lengths to raise money for the charity; their bravery, commitment and determination are humbling and we are so grateful to each and every one of them. Whilst we had a more difficult year in terms of Events fundraising overall, there were still many fantastic fundraising endeavours, including:

- A team of 6 runners, including one of our Endocrine Nurses, took on the Barcelona Marathon in March helping to raise well in excess of £6,000
- We returned to the home of Liverpool Football Club for an 'Anfield Abseil' which saw 15 brave individuals (including our Head of Support Services), scale the heights of the stadium; collectively the team raised more than £5,000
- A team of 8 trekked up Yr Wyddfa (Snowdon) for us in June raising £3,000; thanks as always to Bryn Williams Mountaineering for looking after our team
- The picturesque Brecon Beacons was the setting for our Four Falls trek in May in which we were joined by 30 trekkers including members of our staff team and Medical Committee, collectively raising £6,500. We'd like to thank the team at JT Expeditions for once again guiding our team



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We are so grateful to all the individuals who ran challenges for us in the year, which included the London Marathon (raising over £7,000) and the Brighton Marathon (raising a creditable £3,000 in the process).

**Trusts & Foundations.** We are also grateful to the many grant making Trusts and Foundations which supported our work during the year. These included:

- National Lottery Awards for All England
- The People's Postcode Trust
- The Nationwide Foundation
- The Garfield Weston Foundation
- The Lotus Charitable Foundation
- The William Grant Foundation
- Arnold Clark Community Fund
- The Thompson Family Foundation
- Carmichael Mason Charity Settlement
- Groundworks Trust
- The Gilander Foundation
- William Allen Young Charitable Trust
- The Eveson Trust

**Gifts in Wills:** We are as always very grateful to people who have generously supported our work by leaving gifts in their wills. During the period we were very fortunate to receive donations from the following individuals.:

- Mrs Jane Hickford
- Mr Pavel Simr
- Ms Cheryl Owen
- A Williams
- Ms Ann Shapter

**Donations:** We are truly grateful for the generosity of the pituitary community during the year. The tough economic environment has seen levels of donations reduce over the last 18 months, but we continued to receive generous donations from many donors during the year. We were helped significantly by our first involvement in the Big Give campaign; our Christmas Big Give Campaign 2024 helped to raise an amazing £24,501.

As ever we were very appreciative of the support from the **pharmaceutical industry**. This included:

- Sandoz providing a grant of £10,000 towards Pituitary Life patient resource magazine
- Recordati Rare Diseases who provided sponsorship of £3,000 which related to their staff fundraising efforts by joining us at the Four Falls trek in the Brecon Beacons
- Esteve who provided a grant of £6,000 towards our Pituitary Life patient resource magazine
- Novo Nordisk providing an educational grant of £9,270 towards our Pituitary Life patient resource magazine
- Recordati Rare Diseases who sponsored a Cushing's Disease event that we delivered at St Bartholomew's Hospital in London for health care professionals.

We work in line with good practice guidance in our partnerships with pharmaceutical companies. As reported elsewhere we now disclose our funding from pharmaceutical companies on our website and have a specific policy to cover our work with healthcare companies.

**Earned income** this year included a Summer and Christmas Raffle as well as items that we sell in our online shop. During the year we undertook a review of our shop merchandise and identified new products that proved to be popular with our community and delivered much needed income to help provide for our services.

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**f. Resources expended**

The staff team, alongside the Finance Committee, works hard to keep costs down, seeking competitive quotes and adhering to our financial processes. Within our total costs figure this year we spent £21,000 on our Counselling service provided by Rareminds, £2,900 on a Cushing's video and £4,600 on a Data Protection Audit.

**g. Investment Policy and performance**

We hold general and restricted funds in our main NatWest and RBS bank accounts. We have some modest sums with Monmouthshire Building Society, Julian Hodge Bank and Virgin Money.

We continue to use the excellent CAF Online deposits platform to secure competitive savings rates across a mixture of shorter and longer-term savings investments, which allows us to take advantage of competitive interest rates in an effective manner.

Our investments, which are ethically selected (in line with our Investment Policy) and managed by our investment advisers Evelyn Partners, have improved in value over the course of the year. Our total funds held with Evelyn partners are £103,878 and we have benefitted from an income of £2,693 from these investments during the year.

All of the charity's funds are subject to regular review by our Finance Committee. The Trustees have a confident expectation that The Foundation will continue in operational existence for the foreseeable future and have, therefore, used the going concern basis in preparing the financial statements.

We have an Investment Policy, developed in line with Charity Commission guidelines and reviewed and approved by the Finance Committee and Trustees, to which we continue to adhere. In particular, the Policy:

- excludes direct investment in Oil and Gas, Armaments or Tobacco sectors
- requires that the investment adviser seeks to ensure no investments are made into companies whose activities comprise more than 5% coal extraction, alcohol or tobacco production, gambling or animal testing
- requires, where monies are invested in funds, the investment adviser to target funds with positive ethical and environmental objectives.

**h. Reserves policy**

We plan to retain free operational reserves representing 9 months of our annual operating costs, equating to around £415,000. The remaining free reserves, known as the Development Reserve, are available for investment in the charity's infrastructure, resources and patient support.

Total restricted funds held at the year end were £117,606. Free reserves are calculated as unrestricted reserves less designated funds and fixed assets. The Foundation's free reserves at 30 June 2025 decreased to £585,544 (2024: £646,560). This represented 141% of our free operational reserves target.

As a result, our Development Reserve was £170,544 at the year end. As proposed by the CEO, the Trustees have agreed to planned expenditure of £46,000 from these reserves for the various areas of development outlined in the Plans For Future Periods section above.

The Reserves policy was reviewed in the financial year to ensure it remains appropriate to the Foundations needs and the wider economic landscape.

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## Structure, governance and management

### **a. Governing document**

The Pituitary Foundation was established in 1994 and became a Company Limited by Guarantee and a Registered Charity in 1996. It is, therefore, governed by company law and the Charity Commission.

Its governing documents are its Memorandum and Articles of Association.

Membership is encouraged and is open to all who wish to support The Pituitary Foundation; however, the principal services provided are available to all who require them regardless of membership.

### **b. Recruitment and appointment of Trustees**

At each General Meeting (GM), one third of the Trustees retire by rotation. Any member of The Foundation may nominate any other member for a vacant Trustee position. If there are more nominations than vacancies, members are balloted, and the result announced at the GM. Trustees may be co-opted onto the Board until the next GM at which time they must stand for re-election.

The diversity and skills required by the Board are a key consideration. Trustees recognise the value of ensuring that the composition of the Board is reflective of the community we serve and includes the necessary skills and knowledge to lead and develop The Foundation. Trustees undertook a skills audit in 2024 which helped inform the focus of recruitment during 2025.

### **c. Trustee induction and training**

Trustees are inducted within three months of appointment. They are inducted online or at the Bristol Office. At induction, members of staff and representatives from the Trustees work with the new Trustee to explain the business, services, aims and objectives of The Foundation. At the time of induction, each Trustee is given a Trustee Induction Pack. Trustee training is undertaken on an ad hoc basis. At the time of writing we are planning Trustee training in February 2026. This is planned so as to ensure that new Trustees are properly supported in their roles and to provide for a whole Board development opportunity.

### **d. Trustee tenure and retirement**

Trustees agreed a tenure policy during 2024-25 which brings The Foundation into line with Charity Commission guidance that Trustees can be reappointed up to a maximum of 9 years unless there are exceptional circumstances. The policy means that a number of Trustees retire between 2025-2026 and we are carefully managing this so as not to lose the essential skills and experience required to lead The Foundation. At the General Meeting in May 2025 we expressed our appreciation to four retiring Trustees:

- Professor Stephanie Baldeweg
- Dr Ezra Macdonald
- Claire Thatcher
- Allen Matty

We are immensely grateful to the generous commitment of our retiring Trustees who have dedicated many years to their roles.

### **e. Organisation**

The Board of Trustees governs The Foundation and Trustees are also Directors of the Company. A limit of one third has been set on the number of Trustees who are employed in the health sector to ensure strong representation of patients and carers. During 2024-2025 we had two Trustees employed in the health sector and at 30 June 2025 this reduced to one following retirement.

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The Board of Trustees holds meetings a minimum of three times per year. The Trustees have appointed four Officers of the board. These comprise the Chair, the Vice Chair, the Honorary Secretary and the Honorary Treasurer.

The Trustees debate and make decisions based on recommendations brought to Trustee meetings and informed by the wider staff, relevant Committees and relevant information.

The Foundation is a member of:

- Helplines Partnership
- The National Council for Voluntary Organisation (NCVO)
- Charity Finance Group (CFG)

and is affiliated with the Society for Endocrinology.

The charity is registered with the Fundraising Regulator and the Foundation's Head of Fundraising is a member of The Institute of Fundraising.

The Foundation has three sub-committees, which report to the Trustees through their respective Chairs. Each sub-committee has Terms of Reference:

- Finance Committee
- Lived Experience Committee
- Medical Committee

### **The Finance Committee**

The Finance Committee reports to the Board and assists the Finance and Operations Manager and CEO in the review of budgets and the management of the finances. The Finance Committee is chaired by the Trustee who has the role of Honorary Treasurer, and its members include the Chair and Vice-Chair of Trustees. The CEO and relevant staff members also attend. This Committee reviews the financial position on a quarterly basis and reports to the Board in proposing the budget, reporting on performance, and making other recommendations where there is a financial implication.

### **The Lived Experience Committee**

The Lived Experience Committee was established in June 2024 with the aim of informing and advising on our work, ensuring that the voice of lived experience is at the heart of The Foundation. The committee is chaired by a Trustee, with a report to the Board of Trustees as a standing item. During the year the Committee met three times, including in person, and has established a sub-group to focus on our health information. Also, two of the members help review Pituitary Life magazine, three times a year.

During the year, and given the newness of the Lived Experience Committee, we have considered ways of working which add value to our work and leverage its lived experience expertise. We want to ensure that there is a meaningful collaboration with our Medical Committee and will be building this into the refresh of the terms of reference.

We have also collected 'community insights' for sharing and review twice a year. These insights draw on the feedback and data we collect from our work identifying the top issues and themes. From these, we will discuss and agree any actions so that we are directly responding to what is most important for our community.

Members of the Lived Experience Committee as at 30 June 2025:

- Alexia Deleligné (Trustee, Chair of the Lived Experience Committee)
- Steve Harris
- Ketan Mistry
- Gabrielle de Witte
- Ben Marinic

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- Rachel Lygoe
- Kevin Williams
- Roz Way

We were extremely saddened by the passing of Steve Harris in August 2025. Steve was a long-standing and committed volunteer and member of our community. We would like to acknowledge all his contributions over many years and will always remember him with deep affection.

**The Medical Committee**

Our volunteer Medical Committee is made up of endocrine and related medical professionals who provide invaluable expertise answering more complex patient queries, speaking at our events (online and in person), writing articles for our magazine Pituitary Life, and reviewing and providing content for our website and booklets.

As described above in relation to the Lived Experience Committee, we are now sharing 'community insights' with the Medical Committee twice a year. This provides the opportunity for medical insight and experience to input into the areas of work that we want to prioritise for advocacy and awareness raising.

Also during the year, the chairs of the Medical Committee and the Lived Experience Committee discussed ways of working to ensure that there is a strong connection across the two committees. It was agreed that going forward the Chair of the Lived Experience Committee is invited to the Medical Committee meetings.

Members of the Medical Committee as at 30 June 2025:

<b>Role</b>	<b>Name</b>
Consultant Endocrinologist	Professor Stephanie Baldeweg (Chair)
Consultant Endocrinologist	Dr Rob Murray (deputy Chair)
Endocrine nurse	Darshna Patel
Endocrine nurse	Aldons Chua
Paediatric Endocrinologist	Dr Indi Bannerjee
Consultant Endocrinologist	Dr Niki Karavitaki
Endocrine nurse (Advanced Nurse Practitioner)	Sherwin Criseno
Consultant Endocrinologist	Professor John Wass
Independent Chartered Psychologist	Dr Sue Jackson
Consultant Endocrinologist	Dr Jon Pinkney
Consultant Endocrinologist	Dr Anna Crown
Hon Cons Nurse in Endocrinology	Dr Sofia Llahana
Pituitary surgery specialist nurse	Fiona Cains
Pituitary Surgeon	Mr Omar Pathmanaban
Endocrine nurse	Alison Milne
Consultant Pharmacist	Dr Philip Newland-Jones
Advanced Clinical Pharmacist	Dr Matthew Heppel
Endocrinologist	Professor Miles Levy
Consultant Neurosurgeon	Eleni Maratos
GP	Jennifer Collins
Specialist Neuro Orthoptist & Stroke Lead	Samantha McAuley
Endocrinologist	Antonia Brooke

We've agreed that going forward all Medical Trustees of The Foundation will automatically be members of the Medical Committee. This is to ensure that there is a strong connection between the Board and the Medical Committee.

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The Foundation understands the members of our Medical Committee are professionals with demanding occupations and we are indebted to them for all of their time and effort working with staff and other volunteers to provide support to the pituitary community.

**f. Volunteers**

Volunteers are at the heart of The Foundation. We could not do what we do without the involvement, commitment, and generosity of volunteers. Our volunteer programme has dedicated staff support and we are committed to providing a good quality and rewarding experience of volunteering with us.

During the year over 80 volunteers supported our work. Volunteers gave their time as Trustees, supporting in the office, reviewing our health information, running the information helpline, providing lived experience support, leading support groups (online and in person in local areas), and as ambassadors and fundraisers.

In 2025 we refreshed the volunteer handbooks and guidance and are now confident we're providing volunteers with the best possible resources. We were delighted to welcome ten new volunteers. Two of these volunteers are supporting our Information and Support helpline and enabled the availability of the service to be increased. Towards the end of the year, we delivered online training for all our volunteers, covering safeguarding and data protection essentials.

Volunteers deliver vital roles, generously sharing their lived experience and knowledge of the patient pathway to support people with recent diagnoses, their friends and families, as well as longer term pituitary patients. The Trustees would like to extend heartfelt thanks to each person for their time and kindness being involved in our work.

**f. Staff Employed**

As at 30 June 2025 The Foundation employed eleven members of staff (7 FTE):

Chief Executive Officer	Emma Cooper (from July 2024)
Head of Support Services	Pat McBride
Head of Fundraising	Jay Sheppard
Finance and Operations Manager	Gabrielle Welland
IT and Systems Manager	Martin Cookson
Endocrine Specialist Nurse	Darshna Patel
Endocrine Specialist Nurse	Aldons Chua
Volunteer and Support Services Officer	Nesta Jones (from July 2024)
Communications Officer	Amy Shingler (from January 2025)
Administrator	Isabella Jackson (from October 2024)
Project Officer	Amber Noushad (from June 2025)

During the year, three colleagues left their employment with The Foundation - Stephanie Gauvin, Lottie Storey, and Vanessa Quinn. We are hugely grateful for the contribution each made to our work. Each directly and indirectly supported our community and will be missed by our team and community.

The Charity's staff has a clear line management reporting structure to the CEO, who reports to the Board of Trustees through the Chair. The CEO and the Chairman of the Trustees meet online or in person on a regular basis.

There are clear reporting lines for all staff, and regular 1-1's supporting an annual appraisal process. Staff wellbeing is central to the culture of our work and staff benefits have been reviewed to include access to an EAP programme, 30 days annual leave, flexible working and 5% pension contributions.

Salaries are reviewed annually as well as benchmarked periodically against similar roles either in the Bristol area, or in the NHS for specialised nurse roles. Recommendations for pay increases, or new

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salaries where a role is being advertised, are made by the CEO for review by the Finance Committee and remitted to the Trustee Board for approval.

The team are supremely dedicated and bring a huge amount of energy and commitment to their work, supporting and seeking the best outcomes for all people living with pituitary conditions and their support networks. The Trustees would like to thank them for their ongoing dedication and hard work.

**g. Risk management**

The Foundation maintains a current and comprehensive Risk Register. Management of this register is held jointly by the Chair of the Trustees and the CEO. The Register is reviewed on a regular basis and is reported to the Board of Trustees, including established controls and actions to mitigate relevant identified risks.

**TRUSTEES' RESPONSIBILITY STATEMENT**

The Trustees (who are also the directors of The Pituitary Foundation for the purposes of company law) are responsible for preparing the Trustees' Report and the financial statements in accordance with applicable law and United Kingdom Accounting Standards (United Kingdom Generally Accepted Accounting Practice) including Financial Reporting Standard 102 "The Financial Reporting Standard applicable in the UK and Republic of Ireland".

Company law requires the Trustees to prepare financial statements for each financial year which give a true and fair view of the state of affairs of the charitable company and of the incoming resources and application of resources, including the income and expenditure, of the charitable company for that period. In preparing those financial statements, the Trustees are required to

- select suitable accounting policies and then apply them consistently;
- observe the methods and principles in the Charity SORP;
- make judgements and estimates that are reasonable and prudent;
- prepare the financial statements on the going concern basis unless it is inappropriate to presume that the charitable company will continue in business.

The Trustees are responsible for keeping proper accounting records which disclose with reasonable accuracy at any time the financial position of the charitable company and to enable them to ensure that the financial statements comply with the Companies Act 2006. They are also responsible for safeguarding the assets of the charitable company and hence for taking reasonable steps for the prevention and detection of fraud and other irregularities.

Trustees' report, incorporating a strategic report, approved by order of the board of Trustees, as the company directors, on .....<sup>13/01/2026</sup>..... and signed on the board's behalf by:



.....  
Mr T Sumpster - Trustee

INDEPENDENT EXAMINER'S REPORT TO THE TRUSTEES OF  
THE PITUITARY FOUNDATION

**Independent examiner's report to the trustees of The Pituitary Foundation ('the Company')**

I report to the charity trustees on my examination of the accounts of the Company for the year ended 30 June 2025.

**Responsibilities and basis of report**

As the charity's trustees of the Company (and also its directors for the purposes of company law) you are responsible for the preparation of the accounts in accordance with the requirements of the Companies Act 2006 ('the 2006 Act').

Having satisfied myself that the accounts of the Company are not required to be audited under Part 16 of the 2006 Act and are eligible for independent examination, I report in respect of my examination of your charity's accounts as carried out under Section 145 of the Charities Act 2011 ('the 2011 Act'). In carrying out my examination I have followed the Directions given by the Charity Commission under Section 145(5) (b) of the 2011 Act.

**Independent examiner's statement**

Since your charity's gross income exceeded £250,000 your examiner must be a member of a listed body. I can confirm that I am qualified to undertake the examination because I am a member of the Institute of Chartered Accountants in England and Wales, which is one of the listed bodies.

I have completed my examination. I confirm that no matters have come to my attention in connection with the examination giving me cause to believe:

1. accounting records were not kept in respect of the Company as required by Section 386 of the 2006 Act; or
2. the accounts do not accord with those records; or
3. the accounts do not comply with the accounting requirements of Section 396 of the 2006 Act other than any requirement that the accounts give a true and fair view which is not a matter considered as part of an independent examination; or
4. the accounts have not been prepared in accordance with the methods and principles of the Statement of Recommended Practice for accounting and reporting by charities (applicable to charities preparing their accounts in accordance with the Financial Reporting Standard applicable in the UK and Republic of Ireland (FRS 102)).

I have no concerns and have come across no other matters in connection with the examination to which attention should be drawn in this report in order to enable a proper understanding of the accounts to be reached.

*C Moelwyn-Williams*

C Moelwyn-Williams, BSc FCA

Date: *14th January 2026*



THE PITUITARY FOUNDATION

STATEMENT OF FINANCIAL ACTIVITIES  
(INCORPORATING AN INCOME AND EXPENDITURE ACCOUNT)  
FOR THE YEAR ENDED 30 JUNE 2025

	Notes	Unrestricted fund £	Restricted funds £	2025 Total funds £	2024 Total funds £
<b>INCOME AND ENDOWMENTS FROM</b>					
Voluntary income	2	264,183	9,447	273,630	596,704
<b>Charitable activities</b>	5				
General		65,300	-	65,300	38,760
Restricted		-	56,379	56,379	92,274
Activities for generating funds	3	29,901	-	29,901	39,523
Investment income	4	22,520	-	22,520	12,304
<b>Total</b>		<u>381,904</u>	<u>65,826</u>	<u>447,730</u>	<u>779,565</u>
<b>EXPENDITURE ON</b>					
Costs of Generating Voluntary Income	6	120,831	4,200	125,031	120,331
<b>Charitable activities</b>	7				
General		343,562	54,185	397,747	366,401
<b>Total</b>		<u>464,393</u>	<u>58,385</u>	<u>522,778</u>	<u>486,732</u>
Net gains on investments		<u>2,693</u>	<u>-</u>	<u>2,693</u>	<u>5,943</u>
<b>NET INCOME/(EXPENDITURE)</b>		<u>(79,796)</u>	<u>7,441</u>	<u>(72,355)</u>	<u>298,776</u>
<b>Transfers between funds</b>	21	<u>20,408</u>	<u>(20,408)</u>	<u>-</u>	<u>-</u>
<b>Net movement in funds</b>		<u>(59,388)</u>	<u>(12,967)</u>	<u>(72,355)</u>	<u>298,776</u>
<b>RECONCILIATION OF FUNDS</b>					
Total funds brought forward		648,501	130,573	779,074	480,298
<b>TOTAL FUNDS CARRIED FORWARD</b>		<u><u>589,113</u></u>	<u><u>117,606</u></u>	<u><u>706,719</u></u>	<u><u>779,074</u></u>

**CONTINUING OPERATIONS**

All income and expenditure has arisen from continuing activities.

The notes form part of these financial statements

STATEMENT OF FINANCIAL POSITION

30 JUNE 2025

	Notes	2025 £	2024 £
<b>FIXED ASSETS</b>			
Tangible assets	14	3,569	1,941
Investments	15	103,878	99,344
		<hr/> 107,447	<hr/> 101,285
<b>CURRENT ASSETS</b>			
Debtors	16	9,578	5,458
Investments	17	290,500	119,000
Cash at bank		325,847	571,423
		<hr/> 625,925	<hr/> 695,881
<b>CREDITORS</b>			
Amounts falling due within one year	18	(26,653)	(18,092)
		<hr/>	<hr/>
<b>NET CURRENT ASSETS</b>		<hr/> 599,272	<hr/> 677,789
<b>TOTAL ASSETS LESS CURRENT LIABILITIES</b>		<hr/>	<hr/>
		706,719	779,074
<b>NET ASSETS</b>		<hr/>	<hr/>
		706,719	779,074
<b>FUNDS</b>	21		
Unrestricted funds		589,113	648,501
Restricted funds		117,606	130,573
		<hr/>	<hr/>
<b>TOTAL FUNDS</b>		<hr/> 706,719	<hr/> 779,074

The charitable company is entitled to exemption from audit under Section 477 of the Companies Act 2006 for the year ended 30 June 2025.

The members have not required the company to obtain an audit of its financial statements for the year ended 30 June 2025 in accordance with Section 476 of the Companies Act 2006.

The trustees acknowledge their responsibilities for

- ensuring that the charitable company keeps accounting records that comply with Sections 386 and 387 of the Companies Act 2006 and
- preparing financial statements which give a true and fair view of the state of affairs of the charitable company as at the end of each financial year and of its surplus or deficit for each financial year in accordance with the requirements of Sections 394 and 395 and which otherwise comply with the requirements of the Companies Act 2006 relating to financial statements, so far as applicable to the charitable company.

The financial statements were approved by the Board of Trustees and authorised for issue on 13/01/2026 and were signed on its behalf by:

The notes form part of these financial statements

STATEMENT OF FINANCIAL POSITION - continued

30 JUNE 2025



.....  
Mr T Sumpster - Trustee



.....  
Mr P Rouse - Trustee

17/12/2025

## THE PITUITARY FOUNDATION

### NOTES TO THE FINANCIAL STATEMENTS FOR THE YEAR ENDED 30 JUNE 2025

#### **1. ACCOUNTING POLICIES**

##### **Basis of preparing the financial statements**

The financial statements of the charitable company, which is a public benefit entity under FRS 102, have been prepared in accordance with the Charities SORP (FRS 102) 'Accounting and Reporting by Charities: Statement of Recommended Practice applicable to charities preparing their accounts in accordance with the Financial Reporting Standard applicable in the UK and Republic of Ireland (FRS 102) (effective 1 January 2019)', Financial Reporting Standard 102 'The Financial Reporting Standard applicable in the UK and Republic of Ireland' and the Companies Act 2006. The financial statements have been prepared under the historical cost convention, with the exception of investments which are included at market value, as modified by the revaluation of certain assets.

##### **Financial reporting standard 102 - reduced disclosure exemptions**

The charitable company has taken advantage of the following disclosure exemptions in preparing these financial statements, as permitted by FRS 102 'The Financial Reporting Standard applicable in the UK and Republic of Ireland':

- the requirements of Section 7 Statement of Cash Flows;
- the requirement of paragraph 3.17(d);
- the requirements of paragraphs 11.42, 11.44, 11.45, 11.47, 11.48(a)(iii), 11.48(a)(iv), 11.48(b) and 11.48(c);
- the requirements of paragraphs 12.26, 12.27, 12.29(a), 12.29(b) and 12.29A;
- the requirement of paragraph 33.7.

##### **Income**

All income is recognised in the Statement of Financial Activities once the charity has entitlement to the funds, it is probable that the income will be received and the amount can be measured reliably.

##### **Expenditure**

Liabilities are recognised as expenditure as soon as there is a legal or constructive obligation committing the charity to that expenditure, it is probable that a transfer of economic benefits will be required in settlement and the amount of the obligation can be measured reliably. Expenditure is accounted for on an accruals basis and has been classified under headings that aggregate all cost related to the category. Where costs cannot be directly attributed to particular headings they have been allocated to activities on a basis consistent with the use of resources.

##### **Tangible fixed assets**

Depreciation is provided at the following annual rates in order to write off each asset over its estimated useful life.

Computer equipment - 5 years straight line

##### **Taxation**

The charity is exempt from corporation tax on its charitable activities.

##### **Fund accounting**

Unrestricted funds can be used in accordance with the charitable objectives at the discretion of the trustees.

Restricted funds can only be used for particular restricted purposes within the objects of the charity. Restrictions arise when specified by the donor or when funds are raised for particular restricted purposes.

Further explanation of the nature and purpose of each fund is included in the notes to the financial statements.

## THE PITUITARY FOUNDATION

### NOTES TO THE FINANCIAL STATEMENTS - continued FOR THE YEAR ENDED 30 JUNE 2025

#### **1. ACCOUNTING POLICIES - continued**

##### **Pension costs and other post-retirement benefits**

The charitable company operates a defined contribution pension scheme. Contributions payable to the charitable company's pension scheme are charged to the Statement of Financial Activities in the period to which they relate.

##### **Investments**

Assets held for investment purposes are valued at market value at the balance sheet date. Net gains and losses arising on revaluation and disposals during the year are included in the statement of financial activities.

##### **Cash at bank and in hand**

Cash at bank and in hand includes cash and short-term highly liquid investments with a short maturity of three months or less from the date of acquisition or opening of the deposit or similar account.

##### **Financial instruments**

The charity only has financial assets and financial liabilities of a kind that qualify as basic financial instruments. Basic financial instruments are initially recognised at transaction value and subsequently measured at their settlement value.

##### **Debtors**

Trade and other debtors are recognised at the settlement amount after any trade discount offered. Prepayments are valued at the amount prepaid net of any trade discounts due.

##### **Operating Leases**

Rentals under operating leases are charged to the statement of financial activities on a straight line basis over the lease term.

#### **2. VOLUNTARY INCOME**

	unrestricted funds £	Restricted funds £	2025 Total funds £	2024 Total funds £
Donations	161,909	9,447	171,356	227,224
Gift aid	26,622	-	26,622	35,538
Legacies	42,125	-	42,125	298,095
Membership	33,141	-	33,141	31,699
Sundry income	386	-	386	4,148
	<u>264,183</u>	<u>9,447</u>	<u>273,630</u>	<u>596,704</u>

Included within unrestricted voluntary income is £nil (2024: £nil) of income raised by the local support groups.

Voluntary income in the prior year included £596,222 of unrestricted income and £482 of restricted income.

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

3. ACTIVITIES FOR GENERATING FUNDS

	Unrestricted funds £	Restricted funds £	2025 Total funds £	2024 Total funds £
Lottery & raffle proceeds	14,215	-	14,215	24,537
Merchandise proceeds	15,686	-	15,686	14,986
	<u>29,901</u>	<u>-</u>	<u>29,901</u>	<u>39,523</u>

Activities for generating funds in the prior year included £39,523 of unrestricted income and no restricted income.

4. INVESTMENT INCOME

	Unrestricted funds £	Restricted funds £	2025 Total funds £	2024 Total funds £
Bank interest receivable	20,643	-	20,643	9,927
Dividend Income	1,483	-	1,483	1,994
Investment Income	394	-	394	383
	<u>22,520</u>	<u>-</u>	<u>22,520</u>	<u>12,304</u>

Investment income in the prior year included £12,304 of unrestricted income and no restricted income.

5. INCOME FROM CHARITABLE ACTIVITIES

	General £	Restricted £	2025 Total activities £	2024 Total activities £
Grants	65,150	56,379	121,529	131,034
Publications	150	-	150	-
	<u>65,300</u>	<u>56,379</u>	<u>121,679</u>	<u>131,034</u>

Income from charitable activities in the prior year included £38,760 of unrestricted income and £92,274 of restricted income.

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**6. COSTS OF GENERATING VOLUNTARY INCOME**

**Costs of generating voluntary income**

	Unrestricted funds £	Restricted funds £	2025 Total funds £	2024 Total funds £
Staff costs	88,707	-	88,707	86,878
Event costs	13,169	1,200	14,369	7,215
Online giving fees	3,778	-	3,778	5,273
Printing, postage & promotion costs	4,859	3,000	7,859	6,545
Other staff costs	5,600	-	5,600	5,440
Local Support Group costs	1,398	-	1,398	27
	<u>117,511</u>	<u>4,200</u>	<u>121,711</u>	<u>111,378</u>

Costs of generating voluntary income in the prior year included £110,319 of unrestricted costs and £1,059 restricted costs.

**Fundraising trading: costs of goods sold and other costs**

	Unrestricted funds £	Restricted funds £	2025 Total funds £	2024 Total funds £
Purchases	<u>3,320</u>	<u>-</u>	<u>3,320</u>	<u>8,953</u>

Costs of goods sold and other costs in the prior year included £8,953 of unrestricted costs and no restricted costs.

Aggregate amounts	<u>120,831</u>	<u>4,200</u>	<u>125,031</u>	<u>120,331</u>
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**7. CHARITABLE ACTIVITIES COSTS**

	Direct Costs (see note 8) £	Support costs (see note 9) £	Totals £
General	<u>147,791</u>	<u>249,956</u>	<u>397,747</u>

Charitable activities costs in the prior year were £366,401, made up of £334,989 of unrestricted costs and £31,412 of restricted costs.

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**8. DIRECT COSTS OF CHARITABLE ACTIVITIES**

	2025	2024
	£	£
Staff costs	124,953	114,964
Staff training	2,238	11,815
Publication costs	19,947	12,695
Depreciation	653	485
	<u>147,791</u>	<u>139,959</u>

**9. SUPPORT COSTS**

	Other	Governance	Totals
	£	costs	£
General	<u>212,731</u>	<u>37,225</u>	<u>249,956</u>

Support costs, included in the above, are as follows:

**Other**

	2025	2024
	General	Total
	£	activities
	£	£
Wages	81,438	77,559
Social security	5,868	5,710
Pensions	4,070	3,821
Rent payable under operating leases	23,410	21,866
Insurance	5,944	4,578
Telephone, photocopier, postage & office		
supplies	13,812	13,882
Sundries	708	410
Bank charges	3,662	3,729
Website & online	2,375	9,655
Other staff costs	4,287	12,007
Software licences & expenses	22,437	20,604
Equipment	150	13
Professional fees	28,544	10,473
Finance & HR consultancy	9,685	3,173
Accommodation, travel & subsistence	3,210	3,896
Membership subscriptions	2,709	2,579
PR & Branding	422	422
	<u>212,731</u>	<u>194,377</u>



THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**9. SUPPORT COSTS - continued**  
**Governance costs**

	2025	2024
	General	Total
	£	activities
		£
Wages	24,596	23,230
Social security	1,949	2,084
Pensions	1,228	1,162
Independent Examination fees	2,900	2,850
Other governance costs	6,361	1,419
Trustee training	191	1,320
	<u>37,225</u>	<u>32,065</u>

**10. NET INCOME/(EXPENDITURE)**

Net income/(expenditure) is stated after charging/(crediting):

	2025	2024
	£	£
Depreciation - owned assets	<u>653</u>	<u>485</u>

**11. TRUSTEES' REMUNERATION AND BENEFITS**

There were no trustees' remuneration or other benefits for the year ended 30 June 2025 nor for the year ended 30 June 2024.

**Trustees' expenses**

During the year two Trustees received reimbursement of expenses amounting to £70 (2024: £145).

**12. STAFF COSTS**

	2025	2024
	£	£
Wages and salaries	298,487	281,167
Social security costs	20,617	21,013
Other pension costs	13,705	13,228
	<u>332,809</u>	<u>315,408</u>

The average monthly number of employees during the year was as follows:

	2025	2024
	11	10
Core activities	<u>11</u>	<u>10</u>

No employees received emoluments in excess of £60,000.

The average number of full time equivalent employees during the year was 8.

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**12. STAFF COSTS - continued**

The key management personnel of the charity comprise the Chief Executive Officer, the Head of Patient and Family Services and the Head of Fundraising. The total employee benefits of the key management personnel of the charity were £103,107 (2024: £160,534).

**13. COMPARATIVES FOR THE STATEMENT OF FINANCIAL ACTIVITIES**

	Unrestricted fund £	Restricted funds £	Total funds £
<b>INCOME AND ENDOWMENTS FROM</b>			
Voluntary income	596,222	482	596,704
<b>Charitable activities</b>			
General	38,760	-	38,760
Restricted	-	92,274	92,274
Activities for generating funds	39,523	-	39,523
Investment income	12,304	-	12,304
<b>Total</b>	<b>686,809</b>	<b>92,756</b>	<b>779,565</b>
<b>EXPENDITURE ON</b>			
Costs of Generating Voluntary Income	119,272	1,059	120,331
<b>Charitable activities</b>			
General	334,989	31,412	366,401
<b>Total</b>	<b>454,261</b>	<b>32,471</b>	<b>486,732</b>
Net gains on investments	5,943	-	5,943
<b>NET INCOME</b>	<b>238,491</b>	<b>60,285</b>	<b>298,776</b>
<b>RECONCILIATION OF FUNDS</b>			
Total funds brought forward	410,010	70,288	480,298
<b>TOTAL FUNDS CARRIED FORWARD</b>	<b>648,501</b>	<b>130,573</b>	<b>779,074</b>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**14. TANGIBLE FIXED ASSETS**

	Fixtures and fittings £	Computer equipment £	Totals £
<b>COST</b>			
At 1 July 2024	47,309	3,948	51,257
Additions	-	2,281	2,281
	<hr/>	<hr/>	<hr/>
At 30 June 2025	47,309	6,229	53,538
	<hr/>	<hr/>	<hr/>
<b>DEPRECIATION</b>			
At 1 July 2024	47,309	2,007	49,316
Charge for year	-	653	653
	<hr/>	<hr/>	<hr/>
At 30 June 2025	47,309	2,660	49,969
	<hr/>	<hr/>	<hr/>
<b>NET BOOK VALUE</b>			
At 30 June 2025	-	3,569	3,569
	<hr/>	<hr/>	<hr/>
At 30 June 2024	-	1,941	1,941
	<hr/>	<hr/>	<hr/>

**15. FIXED ASSET INVESTMENTS**

	Unlisted investments £	Cash and settlements pending £	Totals £
<b>MARKET VALUE</b>			
At 1 July 2024	98,430	914	99,344
Additions	16,113	18,819	34,932
Disposals	(16,344)	(16,747)	(33,091)
Revaluations	2,693	-	2,693
	<hr/>	<hr/>	<hr/>
At 30 June 2025	100,892	2,986	103,878
	<hr/>	<hr/>	<hr/>
<b>NET BOOK VALUE</b>			
At 30 June 2025	100,892	2,986	103,878
	<hr/>	<hr/>	<hr/>
At 30 June 2024	98,430	914	99,344
	<hr/>	<hr/>	<hr/>

49% of the investment value is held within the UK. The remainder is held in overseas investments.

Cost or valuation at 30 June 2025 is represented by:

	Unlisted investments £	Cash and settlements pending £	Totals £
Valuation in 2025	2,693	-	2,693
Cost	98,199	2,986	101,185
	<hr/>	<hr/>	<hr/>
	100,892	2,986	103,878
	<hr/>	<hr/>	<hr/>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**16. DEBTORS: AMOUNTS FALLING DUE WITHIN ONE YEAR**

	2025	2024
	£	£
Prepayments and accrued income	9,578	5,458
	<u>9,578</u>	<u>5,458</u>

**17. CURRENT ASSET INVESTMENTS**

	2025	2024
	£	£
Unlisted investments	290,500	119,000
	<u>290,500</u>	<u>119,000</u>

**18. CREDITORS: AMOUNTS FALLING DUE WITHIN ONE YEAR**

	2025	2024
	£	£
Trade creditors	13,566	6,390
Social security and other taxes	6,677	6,889
Other creditors	3,097	1,666
Accruals and deferred income	3,313	3,147
	<u>26,653</u>	<u>18,092</u>

Deferred income in the prior period included income of £nil received in relation to the current financial year ended 30 June 2025.

**19. LEASING AGREEMENTS**

Minimum lease payments under non-cancellable operating leases fall due as follows:

	2025	2024
	£	£
Within one year	25,687	23,975
Between one and five years	20,334	46,375
	<u>46,021</u>	<u>70,350</u>

**20. ANALYSIS OF NET ASSETS BETWEEN FUNDS**

	Unrestricted fund £	Restricted funds £	2025 Total funds £	2024 Total funds £
Fixed assets	3,569	-	3,569	1,941
Investments	103,878	-	103,878	99,344
Current assets	506,088	119,837	625,925	695,881
Current liabilities	(24,422)	(2,231)	(26,653)	(18,092)
	<u>589,113</u>	<u>117,606</u>	<u>706,719</u>	<u>779,074</u>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**21. MOVEMENT IN FUNDS**

	At 1.7.24 £	Net movement in funds £	Transfers between funds £	At 30.6.25 £
<b>Unrestricted funds</b>				
General fund	648,501	(79,796)	20,408	589,113
<b>Restricted funds</b>				
Endocrine Nurse helpline	1,853	3,157	-	5,010
Publications	31,441	(1,254)	-	30,187
Volunteer Development	-	19,787	-	19,787
Society for Endocrinology	615	(615)	-	-
Legacy - Solent & IOW Support Group	7,950	-	-	7,950
Cushing's Disease Day	6,876	200	(7,076)	-
GP Professional Development	11,180	-	(11,180)	-
NI Support Group	152	1,000	(1,152)	-
Paediatric Work	34,876	-	-	34,876
Psychological Support	31,630	(21,450)	-	10,180
Cushings Information Content	4,000	(3,000)	(1,000)	-
Pituitary Life	-	9,616	-	9,616
	<u>130,573</u>	<u>7,441</u>	<u>(20,408)</u>	<u>117,606</u>
<b>TOTAL FUNDS</b>	<u>779,074</u>	<u>(72,355)</u>	<u>-</u>	<u>706,719</u>

Net movement in funds, included in the above are as follows:

	Incoming resources £	Resources expended £	Gains and losses £	Movement in funds £
<b>Unrestricted funds</b>				
General fund	381,904	(464,393)	2,693	(79,796)
<b>Restricted funds</b>				
Endocrine Nurse helpline	9,447	(6,290)	-	3,157
Publications	-	(1,254)	-	(1,254)
Volunteer Development	20,000	(213)	-	19,787
Society for Endocrinology	-	(615)	-	(615)
Cushing's Disease Day	1,700	(1,500)	-	200
NI Support Group	1,000	-	-	1,000
Psychological Support	-	(21,450)	-	(21,450)
Cushings Information Content	-	(3,000)	-	(3,000)
Pituitary Life	33,039	(23,423)	-	9,616
BackSOM Study	640	(640)	-	-
	<u>65,826</u>	<u>(58,385)</u>	<u>-</u>	<u>7,441</u>
<b>TOTAL FUNDS</b>	<u>447,730</u>	<u>(522,778)</u>	<u>2,693</u>	<u>(72,355)</u>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**21. MOVEMENT IN FUNDS - continued**

**Comparatives for movement in funds**

	At 1.7.23 £	Net movement in funds £	Transfers between funds £	At 30.6.24 £
<b>Unrestricted funds</b>				
General fund	410,010	238,491	-	648,501
<b>Restricted funds</b>				
Endocrine Nurse helpline	16,291	(14,438)	-	1,853
Publications	22,487	8,954	-	31,441
Society for Endocrinology	-	615	-	615
2nd Endocrine Nurse Helpline	3,714	(3,714)	-	-
Legacy - Solent & IOW Support Group	7,950	-	-	7,950
Paediatric Endocrine Nurse	19,846	-	(19,846)	-
Cushing's Disease Day	-	6,876	-	6,876
GP Professional Development	-	11,180	-	11,180
NI Support Group	-	152	-	152
Paediatric Work	-	15,030	19,846	34,876
Psychological Support	-	31,630	-	31,630
Cushings Information Content	-	4,000	-	4,000
	<u>70,288</u>	<u>60,285</u>	<u>-</u>	<u>130,573</u>
<b>TOTAL FUNDS</b>	<u>480,298</u>	<u>298,776</u>	<u>-</u>	<u>779,074</u>

Comparative net movement in funds, included in the above are as follows:

	Incoming resources £	Resources expended £	Gains and losses £	Movement in funds £
<b>Unrestricted funds</b>				
General fund	686,809	(454,261 )	5,943	238,491
<b>Restricted funds</b>				
Endocrine Nurse helpline	-	(14,438)	-	(14,438)
Publications	9,899	(945)	-	8,954
Society for Endocrinology	1,000	(385)	-	615
2nd Endocrine Nurse Helpline	-	(3,714)	-	(3,714)
Cushing's Disease Day	7,000	(124)	-	6,876
GP Professional Development	19,680	(8,500)	-	11,180
NI Support Group	152	-	-	152
Paediatric Work	15,030	-	-	15,030
Psychological Support	35,995	(4,365)	-	31,630
Cushings Information Content	4,000	-	-	4,000
	<u>92,756</u>	<u>(32,471)</u>	<u>-</u>	<u>60,285</u>
<b>TOTAL FUNDS</b>	<u>779,565</u>	<u>(486,732 )</u>	<u>5,943</u>	<u>298,776</u>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**21. MOVEMENT IN FUNDS - continued**

A current year 12 months and prior year 12 months combined position is as follows:

	At 1.7.23 £	Net movement in funds £	Transfers between funds £	At 30.6.25 £
<b>Unrestricted funds</b>				
General fund	410,010	158,695	20,408	589,113
<b>Restricted funds</b>				
Endocrine Nurse helpline	16,291	(11,281)	-	5,010
Publications	22,487	7,700	-	30,187
Volunteer Development	-	19,787	-	19,787
2nd Endocrine Nurse Helpline	3,714	(3,714)	-	-
Legacy - Solent & IOW Support Group	7,950	-	-	7,950
Paediatric Endocrine Nurse	19,846	-	(19,846)	-
Cushing's Disease Day	-	7,076	(7,076)	-
GP Professional Development	-	11,180	(11,180)	-
NI Support Group	-	1,152	(1,152)	-
Paediatric Work	-	15,030	19,846	34,876
Psychological Support	-	10,180	-	10,180
Cushings Information Content	-	1,000	(1,000)	-
Pituitary Life	-	9,616	-	9,616
	<u>70,288</u>	<u>67,726</u>	<u>(20,408)</u>	<u>117,606</u>
<b>TOTAL FUNDS</b>	<u>480,298</u>	<u>226,421</u>	<u>-</u>	<u>706,719</u>

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**21. MOVEMENT IN FUNDS - continued**

A current year 12 months and prior year 12 months combined net movement in funds, included in the above are as follows:

	Incoming resources £	Resources expended £	Gains and losses £	Movement in funds £
<b>Unrestricted funds</b>				
General fund	1,068,713	(918,654 )	8,636	158,695
<b>Restricted funds</b>				
Endocrine Nurse helpline	9,447	(20,728)	-	(11,281)
Publications	9,899	(2,199)	-	7,700
Volunteer Development	20,000	(213)	-	19,787
Society for Endocrinology	1,000	(1,000)	-	-
2nd Endocrine Nurse Helpline	-	(3,714)	-	(3,714)
Cushing's Disease Day	8,700	(1,624)	-	7,076
GP Professional Development	19,680	(8,500)	-	11,180
NI Support Group	1,152	-	-	1,152
Paediatric Work	15,030	-	-	15,030
Psychological Support	35,995	(25,815)	-	10,180
Cushings Information Content	4,000	(3,000)	-	1,000
Pituitary Life	33,039	(23,423)	-	9,616
BackSOM Study	640	(640)	-	-
	<u>158,582</u>	<u>(90,856)</u>	<u>-</u>	<u>67,726</u>
<b>TOTAL FUNDS</b>	<u>1,227,295</u>	<u>(1,009,510 )</u>	<u>8,636</u>	<u>226,421</u>

Purpose of funds

**Unrestricted revenue funds:-**

These funds are held for meeting the objectives of the charity, and to provide reserves for future activities, and, subject to charity legislation, are free from all restrictions on their use.

**Restricted funds:-**

Endocrine Nurse Helpline

A specialist endocrine nurse helpline service for pituitary patients, carers, family and the wider pituitary community.

Publications

The charity provides various booklets, factsheets and other information publications.

Volunteer Development

Funding provided for an in-person volunteer training event to be held in June 2026.

Society for Endocrinology

Funds received to cover expenses related to attendance at the BES Conference in November 2024.

Legacy - Solent & IOW Support Group

Legacy income received to specifically support the Solent & IOW support group



THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**21. MOVEMENT IN FUNDS - continued**

Cushing's Disease Day

Funding provided for delivery of a support event in London for Cushing's patients.

GP Professional Development

Funding provided for the provision of an outsourced training event for all UK GP's focused on pituitary illnesses.

NI Support Group

Funds given for use by the Northern Ireland Support Group in providing the Group's services to pituitary patients and their families and carers in Northern Ireland.

Paediatric Work

Funds provided for the charity's work in supporting children and young people, and their families and carers.

Psychological Support

Funds for the provision of counselling and other psychological support services, provided in conjunction with Rare Minds, a charity with expertise in providing psychological support for patients suffering from rare diseases.

Cushings Information Content

Funds received for the creation of informative media regarding Cushing's Disease.

Pituitary Life

Funding provided for the costs of the Pituitary Life magazine produced three times a year by the charity.

BackSOM Study

Funds provided for assistance given in identifying participants for a research study.

**22. EMPLOYEE BENEFIT OBLIGATIONS**

The company operates a defined contributions pension scheme. The assets of the scheme are held separately from those of the company in an independently administered fund. The pension cost charge represents contributions payable by the company to the fund and amounted to £13,705 (2024: £13,228). Contributions totalling £2,155 (2024: £1,968) were payable to the fund at the balance sheet date and are included in creditors.

**23. RELATED PARTY DISCLOSURES**

There were no identified related parties other than the Trustees. Details of any payments to Trustees are detailed in note 11. The total amount of donations received without condition from the Trustees during the year was £225 (2024: £290).

THE PITUITARY FOUNDATION

NOTES TO THE FINANCIAL STATEMENTS - continued  
FOR THE YEAR ENDED 30 JUNE 2025

**24. ULTIMATE CONTROLLING PARTY**

The Pituitary Foundation is a company limited by guarantee governed in accordance with its Memorandum and Articles of Association.

The charitable company is wholly controlled by its members, who are also trustees of the charity.