

Impact Report 2025

CEREBRA

Working wonders for children with brain conditions



Liwsi

Mabli & Liwsi: A Story of Connection, Courage and Possibility

Mabli and Liwsi are just 20 months old, both born with rare genetic conditions. Their mums, Gwenno and Zoe, met by chance at a moment when everything felt overwhelming — two families trying to make sense of a new "normal", unsure what the future would hold for their little girls. They felt alone, frightened, and exhausted.

Today, they are firm friends. And they have found a steadfast ally in Cerebra.

We understand what families are going through, and we're there with targeted support when it's needed most. For these two families, that has meant practical help, emotional reassurance, and the belief that Liwsi and Mabli deserve every chance to play, learn and thrive.

- **Our Parent Guides** helped them secure vital financial support.
- **Our Library Service** provided sensory toys that spark joy and learning.
- **Our Sleep Advice Service** is helping them find rest.
- **And our Innovation Centre** is designing something extraordinary so the girls can explore the world together.

Gwenno, Mabli's mum, says:

"It's so rare to find a charity that not only offers practical help but also makes you feel truly seen, understood and supported. Finding Cerebra has been life-changing.

Mabli absolutely loves the fibre optic sensory tails from the toy library — we simply couldn't have afforded them ourselves. And the Innovation Centre designing a custom sit-ski seat for her means the world to us. It shows people believe in her potential."

Zoe, mum to Liwsi, shares:

"When Liwsi was diagnosed at just three weeks old with two rare chromosome disorders, our world fell apart. We felt so alone.

Cerebra has been there through challenges we never imagined we'd face. Their guides, their sleep support, their kindness — they make the impossible feel possible. We are so grateful."

For Mabli and Liwsi, Cerebra has meant hope, connection and the chance to discover a better life together.

For us, they represent what this year has been all about: standing beside families, believing in their children, and helping them go further.

Who We Are

Cerebra is a charity dedicated to helping children with brain conditions and the families who love them. We listen, we learn, and we stand alongside families as they navigate challenges that no one should face alone. Through expert advice, innovative design, legal support, research and practical help, we work to create a world where every child can discover, explore and thrive — and where families feel supported every step of the way.

Contents

Mabli & Liwsi: A Story of Connection, Courage and Possibility 1

A Message From Our CEO 3

1. Real Lives. Real Change 4

Asking for help should never cause harm 4
Twenty Years of Changing Lives 6
Imogen's Incredible Journey 7
What if no one ever had to ask, "Can I take part?" 8

2. Helping Families Go Further 10

Sleep Advice Service 10
Library Service 12
Research & Information 14
Legal Rights Service 16
Cerebra Innovation Centre 18

3. The Impact of Your Support 21

How we're funded 21
Together, We Go Further 22

A Message From Our CEO

A Year of Impact, Listening and Possibility

When I joined Cerebra at the end of 2024, I was immediately struck by the resilience of the families we support. In every conversation, I saw the extraordinary love that surrounds children with neurological conditions and the unwavering determination of families to create fulfilling, joyful lives for their children. I was reminded why Cerebra's work matters so deeply: no child should have to fight for the opportunity to thrive. We are here to walk alongside families through challenges and uncertainty, offering practical support, trusted expertise and, above all, hope.

2025 has been a year to be proud of. We have continued to strengthen and grow our services, helping more families access the support they need. Whether through our Innovation Centre, Legal Rights Service, Sleep Advice or other specialist services, I have been continually inspired by the dedication, creativity and compassion of our teams. Their work demonstrates the powerful difference that can be made when expertise is combined with genuine understanding.

This year also saw us begin an important conversation about Cerebra's future. Through our strategic review, we took the opportunity to step back, listen and learn. Families told us they want support that is easier to access, more connected and better able to respond to the realities of everyday life. We heard clearly that Cerebra has a vital role to play not only in helping families navigate challenges today, but also in challenging the barriers that can limit opportunities tomorrow. Time and again, we have seen how a small intervention can make a difficult day easier, and how the right support at the right moment can have a lasting impact on a child's life and future.

As we look ahead to 2026, I feel energised by the opportunities before us. We want to reach more children and families than ever before. We also want to raise our voice and campaign with greater urgency for change on the issues that matter most to our community. Our new strategy will see us investing further in our services, strengthening partnerships and creating new ways to connect with families earlier and more effectively. Exciting developments, including our partnership with Thread Fundraising and the introduction of our refreshed brand, will help us build on the strong foundations already in place.

Most importantly, we will continue to listen. The experiences of children, young people and families will remain at the heart of every decision we make and every ambition we pursue. Together, we will keep learning, growing and working towards a future where every child with a neurological condition is able to thrive, achieve their potential and feel fully included.

Thank you to our families, supporters, partners and colleagues for being part of this journey. Your commitment makes our work possible, and I can't wait to see what we can achieve together in the years ahead.

Jess Camburn-Rahmani

Cerebra CEO



1. Real Lives. Real Change

Some stories break your heart. Some lift it. Some remind you what's possible when a child is given the chance to shine. Together, they show the world as it is—and the world it could be. These are the stories that defined our year.

Asking for help should never cause harm

But for thousands of disabled children and their families, it does

In November 2025, Cerebra published a landmark report uncovering a devastating pattern we call Systems Generated Trauma (SGT) - the profound harm caused to disabled children and their families by the very public services designed to support them.

Based on the experiences of more than 1,200 parents, the findings are impossible to ignore.

What is Systems Generated Trauma?

SGT is the severe harm caused when families ask public services for help - and are met with suspicion, blame, bureaucracy and indifference instead.

Families describe it as more traumatic than domestic violence. More distressing than mental health crises. And it is happening every day.

> ***"Trauma has become baked into the system—almost an accepted by-product. It keeps service failures hidden and shifts the blame onto families, turning us into the 'problem to be managed."***
>
> ***"We are not the same people. We are broken and traumatised. We cannot trust anyone anymore."***
>
> ***"I feel like a fish in a goldfish bowl—everything I do is visible, judged, discussed. But no one ever asks what it feels like to live inside the glass."***

Leading the charge for change

Cerebra is demanding better. For every family

This is not just a report. It is a call to action.

Launched at Church House, Westminster in November 2025, the SGT report has already sparked vital conversations at the highest levels - with policymakers, practitioners and leaders across health, education and social care.

Cerebra is proud to be leading this agenda. Because we believe that no family should ever be broken by the systems that are supposed to hold them together.

"It is a painful truth that many disabled children and their families are being traumatised - not by illness, not by accident, but by the very public services that are meant to be their lifeline. Asking for help should never cause harm."

— Jess Camburn Rahmani, Cerebra CEO

"The failure of governments and public bodies to take purposeful remedial action when system defects are identified renders untenable any assertion of blamelessness. In social harm theory, inaction of this kind is best described as 'moral indifference'."

— Professor Luke Clements, Emeritus Professor of Law and Social Justice, University of Leeds (report author)

Cerebra demands:

1. Public services formally acknowledge that their systems cause trauma to disabled children and their families.
2. Policymakers, funders and academic institutions commit to urgent research to understand the scale of SGT and develop evidence-based solutions.
3. A fundamental shift from suspicion and blame to genuine partnership, trust and accountability with families.

"To policymakers, practitioners and leaders: this is your moment. Read this report. Let it move you. Let it challenge you. And then—let it change you. Because our children deserve better. And so do their families."

— Jess Camburn Rahmani, Cerebra CEO

Read the full report and join the campaign at *cerebra.org.uk/systems-generated-trauma*.

1,200+
Parents' voices shaped the report

1,053
Full report downloads

3,302
Campaign page visits

203
People signed up to support

Twenty Years of Changing Lives

Two decades. Two unstoppable teams

Twenty years ago, Cerebra made a promise: that no parent should face the night alone, and no child's potential should go unrealised for want of the right support. That promise is still at the heart of everything we do.

In two decades, we've supported thousands of families through sleep advice and innovative design.

On 4 December 2025, we gathered at the Royal Society to celebrate two decades of impact. Guests explored Innovation Centre products, met our Sleep professionals, and heard from disability rights activist Francesca Martinez and inspiring service users including Imogen Ashwell-Lewis.

"The enthusiasm and sheer pride in the room was unmistakable. The work this organisation does for children with brain conditions is nothing short of remarkable."

— Dani, Motability



Francesca Martinez at our 20 year celebration

The difference twenty years can make:

7,000+

families whose nights were transformed by our Sleep Service

3,000+

families whose children discovered new possibilities through products designed and created by our Innovation Centre

20 years

of expertise, passion and unstoppable commitment

Twenty years down. So much further still to go—together.

Imogen's Incredible Journey

If children can dream big, they can do big

When Imogen Ashwell-Lewis was born with cerebral palsy, some things must have seemed impossible.

They weren't.

Since 2013, Cerebra's Innovation Centre (CIC) team have walked–and climbed, and surfed, and raced–alongside Imogen every step of the way. Designing and creating bespoke equipment, free of charge, that has turned her biggest dreams into reality.

Imogen has:

- Conquered Pen y Fan
- Completed a triathlon
- Surfed the waves
- Gone horse riding

And she's just getting started.

At six, she climbed South Wales' highest peak in a specially-modified mountain bike - pulled and pushed by 40 supporters including army soldiers. That same year, she captained Team Imogen at the Swansea Triathlon, crossing the finish line surrounded by her CIC team, full of smiles.

Then came horse riding. Then a specially-adapted pink surfboard. Then, in December 2025, something perhaps most remarkable of all–Imogen took to the stage at the Royal Society in London, to share her story with the world.

"It's only thanks to the team at Cerebra that she's been able to have these adventures. There have been so many times she hasn't been able to join in. Being able to do things like this is a real boost to her confidence and self-esteem."

— Catherine, Imogen's mum

Imogen is part of the Cerebra family. Her story - and her smile - remind us every day why we do what we do.



Imogen at the top of Pen Y Fan

What if no one ever had to ask, "Can I take part?"

Our Relay Your Way proved it was possible

From 3rd June to 6th July, we ran, cycled, wheeled, surfed, danced and paddled 2,537km across the UK–from Glasgow to Windsor–carrying one baton and one powerful message: that every child deserves to be included, celebrated and given the chance to shine.

Relay Your Way was only possible because thousands of people came together with energy, generosity and joy. Over 33 days, supporters across the UK ran, wheeled, cycled, paddled, danced and cheered — helping us raise more than £60,000 for children with brain conditions.

To everyone who carried the baton, joined a stage, organised a local event, fundraised, donated or supported from the sidelines — thank you. Your enthusiasm turned a national relay into a movement for inclusion, possibility and pride.

You helped create something unforgettable. And you helped children with brain conditions go further.



Relay Your Way map from Glasgow to Windsor

2,537km
from Glasgow to Windsor

33 days
of inclusive adventure

15,000+
people carried the baton

40+
SEN schools visited

£60k+
raised for life-changing services

Every kilometre. Every smile. Every child

The moments that made Relay Your Way unforgettable

Every person who carried the baton had a story. Wolfe, who took the relay onto the water in a custom-built kayak designed by our Innovation Centre. Henry, who led cyclists along the Bridlington promenade with his CIC-made bike trailer - the same Henry whose ventilator backpack gives him freedom in the school playground every day.

And at the finish line in Windsor, eight-year-old twins Aleisha and Larissa carried the baton those final precious metres. Their presence at the finish line reminded us exactly why Relay Your Way exists - to make inclusion possible, joyful and real.

"So many things they can't do like other children… so it's important to show them they can still take part and excel."

— Liang, mum of Aleisha and Larissa

"Just those little things make a massive difference to both the boys being able to be included."

— Debbie, mum of Alfie and George

Highlight moments:

- Penalty shoot-out with Broxi Bear at Ibrox Stadium
- Fun run with political leaders at Scottish Parliament
- Baton on the pitch at Leeds Rhinos vs Warrington Wolves
- Sheffield Steelkings Para Ice Hockey Team take the baton onto the ice



Aleisha and Larissa carry the baton for the final stretch in Windsor



The finish line - 2,537km done

2. Helping Families Go Further

Whatever families are facing, we stand alongside them with practical support that brings relief, strength and the sense that they don't have to do it all alone.

Sleep Advice Service

Supporting families to rest, recover and go further

When a child can't sleep, no one sleeps. Children with brain conditions are far more likely to experience disrupted nights - difficulty falling asleep, frequent waking, not wanting to sleep alone, or rising long before the rest of the household is ready to face the day. Broken nights affect the whole family - health, relationships, and the ability to cope with everything else that comes with caring for a child with a brain condition.

Cerebra's Sleep Advice Service offers practical, personalized, evidence based support - from our Sleep Tips booklet and parent guides, to one-to-one sessions with specialist Sleep Advice Consultants and group workshops and seminars. We also offer two OCN-accredited training courses for professionals, ensuring that expert SEND sleep support reaches further than ever.

466
families supported in 2025

88%
of families reported improved knowledge and confidence in managing their child's sleep

43
professionals completed our sleep training

When families are exhausted and out of options, Cerebra's Sleep Service meets them where they are—and helps the whole family move forward together.

Riley's Story

Three-year-old Riley loves trains, diggers and farm machinery. He's happiest outside, absorbed in sensory activities or carefully organising his toys. But for Riley and his mum Natasha, nights had become a crisis. The two of them were surviving on just a couple of hours of sleep a night.

Riley's condition touched every part of family life. His sister had learned to communicate with him using non-verbal cues. Routines were built around his needs. Every day brought new challenges. But sleep - or the complete lack of it - was pushing the family to breaking point.

When Natasha contacted Cerebra's Sleep Service, everything began to change. Their Sleep Advice Consultant helped create a concrete sleep plan for Riley, and went far beyond that - sending books, links to other organisations and groups, and practical advice for parents navigating life with a child with additional needs. As other professionals became involved with Riley, Cerebra helped bring everyone together around a shared plan.

"It has made a HUGE difference. Without the help from Cerebra, I honestly don't know what I would have done. I so desperately needed help and without the help of their Sleep Team I genuinely don't think I would be in the position I am in now. Their knowledge is fantastic - I felt like they really understood and wanted to help, and that meant a lot to us as a family. Up until this point we had been left to go at it alone."

"You've lifted me up from some really low points of me feeling like a terrible mother for not being able to help or understand what is going on with my child."

— Natasha, Riley's mum



Riley wearing his heaphones with his sister

Library Service

The right book. The right toy. Right when you need it

Every book and toy in Cerebra's library has been specially chosen with children with brain conditions in mind. This isn't a generic collection - it's a carefully curated resource built around the real and varied needs of children and families who often struggle to find the right support elsewhere. Free to use, delivered to your door, and backed by a team who will help you find exactly what you need - our library exists to open doors, spark joy, and help families go further.

387

families borrowed books and toys from our Library in 2025

93%

said borrowing from Cerebra helped them better meet their child's needs

"So many services say they want to support families, but Cerebra actually gets it. You've designed something that works with the reality of our lives."

— Malky, parent

"Instead of labelling himself as 'stupid' or 'clumsy' he now uses the term dyspraxia to explain to others why he might find some things difficult, and to ask for help when he needs it instead of becoming frustrated or feeling ashamed."

— E.H., parent

With every book and toy specially selected for children with brain conditions, our Library brings practical help and moments of joy into family homes.

Finding Calm in Colour

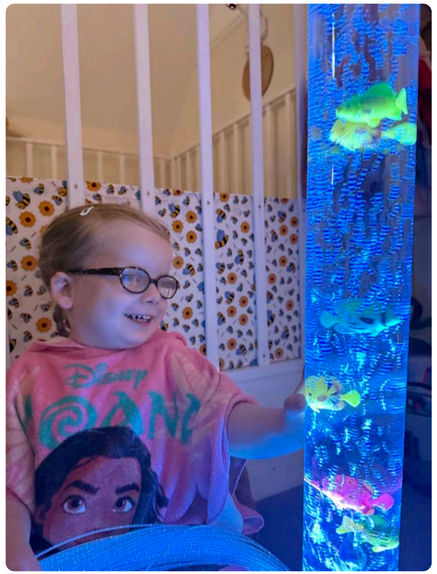
Three-and-a-half-year-old Dominique has Xia-Gibbs Syndrome, a rare genetic condition affecting her development, speech and mobility. Significant sensory processing difficulties mean that staying regulated, calm and engaged with the world around her can be a real challenge. For mum Kat, finding the right equipment felt almost impossible - either too expensive, or simply not the right fit for Dominique's unique needs.

Then she borrowed a fibre optic sensory lamp from Cerebra's Toy Library.

"The first time we set it up, Domi's eyes lit up in a way we hadn't seen in a long time. She immediately reached out to touch it, mesmerised as the colours shifted."

The lamp quickly became part of the family's daily routine - a calming aid before bedtime, a tool to help Dominique refocus after a difficult moment. Meltdowns reduced. Engagement improved. And something unexpected happened too - Dominique's little sister began sitting beside her, exploring the lights together.

"It's not just a piece of equipment - it's a bridge to calmer, happier moments for Domi and for us as a family."



Domi is enraptured by the fibre optic sensory lamp

Research & Information

When you don't know where to turn

When families need answers fast—about diagnosis, rights, education or support—our Research & Information team is there with clear, trusted guidance designed for real life. We translate complex medical, legal and practical information into accessible guides and factsheets that help families take confident action.

We earned the UK's PIF TICK quality mark for clear, accurate and reliable health information.

Your questions, answered

Our most reached-for guides and factsheets in 2025 were:

- Disability Living Allowance: 11,855
- Sleep Guide: 4,621
- Sleep Tips: 3,043
- Sensory Processing: 1,394
- Behaviour Checklist: 1,287

"Your DLA Guide was so helpful when filling in my daughter's forms — and we've now been awarded DLA. This will make a huge difference to our family." – *Parent*

*DLA is financial support for disabled children who need extra care or mobility help.



500,000

families visited our website in 2025



46,087

families received our regular Newsflash newsletter



30,844

times our guides were downloaded



72%

of viewers rated our guides as excellent

Webinars

In 2025, we delivered two national webinars offering research-led guidance on issues families face—supporting siblings, and understanding neurodiversity in communication and social abilities.

PIF TICK accreditation

The PIF TICK is the UK's assessed quality mark for reliable health information. It means our guides are created to strict standards—evidence-based, easy to understand and genuinely helpful for families.

The Cerebra Network for Neurodevelopmental Disorders

We renewed funding for the Cerebra Network, a pioneering collaboration improving understanding and support for children with complex or rare conditions. By developing better diagnostic tools and tailored interventions, the Network is turning cutting-edge research into real-world change.

The Network's researchers are improving how complex neurodevelopmental conditions are identified, understood and supported—especially for children with rare genetic syndromes. Their work is strengthening the evidence behind clinical assessment and interventions, helping professionals make earlier, clearer decisions and giving families access to support that truly fits their child's needs.

Learn more about the Cerebra Network at *cerebranetwork.com*.

Whatever questions families are facing, we make sure they never have to search for answers alone.

Legal Rights Service

When getting support feels like a battle

Public bodies in the UK have a duty to provide health, social care and other services for disabled children. Our Legal Rights Service helps families of children with brain conditions get the right support when they're having difficulty accessing the services they're entitled to.

We can help with issues including assessments, care and support plans, CAMHS, direct payments, school transport, short breaks, carers' rights and more.

97

families supported by our Legal Rights Service

"I am eternally grateful for the time, support and guidance I've received from Cerebra's Legal Rights Team. Without their help, I would still be in a state of despair. They helped guide me through complicated SEN matters with the LA. With their advice, I've been able to navigate the system now and they have helped build my confidence. For context, I suffered a nervous breakdown in November 2025 contributed by systematic failures from the LA towards my disabled little boy. I cannot thank Cerebra enough - it has saved my life."

"The team have offered absolutely nothing but understanding, clear guidance and support. I feel like at every hurdle (and there were so many) that they have provided support and knowledge over and above anything I felt would be possible. It took us 6 years to "fight" the battle of wrong doings and I would absolutely not been able to do it without the support from Cerebra's Legal Rights Service."

"I sought support with integrated care board hours and social care hours/decision making. With regards to the extensive legal paperwork, government policies etc I have neither the capacity nor the technical knowledge to appropriately approach this. The team were professional, clear in taking my initial call and thereafter have been timely, concise and consistent. Their support has also made me feel confident in trying to advocate for my child, who is severely disabled. I am grateful for the support and thankful that vulnerable families like mine have an organisation like CEREBRA to help and support them when they have few other choices, if any."

900,000 children

A basic human right. Denied

Our Legal Rights team has campaigned for appropriate continence care for disabled children. Our landmark research, published in February 2025, exposes a devastating failure of care — and demands urgent action across England, Scotland and Wales.

The findings are stark.

900,000 children and young people in the UK experience bladder and bowel dysfunction. Disabled children are far more likely to need continence support — yet our research found widespread failure: inaccessible services, discriminatory criteria, and a lack of national guidance.

Families are left to cope alone, at enormous cost to their health, finances and dignity.

"We have been made to spend so much money just to let our son go to the toilet hygienically and with dignity. It has had a huge impact on our lives."

"They didn't seem to know what to do with an autistic child with continence issues. I don't have the time or energy to fight for it."

"It beggars belief that such a basic human right is systematically being denied to children who are already some of the most discriminated against in our society. These children deserve much better."

— Jess Camburn Rahmani, Cerebra CEO

Read the full report at cerebra.org.uk/continence-report.

900,000

children and young people affected by bladder and bowel dysfunction in the UK

2,308

campaign page visits

356

downloads of the full report

772

people signed up to support our campaign

This is what our Legal Rights work is for: turning evidence into action, and action into change. Since 2025, drawing on our research and campaign, we've been contributing to the review of nationally adopted guidance for continence provision in the UK, calling for improvements to make it clearer and fairer for families.

Cerebra Innovation Centre

Bright ideas for brilliant kids

The Cerebra Innovation Centre (CIC) designs and builds innovative, bespoke products that help disabled children discover and engage with the world around them.

From iPad cases to rocking horses, Oxy-Gems to surfboards, our award-winning team takes requests from parents, carers and professionals and turns them into beautiful, inventive and above all fun solutions, tailored to the specific needs of each child.

We believe our products should be desirable and exciting, not just functional. Beautiful as well as brilliant. Designed to promote social inclusion, peer acceptance and the chance to join in - because every child deserves that.



James the Scoot horse is a much-loved member of the family



Arthur loves his fish tank



Hydrobeatz was a hit in a music class

122

families supported by the Innovation Centre in 2025

20

brand new bespoke products designed and built in 2025

Skating, confidence and the power of brilliant design

Twelve year old Ollie desperately wanted to try skateboarding — but balance challenges linked to Down Syndrome meant a standard board wasn't safe. His mum, Jilly, reached out to the Cerebra Innovation Centre hoping for a way for him to join in, not sit on the sidelines.

The team designed a bespoke board with two handles, giving Ollie the stability he needed to push off and skate independently. It worked — and his confidence soared.

Seeing how much skating meant to him, the team went further. They developed a world first adaptation for park skating: a harness system suspended from a gantry at Exist Skatepark in Swansea, giving children with reduced mobility the freedom to move safely through the ramps alongside their peers.

For Ollie, who has faced profound anxiety since the loss of his sister, the impact has been transformative. Skating has become a way to connect, socialise and rediscover joy.

"We wanted Ollie to skate with other children, on equipment that looks the same as theirs."

– Dr Ross Head, CIC Lead

"I'm just so grateful. They've given Ollie something he never thought he'd have."

– Jilly, Ollie's mum



Ollie tries out the new skateboard and harness system

The Cerebra Innovation Centre doesn't just build products—it builds possibilities, opening doors for children who are too often told the door is closed. Together, our world class innovation team turn brilliant design into joyful possibility—giving more children the chance to join in, explore and thrive.

In 2025 our design impact grew even further - welcome to the family, MERU!

Two teams. One mission. Bigger impact than ever

The very end of 2025 brought exciting news for Cerebra and for disabled children across the UK. We welcomed the world-class team from specialist charity MERU into the Cerebra family - combining two organisations with a shared belief that brilliant design can bring joy, independence and adventure to every child.

Together, we are stronger than ever.

MERU's expertise perfectly complements our own Innovation Centre - adding their switch-adapted toy service and the remarkable Bugzi, a mini powered wheelchair that gives children as young as one the freedom to explore their world independently. Combined with Cerebra's bespoke design work our expanded team can now reach and support more families than ever before.

The MERU team are now fully part of Cerebra, operating under one name. they continue to be based in Surrey in a brand-new workshop, ready to grow.

"Cerebra and MERU share a steadfast belief that good design can bring inclusion, joy and adventure to disabled children and their families. We are stronger together."

— Jess Camburn Rahmani, Cerebra CEO

"Our missions are so closely aligned that this feels like a natural evolution. Joining the Cerebra family allows us to focus entirely on what we do best: innovating for the children who need us most."

— Tracey Doherty, Sales & Product Lead, MERU



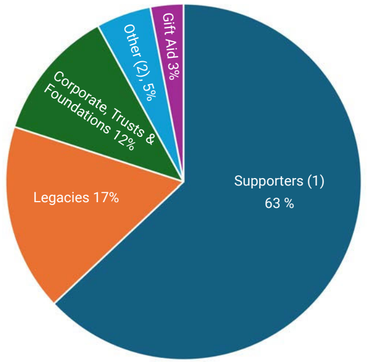
Cerebra and MERU Teams

3. The Impact of Your Support

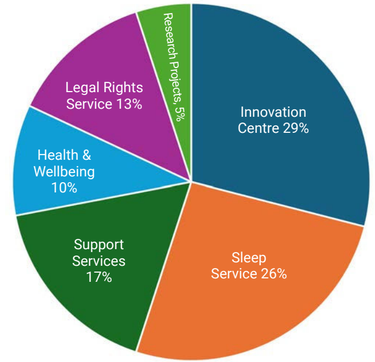
Every pound that reaches Cerebra goes directly towards changing the lives of children with brain conditions — and the families who love them. From sleep advice to life-changing legal support, from custom-built equipment to cutting-edge research, here's how your generosity made a difference in 2025.

How we're funded

Summary of our income streams



Charitable Expenditure 2025



Where your money goes

Your generosity funded the services and support that families relied on most—from rest and reassurance to advocacy, innovation and national change.*

(1) Lottery, Donation Card, Raffle, Regular Giving

(2) CIC Royalties, Sleep Course, Bank Interest

*A note on our finances - Cerebra is committed to transparency. Full audited accounts for 2025 are available at cerebra.org.uk or on request from our team. Registered Charity No. 1089812

Every contribution—however large or small—makes our work possible. Thank you.

Together, We Go Further

Every achievement in this report—every child supported, every family strengthened, every barrier challenged—was made possible by the people who stand with us. The trusts and foundations who fund our work, the legal teams who champion children's rights, the businesses who partner with us, and the many individuals and community groups who give their time, energy and generosity. Together, this extraordinary network of supporters helps families find strength, hope and the chance to go further.

Thread Fundraising Accelerator Partner 2026

Thread Fundraising selected Cerebra as the charity partner for their inaugural Accelerator Programme—a transformational year-long collaboration providing over £100,000 of pro bono strategic fundraising support. Their expertise will help us strengthen supporter engagement, grow income and accelerate our impact in 2026 and beyond.

This partnership marks a pivotal moment in Cerebra's growth—and we are deeply grateful to Thread for their belief in our mission.

THREAD Fundraising

Thanks to everyone who supported Team Cerebra!



Leisuretime by Toureasy Day Trip: "A fantastic way to celebrate St David's Day and support a wonderful cause."



Karoline Klage completed the London Landmarks Half Marathon: "It has been a joy to run for Cerebra".



Twelve incredible volunteers from Valero took on a four-day, 480km cycling challenge — riding from the Pembroke refinery to Valero's London office: "A truly inspiring challenge… cheered on by colleagues every step of the way."

To everyone who stood with us in 2025: thank you. Together, we are building a world where every child with a brain condition can go further, dream bigger and be fully included.

Be A Part Of What Comes Next

Every step forward for Cerebra starts with people like you. Whether you take on a challenge, fundraise, volunteer or support our work, you help create the breakthroughs families tell us they need most.

If you'd like to be part of what comes next, choose the path that suits you best — take on a movement challenge or explore all the ways to get involved and support Cerebra.

Cerebra
The MacGregor Office Suite
Jolly Tar Lane, Carmarthen
SA31 3LW

01267 611711

cerebra.org.uk

Registered Charity no. 1089812 Company no. 4336208

Registered with FUNDRAISING REGULATOR

amrc
ASSOCIATION OF MEDICAL RESEARCH CHARITIES

Trusted Information Creator
Patient Information Forum

OCN London