SHINING A LIGHT ON EPILEPSY
INTERNATIONAL EPILEPSY CONGRESS 2019

From every corner of the world, delegates including Chief Executive Graeme Shears attended the 33rd International Epilepsy Congress in Bangkok, Thailand. Held over five days in June, it was an exciting event with a host of interesting and relevant subjects and world class speakers sharing their knowledge about epilepsy.

There were many highlights including the launch of the World Health Organisation report, “Epilepsy: a public health imperative”. This was the result of great collaboration between the International League against Epilepsy (ILAE) and the International Bureau for Epilepsy (IBE). The Epilepsy Foundation contributed to this most important report and was also an Expert Reviewer. If you would like to read the report visit: www.ibe-epilepsy.org/global-report-epilepsy-a-public-health-imperative or contact the Foundation.

Another wonderful highlight was the Golden Light Award presentation where our own shining star, Scarlett Paige, was one of eight inspiring international young people with epilepsy to be recognized for her contribution to improving the lives of people with epilepsy.

Meetings of the IBE leadership focused on practical projects that would build the capability of epilepsy organisations around the world. Our congratulations to Gweme who was formally elected as a Board member of the IBE and Chair of the capacity building project.

I was diagnosed with focal and generalized convulsive seizures in my early 20s. I was about to embark on my lifelong dream to be an epileptic focal convulsive seizures. Being able to have the opportunity to find a cure for epilepsy is a great success and has had by all.

Scarlett Paige
Epilepsy Support Worker

AND THE GOLDEN LIGHT AWARD GOES TO...

Scarlett Paige has lived with epilepsy since 2006. In June, she was awarded a Golden Light Award in recognition of her work to help people with epilepsy. This is her story.

“My epilepsy journey started in January 2006. I was 18 and had the next two years of my life planned out. I was about to embark on my lifelong dream to become a student. Seizure activity, anti-epilepsy medications, surgical intervention and lifestyles had been part of my life for 10 years.

Once my seizures were under control I started volunteering with the Foundation. In March this year the Foundation nominated me as a shining light for people with epilepsy, due to my community service in Victoria and Tasmania. The nomination covers 11 countries in the Western Pacific region.

In April I received an email congratulating me on my success that I, along with 7 others from around the world, would visit Bangkok to attend our awards at the 33rd International Congress in late June.

On the opening ceremony in front of 2,000 people we were presented with the Golden Light Award. Over the next five days we attended many symposiums from the social impact of epilepsy to the importance of medication. Being able to have the opportunity to meet others and come together in a positive way is something I have never experienced before. Our award was presented to us by the wider public.

In 2021 I am hoping to travel to the next International Congress to discover how my fellow golden lights have been in their quest to end the stigma of epilepsy.

Scarlett Paige (third from right) is one of eight 2018 Golden Light Award recipients.

EPILEPSY SMART SCHOOLS IS NOW NATIONAL

Epilepsy Foundation 367 Canterbury Road Earlwood NSW 2206 General Enquiries: 1300 437 453 Information & Support: 1300 761 487 Education & Training Enquiries: edutra@epilepsyfoundation.org.au Office opening hours: Monday to Friday, 9.30am to 4.30pm Work: epilepsyfoundation.org.au

Epilepsy Foundation of Australia is accredited under the Australian Business Group’s Business Resource Certification System and has met the requirements of the National Australian Accreditation Scheme. We are registered with the ACNC as a Registered Charity. Donors & Supporters: 1300 761 487 Donors & Supporters: 1300 761 487

Around 1 in 200 Australian students attending school have epilepsy. The mean school class has one or more students with epilepsy amongst their population.

Living with epilepsy can have significant impacts on a student’s education. Seizure activity, anti-epilepsy medications, medical attendance appointments and schools. The stigma still associated with epilepsy can have a negative effect on their school performance and social relationships.

Consequently, children miss days at school behind in their schoolwork and can feel isolated.

The Epilepsy Foundation is working hard to change this, which why we developed the Epilepsy Smart Schools program. This program is aimed at a positive impact on education so as to create a culture of change through a better understanding of epilepsy.

Epilepsy Smart School. To learn more visit: www.epilepsysmartschools.org.au

No one should go it alone

No one should go it alone

No one should go it alone

No one should go it alone

No one should go it alone

No one should go it alone

WALK FOR THE ONE YOU LOVE
Walk for Epilepsy 2019

Get your family and friends together, and join other caring Victorians who will be walking for those they love or in memory of a loved one. There was a wonderful community vibe, as people came together to share stories and their personal journeys of living with epilepsy.

This year our goal is to break our record and get 1,000 people joining our Walk for Epilepsy. And we can do this with you!

Get your family and friends together and join other caring Victorians who will be walking for those they love in memory of a loved one. Choose to walk a leisurely 3.5 or 6 km course around the beautiful Princess Park in Carlton.

The walking track is fully accessible to mobility scooters, wheelchairs and prams.

Enjoy free entertainment from 9:20am before the Walk commences at 11am. A coffee van, sausage sizzle and water will be available to purchase. Visit www.walkforepilepsy.com.au to find out more.

Register today at www.walkforepilepsy.com.au

Walk for Epilepsy 2019

SUNDAY, 20 OCTOBER 2019
PRINCES PARK, CARLTON

Enjoy free entertainment from 9:20am before the Walk commences at 11am. A coffee van, sausage sizzle and water will be available to purchase. Visit www.walkforepilepsy.com.au to find out more.

Register today at www.walkforepilepsy.com.au
TOGETHER WE ARE MAKING A DIFFERENCE
At the start of a new financial year, I once again reflect on the impact that your giving has made to people living with epilepsy.

In my 12 years as Chief Executive, I have seen thousands of caring Australians make an incredible difference in the lives of people living with epilepsy. The stories of support, kindness and understanding that our clients will continue to need access to our services and programs makes us all feel very proud to be supported by people like you.

There is so much more we know we can do for people with epilepsy. We are already working with our current clients to develop specific training and awareness programs for staff and for service providers for their support of these people.

In another superb effort, will Mitchell of Maggie Wagge raised $14,000 through a golf day and talent quest. Congratulations and Thanks to all of our community fundraising champions and we look forward to another fabulous Purple Day in 2020!

Thank you for all that you do to support people with epilepsy. I hope you enjoy this update on our issue of fringe and look forward to your continued support in 2020/20.

Heshan and Tara Mendis raised an incredible $40,000 through two fundraising efforts. Thank you Bunnings!

I am truly grateful to you and every one of you. You are the reason around 4,000 people living with epilepsy are supported through over 6,000 interactions, and enabled us to raise more than $2,000,000 to support people with epilepsy.

You are the reason that people with epilepsy never feel that they are alone.

I am truly grateful to each and every one of you. You have made an incredible difference in so many people’s lives.

I hope you enjoy this update on the impact that your giving has made to people living with epilepsy.

Thank you for all that you do to support people with epilepsy.

To find out more about our new store in Berwick, visit www.epilepsyfoundation.org.au/cure-epilepsy

Visit Bianca and hundreds of caring Victorians on Sunday 20 October, 11:00am at Princes Park in Carlton for what will be a fun and wonderful day for people of all ages. Together we can show our support for those who love who have epilepsy and let them know they are not alone.

Join Bianco and hundreds of caring Victorians on Sunday 20 October, 12:00pm at Princes Park in Carlton for what will be a fun and wonderful day for people of all ages.

TOGETHER WE ARE MAKING A DIFFERENCE
You can be a ‘Walk Hero’ by letting your family and friends know they can sponsor you.

You can also become a ‘Walk Hero’ by letting your family and friends know they can sponsor you.

You can raise funds by setting up a fundraising page online and sharing it with your family and friends.

You can also become a ‘Walk Hero’ by letting your family and friends know they can sponsor you.

YOU RAISED VITAL FUNDS. THANK YOU!

Over 700 people came together to walk for those they love, care about, are friends with or who have epilepsy.

TOGETHER WE ARE MAKING A DIFFERENCE
You can get involved and walk for those you love. Register today or visit www.walkforepilepsy.com.au

NEW MEDICAL RESEARCH IS UNDERWAY

Last year the Australian Government announced that they would invest $2 million through the Epilepsy Research Fund to help find a cure for rare genetic epilepsy disorders.

This investment has helped launch the inaugural funding round of the Epilepsy Research Fund, with a particular focus on medical research into the genes of the 80 per cent of people with epilepsy who are undiagnosed. This puts us one step closer to finding a cure for epilepsy.

The very first project to be supported through this Fund is currently underway at the Florey Institute of Neuroscience and Mental Health, examining treatment options for epilepsy caused by mutations in the Synapsin gene. Normally, the Synapsin gene provides instructions for proteins that can play an important role in the function of nerve cells in the brain. These nerve cells act as the “wiring” in the circuitry of the brain, and are critical for learning and memory.

The Epilepsy Foundation looks forward to supporting more exciting and innovative research that will make a difference for people living with epilepsy.

If you would like to contribute to finding a cure for epilepsy, visit www.epilepsyfoundation.org.au/cure-epilepsy

A LITTLE TREASURE WAITING TO BE EXPLORED!

Our new store in Berwick is now open and waiting to be explored by you.

There is a beautiful selection of bric-a-brac, from the old and treasured to the retro and vintage for the building interior decor including a chandelier that makes everything in the store look that much more special.

It’s a new wardrobe you’re looking for but have ladies, men’s and children’s clothes and accessories for the whole family. Newly opened, there are three fitting rooms waiting for you. And for the avid reader, there’s a great selection of books for you to choose from.

Every store is open because of our committed volunteers who give their time, donate their vehicles and equipment and who play an essential role in helping people with epilepsy live better. Together we can show our support for those who love who have epilepsy.

The Foundation thanks each and every one of you for getting involved, increasing awareness and raising essential funds. Special thanks to Giselle Broom whose son has epilepsy and Melissa Lerdal whose son has epilepsy. Both raised well over $10,000 each in support of programs and services to help people with epilepsy.

Bianca and her family will be walking for her husband Adam. Adam has generalised tonic-clonic seizures. We try to live as normal a life as possible. We try to push the fact that he has seizures out of our minds so as a family we can get on with life. Our extended family didn’t realise the impact of Adam’s seizures.

They didn’t understand what was happening to Adam and we readily the seizures were effecting him. They didn’t understand why we would leave early from functions, or why we wouldn’t even go into元境

That was until they saw Adam having a seizure.

Last year participated in the Walk for Epilepsy in New South Wales with magnificent fundraising efforts from the community. This includes a selection of Bunnings stores that got behind our Purple Day efforts. Thank you Bunnings.

Creating a Greater Understanding of Epilepsy with Victoria Police

There has long been a need for specific training for police and other first responders to develop a greater understanding of epilepsy.

Walk for Epilepsy and Show Your Support

Sunday 20 October, Princes Park Carlton, 11.00am start

New Medical Research Is Underway

The very first project to be supported through this Fund is currently underway at the Florey Institute of Neuroscience and Mental Health, examining treatment options for epilepsy caused by mutations in the Synapsin gene. Normally, the Synapsin gene provides instructions for proteins that can play an important role in the function of nerve cells in the brain. These nerve cells act as the “wiring” in the circuitry of the brain, and are critical for learning and memory.

The Epilepsy Foundation looks forward to supporting more exciting and innovative research that will make a difference for people living with epilepsy.

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The Foundation thanks each and every one of you for getting involved, increasing awareness and raising essential funds. Special thanks to Giselle Broom whose son has epilepsy and Melissa Lerdal whose son has epilepsy. Both raised well over $10,000 each in support of programs and services to help people with epilepsy.

You can get involved and walk for those you love. Register today or visit www.walkforepilepsy.com.au

You can be a ‘Walk Hero’ by letting your family and friends know they can sponsor you.

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You can raise funds by setting up a fundraising page online and sharing it with your family and friends.

You can also become a ‘Walk Hero’ by letting your family and friends know they can sponsor you.

YOU RAISED VITAL FUNDS. THANK YOU!

Over 700 people came together to walk for those they love, care about, are friends with or who have epilepsy.

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TOGETHER WE ARE MAKING A DIFFERENCE
At the start of a new financial year, I once again reflect on the impact that your generosity and support made to people living with epilepsy.

In my 12 years as Chief Executive, I have seen thousands of caring people come together to help ensure that those living with epilepsy never feel that they are alone. Their support is the difference between hope and despair for so many individuals who are affected by epilepsy.

I am truly grateful to each and every one of you. You are the reason around $3,000,000 has been supported through over 6,000 interactions, and enabled us to reach more than 3,000 people to help them understand and manage epilepsy better.

I must also thank everyone in our support network. Your ongoing support is critical to our work.

Thank you for helping us to help everyone; thank you for being such a wonderful community.

Heshan and Tara Mendis raised an amazing $60,000 to help the Epilepsy Foundation. They thanked their network of friends and family for all their contributions and support.

There are so many people who we know we can do so for people with epilepsy. We cannot do our work without the support of our current clients and will continue to need access to our staff, services and programs.

It is important we continue to be there for them and for the many Australians who will be diagnosed with epilepsy this year.

The essence of all we exist is to ensure that every person, no matter their age, gender, ethnicity, location or beliefs feel that they are not alone with their epilepsy. To this end and we are now working to ensure that we continue to be available right across Australia through our Epilepsy Australia service partners. Your ongoing support makes this possible.

Thank you for all that you do to support people with epilepsy. I enjoy your stories of progress and look forward to your continued support in 2019/20.

Heshan Mendis
Chief Executive Officer

Creating a Greater Understanding of Epilepsy with Victoria Police

There has been a long need for specific training for police and other first responders to help them gain a greater understanding of epilepsy.

Work has commenced with Victoria Police to develop specific training and awareness programs for these first responders.

In another superb effort, Will Mitchell of Wiggles Wiggles raised selection of Bunnings so that get behind our Purple Day efforts. Thank you Bunnings.

Heshan and Tara Mendis raised an incredible $60,000 through two events – the Mukula Racing “Car Show & Shine” and ‘Beat 2 Beat Epilepsy’ Fundraiser.

Walk for Epilepsy and Show your Support

Sunday 20 October, Princes Park Carlton, 11.00am start

One million Australians will have epilepsy in their lifetime. They are your family, your friends, your community.

Many people with epilepsy experience loneliness, anxiety and isolation. As they are coming to terms with the diagnosis and learning how to better manage their epilepsy, people with epilepsy need to be completely supported.

Although epilepsy is the most common of brain disorders, 90% of Australians do not have access to epilepsy support they need.

The epilepsy Foundation exists so that no one living with epilepsy ever has to go it alone. Our work focuses on helping children with epilepsy get a good education, empowering people to fully participate in the workplace and ensuring they feel safe and connected in the community.

Join the Walk for Epilepsy and show your support for people living with epilepsy or walk in memory of someone. Bianca and her family will be walking for her husband Adam. Adam has generalised tonic clonic seizures. We try to live as normal a life as we possibly can.

They didn’t understand why he has seizures out of no reason as a family we can get on with our life. Our extended family didn’t realise the impact of Adam’s seizures.

They didn’t understand what was happening to Adam and we really the seizures were not affecting him. They didn’t understand why we would leave early from functions, or why we wouldn’t even go to events anymore.

That was until they saw Adam having a seizure.

Last year participated in the Walk. It was a terrible day for us as a family and our local children. We couldn’t do this.

But this year we’re getting as much of our families involved and get everyone together.

I want Adam to know that he is fully supported and loved by all of us, that he is not alone in his epilepsy, that we are there for him as a family.

You can also become a ‘Walk Hero’ by downloading your family and friends can exercise you. Every dollar you raise will support people with epilepsy.

The Foundation thanks each and every one of you for getting involved, increasing awareness and raising essential funds. Special thanks to Toonie Train, being used by Epilepsy Australia, which has now raised over $300,000.

You can get involved and walk for those you love. Register today – visit: www.walkforepilepsy.com.au

NEW MEDICAL RESEARCH IS UNDERWAY

The very first project to be supported through this Fund is currently underway at the Florey Institute of Neuroscience and Mental Health, examining treatment options for epilepsy caused by mutations in the SynGAP gene. Normally, the SynGAP gene provides instructions for building a protein, called SynGAP. Some people have mutations in their SynGAP gene that make the protein unable to function properly, which results in many electrical signals in the brain. These nerve cells act as the “wiring” in the circuitry of the brain, and are critical for learning and memory.

The Epilepsy Foundation looks forward to supporting more exciting and innovative research that will make a difference for people living with epilepsy.

If you would like to contribute to finding a cure for epilepsy go to: www.epilepsyfoundation.org.au/cure-epilepsy

TOGETHER WE ARE MAKING A DIFFERENCE

A LITTLE TREASURE WAITING TO BE EXPLORED!

Our new store in Berwick is now open and waiting to be explored by you.

There is a beautiful selection of bricks, from the old and treasured in the mini and vintage for the budding interior designer, including a chandelier that makes everything in the store look so much more special.

You can get involved and walk for those you love. Register today – visit: www.walkforepilepsy.com.au

Walk for Epilepsy Op Shop, 25 Blackburn Square, Oakleigh South, Monday to Friday 9:30am-4:30pm; Saturday 9:00am-1:00pm.

THANK YOU!

Over 700 people came together to walk for those they feel a special connection and love. They walked, ran or shared heart-warming stories, they all in support of those who live with epilepsy.

The Foundation thanks each and every one of you for getting involved, increasing awareness and raising essential funds. Special thanks to Toonie Train, being used by Epilepsy Australia, which has now raised over $300,000 and is in support of programs and services to help people with epilepsy.

You can get involved and walk for those you love. Register today – visit: www.walkforepilepsy.com.au

YOU CAME, YOU WALKED, AND YOU RAISED VITAL FUNDS. THANK YOU!

You can get involved and walk for those you love. Register today – visit: www.walkforepilepsy.com.au

NEW MEDICAL RESEARCH IS UNDERWAY

Last year the Australian Government announced that they would invest $2 million through the Epilepsy Foundation to help find a cure for rare genetic epilepsy disorders.

This investment has helped launch the inaugural round of the SynGAP Research Fund, with a particular focus on medical research into the genetics of epilepsy to develop treatments and discover new outcomes for the benefit of people living with epilepsy in the future.

The very first project to be supported through this Fund is currently underway at the Florey Institute of Neuroscience and Mental Health, examining treatment options for epilepsy caused by mutations in the SynGAP gene. Normally, the SynGAP gene provides instructions for building a protein, called SynGAP. Some people have mutations in their SynGAP gene that make the protein unable to function properly, which results in many electrical signals in the brain. These nerve cells act as the “wiring” in the circuitry of the brain, and are critical for learning and memory.

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TOGETHER WE ARE MAKING A DIFFERENCE
At the start of a new financial year, I once again reflect on the impact that your generous contributions made to people living with epilepsy.

In my 12 years as Chief Executive, I have seen thousands of caring individuals come together in ongoing acts of generosity and support. Your vibrant community of clients will continue to need access to our programs, resources and programs. It is important we continue to be there for them and for the many Australians who will be diagnosed with epilepsy this year.

Many people with epilepsy experience loneliness, anxiety and isolation. As they are coming to terms with the diagnosis and learning how to better manage their epilepsy, people with epilepsy need to be completely supported.

Although epilepsy is the most common of brain disorders, 80% of Australians do not have access to epilepsy support needed.
The Epilepsy Foundation exists to ensure that no one living with epilepsy ever has to go it alone. Our work focuses on helping children with epilepsy get a good education, empowering people to fully participate in the workplace and ensuring they feel safe and connected in the community.

Join the Walk for Epilepsy and show your support for people living with epilepsy or walk in memory of someone special. Bianca and her family will be walking for her husband Adam. “Adam has generated tons of classic seizures. We try to live a normal life as we possibly can. We try to push the fact that he has seizures out of our mind so as a family we can get on with life.”

Thank you for all that you do to make this possible.

I am truly grateful to each and every one of you.

I am truly grateful to each and every one of you.

YOU CAN AND DID GO PURPLE FOR PURPLE DAY 2019
On Tuesday 26 March, people and communities came together to raise vital funds and much needed awareness of epilepsy by wearing purple, hosting events or going to the Epilepsy Foundation.

Two significant and popular Melbourne landmarks showed their support for Purple Day. The Bolte Bridge and Melbourne Star were illuminated in resplendent purple. We thank Transtar and Melbourne Star for their wonderful support.

This year 28 registered community events were held in Victoria and New South Wales with magnificently generous fundraising efforts from the community. This includes a weekend of Bunnings sales that got behind our Purple Day efforts. Thank you Bunnings.

Thank you to everyone who participated in the Purple Day fundraising campaign.

I am truly grateful to each and every one of you. I am truly grateful to each and every one of you.

CAREERING A GREATER UNDERSTANDING OF EPILEPSY WITH VICTORIA POLICE
There has long been a need for specific training for police and other first responders to develop a greater understanding of epilepsy.

Work has commenced with Victoria Police to develop specific training and awareness programs for that membership. The first resource they will access to begin training is the Fact Sheet that will be shared across all VicPD communication channels.

further work is occurring with VicPD in order to have epilepsy awareness and requirements embedded in their training and education frameworks.
On Saturday 22 June 2019, the Epilepsy Foundation had the privilege of hosting 11 wonderful families to enjoy a day of planned activities, to connect with others in similar situations, to ask questions and to share their journeys with one another.

With 41 children and 34 adults, everyone was able to participate in a range of fun activities. This included the making of slime, being introduced to some native wildlife, balloon designs, face painting and photos from the photo booth to remember the day.

The Family Day was supported by the Epilepsy Support worker team and a strong group of faithful volunteers, who ensured the event was a great success and fun had by all.

### No one with epilepsy should go it alone.

**Walk for Epilepsy 2019**

Last year’s inaugural Walk for Epilepsy was a huge success with over 700 people walking, running or stilting for those they care about who are living with epilepsy, or in memory of a loved one. There was a wonderful community vibe, as people came together to share stories and their personal journeys of living with epilepsy.

This year our goal is to break our record and get 1,000 people joining our Walk for Epilepsy. And we can do this with you!

Get your family and friends together and join other caring Victorians who will be walking for those they love or in memory of a loved one. Choose to walk a 5km, 10km or 15km course around the beautiful Princess Park in Carlton.

The walking track is fully accessible to mobility scooters, wheelchairs and prams.

Enjoy free entertainment from 9:30am while the Walk commences at 11am. A coffee van, sausage sizzle and water will be available! The Epilepsy Foundation’s Education & Training Enquiries will display your own personal fundraising story, show you a pic or two, and motivate others to join in. All funds raised will help provide support for people living with epilepsy.

No one with epilepsy should go it alone. Your participation in this annual event will show those with epilepsy that they are not alone by your side.

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### Walk with Me

Sunday, 20 October 2019 at Princes Park, Carlton

Get your family and friends together, and join other caring Victorians walking for those they love. Registrations and Check In from 9am. Walk starts at 11am.

Register today at www.walkforepilepsy.com.au
SHINING A LIGHT ON EPILEPSY
INTERNATIONAL EPILEPSY CONGRESS 2019

From every corner of the world, delegates including Chief Executive Graeme Shears attended the 33rd International Epilepsy Congress in Bangkok, Thailand. Held on the 29th of June, it was an exciting event with a host of interesting and relevant subjects and world class speakers sharing their knowledge about epilepsy.

There were many highlights including the launch of the World Health Organisation report, “Epilepsy: a public health imperative”. This was the result of great collaboration between the International League against Epilepsy (ILE) and the International Bureau for Epilepsy (IBE). The Epilepsy Foundation contributed to this most important report and was also an Expert Reviewer. If you would like to read the report visit www.ibe-epilepsy.org/global-report-epilepsy-a-public-health-imperative or contact the Foundation.

Another wonderful highlight was the Golden Light Award presentation where our own shining star, Scarlett Paige, was one of eight inspiring international young people with epilepsy to be recognized for her contribution to improvements in the lives of people with epilepsy.

Meetings of the IBE leadership focused on practical projects that would build the capability of epilepsy organizations around the world. Our congratulations to Greeme who was formally elected as a Board member of the IBE and Chair of the capacity building project.

A FUN AND ACTIVE FAMILY DAY OUT

On Saturday 22 June 2019, the Epilepsy Foundation had the privilege of hosting 21 wonderful families to enjoy a day of planned activities, to connect with others in similar situations, to ask questions and to share their journeys with others.

With 41 children and 34 adults, everyone was able to participate in a range of fun activities. This included the making of slime, introduced to some of our first-time participants, tea passtries and a variety of different coloured balls, the art of balancing a feather on your arm, juggling, fabulous balloon designs, face painting and photos from the photobooth to remember the day.

The Family Day was supported by the Epilepsy Support worker team and in particular, a group of fabulous volunteers, who ensured the event was a great success and had fun by all.

Scarlett Paige
Epilepsy Support Worker

Scarlett Paige (third from right) is one of eight 2019 Golden Light Award recipients.

No one should have to go it alone. No one should face epilepsy alone. You are not alone by your side. Enjoy free entertainment from 9:20 am before the Walk commences at 11 am. A coffee van, sausage sizzle and water will be available for you to enjoy. Enjoy the head start for this fun-filled day, and join ‘Walk for the One You Love’.

Walk for Epilepsy Foundation.

No one should go it alone
If you are right by their side.

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Enjoy free entertainment from 9:20am before the Walk commences at 11 am. A coffee van, sausage sizzle and water will be available for you to enjoy. Enjoy the head start for this fun-filled day, and join ‘Walk for the One You Love’.

Walk for Epilepsy Foundation.

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