

How we create an effective member magazine

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What is Your Autism Magazine?

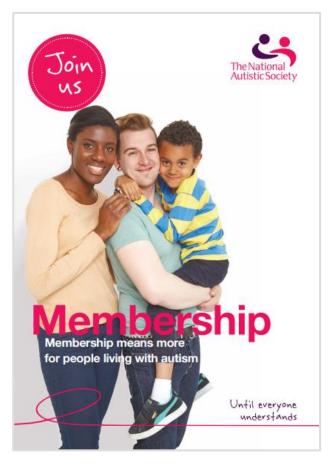






What is Your Autism Magazine?



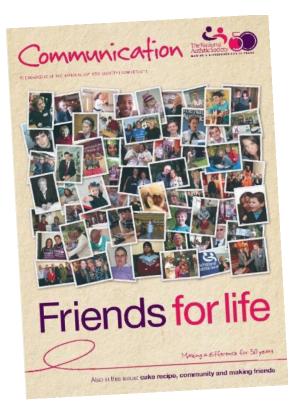






How far we've come

2012





2016





How far we've come

'It put the readers first, focusing on putting their needs at the forefront rather than the needs of the charity. But in doing so, it also met charity objectives."

Judges of the MEMCOM Awards 2014 for Charity magazine of the year







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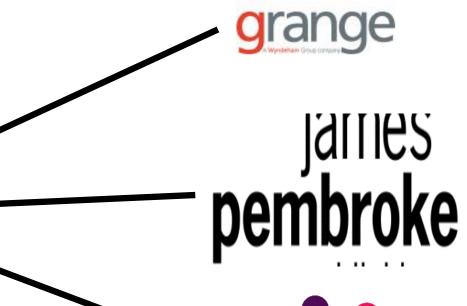
Major research
into describing
autism published

"My wife has pointed out that I may not be sending the best message to my kids, getting paid thousands of pounds for playing with some cardboard."



Putting it all together







Until everyone understands



Principles for a winning read!







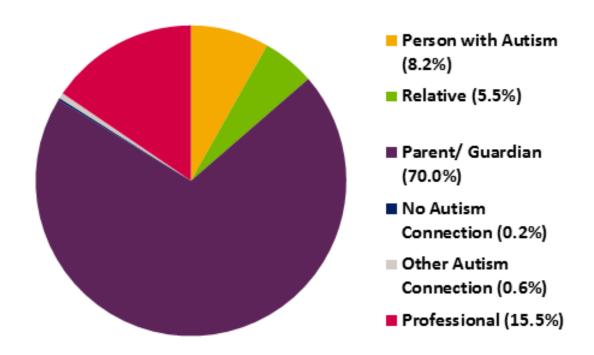
Collaboration







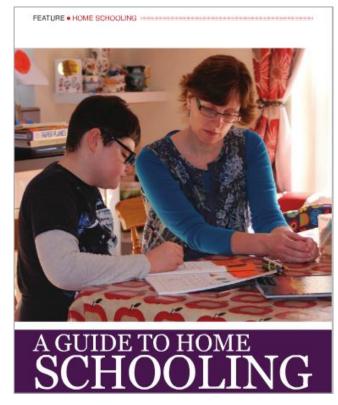
Targeting your readers







Parent/Guardian (70%)





The three Shippey boys: Mason, Owen and Callum



PARENT CAMPAIGNERS . FEATURE

Adding the passion

Kate and Peter Shippey have three suffatic boys. Last year, they successfully campaigned for a sensory room at Sunderland PC so their eldest son, Mason, could enjoy going to matches along with everyone else. We asked them how they did it.

BY SUZANNE WESTBURY

e first heard the term authen when Marion, run edded, was arroard about 18 moreths old—but we had no idea what it was the robus, compiler medical model sook over at the point, and aution only stanted coming up again when he was about five. When he got the diagnosis, we didn't flored in the lost of their as negative. It helped us to flook into it and understand him more.

With our second boy. Owen, we noticed that he was showing similar

your child. So more we use Makazon with the boxs to help them locate and to show other people that we have a particular difficulty. When other people we you using Makazon, that's when their eyes drop and they cop giving you their.

Our advice for other generic would be to beam from your child. I find a led of limes with progression and development, you just need to led them do it as from your washing which your do not never to learn what hothering them, and to understand their icke of a servory norm. The drift added is to do server resized health in see who'd be keen for it. It look a lot of had work and cargoaigness, picharing, support and appraisant and catagogies per a seed to be a few battle be hor inside one of the bare for three people playdest cates. The committee a both the deliver cates. The committee a both the other and public other and public stating, a PEN branch a between and heaving and proper and few the group to put in a freshill-based active with this natures.

We've now trying to get the idea into the premier league and beyond. It's



Until everyone understands



Professional (15.5%)



Bringing you news of the latest research into living and working with autism

How should we describe autism?



When we talk about autism. there are a range of different words and terminologies we can use. The once we prefer might depend on our context - whether that's a medical or research setting. at home with friends and family, or with moses:

The National Authoric Society I'm conducted a piece of imprecedent research to anderstand the stor-people describe against in the UK today. Abuse 4/001 people responded. This hage and representative group reduced adults on the sation. ejectrum, droit parents, families, cares and professionals trorting in summ or a refused 3-dd. You may

hand participated sowerell. We discovered that a wide cariety of tome an usual, and that people have very different views should which one inhor.

10 APRING COL

Looking at all of the responses together, people rated the sorn of the nation spectrum the linghest When the economics from each orgaprofessed the trient on the waters spectrum and nations; powers and cause of individuals on the spectrum preferred than Asperger's or autient

therefore holiene it is reasonable autore: props. When we and on the satient spectrum, and perferenced withing or sating or a school field preferred on the auton spectrum followed shouly conservative with a fixed statement we will provid using the torns that were noted as distillated. is manight deleted by many of our expendents, and we will favour the turn on the autien specture as this was the most saidely professed. sy person with nation or person seith Aspenger's'. Some traditional auties terms are hards make a all any more. Gety a small minimity of purple reported using the term Ramer's syndrome. For when we communicate with a specific group, we will aim to see the specific ferms they prefer.

Katner's satism, satisf, perceins doubpountal disorder, classic within the autism community and we try min bost in reflect show. To find out more about how the At The National Artistic Society National flusters fociety secrets to implement the currentes of the secrets, ledge an eye on the news section of our works. spectrum. Eather, we simile set as a merophore through which the

We know there are many volces

colored those tallogs been up tracked by sales or amplified What we have tessed from our

contach ir ffur their are a variety of professions when a comes to describing sames. We

Epilepsy and autism

Epilepsy is a common, serious neurological condition in which there is a tendency to have recurrent seizures. In the UK, one in every 103 people has epilepsy and 87 people are diagnosed with it every day. We asked a







Parent Dame Stephanie Shirley, who is also an entrepreneur, philanthropist and founder of The Shirley Foundation, explains why she thinks donating the brains. of autistic people for medical research is so important.

BY DAME STEPHANIE SHIPLEY

"WE KNOW THAT autism is brainrelated and researchers know the importance of brain banks allowing access to post mortem brains. But

that my body carrost be used as I.

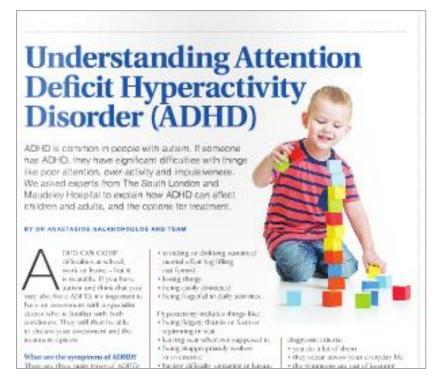
"An awaism basin bank is being planned in Canada, but only the UK. wordment can only be significantly. improved by working with panerolled "Psychological, cultural and practical

BRAIN BANKING - VIEWPOINT





60% join within the first year of getting a formal autism diagnosis









Targeting new audiences



The first generation of people diagnosed with autism in childhood are now reaching middle age. We asked two of our expert service managers for advice on what parents and carers of adults who have autism and more complex needs should consider when helping to prepare someone for later life.

BY SUSANNE LARCOMME AND AMY THORNE

ening older and the prospect of old age meany preparing for major. transition. It can be useful to think of this change as nets of macro transitions. and reicto stansitions. The first category might include changes to housing provision or the death of supporting family members. The second might be more medical appointments or dunges to activities due to a decline in health.



DETERIORATION IN

When it cores to health, there are reveral aspects of acting older which may be more problematic for an auticaic adult. with complex reads. Difficulty with conversation reight recanstare of consens ageing-related Theses are mixed, for example when selling a doctor above pain. Some people was have very bishpain throsholds which could woult in searching rest being pickerd up ural it's really serious. Regularand thorough health checks are therefore vital. Specific checks which talor place as you get often - such as maintreagures - can be hard to environmente, on one rectal stories and visual support to help. explain there in advance. Work with the relevant professionals: to make sure special allowances. are in place. Changes to mobility reight realer a preven's preferred.

How to up your dating game

Unspoken rules, body language, guessing what others are thinking... Dating can be really hard if you're autistic. Luke Jackson, author of bestselling Freaks, Geeks and Asperger syndrome, shares his top tips for how to get started. BY LUKE JACKSON



autism. I was writted I wouldn't be accepted or

he would of dates and relationships is a scary planget feelings for someone. It can turn the most confiden of people into rambling, incoherent. as weeks. But here are some plails you can learn to make it came

in the attenue of francey. The course to come our and ser this grant?





Until everyone understands



Don't limit this to content!









The value of 'look and feel'

FEATURE • MY STORY

Alex and the Marshalls

Ten-year-old actor Alex Marshall is the young star of our Too Much Information film to show the public what a meltdown feels like. At the time of going to press, it had been watched over 40 million times. We asked Alex, his parents Kathryn and Ben. and his little brother Sam, to tell us about real-life meltdowns, and why the public needs to understand more about autism.

EX: "After the day of a bit tired, but it was awesome. It was like winning the lottery twenty guallion

times. My dream is to become an ctor. That's the thing I most want to like I am the actor, it's amazing. The fact that there were actual camera people filming it, actual actors, and director. It felt professional and real and at the end of every take you have this amazing feeling because you've put everything in, and you're so pleased, it doesn't matter if you've smiled too much, or how bad it was -all that matters is that you've acted a take.

I like making films too. In a war it's like acting but different. It's a good way to watch films and get to know people. It can be challenging because y first and last major production didn't really turn out that well, lots of people didn't really want to do it anymore, even people who had already acted in access – and that was a bit of a challenge for continuity. I

but my favourite film is probably Star Wires: Episode III: Recenge of the Sith I watched k when I was six. I liked it then, but I watched it again when I was nine or ten and it blew me away.

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I understood the tragedy in it. I'm a Star Wars fan now. I have to watch all the films, all the episodes, I've got a lot of the comics and I read the books.

MELTDOWNS AND WHAT THEY FEEL LIKE

There are times when I need a lot of room. Sometimes this entire lounge may not be big enough. Sometimes someone brushing past means your room drops to the barest minimum It can be as had as someone ousbing post you. I don't fully know why I find it hard but I think an invasion of space and physical contact when I'm not expecting it and really don't want it.

do intentionally. I have this thing about babies making noise, I don't know why. I just don't like it. That can prove tricky because it can mean I don't want to do something that's planned, because the noises would mean it wouldn't be perfect. When I've been upset and I was going to do something, for me, being upset will have already ruined it. And then I'll get upset because I've missed my chance. And then I get more upset. So the reasons just escalate. If I go ahead and do the thing I was going to do anyway, a lot of my effort logically has to be spent on stopping me from being upset. That's why I'll want to

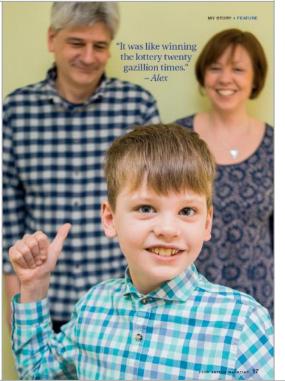
wait till I've calmed down. I don't want

calm down, because a lot of that thing will then be wasted on stopping me being upset, and I'll progress in the chapter of the book or in the level of the game that I like, but I won't

get the full experience.

So instead, I get a comic and flick through it. I have different rules about comics as I do about lots of other things. Once you've read them as much as I have, they are familiar Marvel is my favourite. There is some absolutely mind-blowing Marvel. Sc are average. The thing with comics is sometimes I can find them boring sometimes I find them amazing.

My advice for other people is that if you're upset, it's okay to storm off really angrily, but the important thing with that is the 'off' part, You need to remove yourself from the situation and if there's anything you've got that you can read when you're upset to calm you down the I have my comics), you can do that. And it's okay to be in your room for two and a half hours doing your computer. I'd adviss you not to come down till you're fully calm and fully ready to face the day again. If it takes three hours, that's fine. I'd also advise that if you have ideas for how people can help you t share them. Like making sure you



FEATURE - MY STORY

have the ability to stay in at play times. You don't have to do it all the time, but to have the option is good."

K - Before Alex was diagnosed, things were really difficult at horse and at school for a while.

B - Alex would have meltdowns. I remember sometimes there would be two and a half hours of real. frustration. We used to have to six outside his bedroom door and hold it door shirt because he was so upset. On holiday once the transfer coach was overloaded with people. Alex just went into complete melidown – from the couch to queueing up for check in, it was dreadful.

K−1 had to lock Sam and mpself in the conservatory once because we needed to get away. It was definitely challenging. Aside from that, there was the lack of understanding. I remember late rates you this yebilor no going her girls and wondering why Alex wasn't joining in so much, and pushing him to be part of the children's group without really understanding that actually, he clidn't want to be - and seeing now that, actually, that was fine!

EMBRACING THE DIAGNOSIS

K = Once Alex got the diagnosis, we took the decision that we wanted to fich't really sink in that much initially And it probably wasn't until we got some support from the autism supp unit who went into school to talk to them a bit more about it that it did sun to sink in. It's been something that we've constantly talked about we wanted it to be something that vas there. And I think that's what has made it positive. When people know more, that's when you get more understanding and support. It probably was a relief when we actually got the diagnosis. It helped us understand why Alex was behaving the way be vas and to be able to explain it to

18 SUMMER 2016

Sam (bottom right) says, "I think it's good that Alex really positive and it will help people understand more about autism."

it, the more understanding

ADVICE FOR OTHER PARENTS POST-

DIAGNOSIS

K - Beach out and get as much advice as possible. We looked at all the information on the website from The

National Autistic Society which eally helped. We've also joined the ocal branch. What's great about that is that during that time when you're so alone and so desperate, you can get advice from other parents using the branch's Facebook page. But also secode share tim about what's worked for them and places they've been to which are autism-friendly, so you can get lots of different ideas about what to do. We also went to an anger management seminar – and again just meeting other parents and knowing that although everyone is different.

there are other people in your situation B – If you're over going anywhere on a day out, send them an email. or call them in advance to see what een to places like Bladspool Pleasure Beach where we got to jump every queue. It's also about considering things and realising there may be a reason why something you think is out a solution. Having a diagnosis stally beliped us to think about things in a different way If it doesn't worth, my

something else but there will always be a solution I think. K—You don't suddenly crack it it's constantly a learning curve, and reaching out to different resources is really important.

how he feels and what he tlanks about → Share your story

B - And also to have an idea of what

to put in place and what we needed to

do at home. It also gave us something we could research and find out about

in more detail. Alex is really positive himself about being outsite. And if

he can be positive, then obviously we should be positive too. And our lives

that Alex is outstic and understand

Some of the

out with are

- Ben

astonishing!"

blessom and bearing him talk about

being sutistic is brilliant because he's absolutely funturic. I don't think he

knows how furnistic he is when he

talks about it. Some of the things he comes out with are autonishing!

E - I think it's given us much more

insight into what Alex is thinking and feeling. The more Alex can talk about

his needs. It's probably quite a selfish thing but we want to have as easy a life as we can for him. Seeing Alex

. See Alex in the Too Much information film and join the compaign at www.autism.org.uk/tmi.

 Would you like to share your story? Email yoursuttammagines.org.uk or write to Your Autism Magazine. The National Autistic Society 393 City Road, London EC1V 1NG, and we may feature you in a future issue.







James' story of late diagnosis











Engaging your readers







→ Share your story

- Do you think you could do what David does? Find out about becoming a support worker at www.autism.org.uk/jobs
- Would you like to share your story about any aspect of living or working with autism? Please email YourAutismMag@nas.org.uk or write to Your Autism Magazine, The National Autistic Society, 393 City Road, London EC1V 1NG, and we may feature you in a future issue.





Your Autism Magazine Facebook page







Engage with other organisations

INTERVIEW . KELLY HUNTER

All the world's a stage

Kelly Hunter is an award-winning actor, director and educator. A performer for over 30 years with acclaimed theatre companies such as the RSC and Vesturport, she has been creating drama games for children with autism to improve their communication skills since 2002. This year she directed a

production of The Tempest for children w basis of a longitudinal research study into

BY ELEANOR WHEELER



2 SPECTRUM SPLASHTIME, BRIDGEND BRANCH On Sunday

afternoons

this relaxed pool session is for autistic children and siblings. With water games and floats, there are also lessons available from a parent coach. Find out more at bit.ly/2aeXy3v;



3 AUTISM KIDZ CLUB, RENFREWSHIRE BRANCH

Every other Saturday at 11am-

1pm in Paisley, this group offers arts, crafts, sporting activities, games, baking, toys and other interests. It's a good social opportunity for children and their parents too.

To find out more, email

NASRenfrewshireBranch@nas.org.uk.

→ We'd like to hear about your experiences of diagnosis.

- How long were you waiting?
- Where were you signposted to for help afterwards?
- What advice would you give to others going through a diagnosis at the moment?

Email your answer to

YourAutismMag@nas.org.uk by

16 September 2016 and you'll be entered into a free draw to win some Something Special and Woolly

and Tig goodies, y donated by en Bear Toys.*











1. Aim high and use your charity status







2. Make it everyone's magazine







3. Use your digital offering intelligently







4. Don't be ruled by the \$\$\$







5. Splash your celebrities!







Any questions?

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