

You can make a difference today!

You can make a donation by completing the form below or online through our website:

www.smasupportuk.org.uk/donate

Name

Home Address

Postcode

Signed

Date (DD/MM/YYYY)

Boost your donation by **25p of Gift Aid for every £1 you donate.**

I am a UK taxpayer and understand that if I pay less Income Tax and/or Capital Gains Tax than the amount of Gift Aid claimed on all my donations in that tax year it is my responsibility to pay any difference. I want to Gift Aid my donation of £..... and any donations I make in the future or have made in the past 4 years to SMA Support UK.

Please notify us if you:

- Want to cancel this declaration
- Change your name or home address
- No longer pay sufficient tax on your income and/or capital gains

If you pay Income Tax at the higher or additional rate and want to receive the additional tax relief due to you, you must include all your Gift Aid donations on your Self-Assessment tax return or ask HM Revenue and Customs to adjust your tax code.

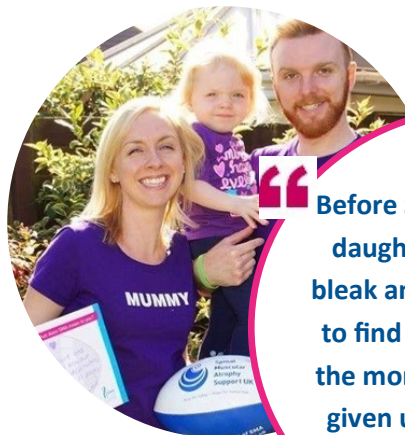
BACS Transfer:

Account: Spinal Muscular Atrophy Support UK

Bank: Barclays

Sort Code: 20-48-08 Account No: 00524514

Please include a reference with your donation.



“Before SMA Support UK, our daughter’s future seemed bleak and we were struggling to find a reason to get up in the morning. The charity has given us something money cannot buy: hope. That is priceless.”

We’d love to hear from you

Spinal Muscular Atrophy Support UK

40 Cygnet Court

Timothy’s Bridge Road

Stratford-upon-Avon, CV37 9NW

Phone: 01789 267520

- Mon – Thurs (9.00 am—3.30 pm)
- Friday (9.00 am—1.00 pm)
- Closed on public holidays.

Email:

fundraising@smasupportuk.org.uk

www.smasupportuk.org.uk



Registered Charity No 1106815
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How You Can Help Us

Let’s make a difference



 **Spinal Muscular Atrophy Support UK**
Help for today • Hope for tomorrow

What is Spinal Muscular Atrophy?

Spinal Muscular Atrophy (SMA) is a rare, genetically inherited neuromuscular condition.

1 in 40-60 
of us carry the gene that causes SMA
which equates to around
1.5 million
 people in the UK.

If two carriers have a baby there is a chance their baby will have SMA.



SMA causes progressive loss of movement and muscle weakness as a result of muscle wasting (atrophy).

What do we do?

We are an established charity that provides anyone affected by SMA with free:

- Information
- Emotional support
- Practical advice and guidance

We are advocates for better services and access to new treatments and we fund research related initiatives.

We raise public awareness and understanding of how SMA affects the lives of babies, children, young people and their families.

We cannot do this without you.

Join an Event

It's a great way to get fit, have fun and socialise as well as raise funds. Our calendar is packed with a wide range of events to suit all ages and abilities, or you can find your own.



Take part in one of our Fundraisers

You can run a Coffee and Cake Day any time, organise a Christmas Jumper Day in December, or get your local school involved in Shoesday in July.



Involve your Employer and Local Businesses

Encourage them to choose SMA Support UK as Charity of the Year, host a collection tin or join in with one of our fundraisers. Every penny counts!



Leave a Legacy

Just as a will brings security to your family's future, a legacy to SMA Support UK will secure a future for our vital work.



Set up a Tribute Fund

All your fundraising efforts can be pooled in the name of someone special affected by SMA.



www.smasupportuk.org.uk/help-us