

Help us work towards a World Free of PSP

## The PSP Association

**There is a lot that can be done to improve the quality of life of people living with PSP, and to support their families and carers.**

The PSP Association was founded in 1994 and is the only organisation in the UK dedicated to helping everyone with PSP, together with their families and carers.

Because PSP is not widely known, we also work with medical, health and social care professionals providing education, training and advice on all aspects of managing the disease. We also work to influence decision makers at national and local levels to raise awareness and improve services.

We fund research into causes, effective treatments and a cure for PSP. To date we have invested £2.5 million. We also actively promote the need for more research worldwide.



## We are here for you

People living with PSP, their families and carers need all the information and advice available to ensure they are fully equipped to access the care and support that's so vital for continued quality of life, and to make informed choices to suit their individual and changing needs.

The PSP Association are here to help. We are the leading experts in PSP and offer a range of information and services designed to cover every aspect of living with PSP, including:

1. A telephone or e-mail link to a **Specialist Nurse**, who will advise on any aspect of living with PSP.
2. A comprehensive **step by step guide** to living with PSP, giving practical advice and information.
3. Local **support groups** providing opportunities to meet others in a similar situation and develop new contacts and networks. Currently there are 70 meetings each year throughout the UK and Ireland. You can also sign up to receive details of other carers in your area.
4. **Local news**, to keep you updated at a national and local level on the support available and how to access it.
5. A **website** - [www.pspeur.org](http://www.pspeur.org) where you can access and download a full range of information. We also have an online **discussion forum** where you can make contact with those in a similar position and share your day to day experiences.
6. Invitations to information events specifically designed to provide you with information and updates on living with PSP, including a day conference with specialist speakers.
7. Opportunities for family and friends to support the work of the Charity via fundraising and volunteering. Whatever you want to do to help we would be keen to support you.

**We are the only organisation in the UK and Ireland specifically for everyone affected by PSP.**

We also support people living with Cortico Basal Degeneration (CBD)



**PSPPA**

**The PSP Association**

PSP House, 167 Watling Street West,  
Towcester, Northamptonshire NN12 6BX

**T:** 01327 322410 **F:** 01327 322412  
[www.pspeur.org](http://www.pspeur.org)

Registered charity numbers: England & Wales 1037087 / Scotland SC041199

Progressive  
Supranuclear Palsy

**PSPPA**

## Membership Form

**“Life would have been a lot harder if The PSP Association hadn't been there providing advice and support to me when dad had PSP.”**



**Working for a  
World Free of PSP**

# Membership

## There are four individual membership options:

1. For people living with PSP there is no fee
2. For carers, spouses and partners of people with PSP there is no fee
3. Individual membership is £10.00
4. Life membership is £250.00

**Overseas membership to people with PSP and carers/spouse/partner is free.**

**The PSP Association receives no Government funding and is wholly dependent on voluntary donations. By becoming a member you are supporting the quest for a 'World Free of PSP' and by strengthening our voice you will help us champion better care and support for patients and carers and fund more research into causes, effective treatments, prevention and ultimately a cure for PSP.**

## Membership includes:

- A welcome pack.
- Three editions a year of our membership magazine PSP Matters, containing updates on care, and the latest research, as well as practical advice on living with PSP.
- Invitations to patients and carers' events, information on fundraising events and initiatives - locally and nationally.
- Carer's Information Pack (if you have PSP or are a Carer).

# Membership application form

Please complete the name of the person holding the membership using **BLOCK CAPITALS**

## Name of person who will hold the membership

I have or care for someone with PSP

Surname

Forename(s)

Title (Mr/Mrs/Ms/Mr & Mrs/other)

Address

Postcode

Telephone

Email

It would be useful to have a date of birth (optional)

Please indicate the membership you would like (full details of which are shown on opposite page)

## Individual membership

Person living with PSP No fee

Carer, spouse or partner of person with PSP No fee

Individual membership £10 PA

Life membership £250

Overseas membership for people with PSP and carers/spouse/partner is free No fee

If taking up Carers' membership please tell us the name of the person you are caring for

My relationship is (tick box):

Spouse  Partner  Other (please state)

You can also join by telephone on 01327 322415

## Additional information

To help us to continue to develop the services we provide to people with PSP we may write to you from time to time asking for your feedback and views. You will also receive PSP Matters, the membership magazine, three times a year.

Please tick the following boxes as appropriate:

I do not wish to be contacted

I do not want to receive PSP Matters

All personal data will remain within the PSP Association and will not be shared with any other organisations or individuals.

## How do I pay?

You can pay your membership by cheque, credit or debit card. Some people also like to make a donation to The PSP Association for which we are always grateful.

My subscription amount is: £  :

I would like to make a donation £  :

I wish to pay by cheque (made payable to The PSP Association)

Credit/Debit card

Mastercard  Visa  Maestro/Switch

Card Number

(shaded boxes required for Maestro/Switch only)

Maestro/Switch Issue No  Valid from  /  Expires  /

For security reasons please write the last 3 unique numbers printed on the reverse of your card.

Cardholder's name

Cardholder's billing address

Postcode

Signature  Date

*giftaid it* I would like the PSP Association to reclaim the tax on this subscription and any donation I have made in the four years prior to this year and any donations I may make in the future from the date of this declaration until I notify you otherwise. I am a UK taxpayer. I am paying at least as much in income and/or capital gains tax each year as The PSP Association reclaims on my donations  Date

I am already signed up to Gift Aid with The PSP Association

I am not a UK taxpayer

Please return your completed form to: **The PSP Association, 167 Watling Street West, Towcester, Northants NN12 6BX.**