

The Atypical Parkinsonism Service

**A guide to the clinic for patients
and families**

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The Team

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(Movement Disorder
Neurophysiotherapist)



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(Neuropsychologists)



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[MSA] Trust Specialist
Nurse)



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(Progressive Supranuclear
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(Speech and Language
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An Information and Support Worker is also part of our team

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You have received this leaflet as you, or someone you care for, has been referred to the Atypical Parkinsonism service.

What is Atypical Parkinsonism?

Atypical Parkinsonism or “Parkinson’s Plus” refers to a group of conditions that can initially look like Parkinson’s disease. These conditions are Multiple System Atrophy (MSA), Progressive Supranuclear Palsy (PSP), Corticobasal Syndrome (CBS) and other similar conditions.

What can we offer you?

We can offer you specialist support in our multidisciplinary service which includes:

- Specialist movement disorder nurse and doctors
- Neurophysiotherapy,
- Neuropsychology
- Links to speech and language therapy
- Community therapy teams.

We have a charity nurse from the MSA (Multiple System Atrophy) Trust and a care navigator from PSPA as well as a carer support worker.

What will happen on the day I attend?

When you attend the clinic you will see a combination of members from the multidisciplinary team including a doctor and a nurse. This will usually be a face to face appointment. You are very welcome to bring a family member, carer or friend.

We will ask questions and examine you and try to identify any problems that we can help with. You will be able to see a neurophysiotherapist (a physiotherapist who specialises in treating people with neurological conditions) and a speech and language therapist if required.

We may offer you an appointment with other members of staff such as a neuropsychologist which will take place on a different day, and will depend on your needs. Neuropsychologists work towards understanding how illnesses of the brain and nervous system affect memory, mood and the ability to function in daily life and how emotions such as anxiety, stress and low mood affect the brain and nervous system.

Your first visit may take up to 2 hours.

We will offer your family member or carer an appointment with the carer support worker.

Some patients are referred for a one off appointment after which their ongoing care will continue with the doctor who referred them. Other patients may continue to attend the service, in this case future appointments are likely to be shorter.

What if I cannot attend face to face?

We will be able to arrange a telephone or video consultation for you.

Where can I get more support?

You can get more support from the following charity organisations

Multiple System Atrophy Trust (MSA) Trust

🌐 <https://www.msatrust.org.uk/>

✉ support@msatrustorg.uk

☎ 0333 323 4591

PSPA (also for people with Corticobasal Syndrome (CBS))

🌐 <https://pspassociation.org.uk/>

✉ helpline@pspassociation.org.uk

☎ 0300 0110 122

Information and support for carers

Carers Information Line: ☎ 0141 353 6504

🌐 <https://www.yoursupportglasgow.org/accesscarerservice>

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