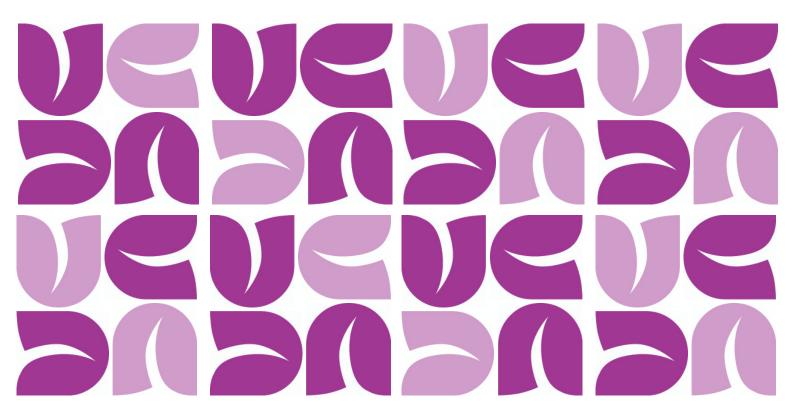


PRE-BUDGET SUBMISSION 2023-24

Proposal for a National Awareness Campaign & GP Education Program



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2023-24 PRE-BUDGET SUBMISSION

Subject:	Proposal for a National Awareness Campaign & GP Education Program		
То:	The Hon Mark Butler, Minister for Health and Aged Care		
From:	Parkinson's Australia	Date:	16/11/22

Parkinson's Australia is the national advocacy body representing people living with Parkinson's, their families and carers, as well as researchers and health professionals in the Parkinson's community.

We promote the best possible quality of life for people with Parkinson's. We advocate for the Parkinson's community on issues of national significance. We work to reduce the impact of Parkinson's by promoting best practice care to ensure that people can maximise their opportunities to live well and maintain their independence.

Our objectives are:

- Raise awareness about early warning signs and the motor and non-motor symptoms of Parkinson's, highlighting that Parkinson's is more than a tremor.
- Challenge the perception that Parkinson's is 'just an old person's condition', as approximately 20% of people diagnosed with Parkinson's are under 50 and are of working age.
- Raise awareness about Parkinson's being the fastest growing condition in Australia.
- Encourage those who have early warning signs and symptoms to seek an earlier diagnosis, which will educate GPs, and other allied health professionals in Australia. This will help them to understand and recognise early warning signs and symptoms which will enable swifter referral pathways to Movement Disorder Specialists.

THE PROBLEM

1 in 260 people in Australia are living with Parkinson's.

Because of its chronic progressive nature, insidious onset, complicated and diverse motor and non-motor symptoms, Parkinson's is one of the most challenging diseases to provide care for. Parkinson's is a neurodegenerative condition that currently has no known cure. Parkinson's is more common than prostate, bowel and many other cancers that are considered National Health Priority Areas.

Parkinson's is the most common Movement Disorder and had the second highest prevalence in 2005 compared to other neurological conditions, exceeded only by dementia (Deloitte Access Economics 2011). People diagnosed with Parkinson's generally have four key clinical features: tremor, muscle rigidity, poverty of movement and loss of normal posture with a tendency for falls. Lesser known is the major impact on non-motor functions such as depression, anxiety, behavioural disturbances and cognitive impairment, sleep disorders, hallucinations, sensory deficits such as loss of smell and visual problems, and impaired speech. Due to its complex symptoms, Parkinson's has a dramatic effect on a whole family.

There are over 100,000 people in Australia with Parkinson's, and another 1,000,000 people directly affected by Parkinson's including spouses/partners, siblings, offspring, friends and colleagues.

Parkinson's prevalence increases threefold after the age of 65. The growth rate in the number of people living with Parkinson's is expected to continue to increase dramatically. It is estimated that the



growth rate will average 4% over the next 20 years. Parkinson's presents a greater immediate pressure on the health system in Australia because of our ageing population. This extends to significant numbers of people with Parkinson's in regional, rural and remote areas which have a higher median age than metro areas.

Living well with Parkinson's requires the expertise and involvement of many allied health professionals which begins with the General Practitioner being able to recognise early warning signs of Parkinson's to enable an early diagnosis.

THE SOLUTION: A NATIONAL AWARENESS CAMPAIGN & GP EDUCATION PROGRAM

An investment by the Federal Government to fund a public awareness campaign and a GP education program is two-fold.

Raising awareness of early warning signs will encourage people to seek professional help, which will lead to an earlier diagnosis. Early intervention leads to better management of the condition, for the individual, their families and loved ones, and the health system more broadly.

Early diagnosis means the person can receive the right treatment and access to support that can help to slow the progression of Parkinson's and improve their overall quality of life. Ultimately this may lead to reduced presentations to emergency departments and can allow people to live independently longer.

A \$2.5 million dollar investment over 3 years (breakdown in table on p5) would ensure the creation and distribution of a National Awareness Campaign and an Education Campaign that will upskill GP's and allied health professionals in how to ensure that someone lives well with Parkinson's.

Evidence-Based Outcomes

<u>Parkinson's disease: patient and general practitioner perspectives on the role of primary care -</u> PubMed (nih.gov)

Conclusion: Patients and GPs see a limited role for the GP in early-stage Parkinson's Disease (PD) care because of patient autonomy and GP's lack of specific knowledge and skills. However, GPs should feel more confident of the added value of their generalist approach to care for patients with a complex chronic disorder as PD. If generalist and specialized care reinforce each other, PD patients benefit.

https://bmchealthservres.biomedcentral.com/articles/10.1186/s12913-022-07503-7

PD diagnosis and treatment practices: The majority of respondents in both geriatricians and GPs groups reported having confidence in making the diagnosis of PD, with no significance difference noted (69.8% of GPs and 78.8% of geriatricians, p = 0.38) (Table 2).

Geriatricians were more comfortable in initiating dopamine replacement therapy (DRT) for patients they had diagnosed with PD, as compared to GPs (OR = 11.8 [95% CI: 3.54–39.3], p < 0.001). When asked further about why they may not feel comfortable with starting the treatment, GPs were more likely to report that they were unfamiliar with the types and/or doses of PD medications as compared to geriatricians (OR 7.14 [2.58–19.8], p < 0.001). GPs were more likely to report being uncomfortable with providing PD education (OR 8.64 [2.84–26.3], p < 0.001), and their patients were more likely to request a neurologist confirmation before initiation of the treatment (OR 6.94 [1.94–24.8], p = 0.003).

Increasing public awareness and GP and allied health education improves the quality of life for people living with Parkinson's. In simple terms, better care and treatment keeps people out of hospitals and residential care. It enables people with Parkinson's to stay in the workforce longer, continue to live independently in their own homes, increase productivity and savings, and reduce the demand for care. The immediate outcomes would be:



- Improve the management of symptoms and quality of life
- · Provide information and education regarding advanced care treatment options
- Provide information and support regarding advanced care directives
- · Reduce hospital admissions and length of stay for necessary admissions
- Delay entry in residential care and reduce the need for costly medical consultations and interventions
- Assisting to manage the very wide, complex and idiosyncratic medication regimes of individuals which is essential as access to a neurologist is usually annual.

The expected outcomes:

- Reduce unplanned admissions
- Increase education regarding treatment options
- Reduce bed days
- Avoid re-admission
- · Improved patient experience for end-of-life care
- · Provide multidisciplinary care
- · Enable self-management
- Support Carers

GP EDUCATION PROGRAM

Parkinson's Australia recognises the need for endorsed, consistent and up-to-date information and education for GPs and Allied Health Professionals. This is to ensure the earliest possible diagnosis of Parkinson's, which leads to the confident implementation of evidence-based treatment and management options.

It is proposed that Parkinson's Australia leads the development of comprehensive education material and resources for health professionals, drawing on the latest scientific evidence and where appropriate, international resources; guided by the needs of people with Parkinson's.

The GP Education Program seeks to:

- Enhance recognition of the early signs of Parkinson's and the benefits of early intervention
- Ensure health professionals are up-to-date with the latest evidence-based management of Parkinson's
- Enhance the understanding and treatment of both motor and non-motor symptoms
- Reduce the lengthy and uncertain pathway to diagnosis for people with Parkinson's
- Improve overall health and wellbeing outcomes for people with Parkinson's

Guiding Principles

The GP Education Program will be guided by:

- Parkinson's Australia Strategic Plan/State Strategic Plans
- Parkinson's educational resources being utilised internationally
- The expertise of Parkinson's Nurse Educators across Australia
- "Champions" working in the individual health professions
- The experiences of people living with Parkinson's



Development of Resources

The development of educational resources for the GP Education Program will draw on a key set of information gathering tools, designed to efficiently consolidate:

- The latest scientific evidence in Parkinson's management
- The latest advancements in online CPD (Continuing Professional Development) health education
- The latest advice from CPD accreditation organisations
- The perspectives and needs of people with Parkinson's
- The perspectives of training bodies including the National Prescribing Service

Workforce Considerations

The GP Workforce

- 37,472 GPs working in Australia in 2019
- 52% of the FTE GP workforce is overseas trained
- 74.5% of GPs work in the major cities
- 90% of the Australian population visits their GP at least once each year
- Australians access general practice more than any other part of the health system (RACGP General Practice Health of the Nation Report 2020)

Royal Australian College of General Practitioners (RACGP)

New processes for approving CPD education activities have been introduced by The Royal Australian College of General Practitioners (RACGP), to strengthen the educational rigour of CPD, and to ensure that activities focus on high quality education suited to the professional development needs of GPs. The advice from RACGP to GPs when choosing an education activity for CPD Points is to: "Consider your current and future scope of practice and choose high-quality education resources or activities that will enhance or develop new skills and knowledge within your defined scope."

Australian College of Rural and Remote Medicine (ACRRM)

The Australian College of Rural and Remote Medicine (ACRRM) offers its Professional Development Program to Australian GPs regardless of location. ACCRM is also utilising new criteria to assess activities for accreditation as follows:

- 1. Does the educational activity help to update knowledge and skills, review performance and/ or measure outcomes?
- 2. Does the activity relate to the College's Primary Curriculum?
- 3. Does the educational event/activity have clear, specific learning objectives?
- 4. Are the planned educational strategies based on adult learning principles?
- **5.** Does the application show that some impact evaluation (changes in knowledge, skill, attitude, practice or patient outcome) is planned?

National Prescribing Service

The National Prescribing Service offers CPD points for GPs, as well as education material on the "Management of Parkinson's Disease," and this is currently based on research papers on Parkinson's medications that are more than 10 years old. These can be found at https://www.nps.org.au/australian-prescriber/articles/management-of-parkinsons-disease



The former CEO of the National Prescribing Service, Professor Steve Morris has recently returned to Adelaide and would be an ideal (and approachable) contact to provide insight into medication prescribing issues that could potentially be addressed with updated educational materials.

NATIONAL AWARENESS CAMPAIGN

Who will build and run the awareness campaign?

Parkinson's Australia has been working on a concept with Dee Madigan and the team from Campaign Edge who have created a detailed proposal for a national awareness campaign. We approached Campaign Edge as they have already developed a very successful Dementia Australia awareness campaign.

We need our target audience to recognise and understand the signs of common symptoms of Parkinson's. We want them to question if they, a family member or close friend are affected by those symptoms, and to seek more information and ultimately advice from a health professional.

Target Audience

- Broad Australian public
- Skewed to 30 60-year-olds
- Even male / female skew
- National campaign across all States and Territories

Audience Campaign Goals

What do we want the target audience to think, feel and do?

Parkinson's is now the fastest growing neurodegenerative condition in Australia with a

broad range of symptoms such as tremor, rigidity, slowness and stooped posture.

Parkinson's can affect people at all ages.

FEEL: Knowledgeable in recognising the symptoms of Parkinson's

DO: Seek help from their GP if they have symptoms

Audience Journey

THINK:

- Exposure to creative TV, Radio, Press & digital environments.
- Click through to a dedicated campaign website with long form content, resources and downloads, plus a dedicated 1800 or 1300 line, either managed by Parkinson's Australia and our associates or outsourced, to answer queries over the phone.
- Develop a remarketing list to keep connected with users through ongoing content.

FUNDING REQUIREMENT

Project Stage	Funding Requirement (inc GST)	Period
Development of GP Education Program	\$800,000	Y1
National promotion of GP Education Program	\$500,000	Y2
National awareness campaign development and production cost	\$296,450	Y2
National awareness campaign promotion and media buy	\$825,000	Y3
TOTAL	\$2,421,450	



Proven Ability to Deliver:

Parkinson's Australia has delivered a successful \$1.1m Information, Linkages and Capacity Building grant and has just been funded an extension of \$470k to continue to enhance the YOP-X app.

The Young Onset Parkinson's Exchange (YOP-X) app and website is a world first information portal and resource hub supporting people with young onset Parkinson's. Developed as a living lab model, the experiences, ideas, knowledge and daily needs of people with young onset Parkinson's have underpinned all app and website content, with more than 250 individual stories and experiences shaping tools and resources. YOP-X also provides a fast track for people with young onset Parkinson's to collect and collate information relevant to their NDIS applications.

The Young Onset Parkinson's Exchange now sits within the top four Google listings for Parkinson's, and has narrowed the gap between diagnosis and access to targeted and appropriate information and support. Prior to the launch of the Young Onset Parkinson's Exchange, a new diagnosis of Parkinson's often meant sifting through websites targeting the over 65 age group, with many of these set up outside of Australia and mostly offering confronting information for later stage Parkinson's. As the Young Onset Parkinson's Exchange has compartmentalised information within pillars for access when a user is ready, YOP-X resources are sitting parallel to stages along the Parkinson's journey to provide targeted assistance, whether it be around maintaining employment, sex and relationship issues, mental health or sleep concerns.

The NDIS Wallet is the first of its kind and is the most utilised function of the YOP-X app, accessed 3861 times and assisting users to consolidate data for their NDIS applications. Aside from the NDIS Wallet, the most accessed areas of the YOP-X app offer daily activities to work through including strength and balance exercise videos accessed 3281 times, mental health tools accessed 3108 times, and sleep tools accessed 2823 times.

Thank you for taking the time to review our submission.

Prof George Mellick

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