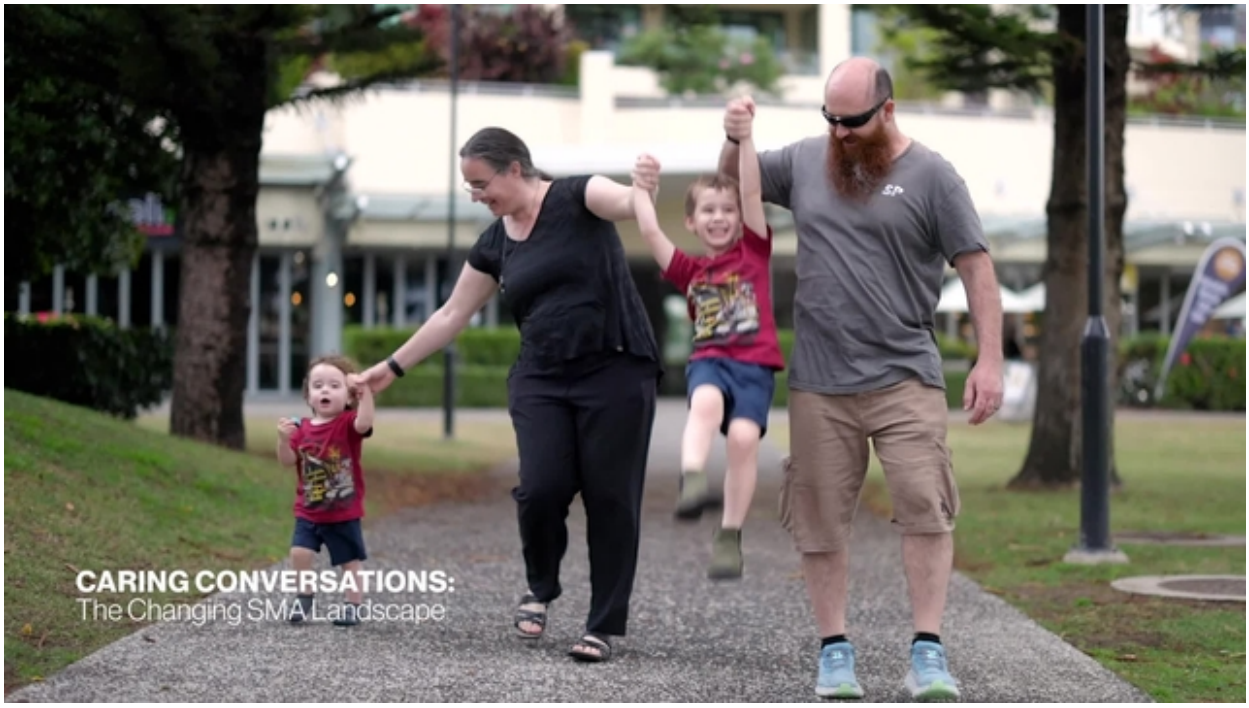
NOVARTIS

Spinal muscular atrophy
Image

Spinal muscular atrophy (SMA) is a rare, inherited monogenic disease characterised by motor neuron degeneration and muscle weakness[4,5]

Listen to mum, Carolyn, describe her family's journey with John's prenatal SMA diagnosis and early treatment at birth.



VIDEO

Image

- Until recently, SMA was a devastating and often fatal disease.[6-8]
- Three PBS-reimbursed disease-modifying therapies for infants with SMA have been available from June 2021.[8-14]
- Reproductive and treatment options should be discussed with appropriate specialists as early as possible.

Where to refer SMA genetic carrier couples

Where to refer SMA genetic carrier couples

Click here for contact details to refer SMA genetic carrier couples to genetic counsellors who have expertise in available treatment options

See more details

Hide details

Interested in receiving more SMA and product specific information/invitations in the future?

Please kindly fill out this quick form with your contact details

Register for Novartis Pro here

PBS, Pharmaceutical Benefits Scheme; **SMA**, spinal muscular atrophy.

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