Our Vision

A future where genetic advances, treatments, connection and care are within reach for every person, no matter how rare.

genetic alliance AUSTRALIA

Our Mission

We connect, support, and advocate alongside people living with rare genetic and undiagnosed conditions. We link families, communities, and systems to improve access to care, build knowledge, peer connection, and self-advocacy.

The Values That Drive Our Work

Listen & Learn

People with lived experience are experts in their own lives.

Champion Strengths

Every person, family and community has unique strengths to recognise and build on.

Collaborate & Amplify

Equity Requires Collaboration. We achieve more when we work together than we ever could alone.

Our Strategic Priorities

Pathways to Care

We help people navigate the uncertainty that often comes with rare genetic and undiagnosed conditions by providing trusted information, personalised navigation and connection to the right supports.

Every interaction helps us identify gaps, improve pathways and advocate for more responsive, equitable systems of care.

Strong, Connected Communities

We facilitate networks for families and communities to share knowledge and support one another. Through our rare contact database we connect families navigating similar journeys.

We support rare condition peer groups - particularly small, volunteer-led organisations - by providing resources, support and shared infrastructure that amplifies their essential work.

Research Partnership & Systems Change

We work with researchers, clinicians, policymakers and communities to ensure lived experience shapes research, supports the translation of new knowledge into practice, and contributes to more responsive, person-centred systems of care.