Palliative Treatment for Children South Africa (PatchSA) is an inclusive and compassionate network that aims to share specialised knowledge, tools, resources and opportunities to ensure the best possible care for children with life-threatening and life-limiting illnesses, and to provide support to healthcare professionals, families and lay caregivers.

PatchSA welcomes any families, lay caregivers and child-caring professionals, including paediatricians, GPs, nurses, social workers, allied health practitioners, psychologists, teachers and clergy to join the network.

Our Vision
To ensure the right of every child to access quality palliative care in South Africa

Our Mission
To mobilise and support a sustainable network of individuals, organisations, professionals and caregivers to provide palliative care for children and their families from diagnosis to bereavement as an essential component of Universal Health Coverage (UHC).

Our Objectives
The Association is a Voluntary Association and non-profit organisation established for the following public benefit objectives:
• To create and sustain a vibrant network of individuals and organisations with an interest in children’s palliative care;
• To be a repository of information on children’s palliative care, resources and service providers in South Africa and to develop and promote local appropriate guidelines and standards for children’s palliative care;
• To provide evidence-based education in children’s palliative care for professionals and parents;
• To be a strong collaborative voice for the right of the child to receive quality palliative care in South Africa;
• To drive the research agenda for evidence-based children’s palliative care; and
• To promote the development of support systems for children, families, professional and lay caregivers.

PatchSA Fact Sheet
Palliative Treatment for Children South Africa

PatchSA is a registered non-profit organisation. NPO no: 124-201