



Patient Support Resources

There are many entities which offer information on pheochromocytoma and paraganglioma, general advice and support, and which connect patients with experienced doctors – and with each other. Many also feature links to studies on pheo/para which are currently recruiting, and stories from patients and caregivers around the world. Some of the most active groups are listed below.

[Pheo Para Alliance](#)

www.pheopara.org

[Pheo Para Project](#)

www.pheoparaproject.org

[SDH-Deficient Cancer Research Advocates](#)

www.sdhcancer.org

[VHLAlliance](#)

www.vhl.org

[AMEND](#)

www.amend.org.uk

[Nederlandse Vereniging voor patiënten met Paragangliomen \(NVPG\)](#) (support for Dutch-speaking patients)

paragangliomen.nl

[Pheipas](#) (support for Spanish-speaking patients)

www.pheipas.org/en/

[pheopara.com](#) (support for Japanese-speaking patients)

[LACNETS](#)

www.lacnets.org

[NorCal CarciNET](#)

www.norcalcarcinet.org



Neuroendocrine Cancer Australia

neuroendocrine.org.au

Care Centers

Healthcare professionals may encounter patients who require more specialized care, or who are already receiving such care at medical facilities outside their local areas. The PheoPara Alliance identifies and certifies such care centers around the world, classifying them according to the level of expert care pheo/para patients can expect to receive. A current list of Centers of Excellence and Clinical Centers can be found on the Pheo Para Alliance web site:

<https://pheopara.org/education/centers-of-excellence>