

Parry Romberg Syndrome (PRS) is a rare disease characterized by progressive degeneration of the tissues under the skin, usually only on one side of the face.

Parry Romberg Factsheet



Parry Romberg Syndrome (PRS) has been described in medical literature for almost 2 centuries, but there are no known causes or cures.



PRS typically first presents in children between the ages of 5-15.



20% of patients have hypo- or hyper-pigmented skin on the affected areas.



45% of patients have neurological issues, including seizure disorders and stroke.



10-20% of patients develop ocular issues



Half of patients develop dental abnormalities



The incidence of PRS is estimated to be 1 in 750,000.



The proportion of women to men with PRS is 3:2