

Hemispherectomy for Rasmussen's encephalitis

Compared with functional hemispherectomy and hemisphere disconnection, anatomical hemispherectomy elicited better seizure outcomes with an acceptable level of complications. Early-stage operations might lead to better cognitive status, but they are associated with a high risk of IQ decline ¹⁾.

Obtaining complete [disconnection](#) is critical for favorable [seizure outcomes](#) from [hemispherectomy](#), and [neurosurgeons](#) should have a low threshold to reoperate in patients with [Rasmussen's encephalitis](#) with [recurrent seizures](#). Rapid [progression](#) of motor [deficits](#) and bilateral [MRI](#) abnormalities may indicate a subpopulation of patients with RE with an increased risk of needing [reoperation](#). Overall, they believe that [hemispherectomy](#) is a curative surgery for the majority of patients with RE, with excellent long-term [seizure outcomes](#) ²⁾.

The majority of pediatric patients undergoing resection or [hemispherectomy](#) for RE achieve good seizure outcome. Although small retrospective cohort studies are inherently prone to bias, the best available evidence utilizing individual participant data suggests hemispheric surgery and younger age at the surgery are associated with good seizure outcomes following epilepsy surgery. Large, multicenter observational studies with long-term follow-up are required to evaluate the risk factors identified in a review ³⁾.

Hemispherotomy remains the gold standard treatment but causes permanent functional impairment. No standardized medical treatment protocol currently exists for patients prior to indication of hemispherotomy, although some immunotherapies have shown partial efficacy with functional preservation but poor antiseizure effect. Some studies suggest a role for tumor necrosis factor alpha (TNF- α) in RE pathophysiology.

¹⁾
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²⁾
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³⁾
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