

Osler-Weber-Rendu syndrome treatment



Osler-Weber-Rendu syndrome treatment focuses on reducing [bleeding](#) from [blood vessel](#) lesions, and sometimes surgery or other targeted [interventions](#) to remove [arteriovenous malformations](#) in organs. Chronic bleeding often requires [iron](#) supplements and sometimes [blood transfusions](#).

Surgery is indicated mainly for the evacuation of [hematoma](#) or diagnosis, especially when favorably located. Also, consider surgery for recurrent hemorrhages (rupture has been reported even after normal angiography) or medically intractable seizures. [Stereotactic radiosurgery](#) has not had a satisfactorily high enough benefit to risk ratio to justify its use ¹⁾.

The risk of treatment of brain [AVMs](#) in patients with Osler-Weber-Rendu syndrome is quite low for appropriately selected patients with treatment individualized to [radiosurgery](#), [microsurgery](#), or a combination of [embolization](#) and microsurgery ²⁾.

Currently, conventional [heparin](#) and [warfarin](#) remain first choice [anticoagulants](#). If newer anticoagulants are considered, although study numbers are small, at this stage [Apixaban](#) appears to be associated with lesser bleeding risk than [Rivaroxaban](#) ³⁾.

References

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