Designed to prevent reherniation

Published Clinical Evidence Overview
These articles contain off-label use information. Please use medical discretion when reviewing these articles.

**Barricaid is approved for the following indications for use:**
Reducing the incidence of reherniation and reoperation in skeletally mature patients with radiculopathy (with or without back pain) attributed to a posterior or posterolateral herniation, and confirmed by history, physical examination and imaging studies which demonstrate neural compression using MRI to treat a large anular defect (between 4-6 mm tall and between 6-10 mm wide) following a primary discectomy procedure (excision of herniated intervertebral disc) at a single level between L4 and S1.

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Full financial disclosures can be found in the respective manuscripts.
Published Peer-Reviewed Journal Manuscripts with Barricaid® Data

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