

# NOW CE-MARKED UNDER EU MDR<sup>†</sup> FOR GENICULAR ARTERY EMBOLISATION (GAE)

## GENESIS Trial:<sup>1</sup> Embosphere Offers Patients Relief

For patients with mild-to-moderate knee osteoarthritis, two-year outcomes showed that GAE with Embosphere delivered durable pain relief, with reduced analgesics use over time, and more.<sup>1</sup>


**87%**  
achieved technical success

**77%**  
positive survey responses

Mean VAS\*  
**36%**  
at 2 years

Mean Total WOMAC\*  
**38%**  
p<0.001

### ORDERING INFORMATION

Catalogue Number 1-mL Syringe	Catalogue Number 2-mL Syringe	Size Range	Packaging Color
S210GH	S220GH	100–300 μm	Yellow 

Embosphere Microspheres are packaged with saline in a 20-mL syringe. Packaged 1 syringe per box.

**WOMAC** = Western Ontario and McMaster Universities Osteoarthritis Index **VAS** = Visual Analogue Score  
\* Percent change from baseline.

1. Little et al. 2024. "Genicular Artery Embolisation in Patients with Osteoarthritis of the Knee (GENESIS) Using Permanent Microspheres: Long-Term Results." *Cardiovasc Intervent Radiol* 47, no. 12 (Dec): 1750–62. doi: 10.1007/s00270-024-03752-7.

**INDICATIONS FOR USE** Embosphere Microspheres are indicated for arterial embolisation in patients undergoing treatment for hypervascular processes, including genicular artery embolisation (GAE) in patients with mild-to-moderate knee osteoarthritis who are refractory to standard-of-care first-line conservative treatment (e.g., physical therapy, medical therapy [NSAIDs, opioids], intra-articular corticosteroid injections).  
<sup>†</sup>Embosphere used in GAE for knee osteoarthritis is CE-marked under the EU Medical Device Regulation (MDR 2017/745).

Before using, refer to Instructions for Use for indications, contraindications, warnings, precautions, and directions for use.

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