

Medtronic

Onyx™ Liquid Embolic System

Features¹



Formulation	Product Code	Viscosity	Volume	Concentration of EVOH	Amount of Tantalum
Onyx-18	105-7200-060	18 cSt	1.5 mL	6%	39%
Onyx-34	105-7200-080	34 cSt	1.5 mL	8%	38%
Onyx-34L	105-7315-080	34 cSt	1.5 mL	8%	23%
Onyx-34L	105-7360-080	34 cSt	6 mL	8%	23%

EVOH: ethylene vinyl alcohol copolymer

cSt: centistokes

Concentration of EVOH calculated using g/mL DMSO

Amount of Tantalum calculated using w/w

1. DS98-005 Rev V Design Spec: DMSO & Onyx 18JUL16.

This material should not be considered the exclusive source of information, it does not replace or supersede information contained in the device manual(s).

Please note that the intended use of a product may vary depending on geographical approvals.

See the device manual(s) for detailed information regarding the intended use, the implant procedure, indications, contraindications, warnings, precautions, and potential adverse events.

For a MRI compatible device(s), consult the MRI information in the device manual(s) before performing a MRI.

If a device is eligible for eIFU usage, instructions for use can be found at Medtronic's website manuals.medtronic.com.

Manuals can be viewed using a current version of any major internet browser. For best results, use Adobe Acrobat® Reader with the browser.

Medtronic products placed on European markets bear the CE mark and the UKCA mark (if applicable).

For any further information, contact your local Medtronic representative and/or consult Medtronic's websites.

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