



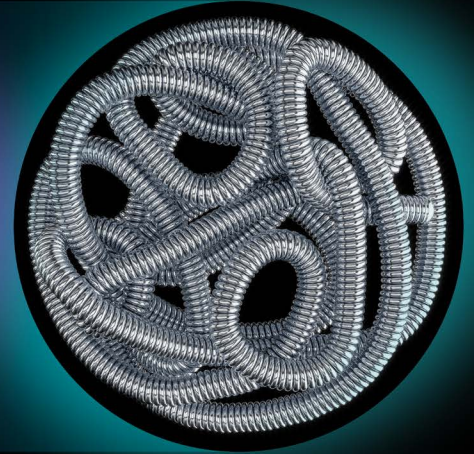
EMBOLD™ Detachable Coil System
SIMPLE BY DESIGN



FIBERED



PACKING



SOFT



**MULTI-CATHETER
COMPATIBILITY**
Accommodates .021" to
.027" ID microcatheters.
Fewer exchanges. Fewer
complications.



**HANDLE-FREE
DETACHMENT**
Provides complete control,
reduces procedure steps
and cost.



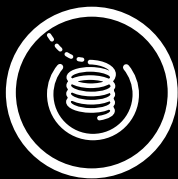
**KINKLESS
PERFORMANCE**
Nitinol delivery system
to ensure kinkless
performance and
resistance to premature
detachment.



**BEST-IN-CLASS
OCCLUSION POWER**
The most occlusion power in
its class, Embold offers the
greatest volume of any .021"
ID microcatheter compatible
coil and is purpose built for
exceptional deliverability.¹

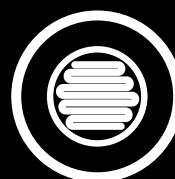
NEW BENEFIT

EMBOLD SOFT COIL
RELIABLE DISTAL OCCLUSION
Easily reach your target in
tortuous anatomy and quickly
embolize when you get there.
Avoid catheter kickback and
premature detachment.
Designed in unique lengths
for procedural flexibility.



NEW BENEFIT

EMBOLD PACKING COIL
PACK-TO-BLACK
A high-volume coil
that packs densely and
conforms to any vessel.²



1. The testing was performed by or on behalf of BSC. Data on file.

2. In clinical use cases, Embold Packing Coils are intended to be used with a backstop like Embold Fibered Coil.

EMBOLD™ Detachable Coil System

EMBOLD™ Fibered Coils			
Diameter (mm)	Length (cm)	Ref Catalog Number	GTIN
2	4	M001394240020040	08714729983521
	6	M001394240020060	08714729983538
3	6	M001394240030060	08714729983545
	12	M001394240030120	08714729983552
4	8	M001394240040080	08714729983569
	15	M001394240040150	08714729983576
5	8	M001394240050080	08714729983583
	15	M001394240050150	08714729983590
6	10	M001394240060100	08714729983606
	20	M001394240060200	08714729983613
8	20	M001394240080200	08714729983620
	20	M001394240100200	08714729983637
10	30	M001394240100300	08714729983644
	50	M001394240100500	08714729983651
	20	M001394240120200	08714729983668
12	30	M001394240120300	08714729983675
	50	M001394240120500	08714729983682
	20	M001394240140200	08714729983699
14	30	M001394240140300	08714729983705
	50	M001394240140500	08714729983712
18	50	M001394240180500	08714729983729
20	50	M001394240200500	08714729983736
22	60	M001394240220600	08714729983743
32	60	M001394240320600	00191506010485

EMBOLD™ Soft Coils			
Diameter (mm)	Length (cm)	UPN	GTIN
2	2	M001394250020020	00191506026363
	4	M001394250020040	00191506026370
	6	M001394250020060	00191506026387
3	8	M001394250020080	00191506026394
	4	M001394250030040	00191506026400
	8	M001394250030080	00191506026417
4	12	M001394250030120	00191506026424
	15	M001394250030150	00191506026431
	8	M001394250040080	00191506026448
5	10	M001394250040100	00191506026455
	15	M001394250040150	00191506026462
	30	M001394250040300	00191506026479
6	10	M001394250050100	00191506026486
	15	M001394250050150	00191506026493
	20	M001394250050200	00191506026509
8	30	M001394250050300	00191506026516
	15	M001394250060150	00191506026523
	20	M001394250060200	00191506026530
8	30	M001394250060300	00191506026547
	60	M001394250060600	00191506026554
	30	M001394250080300	00191506026561
8	60	M001394250080600	00191506026578

EMBOLD™ Packing Coils			
Diameter (mm)	Length (cm)	UPN	GTIN
N/A	15	M001394260000150	00191506026585
	30	M001394260000300	00191506026592
	45	M001394260000450	00191506026608
	60	M001394260000600	00191506026615

EMBOLD™ (SOFT/PACKING/FIBERED) DETACHABLE COIL SYSTEM

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a physician. Rx only. Prior to use, please see the complete "Instructions for Use" for more information on Indications, Contraindications, Warnings, Precautions, Adverse Events, and Operator's Instructions.

INTENDED USE / INDICATIONS FOR USE: The EMBOLD (Soft/Packing/Fibered) Detachable Coil System is indicated to obstruct or reduce rate of blood flow in the peripheral vasculature. This device is not intended for coronary or neurovascular use. **CONTRAINDICATIONS:** There are no known contraindications. **WARNINGS:** The following warning statements provide important information for safe use of EMBOLD (Soft/Packing/Fibered) Detachable Coil System. Observe all warnings and precautions noted throughout these and other instructions relevant to the procedure. Failure to do so may result in patient injury or product damage. • Do not use the EMBOLD (Soft/Packing/Fibered) Detachable Coil System if it is damaged. • To reduce the risk of thromboembolic complications, it is critical that a continuous flow of appropriate flush solution be maintained between a) the microcatheter and guide catheter, and b) the microcatheter and any intraluminal device. • Due to the delicate nature of coils and the tortuosity that can be present in vasculature pathways, as well as various morphologies of vasculature, the coil may become stretched during delivery and repositioning. If stretching is observed or suspected, carefully remove the coil and microcatheter from the vasculature simultaneously. • Do not rotate the delivery wire more than 1 turn (360 degrees) during delivery of the EMBOLD (Soft/Packing/Fibered) Detachable Coil System. Excessive rotation of the delivery wire may damage the EMBOLD (Soft/Packing/Fibered) Detachable Coil System or may result in premature detachment of the coil. **PRECAUTIONS:** The following precaution statements provide important information for safe use of EMBOLD (Soft/Packing/Fibered) Detachable Coil System. Observe all warnings and precautions noted throughout these and other instructions relevant to the procedure. Failure to do so may result in patient injury or product damage. • To ensure safe delivery, never make sudden movements, do not force in case of resistance, and operate gently and with caution. • Ensure the microcatheter in use has the appropriate ID and length for the selected coil. • Do not advance the EMBOLD (Soft/Packing/Fibered) Detachable Coil System if it becomes lodged within the microcatheter. Determine the cause of the resistance and replace the microcatheter and coil if necessary. See the "Operational Instructions - Full System Removal" section for further instructions. • No tools are required to detach the coil. Using a tool could result in damage to the inner wire causing the coil to be difficult or impossible to release. • When in highly tortuous anatomy, increased resistance may be felt when retracting the inner wire and a greater retraction distance may be required to detach the coil. • Multiple embolization coils may be required to achieve the desired occlusion. • Replace microcatheters periodically during delivery of multiple coils or if increased resistance is noted during coil delivery. **ADVERSE EVENTS:** Potential adverse events which may be associated with the use of a coil and/or the peripheral embolization procedure include, but are not limited to: • Allergic reaction (device, contrast, medications, or other) • Arrhythmia • Bleeding/hemorrhage • Cerebrovascular accident (CVA) • Death • Embolism (air, plaque, thrombus, device, tissue, or other) • Hematoma • Infection/sepsis • Ischemia • Necrosis • Need for additional intervention or surgery • Nerve injury • Pain/discomfort • Post-embolization syndrome (PES) • Recanalization • Thrombus/thrombosis • Transient ischemic attack (TIA) • Vasospasm • Vessel trauma (arteriovenous fistula, dissection, injury, perforation, pseudoaneurysm, rupture, or other) **97085821 A.1**

Information for competitive devices excerpted from IFUs. Embold sourced from data on file. All trademarks are the property of their respective owners.

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