



Stryker DBM

Demineralized Bone Matrix Family



Validated: Post-sterilization, product from each lot is subjected to a definitive in vivo test to confirm osteoinductivity.

Handling: DBM utilizes a novel carrier which allows the product to be malleable at room temperature and increase in viscosity after implantation.

Options: DBM is available in three forms: gel, putty, and putty combined with cancellous chips. This allows the graft selection to be tailored to the needs of a given patient.

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Demineralized Bone Matrix Family

stryker®

Trauma & Extremities

DBM GEL

Size	Catalog Number
1cc	3102-1101
5cc	3102-1105
10cc	3102-1110

Note: Gel form comes preloaded in a syringe



DBM PUTTY

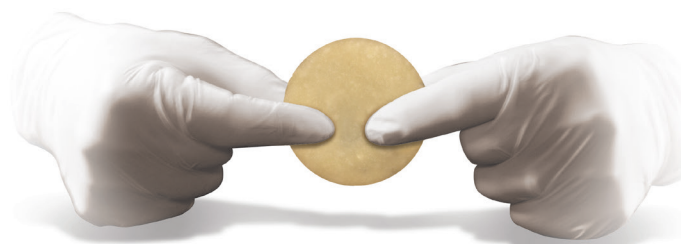
Size	Catalog Number
2.5cc	3102-1002
5cc	3102-1005
10cc	3102-1010



DBM PUTTY PLUS

Size	Catalog Number
5cc	3102-1205
10cc	3102-1210

Note: Putty form is combined with cancellous chips to produce Putty Plus



Stryker Quality. Stryker Experience. Stryker Trauma & Extremities.

This document is intended solely for the use of healthcare professionals. A surgeon must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. Stryker does not dispense medical advice and recommends that surgeons be trained in the use of any particular product before using it in surgery.

The information presented is intended to demonstrate the breadth of Stryker product offerings. A surgeon must always refer to the package insert, product label and/ or instructions for use before using any Stryker product. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your Stryker representative if you have questions about the availability of Stryker products in your area.

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