

Bard Mesh Flats and Pre-shaped

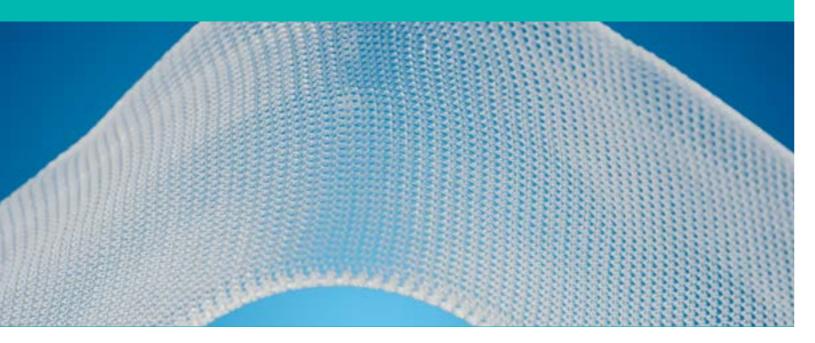
The Gold Standard Monofilament Polypropylene Mesh



With more than 40 years of clinical experience Bard Mesh is the gold standard

With Bard® Mesh, you've got time on your side. Since its introduction in 1962, Bard® Mesh has been considered the gold standard product to be used in a "tension-free" hernia repair technique. The primary cause for most recurrent hernias is the conventional technique of suturing together tissues that are not normally in apposition, which results in tension. Hernia repair with Bard Mesh is a tension-free repair technique that reinforces the weakened area, allowing for tissue ingrowth

and resiliency. Bard® Mesh can be tailored perioperatively and customized to any unique situation. Pre-shaped Bard® Mesh offers unparalleled convenience, while reducing overall waste and operating time. Because the mesh is pre-shaped, the need to trim the mesh perioperatively is minimized. An optional precut opening fits around the spermatic cord and the rounded edges accommodate to the anatomy of the inguinal canal.





Proven

Scientifically proven, with more than 40 years of clinical experience



Reliable

Safe and ideal for a tension-free hernia repair technique



Efficient

Encourages prompt tissue growth



Strong

t Strong repair¹



Functional

Available as flat or pre-shaped mesh

Ordering information

Bard Mesh			
Product code	Qty.	Dimensions	
0112640G	3/cs	1" x 4" (2.5 cm x 10 cm)	
0112650G	3/cs	2" x 4" (5 cm x 10 cm)	
0112660G	1/cs	10" x 14" (25 cm x 35.5 cm)	
0112670G	2/cs	2" x 12" (5 cm x 30.5 cm)	
0112680G	3/cs	3" x 6" (7.5 cm x 15 cm)	
0112720G	3/cs	6" x 6" (15 cm x 15 cm)	

Bard Mesh Pre-shaped

Product code	Qty.	Dimensions	
0112700	3/cs	Pre-shaped, 1.8" x 4" (4.5 cm x 10 cm)	
0112710	3/cs	Pre-shaped with keyhole, 1.8" x 4" (4.5 cm x 10 cm)	
0113700	3/cs	Large Pre-shaped, 2.4" x 5.4" (6 cm x 13.7 cm)	
0113710	3/cs	Large Pre-shaped with keyhole, 2.4" x 5.4" (6 cm x 13.7 cm)	



Bard® Mesh

Indications. Bard® Mesh is indicated for the repair of ventral, incisional and inguinal hernias. Contraindications. 1. Do not use this mesh in infants, children, or pregnant women, whereby future growth may be compromised by use of such mesh material 2. The use of this mesh has not been studied in breastfeeding or pregnant women.
3. Do not use this mesh for the reconstruction of cardiovascular defects. 4. Literature reports there may be a possibility for adhesion formation when the polypropylene is placed in direct contact with the bowel or viscera. **Warnings.** 1. The use of any permanent mesh or patch in a contaminated or infected wound could lead to fistula formation and/or extrusion of the mesh. 2. If an infection develops, treat the infection aggressively. Consideration should be given regarding the need to remove the mesh. An unresolved infection may require removal of the mesh. 3. If unused mesh has been in contact with instruments or supplies used on a patient or contaminated with body fluids, discard with care to prevent risk of transmission of viral infections. 4. To prevent recurrences when repairing hernias, the mesh should be sized with appropriate overlap for the size and location of the defect, taking into consideration any additional clinical factors applicable to the patient. Careful attention to mesh fixation placement and spacing will help prevent excessive tension or gap formation between the mesh and fascial tissue. 5. The mesh is supplied sterile. Inspect the packaging to be sure it is intact and undamaged prior to use. 6. This mesh has been designed for single use only Reuse, reprocessing, resterilization or repackaging may compromise the structural integrity and/or essential material and design characteristics that are critical to the overall performance of the mesh and may lead to mesh failure which may result in injury to the patient. Reuse, reprocessing, resterilization or repackaging may also create a risk of contamination of the mesh and/or cause patient infection or cross infection, including, but not limited to, the transmission of infectious diseases from one patient to another. Contamination of the mesh may lead to injury, illness or death of the patient or end user. 7. To avoid injury, careful attention is required if fixating the mesh in the presence of nerves, vessels or the spermatic cord. Fastener penetration into underlying tissue containing nerves or blood vessels may result in the need for medical/ surgical intervention, cause serious injury or permanent impairment to a body structure. 8. This mesh is not for the use of repair of pelvic organ prolapse via transvaginal approach. 9. This mesh is not for the use of treatment of stress urinary incontinence. **Precautions**. 1. Please read all instructions prior to use. 2. Only physicians qualified in the appropriate surgical techniques should use this mesh.

3. Intact Bard® Mesh exhibits high burst and tensile strength. However, when custom tailoring, in special circumstances where excessive force is placed on the mesh, the following guidelines may be helpful: • When cutting a notch in the mesh, a V-shape with a radiused point will withstand more force than a V-shape which comes to a sharp point. • For best results, it is recommended that the mesh be cut perpendicular to the selvage edge. • The inherent tensile strength of Bard® Mesh is strongest in the direction perpendicular to the selvage edges. Doubling the mesh may also increase the strength of the repair. Note: The selvage edges are recognized as the parallel, finished edges with a smooth appearance and slightly raised contour. **Adverse reactions**. Possible complications may include, but are not limited to, seroma, adhesions hematomas, pain, infection, inflammation, extrusion, erosion, migration fistula formation, allergic reaction and recurrence of the hernia or soft tissue defect.

Bard® Mesh Pre-shaped

Indications. Bard® Mesh Pre-shaped is indicated for the repair of inguinal hernia defects. **Contraindications**. 1. Do not use Bard® Pre-shaped mesh in infants or children, whereby future growth will be compromised by use of such mesh material. 2. Literature reports there may be a possibility for adhesion formation when Bard® mesh is placed in direct contact with the bowel or viscera, **Warnings**. 1. This device must be sterile before use. Please inspect the packaging to be sure it is intact and undamaged. 2. This device is for single use only. Do not resterilize or reuse any portion of Bard® Mesh Pre-shaped. 3. Careful attention to Bard® Pre-shaped mesh handling, fixation, and suture technique is most important in the presence of known or suspected wound contamination or infection. 4. The use of any permanent mesh or patch in a contaminated or infected wound can lead to fistula formation and/or extrusion of the prosthesis. 5. If an infection develops, treat the infection aggressively. Consideration should be given regarding the need to remove the mesh. An unresolved infection may require removal of the device. 6. To prevent recurrences when repairing inguina hernias, the mesh should be large enough to extend beyond the pubic tubercle and should fit securely around the spermatic cord at the internal ring. Many surgeons cut a keyhole in the mesh to allow for easier placement around the cord. 7. Reuse, reprocessing, resterilization or repackaging may compromise the structural integrity and/or essential material and design characteristics that are critical to the overall performance of the device and may lead to device failure which may result in injury to the patient. Reuse, reprocessing, resterilization or repackaging may also create a risk of contamination of the device and/or cause patient infection or cross infection, including, but not limited to, the transmission of infectious diseases from one patient to another. Contamination of the device may lead to injury, illness, or death of the patient or end user. 8. If unused prosthesis has been in contact with instruments or supplies used on a patient or contaminated with body fluids, discard with care to prevent risk of transmission of infection. **Precautions.** 1. Please read all instructions prior to use. 2. Only physicians qualified in the appropriate surgical techniques should use this prosthesis. 3. Intact Bard® Pre-shaped mesh exhibits high burst and tensile strength. However, when custom tailoring, in special circumstances where excessive force is placed on the mesh, the following guidelines may be helpful: When cutting a notch in the mesh, a V-shape with a radiused point will withstand more force than a V-shape which comes to a sharp point. 4. Davol[™] permanent or absorbable fixation devices or nonabsorbable monofilament sutures are recommended to properly secure the prosthesis. If absorbable fixation devices are used, they must be indicated for use in hernia repair. 5. Care should be taken to ensure that the mesh is adequately fixated to the uncompromised tissue of the inguinal floor. If necessary, additional fasteners and/or sutures should be used. **Adverse reactions**. Possible complications include seroma, adhesions, hematoma, inflammation, extrusion, fistula formation and recurrence of the hernia or soft tissue defect.

Please consult package insert for more detailed safety information and instructions for use.

¹ Amid, Shulman, Lichtenstein. Selecting synthetic mesh for the repair of groin hernia. Postgraduate General Surgery. 1992:4:150-155.

² Barnes JP. Inguinal repair with routine use of Marlex mesh. Surgery, Gynecology & Obstetrics. 1987;165:33-37.



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