INDICATIONS / INTENDED USE: The GORE® CARDIOFORM Septal Occluder is a permanently implanted device indicated for the percutaneous, transcatheter closure of ostium secundum atrial septal defects (ASDs).

CONTRAINDICATIONS: The GORE® CARDIOFORM Septal Occluder is contraindicated for use in patients: Unable to take anti-platelet or anticoagulant medications such as aspirin, heparin, or warfarin; with anatomy where the GORE® CARDIOFORM Septal Occluder size or position would interfere with other intracardiac or intravascular structures, such as cardiac valves or peripherally sized, with active endocarditis, or other infections preventing creation of a reliable point of attachment to the heart chamber; with known intracardiac thrombi. Refer to Instructions for Use at goremedical.com for a complete description of all warnings, precautions and adverse events.
The GORE® CARDIOFORM Septal Occluder is designed with two independent discs that span and cover the anatomy, enabling treatment of ASDs, including challenging defects.

- The soft and conformable construction of the frame is designed to reduce wall injury
- The minimal wire frame is designed to provide superior apposition to surrounding anatomy
- Proprietary, thromboresistant ePTFE material allows tissue ingrowth for short- and long-term performance
- A proven legacy of performance, with 15 years of experience and more than 25,000 occluder implants worldwide