• Polycaprolactone (PCL) is a biodegradable polymer that degrades and resorbs fully in vivo by hydrolysis which is then metabolized by the body.

• Osteoplug™ (-C) has a gradual resorption profile, depending on the patient anatomy and metabolism, of approximately 18-24 months.

• Osteoplug™ (-C) possesses optimal resorption rate that maintains mechanical integrity during healing process – minimizing adverse host-implant and inflammatory reactions.

• Osteoplug™ (-C) is manufactured with a porous interconnected micro-architecture that mimics the natural cancellous bone microstructure. It promotes tissue and vascular ingrowth.

• Osteoplug™ (-C) is an integrating implant to restore post-trephination burr hole defects by promoting tissue ingrowth, leading to a shift in burr hole reconstructive surgery from purely cosmesis to functional regeneration of damaged tissues.

• Osteoplug™ (-C) bears the CE mark of compliance, is FDA 510(k) cleared, fabricated in compliance with current Good Manufacturing Practice (cGMP, EN ISO 13485) and provided sterile (gamma irradiation, EN ISO 11137).
PATIENT’S PERSPECTIVE

• No patient developing adverse reactions such as pain, scarring, infections or excessive debris production.
• This implant has shown excellent functional and aesthetic outcomes.

CLINICAL PERSPECTIVE

• Implanted since 2004 with no complications when used according to its approved indications.
• Beyond 2 years of follow up shows host-implant compatibility with no infection and migration of implant.
• Avoided donor site morbidity.
• Designed to perfectly fit into burr holes formed by perforators, allowing easy handling by surgeons and ensured excellent cosmesis.
• Osteoplug™ (-C) were well-integrated into adjacent calvarial bone.

CLINICAL ADVANTAGE

3. INDICATIONS FOR USE

• Osteoplug™ (-C) is intended for use in the repair of neurosurgical burr holes and other cranial defects. It is used to aid closure of post trephination burr hole defects.
• Osteoplug™-C allows the insertion of a catheter after cerebral shunt operations.

4. SURGICAL PROTOCOL

1. SITE PREPARATION
   Prepare the implantation site using standard surgical techniques. Control of active bleeding should be achieved prior to implantation.

2. IMPLANT SELECTION
   Select the plug diameter that best suit perforator drill bit size.

3. IMPLANT PREPARATION
   Osteoplug™ (-C) is soaked with patient’s blood.

4. INSERTION
   Gently lower the Osteoplug™ into the burrhole. The Osteoplug™ would have a snug fit in the defect site. No fixation devices or suturing is required.

5. INSERTION OF CATHETER TUBE (FOR OSTEOPLUG™-C)
   After necessary surgical procedure, insert the catheter tube through hole in Osteoplug™-C.

6. HANDLING ADVANTAGE

• Osteoplug™ (-C) conforms to the defect, thus maximizing direct contact with viable host bone.
• Osteoplug™ (-C) is designed to perfectly fit into burr hole trephined using a perforator, allowing ease of use and streamlining the process of post-trephination cranial reconstruction.
• Osteoplug™ (-C) does not require fixation.

7. OSTEOPLUG™ (-C) SIZE CHART

Recommendation with commercial perforators

<table>
<thead>
<tr>
<th>BRAND</th>
<th>DRILL SIZE, OUTER DIAMETER (MM)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>8</td>
</tr>
<tr>
<td>Acracut</td>
<td>●</td>
</tr>
<tr>
<td>Adeor (Meridian)</td>
<td>●</td>
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<tr>
<td>Aesculap</td>
<td>●</td>
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<tr>
<td>Bojin</td>
<td>●</td>
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<td>Codman</td>
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<td>Aygun</td>
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<tr>
<td>Evonos</td>
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<td>Micromar</td>
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Recommended Osteoplug™ (-C) ● ● ● ● ● ● ● ● ●