

**K822318 ARGYLE MEDIASTINAL DRAIN**Aug 27, 1982  
24 days to decisionK822318 · Product code: **GBS** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k822318/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Ventricular, General & Plastic Surgery (GBS)
Date received	Aug 3, 1982
Decision date	Aug 27, 1982
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Sherwood Medical Co.</b>
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k822318/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 14, 2026