

**K802286 ARGLE LATEX PENROSE TUBING**Oct 23, 1980  
34 days to decisionK802286 · Product code: **KGZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k802286/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter (KGZ)
Date received	Sep 19, 1980
Decision date	Oct 23, 1980
Days to decision	34 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/k802286/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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