

**K882871 CASF DRILL GUIDE DG 170**Sep 23, 1988  
74 days to decisionK882871 · Product code: **LYQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k882871/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Accessories, Fixation, Spinal Intervertebral Body (LYQ)
Date received	Jul 11, 1988
Decision date	Sep 23, 1988
Days to decision	74 days
Third-party review	No

**APPLICANT**

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Company	<b>Terray Manufacturing, Inc.</b>
Location	Ontario K7s 1I8, CA
Contact	JUDY WILSON
510(k) history	9 submissions · 8 cleared · 1987-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882871/>, 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 22, 2026