

K960866 LRM PRODUCES GUIDEWIRES ON AN OEM BASIS FOR MANUFACTURERS, KIT ASSEMBLERS, AND DISTRIBUTORS.

Apr 5, 1996
32 days to decision

K960866 · Product code: **KNY** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k960866/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Mar 4, 1996
Decision date	Apr 5, 1996
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lake Region Mfg., Inc.
Location	Mchenry, IL, US
Contact	KIM E AVES
510(k) history	42 submissions · 42 cleared · 1977-2010

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k960866/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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