

K012023 ANCHOR GUIDE LESION LOCALIZATION DEVICESep 12, 2001
76 days to decisionK012023 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k012023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Jun 28, 2001
Decision date	Sep 12, 2001
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary
Other names	ANCOR GUIDE HOLDER; RF GENERATOR CABLE

APPLICANT

Company	Senorx, Inc.
Location	Irvine, CA, US
Contact	AMY BOUCLY
510(k) history	30 submissions · 26 cleared · 2000-2023

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