FDA 510(k) Premarket Notification

K110739 ENTELLUS MEDICAL SINUS GUIDEWIRE

CLEARED

Jun 14, 2011
89 days to decision

K110739 · Product code: **LRC** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k110739/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ent Manual Surgical (LRC)
Date received	Mar 17, 2011
Decision date	Jun 14, 2011
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Entellus Medical, Inc.**
Location	Maple Grove, MN, US
Contact	KAREN E PETERSON
510(k) history	27 submissions · 27 cleared · 2008-2022