FDA 510(k) Premarket Notification

K921071 DENTOLOCK

CLEARED

Mar 26, 1992
20 days to decision

K921071 · Product code: **KOB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k921071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Suprapubic (and Accessories) (KOB)
Date received	Mar 6, 1992
Decision date	Mar 26, 1992
Days to decision	20 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dentmed , Ltd.**
Location	Chicago, IL, US
Contact	ROBERT W BAUER
510(k) history	2 submissions · 2 cleared · 1992-1994