

K921071 DENTOLOCKMar 26, 1992
20 days to decisionK921071 · Product code: **KOB** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921071/> Data sourced from FDA 510(k) public records (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. - Generated July 2, 2026