

K933941 MENLO CARE URINARY DIVERSION STENTMar 31, 1994
231 days to decisionK933941 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k933941/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Aug 12, 1993
Decision date	Mar 31, 1994
Days to decision	231 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Menlo Care, Inc.
Location	Menlo Park, CA, US
Contact	BOB BLEY
510(k) history	31 submissions · 25 cleared · 1986-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933941/> 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 1, 2026