FDA 510(k) Premarket Notification

K963506 TENET RADIOLUCENT HAND TABLE

CLEARED

Jan 9, 1997
128 days to decision

K963506 · Product code: **FQO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k963506/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	Sep 3, 1996
Decision date	Jan 9, 1997
Days to decision	128 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tenet Medical Engineering**
Location	Calgary, CA
Contact	KEN MOORE
510(k) history	3 submissions · 3 cleared · 1996-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k963506/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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