FDA 510(k) Premarket Notification

K932280 RAE(R) II TRACHEAL TUBE

CLEARED

Jul 8, 1993
57 days to decision

K932280 · Product code: **BTR** · Anesthesiology
Source: https://www.510kdatabase.net/k932280/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tracheal (w/wo Connector) (BTR)
Date received	May 12, 1993
Decision date	Jul 8, 1993
Days to decision	57 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mallinckrodt Medical**
Location	St Louis, MO, US
Contact	LYNN C STIMPSON
510(k) history	43 submissions · 42 cleared · 1990-2008

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k932280/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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