FDA 510(k) Premarket Notification

K850240 KM PRODUCTS SUCTION TIPS

CLEARED

Mar 6, 1985
43 days to decision

K850240 · Product code: **JOL** · General Hospital
Source: https://www.510kdatabase.net/k850240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter And Tip, Suction (JOL)
Date received	Jan 22, 1985
Decision date	Mar 6, 1985
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Km Products**
Location	Tucson, AZ, US
Contact	ALAN R KENT
510(k) history	3 submissions · 3 cleared · 1985-1985

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k850240/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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