FDA 510(k) Premarket Notification

K844873 PENNINE GUEDEL AIRWAY 00,0,1,2,3 & 4

CLEARED

Feb 21, 1985
66 days to decision

K844873 · Product code: **CAE** · Anesthesiology
Source: https://www.510kdatabase.net/k844873/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Airway, Oropharyngeal, Anesthesiology (CAE)
Date received	Dec 17, 1984
Decision date	Feb 21, 1985
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Pennine Medical , Ltd.**
Location	Derby, GB
Contact	TERRY MORGAN
510(k) history	8 submissions · 8 cleared · 1985-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k844873/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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