FDA 510(k) Premarket Notification

K882219 P.E.E.P. VALVE

CLEARED

Aug 30, 1988
96 days to decision

K882219 · Product code: **BYE** · Anesthesiology
Source: https://www.510kdatabase.net/k882219/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Attachment, Breathing, Positive End Expiratory Pressure (BYE)
Date received	May 26, 1988
Decision date	Aug 30, 1988
Days to decision	96 days
Third-party review	No

APPLICANT

Company	**Life Support Products, Inc.**
Location	Anaheim, CA, US
Contact	TOM FANGROW
510(k) history	20 submissions · 20 cleared · 1985-1995