FDA 510(k) Premarket Notification

K880725 SAM(TM) MODEL ONE

CLEARED

Sep 6, 1988
196 days to decision

K880725 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k880725/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Feb 23, 1988
Decision date	Sep 6, 1988
Days to decision	196 days
Third-party review	No

APPLICANT

Company	**Sentel Tech. C/O Criticare Systems, Inc.**
Location	Santee, CA, US
Contact	LAI, PH.D.
510(k) history	1 submissions · 1 cleared · 1988-1988