

K880725 SAM(TM) MODEL ONESep 6, 1988
196 days to decisionK880725 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880725/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026