

**K923402 CRYSTAL TRACE PIEZO RESPIRATORY EFFORT
SENSOR**Dec 30, 1992
173 days to decisionK923402 · Product code: **BZQ** · Anesthesiology
Source: <https://www.510kdatabase.net/k923402/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Jul 10, 1992
Decision date	Dec 30, 1992
Days to decision	173 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Pro-Tech, Inc.
Location	Woodinville, WA, US
Contact	RANDY CARRUTHERS
510(k) history	7 submissions · 6 cleared · 1992-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923402/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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