

K813507 INFALERTJan 12, 1982
28 days to decisionK813507 · Product code: **BZQ** · AnesthesiologySource: <https://www.510kdatabase.net/k813507/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Dec 15, 1981
Decision date	Jan 12, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Briox Technologies, Inc.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1982

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Device record: <https://www.510kdatabase.net/k813507/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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