

**K904432 SLEEP I/T-5 AND SLEEP I/T-8**Jul 10, 1991  
285 days to decisionK904432 · Product code: **BZQ** · AnesthesiologySource: <https://www.510kdatabase.net/k904432/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Sep 28, 1990
Decision date	Jul 10, 1991
Days to decision	285 days
Third-party review	No

**APPLICANT**

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Company	<b>Cns, Inc.</b>
Location	Eden Prairie, MN, US
Contact	JOHN STOLTE
510(k) history	14 submissions · 14 cleared · 1984-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904432/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026