

**K821890 CRITIKON #817 ESOPHAGEAL MULTI-PROBE**Aug 3, 1982  
39 days to decisionK821890 · Product code: **BZT** · AnesthesiologySource: <https://www.510kdatabase.net/k821890/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Esophageal, With Electrical Conductors (BZT)
Date received	Jun 25, 1982
Decision date	Aug 3, 1982
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>Critikon Company, LLC</b>
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821890/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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