FDA 510(k) Premarket Notification

K821890 CRITIKON #817 ESOPHAGEAL MULTI-PROBE

CLEARED

Aug 3, 1982
39 days to decision

K821890 · Product code: **BZT** · Anesthesiology
Source: https://www.510kdatabase.net/k821890/

SUBMISSION DETAILS

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

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