FDA 510(k) Premarket Notification

K842316 CRITIKON TRIPLE-LUMEN CENTRAL VENOUS

CLEARED

Jul 20, 1984
38 days to decision

K842316 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k842316/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 12, 1984
Decision date	Jul 20, 1984
Days to decision	38 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k842316/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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