FDA 510(k) Premarket Notification

K840841 RENEAU UNIT

CLEARED

Aug 7, 1984
165 days to decision

K840841 · Product code: **CBF** · Anesthesiology
Source: https://www.510kdatabase.net/k840841/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Hyperbaric (CBF)
Date received	Feb 24, 1984
Decision date	Aug 7, 1984
Days to decision	165 days
Third-party review	No

APPLICANT

Company	**Reneau, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984