FDA 510(k) Premarket Notification

K811397 ECA

CLEARED

Jul 2, 1981
45 days to decision

K811397 · Product code: **FOA** · General Hospital
Source: https://www.510kdatabase.net/k811397/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Board, Cardiopulmonary (FOA)
Date received	May 18, 1981
Decision date	Jul 2, 1981
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Gillco, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1981-1981