

K823125 CAPIOX IINov 24, 1982
30 days to decisionK823125 · Product code: **DTZ** · Cardiovascular
Source: <https://www.510kdatabase.net/k823125/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oxygenator, Cardiopulmonary Bypass (DTZ)
Date received	Oct 25, 1982
Decision date	Nov 24, 1982
Days to decision	30 days
Third-party review	No

APPLICANT

Company	Terumo Medical Corp.
Location	Elkton, MD, US
510(k) history	143 submissions · 143 cleared · 1980-2011

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k823125/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 28, 2026