FDA 510(k) Premarket Notification

K830994 REDOX CONTROLLER

CLEARED

May 27, 1983
59 days to decision

K830994 · Product code: **KDN** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k830994/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Perfusion, Kidney (KDN)
Date received	Mar 29, 1983
Decision date	May 27, 1983
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**St. Louis Univ. Medical Center**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983