

K830994 REDOX CONTROLLERMay 27, 1983
59 days to decisionK830994 · 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

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Device record: <https://www.510kdatabase.net/k830994/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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