

K810823 ERIKA EXTENSION SETSApr 17, 1981
24 days to decisionK810823 · Product code: **FJK** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k810823/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Tubing, Blood, With And Without Anti-regurgitation Valve (FJK)
Date received	Mar 24, 1981
Decision date	Apr 17, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Erika, Inc.
Location	McHenry, IL, US
Website	https://www.erika.com
510(k) history	43 submissions · 43 cleared · 1976-1985

Erika, Inc. is a medical device company based in McHenry, US. The company specialized in Gastroenterology & Urology devices. Erika, Inc. received FDA 510(k) clearances from total submissions between 1976 and 1985. The company's regulatory focus centered on Gastroenterology & Urology devices, which represented 86% of its submission portfolio. Notable cleared products included infusion pump administration sets, artificial kidney filtration systems, and bicarbonate concentrate formulations. This company is inactive and represents a historical regulatory record. No FDA 510(k)...
