FDA 510(k) Premarket Notification

K830572 CLEAN-FLUSH

CLEARED

Mar 17, 1983
22 days to decision

K830572 · Product code: **KYZ** · General Hospital
Source: https://www.510kdatabase.net/k830572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Irrigating (non Dental) (KYZ)
Date received	Feb 23, 1983
Decision date	Mar 17, 1983
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Burron Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 40 cleared · 1979-1987