FDA 510(k) Premarket Notification

K893803 FEVER LINE AND FEVER LINE PLUS

CLEARED

Aug 21, 1989
91 days to decision

K893803 · Product code: **KPD** · General Hospital
Source: https://www.510kdatabase.net/k893803/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Strip, Temperature, Forehead, Liquid Crystal (KPD)
Date received	May 22, 1989
Decision date	Aug 21, 1989
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**Liquid Crystal Sciences, Inc.**
Location	Atlanta, GA, US
Contact	L BRETT
510(k) history	1 submissions · 1 cleared · 1989-1989