FDA 510(k) Premarket Notification

K791706 BE-SURE

CLEARED

Oct 17, 1979
61 days to decision

K791706 · Product code: **JHJ** · Chemistry
Source: https://www.510kdatabase.net/k791706/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agglutination Method, Human Chorionic Gonadotropin (JHJ)
Date received	Aug 17, 1979
Decision date	Oct 17, 1979
Days to decision	61 days
Third-party review	No

APPLICANT

Company	**Precision Products Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979