FDA 510(k) Premarket Notification

K790646 HELENA QUIPLATE SYSTEM

CLEARED

May 23, 1979
54 days to decision

K790646 · Product code: **JZQ** · Immunology
Source: https://www.510kdatabase.net/k790646/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plates And Equipment, Radial Immunodiffusion (JZQ)
Date received	Mar 30, 1979
Decision date	May 23, 1979
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Helena Laboratories**
Location	Mchenry, IL, US
510(k) history	280 submissions · 280 cleared · 1978-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790646/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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