FDA 510(k) Premarket Notification

K810119 CONTROL SERUMS FOR NONTREPONEMAL TEST

CLEARED

Feb 9, 1981
24 days to decision

K810119 · Product code: **GMP** · Toxicology
Source: https://www.510kdatabase.net/k810119/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Control For Nontreponemal Tests (GMP)
Date received	Jan 16, 1981
Decision date	Feb 9, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Hillside Acres, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981