

K810119 CONTROL SERUMS FOR NONTREPONEMAL TESTFeb 9, 1981
24 days to decisionK810119 · 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

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Device record: <https://www.510kdatabase.net/k810119/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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