

K810301 CRITHIDIA LUCILIAE NATIVE DNA-IFA TESTFeb 25, 1981
21 days to decisionK810301 · Product code: **DHN** · Immunology
Source: <https://www.510kdatabase.net/k810301/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antinuclear Antibody, Indirect Immunofluorescent, Antigen, Control (DHN)
Date received	Feb 4, 1981
Decision date	Feb 25, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Immuno Concepts, Inc.
Location	Mchenry, IL, US
510(k) history	36 submissions · 36 cleared · 1981-2001

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Device record: <https://www.510kdatabase.net/k810301/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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