

K811721 H-101 KAPPA,H-102 LAMBDA,H-107 LYSOZMEJul 16, 1981
28 days to decisionK811721 · Product code: **DFH** · Pathology
Source: <https://www.510kdatabase.net/k811721/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kappa, Antigen, Antiserum, Control (DFH)
Date received	Jun 18, 1981
Decision date	Jul 16, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	ImmuloK, Inc.
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k811721/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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