

K801479 GAMBRO LUNDIA PLATE HIGH FLUX HEMODIAL-Aug 27, 1980
64 days to decisionK801479 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k801479/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jun 24, 1980
Decision date	Aug 27, 1980
Days to decision	64 days
Third-party review	No

APPLICANT

Company	Gambro, Inc.
Location	Mchenry, IL, US
510(k) history	86 submissions · 86 cleared · 1976-2009

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Device record: <https://www.510kdatabase.net/k801479/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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