

**K800556 DIFFERENTIAL REFRACTIVE INDEX DETECTOR**Mar 25, 1980  
13 days to decisionK800556 · Product code: **KIE** · Toxicology  
Source: <https://www.510kdatabase.net/k800556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, High Pressure Liquid Chromatography (KIE)
Date received	Mar 12, 1980
Decision date	Mar 25, 1980
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.hp.com">https://www.hp.com</a>
510(k) history	230 submissions · 229 cleared · 1976-2000

Hewlett-Packard Co. is a technology company headquartered in McHenry, US. The company historically developed medical devices alongside its core computing and printing business. Hewlett-Packard received FDA 510(k) clearances from total submissions, with clearances spanning 1976 to 2000. The company specialized in cardiovascular devices, including defibrillators, telemetry systems, and clinical information systems. Additional cleared devices covered gastroenterology, urology, and radiology applications. This regulatory record reflects the company's historical involvement in...

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Device record: <https://www.510kdatabase.net/k800556/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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