

**K801722 CMV BIO-BEAD TITRATION KIT**Sep 16, 1980  
55 days to decisionK801722 · Product code: **GQH** · Microbiology  
Source: <https://www.510kdatabase.net/k801722/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antigen, Cf (including Cf Control), Cytomegalovirus (GQH)
Date received	Jul 23, 1980
Decision date	Sep 16, 1980
Days to decision	55 days
Third-party review	No

**APPLICANT**

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Company	<b>Litton Bionetics</b>
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1978-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k801722/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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