

**K201606 LacriJet®**Dec 18, 2020  
186 days to decisionK201606 · Product code: **OKS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k201606/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lacrimal Stents And Intubation Sets (OKS)
Date received	Jun 15, 2020
Decision date	Dec 18, 2020
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fci (France Chirurgie Instrumentation) Sas</b>
Location	Paris, FR
Contact	Thierry Fetick
510(k) history	4 submissions · 4 cleared · 2020-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201606/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026