

K031629 CANON NON-MYDRIATIC RETINAL CAMERA, MODEL CR-DGIJun 6, 2003
10 days to decisionK031629 · Product code: **HKI** · Ophthalmic
Source: <https://www.510kdatabase.net/k031629/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	May 27, 2003
Decision date	Jun 6, 2003
Days to decision	10 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Canon USA, Inc.
Location	Mchenry, IL, US
Contact	SHEILA DRISCOLL
510(k) history	48 submissions · 48 cleared · 1984-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k031629/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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