

**K812441 DI-CROWN**Oct 6, 1981  
41 days to decisionK812441 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k812441/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Aug 26, 1981
Decision date	Oct 6, 1981
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Coburn Optical Ind., Inc.</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1978-1989

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

Device record: <https://www.510kdatabase.net/k812441/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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