

**K820877 AUTOMATIC OBJECTIVE REFRACTOR 2000**Jun 25, 1982  
87 days to decisionK820877 · Product code: **HKO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k820877/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Mar 30, 1982
Decision date	Jun 25, 1982
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Dicon, Inc.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1982

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Device record: <https://www.510kdatabase.net/k820877/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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