

K820325 STERICIDALApr 21, 1982
72 days to decisionK820325 · Product code: **HPX** · Ophthalmic
Source: <https://www.510kdatabase.net/k820325/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Contact (polymethylmethacrylate) (HPX)
Date received	Feb 8, 1982
Decision date	Apr 21, 1982
Days to decision	72 days
Third-party review	No

APPLICANT

Company	Breger Mueller West
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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Device record: <https://www.510kdatabase.net/k820325/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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