FDA 510(k) Premarket Notification

# K820325 STERICIDAL

**CLEARED**

Apr 21, 1982
72 days to decision

K820325 · Product code: **HPX** · Ophthalmic
Source: https://www.510kdatabase.net/k820325/

## SUBMISSION DETAILS

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|---|---|
| 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 |