

**K830780 KOALA C99 I.V. CONTROLLER**Apr 8, 1983  
28 days to decisionK830780 · Product code: **LDR** · General Hospital  
Source: <https://www.510kdatabase.net/k830780/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Controller, Infusion, Intravascular, Electronic (LDR)
Date received	Mar 11, 1983
Decision date	Apr 8, 1983
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Chesebrough-Pond&amp;apos;S U.S.A. Co.</b>
Location	Mchenry, IL, US
510(k) history	33 submissions · 30 cleared · 1976-1987

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

Device record: <https://www.510kdatabase.net/k830780/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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