

**K782148 HR UNIT**Jan 8, 1979  
17 days to decisionK782148 · Product code: **IQP** · Physical MedicineSource: <https://www.510kdatabase.net/k782148/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Rotator, Transverse (IQP)          |
| Date received         | Dec 22, 1978                       |
| Decision date         | Jan 8, 1979                        |
| Days to decision      | 17 days                            |
| Third-party review    | No                                 |

**APPLICANT**

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|----------------|-----------------------------------------|
| Company        | <b>Parke-Davis Co.</b>                  |
| Location       | Mchenry, IL, US                         |
| 510(k) history | 47 submissions · 47 cleared · 1976-1986 |

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

Device record: <https://www.510kdatabase.net/k782148/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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