**Appendix Table F1. FDA review of medical devices for physical therapy interventions in adults with knee OA**

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| --- | --- | --- | --- |
| **Review ID** | **Medical device** | **Year** | **Link** |
| K983228 | Bionicare Stimulator System, Model BIO-1000 510(K) Summary | 1998 | http://www.accessdata.fda.gov/cdrh\_docs/pdf/K983228.pdf |
| K971437 | 510(K) Summary - Bionicare Stimulator System, Model BIO-1000 | 1997 | http://www.accessdata.fda.gov/cdrh\_docs/pdf/K971437.pdf |
| K062325 | 510(K) Summary RS-4i Sequential Stimulator | 2007 | http://www.accessdata.fda.gov/cdrh\_docs/pdf6/K062325.pdf |
| K052625 | BioniCare®, Stimulator Model BIG-IOOO TM | 2005 | http://www.accessdata.fda.gov/cdrh\_docs/pdf5/K052625.pdf |
| K042912 | 510(k) Summary- InterX5000 | 2008 | http://www.accessdata.fda.gov/cdrh\_docs/pdf4/K042912.pdf |
| K032652 | 510(K) Summary RS-4i Muscle Stimulator | 2003 | http://www.accessdata.fda.gov/cdrh\_docs/pdf3/K032652.pdf |
| K030332 | 510(K) Summary BioniCare Stimulator, Model BIO- 1OOO | 2003 | http://www.accessdata.fda.gov/cdrh\_docs/pdf3/K030332.pdf |