

Guidant and Medtronic implanted cardiac defibrillators and pacemakers recalled



Nearly 200,000 electronic heart devices, comprised of many different model names and numbers, are being recalled because of manufacturing defects that can cause life-threatening device failure.

Implanted cardiac defibrillators, also called cardioverters or ICD's, made both by Medtronic and Guidant, along with Medtronic pacemakers, are included in the recalls.

Contact us for information about the models covered and about the legal rights of people affected by defects in these devices.

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