

MDL-1708 – In re: Guidant Corp. Implantable Defibrillators Products Liability Litigation

MDL SETTLEMENT CONSIDERATION FORM

EVIDENCE OF INJURY FROM MALFUNCTION – If you believe that your device ever malfunctioned and injured you, please fill out this page. The malfunction that caused you injury (1) may be related to the recall for the device *or* may have led to any explant; and (2) must have resulted in the device failing to monitor or failing to provide therapy prior to explant. Provide a complete explanation of how you were injured and how the device malfunctioned. Provide particular detail if you believe your device malfunctioned through one of the modes of failure that led to a recall. If you are unsure of the mode of failure that led to recall of your device model, please see the information provided at www.guidantsettlement.com.

Please note that an explantation alone is not evidence of injury *due to* malfunction. Nor is the firing of a device, alone, evidence of a malfunction – heart devices are designed to fire and adjustments in how often they fire can be typical and normal. Further, explant for normal battery depletion is not evidence of a malfunction. All devices will be explanted at some point because of battery depletion.

The above information is true and correct to the best of my knowledge.

Signature

Printed Name

Date