

## URGENT FIELD SAFETY NOTICE

## Medtronic Model 5348 and Vitatron MEP3000 Single-Chamber External Pulse Generator

## Medical Device Recommendations

Medtronic/Vitatron Reference FA612

29 May 2014

Dear Health Care Professional (physician, Hospital Administrator, OR Manager, and Risk Manager),

Medtronic has identified a performance issue potentially affecting older Medtronic Model 5348 and Vitatron MEP3000 Single-Chamber External Pulse Generators (EPGs) manufactured between July 1995 and December 2007. You are receiving this communication as Medtronic/Vitatron records indicate your facility has received one or more potentially affected 5348 and/or MEP3000 EPGs. This issue does not affect any other Medtronic or Vitatron EPG models or any Medtronic or Vitatron implantable devices.

Issue Description: Through April 16, 2014, Medtronic has determined 49 events (out of approximately 30,000 potentially affected EPGs, or 0.16 percent), were related to a pacing rate outside of the intended setting, including events of sudden increased pacing rate up to the maximum setting of 180 pulses per minute (ppm). Of these 49 events, one was associated with a patient death with no other reports of critical injuries. An additional 85 reports of pacing rate outside of the intended setting have been received that may be related to this issue, but could not be confirmed due to insufficient data.

**Root Cause:** The root cause of this issue is the development of high resistance on internal electrical connector contacts due to oxidation over time.

Potentially affected Medtronic 5348 and Vitatron MEP3000 EPGs are within the Serial Number Ranges of:

PEP001001P to PEP050019P and PEP001001K to PEP001714K

Malfunction Indications: Due to the unpredictable nature of the oxidation process on multiple electrical contacts, this issue may result in one or more of the following observations:

- Pacing rate outside of the intended setting, potentially including a sudden increase in pacing rate up to the maximum setting of 180 ppm.
- · Output amplitude or sensitivity outside of intended setting.
- Pace, Sense, or Low Battery LED indicators not lighting during power on or reset functions.
- · Rapid Atrial Pacing (RAP) display with intermittent functionality.
- · Intermittent functionality of the On/Off and RAP control buttons.

Recommended Actions: After consulting with our Independent Physician Quality Panel, Medtronic and Vitatron recommends the following actions be taken when using a potentially affected Model 5348 and/or MEP3000 EPG:

- Monitor the EPG function and patient's heart rhythm continuously while the EPG is in use to ensure it is
  operating properly and delivering appropriate therapy to the patient.
- If any malfunction is observed with a 5348 and/or MEP3000 EPG, ensure the patient's condition is stabilized, discontinue use of the 5348 and/or MEP3000 EPG and contact your Medtronic /Vitatron representative.

Medtronic or Vitatron will no longer service or repair EPGs that are more than five years old, including these potentially affected 5348 and MEP3000 EPGs; which is consistent with the five-year service life of the new Model 5392 EPG. Medtronic will separately communicate additional details about this new EPG service policy. If you choose to replace your potentially affected 5348 and/or MEP3000 EPG, please contact your Medtronic /Vitatron representative for assistance with purchasing a replacement device.



## vitatron - The Pace Makers

Medtronic and Vitatron has communicated this information to the appropriate regulatory agencies and is committed to ensuring our products meet the highest quality standards and that our customers are fully supported.

For general questions related to EPG service policy or purchasing a replacement EPG, please contact your Medtronic /Vitatron representative; for technical questions, please call Our local representative Gulf Medical Co. Ltd Tel: +966569888444.

Please share this notification with others in your organization as appropriate. If product within the scope of this notification has been forwarded to another facility, please alert the facility of this notification.

Sincerely,

Fadi Faour CRDM Business Manager , MENA