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Dr Louise Mulroy
Medical Device Specialist
MHRA Adverse Incident Centre
Medicines & Healthcare products Regulatory Agency
Floor 4
151 Buckingham Palace Road
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12th March 2012

MHRA Ref 2012/001/030/031/001

Medical device adverse incident report from Nottingham University Hospitals Trust – City Campus
(their ref: 34655 NUHWEB18288 17/01/2012.

Dear Dr Mulroy

The following is further to my e-mailed letter dated 20th February 2012.

In conclusion:

- ◆ As far as we are aware there isn't anything in regard to the sensor manufacturing materials or manufacturing procedure that could be the cause of the incident.
- ◆ To our knowledge, to-date this is an isolated event; we have not heard of a similar event concerning our sensors.
- ◆ If the sensors are used in line with the concerned instructions for use then there should not be a reoccurrence; whether Viamed sensors are used or whether Draeger sensors are used.

If further information is available from the users - then of course we will freely assist in any further investigations.

In the meantime should you have any queries or require further information, please do not hesitate to contact me.

Yours sincerely

Steve Nixon
Director – Viamed Ltd.