

Medical Device Alert

Action

Ref: MDA/2009/037 Issued: 04 June 2009 at 14:30

Device

Knee replacement implant.
Preservation mobile bearing
unicompartmental knee
manufactured by DePuy
International Limited.



Problem

Revision rate higher than expected.

Action by

Orthopaedic surgeons, orthopaedic departments, and staff involved in the treatment and management of patients with joint replacement implants.

CAS deadlines

Action underway: 03 August 2009

Action complete: 03 December 2009

Action

- Identify patients implanted with affected devices.
- Consider undertaking clinical assessment of affected patients at least annually.

Contact

Manufacturer

Paul Arnott

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Link to full Medical Device Alert

<http://www.mhra.gov.uk/Publications/Safetywarnings/MedicalDeviceAlerts/CON049039>