

Medical Device Alert

Action

Ref: MDA/2010/093 Issued: 01 December 2010 at 14:30

Device

Contact lenses.

1-Day Acuvue TruEye (narafilecon A).

Manufactured by Johnson & Johnson Vision Care (Ireland).

Lot numbers with the first six digits within the ranges 492237 to 492498 inclusive and 502080 to 502269 inclusive.

Problem	Action
<p>Recall of specific product.</p> <p>Lenses from affected lots can cause stinging or pain upon insertion because of higher than expected levels of decanoic acid.</p>	<ul style="list-style-type: none"> Do not supply 1-Day Acuvue TruEye contact lenses as identified in the manufacturer's Field Safety Notice. Return all unused product to Johnson & Johnson Vision Care. Consider contacting end users who have received affected product, requesting that they return unused product to their eye care professional. Report any adverse incidents to the manufacturer and the MHRA.
<h3>Action by</h3> <p>Ophthalmologists, opticians and optometrists who:</p> <ul style="list-style-type: none"> supply contact lenses manage individuals who wear contact lenses. 	
CAS deadlines	Contact
<p>Action underway: 08 December 2010</p> <p>Action complete: 06 January 2011</p>	<p>UK Distributor: Johnson & Johnson Vision Care Tel: 0800 328 9541 Email: trueeyesupport@its.jnj.com</p>

[Link to full Medical Device Alert](#)