

Notice Information: Medical Devices - Advisory 20 June 2006

Part 1. Product Information

a)	Title:	Reusable Needle Holders / Blood Tube Holders
b)	Product Name/Type:	Reusable Needle Holders / Blood Tube Holders
c)	Reference:	SN2006(05)
۹)	Manufacturer/Supplier:	Various

Part 2. Target Audience

a) Target Audience:

Accident and Emergency Department; All Wards; Biochemistry
Departments; Blood Transfusion Laboratories; Directors of Pathology
and Laboratory Services; District Nurses; Haematology Laboratories;
Immunology Laboratories; Infection Control Teams; General Medical
Practitioners; Medical Directors; Microbiology Laboratories; Outpatient
Departments; Phlebotomists; Risk Managers; Supplies Managers;
Theatre Managers

Part 3. Problem/Issue

a) Problem/Issue:

The risk of needle stick injuries and the risk of cross contamination of patient blood with blood borne pathogens from the use of reusable blood tube / needle holders.

Part 4. Background Information

a) Background Information:

It has been brought to the attention of the Irish Medicines Board (IMB) that in addition to the risk of needle stick injuries associated with reusable blood tube / needle holders there is a potential risk of crossinfection of patients with blood borne pathogens in circumstances where reusable blood tube / needle holders become contaminated and are not decontaminated before re-use. This risk has also been identified by professionals and manufacturers and has been highlighted by the US based Occupational Safety and Health Administration, US National Association of Phlebotomists, ECRI (Emergency Care Research Institute) amongst others. The IMB has determined that both single use and reusable devices are used in the variety of healthcare settings in Ireland e.g. GP surgeries, hospitals, clinics and Information obtained by the IMB from users nursing homes. suggests that, in the case of some reusable devices: The written information provided by the manufacturer is limited and does not always provide sufficient details of how the device can be adequately decontaminated between use. The practice of visual inspection between use of reusable devices for macroscopic evidence of contamination may not be sufficient in the prevention of cross contamination. Some reports suggest that manufacturers recommend devices be used on a finite number of occasions. The control and management of this recommendation may not be realistic in a healthcare setting. The IMB is continuing to following up these concerns with the relevant manufacturers.

Part 5. Action to be taken

a) Action to be taken:

The IMB recommends that all healthcare facilities conduct a review and analysis of the reusable blood tube / needle holders that are in use in their facility. In General: Determine the type of devices that are in use in your facility e.g. single use or reusable Examine the recommendations outlined by the devices. manufacturer in the instructions for use. Review / implement local policy for the use / decontamination of such devices. If necessary consult with the local / regional infection control team to ensure compliance with local policy on the use, safety and sterilisation of reusable blood tube / needle holders. For Single Use Devices: Ensure that procedures are in place to ensure that single use devices are only used once. Ensure that facilities are in place ensure the safe disposal of the device. For Reusable Devices: Ensure that all staff using these devices are familiar with the manufacturer's recommendations for decontamination. Ensure that the manufacturer's recommendations for decontamination are completed following every use. Ensure that your local / regional infection control team are satisfied with the level of decontamination between uses of reusable blood tube / needle holders. Ensure that procedures are in place to ensure the safe disposal of the reusable device.

Part 6. Enquiries

a) All enquiries should be made to:

Adverse incidents may be reported via the on-line vigilance reporting system, which can be accessed in the medical devices section of the IMB website www.imb.ie, or by completing a Medical Device User Adverse Incident Report Form, which is available on request from the IMB or may be downloaded from the IMB website. All adverse incidents relating to a medical device should be reported to the Medical Devices Department at the IMB: Irish Medicines Board Medical Devices Department Kevin O'Malley House Earlsfort Centre Earlsfort Terrace Dublin 2 If you have any enquiries, you may contact the Medical Devices Department at: Telephone: +353-1-6764971 Fax: +353-1-6344033 Email: medicaldevices@imb.ie Website: www.imb.ie SN2006(05): Reusable Needle Holders / **Blood Tube Holders**

Part 7. Keywords

a) Keywords:

Reusable Needle Holders / Blood Tube Holders