

## **Guidelines for the Blood Transfusion Services**

### **6.12: Component recall and traceability**

<http://www.transfusionguidelines.org/red-book/chapter-6/6-12>

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There must be a documented system available in each Blood Establishment whereby adverse effects caused by the administration of any component, or the identification of a component quality problem, can enable the recall, if appropriate, of all unused components derived from that donation or all donations which are a constituent of a component pool. Similarly, there must be a documented system in each Blood Centre for the recall of any component or constituent of a component pool where reasonable grounds exist for believing it could cause adverse effects.

Any recall of a component should lead to a thorough investigation with a view to preventing a recurrence.

A system must be in place that ensures that any transfused (or discarded) blood component can be linked to the original donation and donor from which it was derived.<sup>7</sup>