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HESSAGE FROM DR J HETTERS, DEFUTY CHIEF MEDICAL OFFICER DEPARTMENT OF REALTH
25 February 1998 CEK/CHO/98/5

> FURTHER PRECAUTIONARY MEASURES ON BLOOD PRODUCTS announced

Further precautionary measures to protect patients against the theoretical risk of contracting new variant cin from blood products were announced today.

The action follows three recalls of blood products last November because donors contributing to the plasma used in them subsequently developed notion. It was made clear then that the Government would follow the recommendations of the relevant scientific committees to maximise the safety of the blood supply.

The further precautionary measures were announced after advice today from the UK Committee on Safety of Medicines (CSM), which considered all the current data, including the conclusions of this week's Committee on Proprietary Medicinal Products (CPMP). The CEM advice offectively signals a review of the use of UK-squrged plasma, a component of blood used in the manufacture of a variety of blood products. The CSN will accordingly be looking at all products individually to ensure a safe and sufficient supply of blood products to the NHS.

The CEM also advised an extension of blood product recalls to include donors subsequently identified as being strongly suspected of having notif. Previous recalls were based on confirmed cases only.

THE MEASURES RECOMMENDED BY THE CSM ARE PRECAUTIONARY. They do NOT mean that UK blood and blood products are uncase. Doctors, HEALTH PROPESSIONALS AND PATIENTS CAN CONTINUE TO USE UK SOURCED PRODUCTS WITH CONFIDENCE. There is no evidence to show that nvcID can be transmitted via blood products or blood - the risk remains theoretical only.

Some blood products may now be manufactured from imported plasma including the clotting agent Factor VIII, used in the treatment of haemophilia, immunoglobuling which are used in the treatment of a range of diseases such as totanus and the prevention of harmolytic disease of the newborn, and albumin, used in the treatment of burns and serious accidents, and as a stabiliser in some vaccines. VACCINES CURRENTLY USED IN THE UK CHILDHOOD IMMUNISATION PROGRAMME DO NOT CONTAIN UK ALBUMIN.

The action is supported by all the UK Chief Hedical Officers.

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WHOLE BLOOD, USED IN SLOOP TRANSFUSIONS, IS NOT AFFECTED BY TODAY'S MEASURES but as a further precautionary measure, the National Blood Authority has already been instructed to prepare a strategy for the possible removal of white blood rells from donations by the process of leucodeplation, should it be required.

There is absolutely no question of any risk to blood donors of contracting aveJU through the donation process. Donated blood is vital to the work of the NHS and all donors are urged to continue their life-saving work.

The autcome of a review of the NHS's provision of the blood product Factor VIII, used in the treatment of haemophilia, was also announced today.

The MRS Executive will how be writing to all health authorities setting out arrangements to ensure that recombinant Factor VIII is made available to those children under the age of 16 who are not already receiving it, and to new patients.

If you have any quaries about this message further information can be obtained from Dr Kike McGovern (Tel:

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