Yellowcard

Medicines and Healthcare products Regulatory Agency (MHRA) is the executive Agency of the Department of Health and Social Care that acts on behalf of the Ministers to protect and promote public health and patient safety, by ensuring that medicines and medical devices meet appropriate standards of safety, quality and efficacy.

The Yellowcard Scheme is a system built for collecting and monitoring any information which concerns any safety concerns or incidents involving medicines and medical devices.

Reports can be made for all medicines including vaccines, blood factors and immunoglobulins, herbal medicines and homeopathic remedies, and all medical devices available on the UK market.

https://yellowcard.mhra.gov.uk/

Medical device adverse incidents

The term 'medical device' covers a broad range of products that are used in healthcare. They can be physical items or software which are used for the diagnosis, prevention, monitoring or treatment of illness or disability.

An adverse incident is an event that caused, or almost caused, an injury to a patient or other person, or a wrong or delayed diagnosis and treatment of a patient.

If your equipment has a fault you should let yellowcard know as soon as you can.

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