



Date: Thursday, 19th November 2020

Our Ref: MB/SH FOI 4502

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Re: Freedom of Information Request FOI 4502

We are writing in response to your request submitted under the Freedom of Information Act, received in this office on 22nd October 2020.

Your request was as follows:

I would like to obtain specific information regarding the products used for the purposes of antiseptic skin preparation prior to invasive surgery on patients undergoing procedures in The Walton Centre NHS Foundation Trust.

- 1) Please confirm if The Walton Centre NHS Foundation Trust follows NICE guidance 125 (Surgical Site Infections: Prevention and Treatment) when carrying out antiseptic skin preparation?
- a. If yes, please provide a copy of The Walton Centre NHS Foundation Trust s most up-to-date surgical site infection prevention protocol.
- 1) NICE Guidance 125 has not yet been fully implemented at The Walton Centre.
- a. N/A
- 2) How does The Walton Centre NHS Foundation Trust communicate the category difference between antiseptic products that are medicines vs biocides and their intended use?
- a. Please share any communication materials.
- 2) The Walton Centre do not specifically communicate the category difference between antiseptic products that are medicines vs biocides and their intended use
- a. N/A
- 3) Can you please specify which antiseptic skin preparation products are used in The Walton Centre NHS Foundation Trusts operating theatres for the purposes of prevention of surgical site infections?

Alcoholic and Aqueous Betadine
Alcoholic and Aqueous Chlorhexidine

- 4) Does The Walton Centre NHS Foundation Trust obtain patient consent prior to surgical procedures?
- a. If yes, please provide the written material that The Walton Centre NHS Foundation Trust uses to obtain consent
- 4) Yes
- a. Please find attached the consent forms (x3) used by The Walton Centre. In addition, The Walton Centre also uses materials such as procedure specific patient information leaflets and documented clinical consultations as part of the process of obtaining patient consent.









Please see our response above in blue.

Re-Use of Public Sector Information

All information supplied by the Trust in answering a request for information (RFI) under the Freedom of Information Act 2000 will be subject to the terms of the Re-use of Public Sector Information Regulations 2005, Statutory Instrument 2005 No. 1515 which came into effect on 1st July 2005.

Under the terms of the Regulations, the Trust will licence the re-use of any or all information supplied if being used in a form and for the purpose other than which it was originally supplied. This license for re-use will be in line with the requirements of the Regulations and the licensing terms and fees as laid down by the Office of Public Sector Information (OPSI). Most licenses will be free; however the Trust reserves the right, in certain circumstances, to charge a fee for the re-use of some information which it deems to be of commercial value.

Further information can be found at www.opsi.gov.uk where a sample license terms and fees can be found with guidance on copyright and publishing notes and a Guide to Best Practice and regulated advice and case studies, at www.opsi.gov.uk/advice/psi-regulations/index.htm

If you are dissatisfied with the handling of your request, you have the right to ask for an internal review. Internal review requests should be submitted within two months of the date of receipt of the response to your original letter and should be addressed to the Freedom of Information Office at the address above.

Please remember to quote the reference number, FOI 4502 in any future communications.

If you are not content with the outcome of the internal review, you have the right to apply directly to the Information Commissioner for a decision. The Information Commissioner can be contacted at: Information Commissioners Office, Wycliffe House, Water Lane, Wilmslow, Cheshire, SK9 5AF.

Yours sincerely

Mike Burns

Mr. Mike Burns, Executive Lead for Freedom of Information



