Yorkshire Specialist Register of Cancer in Children and Young People Research Database Privacy Notice and Fair Processing Statement

This statement explains how we use any personal information we collect about you / the child you care for.

*What is the legal basis for you collecting this information?*

As set out under the General Data Protection Regulation (GDPR), our lawful basis for processing your data under Article 6(e) and special category data under Article 9(2)(j) is “Public Task”. This means the processing of your/ your child’s data is necessary for us to perform a task in the public interest. This task has a clear basis in law in terms of relating to cancer registration.

A summary of the entry for YSRCCYP is available on the register of approvals on the HRA website (YSRCCYP appears as app no. 0062 and its reference is: CAG 1-07(b)/2014).

Furthermore, ethical approval for the register has been granted by the Northern and Yorkshire Multi Centre Research Ethics Committee (YSRCCYP reference MREC/00/3/001).

*What data are being collected?*

The Yorkshire Specialist Register of Cancer in Children and Young People, or YSRCCYP, aims to collect data on every child and young person aged <30 years diagnosed with cancer within the Yorkshire and Humber region. Data are collected about each individual for the whole period of their care; this includes any cancer diagnoses (malignant, benign and relapses), hospital visits, operations and details of chemotherapy and radiotherapy treatments.

Personal identifiers include:

* Full and previous names;
* Address history;
* Date of birth;
* NHS number.

*Where does this data come from?*

The data originates from a variety of sources, including:

* Hospital case notes (from National Health Service Trust hospitals);
* National Cancer Registration and Analysis Service (from Public Health England);
* Systemic Anti-Cancer Therapy dataset (from Public Health England);

Additional linked data sources include:

* Hospital episode statistics (from NHS Digital);
* Primary Care (from The Phoenix Partnership and EMIS Health);
* Social outcomes (Department for Work and Pensions);
* Educational outcomes (Department for Education).

In order to receive data from these sources, YSRCCYP may need to share an appropriate identifier(s) to enable data linkage, such as NHS number, name and date of birth.

*What are the data being used for?*

The database will be used to carry out a programme of epidemiological and applied health research investigating incidence, survival, aetiology and long-term health and social outcomes of cancer occurring in children and young people. No personal information will ever be made public in any report or publication.

*Where are the data held and for how long?*

All of the data are held locally at the University of Leeds, on encrypted hard drives, in a highly secure environment. As an ongoing register of cancer, we keep these data permanently so we can study the impact of treatment and care on long-term outcomes for survivors.

*Automated-decision making and profiling.*

YSRCCYP does not use automated-decision making. YSRCCYP does not use profiling to make a decision by automated means.

*Sharing data with other organisations*

YSRCCYP may share data held on its database with researchers (in the UK and worldwide), other registers and other health or social care providers in order to help improve patient care and examine health and social outcomes. Occasionally, a very limited set of personal data may be processed by an authorised third party such as NHS Digital to enable us to study health and social outcomes, but only with the appropriate regulatory permissions.

Identifiable data will not be shared outside of England and Wales.

*What rights do I have over my/my child’s data?*

You have the right of access, the right of rectification, the right to restriction of processing and the right to object to processing. If you wish to enact any of these rights, please contact the YSRCCYP team using the contact details below.

You may also wish to visit the national data opt-out website for further information about removing your personal information from national datasets: <https://digital.nhs.uk/services/national-data-opt-out>

**Contact Details**

**The Yorkshire Register key contacts:**

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Tel +44 (0)113 343 4841

https://medicinehealth.leeds.ac.uk/leeds-institute-cardiovascular-metabolic-medicine/doc/yorkshire-specialist-register-cancer-children-young-people

**Email:** [r.g.feltbower@leeds.ac.uk](mailto:r.g.feltbower@leeds.ac.uk)

**Professor Adam Glaser**

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LS1 3EX

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*Who is responsible for the data and who processes it?*

The **Data Controller and Data Processor** for data collected is:

**University of Leeds**

Woodhouse Lane

Leeds

LS2 9JT

**Data Protection Officer**: Alice Temple

*What if I have any other concerns about the way my personal data are processed?*

If you have any concerns with regard to the way your personal data are being processed or have a query with regard to this Notice, please contact our Data Protection Officer, Alice Temple via email at a.c.temple@leeds.ac.uk.

Our general postal address is University of Leeds, Woodhouse Lane, Leeds LS2 9JT, UK.

Our postal address for data protection issues is University of Leeds, Room 11.72 EC Stoner Building, Leeds, LS2 9JT. Our telephone number is +44 (0)113 2431751.

Our data controller registration number provided by the Information Commissioner's Office is Z553814X.

If you are unhappy with the way you or your child’s data are being processed you have the right to complain to the Information Commissioner’s Office (ICO).

<https://ico.org.uk/>