

GENERAL PRACTICE DATA FOR PLANNING AND RESEARCH (GPDfPR)

There has been much negativity in the recent press regarding a new initiative from NHS Digital to collect data from your GP practice.

The NHS needs data about the patients it treats to plan and deliver its services and to ensure that care and treatment provided is safe and effective. The General Practice Data for Planning and Research data collection will help the NHS to improve health and care services for everyone by collecting patient data that can be used to do this. For example, patient data can help the NHS to:

- Monitor the long-term safety and effectiveness of care
- Plan how to deliver better health and care services
- Prevent the spread of infectious diseases
- Identify new treatments and medicines through health research

GP practices already share patient data for these purposes, but this new data collection will be more efficient and effective.

This means that GPs can get on with looking after their patients, and NHS Digital can provide controlled access to patient data to the NHS and other organisations who need to use it, to improve health and care for everyone.

Contributing to research projects will benefit us all as better and safer treatments are introduced more quickly and effectively without compromising your privacy and confidentiality.

NHS Digital has engaged with the <u>British Medical Association (BMA)</u>, <u>Royal College of GPs</u> (<u>RCGP</u>) and the <u>National Data Guardian (NDG</u>) to ensure relevant safeguards are in place for patients and GP practices.

NHS DIGITAL PURPOSES FOR PROCESSING PATIENT DATA

Patient data from GP medical records kept by GP practices in England is used every day to improve health, care and services through planning and research, helping to find better treatments and improve patient care. The NHS is introducing an improved way to share this information - called the General Practice Data for Planning and Research data collection.

NHS Digital will collect, analyse, publish and share this patient data to improve health and care services for everyone. This includes:

- Informing and developing health and social care policy
- Planning and commissioning health and care services
- Taking steps to protect public health (including managing and monitoring the coronavirus pandemic)
- In exceptional circumstances, providing you with individual care

• Enabling healthcare and scientific research

Any data that NHS Digital collects will only be used for health and care purposes. It is never shared with marketing or insurance companies.

WHAT DOES THIS MEAN?

This collection will start from 1 September 2021. Patient data will be collected from GP medical records about:

- Any living patient registered at a GP practice in England when the collection started this includes children and adults
- Any patient who died after the data collection started, and was previously registered at a GP practice in England when the data collection started

NHS Digital will not collect your name or where you live. Any other data that could directly identify you, for example NHS number, General Practice Local Patient Number, full postcode and date of birth, is replaced with unique codes which are produced by de-identification software before the data is shared with NHS Digital.

This process is called pseudonymisation and means that no one will be able to directly identify you in the data. The diagram below helps to explain what this means. Using the terms in the diagram, the data we collect would be described as de-personalised.



Image provided by Understanding Patient Data <u>under licence</u>.

NHS Digital will be able to use the same software to convert the unique codes back to data that could directly identify you in certain circumstances, and where there is a valid legal reason. Only NHS Digital has the ability to do this. This would mean that the data became personally identifiable data in the diagram above. An example would be where you consent to your identifiable data being shared with a research project or clinical trial in which you are participating, as they need to know the data is about you.

THE DATA NHS DIGITAL COLLECT

NHS Digital will only collect structured and coded data from patient medical records that is needed for specific health and social care purposes explained above.

Data that directly identifies you as an individual patient, including your NHS number, General Practice Local Patient Number, full postcode, date of birth and if relevant date of death, is replaced with unique codes produced by de-identification software before it is sent to NHS Digital. This means that no one will be able to directly identify you in the data.

NHS Digital will collect:

- Data on your sex, ethnicity and sexual orientation
- Clinical codes and data about diagnoses, symptoms, observations, test results, medications, allergies, immunisations, referrals and recalls, and appointments, including information about your physical, mental and sexual
- Data about staff who have treated you

More detailed information about the patient data we collect is contained in the <u>Data Provision</u> <u>Notice issued to GP practices</u>.

Any other data that could directly identify patients (such as NHS number, date of birth, full postcode) is replaced with unique codes which are produced by de-identification software before the data is shared with NHS Digital.

This process is called pseudonymisation and means that patients will not be identified directly in the data. NHS Digital will be able to use the software to convert the unique codes back to data that could directly identify patients in certain circumstances, and where there is a valid legal reason.

NHS Digital Does not collect:

- Your name and address (except for your postcode in unique coded form)
- Written notes (free text), such as the details of conversations with doctors and nurses
- Images, letters and documents
- Coded data that is not needed due to its age for example medication, referral and appointment data that is over 10 years old
- Coded data that GPs are not permitted to share by law for example certain codes about IVF treatment, and certain information about gender re-assignment

HOW DOES NHS DIGITAL USE PATIENT DATA?

NHS Digital will analyse and link the patient data we collect with other patient data we hold to create national data sets and for data quality purposes.

NHS Digital will be able to use the de-identification software to convert the unique codes back to data that could directly identify you in certain circumstances for these purposes, where this is necessary and where there is a valid legal reason. There are strict internal approvals which need to



be in place before we can do this and this will be subject to independent scrutiny and oversight by the <u>Independent Group Advising on the Release of Data (IGARD)</u>.

These national datasets are analysed and used by NHS Digital to produce national statistics and management information, including public dashboards about health and social care which are published. We never publish any patient data that could identify you. All data we publish is anonymous statistical data.

For more information about data we publish see <u>Data and Information</u> and <u>Data Dashboards</u>.

We may also carry out analysis on national data sets for data quality purposes and to support the work of others.

WHO DOES NHS DIGITAL SHARE PATIENT DATA WITH?

All data which is shared by NHS Digital is subject to robust rules relating to privacy, security and confidentiality and only the minimum amount of data necessary to achieve the relevant health and social care purpose will be shared.

All requests to access patient data from this collection, other than anonymous aggregate statistical data, will be assessed by NHS Digital's <u>Data Access Request Service</u>, to make sure that organisations have a legal basis to use the data and that it will be used safely, securely and appropriately.

These requests for access to patient data will also be subject to independent scrutiny and oversight by the <u>Independent Group Advising on the Release of Data (IGARD)</u>. Organisations approved to use this data will be required to enter into a data sharing agreement with NHS Digital regulating the use of the data.

There are a number of organisations who are likely to need access to different elements of patient data from the General Practice Data for Planning and Research collection. These include but may not be limited to:

- The Department of Health and Social Care and its executive agencies, including Public Health England and other government departments
- NHS England and NHS Improvement
- Primary care networks (PCNs), clinical commissioning groups (CCGs) and integrated care organisations (ICOs)
- Local authorities
- Research organisations, including universities, charities, clinical research organisations that run clinical trials and pharmaceutical companies

If the request is approved, the data will either be made available within a secure data access environment within NHS Digital infrastructure, or where the needs of the recipient cannot be met this way, as a direct dissemination of data. We plan to reduce the amount of data being processed outside central, secure data environments and increase the data we make available to be accessed via our secure data access environment. For more information read about improved data access in <u>improving our data processing services</u>.



Data will always be shared in the uniquely coded form (de-personalised data in the diagram above) unless in the circumstances of any specific request it is necessary for it to be provided in an identifiable form (personally identifiable data in the diagram above). For example, when express patient consent has been given to a researcher to link patient data from the General Practice for Planning and Research collection to data the researcher has already obtained from the patient.

It is therefore possible for NHS Digital to convert the unique codes back to data that could directly identify you in certain circumstances, and where there is a valid legal reason which permits this without breaching the common law duty of confidentiality. This would include:

- Where the data was needed by a health professional for your own care and treatment
- Where you have expressly consented to this, for example to participate in a clinical trial
- Where there is a legal obligation, for example where the COPI Notices apply see <u>Our legal</u> <u>basis for collecting, analysing and sharing patient data</u> above for more information on this
- Where approval has been provided by the <u>Health Research Authority</u> or the Secretary of State with support from the <u>Confidentiality Advisory Group (CAG)</u> under Regulation 5 of the Health Service (Control of Patient Information) Regulations 2002 (COPI) - this is sometimes known as a 'section 251 approval'

This would mean that the data was personally identifiable in the diagram above. Re-identification of the data would only take place following approval of the specific request through the Data Access Request Service, and subject to independent assurance by IGARD and consultation with the Professional Advisory Group, which is made up of representatives from the BMA and the RCGP. If you have registered a National Data Opt-out, this would be applied in accordance with the National Data Opt-out policy before any identifiable patient data (personally identifiable data in the diagram above) about you was shared. More about the National Data Opt-out is in the section below.

WHERE DOES NHS DIGITAL STORES PATIENT DATA?

NHS Digital only stores and processes patient data for this data collection within the United Kingdom (UK).

Fully anonymous data (that does not allow you to be directly or indirectly identified), for example statistical data that is published, may be stored and processed outside of the UK. Some of our processors may process patient data outside of the UK. If they do, we will always ensure that the transfer outside of the UK complies with data protection laws.

OPTING OUT OF NHS DIGITAL COLLECTING YOUR DATA (TYPE 1 OPT-OUT)

If you do not want your identifiable patient data to be shared outside of your GP practice for purposes except for your own care, you can register an opt-out with your GP practice, not Taurus Healthcare as we are not your registered practice. This is known as a Type 1 Opt-out.

Type 1 Opt-outs were introduced in 2013 for data sharing from GP practices, but may be discontinued in the future as a new opt-out has since been introduced to cover the broader health and care system, called the National Data Opt-out. If this happens people who have registered a Type 1 Opt-out will be informed.



NHS Digital will not collect any patient data for patients who have already registered a Type 1 Optout in line with current policy. If this changes patients who have registered a Type 1 Opt-out will be informed.

If you do not want your patient data shared with NHS Digital, you can register a Type 1 Opt-out with your GP practice. You can register a Type 1 Opt-out at any time. You can also change your mind at any time and withdraw a Type 1 Opt-out.

Data sharing with NHS Digital will start on **1 September 2021**.

If you have already registered a Type 1 Opt-out with your GP practice your data will not be shared with NHS Digital.

If you wish to register a Type 1 Opt-out with your GP practice before data sharing starts with NHS Digital, this should be done by <u>returning this form</u> to your GP practice by **23 June 2021** to allow time for processing it. If you have previously registered a Type 1 Opt-out and you would like to withdraw this, you can also use the form to do this. You can send the form by post or email to your GP practice or call 0300 3035678 for a form to be sent out to you.

If you register a Type 1 Opt-out after your patient data has already been shared with NHS Digital, no more of your data will be shared with NHS Digital. NHS Digital will however still hold the patient data which was shared with us before you registered the Type 1 Opt-out.

If you do not want NHS Digital to share your identifiable patient data (personally identifiable data in the diagram above) with anyone else for purposes beyond your own care, then you can also register a <u>National Data Opt-out</u> which CANNOT be done through the GP practice. You can do this via the website, by phone, email or post.

- Telephone: 0300 303 5678
- Email: <u>enquiries@nhsdigital.nhs.uk</u>
- Postal address: National Data Opt Out, Contact Centre, NHS Digital, HM Government, 7 and 8 Wellington Place, Leeds. LD1 4AP
- Website: <u>National Data Opt Out</u>

If you wish to discuss this initiative further, do please ask at your registered GP Practice in the first instance.