Park Lane Surgery

Data Protection Impact Assessment

Document History

Document Reference:				
Document Purpose:	 The purpose is to have the potential to detect and mitigate information risks, as well as to modify plans accordingly. A DPIA should be completed when the following activities occur: Developing or procuring any new programme, policy, procedure, service, technology or system ("project") that handles or collects information relating to individuals. Developing revisions to an existing programme, policy, procedure, service, technology or system which significantly change how information is managed. 			
Date Approved:	6 April 2021			
Version Number:	4.0			
Status:	FINAL			
Next Revision Due:	April 2022			
Developed by:	Paul Couldrey – IG Consultant			
Policy Sponsor:	Practice Manager			
Target Audience:	This policy applies to any person directly employed, contracted, working on behalf of the Practice or volunteering with the Practice.			
Associated Documents:	All Information Governance Policies and the Information Governance Toolkit, and Data Security and Protections Toolkit 2020			

Data Protection Impact Assessment (DPIA)

When to carry out a DPIA

The DPIA identifies and assesses privacy implications where information (data) about individuals is collected, stored, transferred, shared, and managed. It should be process rather than output orientated.

The purpose is to have the potential to detect and mitigate information risks, as well as to modify plans accordingly.

A PIA should be completed when the following activities occur:

- Developing or procuring any new programme, policy, procedure, service, technology or system ("project") that handles or collects information relating to individuals.
- Developing revisions to an existing programme, policy, procedure, service, technology or system which significantly change how information is managed.

The UK General Data Protection Regulation (UK GDPR) became law on 25th May 2016, with the Data Protection Act 2018 replacing all previous data protection laws in the UK.

The Regulation in Article 35 (recitals **84**, **89**, **90**, **91**, **92**, **93**, **95**) makes it obligatory to perform a Data Protection impact assessment in case of large scale processing of special categories of data (as in this case health data and genetic data see article **9(1)**. This could help to ascertain the legal basis for processing, which will be helpful for public authorities now that the open door of 'legitimate interests' is closed. It is also important to note that "a single assessment may address a set of similar processing operations that present similar high risks". This could significantly help in reducing the administrative burden for hospitals and health and care providers when performing such an \\assessment.

A data protection impact assessment shall in particular be required in the case of:

- (a) a systematic and extensive evaluation of personal aspects relating to natural persons which is based on automated processing, including profiling, and on which decisions are based that produce legal effects concerning the natural person or similarly significantly affect the natural person;
- (b) processing on a large scale of special categories of data referred to in **Article 9(1)**, or of personal data relating to criminal convictions and offences referred to in Article 10; or

(c) a systematic monitoring of a publicly accessible area on a large scale.

This DPIA has been designed to meet the requirements of current legislation and common law duties and the expanded requirements of the UK GDPR as above, however Consent modelling / Fair Processing modification should be addressed by separate Trust UK GDPR action plans and strategies as several of the policies currently in use will need to be updated to reflect legislative changes.

Step 1 – I	Project Details
Project name/title	COVID Vaccine Invitation Project
initiative.	nclude how many individuals will be affected by the heir Coronavirus vaccine, once they become eligible
Details of any link to any wider initiative (if applicable)	This project will go ahead across the whole of Greater Derby and Derbyshire.
Stakeholder Analysis List those who may be affected (stake holder have been consulted prior to project start), eg. Service Users, Clients, Staff-managers and practitioners, Trade Unions, Visitors, Professional organisations, IT providers, Regulators and inspectorial bodies, MPs, Councillors, Partner organisations, Media, Carers	Internal: All staff External: Nil
Does the initiative involve the use of existing personal and/or confidential data: • For new purposes? • In different ways? If so, please explain (if not already covered above)	Uses existing personal data
Are potential new purposes likely to be identified as the scope of the initiative expands?	Likely to lead to a vaccine denied audit
What is already available? Any Previous PIA, Research or Consultation undertaken.	

Stop 1 - Project Details

Step 2 – Contacts

Who is completing this assessment?				
Name	Louis Wood			
Job Title	Assistant Practice Manager			
Department/Directorate name				
Contact address	2 Park Lane, DE22 2DS			
Email address	Louis.wood1@nhs.net			
Telephone number	01332 552461			
Connection to Project	Practice Project Lead			

Other person(s) with responsibility for this initiative e.g. Project Manager/Director, Senior Responsible Officer (SRO)					
Name	Lesley Hutchinson				
Job Title	Practice Manager				
Department/Directorate name					
Contact address	2 Park Lane, DE22 2DS				
Email address	Lesley.hutchinson@nhs.net				
Telephone number	01332 552461				
Connection to project	Vaccination Centre Lead Admin				

Technical Lead(s) (if relevant)					
Name					
Email address					
Telephone number					

Step 3 – Screening Questions

The	The purpose of these questions is to establish whether a full Privacy Impact Assessment is								
	necessary and to help to draw out privacy considerations								
		Yes	No	Unsure	Comments - document initial comments on privacy impacts or clarification for why this is not an issue or why you are unsure				
Ι	Is the information about	100		Uniouro	loode of why you are alloure				
	individuals likely to raise privacy concerns or expectations e.g. health records, criminal records or other information people would consider particularly private?								
ii	Will the initiative involve the collection of new information about individuals?				Collection of COVID vaccination refusals				
iii	Are you using information about individuals for a purpose it is not currently used for, or in a way it is not currently used?		\checkmark						
iv	Will the initiative require you to contact individuals in ways which they may find intrusive ¹ ?		\checkmark						
V	Will information about individuals be disclosed to organisations or people who have not previously had routine access to the information?				Sharing of patient information with Swiftqueue – a booking system for patients attending COVID vaccination appointments				
vi	Does the initiative involve you using new technology which might be perceived as being privacy intrusive e.g. biometrics or facial recognition?		\checkmark						
vii	Will the initiative result in you making decisions or taking action against individuals in ways which can have a significant impact on them?		~						
viii	Will the initiative compel individuals to provide information about themselves?	\checkmark			Provide information to input into Swiftqueue booking system				

If you answered **No** to <u>all</u> of the above screening questions, and you can evidence/justify your answers in the comments box above, you do not need to continue with the PIA.

Should the project at any point in the future use personal information you will need to revisit the screening questions and the PIA.

If you answered or **Unsure** to any of the above, please continue with the PIA.

¹ Intrusion can come in the form of collection of excessive personal information, disclosure of personal information without consent and misuse of such information. It can include the collection of information through the surveillance or monitoring of how people act in public or private spaces and through the monitoring of communications whether by post, phone or online and extends to monitoring the records of senders and recipients as well as the content of messages

Step 4 – Data Collection Please mark all information to be collected

	Please mark all informat	
Description	Specific data item (s)	Justification
		Reason that the data item(s) is/are needed
Personal Details	Name DOB NHS No. Mobile Number Gender	These specific details are required to ensure the patient is matched with the correct medical record, and that the COVID vaccination information transferred to the corresponding GP surgery.
Family, lifestyle		N/A
and social circumstances		
Education and		N/A
training details		
	Not applicable	N/A
Employment details	Not applicable	
Financial details	Not applicable	N/A
Sensitive Data: Racial or ethnic origin		N/A
Sensitive Data: Physical or mental health or condition	Not applicable	N/A
NB. Includes treatment if applicable. Include Mental		
Health status eg. whether detained or voluntary under the Mental Health Act if applicable.		
Sensitive Data: Sexual identity and life	Gender	For adverse effects monitoring
Sensitive Data: Religious or other beliefs of a similar nature	Not applicable	N/A
Sensitive Data: Trade union membership	Not applicable	N/A
Sensitive Data: Offences including alleged offences	Not applicable D	N/A

Description	Specific data item (s)	Justification Reason that the data item(s) is/are needed
Sensitive Data: Criminal proceedings, outcomes and sentences	Not applicable D	N/A

Step 3 – The Information Asset

How will the data be obtained and from where?	Information will be obtained either from our existing records or provided by the patient themselves via the			
	internet or telephone			
How will the data be used?	The data will be used to book an appropriate			
	appointment and to transfer vaccination information into our clinical system to match with the patient's			
	record			
Will the data be used locally or nationally?	Locally			
If National, list any available guidance				
Who will be the owner of the information?	Park Lane Surgery			
Ie. the Information Asset Owner (IAO)				
This is usually the Director or Service Lead under				
which this asset sits				
Who will be the Information Asset	Park Lane Surgery			
Administrator? (IAA)				
This is usually the Business Manager or person				
with day-to-day access and control				
Will a Third Party have access to the	No			
information?				
If so, name the third party, the circumstances and				
details of how the data will be accessed				
Will the data be shared with any other team or organisation?	Data will be shared between us and outcomes for health – the vaccination recording software –			
If so, name the organisation and the circumstances	consent for this is gained at the time of the			
If so, is there a data sharing agreement in place?	vaccination at their vaccination site.			
in so, is there a tract sharing agreement in place!				

Step 8 – Data Flows

Please provide a process map or diagram if available, or complete the table below

The answer to most the questions for the data flows are the same, as described below.

Name of Flow	What is the purpose of the data flow?	Will you be receivi ng data or sendin g it or both?	Where will you be receiving it from and/or sending it to?	Is the data anony mised ?	Is the data electronic or paper?	How is the data to be transferred? Eg. via a system, email, fax, post, by hand	How will the data be secured in transit? Eg. nhs.net to nhs.net	How often will data be transferred?	How many records in each transfer?	Where will the data be stored?	How will the data in storage be secured?
COVID Vaccine	Vaccination Status	Both	Received from pt > to Swiftqueue > to Outcomes for health > to Us	No	Eelectroni c	Via secure internet – System to system	Secure NHS web	Once per vaccination	Varies from 1- 1000 per day	Local secure medical system	Via smartcard + password protection

Step 9 – Data Protection Act Compliance

Name the data controller(s)	Park Lane Surgery
The data controller is the organisation which, alone or jointly or in common with other organisations, determines the purposes for which and the manner in which any personal data are, or are to be, processed.	
The data controller takes responsibility for complying with the UK GDPR.	
Name any data processors and provide contact details	Outcomes For Health
	Outcomes For Health
contact details A data processor means any organisation which processes the data on behalf of the	Outcomes For Health Consent

Data Protection Act Principles

Principle	Response	Actions required					
Principle 1: Personal data shall be processed lawfully, fairly and in a transparent manner.							
Individuals affected by the project must be informed about the processing of their data. Has a fair processing notice been provided or is a new or revised communication needed?	Consent gained from pt on booking						
What processes are in place to ensure that data required for secondary purposes is pseudonymised (or anonymised)?	Nil – personal data isn't used for secondary purposes						
If you are relying on consent to process personal data, how will consent be obtained and recorded, what information will be provided to support the consent process and what will you do if permission is withheld or given but later withdrawn?	Consent is gained either verbally or via SMS message and recorded in their medical record if processed by us.						
Principle2: Personal data sha purposes	all be collected for specified, e	explicit and legitimate					
What procedures are in place to ensure that privacy implications are considered prior to using data for a different purpose to that originally specified?	Any reports conducted within our clinical system do not use personal data						
Principle 3: Personal data shall be adequate, relevant and limited to what is necessary							
What procedures are in place for ensuring that data collection is adequate, relevant and not excessive in relation to the purpose for which data are being processed?	Collect only the information required by the external software						

Principle	Response	Actions required		
How will you ensure that the data you are using is likely to be of good enough quality for the purposes it is used for?	Compare information given to that recorded in our clinical system			
Principle 4: Personal data shall be accurate and where necessary kept up to date.				
What procedures are in place for ensuring that data collection is accurate?	Data given verbally should match with that in our clinical system			
What procedures are in place for ensuring that data collection is kept up to date?	Any contact with patients confirms their most up to date information			
What procedures are in place to correct inaccurate data when requested to do so by a data subject?	Staff are adequately trained to make any necessary amendments			
Principle 5: Personal data sh data subject for no longer the	all be kept in a form which pe an is necessary	rmits identification of the		
How long is the data to be retained for?	Unknown			
What procedures are in place for archiving / anonymisation / deletion / destruction of the data?	Use of PCSE / Outcomes for health Records retention policy			
Are there likely to be any exceptional circumstances for retaining certain data for longer than the normal period(s)?	Nil foreseeable			
What procedures are in place to provide data subjects access to their records?	NHS App / Systmonline			
What procedures are in place to prevent the processing of data which may cause damage or distress?	Redaction Software used upon request of records should it be necessary			
What procedures are in place for data subjects who may require the rectification, blocking, erasure or destruction of inaccurate data?	Staff adequately trained to action any inaccurate data			

Principle	Response	Actions required		
Principle 6: Appropriate technical & organisation measures shall be taken against unauthorised or unlawful processing of personal data & against accidental loss destruction or damage				
What procedures are in place to ensure that all staff who have access to the data undertake information governance training? What procedures are in place to ensure that data, whether	System warnings from our training portal to remind management and staff Managed by NHSE and our IT contractor NECS			
at rest or in transit, is secured? What procedures are in place to prevent the unauthorised disclosure of data to third parties?	End to end encryption Password and smartcard access only to the data			
Please ensure that the Checklist for Third Party Supplier of Services				

is completed where any new system is being introduced

Common Law Duty of Confidentiality

	Assessment of Compliance
Has the individual to whom the information relates given consent?	Y
Is the disclosure in the overriding public interest?	N/A
Is there a legal duty to do so, for example a court order	Ν
Is there a statutory basis that permits disclosure such as approval under Section 251 of the NHS Act 2006	Ν

Human Rights Act 1998

The Human Rights Act establishes the right to respect for private and family life. Current understanding is that compliance with the Data Protection Act and the common law of confidentiality should satisfy Human Rights requirements.

Will your actions interfere with the right to privacy under Article 8? – have you identified the social need and aims of the project? Are your actions a proportionate response to the social need?

N/A

Step 10 – Privacy issues identified and risk analysis

Any privacy issues which have been identified during the PIA process (for example: no legal basis for collecting and using the information; lack of security of the information in transit, etc.) should be documented in the risk register template embedded below. This risk register will enable you to analyse the risks in terms of impact and likelihood and document required action(s) and outcomes.

Note that where it is proposed that a privacy risk is to be 'accepted', approval for such acceptance should be sought from the Caldicott Guardian where patient data is concerned and the SIRO for all information risks.

The PMO holds the formal project risk register each IG lead should identify and records IG risks via the PMO.

Step 11 – Data Protection Principles Compliance and Authorisation

Please provide a summary of the conclusions that have been reached in relation to this project's overall compliance with the DPPs. This could include indicating whether some changes or refinements to the project might be warranted.

Information Asset Owner	Name: Louis Wood Date: 22/04/2021		
	Signature: Louis Wood		
Reasoning behind the decision to accept or reject the identified privacy risks			
Caldicott Guardian	Name: Lesley Hutchinson		
(only where the personal data are about patients)	Date: 22/04/2021		
	Signature:		
Reasoning behind the decision to accept or reject the identified privacy risks			
Senior Information Risk Owner	Name:		
(where the identified privacy risks are significant)	Date: Signature:		
	Signature.		
Reasoning behind the decision to accept or reject the identified privacy risks			
Information Governance Lead	Name: Lesley Hutchinson		
	Date: 22/04/2021		
	Signature:		
Reasoning behind the decision to accept or reject the identified privacy risks			

References

- · Data Protection Act 2018;
- · General Data Protection Regulations 2016
- · The Caldicott Principles;
- · Common Law Duty of Confidentiality;
- The Freedom of Information Act 2000;
- The Mental Capacity Act 2005;

• Section 251 of the NHS Act 2006 (originally enacted under Section 60 of the Health and Social Care Act 2001);

- Public Health (Control of Disease) Act 1984;
- · Public Health (Infectious Diseases) Regulations 1988;
- The Gender Recognition Act 2004;
- · Confidentiality: NHS Code of Practice 2003;
- · IGA Records Management Code of Practice for Health and Social Care 2016;
- · Reporting of Injuries, Diseases and Dangerous Occurrences Regulations 2013;
- Abortion Regulations 1991;
- · Road Traffic Act 1988;
- · ICO Data Sharing Code of Practice;
- · Confidentiality and Disclosure of Information Directions 2013;
- · Health and Social Care Act 2012;
- The Criminal Justice Act 2003;
- The NHS Information Security Management Code of Practice 2007;
- The Computer Misuse Act 1990;
- The Electronic Communications Act 2000;
- The Regulation of Investigatory Powers Act 2000;
- The Prevention of Terrorism Act 2005;
- The Copyright, Designs and Patents Act 1988;
- The Re-Use of Public Sector Information Regulations 2005;
- The Human Rights Act 1998;
- · The NHS Care Record Guarantee 2007; and
- · Anonymisation Standard for Publishing Health and Social Care Data Code of Confidentiality.