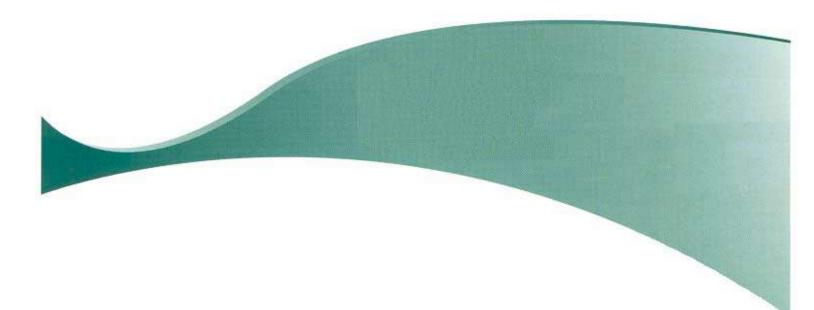


Privacy Impact Assessment (PIA) Form

Private & Confidential



This form should be completed with reference to the HSE Privacy Impact Assessment Process Guidance Document

> Version2.0 June 2019



Document Information

Title:	HSE Privacy Impact Assessment (PIA) Form
Purpose:	A PIA is a process to help identify and minimise the data privacy risks of a project or activity so as to ensure that patients and service users' rights to privacy and confidentiality are appropriately protected.
Author:	Joe Ryan
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Document History

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Privacy Impact Assessment Form

Section 1 – Initial Details (Threshold Assessment)		
Title of the activity:		
Name of person completing this form:		
Title:		
Service Area:		
Is personal data being collected or used?	Yes 🗌 No 🔲	
Are special categories of personal data being collected or used? (as listed below)	Yes 🗌 No 📋	
If yes, indicate the categories involved:	 Health data Data revealing racial or ethnic origin Political opinions Religious or philosophical beliefs Trade union membership Sex life data Genetic data Biometric data 	
If you answered 'No' to both of the ques remainder of the form as a PIA is not rec	tions above you do not need to complete the quired	
If you answered 'Yes' to any of the ques answer the questions below to establish	tions above you may need to complete a PIA - n if a PIA is required:	
Does the processing include the processing	g on a large scale of special category data?	
Yes 🗌 No 🗌		
Could the processing likely result in a high risk to the rights and freedoms of data subjects? Yes No		
Does the processing include a systematic r scale e.g. CCTV?	monitoring of a publicly accessible area on a large	
Yes 🗌 No 🗌		
Does the processing involve the automated processing, including profiling, on which decisions are based that produce legal effects concerning the data subjects? Yes No		
If you have answered 'Yes' to any of the complete the remainder of this form as a		
In order to complete this form please no completed the HSELanD GDPR/Data Pro that you have completed this training:	otection Awareness training. Please confirm	



Briefly outline the activity (name, purposes, c	ontext of use, etc.)
Describe how the activity generally works (fro stages, storage etc.) give a detailed descriptio	om data collection to data destruction, different processing on of each of the processes carried out.
What is the legal basis for processing the	Consent from the data subject.
data?	Processing is necessary for the performance of a contract.
	Processing is necessary for a legal obligation to which the HSE subject.
	Processing is necessary to protect the vital interests of the data subject.
	Processing is necessary for the performance of a task carried out in the public interest or in the exercise of official authority vested in the HSE.
f processing special categories of data what	Explicit Consent
If processing special categories of data what is the legal basis?	For the purposes of preventative or occupational
s the legal basis?	medicine, for the assessment of the working capacity of an
s the legal basis?	employee, for medical diagnosis, for the provision of medica care, treatment or social care, for the management of health or social care systems and services. Or pursuant to a contract with a health practitioner.



Name the Data Controller(s) involved in this activity:

Describe the Role of the data controller(s) for this activity:

Provide details of all data processors involved in this activity:

Describe the role of data processor(s) as relevant to this activity:

For each data processor, describe their responsibilities (duration, scope, purpose, documented processing instructions, prior authorisation, contracts in place) for this activity:

	e automated decision making? ibe the automated decision making	Yes 🗌 No 🗌
If 'Yes' what are the	consequences of the automated decision r	naking for the data subject:
	onal data collected is necessary for the pu	rpages of your processing
	onal data conected is necessary for the pu	rposes of your processing.
List the data suppor	ting assets (hardware, software, networks,	people, paper or paper transn
channels):		

	being sourced from				me from e mubli
accessible	se state where the d source:	ata originates fi	om and if appi	icable, did it co	ome from a public
What is the	retention period for	the different it	ame of poreone	l data:	
Describe th	e steps taken to ens	sure that the pe	rsonal data is l	ept up to date	and accurate:
Describe th	ne steps taken to ens	sure that the pe	rsonal data is I	ept up to date	and accurate:
Describe th	e steps taken to ens	sure that the pe	rsonal data is I	ept up to date	and accurate:
Describe th	e steps taken to ens	sure that the pe	rsonal data is I	ept up to date	and accurate:
Describe th	e steps taken to ens	sure that the pe	rsonal data is I	ept up to date	and accurate:
	ne steps taken to ens			ept up to date	and accurate:



How can data subjects exercise their right to access and to data portability under Article 15 and Article 20 of the GDPR?

How can data subjects exercise their right to rectification and erasure under Articles 16 & 17 of the GDPR?

How can data subjects exercise their right to restriction and object under Article 18 and Article 21 of the GDPR?

Is the personal data being transferred outside of the Republic of Ireland? Yes 🗌 No 🗍

If yes, list the countries where the personal data is to be transferred:

For each country outside of the EEA (European Economic Area) where data is stored or processed, name it and describe the provisions concerning the transfer:

Describe the organisational security measures associated with this activity:

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	-	arondant	Service

Describe the technical security measures associated with this activity:

Describe the additional measures taken to ensure data security for this activity:

Section 3 – Research

Please complete the following section only if you are completing this PIA as part of a research proposal.

If you are not completing this PIA as part of a research proposal you can go immediately to Section 4.

Please specify what arrangements are in place to ensure that personal data will be processed as is necessary;

(a) to achieve the objective of the health research and;

(b) to ensure that data shall not be processed in such a way as to damage or distress the data subject:

The provision of training in data protection law and practice to anyone involved in carrying out the health research is a mandatory legal requirement. Please specify the data protection training undertaken by those involved in this research:



Please specify the controls in place to log whether and by whom personal data has been consulted, altered, disclosed or erased:

Please specify the arrangements to anonymise, archive or destroy personal data once the health research has been completed and how this will be carried out:

Please specify other technical and organisational measures designed to ensure that processing is carried out in accordance with the Data Protection Regulation, together with process for testing and evaluating the effectiveness of such measures:



Section 4 – Risks and Ri	isk Mitigation
Is there a risk of:	 a. Illegitimate access to personal data b. Unwanted modification to personal data c. Personal data disappearance d. Other (please state)

Section 4 (a) – Illegitimate access to personal data Complete the following questions if you selected a. (Illegitimate access to personal data) What are the main threats that could lead to the risk? What are the potential impacts on data subjects arising from the risk? What are the risk sources? What controls are in place to address the risk and are these controls adequate?

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Níos Fearr	Better Health
á Forbairt	Service

How do you estimate the likelihood of the risk, especially in respect of threats, sources of risk and planned controls?	 1 – Rare 2 – Unlikely 3 – Possible 4 – Likely 5 – Highly Certain
How do you estimate the potential impact of the risk on data subjects?	 1 – Negligible 2 – Minor 3 – Moderate 4 – Major 5 – Critical
What is the overall risk rating (likelihood x impact)?	 Low Medium High

Section 4 (b) – Unwanted modification to personal data <u>Complete the following questions if you selected b. (Unwanted modification to personal data)</u> What are the main threats that could lead to the risk?

What are the potential impacts on data subjects arising from the risk?

What are the risk sources?

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~	Níos Fearr	Better Health
~	á Forbairt	Service

What controls are in place to address the risk and are these c	ontrols adequate?
How do you estimate the likelihood of the risk, especially in respect of threats, sources of risk and planned controls?	 1 – Rare 2 – Unlikely 3 – Possible 4 – Likely 5 – Highly Certain
How do you estimate the potential impact of the risk on data subjects?	 1 – Negligible 2 – Minor 3 – Moderate 4 – Major 5 – Critical
What is the overall risk rating (likelihood x impact)?	 Low Moderate High

Section 4 (c) – Personal data disappearance

Complete the following questions if you selected c. (Personal data disappearance)

What are the main threats that could lead to the risk?

What are the potential impacts on data subjects arising from the risk?

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What are t	he risk sourc	es?		
What cont	rols are in pla	ace to address the ris	sk and are these c	controls adequate?
How do y	u octimata th	a likelihaad of the riv	sk osposially in	🗌 1 – Rare
respect of	threats, sour	ne likelihood of the ris rces of risk and plann	ned controls?	2 – Unlikely
				\square 3 – Possible
				 □ 4 – Likely
				5 – Highly Certain
How do yo subjects?	ou estimate th	ne potential impact of	the risk on data	 1 – Negligible 2 – Minor
				3 - Moderate
				\square 4 – Major
				☐ 5 – Critical
What is th	e overall risk	rating (likelihood x in	mpact)?	Low
				Moderate



Section 4 (d) – Other

Complete the following questions if you selected d. (Other)

Describe in detail the risk

What are the main threats that could lead to the risk?

What are the potential impacts on data subjects arising from the risk?

What are the risk sources?

What controls are in place to address the risk and are these controls adequate?

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	e likelihood of the risk, especially burces of risk and planned	 1 – Rare 2 – Unlikely 3 – Possible 4 – Likely 5 – Highly Certain
How do you estimate th data subjects?	e potential impact of the risk on	 1 – Negligible 2 – Minor 3 – Moderate 4 – Major 5 – Critical
What is the overall risk	rating (likelihood x impact)?	 Low Medium High

Section 5 – Data Subject Consultation			
Were data subjects (or a representative) consulted as a part of the PIA process? Yes No			
If Yes, state the number of data subjects consulted, method of consultation and describe the outcome of the consultation:			
If No, explain the reasons for not consulting data subjects:			

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Section 6 – DPO/DDPO Consultation

DPO opinion (please ensure the previous questions are completed fully before the DPO can provide an opinion):

Section 7 – Approval To be completed by the data controller				
Outcome:				
	DPC Consultation Needed			
	Further Updates Needed			
Signed:	Date:			