

# **Data Protection Impact Assessment (DPIA) Template**

A DPIA is designed to describe your processing and to help manage any potential harm to individuals in the use of their information. DPIAs are also important tools for demonstrating accountability, as they help you as a Controller to comply with the requirements of the Data Protection Legislation. Non-compliance with DPIA requirements can lead to fines imposed by the Information Commissioners Office (ICO); this includes not carrying out a DPIA at all, carrying out a DPIA in an incorrect way or failing to consult the ICO where required.

DPIA's are not new; the use of Privacy Impact Assessments has become common practice in the NHS and can provide evidence of compliance within the Data Security and Protection toolkit (DSPT); DPIAs build on that practice.

It is not always clear whether you should do a DPIA or not but there are a number of situations where a DPIA **should** be considered or where a DPIA is a **legal requirement**. If you can tick against the criteria below it is highly recommended that you undertake a DPIA and if you decide not to, ensure that you document the reasons for your decision.

You as Controller MUST carry out a DPIA where you plan to:	Tick or
	leave
	blank
Use <b>profiling or automated decision-making</b> to make significant decisions about people or their access to a	📙
service, opportunity or benefit;	
Process special-category data or criminal-offence data on a large scale;	
Monitor a publicly accessible place on a large scale;	
Use innovative technology in combination with any of the criteria in the European guidelines;	
Carry out <b>profiling</b> on a large scale;	
Process biometric or genetic data in combination with any of the criteria in the European guidelines;	
Combine, compare or match data from multiple sources;	
Process personal data without providing a privacy notice directly to the individual in combination with any of the	
criteria in the European guidelines;	
Process personal data in a way that involves tracking individuals' online or offline location or behaviour, in	
combination with any of the criteria in the European guidelines;	
Process children's personal data for profiling or automated decision-making or for marketing purposes, or offer	
online services directly to them;	
Process personal data that could result in a risk of physical harm in the event of a security breach.	
You as Controller should <b>consider</b> carrying out a DPIA where you	Tick or
, ,	leave
	blank
Plan any major project involving the use of personal data;	Ш
Plan to do evaluation or scoring;	
Want to use systematic monitoring;	✓
Process sensitive data or data of a highly personal nature;	
Processing data on a large scale;	
Include data concerning vulnerable data subjects;	
Plan to use innovative technological or organisational solutions;	

A new DPIA should be carried out if you decide that there is a significant enough change to what you originally intended but it is good practice for DPIAs to be kept under review and revisited when necessary.

There is guidance to help you. Your Data Protection Officer (DPO) can be consulted before completing a DPIA in order to provide specialist advice and guidance or simply to talk things through with you.



Background Information			
Date of your DPIA:	09/03/2023		
Title of the activity/processing:	Data feed into regional Demand and Capacity database		
Who is the person leading this work?	, BOB ICB		
Who is the Lead Organisation?	BOB ICB		
Who has prepared this DPIA?	, BOB ICB		
Who is your Data Protection Officer (DPO)?			
Describe what you are proposing to do: (Include as much background information as you can about why the new system/change in system/sharing of information/data processing is required).	To implement an ICB wide data system to monitor real time (or as close to real time as possible) data from all general practices to display the real time demand and capacity situation, enabling operational management decision making. The regional Demand and Capacity dashboard, developed by suppliers Edenbridge, will display aggregated data from each general practice in BOB ICB (with the exception of practices who choose to opt-out), in one dashboard. The Demand and Capacity dashboard will not collect any PID data or data that can be used to identify any individuals and will not collect any sensitive data. The data collected is related to appointment activity. For those practices in BOB who are using the TVS shared care record, their data will be extracted from the software used for this platform (Graphnet). Manual submissions to the regional dashboard will be made until such a time that a practice has adopted TVS.		
Are there multiple organisations involved? (If yes – you can use this space to name them, and who their key contact for this work is).	Yes, all general practices across BOB (excluding those who opt out). The key contact for coordinating their involvement is Sanjay Desai, Head of Operations for Primary Care at BOB ICB		
Can you think of any other Key Stakeholders that should be consulted or involved in this DPIA? (If so then include the details here).	The GP DPO for BOB ICB has been consulted		
Detail anything similar that has been undertaken before?	Multiple deployments across the NHS. The majority of ICBs submitting to the Edenbridge regional dashboard are using Apex, an insights and analytics platform developed by Edenbridge. All deployments use aggregated and/or summarised data and no PID or sensitive data has ever been collected. All deployments have concluded that there is no risk to patient data and are fully compliant with all UK Data Protection and associated laws and guidelines. It has been commissioned by NHS England.		

1. Categories, Legal Basis, Responsibility, Processing, Confidentiality, Purpose, Collection and Use				
1.1.				
What data/information will be used?	Tick or leave	Complete		
Tick all that apply.	blank			
Personal Data		1.2		
Special Categories of Personal Data		1.2 AND 1.3		
Personal Confidential Data		1.2 AND 1.3 AND 1.6		



Sensitive Data (usually criminal or law enforcement data )		1.2 but speak to your IG adviso	r first	
Pseudonymised Data		1.2 and consider at what point	the data	
		is to be pseudonymised		
Anonymised Data	✓	Consider at what point the data	a is to be	
Commentally Confidentially Constitution		anonymised		
Commercially Confidential Information		Consider if a DPIA is appropriat		
Other	Ш	Consider if a DPIA is appropriat	:e	
1.2.  Processing has to be lawful so identify which of the following do and include an explanation as to why in the relevant box				
Article 6 (1) of the GDPR includes the following:				
a) THE DATA SUBJECT HAS GIVEN CONSENT			Tick or leave blank	
Why are you relying on consent from the data subject? Click here to enter text.				
What is the process for obtaining and recording consent from	m the	Data Subject? (How, where, when, by v	vhom).	
Click here to enter text.				
Describe how your consent form is compliant with the Data	Prote	ction requirements? (There is a chec	klist that	
can be used to assess this). Click here to enter text.				
b) IT IS NECESSARY FOR THE PERFORMANCE OF A CONTR	ACT TO	WHICH THE DATA SUBJECT IS	Tick or leave blank	
(The contract needs to be between the Controller and the individual and not conce a contract with the Controller. Processing can happen before the contract is enteroprivate or cosmetic procedure that is a paid for service with the delivery of that can Practitioner).	ed into e.g	processing a pre-health assessment for a		
What contract is being referred to?				
Click here to enter text.				
c) IT IS NECESSARY UNDER A LEGAL OBLIGATION TO WHI	CH THE	CONTROLLER IS SUBJECT	Tick or leave	
(A legal obligation mandates processing of data as a task in itself where there are li			blank	
e.g. an Employer has a legal obligation to disclose salary information to HMRC).				
Identify the legislation or legal obligation you believe required click here to enter text.	res you	to undertake this processing.		
			Tick or	
d) IT IS NECESSARY TO PROTECT THE VITAL INTERESTS OF NATURAL PERSON	THE DA	ATA SUBJECT OR ANOTHER	leave blank	
(This will apply only when you need to process data to protect someone's life. It m individual whose data is being processed. It can also apply to protect another pers category but planned care would not. You may need to process a Parent's data to is unlikely to be able to provide consent physically or legally; if you are able to gain	on's life. E protect th	Emergency Care is likely to fall into this e life of a child. The individual concerned		
How will you protect the vital interests of the data subject			ng this	
activity? Click here to enter text.				
e) IT IS NECESSARY FOR THE PERFORMANCE OF A TASK CONTURNED OF CONT			Tick or leave blank	
(This is different to 6 c). If you are processing data using this basis for its lawfulnes function or power that is set out in law. The processing must be necessary, if not ti	•	· · ·	<b>√</b>	
What statutory power or duty does the Controller derive th	eir offi	cial authority from?		
NHS Act 2006, Health and Social Care Act 2012				



_				
f)	IT IS NECESSARY FOR THE LEGITIMATE INTERESTS OF THE CONTROLLI	R OR THIRD PARTY	Tick or leave	
•	c authorities can only rely on legitimate interests if they are processing for a legitimate reason oth ublic authority. See the guidance for more information about the legitimate interest test).	ner than performing their tasks	blank	
	at are the legitimate interests you have?			
	k here to enter text.			
1.3.				
If usi	ng special categories of personal data, a condition for processing und	der Article 9 of the GD	PR must be	
satisf	ied in addition to a condition under Article 6. You must select at leas	st 1 from a) to c) or g)	to i). NOTE:	
d), e)	and f) are not applicable			
Arti	cle 9 (2) conditions are as follows:			
a)	THE DATA SUBJECT HAS GIVEN EXPLICIT CONSENT		Tick or leave	
(Requ	irements for consent are the same as those detailed above in section 1.2, a))		blank	
b)	FOR THE PURPOSES OF EMPLOYMENT, SOCIAL SECURITY OR SOCIAL	PROTECTION	Tick or leave	
(Sche	dule 1, part 1 of the Data Protection Act 2018 gives more detail on when this can apply to proce	ssing and further guidance is	blank	
availa		g		
c)	IT IS NECESSARY TO PROTECT THE VITAL INTERESTS OF THE DATA SU		Tick or leave blank	
	NATURAL PERSON WHERE THEY ARE PHYSICALLY OR LEGALLY INC	CAPABLE OF GIVING		
	CONSENT			
	irements for this are the same as those detailed above in section 1.2, d))		2/4	
d)	It is necessary for the operations of a not-for-profit organisation such a	•	NA	
۵)	philosophical, trade union and religious body in relation to its member	S	NA NA	
<u>e)</u> f)	The data has been made public by the data subject  For legal claims or courts operating in their judicial category		NA NA	
g) SUBSTANTIAL PUBLIC INTEREST				
			blank	
(Sche	dule 1, part 2 of the Data Protection Act 2018 gives more detail on when this can apply to process ble).	ing and further guidance is		
h)	PROCESSING IS NECESSARY FOR THE PURPOSES OF PREVENTIVE OR OCCU		Tick or leave blank	
	FOR THE ASSESSMENT OF THE WORKING CAPACITY OF THE EMPLOYEE, MEI PROVISION OF HEALTH OR SOCIAL CARE OR TREATMENT OR THE MANAGI	-		
	SOCIAL CARE SYSTEMS AND SERVICES ON THE BASIS OF UNION OR ME			
	PURSUANT TO CONTRACT WITH A HEALTH PROFESSIONAL AND SUBJECT	TO CONDITIONS AND		
	SAFEGUARDS			
(Scheavaila	dule 1, part 1 of the Data Protection Act 2018 gives more detail on when this can apply to procest ble).	ssing and further guidance is		
i)	PROCESSING IS NECESSARY FOR REASONS OF PUBLIC INTEREST IN THE ARI	-	Tick or leave blank	
	SUCH AS PROTECTING AGAINST SERIOUS CROSS-BORDER THREATS TO HEAL STANDARDS OF QUALITY AND SAFETY OF HEALTH CARE AND OF MEDI			
	MEDICAL DEVICES, ON THE BASIS OF UNION OR MEMBER STATE LAW V			
	SUITABLE AND SPECIFIC MEASURES TO SAFEGUARD THE RIGHTS AND FRE			
	SUBJECT, IN PARTICULAR PROFESSIONAL SECRECY			
(Scheavaila	dule 1, part 1 of the Data Protection Act 2018 gives more detail on when this can apply to procestle).	ssing and further guidance is		
1.4.				
Confi	rm who the Controller/s is/are and whether solely or jointly responsi	ble for any data proce	ssed?	
	y any other parties who will be included in the agreements and who will have involvement/share project/activity. Use this space to detail this but you may need to ask your DPO to assist you. Co			
	project/activity. Use this space to detail this but you may need to ask your DPO to assist you. Co ganisations where required (the text has been left unlocked for this purpose on that row only).	py and paste the last empty ro	ww iii tile table to	
Nan	ne of Organisation	Role		
		Sole Controller		



Click here to enter text.	Choose an item.
Click here to enter text.	Choose an item.
Click here to enter text.	Choose an item.
Click here to enter text.	Choose an item.
Click here to enter text.	Choose an item.

#### 1.5.

# Describe exactly what is being processed, why you want to process it and who will do any of the processing?



NHSE%20South%20

Appointment data as detailed in the following schema: East%20-%20Regiona (see attached as an appendix to the document) as recorded by clinical and non-clinical staff (see Intro tab of document) which is anonymous and aggregated at ICB level, not PCN or Practice level. There is no PID data or data that can be used to identify any individuals and sensitive data is not being collected.

The data collected is for operational review purposes to understand pressures in general practice and has been requested by the NHS England South East regional team. More information can be found here:

# Demand & Capacity tooling in the Region

Why are we implementing T&C tooling?



#### Why we're implementing D&C tooling

Acknowledging the sustained pressures that Primary Care are under, and the current reporting source (GPAD) underrepresenting the true activity of general practice, regional investment in 2022 has enabled the development and implementation of bespoke Demand & Capacity tooling, to:

- Principally to help practices make operational decisions to deliver the best possible service to their patients, and use their finite resources in the most
- efficient ways possible.

  To enable the provision of support / mutual aid / funding from the ICB.

  Enable improved reporting of general practice activity (GP appointment data).

  Support operational decision making.

#### Furthermore, several drivers sit behind the delivery of these bespoke D&C tools

- of these bespoke D&C tools:

  Increased focus from Ministers and national priorities in providing patients with data

  Publication of the GPAD data set D&C tooling, which more accurately captures and reports on GP appointment data, will be a vital mitigation to inaccurate narratives derived from GPAD.

  Supporting practices to in rebutting negative publicity around pressures.

#### What a system level data feed will be for?

With these drivers in mind, ICBs have agreed with the region the need for system level data within an aggregated regional dashboard, to bring about a set of benefits in the interest of Primary Care.

As a Practice, what would the regional dashboard

Building on the progress that has been made by Practices, PCNs and ICBs, the System level data will be used for:

- Provide a more accurate, reliable and timely articulation of demand pressures for the region as a
- wnoie.
  Ensure a consistent and comparable (interoperable)
  GP appointment data set across the region.
  Enable the provision of support, as such mutual aid
  and the direction of funding based on need, within
  and across geographic boundaries.
- and across geographic ocunianes.

  By providing aggregated System level data into the regional dashboard, ICSs and Region will be able to more accurately respond and/or influence future or
- more accurately respond and a secondary potential policy.

  Ensure that General Practice data is as timely and robust as Secondary Care data to make sure system-wide conversations on demand are as well-informed as possible

#### What System level data won't be for?

Importantly, we want to be clear on what System level data will not be used for:

- The system level data will not be used for performance management purposes. The region has not asked for and will therefore be unable to access data on individual practices or PCNs.
- Its not the remit or responsibility of NHS England

CSs and regions have agreed that aggregated data will feed into the overarching regional demand

This system level data feed would not identify individual practices, and would be an aggregated totality of all practices' data within the System:

Frimley ICB are undertaking the development of data feeds into the Edenbridge regional dashboard as part of work commissioned by BOB ICB. Practices who have not opted out of the initiative and are part of the TVS shared care record dashboard will have the relevant data extracted directly from Graphnet for submission to the regional dashboard. BOB ICB will establish contingencies for those practices not yet using TVS to determine a manual submission process. Data will be aggregated with no data identifying the practice submitted to Edenbridge.

#### 1.6.



Tick here if you owe a duty of confidentiality to any information.

If so, specify what types of information. (e.g. clinical records, occupational health details, payroll information)

No personally identifiable data is being processed within this project

1.7.

How are you satisfying the common law duty of confidentiality?

No disclosure due to anon/pseudo actions

If you have selected an option which asks for further information please enter it here

No further actions as no PID data or sensitive data is being received

1.8.

Are you applying any anonymisation/pseudonymisation technique or encryption to any of the data to preserve the confidentiality of any information?

No

If you are then describe what you are doing.

Not applicable as no PID/Sensitive data is being received

If you don't know then please find this information out as there are potential privacy implications with the processing.

1.9.

Tick here if you are intending to use any information for a purpose that isn't considered as direct patient care.  $\checkmark$ 

# If so describe that purpose.

All of the data displayed on Demand and Capacity regional dashboard is with regards to pressure and flow and therefore all data in the dashboard would be considered as not directly for patient care. The real time pressures give providers and commissioners an overview of the situation rather than the granular level detail that PID contains.

1.10.

Approximately how many people will be the subject of the processing?

CCG population

1.11.

**How are you collecting the data?** (e.g. verbal, electronic, paper (if you need to add more selections then copy the last 'choose an item' and paste, the text has been left unlocked for you to do this.)

Other method not listed

Electronic form

By e-mail

Choose an item.

Choose an item.

If you have selected 'other method not listed' describe what that method is.

Submission into the Edenbridge data warehouse

1.12.

How will you edit the data?

No data editing occurs

1.13.

#### How will you quality check the data?

We have a live dashboard. Therefore, data quality checks are carried out by technical members of staff reviewing data with a technical and operational member of staff at each provider to review the data being provided to ensure that it is an accurate reflection of the real situation.

1.14.



# Review your business continuity or contingency plans to include this activity. Have you identified any risks?

No

If yes include in the risk section of this template.

1.15

#### What training is planned to support this activity?

Training will be developed where required to support those practices who are not subject to the data extraction from Graphnet to retrieve and submit the data manually.

## 2. Linkage, Data flows, Sharing and Data Opt Out, Sharing Agreements, Reports, NHS Digital

#### 2.1.

Are you proposing to combine any data sets?

No

If yes then provide the details here.

Click here to enter text.

2.2.

What are the Data Flows? (Detail and/or attach a diagram if you have one).

Where practices are already feeding into the Thames Valley and Surrey (TVS) shared care record, extracts of pre-mapped appointment data in accordance with the requirements from NHS England NHSE South East - Regional Dashboard Schema v1-0 (1).xlsx are automatically collated and submitted into the regional dashboard.

Practices who are not currently live on TVS have the option to generate a manual submission in conjunction with the ICB where the submission is extracted directly from a GP IT clinical foundation system in the form of a .cvs report, then sent to Edenbridge for inclusion in the BOB ICB dataset.

## 2.3.

#### What data/information are you planning to share?

Summarised and/or aggregated incident and performance data

2.4.

#### Is any of the data subject to the National Data Opt Out?

No - it is not subject to the national data opt out

If your organisation has to apply it describe the agreed approach to this

Click here to enter text.

#### If another organisation has applied it add their details and identify what data it has been applied to

Data sent to the regional demand and capacity dashboard is anonymous at the point of sending, directly from GP IT foundation systems via Graphnet or through manual submission. No PID data is ever sent to the regional dashboard and therefore no anonymisation processes occur within the demand and capacity dashboard system.

If you do not know if it applies to any of the data involved then you need to speak to your Data Protection Officer to ensure this is assessed.

2.5.

#### Who are you planning to share the data/information with?

The data is shared with NHS England regional and national colleagues who have access to the regional demand and capacity dashboard. However, this is aggregate data and no PID is shared.

2.6.

Why is this data/information being shared?



To provide an overview of the real time pressures across general practices in BOB ICB to give better and up to date information so that providers and commissioners can make better and quicker decisions based on patient flow and real time pressures and capacity

#### 2.7.

How will you share it? (Consider and detail all means of sharing)

Aggregate data will be shared via the regional demand and capacity dashboard only.

Tick if you are planning to use Microsoft Teams/Skype or another similar online networking/meeting solution that may have the facility to store or record conversations or related data as part of the sharing arrangements ✓

#### Provide details of how you have considered any privacy risks of using one of these solutions

As part of the N365 tenant, BOB ICB is subject to the information governance and data protection protocols in place as part of its implementation. Meetings are only recorded where necessary. As no PID or sensitive data is being shared, the topics of discussion in meetings are strictly limited to the appointment activity data required for the regional demand and capacity dashboard.

#### 2.8.

#### What data sharing agreements are or will be in place?

Due to the nature of the data (non PID), BOB ICB information governance leads have determined that a data sharing agreement is not required for this project.

#### 2.9.

#### What reports will be generated from this data/information?

None directly but aggregate data on the dashboard.

#### 2.10.

Are you proposing to use Data that may have come from NHS Digital (e.g. SUS data, HES data etc.)? No

#### If yes, are all the right agreements in place?

Choose an item.

#### Give details of the agreement that you believe covers the use of the NHSD data

Click here to enter text.

If no or don't know then you need to speak to your Data Protection Officer to ensure they are put in place if needed.

#### 3. Data Processor, IG Assurances, Storage, Access, Cloud, Security, Non-UK processing, DPA

#### 3.1

Are you proposing to use a third party, a data processor or a commercial system supplier? Yes

If yes use these spaces to add their details including their official name and address. If there is more than one then include all organisations. If you don't know then stop and try and find this information before proceeding.

Edenbridge Healthcare are incorporated and registered in England and Wales as Egton Medical Information Systems Ltd with company number 02117205 whose registered office is at Fulford Grange Micklefield Lane, Rawdon, Leeds, England LS19 6BA

Click here to enter text.

3.2



**Is each organisation registered with the Information Commissioner?** Copy and paste the last empty row in the table to add organisations where required (the text has been left unlocked for this purpose on that row only)

Name from 3.1 above	Registered	Registration details or comments if not registered	
Egton Medical Information Ltd	Yes	Data Protection Registration Number:Z5514037	
NHS Buckinghamshire Oxfordshire and Berkshire West Integrated Care Board	Yes	Data Protection Registration Number: ZB343068	
Click here to enter text.	Choose an item.	Click here to enter text.	
Click here to enter text.	Choose an item.	Click here to enter text.	
Click here to enter text.	Choose an item.	Click here to enter text.	
Click here to enter text.	Choose an item.	Click here to enter text.	

#### 3.3

# What IG assurances have been provided to you and does the contract contain IG clauses that protect you as the Controller? (e.g. in terms and conditions their contract, their tender submission). Convend nate the last empty row in the table to add

**the Controller?** (e.g. in terms and conditions, their contract, their tender submission). Copy and paste the last empty row in the table to add organisations where required (the text has been left unlocked for this purpose on that row only)

Name from 3.1 above	Brief description of assurances obtained
Egton Medical Information Ltd	Contract. ICO registered and DSP Toolkit compliant.
Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.

#### 3.4

#### What is the status of their Data Security Protection Toolkit?

#### **DSP Toolkit**

copy and paste the last empty row in the table to add organisations where required (the text has been left unlocked for this purpose on that row only)

Name from 3.1 above	ODS Code	Status	Published date
Egton Medical Information Ltd	YGM06	22/23 Standards Exceeded	12/06/2023
BOB ICB	QU9	Standards Exceeded	27/06/2023
Click here to enter text.	Click here to enter text.	Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.	Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.	Click here to enter text.	Click here to enter text.
Click here to enter text.	Click here to enter text.	Click here to enter text.	Click here to enter text.

#### 3.5

**How and where will the data/information be stored?** (Consider your answer to 2.7 and the potential storage of data in any online meeting or networking solution).

Hosted by AWS UK (London) instance. Fully compliant with all data storing and sharing laws regarding UK NHS data. This includes support and technical staff that may need to access the system.

#### 3 6

#### How is the data/information accessed and how will this be controlled?

The regional demand and capacity dashboard is accessible through a login to the online tool. Logins are controlled by the NHS England south east regional and national teams.



#### 3.7

Is there any use of Cloud technology?

Yes

If yes add the details here.

Please see details in section 3.5 above.

#### 3.8

#### What security measures will be in place to protect the data/information?

Please see section 3.5 and section 3.7 for details. Also, the Demand and Capacity dashboard will not collect any PID data or data that can be used to identify any individuals and will not collect any sensitive data. The data collected is related to appointment activity. For those practices in BOB who are using the TVS shared care record, their data will be extracted from the software used for this platform (Graphnet). Manual submissions to the regional dashboard will be made until such a time that a practice has adopted TVS.

#### Is a specific System Level Security Policy needed?

Choose an item.

If yes or don't know then you need to speak to your Data Protection Officer to ensure one is put in place if needed.

3.9

Is any data transferring outside of the UK? (you must determine this so only select don't know if you have further investigations to make but the DPIA will not be approved without this information)

If yes describe where and what additional measures are or will be in place to protect the data.

Click here to enter text.

#### 3.10

# What Data Processing Agreement is already in place or if none, what agreement will be in place with the organisation and who will be responsible for managing it?

No process in place as no PID data ever received. The technical set up of the system means that Edenbridge are unable to receive PID data as we only accept numerical figures in the form of pure numbers or minutes which are already in non-PID form as described in previous segments.

#### 4. Privacy Notice, Individual Rights, Records Management, Direct Marketing

#### 4.1

Describe any changes you plan or need to make to your Privacy Notice and your proposed completion date? (There is a checklist that can be used to assess the potential changes required or if you wish for it to be reviewed then add the link below). None required as no impact or data to be retained.

#### 4.2

**How will this activity impact on individual rights under the GDPR?** (Consider the right of access, erasure, portability, restriction, profiling, automated decision making).

There will be no impact on individual rights. Information is made available in anonymised form. In this case, the data is always anonymous aggregated counts of information. NHS National data opt-outs are not applied where information is made available in anonymised form. In this case, the data is always anonymous aggregated counts of information.

#### 4.3

# How long is the data/information to be retained?

No PID data but any summary data can be kept for up to 20 years, or to the end of contract.

#### 4.4

## How will the data/information be archived?

No current archive approach

#### 4.5

#### What is the process for the destruction of records?

Deletion of database and decommission of services in AWS



4.6

### What will happen to the data/information if any part of your activity ends?

Database and any backups to be deleted at the end of any contract

4.7

Will you use any data for direct marketing purposes? (you must determine this so only select don't know if you have further investigations to make but the DPIA will not be approved without this information)

No

If yes please detail.

Click here to enter text.

#### 5. Risks and Issues

5.1

What risks and issues have you identified? The DPO can provide advice to help complete this section and consider any measures to mitigate potential risks.

Describe the source of risk and nature of potential impact on individuals.  (Include associated compliance and corporate risks as necessary and copy and paste the complete bottom row to add more risks (the text has been left unlocked in both tables to enable you to do this)).	Likelihood of harm	Severity of harm	Overall risk
Access to data by unauthorised individuals	Remote	Minimal	Low
Data lost or misappropriated	Possible	Minimal	Low
Processing of incorrect data	Possible	Minimal	Low
System Failure	Possible	Minimal	Medium

5 2

Identify additional measures you could take to reduce or eliminate risks identified as medium or high risk in 5.1

Risk	Options to reduce or eliminate risk	Effect on risk	Residual risk	Measure approved (SIRO)
System Failure		Tolerated	Medium	Choose an
				item.
Click here to enter	Click here to enter text.	Choose an item.	Choose an	Choose an
text.			item.	item.
Click here to enter	Click here to enter text.	Choose an item.	Choose an	Choose an
text.			item.	item.
Click here to enter	Click here to enter text.	Choose an item.	Choose an	Choose an
text.			item.	item.

5.3

What if anything would affect this piece of work?

Nothing of note

5.4

Please include any additional comments that do not fit elsewhere in the DPIA?

Nothing of note

## 6. Consultation

6 1

Have you consulted with any external organisation about this DPIA?

No

If yes, who and what was the outcome? If no, detail why consultation was not felt necessary.



Click here to enter text.

6.2

Will you need to discuss the DPIA or the processing with the Information Commissioners Office? (You may need the help of your DPO with this)

Nο

If yes, explain why you have come to this conclusion.

There is no compelling reason to do this

#### 7. Data Protection Officer Comments and Observations

7.1

**Comments/observations/specific issues** 

#### 8. Review and Outcome

Based on the information contained in this DPIA along with any supporting documents, you have determined that the outcome is as follows:

A) There are no further actions needed and we can proceed

If you have selected item B), C) or D) then please add comments as to why you made that selection Click here to enter text.

#### We believe there are

Choose an item.

If you have selected item B) or C) then list these in the amber boxes below and then consider additional measures you could take and include these in the green boxes below

Residual risks and nature of potential	Likelihood of harm	Severity of harm	Overall risk
impact on individuals. (Include associated compliance and corporate risks as necessary and copy and paste the complete bottom row to add more risks (the text has been left unlocked in both tables to enable you to do this)).			
Click here to enter text.	Choose an item.	Choose an item.	Choose an item.
Click here to enter text.	Choose an item.	Choose an item.	Choose an item.
Click here to enter text.	Choose an item.	Choose an item.	Choose an item.
Click here to enter text.	Choose an item.	Choose an item.	Choose an item.

Additional measures you could take to reduce or eliminate residual risks identified as medium or high ris above (B and C)					
Risk	Options to reduce or eliminate risk	Effect on risk	Residual risk	Measure approved (SIRO)	
Click here to enter text.	Click here to enter text.	Choose an item.	Choose an item.	Choose an item.	
Click here to enter text.	Click here to enter text.	Choose an item.	Choose an item.	Choose an item.	
Click here to enter text.	Click here to enter text.	Choose an item.	Choose an item.	Choose an item.	
Click here to enter text.	Click here to enter text.	Choose an item.	Choose an item.	Choose an item.	



Signed and approved on behalf of Buckinghamshire Oxfordshire and Berkshire West Integrated Care Board

Date: 23/05/2024

Name:

Job Title: Data Protection Officer

Signature:

#### Please note:

You should ensure that your Information Asset Register and Data Flow Mapping Schedules are updated where this is relevant.

This DPIA can be disclosed if requested under the Freedom of Information Act (2000). If there are any exemptions that should be considered to prevent disclosure detail them here:

Click here to enter text.

# Schema for NHSE South East - Regional Dashboard

V1.0

Commercial in Confidence

# Introduction

This document lists the appointment data items required to populate the NHSE South East Regional Appointment Dashboard.

The data is structured into a single file/table, exported as CSV on a daily basis, for each of the following clinician groups.

- 1. All Clinician Groups
- 2. GP
- 3. Nurse
- 4. Other Healthcare Professional
- 5. Admin/Clinical Support

# **Change Log**

Date	Description	Version
07-Feb-24	Initial Version	1.0

Column Name	Description
Date	Date in format yyyy-mm-dd (e.g. 2024-01-31)
Population	Number of Regular registered patients
Appointment_Count	Number of all booked appointments
Appointment_Duration	Total planned duration of all booked appointments in minutes
SameDay_Count	Number of appointments booked on the day of the appointment
SameDay_Duration	Total planned duration of appointments booked on the day of the appointment in minutes
BookedInAdvance_Count	Number of appointments booked in advance of the day of the appointment
BookedInAdvance_Duration	Total planned duration of appointments booked in advance of the day of the appointment in minutes
DNA_Count	Number of appointments where the patient did not attend
DNA_Duration	Total planned duration of appointments where the patient did not attend in minutes
EnhancedAccess_Count	Number of booked appointments classified as 'Enhanced Access' i.e., booked appointments between the hours of 6.30pm to 8pm Mondays to Fridays and between 9am and 5pm on Saturdays
EnhancedAccess_Duration	Total planned duration of booked appointments classified as 'Enhanced Access' in minutes i.e., booked appointments between the hours of 6.30pm to 8pm Mondays to Fridays and between 9am and 5pm on Saturdays
ModeOfContact_FaceToFace_Count	Number of booked appointments delivered face-to-face
ModeOfContact_FaceToFace_Duration	Total planned duration of booked appointments delivered face-to-face in minutes
ModeOfContact_Telephone_Count	Number of booked appointments delivered by telephone
ModeOfContact_Telephone_Duration	Total planned duration of booked appointments delivered by telephone in minutes
ModeOfContact_HomeVisit_Count	Number of booked appointments delivered by home visit
ModeOfContact_HomeVisit_Duration	Total planned duration of booked appointments delivered by home visit in minutes
ModeOfContact_Digital_Count	Number of booked appointments delivered by digital means
ModeOfContact_Digital_Duration	Total planned duration of booked appointments delivered by digital means in minutes
ModeOfContact_NonCategorised_Count	Number of booked appointments where the mode-of-contact cannot be categorised i.e. not categorised as face-to-face, telephone, home visit or digital
ModeOfContact_NonCategorised_Duration	Total planned duration of booked appointments where the mode-of-contact cannot be categorised in minutes i.e. not categorised as face-to-face, telephone, home visit or digital
BookedToSeen_OnTheDay	Number of appointments booked on the day of the appointment
BookedToSeen_1Day	Number of appointments booked 1 day prior to the appointment date
BookedToSeen_2to3Days	Number of appointments booked 2 to 3 days prior to the appointment date
BookedToSeen_4to5Days	Number of appointments booked 4 to 5 days prior to the appointment date
BookedToSeen_6to7Days	Number of appointments booked 6 to 7 days prior to the appointment date
BookedToSeen_8to13Days	Number of appointments booked 8 to 13 days prior to the appointment date
BookedToSeen_14plusDays	Number of appointments booked 14 days or more prior to the appointment date