

Data Protection Impact Assessment (DPIA) For Extended Access Services Provided by Leeds GP Confederation (includes all activity hosted by the Confederation Network Infrastructure)

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Date approved by Quality, Performance and Finance Committee	Operationally Approved by IG Project Team on 22 nd July 2019 pending Committee Approval
Version	1.1
Review date	July 2020

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Data Protection Impact Assessment (DPIA)

Section 1

System/Project General Details

System/project/process (referred to thereafter as 'project') title: Objective: Detail: Why is the new system/change in system required? Is there an	Confederation Extended Access Booking Platform (use by Confed Staff), 2. Confederation Staff Access of GP Clinical Systems, 3. Use of Confederation Clinical System by GP Practice / PCN Staff. — To be known Collectively as Confederation Clinical System Use. To have an IG compliant citywide service infrastructure managed by the Confederation for the benefit of the patients of Leeds. With the establishment of the PCN's from 1st July, services will be delivered at scale to achieve the outcomes of the Network DES of the GMS/PMS Contract.		
approved business case? Stakeholders/Relationships /Partners: Please outline the nature of such relationships and the corresponding roles of other organisations.	GP Practices, PCNs, Federations, LCH, Leeds CCG		
Other related projects:	CAS pilot		
Project lead:	Name: Title: Department: Telephone: Email	Simon Boycott Head of Development and Governance Executive Team 07717 841665 simon.boycott@nhs.net	
Information Asset owners/Ad	ministrators (if a	applicable)	
All information systems/assets must have an Information Asset Owner (IAO). IAO's should normally be a Head of Department/Service. Information Asset Administrator: Information systems/assets may have an Information Asset Administrator (IAA) who reports	Title: Department: Telephone: Email Name: Title: Department: Telephone: Email	Ciara O'Grady Business Transformation Lead Digital 07896020876 ciara.dunne@nhs.net Matt Walker Business and Operations Manager Operations 0113 887 3899 Matthew.walker15@nhs.net	
the IAO. IAA's are normally System Managers/Project Leads.	LIIIdii	iviattiiew.waikei 13@iiii5.iiet	

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<u>Section 2A – Confederation Extended Access Booking Platform (use by Confed Staff)</u>

Data Protection Impact Assessment Key Questions

	Question	Response
1.	Will the project use identifiable or potentially identifiable data in any way? If answered 'No' then a DPIA is not normally suggested.	 ✓ Yes ✓ No If yes, who will this data relate to: ✓ Patient ✓ Staff ✓ Other: Click here to enter text.
2.	Please state purpose for the	Patient care
	For example, patient care, commissioning, research, audit, evaluation.	
3.	Please tick the data items that	
	Personal Special categories	 Name Post Code Date of Birth GP Practice Date of Death NHS Number Passport Number Pseudonymised Data Online Identifiers (e.g. IP Number, Mobile Device ID) Health Data Trade Union membership
	of personal data (sensitive data)	 ☐ Political opinions ☐ Racial or Ethnic Origin ☐ Sex life and sexual orientation ☐ Genetic Data ☐ Other:
4.	The data of approximately	□1-10
	how many individuals will be affected?	□10-100 □100-1000 □1000-10 000 □10 000-100 000 ⊠100 000+ □Unable to ascertain Click here to enter text.
5.	Have the individuals been	☐ Yes
	informed of this Data Processing activity	If yes, please specify: Click here to enter text.
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	Question	Response
6.	Will this activity create a new Information Asset for the Practice?	
		Has an Information Asset Owner been identified and does the Information Asset and Data Flow Register require updating?
		Does this project constitute a change to existing Information Asset(s) or is this a new Information Asset? ☑ Yes □ No
		If yes, include the completed Information Asset Register and Data Flow Mapping Form for risk review.
7.	Who will be the Data Controller for this activity? The data controller is the individual or organisation who is responsible for determining the reason for the data processing activity, who may not be the "holder" of the data	Simon Boycott
8.	Will a third party be processing data as part of this activity	☐ Yes ☐ No If "Yes" please ensure that the Data Protection Impact Assessment Key Questions for Providers/Processors section of this document is filled in by the Provider. Also ensure that either a) the third party/supplier contract(s) include all the necessary Information Governance clauses regarding Data Protection and Freedom of Information b) Is the contract based on or utilises the NHS standard
		contract If neither are done, please records as a risk

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	Question	Response
9.	What legal basis enables this data processing? For more information about conditions for processing, please see the ICO's GDPR website.	Personal data (identifiers and potentially identifiable data): Consent: Click here to enter text. Relating to a contract: APMS Contract (GPAF) Legal obligation: APMS Contract to provide Primary Medical Care Vital interests: Click here to enter text. Public task: Click here to enter text. Other: Click here to enter text. Special categories of personal data (sensitive data), if applicable: Consent: Click here to enter text. Medical related: Click here to enter text. Public Health: Click here to enter text. Employment related: Click here to enter text. Already public: Click here to enter text. Legal claim related: Click here to enter text. Substantial public interest: Click here to enter text. Other: Click here to enter text.
10.	Are you relying on individuals (patients/staff) to explicit consent to the processing of personal identifiable or sensitive data? Please provide copies of any consent documentation that will be used, including patient information leaflets	 ☐ Yes ☐ No (skip next question) How will consent be obtained and by whom? Click here to enter text. Will the consent cover all proposed processing and sharing/disclosures? ☐ Yes ☐ No If no, please detail: Click here to enter text.
11.	If you are relying only on consent, did you consider any other legal basis? Please be aware that consent may not be the best legal basis to use under many circumstances due to the strengthened rights it gives individuals over their data.	☐ Yes ☐ No ☐ N/A If no, please detail why: Click here to enter text.
12.	Who will have access to the data within the project? Please refer to roles/job titles/organisations.	Confederation Service Delivery Staff including employed, seconded and non-contracted aligned staff. Will include receptionists, administrators, GP's Nurses and other allied health professionals.

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	Question	Response
13.	Have consultation/checks have been made regarding the adequacy, relevance and necessity for the processing of the data for this project?	☐ Yes ☐ No (please record as a risk) If yes, please specify: No consultation made – service and contract is inoperable without the use of the data in this way and consultation would not materially change the use of data, nor the approach to service delivery.
14.	Does the project involve linkage of personal data with data in other collections, or significant change in data linkages? The degree of concern is higher where data is transferred out of its original context (e.g. the sharing and merging of datasets can allow for a collection of a much wider set of information than needed and identifiers might be collected/linked which prevents personal data being kept anonymously)	☐ Yes (please record as a risk) ☐ No If yes, please provide a data flow diagram showing how identifiable information would flow and ensure this is added to the practice Information Asset and Data Flow Register (see Information Assets and Data Flows section).
15.	Has stakeholder engagement taken place?	 ✓ Yes ☐ No (please record as a risk) If yes, how have any issues identified by stakeholders been considered? General Information Governance concerns around data sharing which are addressed by the Confed IG Management Framework If no, please outline any plans in the near future to seek stakeholder feedback: Engagement will be ongoing through the Citywide EA Steering Group.
16.	Does the project involve any new data sharing between stakeholder organisations?	Yes (consider if this will be a risk) ⊠ No If yes, please describe: Click here to enter text. Please provide a high level data flow diagram showing how identifiable information would flow.
17.	Does the project involve the collection of data that may be unclear or intrusive? Are all data items clearly defined? Is the data collected limited to a specific set of predefined categories?	☐ Yes (please record as a risk) ☐ No If yes, please explain: Click here to enter text.

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	Question	Response
18.	What are the specific	Health Records – 10yrs after the patient is deceased.
	retention periods for this data? Please refer to the Records Management Code of Practice for Health and Social Care 2016 and list the retention period for identifiable project datasets.	If no retention period is specified, please record as a risk
19.	Will the data be securely	
	destroyed when it is no longer	
	required?	If no, please detail: Click here to enter text.
20.	Will identifiable/potentially	\square Yes (please record as a risk) \boxtimes No
	identifiable from the project	Maria da ancida da Citababa da antanta da Citababa da Citaba da Citababa da Citababa da Citababa da Citababa da Citababa da Citaba da Citaba da Citaba da Citababa da Citaba da Citaba da
	be released as Open Data (placed in to the public	If yes, please describe: Click here to enter text.
	domain)?	
21.	Will any personal and/or	☐ Yes (please record as a risk) ☐ No
	sensitive data be transferred	
	to a country outside the UK?	If yes, which data and to which country?
		Click here to enter text.
22.	Will identifiable data only be handled within the patients'	
	direct care team (in	a risk)
	accordance with the Common	If no, please detail:
	Law Duty of Confidentiality)?	Click here to enter text.
23.	Will an evaluation of the	⊠ Yes □ No
	activity be required?	
		If yes, has a suitable data set been decided, that specifies what
		data will be used, where it will be extracted from and what measures are in place (anonymization, pseudonymisation etc)
		to protect personal data
		oxtimes Yes $oxtimes$ No (please record as a risk)

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<u>Section 3A – Confederation Extended Access Booking Platform (use by Confed Staff)</u>

Data Protection Impact Assessment Key Questions for Providers/Processors

	Question	Response
1.	Is the Provider/Data	☐ Yes ☐ No (please record as a risk)
	Processor registered with the	· ·
	Information Commissioner?	Organisation: Click here to enter text.
		Data Protection Registration Number: Click here to enter text.
2.	Has the Provider/Data Processor completed and published a satisfactory Data Security and Protection Toolkit submission? Please note that the Data Security and Protection Toolkit replaced the IG Toolkit from 1 April 2018.	☐ Yes ☐ No (please record as a risk) If yes, please give organisation code and percentage score: DSP Toolkit to be completed in respect of FY19/20 upon completion of IG Management Framework Implementation DSP/IG Toolkit Score: ☐ Satisfactory ☐ Not satisfactory (please record as a risk) ☐ Satisfactory with Improvement Plan If satisfactory with an improvement plan, please request a copy of the plan and enclose it with this assessment. If not satisfactory, please explain how the service has been procured: Click here to enter text.
3.	Will other third parties (not already identified) have access to the data, or act as Provider/Data Processors? Include any external organisations. Please ensure any third party organisation that will have access to this data also complete a DPIA.	☐ Yes (please consider as a risk) ☐ No If so, for what purpose? Click here to enter text. Please list organisations and by what means of transfer: Click here to enter text.
4.	Where will the data be kept/stored/accessed? Where applicable, please refer to data flow diagram.	Only available electronically
5.	Please indicate all methods in which data will be transferred	☐ Fax ☐ Email (Unsecure/Personal) ☐ Email (Secure/nhs.net) ☐ Internet (unsecure – e.g. http) ☐ Telephone ☐ Internet (secure – e.g. https) ☐ By hand ☐ Courier ☐ Post – track/traceable ☐ Post – normal ☐ Software ☐ Mobile app ☐ Other: Click here to enter text.
6.	How will the data be kept up to date and checked for accuracy and completeness?	Links to the NHS spine

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7.	Please outline how individuals will be informed and kept informed about how their data will be processed. A copy of the privacy notice and/or leaflets must be provided.	Our Privacy Notice will be made available on the confederation website and within each physical service location.
8.	How will consent/non- consent (if applicable), objections or opt-outs be recorded and respected?	As per the Confederation Consent Policy and via patient medical records.
9.	What arrangements are in place to process Subject Access Requests? Please include a copy of the SAR procedure if one exists	As per the Confederation Individual Rights and SAR Procedue.
10.	What process is in place for rectifying/blocking data? What would happen if such a request were made?	As at 9.
11.	Will the processing of data be automated? Will the proposed processing of data involved automated means of processing to determine an outcome for the individual?	 ☐ Yes(please record as a risk ☐ No ☑ Not applicable If yes, please outline what arrangements are available to enable the individual access and to extract data (in a standard file format). Please also detail any profiling that may take place as part through automated processing: Click here to enter text.
12.	Is there a useable audit trail in place for the project? For example, to identify who has accessed a record?	 ✓ Yes ☐ No (please record as a risk) ☐ Not applicable If yes, please outline the audit plan: Quarterly User Access Audits
13.	Is there an Access Control Policy in place for the Data/the systems the data is held within?	 ✓ Yes ☐ No (please record as a risk) ☐ Not applicable If yes, please outline the policy and how it is implemented: as per the Confederation Network Security Policy and Confederation Standard Operating Procedure for Hub Staff Registration.
14.	Does the project involve privacy enhancing technologies? New forms of encryption, two factor authentication and/or pseudonymisation.	☐ Yes ☐ No If yes, please give details: Click here to enter text.

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15.	Will the project involve the sending of unsolicited	oxtimes Yes (please record as a risk) $oxtimes$ No
	marketing messages	If yes, what communications will be sent?
	electronically such as	Appointment reminder and Friends and Family Friends Test
	telephone, fax, email and	feedback request SMS Messages
	text?	
	Please note that seeking to influence	Will consent be sought prior to this?
	an individual is considered to be marketing.	
		If no, please explain why consent is not being sought first: Click here to enter text.
16.	Have the business continuity	
	requirements been	☐ Business Continuity is not applicable
	considered?	
		Diago cyplain and aither reference how such plans link with
		Please explain and either reference how such plans link with
		the organisational plan or why there are no business
		·
		the organisational plan or why there are no business

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<u>Section 2B – Confederation Staff Access of GP Clinical Systems</u>

Data Protection Impact Assessment Key Questions

	Question	Response
1.	Will the project use identifiable or potentially identifiable data in any way? If answered 'No' then a DPIA is not normally suggested.	 ✓ Yes If yes, who will this data relate to: ✓ Patient ✓ Staff ☐ Other: Click here to enter text.
2.	Please state purpose for the	Patient care.
	processing of the data: For example, patient care, commissioning, research, audit, evaluation.	
3.	Please tick the data items that	
	Personal	 Name Post Code Date of Birth GP Practice NHS Number Passport Number Pseudonymised Data Online Identifiers (e.g. IP Number, Mobile Device ID)
	Special categories of personal data (sensitive data)	 ☑ Health Data ☐ Political opinions ☑ Religion ☑ Racial or Ethnic Origin ☑ Biometric Data ☐ Other:
4.	The data of approximately	□1-10
	how many individuals will be affected?	□10-100 □100-1000 □1000-10 000 □10 000-100 000 ⊠100 000+ □Unable to ascertain Click here to enter text.
5.	Have the individuals been	☐ Yes ☐ No (please record as a risk)
	informed of this Data Processing activity	If yes, please specify:
	0 1	

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	Question	Response
6.	Will this activity create a new Information Asset for the Practice?	☐ Yes
		Has an Information Asset Owner been identified and does the Information Asset and Data Flow Register require updating?
		☐ Yes ☐ No If yes, include the completed Information Asset Register New Entry Form.
		Does this project constitute a change to existing Information Asset(s) or is this a new Information Asset? ☐ Yes ☐ No
		If yes, include the completed Information Asset Register and Data Flow Mapping Form for risk review.
7.	Who will be the Data Controller for this activity? The data controller is the individual or organisation who is responsible for determining the reason for the data processing activity, who may not be the "holder" of the data	caldicott guardian for the GP practice
8.	Will a third party be processing data as part of this	☐ Yes ⊠ No
	activity	If "Yes" please ensure that the <u>Data Protection Impact</u> <u>Assessment Key Questions for Providers/Processors</u> section of this document is filled in by the Provider. Also ensure that either c) the third party/supplier contract(s) include all the necessary Information Governance clauses regarding Data Protection and Freedom of Information d) Is the contract based on or utilises the NHS standard
		contract If neither are done, please records as a risk

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	Question	Response
9.	What legal basis enables this data processing? For more information about conditions for processing, please see the ICO's GDPR website.	Personal data (identifiers and potentially identifiable data): Consent: Click here to enter text. Relating to a contract: Click here to enter text. Legal obligation: APMS Contract to Provide Primary Medical Care Vital interests: Click here to enter text. Public task: Click here to enter text. Other: Click here to enter text. Special categories of personal data (sensitive data), if applicable: Consent: Click here to enter text. Medical related: APMS Contract for Primary Care Services Public Health: Click here to enter text. Employment related: Click here to enter text. Vital interests: Click here to enter text. Already public: Click here to enter text. Legal claim related: Click here to enter text. Substantial public interest: Click here to enter text. Other: Click here to enter text.
10.	Are you relying on individuals (patients/staff) to explicit consent to the processing of personal identifiable or sensitive data? Please provide copies of any consent documentation that will be used, including patient information leaflets	 ☐ Yes ☐ No (skip next question) How will consent be obtained and by whom? Will the consent cover all proposed processing and sharing/disclosures? ☐ Yes ☐ No If no, please detail: Click here to enter text.
11.	If you are relying only on consent, did you consider any other legal basis? Please be aware that consent may not be the best legal basis to use under many circumstances due to the strengthened rights it gives individuals over their data.	☐ Yes ☐ No ☑ N/A If no, please detail why: Click here to enter text.
12.	Who will have access to the data within the project? Please refer to roles/job titles/organisations.	Clinical staff with Smartcard access working in Extended Access Services on behalf of the GP practice, Confederation RA Sponsors, GP Practice Staff with Smartcard access.

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	Question	Response
13.	Have consultation/checks have been made regarding the adequacy, relevance and necessity for the processing of the data for this project?	☐ Yes ☐ No (please record as a risk) If yes, please specify: The data to be processed is mandated as part of the APMS contract and therefore the legal basis for processing.
14.	Does the project involve linkage of personal data with data in other collections, or significant change in data linkages? The degree of concern is higher where data is transferred out of its original context (e.g. the sharing and merging of datasets can allow for a collection of a much wider set of information than needed and identifiers might be collected/linked which prevents personal data being kept anonymously)	☐ Yes (please record as a risk) ☐ No If yes, please provide a data flow diagram showing how identifiable information would flow and ensure this is added to the practice Information Asset and Data Flow Register (see Information Assets and Data Flows section).
15.	Has stakeholder engagement taken place?	 ✓ Yes ☐ No (please record as a risk) If yes, how have any issues identified by stakeholders been considered? General Information Governance concerns around data sharing which are addressed by the Confed IG Management Framework If no, please outline any plans in the near future to seek stakeholder feedback: Engagement will be ongoing through the Citywide EA Steering Group.
16.	Does the project involve any new data sharing between stakeholder organisations?	☐ Yes (consider if this will be a risk) ☐ No If yes, please describe: Click here to enter text. Please provide a high level data flow diagram showing how identifiable information would flow.
17.	Does the project involve the collection of data that may be unclear or intrusive? Are all data items clearly defined? Is the data collected limited to a specific set of predefined categories?	☐ Yes (please record as a risk) ☐ No If yes, please explain: Click here to enter text.

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	Question	Response
18.	What are the specific retention periods for this data? Please refer to the Records	GP Records retain for 10 years after death. Electronic patient records (EPRs) must not be destroyed, or deleted, for the foreseeable future.
	Management Code of Practice for Health and Social Care 2016 and list the retention period for identifiable project datasets.	If no retention period is specified, please record as a risk
19.	Will the data be securely	oxtimes Yes $oxtimes$ No (please record as a risk)
	destroyed when it is no longer required?	If no, please detail: Click here to enter text.
20.	Will identifiable/potentially identifiable from the project	\square Yes (please record as a risk) \boxtimes No
	be released as Open Data (placed in to the public domain)?	If yes, please describe: Click here to enter text.
21.	Will any personal and/or sensitive data be transferred	☐ Yes (please record as a risk) ☐ No
	to a country outside the UK?	If yes, which data and to which country? Click here to enter text.
22.	Will identifiable data only be handled within the patients'	oximes Yes $oximes$ No (please consider if this will be a risk)
	direct care team (in	
	accordance with the Common common common common color: blue; the common color: blue;	If no, please detail: Click here to enter text.
23.	Will an evaluation of the activity be required?	⊠ Yes □ No
		If yes, has a suitable data set been decided, that specifies what data will be used, where it will be extracted from and what measures are in place (anonymization, pseudonymisation etc) to protect personal data

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<u>Section 3B – Confederation Staff Access of GP Clinical Systems</u>

Data Protection Impact Assessment Key Questions for Providers/Processors

	Question	Response
1.	Is the Provider/Data Processor registered with the Information Commissioner?	☐ Yes ☐ No (please record as a risk)
2.	Has the Provider/Data Processor completed and published a satisfactory Data Security and Protection Toolkit submission? Please note that the Data Security and Protection Toolkit replaced the IG Toolkit from 1 April 2018.	☐ Yes ☐ No (please record as a risk) If yes, please give organisation code and percentage score: DSP Toolkit to be completed in respect of FY19/20 upon completion of IG Management Framework Implementation DSP/IG Toolkit Score: ☐ Satisfactory ☐ Not satisfactory (please record as a risk) ☐ Satisfactory with Improvement Plan If satisfactory with an improvement plan, please request a copy of the plan and enclose it with this assessment.
		If not satisfactory, please explain how the service has been procured:
3.	Will other third parties (not already identified) have access to the data, or act as Provider/Data Processors? Include any external organisations. Please ensure any third party organisation that will have access to this data also complete a DPIA.	 ☐ Yes (please consider as a risk) ☐ No If so, for what purpose? Click here to enter text. Please list organisations and by what means of transfer: Click here to enter text.
4.	Where will the data be kept/stored/accessed? Where applicable, please refer to data flow diagram.	GP Clinical System
5.	Please indicate all methods in which data will be transferred	 ☑ Fax ☐ Email (Unsecure/Personal) ☑ Email (Secure/nhs.net) ☐ Internet (unsecure – e.g. http) ☑ Telephone ☐ Internet (secure – e.g. https) ☑ By hand ☑ Courier ☑ Post – track/traceable ☑ Post – normal ☑ Software ☑ Mobile app ☐ Other: Click here to enter text.
6.	How will the data be kept up to date and checked for accuracy and completeness?	Local GP practices policies as part of their internal IG processes.

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7.	Please outline how individuals will be informed and kept informed about how their data will be processed. A copy of the privacy notice and/or leaflets must be provided.	Our Privacy Notice is available on our website and will be displayed in each service location
8.	How will consent/non- consent (if applicable), objections or opt-outs be recorded and respected?	As per the Confederation Consent Policy and via the GP Practice Clinical System
9.	What arrangements are in place to process Subject Access Requests? Please include a copy of the SAR procedure if one exists	As per the Confederation Individual Rights and Subject Access Request Procedure.
10.	What process is in place for rectifying/blocking data? What would happen if such a request were made?	As at 9.
11.	Will the processing of data be automated? Will the proposed processing of data involved automated means of processing to determine an outcome for the individual?	☐ Yes(please record as a risk ☐ No ☐ Not applicable If yes, please outline what arrangements are available to enable the individual access and to extract data (in a standard file format). Please also detail any profiling that may take place as part through automated processing: Click here to enter text.
12.	Is there a useable audit trail in place for the project? For example, to identify who has accessed a record?	 ✓ Yes ☐ No (please record as a risk) ☐ Not applicable If yes, please outline the audit plan: Audit log in EMIS and systmone.
13.	Is there an Access Control Policy in place for the Data/the systems the data is held within?	 ✓ Yes ☐ No (please record as a risk) ☐ Not applicable If yes, please outline the policy and how it is implemented: Data sharing agreement in place issued by the Confederation; Confederation Network Security Policy and SOP's
14.	Does the project involve privacy enhancing technologies? New forms of encryption, two factor authentication and/or pseudonymisation.	☐ Yes No If yes, please give details: Click here to enter text.

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15.	Will the project involve the sending of unsolicited	\square Yes (please record as a risk) \boxtimes No
	marketing messages	If yes, what communications will be sent?
	electronically such as	Click here to enter text.
	telephone, fax, email and	
	text?	Will consent be sought prior to this?
	Please note that seeking to influence an individual is considered to be	☐ Yes ☐ No (please record as a risk)
	marketing.	If no, please explain why consent is not being sought first:
		Click here to enter text.
16.	Have the business continuity	
	requirements been	☐ Business Continuity is not applicable
	considered?	
		Please explain and either reference how such plans link with
		the organisational plan or why there are no business
		continuity considerations that are applicable for this project:
		Business Continuity plans are managed via cloud-based
		software and approved by the QPF Committee.

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<u>Section 2C – Use of Confederation Clinical System by GP Practice / PCN Staff</u>

Data Protection Impact Assessment Key Questions

	Question	Response
1.	Will the project use identifiable or potentially identifiable data in any way? If answered 'No' then a DPIA is not normally suggested.	 ✓ Yes If yes, who will this data relate to: ✓ Patient ✓ Staff ☐ Other: Click here to enter text.
2.	Please state purpose for the processing of the data: For example, patient care, commissioning, research, audit, evaluation.	Patient Care; delivery of medical services at scale within Primary Care Networks; operation of a centrally supported, scaled appointments system.
3.	Please tick the data items that are held in the system Personal Special categories of personal data (sensitive data)	☑ Name ☑ Address ☑ Post Code ☑ Date of Birth ☑ GP Practice ☐ Date of Death ☑ NHS Number ☐ NI Number ☐ Passport Number ☐ Pseudonymised Data ☐ Online Identifiers (e.g. IP Number, Mobile Device ID) ☐ Health Data ☐ Trade Union membership ☐ Political opinions ☐ Religion ☐ Racial or Ethnic Origin ☐ Sex life and sexual orientation ☐ Biometric Data ☐ Genetic Data
 4. 5. 	The data of approximately how many individuals will be affected? Have the individuals been informed of this Data	☐ Other: ☐ 1-10 ☐ 10-100 ☐ 100-1000 ☐ 1000-10 000 ☐ 10 000-100 000 ☒ 100 000+ ☐ Unable to ascertain Click here to enter text. ☐ Yes ☒ No (please record as a risk) If yes, please specify: Click here to enter text.
	Processing activity	

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6.		Response
	Will this activity create a new Information Asset for the Practice?	☐ Yes No If yes
		Has an Information Asset Owner been identified and does the Information Asset and Data Flow Register require updating?
		☐ Yes ☐ No If yes, include the completed Information Asset Register New Entry Form.
		Does this project constitute a change to existing Information Asset(s) or is this a new Information Asset? ☐ Yes ☐ No
		If yes, include the completed Information Asset Register and Data Flow Mapping Form for risk review.
7.	Who will be the Data Controller for this activity? The data controller is the individual or organisation who is responsible for determining the reason for the data processing activity, who may not be the "holder" of the data	Simon Boycott
8.	Will a third party be processing data as part of this activity	☐ Yes ☐ No If "Yes" please ensure that the Data Protection Impact Assessment Key Questions for Providers/Processors section of this document is filled in by the Provider. Also ensure that either e) the third party/supplier contract(s) include all the necessary Information Governance clauses regarding Data Protection and Freedom of Information f) Is the contract based on or utilises the NHS standard contract If neither are done, please records as a risk

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i	Question	Response
9.	What legal basis enables this data processing? For more information about conditions for processing, please see the ICO's GDPR website.	Personal data (identifiers and potentially identifiable data): Consent: Click here to enter text. Relating to a contract: Click here to enter text. Legal obligation: GMS/PMS Contract – Network DES, delegated processing under GP Confederation Data Security and Protection Framework Agreement. Vital interests: Click here to enter text. Public task: Click here to enter text. Other: Click here to enter text. Special categories of personal data (sensitive data), if applicable: Consent: Click here to enter text. Medical related: Click here to enter text. Public Health: Click here to enter text. Employment related: Click here to enter text. Vital interests: Click here to enter text. Already public: Click here to enter text. Legal claim related: Click here to enter text. Substantial public interest: Click here to enter text. Other: Click here to enter text.
10.	Are you relying on individuals (patients/staff) to explicit consent to the processing of personal identifiable or sensitive data? Please provide copies of any consent documentation that will be used, including patient information leaflets	 ☐ Yes ☐ No (skip next question) How will consent be obtained and by whom? Click here to enter text. Will the consent cover all proposed processing and sharing/disclosures? ☐ Yes ☐ No If no, please detail: Click here to enter text.
11.	If you are relying only on consent, did you consider any other legal basis? Please be aware that consent may not be the best legal basis to use under many circumstances due to the strengthened rights it gives individuals over their data.	☐ Yes ☐ No ☑ N/A If no, please detail why: Click here to enter text.
12.	Who will have access to the data within the project? Please refer to roles/job titles/organisations.	GP Practice and Primary Care Network based staff, GP Confederation based staff.

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	Question	Response
13.	Have consultation/checks have been made regarding the adequacy, relevance and necessity for the processing of the data for this project?	☑ Yes ☐ No (please record as a risk) If yes, please specify: The need for, and form of information to be processed has been evaluated and no alternative option for the delivery of the service is available to the Confederation and its members for whom the Confederation will process data.
14.	Inkage of personal data with data in other collections, or significant change in data linkages? The degree of concern is higher where data is transferred out of its original context (e.g. the sharing and merging of datasets can allow for a collection of a much wider set of information than needed and identifiers might be collected/linked which prevents personal data being kept anonymously)	☐ Yes (please record as a risk) ☐ No If yes, please provide a data flow diagram showing how identifiable information would flow and ensure this is added to the practice Information Asset and Data Flow Register (see Information Assets and Data Flows section).
15.	Has stakeholder engagement taken place?	 ✓ Yes ☐ No (please record as a risk) If yes, how have any issues identified by stakeholders been considered? Issues of accessibility and IG fed back through Primary Care Development Team, addressed through IG Management Framework. If no, please outline any plans in the near future to seek stakeholder feedback: PCD Team work with PCN's on an ongoing basis
16.	Does the project involve any new data sharing between stakeholder organisations?	☐ Yes (consider if this will be a risk) ☒ No If yes, please describe: Click here to enter text. Please provide a high level data flow diagram showing how identifiable information would flow.
17.	Does the project involve the collection of data that may be unclear or intrusive? Are all data items clearly defined? Is the data collected limited to a specific set of predefined categories?	☐ Yes (please record as a risk)

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	Question	Response
18.	What are the specific	Health Care Record Retention Period.
	retention periods for this data? Please refer to the Records Management Code of Practice for Health and Social Care 2016 and list the retention period for identifiable project datasets.	If no retention period is specified, please record as a risk
19.	Will the data be securely	
	destroyed when it is no longer	
	required?	If no, please detail: Click here to enter text.
20.	Will identifiable/potentially	\square Yes (please record as a risk) \boxtimes No
	identifiable from the project be released as Open Data	If you place describe. Click have to enter tout
	(placed in to the public	If yes, please describe: Click here to enter text.
	domain)?	
21.	Will any personal and/or	☐ Yes (please record as a risk) ☐ No
	sensitive data be transferred	
	to a country outside the UK?	If yes, which data and to which country? Click here to enter text.
22.	M/ill identifiable data only be	
22.	Will identifiable data only be handled within the patients'	oxinesize Yes $oxinesize$ No (please consider if this will be a risk)
	direct care team (in	a risky
	accordance with the Common	If no, please detail:
	Law Duty of Confidentiality)?	Click here to enter text.
23.	Will an evaluation of the	☐ Yes
	activity be required?	
		If yes, has a suitable data set been decided, that specifies what data will be used, where it will be extracted from and what
		measures are in place (anonymization, pseudonymisation etc)
		to protect personal data
		□ No (places passed as a girl)
		\square Yes \square No (please record as a risk)

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Section 3C Data Protection Impact Assessment Key Questions for Providers/Processors

	Question	Response
1.	Is the Provider/Data	
	Processor registered with the	,,
	Information Commissioner?	Organisation: GP Practices in Leeds
		Data Protection Registration Number: Requirement to be
		registered signed up to in the DSP Framework Agreement.
2.	Has the Provider/Data	
	Processor completed and	If yes, please give organisation code and percentage score:
	published a satisfactory Data	Self-declaration by practices in the DSP Framework
	Security and Protection	Agreement.
	Toolkit submission?	
	Please note that the Data Security	DSP/IG Toolkit Score:
	and Protection Toolkit replaced the	☐ Satisfactory ☐ Not satisfactory (please record
	IG Toolkit from 1 April 2018.	as a risk)
		☐ Satisfactory with Improvement Plan
		If satisfactory with an improvement plan, please request a
		copy of the plan and enclose it with this assessment.
		If not satisfactory, please explain how the service has been
		procured:
		Click here to enter text.
3.	Will other third parties (not	☐ Yes (please consider as a risk) ☐ No
	already identified) have	,
	access to the data, or act as	If so, for what purpose?
	Provider/Data Processors?	Click here to enter text.
	Include any external organisations.	
	Please ensure any third party	Please list organisations and by what means of transfer:
	organisation that will have	Click here to enter text.
	access to this data also	
	complete a DPIA.	
4.	Where will the data be	Confederation Clinical System.
	kept/stored/accessed?	
	Where applicable, please refer to	
	data flow diagram.	
5.	Please indicate all methods in	Fax Email (Unsecure/Personal)
	which data will be transferred	☐ Email (Secure/nhs.net) ☐ Internet (unsecure – e.g. http)
		☐ Telephone ☐ Internet (secure – e.g. https)
		☐ By hand ☐ Courier
		☐ Post – track/traceable ☐ Post – normal
		oxtimes Software $oxtimes$ Mobile app
		☐ Other: Click here to enter text.

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6.	How will the data be kept up to date and checked for accuracy and completeness?	Data stored is verified as accurate when stored through the clinical system link to the NHS spine		
7.	Please outline how individuals will be informed and kept informed about how their data will be processed. A copy of the privacy notice and/or leaflets must be provided.	Privacy Notice is available on the confederation website and displayed in service locations.		
8.	How will consent/non- consent (if applicable), objections or opt-outs be recorded and respected?	As per Confederation Consent policy.		
9.	What arrangements are in place to process Subject Access Requests? Please include a copy of the SAR procedure if one exists	Confederation Individual Right and SAR Procedure		
10.	What process is in place for rectifying/blocking data? What would happen if such a request were made?	As at 9.		
11.	Will the processing of data be automated? Will the proposed processing of data involved automated means of processing to determine an outcome for the individual?	☐ Yes(please record as a risk ☐ No ☐ Not applicable If yes, please outline what arrangements are available to enable the individual access and to extract data (in a standard file format). Please also detail any profiling that may take place as part through automated processing: Click here to enter text.		
12.	Is there a useable audit trail in place for the project? For example, to identify who has accessed a record?	✓ Yes☐ No (please record as a risk)☐ Not applicableIf yes, please outline the audit plan: Click here to enter text.		
13.	Is there an Access Control Policy in place for the Data/the systems the data is held within?	 ✓ Yes ☐ No (please record as a risk) ☐ Not applicable If yes, please outline the policy and how it is implemented: GP Confederation Network Security Policy is in place and supported by the DSP Framework Agreement. 		
14.	Does the project involve privacy enhancing technologies? New forms of encryption, two factor authentication and/or pseudonymisation.	☐ Yes		

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15.	Will the project involve the sending of unsolicited	oxtimes Yes (please record as a risk) $oxtimes$ No			
	marketing messages	If yes, what communications will be sent?			
	electronically such as	Appointment reminder SMS Messages			
	telephone, fax, email and				
	text?	Will consent be sought prior to this?			
	Please note that seeking to influence an individual is considered to be				
	marketing.	If no, please explain why consent is not being sought first:			
		Click here to enter text.			
16.	Have the business continuity				
	requirements been	☐ Business Continuity is not applicable			
	considered?				
		Please explain and either reference how such plans link with			
		the organisational plan or why there are no business			
		continuity considerations that are applicable for this project:			
		Business Continuity plans are managed via cloud-based			
		software and approved by the QPF Committee.			

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Section 4: Data Protection Impact Assessment Information Governance Review

Information Governance Review				Response (for completion by project lead)		
	Issue	Potential Risk	Recommendation	Agreed Action	Completion (Date and Initials)	
2A-13	No consultation undertaken regarding adequacy, relevance and necessity for data processing.	The Confed processes data without justification or legal basis	None – Project & dataset is not complex or ambiguous and adequacy, relevance and necessity is self-evident	None	N/A	
3A-1	Confed not registered with ICO	Confed faces regulatory action	Confed to register with the ICO	MW has registered and waiting for confirmation from ICO	Complete July19	
3A-2	Confed has not completed DSP Toolkit	Confederation lacks member confidence and/or faces regulatory action for not meeting required standards	DSP Toolkit is completed as part of the implementation of the IG Management Framework.	DSP Toolkit completed by 31 st August 2019	31 st August 2019	
3A-15	Data processing will involve unsolicited SMS Messaging	Complaint re privacy or data breach through poor IG connected with use of SMS Messaging.	Apply an implied consent model for SMS messaging and update the Privacy Notice to inform Service Users. Monitor the use of SMS messaging through quality reporting to the QPF Committee – if a theme emerges in feedback, management action to be taken	Implied consent will be applied to all service users and the privacy notice updated to inform service users of this policy.	31 st July 2019	
2B-5	Individuals have not been informed of the data processing	Individual rights of data subjects are breached	Clause to be added to DSP Framework Agreement requiring parties to keep privacy notices updated with relevant information	Add relevant clause to Framework Agreement	Complete July19	

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2B-13	No consultation undertaken regarding adequacy, relevance and necessity for data processing.	The Confed processes data without justification or legal basis	None – Project & dataset is not complex or ambiguous and adequacy, relevance and necessity is self-evident	None	N/A
3B-1	Confed not registered with ICO	Confed faces regulatory action	Confed to register with the ICO	MW has registered and waiting for confirmation from ICO	Complete July19
3B-2	Confed has not completed DSP Toolkit	Confederation lacks member confidence and/or faces regulatory action for not meeting required standards	DSP Toolkit is completed as part of the implementation of the IG Management Framework.	DSP Toolkit completed by 31 st August 2019	
3C-15	Data processing will involve unsolicited SMS Messaging	Complaint re privacy or data breach through poor IG connected with use of SMS Messaging.	Apply an implied consent model for SMS messaging and update the Privacy Notice to inform Service Users. Monitor the use of SMS messaging through quality reporting to the QPF Committee – if a theme emerges in feedback, management action to be taken	Implied consent will be applied to all service users and the privacy notice updated to inform service users of this policy.	31 st July 2019

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For completion by IG:

	Residual Risk	Main Risk Sources	Main Threats	Main Potential Impacts	Main Controls Reducing the Severity and Likelihood	Severity	Likelihood
1	Appropriate controls for user access across the Confederation network infrastructure	Number of Users, lack of consistent organisational protocols across all practices, lack standardised process	Risk of Privacy Breach, risk of malicious activity, risk inappropriate access being granted through poor process	Reputational damage, enforcement action by regulators, loss of partner and commissioner confidence	IG Management Framework including policies, Framework agreement to formalise controls and relationship between Confed & Practices	Moderate (3) – Severity is the same before and after management action to mitigate risk	Inherent – Likely (4) Residual - Unlikely (2)
3							

IG review completed by: Simon Boycott- Head of Development and Governance Review date: 31st July 2019

Date complete and risk assessed: 22nd July 2019

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