Devices Patient Level Contract Monitoring (DePLCM)

Requirements Specification

NHS England and NHS Improvement

Devices Patient Level Contract Monitoring (DePLCM): Requirements Specification

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Prepared by: Martin Hart, NHS England

Ceri Townley, NHS England

This information can be made available in alternative formats, such as easy read or large print, and may be available in alternative languages, upon request. Please contact 0300 311 22 33 or email england.contactus@nhs.net.

Data Coordination Board

This information standard (DCB3002) has been approved for publication by the Department of Health and Social Care under section 250 of the Health and Social Care Act 2012.

Assurance that this information standard meets the requirements of the Act and is appropriate for the use specified in the specification document has been provided by the Data Coordination Board (DCB), a sub-group of the Digital Delivery Board.

This information standard comprises the following documents:

- Requirements Specification
- Implementation Guidance.

An Information Standards Notice (DCB3002 Amd 13/2017) has been issued as a notification of use and implementation timescales. Please read this alongside the documents for the standard.

The controlled versions of these documents can be found on the NHS Digital website. Any copies held outside of that area, in whatever format (e.g. paper, email attachment), are considered to have passed out of control and should be checked for currency and validity.

Date of publication: 29 April 2019

Glossary of terms

Term	Acronym	Definition
Aggregate Contract Monitoring	ACM	Aggregate Contract Monitoring provides a summary of the volume of clinical activity performed by a healthcare provider and associated costs chargeable to the commissioner for that activity. This report serves the contractual requirement for the aggregate finance and activity report, submission of which is required under Schedule 6 of the NHS Standard Contract.
Clinical Commissioning Group	CCG	An organisation responsible for implementing the commissioning roles as set out in the Health and Social Care Act 2012. They are comprised of groups of GP practices that are responsible for commissioning most health and care services for patients.
Commissioning Data Sets	CDS	Commissioning Data Sets (CDS) are maintained and developed by NHS Digital, in accordance with the needs of the NHS and the Department of Health and Social Care. They form the basis of data on activity carried out by organisations reported centrally for monitoring and payment purposes.
Commissioning Support Unit	CSU	An organisation that provides commissioners with external support, specialist skills and knowledge to support them in their role.
Data Landing Portal	DLP	A system, developed by NHS Digital that allows data to be securely transferred between organisations. The system enables Data Services for Commissioners Regional Offices to set up data specifications, against which incoming data from Providers is validated.
Data Services for Commissioners Regional Office	DSCRO	Regional offices staffed by NHS Digital that support the data management needs of commissioners with the provision of appropriate technical and procedural controls and legal basis to store and process personal confidential data.
Information Governance	IG	The set of multi-disciplinary structures, policies, procedures, processes and controls implemented to manage information at an enterprise level, supporting an organisation's immediate and future regulatory, legal, risk, environmental and operational requirements.

Glossary of terms (cont/...)

Term	Acronym	Definition
Information Standards Notice	ISN	A publication that announces new or changes to information standards published under section 250 of the Health and Social Care Act 2012.
Information Technology	IT	The use of any computers, storage, networking and other physical devices, infrastructure and processes to create, process, store, secure and exchange all forms of electronic data.
National Information Board	NIB	A partnership group with membership from organisations across the health and care system.
Patient Level Contract Monitoring	PLCM	Patient Level Contract Monitoring is a means to enable the interchange, in a uniform format, of monthly patient-level contract monitoring data between commissioners and providers of healthcare.
Secondary Uses Service	SUS+	SUS+ is a comprehensive repository for commissioning data sets in England. It is held by NHS Digital and it enables a range of reporting and analyses to support the NHS in the delivery of healthcare services.

Contents

1. Background and context

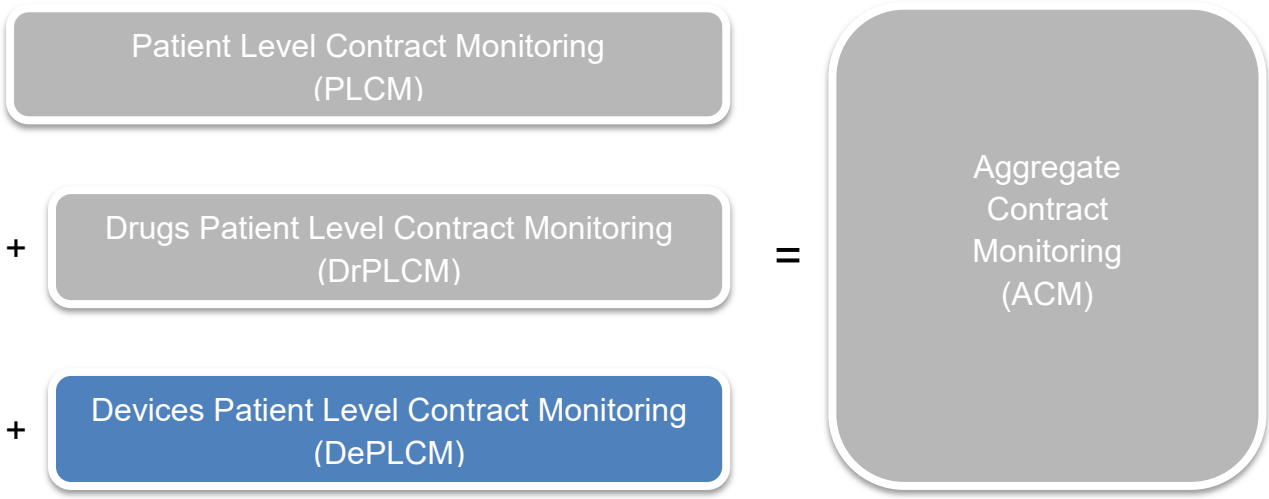
The purpose of the Devices Patient Level Contract Monitoring (DePLCM) Information Standard (hereafter the Standard) is to enable the interchange, in a uniform format, of monthly patient level device contract monitoring data between commissioners and providers of healthcare. This will ensure that device contract monitoring and reporting is consistent and comparable across all commissioning organisations and their footprints.

The Devices Patient Level Contract Monitoring (DePLCM) is a patient level report containing patient identifiers relating to high cost devices. Its purpose is to substantiate and provide further details of the 'DEVICE' aggregate reporting line shown in the Aggregate Contact Monitoring (ACM) and is already widely used by most providers to report NHS England directly-commissioned services.

The ACM is the Activity and Finance Report which each provider is required to submit to its commissioners as a requirement of Schedule 6A of the NHS Standard Contract. It demonstrates the volume of activity and the aggregated cost of commissioned clinical care provided to patients as well as financial adjustments not attributed directly to clinical care. Although the financial reconciliation process in Service Condition 36 of the NHS Standard Contract allows for changes to be made to the amount charged after its submission, the ACM should be a very good indication of the amount a commissioner will be expected to pay for the period and therefore provides a good basis for validation. It is already widely used by most providers to report NHS England directly-commissioned services.

Diagram 1 below shows how the DePLCM relates to the three other contract monitoring data flows, each of which is covered by a separate data standard. The DePLCM **should not** contain any patient level **activity** information.

Diagram 1.



1.1 Relationship to other policies, programmes, projects and services

This new information standard is aligned to the National Information Board's (NIB) Domain H (Data Outcomes for Research and Oversight) and the high-level rationale for modular data. It is designed to collect data more efficiently and includes activities not recorded by Commissioning Data Sets (CDS). This information is essential to the efficient running, planning and development of the NHS and enables data to be analysed in new and different ways for the health and social care system.

1.2 Supporting information

This Standard should be read alongside the following supporting documents or information resources contained within the following websites:

#	Name	Summary
1.	Devices Patient Level Contract Monitoring (DePLCM): Implementation Guidance	Implementation guidance for users of the Standard.
2.	Devices Patient Level Contract Monitoring (DePLCM): User Guidance for Providers and Commissioners	Guidance for users of the Standard including population guidance for individual data elements and answers to a number of frequently asked questions (FAQs).
3.	NHS Data Model and Dictionary v3	Includes definitions for many of the data elements contained within the Standard
4.	NHS Digital Data Landing Portal	Resources and user guides relating to the Data Landing Portal (DLP) – the means by which providers can securely transfer data to Data Services for Commissioners Regional Offices (DSCROs).

2. Purpose and scope

2.1 Users of the Standard

The Devices Patient Level Contract Monitoring (DePLCM) is to be used across the NHS and Independent Sector organisations in England, primarily within commissioning functions. The main users of this are:

- Staff in providers responsible for contracting, finance and business intelligence (informatics);
- National bodies which support the delivery of Health and Social Care such as NHS Digital, NHS Improvement, the Care Quality Commission and Public Health England (PHE);
- NHS England, its commissioning regions and local offices;
- All NHS England direct commissioning functions, clinical commissioning groups (CCGs), Data Services for Commissioners Regional Offices (DSCROs) and organisations providing a commissioning support unit (CSU) service;
- Any other NHS organisations that replace any of the above and take on their functions in future.

2.2 Scope

The scope of the Standard is **all NHS-funded devices not reimbursed through National Tariff prices, as defined in the NHS Improvement National Tariff Payment System High Cost Devices List and any high cost devices not associated with a National Tariff, provided to patients for all NHS commissioners**. The total charged to a commissioner in the Devices Patient Level Contract Monitoring must be equivalent to the aggregate monetary value shown relating to tariff-excluded devices in the ACM for a particular commissioner.

This covers:

- All NHS and independent sector acute providers operating under the full-length version of the NHS Standard Contract – see table below, but not primary care from whom the NHS commissions healthcare;
- All NHS commissioners;
- All NHS-funded devices not reimbursed through National Tariff prices.

The table below is a detailed list of the scope of the Standard for providers.

Provider Type and NHS Standard Contract version	Devices Patient Level Contract Monitoring (DePLCM)
NHS or Independent Sector provider commissioned to provide acute services under the full-length version of the NHS Standard Contract	Mandatory
NHS or Independent Sector provider commissioned to provide mental health services under the full-length version of the NHS Standard Contract	
NHS or Independent Sector provider commissioned to provide community services under the full-length version of the NHS Standard Contract	Recommended (where applicable)
NHS provider commissioned to provide ambulance services under the full-length version of the NHS Standard Contract	
NHS or Independent Sector provider commissioned to provide services of any type under the shorter-form version of the NHS Standard Contract	

2.3 Rationale

Currently, local providers and commissioners can agree amongst themselves the content and format of a contract monitoring data set. For providers this can result in a range of different formats for different commissioners and when multiplied by the number of providers across the country this can become a large number of differing formats.

Where an individual provider is required to generate a different reporting format for each commissioning function it increases the data collation and reporting burden for the provider.

A requirement under the current Schedule 6 of the NHS Standard Contract is the production of an Activity and Finance Report and that *"…this report may also serve as the reconciliation account to be sent by the Provider by the First Reconciliation Date under SC36.28, or under SC36.31)*". Aggregate Contract Monitoring (ACM) submissions can therefore be a means by which the initial monthly financial value claimed by the provider can be validated by the commissioner.

The Devices Patient Level Contract Monitoring is a patient level report, containing patient identifiers. Its purpose is to substantiate and provide detail of the aggregate value in the ACM relating to devices not reimbursed through National Tariff prices as defined by the NHS Improvement National Tariff Payment System High Cost Devices List. It also covers high cost devices not associated with a National Tariff. It contains details relating to the implementation of these devices that are not found in standard CDS flows submitted to SUS+.

In order for a commissioning organisation to have a total view of its high cost devices expenditure, there is a need to aggregate its reporting. In many instances this requires the re-mapping of differing provider returns into a common format, resulting in an additional administrative burden.

2.4 Benefits

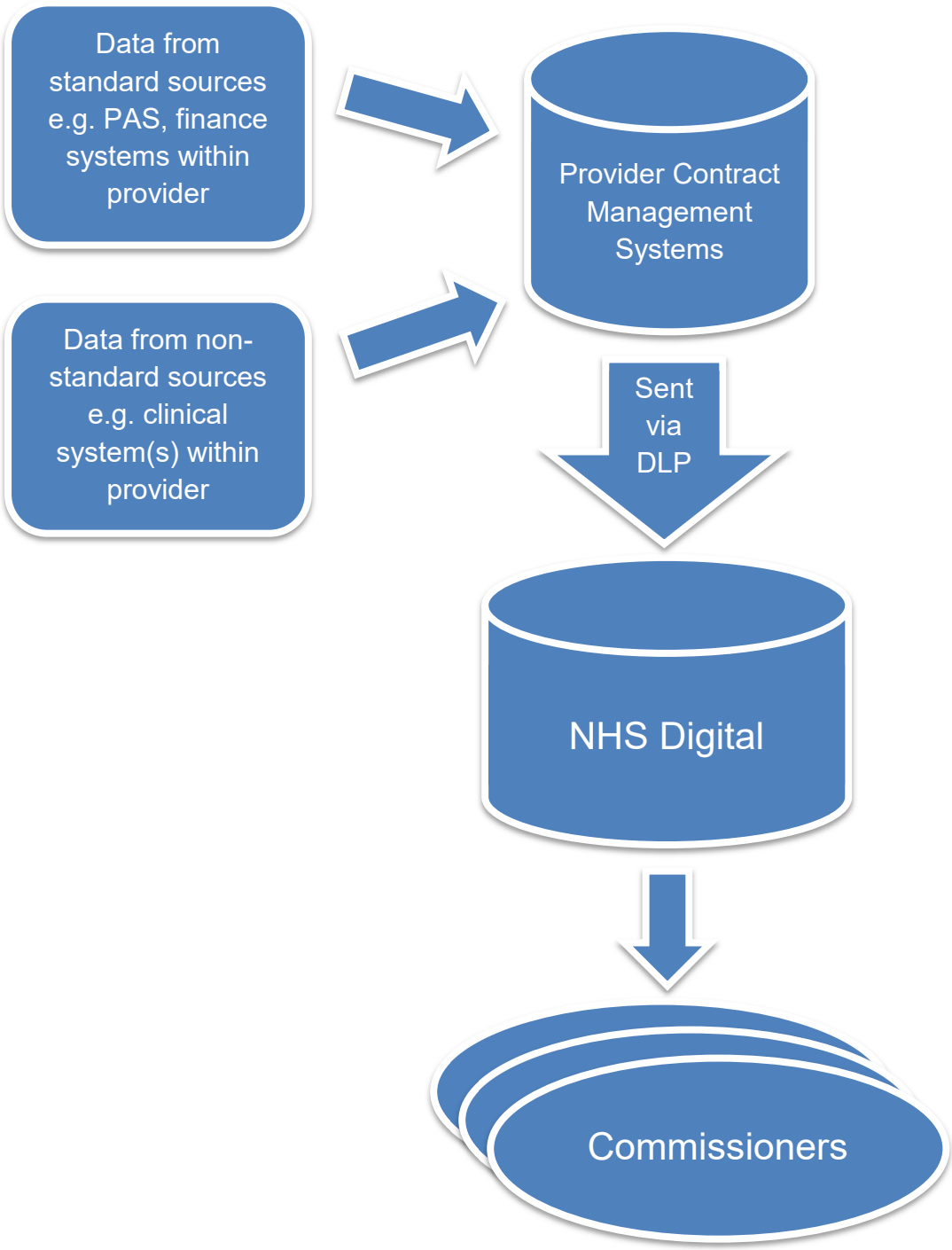
The Standard will ensure that monthly device contract monitoring data that flows from providers to commissioners via NHS Digital will contain a consistent set of data items of sufficient quality. This will:

- Minimise the burden on providers through convergence to a single report format for use by all commissioning functions regardless of organisation;
- Reduce the burden on commissioners and their CSUs through convergence to a single report format from all providers;
- Allow the development of a standard automated reconciliation process for secondary care device implementation and finance which will increase efficiency through removal of manual validation checks;
- Improve year-end forecasting and forecasting against plan of device usage for commissioners;
- Provide greater assurance that the right patients are receiving the right devices at the right place for the correct price;
- Improve the regional and national consistency of reporting of NHS England directly-commissioned services, resulting in national economies of scale.

2.5 High level process

Diagram 2 provides a high-level overview of the data flows associated with the production and submission of the Standard.

Diagram 2.



3. Requirements

3.1 Definitions

The definitions of the key words MUST, SHOULD and MAY are taken from the Internet Engineering Task Force Best Current Practice Document. Other terms used below and elsewhere in this Specification are defined as follows:

Term	What it stands for
Organisations	Organisations required to implement and comply with the Standard, that is: • All NHS and independent sector acute providers operating under the full-length version of the NHS Standard Contract, but not primary care from whom the NHS commissions healthcare; • All NHS commissioners.
Relevant Staff	Employees or contractors of organisations to which the Standard applies who have a contracting, commissioning, performance, finance, business informatics or IT role.
Systems	Any clinical, contracting, financial, administrative or contract management system used in the capture of data for, or in the production of, the Standard.

- All MUST requirements must be met;
- All SHOULD requirements must be met or there be a credible and legitimate reason for why they have not been;
- Any MAY requirements are purely optional.

3.2 General requirements

#	Requirement
Implementing the Standard: Procedures, Systems and Governance	
1.	Organisations MUST prepare for the implementation of the Standard, by assessing their current systems and processes, developing a local implementation plan and the subsequent roll-out of this plan.
2.	Organisations SHOULD refer to and utilise the Implementation Guidance accompanying this Standard to help steer decisions.
3.	Organisations MUST review their current systems used in the production of the Standard to ensure that the necessary data items are held and are in the correct format. In cases where data items used in the Standard are missing IT systems MUST be suitably adapted.
4.	Information governance leads MUST review the information governance implications of implementation of the Standard within their organisation and if necessary plan to implement mitigating actions to address any identified risks such that they are as low as reasonably possible.
Implementing the Standard: Workforce	
5.	Organisations MUST provide, arrange and/or support relevant staff to receive any training which is identified as locally necessary to enable effective implementation of and conformance to the Standard.
Ongoing Compliance with the Standard: Accuracy of Data	
6.	Organisations MUST ensure that data recorded for compliance to the Standard is accurate. Systems to quality assure data MUST be put in place.

#	Conformance Criteria
Implementing the Standard: Procedures, Systems and Governance	
1.	Organisations prepared effectively for implementation of the Standard, assessed their current systems and processes and developed a local roll-out plan. Measurement beginning from the publication of the Information Standards Notice (ISN) in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
2.	Implementation Guidance accompanying this Standard was read and used to inform local decision-making. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
3.	Systems used in the production of the Standard were reviewed and, in cases where data items used in the Standard were missing these systems were suitably adapted, to allow production of the Standard. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the

	implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
4.	Information governance risks associated with implementation of the Standard were identified and mitigating actions completed such that residual risks were as low as reasonably possible. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
Implementing the Standard: Workforce	
5.	Staff competency/training records indicated that relevant staff received any training identified as locally necessary that enabled implementation of and conformance to the Standard. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
Ongoing Compliance with the Standard: Accuracy of Data	
6.	Quality assurance processes were in place to enable verification of the accuracy of data recorded for production of the Standard. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.

3.3 Those with responsibility for data capture and IT solutions

#	Requirement
Overview	
	Those responsible for IT systems used in the capture of data for, or in the production of the Devices Patient Level Contract Monitoring MUST update, change or replace those systems so that they allow conformance to the Standard.
Functionality: Data Items	
1.	Systems MUST enable recording of all data items contained within the Standard, in their specified format. Local systems MAY hold more information than is required by the Standard.
2.	Systems MUST allow for changes to the data items associated with the Standard over time, including the release of new or amended codes.
Functionality: Timeliness	
3.	Systems MUST enable recording of all data items contained within the Standard, in a timely fashion in order to allow production of the Devices Patient Level Contract Monitoring in line with national reporting timetables.

#	Conformance Criteria
Functionality: Data Items	
1.	Systems enabled recording of all data items defined in the Standard and in their specified formats. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
2.	Systems allowed for changes to data items associated with the Standard over time, including the release of new or amended codes (where used by relevant systems). Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
Functionality: Timeliness	
3.	Systems allowed for recording of all data items contained in the Standard, in a timely fashion in order to allow production of the Standard in line with national reporting timetables. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.

3.4 Those with responsibility for the production/submission of the Standard

#	Requirement
Overview	
	Those responsible for contract management reporting and the production of the Devices Patient Level Contract Monitoring MUST ensure that routine submissions are made that conform to the Standard.
Functionality: Data Items	
1.	Systems MUST be populated with data items required by the Standard, in their specified format. Local systems MAY hold more information that is required by the Standard.
2.	Systems MUST allow for changes to the data items associated with the Standard over time, including the release of new or amended codes.
Functionality: Timeliness	
3.	Systems MUST be populated with all data items (where relevant) contained within the Standard, in a timely fashion in order to allow production and submission of the Devices Patient Level Contract Monitoring in line with national contracting/commissioning timetables.

#	Conformance Criteria
Functionality: Data Items	
1.	Systems were populated with data items defined in the Standard and in their specified formats. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
2.	Systems allowed for changes to data items associated with the Standard over time, including the release of new or amended codes (where used by relevant systems). Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.
Functionality: Timeliness	
3.	Systems were populated with all data items (where relevant) contained in the Standard, in a timely fashion in order to allow production and submission of the Standard line with national contracting/commissioning timetables. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.

3.5 Those who are other users of the data

#	Requirement
Overview	
	Relevant staff SHOULD be familiar with the contents of the Devices Patient Level Contract Monitoring and have an understanding of its uses and relevance.
Implementing the Standard: Workforce	
1.	Organisations SHOULD provide, arrange and/or support relevant staff to receive any training and/or awareness programme which is identified as locally necessary to enable effective implementation of and conformance to the Standard.

#	Conformance Criteria
Implementing the Standard: Workforce	
1.	Staff competency/training records indicate that relevant staff have received training identified as locally necessary to enable effective implementation of and conformance to the Standard. Measurement beginning from the publication of the ISN in May 2019: 1. The number of queries received by NHS Digital regarding the implementation of the changes may denote non-conformity. 2. The number of queries/complaints received regarding the Standard will be monitored to inform compliance.

4. When should the Standard be submitted?

The submission of the Standard is an NHS Standard Contract requirement and must be in line with the timescale indicated in the National Requirements Reported Locally section within Schedule 6 of the NHS Standard Contract.

5. How should the Standard be submitted?

All submissions up to the agreed submission date must be on a bulk replacement/update basis i.e. each submission/resubmission will overwrite and replace in full any previous submissions for the same reporting period or periods.

The completed monthly Devices Patient Level Contact Monitoring should be transmitted using the NHS Digital Data Landing Portal (DLP). The DLP allows data to be transferred securely between organisations using a centrally managed system. It also facilitates the standardisation of local data transfers nationally.

Before first submission, users MUST alert their DSCRO so that the necessary loading files for the Standard can be created prior to use.

The DLP currently accepts files in a comma-separated value (CSV) format, or CSV files compressed using the gzip format. It has a maximum allowable file size of 1Gb for uncompressed CSV files (or 100Mb for compressed files). The first row must contain column headers, the names of which must match those in the specification being used when submitting the file. **Spaces used in the field names of the Specification must be replaced by underscores.**

For more detailed guidance on submission of data using the DLP please refer to guidance on the NHS Digital Data Landing Portal (DLP) site. Users should be aware that the DLP interface is accessed using Google Chrome installed with the NHS Digital Chrome Extension or using Internet Explorer 11.

If using Google Chrome please refer to the Google Chrome Installation Guide which can be downloaded from the NHS Digital DLP webpage. The guide provides full instructions on installing Google Chrome and the required NHS Digital Chrome Extension.

6. How should the Standard be completed?

Providers must use a consistent method of completion to populate the Standard with data for each submission/resubmission.

The Standard must be completed in such a manner that it contains data relating to the current reporting month and all previous months, with all previous months shown individually. Each submission must contain data for each of the submission periods prior to the current submission period i.e. the submission relating to devices implemented in June 2019 must contain data for devices implemented in April 2019, May 2019 and June 2019 **all shown separately**.

7. Specific data requirements

The table below defines the detailed data requirements of the Standard, listing each data element and its format. All data elements listed below MUST be included, their completion being determined by the completion criteria (M/R/O) shown in the final column.

For a detailed technical specification of the Standard showing individual data elements and lists of valid codes (where these are not contained within the NHS Data Model and Dictionary v3) please refer to the Devices Patient Level Contract Monitoring (DePLCM): User Guidance for Providers and Commissioners document.

Data Element	Format and Length	Population Guidance	Notes	Mandatory (M), Mandatory Where Relevant (R) or Optional (O)
FINANCIAL MONTH	max an2	1=April, 2= May, 3 June….12=March, with no leading zeros.	The month in which the device was implemented. For non-activity items this should be the financial month to which the charge relates.	M
FINANCIAL YEAR	an6	201920=2019/20, 202021=2020/21 etc. The slash (/) symbol must not be included.	The financial year in which the device was implemented. For non-activity items this should be the financial month to which the charge relates.	M
DATE AND TIME DATA SET CREATED	an19 CCYY-MM-DD hh:mm:ss	Valid date and time format - as shown in the Specification.	The date and time that the file was created prior to its submission to the DLP. This timestamp will be used to ascertain the latest version of the submission.	M

ORGANISATION IDENTIFIER (CODE OF PROVIDER)	min an3 max an6	Valid ODS code – see the NHS Digital ODS Portal for valid codes. NHS Providers must complete this data element with their valid national 3-character Trust code with no trailing zeros (i.e. RNA not RNA00). Non-NHS providers should complete this data element with their valid national full 5-character code. Only where a hospital site is required for specific contract monitoring purposes should NHS providers use a valid national 5-figure code.		M
ORGANISATION IDENTIFIER (GP PRACTICE RESPONSIBILITY)	min an3 max an5	Valid ODS code – see the NHS Digital ODS Portal for valid codes. Where the POINT OF DELIVERY CODE is one of the non-activity codes this should be populated with Q99.	This is the responsible CCG code. The organisation responsible for the GP Practice where the patient is registered, irrespective of whether they reside within the boundary of the Clinical Commissioning Group (CCG).	M
ORGANISATION IDENTIFIER (CODE OF COMMISSIONER)	min an3 max an5	Valid ODS code – see the NHS Digital ODS Portal for valid codes.	The derived commissioner as derived with reference to the NHS England Commissioner Assignment Method (CAM) and hierarchy for assigning NHS England directly-commissioned services.	M
GENERAL MEDICAL PRACTICE CODE (PATIENT REGISTRATION)	an6	Valid ODS code – see the NHS Digital ODS Portal for valid codes. Where the POINT OF DELIVERY CODE is one of the non-activity codes this should be populated with V81998.		M

WITHHELD IDENTITY REASON	an2	Valid code - see the NHS Data Model and Dictionary website for valid codes. To be populated where any of the patient identifiable fields are not provided due to withheld identity reasons and the POINT OF DELIVERY CODE is not one of the non-activity codes. Where the ACTIVITY TREATMENT FUNCTION CODE or SPECIALISED SERVICE CODE indicate activity relating to a sensitive data item e.g. HIV or G-U Medicine no patient identifiable fields should be populated and the appropriate WITHHELD IDENTITY REASON code used.		R
NHS NUMBER	n10	If the NHS NUMBER does not exist the LOCAL PATIENT IDENTIFIER (EXTENDED) must be populated. The population of this data element is not required where the WITHHELD IDENTITY REASON is populated or for records that do not relate to patient level activity.		R
LOCAL PATIENT IDENTIFIER (EXTENDED)	max an20	To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.		R

POSTCODE OF USUAL ADDRESS	max an8	The population of this data element is not required where the WITHHELD IDENTITY REASON is populated or for records that do not relate to patient level activity. Where the POSTCODE OF USUAL ADDRESS is not known, (for example, the patient has no fixed abode, the patient is an overseas visitor etc.) the appropriate ODS pseudo postcode must be used.		R
PERSON BIRTH DATE	an10 CCYY-MM-DD	The population of this data element is not required where the WITHHELD IDENTITY REASON is populated or for records that do not relate to patient level activity.		R
AGE AT ACTIVITY DATE (CONTRACT MONITORING)	max n3	To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.	Age of the patient when device was fitted.	R
PERSON STATED GENDER CODE	an1	Valid code - see the NHS Data Model and Dictionary website for valid codes. To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.		R
HOSPITAL PROVIDER SPELL NUMBER	max an12	To be populated where a device was implemented in an admitted patient care (inpatient) setting and the POINT OF DELIVERY CODE is not one of the non-activity codes.	The format of this data element should be in the same format as that submitted to SUS+ in order to enable accurate linkage between the data sets.	R

ATTENDANCE IDENTIFIER	max an12	To be populated where a device was implemented in a non-admitted patient care (outpatient) setting and the POINT OF DELIVERY CODE is not one of the non-activity codes.	The format of this data element should be in the same format as that submitted to SUS+ in order to enable accurate linkage between the data sets.	R
CLINICAL INTERVENTION DATE (MEDICAL DEVICE IMPLEMENTATION)	an10 CCYY-MM-DD	To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.	Date the device was implemented.	R
HIGH COST TARIFF EXCLUDED DEVICE CODE (SNOMED CT DM+D)	min an6 max an20	Valid SNOMED CT code. To be populated at providers with electronic patient health records where the POINT OF DELIVERY CODE is not one of the non-activity codes.		O - until dm+d enabled system in place, thereafter R
HIGH LEVEL CODE (HIGH COST TARIFF EXCLUDED DEVICE)	max an5	Valid code - as listed in the User Guidance of the Specification. To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.		R
SUBSIDIARY LEVEL CODE (HIGH COST TARIFF EXCLUDED DEVICE)	max an20	Valid code - as listed in the User Guidance of the Specification. To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes and the SUBSIDIARY LEVEL CODE (HIGH COST TARIFF EXCLUDED DEVICE) code exists in the taxonomy.		R
LOCAL CODE (HIGH COST TARIFF EXCLUDED DEVICE)	max an20	Free text for the local code of the device.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O

MEDICAL DEVICE PROCUREMENT ROUTE (HIGH COST TARIFF EXCLUDED DEVICE)	an2	Valid code - as listed in the User Guidance of the Specification. To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.		R
MEDICAL DEVICE MANUFACTURER (HIGH COST TARIFF EXCLUDED DEVICE)	max an50	Free text for the name of the manufacturer of the device.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R
MEDICAL DEVICE NAME (HIGH COST TARIFF EXCLUDED DEVICE)	max an50	Free text for the manufacturer's name of the device.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R
MEDICAL DEVICE SERIAL NUMBER (HIGH COST TARIFF EXCLUDED DEVICE)	max an50	Free text for the serial number of the device.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R
MEDICAL DEVICE SIZE (HIGH COST TARIFF EXCLUDED DEVICE)	max an10	Free text for size of the device where applicable. If no specific size is given or applicable then populate with the word STANDARD.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R
MEDICAL DEVICE QUANTITY (HIGH COST TARIFF EXCLUDED DEVICE)	max n4	To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.		R
PROVIDER REFERENCE NUMBER	max an17	To be populated where the POINT OF DELIVERY CODE is not one of the non-activity codes.	To be used for a specific locally-agreed purpose between a provider and a commissioner e.g. where treatment requires prior approval.	R

COMMISSIONED SERVICE CATEGORY CODE	an2	Valid code - as listed in the User Guidance of the Specification.	This should be derived with reference to the NHS England Commissioner Assignment Method (CAM) and hierarchy for assigning NHS England directly-commissioned services.	M
SPECIALISED SERVICE CODE	max an12	Valid code - as listed in the User Guidance of the Specification. This is only required where NHS England - Specialised Services is the commissioner.		R
POINT OF DELIVERY CODE	max an10	Valid code - as listed in the User Guidance of the Specification.	DEVICE for each patient-level device recorded, otherwise BLOCK, ADJUSTMENT or NAOTHER. No other POINT OF DELIVERY CODE must be used.	M
POINT OF DELIVERY FURTHER DETAIL CODE	max an10	This data element must be populated where the point of delivery patient type taxonomy is starred (**), indicating that it requires more detail. Where the POINT OF DELIVERY CODE is indicated as requiring the completion of the POINT OF DELIVERY FURTHER DETAIL CODE data element it is suggested that providers show the local code used for this service.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R

POINT OF DELIVERY FURTHER DETAIL DESCRIPTION	max an100	This data element must be populated where the point of delivery patient type taxonomy is starred (**), indicating that it requires more detail. Where the POINT OF DELIVERY CODE is indicated as requiring the completion of the POINT OF DELIVERY FURTHER DETAIL DESCRIPTION data element it is suggested that providers show the local description and/or measure used for this service. It is advised that where providers and commissioners need to capture structured data in the POINT OF DELIVERY FURTHER DETAIL DESCRIPTION data element, a delimiter is used by local agreement. Where it may add value e.g. where the POINT OF DELIVERY CODE is CQUIN, the POINT OF DELIVERY FURTHER DETAIL DESCRIPTION data element may be used to provide further information.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	R

CONTRACT MONITORING ADDITIONAL DETAIL (FIRST)	max an50	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DESCRIPTION (FIRST)	max an100	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DETAIL (SECOND)	max an50	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DESCRIPTION (SECOND)	max an100	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DETAIL (THIRD)	max an50	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DESCRIPTION (THIRD)	max an100	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DETAIL (FOURTH)	max an50	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O

CONTRACT MONITORING ADDITIONAL DESCRIPTION (FOURTH)	max an100	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DETAIL (FIFTH)	max an50	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
CONTRACT MONITORING ADDITIONAL DESCRIPTION (FIFTH)	max an100	Very general purpose (VGP) data element. The population of this data element can be used for differing purposes by local agreement.	**No Patient Identifiable Data (PID) or sensitive data should be recorded in this data element.**	O
UNIT PRICE (SUPPLIER)	max n18.max n8	Unit price paid to a supplier excluding VAT.		M
UNIT PRICE (COMMISSIONER)	max n18.max n8	Unit price charged to a commissioner excluding VAT, including agreed on costs.		M
VALUE ADDED TAX CHARGED INDICATOR (CONTRACT MONITORING)	an1	Y=VAT charged, N=VAT not charged or exempt.		M

TOTAL COST	max n18.max n8	Where the POINT OF DELIVERY CODE is not one of the non-activity codes this is UNIT PRICE (COMMISSIONER) * MEDICAL DEVICE QUANTITY (HIGH COST TARIFF EXCLUDED DEVICE). In cases where VALUE ADDED TAX CHARGED INDICATOR (CONTRACT MONITORING) is charged (Y) VAT should also be added to this amount.		M