

Health Records Policy

Version number:	2.5
Consultation Groups	Information Governance Steering Group
Approved by (Sponsor Group)	Information Governance Steering Group
Ratified by:	Quality Committee
Date ratified:	September 2019
Name of originator/author:	Information Governance Manager
Executive Director lead :	Director of Planning and Performance
Implementation Date :	August 2019
Last Review Date	July 2019
Next Review Date :	July 2022

Services	Applicable
Trust wide	√

Version Control

Version	Date	Author	Status	Comment
1.0	October 2011	Head of Information Governance	Final	Condenses previous versions of the policies and procedures below into one document: Clinical Records Management Procedures, Health Record Keeping Policy, Procedure for Health Record Archiving and Destruction
1.1	January 2012	Head of Information Governance	Final	Incorporates NHSLA Level 3 requirements
1.2	March 2012	Head of Information Governance	Final	Incorporates revised NHSLA Level 3 requirements published end January 2012
1.3	May 2012	Head of Information Governance	Final	Sets definitive timescales for entering information on to electronic record keeping systems
1.4	July 2012	Head of Information Governance Clinical Records Development Manager	Final	Revised to reflect access / tracer card practice in locations without a dedicated Records Manager
1.5	August 2012	Head of Information Governance	Final	Includes updated guidance on filing incident reports. Outlines clinical responsibility for ensuring new information about inpatients is seen and acted on
1.6	November 2012	Clinical Records Development Manager	Final	Strengthened guidelines to address queries related to errors identified in the entries made in clinical records. Clarified guidance on electronic health records.
1.7	January 2014	Head of Information Governance	Final	Revised guidance to transfer records from CAMHS to adult services

2.0	January 2016	Associate director of Assurance	Final	Revised as the primary health record is now electronic. Incorporates previous Clinical RiO policy
2.1	February 2016	Head of Information Governance	Final	Revised validations section following IGSG/IMTAC review
2.2	March 2017	Interim Head of Information Governance	Final	Addition of defining paragraph on responsibilities for the validation of collaborative entries to Validation Section 6
2.3	March 2017	Interim Head of Information Governance	Final	Added Godard Enquiry hold on deletion of all records until further notice
2.4	October 2018	Information Rights Manager	Final	Revised to comply with GDPR
2.5	July 2019	Information Governance Manager	Final	Include appendix about Arts Therapies and Identification Criteria for Unknown Patients. DPO role made explicit

Contents

1.	Introduction	7
2.	Purpose	7
3.	Duties	7
4.	Clinicians	8
5.	Validation	8
6.	Local Records Managers	8
7.	Duty Senior Nurse	9
8.	Data Controller	9
9.	Data Protection Officer	9
10.	Caldicott Guardian	9
11.	Service Directors as Information Asset Owners	9
12.	Information Asset Administrators	9
13.	General Principles	9
13.1	Health Records Use	9
13.2	Health Records Standards	10
13.3	Confidentiality	10
13.4	Format	10
13.5	Records Retention	11
13.6	Mental Health Act Documentation	12
13.7	Incident Forms	13
13.8	Complaints Records	13
13.9	Child Health Records - Transfer of Records from CAMHS to AMHS	13
13.10	Research and Teaching	13
13.11	Service Closure or Transfer	14
13.12	Criteria for safer temporary identification of unknown patients	14
14.	Electronic Records	14

14.1	Principles	14
14.2	Temporary Paper Folder Actions	16
15	Paper Records	17
15.1	Principles	17
15.2	Tracking Procedure	17
15.3	Filing Area Standards	17
15.4	Paper Records in Transit	18
15.5	Paper Records Review	18
	Appendix: Arts Therapies	20
	Appendix: Identification Criteria for Unknown Patients	22

1. Introduction

High standards of record keeping underpin the delivery of high quality, evidence based healthcare. Records should be up to date, accurate and accessible when required. This policy provides a framework for achieving high quality safe record keeping based on the principle that records are electronic.

2. Purpose

A health records can be summarised as ‘one which relates to the physical or mental health of an individual, made by or on behalf of a health professional in connection with the case of that individual’. Thus with the exception of the anonymised information most if not all NHS information concerning patients whether held electronically or on paper will fall within the scope of the General Data Protection Regulation / Data Protection Act 2018.

This policy sets out the standards and processes required for maintaining high quality health record keeping standards for all service users.

The electronic record has replaced the paper record as the primary record for both community and mental health service users unless a service does not yet have access to an electronic clinical system.

3. Duties

All individuals

All individuals must ensure confidentiality, integrity, accuracy and appropriate availability of records. All individuals are personally responsible for the records they create or use and will not be allowed access to RiO or any other electronic clinical system until they have completed training at this Trust.

All NHS bodies and those carrying out functions on behalf of the NHS have a common law duty of confidentiality. Everyone accessing the health information of service users has a common law duty of confidentiality to both the service user and the Trust. Confidentiality extends even after the death of a service user.

Individuals should be aware that information recorded in a service user’s record is legally binding. The record should therefore be non-judgmental and contain only information the individual would be willing to be accountable for in Court.

Any notes about a service user form part of the clinical record and should be uploaded to their clinical record e.g. during ward rounds, during CPA meetings, where Duty of Candour is invoked etc.

All teams should have an internal process for ensuring the relevant clinician / care worker is aware of all new documentation / correspondence relating to a service user.

All individuals should be familiar with their patient system crib sheets which guide the user through particular tasks.

Individuals should never share Smartcards. If information is accessed / added on behalf of another individual this should be made clear within the record. If an individual accesses the electronic record of a patient not under their direct care they will be prompted to state why they are accessing this information. This will be recorded and audited. Access without a legitimate reason may result in disciplinary action or dismissal and legal proceedings.

Line managers

All ward, team or service managers, supported by the Associate Director of Information Governance, are responsible for ensuring the principles set out in this policy are followed together with any local supporting procedures.

All line managers and supervisors must ensure that their staff, whether administrative or clinical, are adequately trained in the management of health records and apply the appropriate guidelines and crib sheet procedures.

Line managers will be responsible for authorising access to RiO and other systems requiring Smartcards through the Registration Authority (RA) who will issue smartcards on receipt of that authorisation.

4. Clinicians

Clinicians must ensure they adhere to the records management standards of their professional body in addition to the requirements of this policy.

5. Validation

All qualified staff are able to validate their own records. The only exceptions to this are those staff with identified capability issues or students whose need for validation should be highlighted on the access application by line managers/supervisors.

There is an expectation that where a staff member enters a patient record of contact involving multiple staff, that all the staff involved check the record made and ensure the accuracy of it and where needed make corroborative entries or corrections.

6. Local Records Managers

Each directorate will have a local Records Manager under whose guidance designated staff will:

For paper records:

- Maintain safe and structured records management stores
- Ensure there is a process for the retrieval of case notes and update tracer cards.
- Organise the safe transportation of records from one location to another
- Deal with requests for access to health records within statutory time limits. In some services there is designated access to records leads. This is covered in detail in the Trust's Access to Records policy.

7. Duty Senior Nurse

Out of hours, the Duty Senior Nurse / Manager in charge is the only individual permitted to access a paper records store to retrieve case notes and update tracer cards. Out of hours, the Duty Senior Nurse / Manager in charge must enter inpatient events on to the relevant electronic clinical recording system within four hours of the occurrence of the event.

8. Data Controller

The Chief Executive is the Trust's Data Controller and has an overarching duty to make arrangements for the safety and integrity of the Trust's health records.

9. Data Protection Officer

The Associate Director of Information Governance is the Trust's registered Data Protection Officer and has responsibility for ensuring health records meet statutory requirements. A Data Protection Officer is a legal requirement under Article 37 of the General Data Protection Regulation. The Data Protection Officer monitors internal compliance with data protection matters, provides advice and information on data protection obligations, acts as a contact point for data subjects and the Information Commissioner's Office. The Data Protection Officer is independent and has direct communication with the Board.

10. Caldicott Guardian

The Medical Director is the Caldicott Guardian and has responsibility for matters relating to patient confidentiality.

11. Service Directors as Information Asset Owners

Service Directors are Information Asset Owners for the health records within their service and must ensure the quality, integrity and security of records in their directorates.

12. Information Asset Administrators

IAs support the IAOs by managing information assets on a day to day basis.

13. General principles

13.1 Health Records Use

Health records must be reliable, accurate and timely to support:

- Continuity of care
- Multi-disciplinary working
- Defence in cases of litigation or complaints
- Evidence based clinical practice
- Administrative and managerial decision making
- Legal requirements including requests for access to records

- Clinical audit and effectiveness
- Statutory and contractual reporting requirements

13.2 Health Records Standards

Health records must:

- Be factual, consistent and be written according to Trust policy and accepted professional standards
- Contain as a minimum the service user's name, date of birth and NHS number. Where the NHS number is not known it should be verified via SCR / PDS. To ensure continuing accuracy of records the service user's full name, address and key contact details (including GP) should be verified at each contact and differing records synchronised.
- Completed as soon as possible after the event. Services should check local CQUINS / KPIs adhering to any shorter timescales where required. In particular, Trust timescales are:
 - Referrals within 24 hours of receipt
 - Outpatient / community entries within 24 hours of event
 - Inpatient entries within four hours of occurrence e.g. admission, discharge, transfer, AWOL, leave etc.
 - Physical health forms within 24 hours

13.3 Confidentiality

Information in a service user's record must be held in complete confidence and only viewed by those directly involved in the provision of care or who are otherwise authorised by the Trust to do so. Unauthorised disclosure and misuse of information constitutes a breach of confidentiality and breach of this policy which may lead to disciplinary action, dismissal and legal action.

Information contained in a service user's record should only be accessed on a 'need to know' basis. Individuals are specifically not permitted to access the records of people they know via a work, social or family connection or in response to media interest.

The Trust will undertake regular and targeted audits to identify instances where confidentiality is breached and take action against any individual found to have inappropriately accessed a record.

Where copies of records are taken outside the Trust (e.g. to visit a service user) they should not be opened / looked at whilst on public transport, left in a visible location (e.g. the back seat of a car) & stored securely whilst not in use.

13.4 Format

Health records will be in a number of formats including emails, x-rays, photographs, audio visual recordings. The same principles apply to all formats. Where paper copies are received note that the primary health record in the Trust is electronic. Paper copies should therefore routinely be scanned and held electronically unless the format prevents this.

13.5 Records Retention

Records in any media (electronic or paper) must be retained according to the Trust's [Record Retention and Disposal Schedule](#). This is based on the NHS Records Management Code of Practice.

Inactive records that have reached the minimum retention period should be reviewed annually to identify the need for extended retention or destruction. After review the list should be updated to show those identified for destruction and those identified for extended retention. Paper records identified for destruction should be destroyed by the relevant Trust approved company and a destruction certificate must be obtained from the company.

Audit/check list – shredding/destruction companies

- 1 – Are there secure receptacles or bins where members of staff can place confidential documents to be shredded/destroyed?
- 2 – How is security of these receptacles or bins ensured so that documents cannot be accessed?
- 3 – How is the material in the bins collected?
- 4 – Where are they shredded / destroyed?
- 5 – Is there a risk that paper may escape from the receptacles or bins between these being collected and the actual shredding/destruction operation?
- 6 – What type of shredding/destruction is used? Is it enough to ensure that information cannot be put back together?
- 7 – What documents does the company have to show compliance with security best practices?
- 8 – Are members of staff dully checked before they start working for the company?
- 9 – What is done with the shredded/destroyed material?

Goddard Enquiry

This Independent Inquiry into Child Sexual Abuse was set up because of serious concerns that some organisations had failed and were continuing to fail to protect children from sexual abuse.

Under Section 21 of the Inquiries Act 2005 the Inquiry has the power to order the production of documents. Failure to comply with such an order without reasonable excuse is an offence punishable by imprisonment (Section 35 of the Inquiries Act 2005). It is also an offence for a person, during the course of an Inquiry, to destroy, alter or tamper with evidence that may be relevant to an Inquiry, or deliberately to do an act with the intention of suppressing evidence or preventing it being disclosed to the Inquiry. Trusts therefore have an obligation to preserve records for the Inquiry for as long as necessary to assist the Inquiry. Prolonged retention of personal data by an organisation at the request of the Inquiry would not therefore contravene data protection legislation, provided such information is restricted to that necessary to fulfil any potential legal duties that organisation may have in relation to the Inquiry. An organisation may have to account for its previous activities to the Inquiry so retention of the data will be regarded as necessary for this purpose. The obligation to the Inquiry to retain documents will remain throughout its duration.

Trusts may also incur separate legal obligations to retain documents during the course of the Inquiry, for example in relation to other legal proceedings.

Services should therefore retain any health records that may be relevant.

Confidential waste

All confidential waste must be securely destroyed. Waste material should be sent to the designated confidential destruction site or shredded. All waste paper baskets must be properly labelled to read 'No Confidential Waste'. In the absence of a shredder, confidential waste must be kept in a 'Confidential Bag' which should be kept in secure location and send to the confidential destruction site. Many services in the Trust will have locked bins from companies authorised to shred confidential information. The bins will be collected by these companies which will securely shred/destroy the bins contents.

Information sharing

Generally, the following principles apply:

- The Trust does not need to seek consent to share information for health purposes with health and social care agencies. Where RiO is used, the Additional Personal Information form should be completed with contact preferences and preferences for sharing information where there is a choice (family, school, voluntary agencies etc.). The form should be updated at regular intervals. Service users should also be given a copy of the "Your records and you" leaflet that explains how their information is processed, and advised that in some circumstances it may not be possible to care for them if their information is not shared.
- Service user information may not be passed on to others without the service user's consent except for healthcare purposes, or when there is a legal requirement, including where there is a risk to the safety of the individual or others.
- Explicit consent is required if identifiable service user information is used in any publication.
- Where identifiable information is disclosed a record of the disclosure and the reasons for doing so should be recorded in the notes. If a decision not to disclose is made, this should similarly be recorded.
- In circumstances where information is shared without the consent of the service user the responsible clinician should make this decision. The Information Rights Manager, Associate Director of Information Governance or Caldicott Guardian can also advise.
- When a child or vulnerable adult is believed to be at risk then relevant information should be shared without delay. This requirement will always override all other confidentiality considerations. Conversely, if it is believed disclosure may compromise an individual's safety, or that it is not in the public interest to disclose, then the information should not be shared. Reasons must always be documented in the record.

13.6 Mental Health Act Documentation

There are specific requirements regarding the receipt and scrutiny of documents under the Mental Health Act 1983:

- Receipt of section papers - only Mental Health Lawstaff and clinical staff at Band 5 or above

(or equivalent) who have at least one year's experience at that level and have attended the relevant Trust training are authorised to formally receive section papers and scrutinise them

- Leave periods - the granting of leave and the conditions attached to it should be recorded in the notes and on the Trust's section 17 form. Copies of the form should be given to the service user, any appropriate relatives or friends and any professionals in the community who need to know
- The service user's legal status in respect of their detention, leave and consent to treatment
- should be immediately apparent from the medical notes

13.7 Incident Forms

- Where an incident relates to a service user the incident form must be added to the health record or as a minimum the Datix reference logged in the health record.
- Where incidents relate to more than one service user they may be added / logged in each health record provided person identifiable information relating to the other service user(s) is redacted. Otherwise use the Datix reference number filed in each service user's progress notes together with a summary of the incident.
- Any Duty of Candour correspondence or conversation should routinely be uploaded to the clinical system

13.8 Complaints Records

Complaints records must be filed separately from the service user's health record unless there is a need to record the complaint (for example where it is directly relevant to the service user's health and failure to recognise this when caring for the service user could have a detrimental effect on the health and well-being of that individual).

13.9 Child Health Records

Transfer of records from CAM HS to adult mental health services

For service users admitted to an adolescent inpatient service within the Trust or elsewhere (e.g. the private sector), who are approaching 18 years of age and are known to community CAMHS, it will usually be the responsibility of community CAMHS to initiate the transfer process. All CAMHS records are now electronic. Detailed information is available in the [CAMHS Transition policy](#)

13.10 Research and Teaching

The Trust has teaching hospital status for students in a number of disciplines. All students are bound by the principles of confidentiality. The following standards apply:

- Service user records may be used for teaching purposes and clinical supervision within the clinical area
- If records are to be used for teaching purposes outside of the clinical area then service user details must be made completely anonymous.
- The principles of access and confidentiality remain the same and the right of the patient to refuse access to their records should be respected and documented in their notes

- The Local Research Ethics Committee must approve the use of service user records for research
- Any use of service users' records for research purposes must comply with the Department of Health's Research Governance Framework

13.11 Service Closure or Transfer

- If a service or site is closed, split or amalgamated with or from another service or Trust, the service manager should advise the Associate Director of Information Governance who will coordinate work with the electronic clinical records team, access to records leads / local Records managers and Estates to ensure appropriate transfer, access, storage and retention of the relevant records.
- In some circumstances, an information sharing agreement or SLA may be required. In such cases the Associate Director of Information Governance should be consulted.

13.12 Criteria for Safer Temporary Identification of Unknown Patients – see appendix 2.

14. Electronic Records

The Trust's primary record keeping systems are electronic. Duplicate paper records systems must not therefore be used.

14.1 Principles

The following principles apply:

- Only Trust approved scanners will be used for scanning documents
- All information about a service user received electronically will be uploaded to the clinical system
- All information about a service user received in paper format will be scanned, subsequently uploaded to the clinical system and the original securely destroyed
- All services will set up team based shared folders on the K drive. Folders and files / documents will be locked down to those individuals with a specific need to access the information. Standardised naming conventions will be used. Folders will be used only for drafting, processing and auditing information prior to uploading to the clinical system and will not become a secondary clinical system.
- When the clinical system is unavailable paper records should be scanned and temporarily held in the team folder on the K drive. Electronic records should similarly be temporarily held on the K drive rather than individual mailboxes or personal drives. When the clinical system becomes available, priority should be given to uploading documents.
- The electronic clinical system record is the primary record. No other current records will be kept other than a small temporary folder used by some services. All original paper and electronic information will therefore be deleted once the scanned copy on the electronic clinical system has been verified as attaining the same standard as the originating copy. This is to prevent duplication of systems and information and the potential for information to be missed, incorrectly added to or otherwise inappropriately processed.

- Teams will set up a systematic process for alerting all members of the treating team to newly received and uploaded information including incoming electronic and paper documents
- Where teams or individuals outside the treating team add information to a clinical system they will routinely alert the treating team to raise awareness of any immediate or significant issues
- Clinicians will routinely check the clinical system for newly received information prior to an appointment / intervention with a service user
- Correspondence must not be filed in the clinical system unsigned. Typed correspondence must always contain an electronic signature. Where the clinician prefers to delegate authority to an administrator for adding the signature, this should be confirmed by email and the email also retained in the clinical system
- Electronic records are subject to the same retention and deletion periods as paper records (i.e. the retention period does not alter simply because the format is electronic rather than paper)
- Electronic records are subject to regular audit including record keeping standards and legitimate relationship access to records. This may include targeted audits
- Records pre-dating clinical system implementation will be retained in paper format and will not currently be uploaded to the clinical system except in instances where the Responsible Clinician believes there is distinct value in doing so
- Where records are recalled from archiving for subject access request purposes these may be attached to the clinical system and the original destroyed in accordance with the guidelines set out for deletion of records
- Requests for access to records will be centrally managed by the Access to Records leads.
 - Where solicitors require access to the service user's electronic record this should be printed out and filtered for third party information. This should comply with the one month timescales referred to within the GDPR.
- Services should always access Reporting Services for contemporaneous clinical information when RiO is unavailable
- The electronic record is the primary record. No other current records will be kept other than the temporary folder outlined below. Original documents will be deleted once the scanned copy has been verified as attaining the same standard as the original document. This is to prevent duplication of systems and information and the potential for information to be missed, incorrectly added to or otherwise inappropriately processed

14.2 Temporary Paper Folder Actions

Document type	Retain in paper format once episode of care complete?	Action on completion of episode of care
'This is me' first person care plan	No	Stamp and upload to clinical system
Prescription / depot charts	Yes	
Observation forms	Yes	Also upload to clinical system
Outcome measures & other questionnaires	No	Upload to clinical system
Written copy of CPA form with signatures	Yes, temporarily until next review	Upload to clinical system
Sticky labels with patient details (for ordering investigations)	No	
Registration form printout	Yes	
Current care plan	No	Upload to clinical system
Current risk assessment	No	Upload to clinical system
Patient property receipts	No	Upload to clinical system
MHA documents (excluding police documentation which should be uploaded to RiO)	Yes	
Alerts with an immediate impact on patient care	Yes	Also upload to clinical system
Do not resuscitate forms	Yes	Also upload to clinical system
Results / investigations / bloods	No	Upload to clinical system
Audio visual recordings	No	Transcribe and upload to clinical system
Artwork / non standard scan size documents	Check if patient wants	Take photo and upload to clinical system
Process notes	No	Destroy. Do not scan / upload unless Therapist advises should be kept
Seclusion documents	No	Scan and upload to clinical system
ECGs	Yes. Store in the resealable wallets provided specifically for this purpose	
Growth charts	No	Scan and upload to clinical system. Download back to hardcopy if patient presents again.

For services using RiO the processes on the RiO user guide pages on the intranet must be

followed at all times

15. Paper Records

15.1 Principles

Although the primary record is electronic, the Trust still holds historic paper records. The following principles apply:

Paper records must be stored in an area where service users, members of the public and unauthorised staff are unable to gain access to them

- Requests for records held in an external repository (e.g. Iron Mountain) must be made via the Access to Records Leads / local Records Managers.
- Trust individuals are only allowed access to records stores during normal working hours with the permission of the relevant local Records Manager / Access to Records Leads and on production of valid identification. Out of Hours, the only access is via the Duty Senior Nurse / Manager in charge.
- Where records are stored in areas that do not have a 24/7 staff presence then they must be secured in an area that is securely locked when the premises are unstaffed.
- Missing or part missing health records must be reported via the Trust's incident reporting process and as a result will be investigated.

15.2 Tracking Procedure

- Tracer cards should be used to document the movement of paper records unless there is an electronic tracking system.
- Only medical records staff or the Duty Senior Nurse / Manager in charge are authorised to retrieve records and update tracer cards. In smaller centres where there are no dedicated medical records staff, the service manager will assign responsibility to relevant staff members to ensure compliance with the record tracking procedure.

15.3 Filing Area Standards

The following standards apply at all times:

- Place files on the shelf in the correct numerical or alphabetical order with the spine underneath and the case note number facing outwards
- Leave the tracer card in situ at all times. Complete movements and mark RETURNED when case notes are filed back. Do not add any other documentation
- The file storage area should be regularly audited utilising the above standards
- Separate records over three inches thick into separate volumes
- Do not eat, drink or smoke near records

15.4 Paper Records in Transit

If paper records need to be transported from one clinical area to another it is the personal responsibility of the individual transporting the records to ensure their safety and security whilst in transit and ensure they cannot be accessed by an unauthorised individual at any time during transit. The following standards apply:

- Records should be handled carefully - never thrown, transported with materials that could cause risk to the records (e.g. chemicals) or exposed to weather, excessive light or risk of theft
- Records must not be transferred internally using the internal mail service as this service does not meet Trust confidentiality standards. They should be transferred using the internal courier service (man with a van / man with a car service)
- An approved courier must be used when transferring health records externally (normally TNT, Loomis or Royal Mail Special Delivery) using 'track and trace' including the provision of a signature on delivery
- Couriers transporting records for archiving / off site storage purposes must meet information governance standards
- Large quantities of health records should be boxed in suitable crates or boxes that give adequate protection. Otherwise, tamper proof envelopes or padded envelopes should be used
- All packaging should be clearly marked with the recipient's name and address. The Trust's
- PO Box Return to Sender address must be included on the reverse of the envelope
- All packages should be marked 'Private and Confidential. To be opened by the addressee only'
- Records in transit must be stored securely e.g. in the locked boot of a car rather than the back seat
- Where practical records should not be left unattended in a vehicle
- If it is absolutely necessary to retain records overnight in no circumstances should they be left in a vehicle. They must be kept under the same conditions of security as on Trust premises i.e. in a locked cupboard within a locked building
- Records should not be looked at in a public place

15.5 Paper Records Review

Inactive records that have reached the minimum retention period should be reviewed annually to identify the need for extended retention or destruction.

- A sticky year label should be affixed in the 'year label' box on the folder front cover at the service user's first contact with the service
- The sticker should be updated if the service user has subsequent contacts. The last contact is the date used to determine retention or destruction

- Deceased patients' folders should be marked 'Deceased' with marker pen on the front cover of the folder
- An electronic clinical system search should be undertaken annually to identify inactive and deceased patients' records. The search should be verified with the paper records before any decision regarding destruction is taken. In the same way all information about a service user should be drawn together to assist the review
- Records that have been identified as potentially reaching the end of their retention period should be listed by the local Records Manager to include:
 - Name
 - Date of birth
 - NHS number
 - Address
 - Last contact date
 - Record status – inactive or deceased
- After review the list should be updated to show those identified for destruction and those identified for extended retention (e.g. research or legal value, historical retention at the Public Records Office, identified familial illness). This should be approved by a senior clinician and documented in the record and a 'Do Not Destroy' sticker attached to the record.
- The services' local clinical governance committee will agree destruction / extended retention. Prior to destruction, this should be agreed by the Information Governance Steering Group
- On notification of final approval from IGSG, record destruction should be undertaken in a secure manner. Approved contractors should be used to destroy records under secure conditions. A certificate of destruction must be obtained and permanently kept with the locality records manager

Appendix 1: Arts Therapies

Art, drama, dance, and music therapies' record keeping

This section applies to the creative psychological therapies: art therapy, drama therapy; dance movement therapy and music therapy. It might also apply where similar techniques are used in clinical practice by clinical staff who are not registered therapists registered in these fields. Basic record keeping is integrated into clinical practice and is embedded in pre-registration training. However, the Records Management Code of Practice tends to assume that the narrative in records will be added by clinicians based on accounts of interactions with service users, and whilst this is largely true for the creative psychological therapies, the rich diversity of media and approach yields a different output that needs consideration of the context and purpose that will help clinicians decide what material might be useful to record in the patient's record.¹

Principles guiding practice

1. Ultimately the ownership of the material remains with the service user, as does the manner of its use and disposal.
2. It is a clinical decision as to what should be recorded in the clinical records, subject to the practical limitations of the system. Where it is not possible to record material that is deemed clinically important, local arrangements should only be made in conjunction with Clinical Systems and Information Governance colleagues.
3. Material may be used confidentially by therapists and trainees for clinical supervision of therapists and therapists in training.

In practice

1. Members must record the service user's attendance for therapy
2. Material produced during the therapy session should be named, dated, and ideally safely stored throughout the therapeutic relationship within the therapeutic space.²
3. Material will usually be stored for the duration of the clinical engagement; however, therapists may consider that certain material is of such significance that it needs to be kept so it can be referred to in future, with explicit consent sought from service users.
4. In general, the disposal of material not to be kept should be negotiated with the service user.
5. Therapists will decide what information, in addition to attendance, will be recorded in the Trust's designated clinical record system. Typically this will be a short narrative on the experience, but may include visual material. Any such material must be scanned into PDF format.

Reproduction, exhibition, and removal by the therapist or service user of service user's material

1. Always obtain permission before publishing or using a service user's material publically whether or not such use of the material is to a public audience, or to a limited audience or forum comprised of fellow health professionals from the extended multi-disciplinary team for therapeutic discussion or training purposes.
2. The effects of material to be seen or heard publically must be considered clinically in order that the sequelae of any display or performance can be sensitively explored. Requests by service users to take material home should be considered on a similar basis. There may be cases where the therapist will need to explain the risks to service users of taking material away. Ultimately the decision is the service user's as owner of the material, unless the action of taking material away could put the service user, or any other person, at risk, e.g. where there are safeguarding issues to consider.

3. Permission for reproduction or exhibition or performance must be obtained following a discussion on informed consent:
 - (i) obtain this in a written agreement signed by the service user and scan this into the main record
 - (ii) include about how the material will be used and whether it will be returned
 - (iii) whenever possible, seek service users' views, feedback or participation in the event or activity at all stages, which may require seeking written signed consent more than once
4. Therapists must never seek to profit financially from the sale of artistic expressions and material produced in the therapeutic relationship.

Further information and support

Contact your line manager for guidance on Trust procedures, or your clinical supervisor in applying guidance to clinical situations. For technical advice on using electronic systems, log a call with the IT Helpdesk in the first instance. For advice about data protection contact the IG team via elft.information.governance@nhs.net

Notes

¹ 'Material' pertains to the original visual, musical, physical and non-verbal artistic expressions created by the client during therapy and within the therapeutic relationship.

² Storage of material includes original hard copy (e.g. 2D and 3D artworks) and any digital recording or reproduction made of client's original visual, musical, physical and non-verbal artistic expressions during therapy and within the therapeutic relationship (e.g. photographs, tape recordings, videos, DVDs, etc.)

Appendix 2 – Safer temporary identification criteria for unknown or unidentified patients

Emergency departments including Mental Health often care for patients unable or unwilling to give their identity including people who are unconscious or who have a critical illness, people with a mental health condition or delirium, and people affected by drink or drugs. Several unidentified patients may arrive together after an accident, or in mass casualty situations. Giving a unique identity to each unknown patient ensures safe and prompt diagnostic testing and treatment. For example, it helps prevent allocating the wrong results to the wrong patient.

Temporary identification (ID) systems can have high potential for error if they use:

- the same or similar names, eg unknown male, unknown female
- pre-allocated numbers that differ sequentially by one digit, eg T0000123, T0000124
- identical dates of birth (DOB), eg 01.01.1900.

These systems create a risk of misidentification compared to other patients for whom first name and surname, unique NHS number and individual date of birth are all used. Also, temporary numbers that are unique locally may not be suitable when a patient transfers between hospitals. While many EDs have created less error-prone combinations of identifiers, their differing practices can confuse staff when changing jobs and moving between hospitals.

For DOB, the convention of using 01 Jan 1900 for adults and 01 Jan 2000 for children has become impractical: pathology systems can reject 1900 as implausible and 2000 no longer indicates a child. Using the same DOB for any unidentified patient may also lead to misinterpretation of pathology results because normal ranges are given by age and does not meet age-related transfusion guidelines.

When registering unidentified patients the following system must be implemented:

For **names**:

a distinctive method is to randomly generate combinations of first and surname from an edited phonetic alphabet eg Foxtrot Whisky. A list is available for each site held by the admin lead, the list must be maintained and indicating that the name has been taken from the list and the date used.

For **temporary numbers**:

use the RiO locally generated number. If the patient is admitted directly from A&E record any temporary acute hospital numbers, that have already been allocated, in the RiO 'alternative id' fields via the demographic screen.

For **DOB**:

Staff are asked to combine 1 Jan with an estimated year of birth, eg 01 Jan 1950, 01 Jan 2015. While unlikely to be the patient's true age, this approach is safer than using a standard DOB.

Merging medical records, once the patients identify is known.

If this is a new patient simply correct the RiO record, if the patient record already exists in RiO please raise a request to merge the two records with the Clinical Systems team via the IT Servicedesk Portal.

Source

Patient Safety Alert - Alert reference number: NHS/PSA/RE/2018/008
Joanne Sims
Risk and Datix Manager June 2019