

Standard Operating Procedure - Transcribing within Integrated Nursing and Conditions Service (Adults)

Reference No:	933				
Version:	4				
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First Issued On:	1.8.2019				
Latest Issue Date:	December 2024				
Review Date:	30th November 2026				
Referenced	See reference section				
Documents:	See reference section				
Ratified By:	Therapeutics & Pathway Group				
Distribution:	MyCompliance, Community Nursing Teams, GP Practice				
Distribution.	Staff, Local Medical Committee, The Local				
	Pharmaceutical Committee				

Document Revisions									
Date	Author	Nature of Change	Reference						
August 2019	Emma Baggaley	New SOP							
March 2020	Emma Baggaley	Inclusion of further GP surgeries and how to view patient details for EMIS practices	Appendix 2						

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April 2020	Emma Baggaley	Update to include transcribing of	Appendix 1
		Insulin charts during COVID 19	
		pandemic	
May 2021	Emma Baggaley/	Review and update to remove	
	Andrew Powell	COVID 19 as standard practice	
		now	
May 2022	Emma Baggaley	Review of SOP prior to roll out of	Version 2
		transcribing across Hull and ER	
October 2023	Emma Baggaley	Review of SOP prior to roll out of	
	Tracy Turner	transcribing across ER PCNs	
	Jane Jennison		
August 2024	Emma Baggaley	Remove S1 label as service now	Version 3.1
		paperlight	
		Expand the number of	
		medications that can be	
		transcribed	
December	Tracy Turner	Addition of Palliative Care	Version 4
2024		Medication.	

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1. INTRODUCTION

Transcribing can be defined as the act of making an exact copy, usually in writing. In the context of this Standard Operating Procedure (SOP), transcribing is the copying of previously prescribed medicines details to enable their administration in line with legislation (i.e., in accordance with the instructions of a prescriber) (RCN 2019).

This SOP must be read in conjunction with CHCP Guide to Safe and Secure Handling of Medicines.

2. PURPOSE

The purpose of this document is to describe the standard operating procedures for Transcribing within Integrated Nursing and Conditions Services (Adults). This SOP also describes the process for transcribing of palliative care medication , for the purposes of recording administration, by community nurses and the palliative care team. This process will be piloted with GP surgeries and then further rolled out across Hull and East Riding.

3. SCOPE

To provide an agreed framework, with specified parameters, which allows the process of transcribing to take place within a safe and supported environment. This SOP will be used to allow transcribing all injectable medication by community nurses and palliative care teams across Hull and East Riding for the purposes of recording administration of the medication.

4. FAIRNESS, RESPECT, EQUALITY, DIVERSITY, INCLUSION & ENGAGEMENT

CHCP promotes the principles of FREDIE (Fairness, Respect, Equality, Diversity, Inclusion and Engagement) throughout the organisation and beyond. Whilst supporting and sustaining an inclusive and diverse workforce that is representative of the community it serves, equally we are committed to the provision of services that not only respect our increasingly diverse population but also which promotes equity of access and care.

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This document has been developed with due consideration to the principles of FREDIE including completion of an equality impact assessment (EIA).

5. ABBREVIATIONS & DEFINITIONS

JIC Just in Case

SOP Standard Operating Procedure

RCN Royal College of Nursing

FREDIE Fairness, Respect, Equality, Diversity, Inclusion & Engagement

EIA Equality Impact Assessment

CHCP City Health Care Partnership

SCR Summary Care Record

SystmOne Electronic Patient Record System

MAR Medication Administration Record

NHS National Health Service

g Grams

mg Milligram

ml Millilitres

IV Intravenous

INH By Inhalation

IM Intramuscular

SC Subcutaneous

PR Per Rectum

PV Per Vagina

PO By Mouth

NEB By Nebuliser

Oral By Mouth

Gastro/PEG By Gastrostomy

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Top Topical

T Transcribing

CN Community Nursing

DSN Diabetic Specialist Nurse

GP General Practitioner

Rx Prescription

PRN As needed (pro re nata)

EMIS Electronic Management Information System

6. PROCESS

Staff members who transcribe are accountable for their actions and omissions. To be considered for the transcribing role a nurse must:

- be in a Band 5 position or above, assessed and recorded as competent in the administration of medicines
- undertake training in transcribing and be assessed as competent to transcribe by their line manager. Assessment of competency will be reviewed annually.
- have completed their initial probation period.

Transcribing cannot be undertaken within the electronic system. Transcribing can only be undertaken by hand onto the CHCP medication chart for the purposes of recording administration of a medicine.

Transcribing the information is copying from the sources below without any alterations or additions. Changes may not be made based on information provided by the patient, family member or carer. If information is provided by the patient regarding any changes to their medication, then the nurse must liaise with the prescriber for clarification.

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In most instances the community/hospital pharmacy label is the primary source used to transcribe, however, to ensure safety and reduce risk, the details on the label must be checked against a second source from the following list:

- List of medication obtained from the patients electronic medical record (Systm1 or EMIS). For guidance on how to view the EMIS record, see Appendix 2.
- List of medication obtained from the patients Summary Care Record (SCR)
 following patient consent
- Discharge prescription/immediate discharge summary written in the hospital where the service user has been

The second source used must be documented in the patient's electronic S1 record.

Only use recently dispensed medicines with a pharmacy label as evidence of current drug treatment (e.g., dispensed within the last month unless a patient was given more than one month's supply at time of dispensing or if a patient has been prescribed JIC palliative care medication in advance). The dose on the pharmacy label (i.e. prescribed dose) must match the dose that is transcribed on the medication chart.

Nurses MUST not amend or write on the pharmacy label to update a dose. It is permissible to cross through the pharmacy label for JIC medications when the prescriber has provided written evidence of a dose change, see section 6.8.

All patients referred into community nursing teams must give consent to their medical records sharing, those deemed not to have the mental capacity to consent must have the appropriate authorisation for sharing through Lasting Power of Attorney (Health and Wellbeing), Best Interest Assessment or Court Order.

6.1 Transcribing Process

- Fill in all required fields on the CHCP Medication Administration Record (MAR) relating to medicines and patient details. Partial completion can lead to delays and errors.
- Write legibly (in capitals) using a ballpoint pen, in black ink

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- It is the transcriber's responsibility to enter the following in block capitals of Patient's full name & address of NHS number
 - Date of Birth
 Recent weight (and date recorded)
- Enter details of drug and other allergies in the appropriate section when initially completing the MAR. If none are known then this must also be indicated. Information added on allergy status must be signed and dated at the time of entry or amendment.
- All medication details should be clearly legible to include:
 - o medication name
 - o form of medicine e.g. tablet, capsule, powder for injection, solution
 - strength e.g. 10mg. The following units may be used for expressing strength or dosage:
 - g = grams
 - mg = milligram
 - ml = milliliters
 - micrograms must be written in full
 - nanograms must be written in full
 - `units' must be written in full
 - dose and frequency e.g. 10mg TWICE A DAY. Write out the frequency in words and not figures e.g. THREE TIMES A DAY or THREE x DAILY and not 3 x daily or 3 times a day.
 - o Indicate the route of administration clearly. Accepted abbreviations are:

IV =Intravenous

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SC = Subcutaneous

PR = Per Rectum

PV = Per Vagina

INH= By Inhalation

PO = By Mouth

IM= Intramuscular

NEB = By Nebuliser

Oral = By Mouth

Gastro/PEG = By Gastrostomy

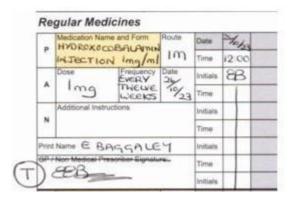
Top = Topical

All other routes should be written out in full e.g. Sublingual, Buccal. Only one route should be indicated for a given administration time.

- The date of the transcribing. The transcribing nurse must print their name, sign and date the MAR and annotate that they have transcribed the medication using a capital T circled.
- Any additional directions or information in the special instructions box
 e.g. with food
- The use of decimal points should be avoided where possible e.g. transcribe as 200 micrograms and not 0.2mg, or transcribe as 2mg and not 2.0mg.
- If small volumes are prescribed (less than 1ml) write as 0.5ml and not .5ml
- Medication labelled as "as directed" must not be transcribed onto the CHCP MAR chart. Contact the independent prescriber who will need to produce another FP10 with clear dosage instructions on or complete a CHCP MAR chart with clear prescribed instructions to allow a nurse to administer the medication.
- Ensure any indications for `as required drugs` are copied. The dose interval should be specified (e.g. every 4 hours) as well as the maximum quantity that could be administered (e.g. max 30mg in 24 hours).

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- Multiple CHCP MAR charts must be condensed onto one chart whenever it is possible. If the patient requires more than one CHCP MAR chart, mark clearly on the front of the card `1 of 2', or `2 of 2' etc.
- The whole CHCP MAR chart must be re-written when it becomes messy or illegible, especially after several medications have been stopped or changed or when the CHCP MAR chart is full. If the CHCP MAR is full and the current medication the patient is administered is still in use then the prescribed dosage instructions can be transcribed on to a new CHCP MAR. When a new CHCP MAR sheet is written the old one should be cancelled by drawing a diagonal line across it and writing 're-written'. The nurse must then print their name, sign and date the sheet.
- When a new item is prescribed midway through a CHCP MAR then the signature and time administered cells should be scored through for the dates prior to the additional item being added
- At the present time, we will continue to use the CHCP Medication Authorisation and Administration Record Charts. When transcribing, the nurse must print their name, sign the chart, and annotate the transcription with a T.



6.2 Cancellation of Treatment

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- Do not discontinue any prescribed treatment unless documented evidence from the independent prescriber is available. The source of the evidence must be documented in the patient's record.
- If the medication is discontinued, the CHCP MAR chart should be annotated
 with the reason for the discontinuation, name of nurse annotating the CHCP
 MAR and the date. A single bold line must be drawn diagonally across the
 details of the medication on the CHCP MAR chart and any remaining unused
 administration record on it.

6.3 Change to Treatment Dose

• If the change in dose is to be transcribed then the prescriber will need to produce a new FP10, with clear dosage instructions, which will be dispensed by a pharmacy who can then add the prescribers instructions on the pharmacy label. These changes need to be in place to allow transcription of the new dose onto the MAR before changes are made to administration. Amended doses should be cancelled as above and then re-written. Any boxes which contain a pharmacy label with the "old" dose must be discarded, medication that is still in date can continue to be used to administer the updated dose to the patient (for diabetic medication see administering medication section, diabetic medication (injectable)).

Immediate and necessary change to a treatment dose

• In exceptional circumstances, where medication has been previously prescribed and the prescriber is unable to issue a new prescription, but where changes to the dose are considered immediate and necessary, the use of information technology may be used but must confirm any change to the original prescription. A verbal order on its own is NOT acceptable. A new prescription should be generated and signed by the prescriber who confirmed the changes within normally a maximum of 24 hours (72 hours maximum – bank holidays and weekends). When written confirmation to the dose change is received, the new dose can be transcribed onto the CHCP MAR chart.

Acceptable written confirmation which can be viewed and acted upon by the transcribing nurse is as below:-



- For GP's using SystmOne and the GP Out of Hours Service, the prescriber should add an entry into the SystmOne patient record detailing the dose change.
- For GP's using EMIS, the prescriber should issue an FP10 via electronic prescribing which can be viewed by the transcribing nursing using third party record (see Appendix 1).
- The transcriber must document within S1 that they have amended the dose based on written authorisation and a new prescription has been requested. There will be a difference between the MAR Chart and pharmacy label for a short period of time until the new prescription (with the updated dose) is received. This information should be added to the visit list within S1 to inform the next visiting nurse.

Controlled Drugs

Any change in dose to a prescribed controlled drug needs to be covered by a new direction to administer that needs to be in writing. If the GP/prescriber record can be accessed and a new prescription can be seen to have been issued with an increased dose, then this is sufficient written evidence that there is a new direction to administer.

Acceptable written confirmation which can be viewed and acted upon by the transcribing nurse is as below:-

- For GP's using SystmOne and the GP Out of Hours Service, the prescriber should add an entry into the SystmOne patient record detailing the dose change.
- For GP's using EMIS, the prescriber should issue an FP10 via electronic prescribing which can be viewed by the transcribing nursing using third party record (see Appendix 1).

The transcriber must document within S1 that they have amended the dose based on written authorisation and a new prescription has been requested. There will be a difference between the MAR Chart and pharmacy label for a short period of time until the new prescription (with the updated dose) is

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received. This information should be added to the visit list within S1 to inform the next visiting nurse.

6.4 Medicines which can be Transcribed

The table below lists the most common medications which community nursing teams will transcribe. If a prescriber is unable to provide a CHCP MAR chart and they have issued a prescription with clear specific instructions on how to administer a medication and a second source can be checked, then these medicines can be transcribed onto a CHCP MAR chart for the purposes of recording administration.

Medicine	GP/Prescriber	Nurse	
	responsibility	responsibility	
Hydroxocobalam	GP to issue	Nurse to use	If the
in	prescription for	pharmacy	prescription
(Vitamin B12)	hydroxocobalam	label as	is issued with
	in injection with	primary source	"as directed",
To be	clear dose	from which to	this cannot
administered	instructions	transcribe.	be
intramuscularly	e.g.		transcribed
	Loading dose –	Second check	on the
	X doses to be	the dose	CHCP
	given over	against the	MAR.
	TWO weeks	patient's	Nurse will
		electronic	need to
	Or	medical record	request a
	Maintena	or patient's	new
	nce dose	SCR.	prescription
			from the GP
	1mg to be	Administer the	with clear
	administered	medication in	dosage
	once every 12	accordance	instructions
	weeks	with the	or GP will
		prescrib	need to

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		er's	complete
		instructions.	Medicines
			Administratio
			n and Record
			Chart to
			authorise
			administratio
			n.
Dalteparin	GP to issue	Nurse to	If the
	prescription for	use	prescription
To be	Dalteparin with	pharmacy	is issued with
administered by	clear dose	label as	"as directed",
Subcutaneous	instructions e.g.	primary source	this cannot
injection		from which	be
y	5000 units to be	to	transcribed
	administered	transcribe.	on the
	DAILY	Second check	CHCP
		the dose	MAR.
		against the	Nurse will
	GP to specify	patient's	need to
	the duration of	electronic	request a
	treatment.	medical record	new

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or patient's	prescription
SCR.	from the GP
Administer the	with clear
medication in	dosage
accordance	instructions
with the	or GP will
prescribers	need to
instructions	complete
	Medicines
	Administratio
	n and Record
	Chart to
	authorise
	administratio
	n

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Enoxaparin	GP to issue	Nurse to use	If the
	prescription for	pharmacy	prescription is
To be	Enoxaparin	label as	issued with
administered	with clear dose	primary	"as directed",
by	instructions	source from	this cannot be
Subcutaneous	e.g.	which to	transcribed on
injection		transcribe.	the CHCP
	2000 units to	Second	MAR.
	be	check the	Nurse will
	administered	dose against	need to
	DAILY	the	request a new
		patient's	prescription
	GP to specify	electronic	from the GP
	the duration	medical	with clear
	of treatment.	record or	dosage
		patient's	instructions or
		SCR.	GP will need
			to complete
		Administer	Medicines
		the	Administration
		medication in	and Record
		accordance	Chart to
		with the	authorise
		prescribers	administration.
		instructions	

Diabetic			
Medication	GP to issue	Nurse to use	If prescribed as
(injectable)	prescription	pharmacy	'as directed' the
e.g. Insulin	for diabetic	label as	nurse is unable to
and GLP –	injectable	primary	transcribe the
1RA (i.e.	medication	source from	dose and will
liraglutide,	with clear	which to	require another
semaglutide,	dose	transcribe.	prescription with
dulaglutide,	instructions		clear dosage
lixisenatide	e.g. 6 units in	Second	instructions or GP
and	a morning	check the	will need to
exenatide)	and 10 units	dose against	complete
	at teatime.	the patient's	Medicines
To be		electronic	Administration
administered	Change of	medical	Record Chart to
by	dose. If the	record or	authorise
Subcutaneous	prescribed	patient's	administration.
injection	dose of	SCR.	
	diabetic		
	medication is	Administer	
	changed,	the	
	then a new	medication	
	prescription	in ccordance	
	is required to	with the	
	be generated	prescriber's	
	by the GP	instructions.	
	with clear		
	dose and	Upon receipt	
	instructions	of diabetic	
	and inform	medication	
	Community	with	
	Nursing via	pharmacy	
	247111 of a	label with	

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change of the dose. amended dose, the To reduce the nurse will amount of cancel the old dose waste from the medication, nurses will **CHCP MAR** request the smallest The new number of diabetic medication diabetic pen dose will be devices e.g. transcribed 2 pens rather than a box of onto the CHCP MAR 5 pens Nurse to amend the title of the diabetic care plan within S1 as 'Transcribed' See also flow chart 'Diabetic Injectable Medication change of dose'

For injectable palliative care medication, see palliative care section

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6.5 Administering Medication

When a member of staff is administering medication from a CHCP MAR chart which a nurse has transcribed, before administering the medication they must ensure that the directions on the pharmacy label of each medication matches the transcription.

If there are any discrepancies, investigation into the reasons for this must be undertaken before administration takes place.

6.6 Diabetic Medication (Injectable)

The current diabetic medication pen device that is in use by the nursing team (and kept out of the fridge) will be stored with the CHCP MAR chart and the medication box with the pharmacy label that states the current dose of medication. This pharmacy label can then be checked against the CHCP MAR chart prior to administration of diabetic medication. A photograph of the current pharmacy label will also be uploaded on S1 (see flow chart below).

The remaining diabetic medication will be stored in the fridge in a sealable container e.g. plastic Tupperware container.

If a patient has more than one type of diabetic medication device, a separate box will be needed for each device.

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6.7 Diabetic Injectable Medication: Change of Dose (non-urgent change)

CHCP Community Nursing (CN) Team informed by Diabetic Specialist Nurse (DSN) who is a prescriber and/or GP of dose change. DSN will need to send a treatment request to the GP who will issue a new FP10 CN requests new Rx from GP, highlighting the change of dose and advise quantity of diabetic medication needed e.g. 2 pens. CN to add note to visit list that new prescription has been requested. Continue to administer existing dose of diabetic medication until the new medication (with the correct dose on the pharmacy label) is obtained. CN to take a photograph of the new community pharmacy label with a CHCP encrypted mobile phone and upload on patients SystmOne (S1) record - amend jpeg title to 'Diabetic Medication label, DATE, PATIENT NAME'. This is available to view in record of attachments. Transcribe the new dose onto the CHCP MAR Chart (cancel the old dose on the chart). Decant the diabetic medication from the pharmacy box and store in the patients container in the fridge. Discard the old medication box with the pharmacy label and keep the medication box with the new pharmacy label with current pen device in use and the CHCP MAR Chart.

NB If the pharmacy has dispensed and labelled each individual insulin pen device with dosage and frequency, then these pens will need to be discarded if there is a change in dose.

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6.8 Palliative Care Anticipatory Medications PRN injection

The example doses given below would be suitable for an opioid and benzodiazepine naive patient only and are taken from the area prescribing committee guidelines

Pain	*Morphine 2.5mg s/c 2 hourly PRN prescribe 10mg/ml ampoules 5(five) ampoules maybe repeated after 60 minutes if needed	In renal impairment eGFR <30ml/min, please use Oxycodone 1.5mg s/c 4 hourly PRN
Agitation/restlessness	*Midazolam 2.5mg s/c 2 hourly PRN prescribe 10mg/2ml injection 5(five) ampoules	(if patient in last days of life manifests features suggestive of delirium
100	maybe repeated after 30 minutes if needed	consider haloperidol +/- midazolam)
	Please ensure the 10mg/2ml injection is prescribed, very uncomfortable for patients as a s/c injection, du	
Nausea/Vomiting	Haloperidol 1mg s/c 4 hourly PRN prescribe 5mg/ml injection 5 ampoules	For patients with Parkinson's disease use cyclizine 25mg 4 hourly prn
Excess secretions/ Bowel colic	Hyoscine Butylbromide 20mg s/c 4 hourly PRN prescribe 20mg/ml injection 5 ampoules	If TWO doses are required in 24 hours consider a syringe pump containing 60mg over 24 hours

Initial prescriptions for PRN injections will be provided by the prescriber and may be transcribed onto the CHCP MAR chart. The prescription must specify clear dosage instructions including rationale e.g. pain/agitation.

NB: Prescriptions with a sliding scale dose cannot be transcribed.

If a sliding scale prescription is issued, the community nurse will contact the prescriber and request a specific dosage direction.

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For a non urgent change of a dose for PRN medication then the following process will be followed (see section 6.3 for the urgent change of dose process)

Community nurse will communicate with the appropriate prescriber to discuss review of dose for the patient. Prescriber will issue a new FP10 with the updated specific dose. Community nurse will document in the patient's electronic record and add a note to the visit list that a new prescription has been requested



Continue to administer existing doses of palliative care medication until the new medication (with the correct doses on the pharmacy labels) is obtained



CN to take a photograph of the new community pharmacy label with a CHCP encrypted mobile phone and upload to patient's SystmOne record – amend jpeg title to 'Palliative Care Medication label, DATE, PATIENT NAME'. This is available to view in record of attachments. (Delete photograph from phone once uploaded into SystmOne. Transcribe the new dose onto the CHCP MAR Chart (second source must be checked and documented and the old dose cancelled see section 6.2)).



Cross through the pharmacy label on the old medication box and keep the new medication box with the new pharmacy label which details the new dose and directions and the CHCP MAR chart.

6.9 Palliative Care Medication via sub-cutaneous Continuous Infusion

When transcribing palliative care medications for administration via sub-cutaneous continuous infusion (syringe pump), the prescriber must issue an FP10 prescription for the drug(s) required according to the patients' symptoms. The prescription must specify clear dosage instructions including rationale e.g. pain/agitation.

NB: the prescriber will also need to issue a prescription for the appropriate diluent i.e. water for injection

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If the above is in place, then the palliative care medications for sub-cutaneous continuous infusion can be transcribed on to the CHCP MAR chart by the community nurse as per the process above. When there is a change in dose for sub-cutaneous continuous infusion then the process below should be followed.

NB: Prescriptions with a sliding scale dose cannot be transcribed.

If a sliding scale prescription is issued, the community nurse will contact the prescriber and request a specific dosage direction.

6.10 <u>Palliative Care Medication via sub-cutaneous Continuous Infusion: Change of Dose</u>

CHCP Community Nursing (CN) Team informed by Specialist Palliative Care Nurse who is a prescriber and/or GP of dose change. The prescriber will need to issue an FP10 with new details specified. This information must also be visible within the patient electronic record (see section 6.3 for urgent changes to a treatment dose for a controlled drug)

Continue to administer existing doses of palliative care medication until the new medication (with the correct doses on the pharmacy labels) is obtained.

CN to take a photograph of the new community pharmacy label with a CHCP encrypted mobile phone and upload to patient's SystmOne (S1) record – amend jpeg title to 'Palliative Care Medication label, DATE, PATIENT NAME'. This is available to view in record of attachments. Delete photo from phone once uploaded onto SystmOne. Transcribe the new dose onto the CHCP MAR Chart (cancel the old dose on the chart see section 6.2).

Cross through the pharmacy label on the old medication box and keep the new medication box with the new pharmacy label which details the new dose and directions and the CHCP MAR chart.

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7. APPROVAL

This SOP has been reviewed and approved by the stakeholders identified on the document checklist submitted to the Therapeutics & Pathway Group which reviewed the checklist and ratified this document.

8. MONITORING & COMPLIANCE

Incidents relating to transcribing will be recording on CHCP incident reporting system, and will be reviewed and reported to Therapeutics and Pathway Group.

9. REVIEW

This SOP will be reviewed every 2 years or sooner as needed as the transcribing programme is rolled out across Hull and East Riding.

10. ASSOCIATED DOCUMENTATION

- CHCP Medicines Policy
- CHCP Guide to the Safe and Secure Handling of Medicines
- CHCP Diabetic Protocol
- Referral & review requests for the Diabetes Specialist Nursing Teams HUTH,
 Hull and ER
- Microsoft Word Herpc April 2017

11. APPENDIX 1 - Viewing EMIS shared care record through SystmOne

SystmOne and EMIS Integration – Introduction

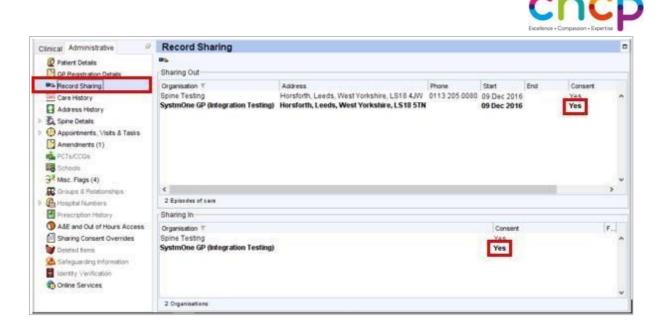
Currently SystmOne has direct shares with other GP Practices that deliver care through SystmOne, allowing Clinicians to view a patient's GP record, where relevant shares are in place. CHCP have recently joined a pilot that will also allow viewing into EMIS records, meaning we will also have visibility of those patients registered within EMIS Practices. As with SystmOne this will be dependent on the relevant shares being available.

This document will outline:

- Which records will be shared with EMIS
- How to view a patients EMIS record

Which records will be shared with EMIS?

The sharing consent for the patient at your organisation is highlighted in bold. To share data from SystmOne to an EMIS organisation, the patient must have consented to share out. This is indicated by a **Yes** or a **Not asked – Record shared** in the Consent column.



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In order to view EMIS records in SystmOne, the patient must consent to sharing in at the SystmOne organisation.

Before the record is sent to EMIS, the **EMIS user** will also need to record consent from the patient to view the SystmOne patient record.

Viewing a Patient Record sent from EMIS

Log on with a smartcard so you get the latest details from PDS for the patient

Note: If you are not logged on with a smartcard, you will not be permitted to use the integration. If a patient is flagged as Spine Sensitive, then you will not be able to use the integration for this patient

- 1. Select Third Party Patient Record
- 2. Select Check for Records this will send a message to participating EMIS organisations querying whether EMIS has the patient registered and if information for the patient is available to be shared. The name of the organisation will appear as a new sub-node within the clinical tree

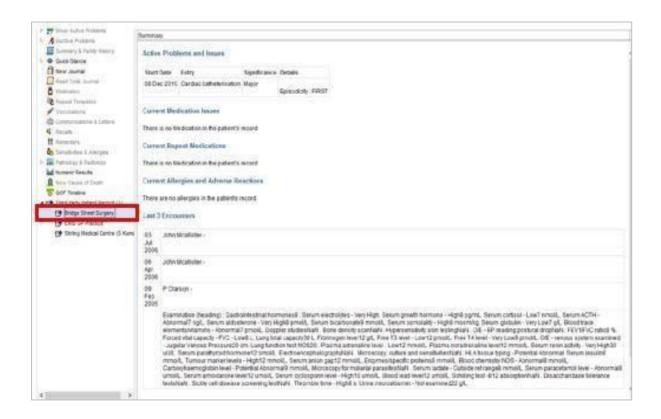


3. Select the EMIS Practice from the list



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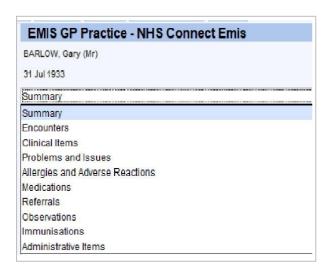
Clicking on the name of the Practice from the Clinical Tree will retrieve the record from that organisation for the patient. The Summary view will be displayed by default.



EMIS does not support the sending of data for three categories within the Summary. This data may be present in the patient's EMIS record. These categories are:

- Warnings
- Key Indicators
- Current Recalls
- Historical Allergies and Adverse Reactions
- Investigations
- Administrative Items

After retrieving the record, you can change the view using the drop-down menu



Some views have a date filter, which will default to show the last 6 months of information. Users can then change this date filter to request the information that they are interested in.



Pressing Ctrl+F on the keyboard will bring up a search dialog. This will make finding specific information in the record quicker and easier, particularly on screens that have a lot of information e.g. the Encounters view.

Signature sheet:

For staff to sign when they have read and accepted the standard operating procedure.

Date	Print Name	Signature

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