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How timely is access to palliative care medicines in the community? A mixed methods study in a UK city

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SCHOLARONE™ Manuscripts Title: How timely is access to palliative care medicines in the community? A mixed methods study in a UK city

Elizabeth Miller¹, Julie D. Morgan², Alison Blenkinsopp²

- ¹ Sheffield Teaching Hospitals NHS Foundation Trust, Sheffield, UK
- ² University of Bradford, Bradford, UK

Corresponding author: <u>elizabeth.miller@sth.nhs.uk</u>

Pharmacy Department, Royal Hallamshire Hospital, Sheffield S10 2JF

Tel: 0114 2711900

ABSTRACT

Objective: To investigate timeliness of access to palliative medicines (PMs) from community pharmacies to inform palliative care service delivery.

Design: Mixed methods in two sequential phases: 1. Prospective audit of prescriptions (Px) and concurrent survey of patients/representatives collecting PMs from pharmacy; 2. Interviews with Community Pharmacists (CPs) and other healthcare professionals (HCPs).

Setting: Five community pharmacies in Sheffield, UK and healthcare professionals that deliver palliative care in that community.

Participants: Phase 1: Five CPs: two providing access to PMs within a Locally Commissioned Service (LCS) and three not in the LCS; 55 patients/representatives who completed the survey when accessing PMs; Phase 2: 16 HCPs, including 5 Phase 1 CPs, were interviewed.

Results: The Px audit collected information on 75 prescriptions (75 patients) with 271 individual PMs; 55 patents/representatives (73%) completed the survey. Patients/representatives reported 73% of PMs were needed urgently. In 80% of cases patients/representatives received all PMs on the first pharmacy visit. One in five had to travel to more than one pharmacy to access PMs. The range of PMs stocked by pharmacies was the key facilitating factor. CPs reported practical issues causing difficulty keeping PMs in stock and playing a reactive role with palliative prescriptions. Confidentiality concerns were cited by other HCPs who were reluctant to share key patient information proactively with pharmacy teams. Inadequate information transfer, lack of CP integration into the care of palliative patients, and poor HCP knowledge of which pharmacies stock PMs meant patients and their families were not always able to access PMs promptly.

Conclusions: Consistent routine information transfer and integration of pharmacy teams in the care of palliative patients are needed to achieve timely access to PMs. Commissioners of PM access schemes should review and monitor access. HCPs need to be routinely made aware and reminded about the service and its locations.

Key Words: palliative care; community pharmacy services; pharmacists; prescriptions; interprofessional issues

Word Count 5038

Article Summary

Strengths and limitations of this study

This is the first published study to identify the relative impact of factors contributing to non-timely access to PMs.

This paper is the first in the UK to examine perspectives of community pharmacists (CPs), general practitioners (GPs) and other community healthcare professionals (HCPs) on factors supporting and hindering access to PMs.

Fewer pharmacies participated in the prescription data collection than anticipated and had fewer PM prescriptions than previous audits and prescribing data would suggest.

This is the first study to examine customer experience of accessing PMs and the survey achieved a high response rate and generated valuable information for HCPs and commissioners.

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INTRODUCTION

Population aging and an increase in those dying with complex multi-morbidity will increase the need for palliative care with predictions suggesting End-of-Life Care (EOLC) provision in the community and care homes needs to double by 2040. UK strategy aims to improve accessibility to palliative care in the community in alignment with research suggesting most people would prefer a home death with 80% not changing their preference as their illness progresses, however resource issues in primary care can make attaining this very challenging. In particular, accessing medicines for symptom control towards the end-of-life is imperative in controlling pain and distress and to prevent unnecessary hospital admissions. Difficulties in predicting the end-of-life, especially in those with chronic illnesses who have an uncertain disease trajectory, can lead to an unpredictable yet urgent demand for Palliative Medicines (PMs) in the community.

For most patients in primary care the source of medicines is from their community pharmacy (retail pharmacy or "chemist shop") however previous research and service audits show access to PMs such as injectable medicines used for symptom control towards the end-of-life may not be as timely as patients and their families may need and wish.⁴⁻⁶ Pharmacies cannot stock every possible PM; local formularies help to address this in the UK. However knowledge on which PMs are listed in the formulary and those pharmacies holding stocks may be lacking among prescribers which could lead to Prescriptions (Px) being issued for 'non-formulary' items not on the approved local palliative care stock list and/or Px being presented to pharmacies that do not routinely hold PM stock. 4 6-11 Delays may also be caused by legal errors on Px necessitating the pharmacist make professional and ethical judgements in supporting patient care especially in the out-of-hours period.⁴⁶⁸ (Stuart, J. 2013. 'Investigating the prevalence and nature of controlled drugs prescribing errors identified in community pharmacies.' Unpublished MSc dissertation, University of Strathclyde, UK.) There is a suggestion that hand-written Px may be particularly problematic due to higher Px error rate and out-of-hours presentation, and they are still in use in the UK for home visits. 4612 (Stuart, J. 2013. 'Investigating the prevalence and nature of controlled drugs prescribing errors identified in community pharmacies.' Unpublished MSc dissertation, University of Strathclyde, UK.)

Australian research on a proposed core set of PMs found that pharmacies stocked on average three out of the list of 12,¹⁰ while a systems analysis in Ireland found that not stocking PMs in the pharmacy was the most likely factor leading to delays⁹ and this has also been found in the UK.⁴⁶⁸¹¹ Reported contributory factors include: the unpredictable nature of PM Px requests; national stock shortages; the Px of PMs or strengths not on the recommended palliative care stock list; unlicensed medicines; errors on Controlled Drug (CD) Pxs and the inability to contact the prescriber, for instance outside GP (family doctor) practice opening hours.⁴⁸¹²

Community pharmacies may take part in local or nationally commissioned services to support access to PMs in the community. In England, a Locally Commissioned Service (LCS) can be provided by the local clinical commissioning group (CCG) or a Local Enhanced Service (LES) can be commissioned by NHS England Area Teams in response to public need. Such services differ across geographical regions and are not commissioned from all pharmacies, causing confusion for patients and their caregivers who are often involved in Px collection and medicines management when a patient's condition deteriorates.^{7 13-15} Furthermore, a lack of monitoring of PM availability against those prescribed both within the pharmacy and by the commissioning body could mean PMs are not available when needed. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK)

There is little research internationally on community pharmacists' (CPs) involvement in supporting timely access to PMs. Hence this study seeks to answer the question 'What is the community pharmacist's role in the delivery of timely access to palliative care medicines.' Due to the dearth of published research particularly in the context of community pharmacy services in England the aim of this study was to evaluate timely access to PMs in the community pharmacy setting and make recommendations to inform the commissioning of services and future practice. The objectives were to:

- determine the timeliness of access to PMs in the community;
- investigate the prevalence and nature of prescribing errors on Pxs for PMs presented to community pharmacies and determine whether errors impact on access to urgent PMs;

- investigate processes for accessing PMs from pharmacies where a locally commissioned service (LCS) operates including referrals when PMs are not available;
- explore the views and experiences of CPs and other stakeholders on accessing PMs from community pharmacies.

METHODS

This study used mixed methods across two sequential phases (See Table 1 for study overview) conducted in Sheffield, UK. Participants in both phases gave informed consent before taking part. Ethical approval was obtained from the University of Bradford.

Table 1 Overview of study phases

Phase 1:

Audit of palliative prescriptions meeting inclusion criteria in participating pharmacies from May - October 2016

Customer survey for those collecting palliative prescriptions in participating pharmacies from May - October 2016

Phase 2:

Semi-structured face-to-face interviews with pharmacists participating in Phase 1 and other healthcare professionals involved in palliative care in the community from September 2016 - March 2017

Phase 1: Audit of PM supplies over six-month data collection period. Sheffield pharmacies were recruited through e-bulletin sent by the Local Pharmaceutical Committee (LPC), fax invitation to LCS PM pharmacies (19 of the 128 in the city), and verbal invitation at a local pharmacy practice development event. CPs expressing interest in taking part were given an information leaflet and consent form via email providing further information and the study inclusion criteria. Eligible CPs participated in the LCS or usually dispensed thirty or more PM Pxs in a month based on NHS Digital Px data for opioid analgesics and midazolam dispensed in pharmacies in the region. Exclusion criteria were: i) pharmacists who had worked in the UK for less than 12 months (to ensure participants were familiar with UK and local community pharmacy services), and ii) if the company or manager did not give

permission for participation. None of the interested pharmacies had to be excluded based on these criteria.

A pragmatic approach to sample size was taken; the intention was to recruit up to 15 pharmacies however only five CPs consented to participate, partly due to the unexpectedly low level of PM Pxs reported. Informed consent was obtained. EM personally visited each participating pharmacy to brief them on the project, data collection forms and answer any questions to enhance consistency of data collection.

Consenting pharmacies collected data on thirty consecutively presented Pxs which contained medicines likely to be prescribed for patients who are palliative i.e. in the last year of life, using criteria provided by the researcher to identify prescriptions. Px data was intended to be collected for a four-week period in May 2016 but due to the low level of palliative care Pxs all consenting pharmacies agreed to continue until October 2016. Pharmacy data collection forms were developed by EM and reviewed by JDM and piloted in one community pharmacy. The form recorded anonymised Px data including: first part of patient's postcode; names of medications on the Px; whether there was a legal or non-legal error on the Px and further information on how that error was resolved. Legal and non-legal Px errors were identified by the CPs and non-legal errors were classified by EM according to criteria within the PRACtICe study.¹⁷ Further details on non-legal errors were completed on a separate form to allow EM to verify the classification. Previous research suggested that delays may be caused by doctors prescribing products not on the local stock list^{6 8} hence where prescribers issued legally correct Pxs for products not recommended on the LCS PM stock list these were classified as non-legal errors. Subcutaneous items in the audit were checked by EM against the LCS stock list to identify non-formulary items in both LCS and non-LCS pharmacies.

Prescriptions were classified as urgent when i) the survey respondent stated it was urgent, ii) they included anticipatory subcutaneous medicines and PMs to be given by a syringe pump iii) they were from an out-of-hours provider. The date/time a Px was received by the pharmacy and the date/time when it was ready for collection were recorded by pharmacy staff.

Survey of patients/representatives collecting PM prescriptions. The national pharmacy contracting organisation the Pharmaceutical Services Negotiating Committee (PSNC) Community Pharmacy Patient Questionnaire (CPPQ)¹⁸ was used as a basis to develop a short customer survey of experiences of patients and their representatives of collecting Pxs with PMs from the community pharmacy. Questions included the perceived urgency of the Px, the customer's previous use of the pharmacy, whether they were the patient or the patient's representative, whether they were able to access all required PMs, whether they had been referred to the pharmacy (e.g. by another healthcare professional) and whether they had to visit more than one pharmacy to access the PMs on the Px. When patients/representatives indicated that not all items were available they were given the option of completing a free text section to explain how they intended to get these items. A free text section allowed respondents to record their answer to 'are there any things that could have been improved to make your visit better?' The customer survey was developed by EM with input from JDM, AB, a hospice patient user coordinator and risk manager. It was piloted in one pharmacy, further refined and piloted with patients within a hospice day centre. Pharmacy teams were provided with a written briefing on how to introduce the survey to patients/representatives. Individuals collecting Pxs for PMs were invited to participate by pharmacy counter staff or the CP depending on the procedure decided upon by the pharmacy. A unique number was used to match the customer survey to the pharmacy data form to allow verification of the data and assess any discrepancies.

Patients/representatives not attending the pharmacy, e.g. home deliveries and care home residents, did not complete the customer survey.

Phase 2: Semi-structured interviews with CPs and other HCPs involved in care of palliative patients. EM conducted interviews with the five CPs participating in phase 1 and with a purposive sample of other HCPs involved in palliative care in the community including GPs, community specialist palliative care team, community nurses, district nurses and intermediate care team members. HCPs were invited to participate via e-bulletin, email and through gate-keepers (practice managers and team leaders). Interviews were audio-recorded with consent and transcribed verbatim by EM. The interviews explored views and experiences of accessing PMs

in the community; factors that supported or hindered access and their knowledge of the LCS. The interview schedule is available on request from the authors.

Data analysis

Prescription data were entered into and analysed using IBM SPSS[®] V.23 statistical software by EM. Frequencies and percentages were calculated for all categorical variables with mean and standard deviation calculated for time to process Pxs.

Interview transcripts were read by EM for content familiarisation then annotated and coded manually using a priori themes from the study objectives. Following development of an analytical framework, two over-arching themes were then used to 'chart' the coded data: (1) timely access to PMs, (2) the community pharmacist's role in palliative care, using the Framework Method. The framework was revised and iteratively refined with CW and AB against the coded interview transcripts with emergent themes and subthemes applied across the whole data set. Summaries of data were added within the framework to capture participant's views. Mapping and interpretation of findings compared similarities and contrasts between and across professional groups and was supported through discussion and reflection with AB and CW.

Data from both phases were then triangulated where two or more sources agreed or contrasted with each other to help explain the quantitative results of the study. Triangulation enhanced the validity and reliability of the results and enabled integration of the findings, such that it was possible to make recommendations for practice improvement and identify issues for service commissioners to consider.

Patient Involvement

The study was informed by research priorities in palliative care²⁰ and through EM's professional experience including discussion with patients and carers experiencing medicines access problems following admission to a hospice. The customer survey tool was developed and piloted with patients in a hospice day setting.

RESULTS

Participants in each phase of the study

Phase 1 CP audit: Participating pharmacies were diverse in that they included pharmacies classified as independent (having less than five branches) and multiple (having five or more branches); two provided access to PMs under an LCS and three did not. Pharmacy sites were a combination of high street/local parade of shops (3), and suburban (2) with both suburban pharmacies co-located with a GP practice. For pharmacies not consenting to take part, the main reason cited was small numbers of palliative care Pxs dispensed in the pharmacy.

Customer survey: Customer surveys were completed against 55/75 CP audit forms; response rate 73.3%. Non-completion related primarily to home deliveries and care home Pxs.

Phase 2 CP and other HCP Interviews: 16 individuals participated: CPs (5), GPs (3), Specialist Palliative Care Team (2), Community Nurses (5), and Intermediate Care Team (1). The five CPs were also involved in phase 1. Median interview durations were 51 minutes for CPs and 18.5 minutes for GPs and other HCPs.

Phase 1: Prescription characteristics

A total of 271 Px items on 75 Px forms was recorded (range 2 to 33 per pharmacy, median 14) over the 6-month audit period with a mean number of 3.6 Px items per form. This included 68.3% (n = 185) of PMs identified as urgent, 49.8% (n = 135) containing subcutaneously administered PMs and 24.7% (n = 67) containing subcutaneously administered CDs. In 91.1% (n = 123) of cases, subcutaneous items were chosen from the LCS formulary stock list. Non-formulary choices were either different presentations of formulary items or items not on the LCS stock list (see Table 2). Varying strengths of midazolam ampoules accounted for 41.7% (n = 5) of non-formulary choices.

Table 2: Subcutaneous items and formulary status on the locally commissioned palliative service stock list

Formulary or non- formulary item	Frequency	Percentage prevalence
Item included in LCS stock	123	91.1%
Non-formulary item chosen was a different strength, size or presentation to the item on the LCS stock list	8	5.9%
Non-formulary item not on LCS formulary stock list	4	3.0%

Prescriptions were computer-generated (n = 245, 90.4%) or hand-written (n = 22, 8.1%), with no Pxs delivered electronically via the Electronic Prescription Service (EPS); missing data (n = 4, 1.5%). Most Pxs were written by NHS GPs providing inhours services (n = 233, 86%), with out-of-hours GPs (n = 33, 12.2%) or specialist palliative care team (n = 5, 1.8%) writing the remainder. There were no non-medical prescriber (NMP) Pxs within the sample. Prescriptions were presented to the pharmacy during GP opening hours (from 9-6pm Monday to Friday) (n = 176, 64.9%) or outside GP hours (evenings and weekends) (n = 77, 28.4%); missing data on 6.6% of forms (n = 18).

Phase 1: Prescription audit

Legal problems arose in 1.1% (n = 3) of Px items; all of which were computergenerated, not specifying a dose on a controlled drug (CD) given via infusion. There were no legal errors on handwritten Pxs for PMs. There was insufficient evidence of a difference between Px generation method and legal errors (Fisher's Exact 2-sided test, p = 0.052). Other non-legal prescribing errors such as incomplete information, dose/ strength error, generic/ brand error, allergy, and quantity error occurred in

3.0% (n = 8) of items. Table 3 summarises prevalence of different medication problems on Pxs and table 4 indicates types of prescribing errors using categories as in the PRACtICe study.¹⁷

Table 3: Prevalence of medication problems

Type of medication problem	Total number of prescription items	Frequency of problem	Percentage prevalence
Legal problems	271	3	1.1%
Prescribing errors (see table 4)	271	8	3.0%
Out of stock with supplier	271	1	0.4%
Non-formulary LCS item requested	135	12	8.9%
		4	

Table 4: Prescribing errors (n=271)

Type of prescribing error	Frequency	Percentage prevalence
Incomplete information on prescription	2	0.7%
Dose / strength error	2	0.7%
Generic / Brand error	2	0.7%
Allergy	1	0.4%
Quantity error	1	0.4%

Phase 1: Time to access urgent palliative care medicines

Valid time data was available for 57.8% (n = 107) of 185 urgent items (n = 73 missing data; n = 5 excluded where PMs unavailable and Px taken elsewhere and recorded as 0 minutes). Median time to process (time of Px receipt to time of complete supply of PMs) urgent PMs was 2 hours (10 minutes in LCS pharmacies and 5 hours in non-LCS pharmacies). The maximum time to process urgent PMs was 3 hours and 39 minutes within LCS pharmacies, and 47 hours and 15 minutes within non-LCS pharmacies (see figure 1).

The median time taken to access urgent medications (107) between pharmacies participating in the LCS and pharmacies not participating in the service was significantly different (independent samples median test p = 0.002 at 95% confidence level); with pharmacies not participating in the LCS taking significantly longer than pharmacies in the LCS.

Figure 1: Time taken for urgent palliative medicines between pharmacies (see separate file)

Legal errors had minimal effect on access as all urgent PMs with legal errors were available within 30 minutes of presentation. Legal errors were resolved by: contacting the nursing home to specify the dose to be given on a Px for PMs via a syringe driver using a community medicines administration record, using the pharmacy Patient Medication Record (PMR) to access information on a previously issued Px, and contacting the prescriber. The Summary Care Records (SCR) was not used to resolve errors in the Px audit sample.

Phase 1: Customer Survey

Survey responses showed that representatives collected PMs on behalf of the patient (65.5%); for both themselves and the patient (1.8%); and patients collected their own PMs (32.7%); 72.9% of surveys overall indicating the Px included urgent item(s). All cases for urgent subcutaneous medications were collected by a representative on behalf of a patient. In 42.6% of cases the patient attended their usual pharmacy. Patients/representatives also indicated the pharmacy was: convenient (14.8%); one of several pharmacies used (20.4%), or that they had been referred to the pharmacy for the medications (21.8%).

In 80% of cases patients/representatives received all medications against the Px at the first pharmacy they visited. In 20% (11/55) one or more items on the Px was not available, in five of these the item(s) were urgent. Free text sections were completed for six of the 11 cases of unavailable items. Four indicated they would return to collect the item from the pharmacy and two said they would try another pharmacy to obtain the items.

Overall one in five patients/representatives had to go to more than one pharmacy to get urgently needed PMs, increasing to one in three for urgent subcutaneous injection Px items. One in every two patients/representatives referred to the pharmacy by another healthcare professional had to go to more than one pharmacy.

Thirteen respondents made additional comments on whether their experience could have been better. Comments were mostly positive: six indicated 'no', 'none', 'no fine' or similar phrase; five made comments on the staff or service: 'friendly services under difficult circumstances', 'no - staff really friendly and helpful, service was quick and efficient', 'Nothing – excellent and quick service'; and one explained 'nothing

much that would make it better, but I phone in advance to make sure my items are in stock'. One respondent requested to 'keep a stock of all required items'.

Phase 2: Interview findings

Timely access

Community nurses and palliative care team staff described how they met the need for advanced planning to prescribe anticipatory medicines prior to the last days of life for patients where appropriate. They reported conducting an end of week check and balance to ensure sufficient stock for over the weekend when fewer staff were available. Specialist palliative care team staff also described making do with the medicines already available in the house for a syringe driver and then ordering medication for the next day. CPs recognised that the GP may be open to change the Px to an item that is available due to the need for timely access rather than waiting for the 'perfect combination' to be in stock.

Challenges

CPs described practical issues in supplying PMs (Box 1) for example: stock ordering processes; CD cabinet size (to meet UK legal requirements for storage); an inability to return CD items to suppliers due to legal restrictions and wholesaler cut-off times for same-day deliveries. Furthermore, patient records and charts to check opioid dose changes and syringe drivers were often not accessible to them. Restrictions under the NHS England pharmacy contract for Medicines Use Reviews (MURs) meant pharmacists were unable to routinely see and review palliative patients unless they could physically attend the pharmacy.

Knowledge of LCS

The three CPs who were not LCS providers knew of the LCS and how to refer a patient/carer if they did not have the requested medication available. Usually they would phone ahead to the LCS pharmacy to check the medication was available before making a referral. Often making a referral depended on whether the carer had access to a car. Other HCPs had little knowledge of either the LCS or the pharmacies commissioned to provide it (Box 2) but knew which pharmacies were likely to keep some PMs in stock. GPs generally thought that all pharmacies kept

some injectable PMs in stock but said they might ring in advance to check the medication was available if a supply was needed urgently.

GP practice systems

GPs indicated that the GP prescribing system was helpful in ensuring a legally correct Px and ensuring the correct medication was prescribed according to local guidelines. The local CCG had implemented a template on the GP prescribing system to provide a 'suite' of PMs according to local last days of life algorithms which included some of the injectable medicines listed on the LCS formulary. Even so, in phase 1 several 'non-formulary' medications not on the local CCG stock list were prescribed and in phase 2 CPs in LCS pharmacies described non-compliance with the local formulary as a reason for a lack of timely access to PMs.

Communication and Collaboration

Two pharmacies not in the LCS had worked more closely with GP practices to discuss and agree to stock a smaller subset of the LCS PMs; such pharmacies had similar response times to pharmacies in the LCS which stocked a wider range of medications. CPs reported that some patients/carers worked with pharmacy processes by contacting the pharmacy when they ordered a Px for a CD that might not be stocked. Community and specialist palliative care team staff described how they would suggest the pharmacist kept sufficient stock in when they had someone on a syringe driver or enormous quantities of injectable medications. There appeared to be some examples where excellent communication and collaboration existed between GPs, HCPs and CPs which resulted in more timely access for PMs. However, concerns around patient confidentiality by GPs and other HCPs meant that more often this information was not shared with the pharmacy team in advance of receiving the Px.

Box 1 Community pharmacist interviews

Timely access

We can normally order things for the same day, if it's before midday we can get them [medicines] for 4 o'clock that afternoon. (P2, Community Pharmacist)

Challenges

We don't have an ability to be able to keep a lot [controlled drugs] and so we have a particular issue with the quantities that they write on the prescriptions sometimes which can impact on the next patient. (P4, Community Pharmacist)

We've only got very small CD cabinets...the more controlled drugs you keep the more issues you are going to have (P1, Community Pharmacist)

We've got three different strengths of oxycodone injection, and they [GPs] prescribe all three, and you might not have one, you might have the other...it's just so frustrating...you don't want to delay treatment for what is a really difficult time for the patient and the family...but unfortunately our hands are tied by the legislation and our ability to be able to alter any of these prescriptions (P4, Community Pharmacist)

Knowledge of Locally Commissioned Service / GP Practice Systems

The big problem is midazolam...so many strengths...volumes of ampoules...the GPs just pick one. (P5, Community Pharmacist)

Communication and Collaboration

So, the surgery down the road...one GP...rang us and said well what have you got in stock and what can you get, which I found really, really useful because as the prescription came in the stock came in and this thing was completely seamless (P3, Community Pharmacist)

The logistics of community and primary care don't support that [multidisciplinary working] as well with regards to the geographical locations of these people and with responsible pharmacist regulations. (P1, Community Pharmacist)

Box 2 GP and other healthcare professional interviews

Timely access

...I could go in now and say, 'I need these drugs' (and the CP might say) 'Oh I can get them in for 11 o'clock tomorrow morning' [exasperated laugh] it's like that's not really very helpful, I need them now (HCP7, Community Healthcare Professional)

I think if just more chemists had the bog-standard stuff in. (HCP11, Community Healthcare Professional)

I'd be enquiring what medicines were available...I might know what I want to prescribe but there's no point if it's not there and it's going to lead to a delay (HCP2, GP)

Challenges

A GP won't prescribe a syringe driver ahead of time...but that means we are always being [sic] having to do it now not in a more considered way (HCP4, Community Healthcare Professional)

Knowledge of Locally Commissioned Service / GP Practice Systems

I don't know who's commissioned we just basically know which ones we go to that are more likely to have it. (HCP4, Community Healthcare Professional)

...relatives who are running right left and centre trying to get hold of these meds...there is a commissioned service...but we don't know who they are. (HCP1, Community Healthcare Professional)

Communication and Collaboration

When we were down at [previous community nurse location] ...there was a pharmacy next door so...if we had any quick questions, we would go and talk to them...they were more like part of the team (HCP7, Community Healthcare Professional)

- ...so sometimes by sharing knowledge with pharmacists I think we could get better results for patients (HCP4, Community Healthcare Professional)
- ...but you're limited by what you can tell them [pharmacists] obviously from a confidentiality point of view... (HCP11, Community Healthcare Professional)

We don't communicate with them [community pharmacist] what the problem with the patient is we just prescribe the drugs... sometimes they can obviously work it out. (HCP3, GP)

I do have some slight reservations about them [pharmacists] knowing all those ins and outs...I'm not sure how wide that circle is in there [pharmacy]...I'd prefer it ...on just a case by case basis...to an identified clinician... (HCP10, GP)

DISCUSSION

Timeliness of access was found to primarily relate to medicines stocks held by CPs with legal errors playing a much smaller role having little impact on access to PMs in this study. Stock availability as a significant factor to support timely access has also been seen in previous studies.⁴ 8-10

Study results indicate a low prevalence of legal errors on palliative care Pxs compared to previous unpublished UK audit data, 12 (Stuart, J. 2013. 'Investigating the prevalence and nature of controlled drugs prescribing errors identified in community pharmacies.' Unpublished MSc dissertation, University of Strathclyde, UK.) in particular for handwritten Pxs and those issued by out-of-hours providers. All legal errors encountered related to the requirement to have a specific dose on a computer-generated Px for a controlled drug to be administered subcutaneously via a syringe pump. In this study and in Stuart, errors were four times more likely for injectable products compared to non-injectable products (Stuart, J. 2013. 'Investigating the prevalence and nature of controlled drugs prescribing errors identified in community pharmacies.' Unpublished MSc dissertation, University of Strathclyde, UK.) Legal errors relate to the statutory CD Px writing requirements in the Misuse of Drugs Act 1971²¹ (and subsequent amendments) and the Medicines Act 1968²² that specify that the Px must include a specific dose. As a patient deteriorates towards the end-of-life, frequent dose changes may be required for medication administered in a syringe pump; prescribers can be reluctant to include a specific dose on the Px in case this subsequently causes an error or confusion for those administering the medication. The continuation of legal errors on

subcutaneous Pxs in this study and previous studies^{4 8}(Stuart, J. 2013. 'Investigating the prevalence and nature of controlled drugs prescribing errors identified in community pharmacies.' Unpublished MSc dissertation, University of Strathclyde, UK.) suggests a review of the legal requirements should be undertaken as it is questionable whether legislation set in 1971 is relevant to clinical practice today. The PM prescribing template introduced by the primary care organisation may have impacted positively, minimising the number of errors compared to previous studies. It is not possible to ascertain the effect of EPS on error rate or timeliness as no Pxs were delivered via EPS in this study. At the time of the data collection, CD Pxs could not be transferred via EPS; pilots within the first ten GP practices started in October 2018.²³ Further studies should assess the impact of EPS on CD Pxs and access to PMs within community pharmacies.

Our findings of differential time to access PMs between community pharmacies participating in a LCS for PMs and those that did not, indicates that a local scheme can enhance access. Those pharmacies working with local GP practices to keep a small range of PMs in stock had similar access times to those within the LCS, suggesting that such collaboration can also support more timely access and improve patient and carer experience. Such wider collaboration has been advocated within national policy drivers and enables greater integration of pharmacy teams in improving patient care.²⁴⁻²⁷

Palliative patients often rely on family members and friends to support them with managing their medication especially towards the end-of-life.⁷ ¹³ ¹⁴ Our findings show that some families have to obtain urgently required medicines from a pharmacy different than the one that usually supplied the patient's medicines. CPs cannot access other pharmacies' medicines supply records, so the patient's regular pharmacy may be unaware of supply requirements; it is unclear what effect these changes in continuity of care between pharmacies might have towards the end-of-life. As patients and their representatives are free to use any pharmacy, not needing to register like they do with GP practices, accessing information to allow fast access to the required medication could be difficult. A potential solution could be through having read and write access to SCR allowing the CP including the patient's regular, out-of-hours or LCS pharmacy to record patient care scenarios to ensure safe,

continuity of care of PMs. Variable accessibility and difficulties in use of SCR by CPs²⁸ may suggest wider access to patient records is required.

One in five patients/representatives accessing PMs had to go to more than one pharmacy. This is the first study to quantify the number of patients/representatives who had to visit more than one pharmacy to access PMs. The high number could be explained by a lack of awareness of the LCS since this is not advertised to the public and there was low awareness amongst HCPs and GPs in the interviews. There was a belief by some GPs that most pharmacies kept stocks of PMs and so they did not always ring in advance of writing a Px. An alternative explanation could be that pharmacies do not regularly check their stocks or carry enough PM stocks, causing difficulties if more than one Px is presented on the same day. Monitoring of LCS/LES services by commissioners may be deficient as demonstrated in an unpublished audit across a network of commissioned pharmacies in Birmingham. In this audit it was reported only one pharmacy out of nineteen held all PMs on the formulary list and some CPs were not aware that the scheme was active. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK) There was also evidence in phase 1 of Pxs being written for items not on the LCS list which would not be usually stocked in the pharmacies. Further investigation of referral patterns from pharmacies not within the LCS and monitoring of pharmacies in the LCS may improve practice and caregivers' experience.

There are a few limitations which affect the interpretation/generalisability of the findings of this study. The small sample of participating pharmacies, missing data, reliance on CPs to identify Pxs and confounding factors such as time of day, number and type of staff working in the pharmacy may limit interpretation of the results and introduce a degree of bias. Also, it is possible that the timing of the study following the government announcement to make cuts in community pharmacy funding²⁹ could have had a negative impact on pharmacist motivation and recruitment to the study with pharmacists unwilling to undertake additional non-service tasks or research. Differences in the commissioning of access to PMs within England also may limit the findings as some services are commissioned by NHS England as Local Enhanced Services and others are commissioned locally by CCGs, with no standard service

specification stating the outcomes to be measured. Furthermore, the geographical restriction with data only collected in one city could limit application to other areas including those in remote locations, with different out-of-hours providers, and access to palliative care support in the community. Nevertheless this is the first published study to our knowledge to bring together Px data with stakeholder views and experiences on accessing PMs through community pharmacies in England and the methodology has enabled new insights into factors contributing to timely access.

CONCLUSION

The findings of this study suggest that legal prescribing errors may now have only a small impact on access to urgent PMs from community pharmacies compared to stock availability and supply chain factors. The CPs decisions about which PMs to stock in the pharmacy impact on timeliness of access with participation in a locally commissioned scheme or collaboration with local prescribers likely to improve access to PMs. There are likely to be advantages to GP practices working with local pharmacies to keep a small range of PMs available. Likewise, local integration and collaboration including CP integration into the primary healthcare team is important to ensure timely access to PMs. Improved communication between pharmacies and other HCPs around pharmacy opening times and cut-off times for same-day delivery of medicines could also support access to PMs. Moving forward, NHS England will be supporting development and integration of CP services into primary care through its Pharmacy Integration Fund³⁰ and this may also improve interprofessional communication and access to PMs.

Further studies are needed investigating the effect of EPS and referral patterns on access to PMs and the effect on patient's continuity of care at the end-of-life.

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Parts of this study have been previously presented at conferences and published as conference abstracts.

Author Contributions

EM implemented the study, designed and piloted data collection tools, monitored data collection, wrote the analysis plan, analysed the data, wrote and piloted interview schedule, transcribed interviews, developed the thematic framework and drafted and revised the manuscript. JDM and AB supervised EM in planning and undertaking the study including study design, development of data collection tools, data analysis, drafting and revision of the manuscript.

Dr Christina Wong (CW), piloted the interview schedule, supported the qualitative data analysis and provided workplace supervision for EM as part of the study. Statistical input provided by Dr Jon Silcock, University of Bradford.

Competing interests

EM received research funding from Pharmacy Research UK and Sheffield Teaching Hospitals NHS Foundation Trust as well as support from St Luke's Hospice, Sheffield; JDM and AB are employees of University of Bradford, all these organisations might have an interest in the submitted work – in the previous three years. EM is Treasurer of the Association of Supportive and Palliative Care Pharmacy. AB and JDM report grants from Pharmacy Research UK during the conduct of the study. EM, JDM, CW and AB are all pharmacists registered with the General Pharmaceutical Council.

Ethics approval

Ethical approval obtained 17th December 2015 from the Chair of the Biomedical, Natural, Physical and Health Sciences Ethics Panel, University of Bradford (approval reference E493).

Data sharing statement

No additional data are available.

Patient Involvement

The design of the study was based on EM's experience as a clinical pharmacist including discussions with patients and their families on accessing medicines. The research question was therefore derived from patients' and family carers' experience on accessing medicines towards the end-of-life. A Hospice Patient User Co-ordinator provided support with the customer survey based on their experience of conducting surveys. Furthermore, patients within a hospice day centre supported the piloting of the customer survey.

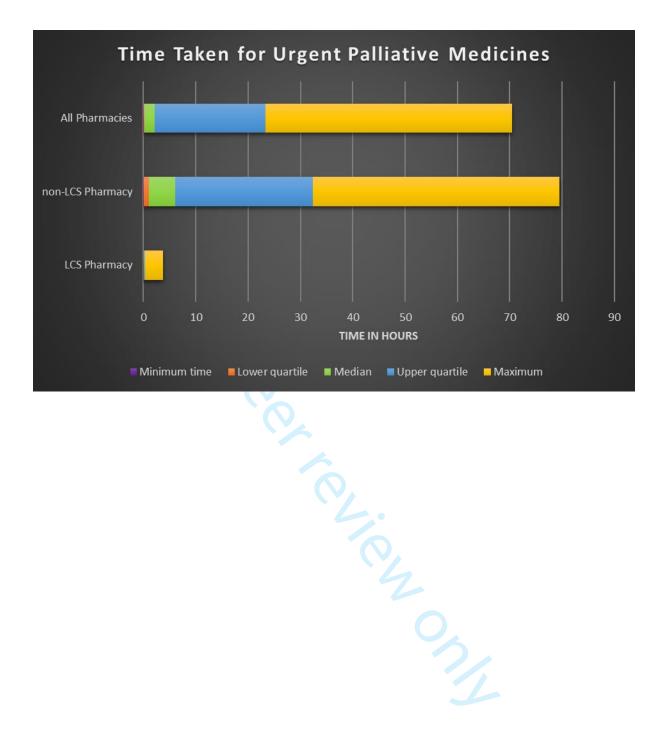
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SCHOLARONE™ Manuscripts Title: How timely is access to palliative care medicines in the community? A mixed methods study in a UK city

Elizabeth Miller¹, Julie D. Morgan², Alison Blenkinsopp²

- ¹ Sheffield Teaching Hospitals NHS Foundation Trust, Sheffield, UK
- ² University of Bradford, Bradford, UK

Corresponding author: elizabeth.miller@sth.nhs.uk

Pharmacy Department, Royal Hallamshire Hospital, Sheffield S10 2JF

Tel: 0114 2711900

ABSTRACT

Objective: To investigate timeliness of access to palliative medicines (PMs) from community pharmacies to inform palliative care service delivery.

Design: Mixed methods in two sequential phases: 1. Prospective audit of prescriptions and concurrent survey of patients/representatives collecting PMs from pharmacy; 2. Interviews with Community Pharmacists (CPs) and other healthcare professionals (HCPs).

Setting: Five community pharmacies in Sheffield, UK and healthcare professionals that deliver palliative care in that community.

Participants: Phase 1: Five CPs: two providing access to PMs within a Locally Commissioned Service (LCS) and three not in the LCS; 55 patients/representatives who completed the survey when accessing PMs; Phase 2: 16 HCPs, including 5 Phase 1 CPs, were interviewed.

Results: The prescription audit collected information on 75 prescriptions (75 patients) with 271 individual PMs; 55 patients/representatives (73%) completed the survey. Patients/representatives reported 73% of PMs were needed urgently. In 80% of cases patients/representatives received all PMs on the first pharmacy visit. One in five had to travel to more than one pharmacy to access PMs. The range of PMs stocked by pharmacies was the key facilitating factor. CPs reported practical issues causing difficulty keeping PMs in stock and playing a reactive role with palliative prescriptions. Confidentiality concerns were cited by other HCPs who were reluctant to share key patient information proactively with pharmacy teams. Inadequate information transfer, lack of CP integration into the care of palliative patients, and poor HCP knowledge of which pharmacies stock PMs meant patients and their families were not always able to access PMs promptly.

Conclusions: Consistent routine information transfer and integration of pharmacy teams in the care of palliative patients are needed to achieve timely access to PMs. Commissioners of PM access schemes should review and monitor access. HCPs need to be routinely made aware and reminded about the service and its locations.

Key Words: palliative care; community pharmacy services; pharmacists; prescriptions; interprofessional issues

Word Count 5302

Article Summary

Strengths and limitations of this study

This is the first published study to identify the relative impact of factors contributing to non-timely access to PMs.

This paper is the first in the UK to examine perspectives of different (HCPs) on factors supporting and hindering access to PMs.

The study is also novel in its examination of customer experience of accessing PMs and the survey achieved a high response rate.

The study is possibly limited by the low number of sites but adds value to the literature in terms of barriers that need to be considered if more timely access to PMs is to be more widely implemented.

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INTRODUCTION

Population aging and an increase in those dying with complex multi-morbidity will increase the need for palliative care with predictions suggesting End-of-Life Care (EOLC) provision in the community and care homes needs to double by 2040.[1] In England, government strategy aims to improve accessibility to palliative care in the community[2] in alignment with research suggesting most people would prefer a home death with 80% not changing their preference as their illness progresses;[3] however resource issues in primary care can make attaining this very challenging. In particular, accessing medicines for symptom control towards the end-of-life is imperative in controlling pain and distress and to prevent unnecessary hospital admissions. Difficulties in predicting those who may die within 12 months who require EOLC[4] can lead to an unpredictable yet urgent demand for medicines to relieve symptoms towards the end-of-life in the community.

For most patients in primary care the source of medicines is from their community pharmacy (retail pharmacy or "chemist shop"); however previous research and service audits show access to medicines such as injectable medicines used for symptom control towards the end-of-life, which are referred to in this study as Palliative Medicines (PMs), may not be as timely as patients and their families may need and wish.[5–9] Pharmacies cannot stock every possible PM; local formularies which provide lists of preferred medicines help to address this in the UK.[7,8] However knowledge on which PMs are listed in the formulary and those pharmacies holding stocks may be lacking among prescribers[7,8] which could lead to prescriptions being issued for 'non-formulary' items not on the approved local PM list and/or prescriptions being presented to pharmacies that do not routinely hold PMs.[5–12] Delays may also be caused by legal errors on prescriptions where the prescription does not comply with legislation necessitating the pharmacist making professional and ethical judgements in supporting patient care especially in the outof-hours period.[5,7,8] There is a suggestion that hand-written prescriptions may be particularly problematic due to higher prescription error rate and out-of-hours presentation[7,13,14] and they are still in use in the UK for home visits.[5,7,14]

Australian research on a proposed core set of PMs found that pharmacies stocked on average three out of the list of 12,[11] while a systems analysis in Ireland found that not stocking PMs in the pharmacy was the most likely factor leading to

delays[10] and this has also been found in the UK.[5,7,8,12] Reported contributory factors include: the unpredictable nature of PM prescription requests; national stock shortages; the prescription of PMs or strengths not on the recommended list; unlicensed medicines; errors on Controlled Drug (CD) prescriptions and the inability to contact the prescriber, for instance outside GP (family doctor) practice opening hours.[5,8,14]

Community pharmacies may take part in local or nationally commissioned services to support access to PMs in the community. In England, a Locally Commissioned Service (LCS) can be provided by the local clinical commissioning group (CCG) or a Local Enhanced Service (LES) can be commissioned by NHS England Area Teams in response to public need.[15] Such services differ across geographical regions and are not commissioned from all pharmacies, causing confusion for patients and their caregivers who are often involved in prescription collection and medicines management when a patient's condition deteriorates.[9,16–18] Furthermore, a lack of monitoring of PM availability against those prescribed both within the pharmacy and by the commissioning body could mean PMs are not available when needed. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK)

There is little research internationally on community pharmacists' (CPs) involvement in supporting timely access to PMs. Hence this study seeks to answer the question 'What barriers are encountered by community pharmacists in delivering timely access to palliative care medicines.' Due to the dearth of published research particularly in the context of community pharmacy services in England the aim of this study was to evaluate timely access to PMs in the community pharmacy setting and make recommendations to inform the commissioning of services and future practice. The objectives were to:

- determine the timeliness of access to PMs in the community;
- investigate the prevalence and nature of prescribing errors on prescriptions for PMs presented to community pharmacies and determine whether errors impact on access to urgent PMs;

- investigate processes for accessing PMs from pharmacies where a locally commissioned service (LCS) operates including referrals when PMs are not available:
- explore the views and experiences of CPs and other stakeholders on accessing PMs from community pharmacies.

METHODS

This study used mixed methods across two sequential phases (See Table 1 for study overview) conducted in Sheffield, UK. Participants in both phases gave informed consent before taking part. Ethical approval was obtained from the University of Bradford.

Table 1 Overview of study phases

Phase 1:

Audit of palliative prescriptions meeting inclusion criteria in participating pharmacies from May - October 2016

Customer survey for those collecting palliative prescriptions in participating pharmacies from May - October 2016

Phase 2:

Semi-structured face-to-face interviews with pharmacists participating in Phase 1 and other healthcare professionals involved in palliative care in the community from September 2016 - March 2017

Phase 1: Audit of PM supplies over six-month data collection period. Sheffield pharmacies were recruited through e-bulletin sent by the Local Pharmaceutical Committee (LPC), fax invitation to LCS PM pharmacies (19 of the 128 in the city), and verbal invitation at a local pharmacy practice development event. CPs expressing interest in taking part were given an information leaflet and consent form via email providing further information and the study inclusion criteria. Eligible CPs participated in the LCS or usually dispensed thirty or more PM prescriptions in a month based on NHS Digital prescription data for opioid analgesics and midazolam dispensed in pharmacies in the region.[19] Exclusion criteria were: i) pharmacists who had worked in the UK for less than 12 months (to ensure participants were familiar with UK and local community pharmacy services), and ii) if the company or

manager did not give permission for participation. None of the interested pharmacies had to be excluded based on these criteria.

A pragmatic approach to sample size was taken; the intention was to recruit up to 15 pharmacies however only five CPs consented to participate, partly due to the unexpectedly low level of PM prescriptions reported. Informed consent was obtained. EM personally visited each participating pharmacy to brief them on the project, data collection forms and answer any questions to enhance consistency of data collection.

Consenting pharmacies collected data on thirty consecutively presented prescriptions which contained medicines likely to be prescribed for EOLC patients using criteria developed by EM. Eligible prescriptions were for adults aged 18 years or over and included one or more of the following: a long acting oral or transdermal strong opioid co-prescribed with a short acting opioid; fast acting fentanyl product; prescription of subcutaneous or syringe pump PMs, specified unlicensed medicines used in palliative care, as well as any prescription issued by the palliative care team. Prescription data was collected for six months between May to October 2016. Pharmacy data collection forms were developed by EM and reviewed by JDM and piloted in one community pharmacy. The form recorded anonymised prescription data including: names of medications on the prescription; whether there was a legal or non-legal error on the prescription and further information on how that error was resolved including whether the Summary Care Record (SCR) was accessed to resolve an error. Legal and non-legal prescription errors were identified by the CPs and non-legal errors were classified by EM according to criteria within the PRACtICe study [20] Further details on non-legal errors were completed on a separate form to allow EM to verify the classification. Previous research suggested that delays may be caused by doctors prescribing products not on the community pharmacy PMs list[7,8] (e.g. midazolam 5mg/5ml prescribed when midazolam 10mg/2ml on stock list) hence where prescribers issued legally correct prescriptions for products not recommended on the LCS PM list these were classified as non-legal errors. Subcutaneous items in the audit were checked by EM against the LCS list to identify non-formulary items in both LCS and non-LCS pharmacies.

Prescriptions were classified as urgent when i) the survey respondent stated it was urgent, ii) they included anticipatory subcutaneous medicines and/or PMs to be given by a syringe pump or iii) they were from an out-of-hours provider. The date/time a prescription was received by the pharmacy and the date/time when it was ready for collection were recorded by pharmacy staff.

Survey of patients/representatives collecting PM prescriptions. The national pharmacy contracting organisation the Pharmaceutical Services Negotiating Committee (PSNC) Community Pharmacy Patient Questionnaire (CPPQ)[21] was used as a basis to develop a short customer survey of experiences of patients and their representatives of collecting PM prescriptions from the community pharmacy. Questions included the perceived urgency of the prescription, the customer's previous use of the pharmacy, whether they were the patient or the patient's representative, whether they were able to access all required PMs, whether they had been referred to the pharmacy (e.g. by another HCP) and whether they had to visit more than one pharmacy to access the PMs on the prescription. When patients/representatives indicated that not all items were available they were given the option of completing a free text section to explain how they intended to get these items. A free text section allowed respondents to record their answer to 'are there any things that could have been improved to make your visit better?' The customer survey was developed by EM with input from JDM, AB, a hospice service user coordinator and risk manager. It was piloted in one pharmacy, further refined and piloted with patients within a hospice day centre. Pharmacy teams were provided with a written briefing on how to introduce the survey to patients/representatives. Individuals collecting prescriptions for PMs were invited to participate by pharmacy support staff or the CP depending on the pharmacy. A unique number was used to match the customer survey to the pharmacy data form to allow verification of the data and assess any discrepancies. Patients/representatives not attending the pharmacy, e.g. home deliveries and care home residents, did not complete the customer survey.

Phase 2: Semi-structured interviews with CPs and other HCPs involved in care of palliative patients. EM conducted interviews with the five CPs participating in phase 1 and with a purposive sample of other HCPs involved in palliative care in the community including GPs, community specialist palliative care team, community

nurses, district nurses and intermediate care team members. HCPs were invited to participate via e-bulletin, email and through gatekeepers (practice managers and team leaders). Interviews were audio-recorded with consent and transcribed verbatim by EM. The interviews explored views and experiences of accessing PMs in the community; factors that supported or hindered access and their knowledge of the LCS. The interview schedule is available on request from the authors.

Data analysis

Prescription data were entered into and analysed using IBM SPSS® V.23 statistical software by EM. Frequencies and percentages were calculated for all categorical variables with mean and standard deviation calculated for time to process prescriptions. Crosstabs was used to check relationship between legal or clinical errors and prescription generation method.

Interview transcripts were read by EM for content familiarisation then annotated and coded manually using a priori themes from the study objectives. Following development of an analytical framework, two over-arching themes were then used to 'chart' the coded data: (1) timely access to PMs, (2) the community pharmacist's role in palliative care, using the Framework Method.[22] The framework was revised and iteratively refined with CW and AB against the coded interview transcripts with emergent themes and subthemes applied across the whole data set. Summaries of data were added within the framework to capture participants' views. Mapping and interpretation of findings compared similarities and contrasts between and across professional groups and was supported through discussion and reflection with AB and CW.

Data from both phases were then triangulated where two or more sources agreed or contrasted with each other to help explain the quantitative results of the study. Triangulation enhanced the validity and reliability of the results and enabled integration of the findings, such that it was possible to make recommendations for practice improvement and identify issues for service commissioners to consider.

Patient Involvement

The study was informed by research priorities in palliative care[23] and through EM's professional experience including discussion with patients and carers experiencing

medicines access problems following admission to a hospice. The customer survey tool was developed and piloted with patients in a hospice setting.

RESULTS

Participants in each phase of the study

Phase 1 CP audit: Participating pharmacies were diverse in that they included pharmacies classified as independent (having fewer than five branches) and multiple (having five or more branches); two provided access to PMs under an LCS and three did not. Pharmacy sites were a combination of high street/local parade of shops (3), and suburban (2) with both suburban pharmacies co-located with a GP practice. For pharmacies not consenting to take part, the main reason cited was small numbers of palliative care prescriptions dispensed in the pharmacy.

Customer survey: Customer surveys were completed against 55/75 CP audit forms; response rate 73.3%. Non-completion related primarily to home deliveries and care home prescriptions.

Phase 2 CP and other HCP Interviews: 16 individuals participated: CPs (5), GPs (3), Specialist Palliative Care Team (2), Community Nurses (5), and Intermediate Care Team (1). The five CPs were also involved in phase 1. Median interview durations were 51 minutes for CPs and 18.5 minutes for GPs and other HCPs.

Phase 1: Prescription characteristics

A total of 271 prescription items on 75 prescription forms was recorded (range 2 to 33 per pharmacy, median 14) over the 6-month audit period with a mean number of 3.6 prescription items per form. This included 68.3% (n = 185) of PMs identified as urgent, 49.8% (n = 135) containing subcutaneously administered PMs and 24.7% (n = 67) containing subcutaneously administered CDs. In 91.1% (n = 123) of cases, subcutaneous items were chosen from the LCS formulary list. Non-formulary choices were either different presentations of formulary items (5.9%, n = 8) or items not on the LCS list (3.0%, n = 4)). Varying strengths of midazolam ampoules accounted for 41.7% (n = 5) of non-formulary choices.

Prescriptions were computer-generated (n = 245, 90.4%) or hand-written (n = 22, 8.1%), with no prescriptions delivered electronically via the Electronic Prescription Service (EPS); missing data (n = 4, 1.5%). Most prescriptions were written by NHS GPs providing in-hours services (n = 233, 86%), with out-of-hours GPs (n = 33, 12.2%) or specialist palliative care team (n = 5, 1.8%) writing the remainder. There were no non-medical prescriber prescriptions within the sample. Prescriptions were presented to the pharmacy during GP opening hours (from 9-6pm Monday to Friday) (n = 176, 64.9%) or outside GP hours (evenings and weekends) (n = 77, 28.4%); missing data on 6.6% of forms (n = 18).

Phase 1: Prescription audit

Legal errors were present in 1.1% (n = 3) of prescription items; all of which were computer-generated, not specifying a dose on a controlled drug (CD) given via infusion. There were no legal errors on handwritten prescriptions. There was insufficient evidence of a difference between prescription generation method and legal errors (Fisher's Exact 2-sided test, p = 0.052). Other non-legal prescribing errors such as incomplete information, dose/ strength error, generic/ brand error, allergy, and quantity error occurred in 3.0% (n = 8) of items. Table 2 summarises prevalence of different medication problems on prescriptions and table 3 indicates types of prescribing errors using categories as in the PRACtICe study.[20]

Table 2: Prevalence of medication problems

Type of medication problem	Frequency of problem	Total number of prescription items	Percentage prevalence
Legal errors	3	271	1.1%
Prescribing errors (see table 3)	8	271	3.0%
Out of stock with supplier	15	271	0.4%
Non-formulary LCS item requested	12	135	8.9%

Table 3: Prescribing errors (n=271)

Type of prescribing error	Frequency	Percentage prevalence
Incomplete information on prescription	2	0.7%
Dose / strength error	2	0.7%
Generic / brand error	2	0.7%
Allergy ¹	1	0.4%
Quantity error	1	0.4%

¹ Allergy ascertained from patient or patient's medical record on pharmacy dispensing system

Phase 1: Time to access urgent palliative care medicines

Valid time data was available for 57.8% (n = 107) of 185 urgent items (n = 73 missing data; n = 5 excluded where PMs unavailable and prescription taken elsewhere and recorded as 0 minutes). Median time to process urgent PMs (time of prescription receipt to time of complete supply of PMs) was 2 hours (10 minutes in LCS pharmacies and 5 hours in non-LCS pharmacies). The maximum time to process urgent PMs was 3 hours and 39 minutes within LCS pharmacies, and 47 hours and 15 minutes within non-LCS pharmacies (see figure 1).

The median time taken to access urgent medications (n = 107) between pharmacies participating in the LCS and pharmacies not participating in the service was significantly different (independent samples median test p = 0.002 at 95% confidence level); with pharmacies not participating in the LCS taking significantly longer than pharmacies in the LCS.

Figure 1: Time taken to access urgent palliative medicines from community pharmacies (see separate file)

Legal errors had minimal effect on access as all urgent PMs with legal errors were available within 30 minutes of presentation. Legal errors were resolved by: contacting the nursing home to specify the dose to be given on a prescription for PMs via a syringe pump using a community medicines administration record, using the pharmacy Patient Medication Record (PMR) to access information on a previously issued prescription, and contacting the prescriber. The SCR was not used to resolve errors in the prescription audit sample.

Phase 1: Customer Survey

Survey responses showed that representatives collected PMs on behalf of the patient (65.5%); for both themselves and the patient (1.8%); and patients collected their own PMs (32.7%); 72.9% of surveys overall indicating the prescription included urgent item(s). All cases for urgent subcutaneous medications were collected by a representative on behalf of a patient. In 42.6% of cases the patient attended their usual pharmacy. Patients/representatives also indicated the pharmacy was: convenient (14.8%); one of several pharmacies used (20.4%), or that they had been referred to the pharmacy for the medications (21.8%).

In 80% of cases patients/representatives received all medications against the prescription at the first pharmacy they visited. In 20% (11/55) one or more items on the prescription was not available, in five of these the item(s) were urgent. Free text sections were completed for six of the 11 cases of unavailable items. Four indicated they would return to collect the item from the pharmacy and two said they would try another pharmacy to obtain the items.

Thirteen respondents made additional comments on whether their experience could have been better. Comments were mostly positive: six indicated 'no', 'none', 'no fine' or similar phrase; five made comments on the staff or service: 'friendly services under difficult circumstances', 'no - staff really friendly and helpful, service was quick and efficient', 'Nothing – excellent and quick service'; and one explained 'nothing much that would make it better, but I phone in advance to make sure my items are in stock'. One respondent requested to 'keep a stock of all required items'.

Overall one in five patients/representatives had to go to more than one pharmacy to get urgently needed PMs, increasing to one in three for urgent subcutaneous injection prescription items. One in every two patients/representatives referred to the pharmacy by another HCP had to go to more than one pharmacy. Data from the prescription audit and customer survey were triangulated to verify the validity of the information in phase 1.

Phase 2: Interview findings

Findings from the interviews are presented in four sections: timely access; challenges; knowledge of LCS; and communication and collaboration. The findings are illustrated by verbatim quotes from the interviews where appropriate.

Timely access

Anticipating need and forward planning were key themes to ensure timely access to PMs.

... I could go in now and say, 'I need these drugs' (and the CP might say) 'Oh I can get them in for 11 o'clock tomorrow morning' [exasperated laugh] it's like that's not really very helpful, I need them now (HCP7, Community Healthcare Professional)

Community nurses and palliative care team staff reported strategies to enhance access including conducting an end of week check and balance for those on syringe pumps to ensure sufficient stock for over the weekend when fewer staff were available. Specialist palliative care team staff also described making do with the medicines already available in the house and then ordering medication for the next day. CPs perceived that patients/HCPs phoning ahead for large quantities would be helpful. Insufficient quantities of PMs could adversely impact patient symptom management and had consequences for staff resources, however not all situations could be taken into consideration.

... a GP won't prescribe a syringe driver ahead of time...we are always...having to do it now not in a more considered way (HCP4, Community Healthcare Professional)

... I remember having to go [to the pharmacy] in the middle of doing a [syringe] driver because there weren't enough drugs (HCP8, Community Healthcare Professional)

The local CCG had implemented a template on the GP prescribing system to provide a 'suite' of PMs according to local last days of life algorithms which included some of the injectable medicines listed on the LCS formulary. Even so, in phase 1 several 'non-formulary' medications not on the local CCG PM list were prescribed and in phase 2 CPs in LCS pharmacies described non-compliance with the local formulary as a reason for a lack of timely access to PMs.

We've got three different strengths of oxycodone injection, and they [GPs] prescribe all three, and you might not have one, you might have the other...it's just so frustrating... (P4, Community Pharmacist)

The big problem is midazolam...so many strengths...volumes of ampoules...the GPs just pick one. (P5, Community Pharmacist)

Challenges

CPs described practical issues in supplying PMs for example: stock ordering processes including: timing of deliveries and the inability to return CD items to

suppliers due to legal restrictions; CD cabinet size (to meet UK legal requirements for storage); and quantities on prescriptions.

We don't have an ability to be able to keep a lot [controlled drugs] and so we have a particular issue with the quantities that they write on the prescriptions sometimes which can impact on the next patient (P4, Community Pharmacist)

We've only got very small CD cabinets...the more controlled drugs you keep the more issues you are going to have (P1, Community Pharmacist)

Furthermore, patient records and charts to check opioid dose changes and syringe pumps were often not accessible to CPs.

With regards to changing doses or monitoring, I think that would be difficult for a community pharmacist. We have the summary care [record]... with palliative care the dose can change, you've got the pink card...sometimes we see [the pink card] and sometimes we don't (P2, Community Pharmacist)

... they [CPs] don't get the pink card [syringe pump chart] they simply get the prescription (HCP3, GP)

I might only need...extra diamorphine...whereas I might be using midazolam and haloperidol...but I've still got a supply of those, so they [CPs] don't always know what's in the [syringe] driver (HCP4, Community Healthcare Professional)

Knowledge of LCS

HCPs had little knowledge of either the LCS or the pharmacies commissioned to provide it but knew which pharmacies were likely to keep some PMs in stock.

I don't know who's commissioned we just basically know which ones we go to that are more likely to have it. (HCP4, Community Healthcare Professional)

... relatives who are running right left and centre trying to get hold of these meds...there is a commissioned service...but we don't know who they are. (HCP1, Community Healthcare Professional)

GPs generally thought that all pharmacies kept some injectable PMs in stock but said they might ring in advance to check the medication was available if a supply was needed urgently. Non-LCS pharmacist providers knew of the service and how to refer a patient/carer if they did not have the requested medication available. Usually they would phone ahead to the pharmacy to check the medication was available before making a referral. Being able to make a referral depended on whether the carer had access to a car.

... if they haven't got a car it's sometimes a bit tricky (P1, Community Pharmacist)

Communication and Collaboration

Participants described strategies to enhance access to PMs. Two non-LCS pharmacies had worked with GP practices to discuss and agree to stock a subset of the LCS PMs; they had similar response times to pharmacies in the LCS.

... we went to them [GPs] and said, what are the most common drugs you would prescribe in palliative care...they [GPs] came back with a list...so we would try to keep the stock in for what they specified (P3, Community Pharmacist)

CPs reported that some patients/carers contacted the pharmacy when they ordered a prescription for a CD that might not be stocked. Community and specialist palliative care team staff described how they would suggest the pharmacist kept sufficient stock when they had someone on a syringe pump or large quantities of injectable medications. There appeared to be some examples where excellent communication and collaboration existed between GPs, HCPs and CPs which resulted in more timely access for PMs.

... one GP...rang us and said well what have you got in stock and what can you get, which I found really, really useful because as the

prescription came in the stock came in and this thing was completely seamless (P3, Community Pharmacist)

When we were down at [previous community nurse location] ...there was a pharmacy next door so...if we had any quick questions, we would go and talk to them...they were more like part of the team (HCP7, Community Healthcare Professional)

However, concerns around patient confidentiality by GPs and other HCPs meant that more often this information was not shared with the pharmacy team in advance of receiving the prescription.

... but you're limited by what you can tell them [pharmacists] obviously from a confidentiality point of view... (HCP11, Community Healthcare Professional)

We don't communicate with them [community pharmacist] what the problem with the patient is we just prescribe the drugs... sometimes they can obviously work it out. (HCP3, GP)

I do have some slight reservations about them [pharmacists] knowing all those ins and outs...I'm not sure how wide that circle is in there [pharmacy]...I'd prefer it ...on just a case by case basis...to an identified clinician... (HCP10, GP)

DISCUSSION

The sequential use of mixed methods to firstly quantify the "problem" and then qualitative methods to provide context to the barriers to timely supply of PMs generated new insights into a longstanding problem. Timeliness of access was found to primarily relate to a mismatch between medicines stocks held by CPs and the PMs that GPs prescribed. Legal errors played a much smaller role and had little impact on access to PMs in this study. Stock availability as a significant factor to support timely access has also been seen in previous studies.[5,8,10,11]

Study results appear to indicate a low prevalence of legal errors on palliative care prescriptions taking into account the high volume of controlled drugs prescribed.

Previous studies suggest legal errors on prescriptions can range from less than 1% up to approximately 1.9% [14,20,24–28] though data does not specifically focus on palliative care prescriptions. Legal errors in the current study arose on computergenerated prescriptions for controlled drugs administered by a syringe pump. The proportion of errors is lower than may be anticipated given that 42% of these prescriptions were for controlled drugs; the primary care organisation prescribing template for PMs may have impacted positively, minimising the number of errors.

Our findings of differential time to access PMs between community pharmacies participating in the LCS for PMs and non-participants, indicates that a local service can enhance access. Those pharmacies working with local GP practices to keep a small agreed range of PMs in stock had similar access times to those within the LCS, suggesting that such collaboration can also support more timely access and improve patient and carer experience. Such wider collaboration has been advocated within national policy drivers and enables greater integration of pharmacy teams in improving patient care. [29-32] Some HCPs considered CPs "part of the team" but others saw them only in their "supply" role. Together with concerns about confidentiality this prevented many HCPs from communicating with CPs about available stock or giving advance warning to allow stock to be obtained in a more timely way. A particular concern seemed to be who else, in addition to the pharmacist, might have access to sensitive information. Commissioners could remind primary care staff of community pharmacy ethical and information governance practices to correct any misunderstandings. They could also encourage HCPs to be proactive in checking stock availability with the patient's usual CP.

Palliative patients often rely on family members and friends to support them with managing their medication especially towards the end-of-life.[9,16-18] Our findings show that some families obtain urgently required medicines from a pharmacy different than the one that usually supplied the patient's medicines. It is unclear what effect these changes in continuity of care between pharmacies might have towards the end-of-life, but this study shows that some CPs enhance access by calling other pharmacies to ascertain they have the PMs in stock. A potential solution could be through CPs having read and write access to SCR allowing them, whether they are the patient's regular, out-of-hours or LCS pharmacy, to record patient care scenarios

to ensure safe, continuity of care of PMs. Variable accessibility and difficulties in use of SCR by CPs[33] may suggest wider access to patient records is required.

One in five patients/representatives accessing PMs had to go to more than one pharmacy. This is the first study to quantify the number of patients/representatives who had to do so, with associated inconvenience, wasted time and stress. This finding could be explained by a lack of awareness of the LCS since this is not advertised to the public and there was also low awareness amongst HCPs and GPs in the interviews. Monitoring of LCS/LES services by commissioners may not always be effective. One audit found only one of nineteen pharmacies held all PMs on the formulary list and some CPs were not aware that the scheme was active. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK) There was also evidence in phase 1 of prescriptions being written for items not on the LCS list which would not usually be stocked in the pharmacies. Commissioners' service audits could investigate referral patterns from pharmacies not within the LCS, and monitoring of LCS pharmacies may improve practice and caregivers' experience. Commissioners could also act to improve awareness of LCS among local HCPs.

Some limitations affect the interpretation/generalisability of the findings of this study. The small sample of participating pharmacies, missing data, reliance on CPs to identify prescriptions and confounding factors such as time of day, number and type of staff working in the pharmacy may limit interpretation of the results and introduce a degree of bias. Differences in the commissioning of access to PMs within England also may limit the findings as there is no standard service specification stating the outcomes to be measured. Furthermore, the geographical restriction with data only collected in one city could limit application to other areas including those in remote locations, with different out-of-hours providers, and access to palliative care support in the community. Nevertheless, the study adds value to the literature in terms of barriers that need to be considered if more timely access to PMs is to be more widely implemented and the methodology has enabled new insights into factors contributing to timely access.

CONCLUSION

The findings of this study suggest that legal prescribing errors may now have a smaller impact on access to urgent PMs from community pharmacies compared to mismatches between stock availability and PMs prescribed. Both participation in a LCS or collaboration with local prescribers are likely to improve access to PMs.

Recommendations for future commissioning and practice:

Commissioners should:

- i) encourage GP practices to work with local pharmacies to keep a small range of PMs available
- ii) remind HCPs of the ethical and information governance requirements of community pharmacies and encourage early contact to check stock availability
- iii) involve patients and the public in designing audits of LCS.

Community pharmacies should improve their communication with HCPs around pharmacy opening times and cut-off times for same-day delivery of medicines.

Moving forward, NHS England will be supporting development and integration of CP services into primary care through its Pharmacy Integration Fund[34] and this may also improve interprofessional communication and access to PMs.

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Parts of this study have been previously presented at conferences and published as conference abstracts.

Author Contributions

EM implemented the study, designed and piloted data collection tools, monitored data collection, wrote the analysis plan, analysed the data, wrote and piloted interview schedule, transcribed interviews, developed the thematic framework and drafted and revised the manuscript. JDM and AB supervised EM in planning and undertaking the study including study design, development of data collection tools, data analysis, drafting and revision of the manuscript.

Competing interests

EM received research funding from Pharmacy Research UK and Sheffield Teaching Hospitals NHS Foundation Trust as well as support from St Luke's Hospice, Sheffield; JDM and AB are employees of University of Bradford, all these organisations might have an interest in the submitted work – in the previous three years. EM is Treasurer of the Association of Supportive and Palliative Care Pharmacy. AB and JDM report grants from Pharmacy Research UK during the conduct of the study. EM, JDM, and AB are all pharmacists registered with the General Pharmaceutical Council.

Ethics approval

Ethical approval obtained 17th December 2015 from the Chair of the Biomedical, Natural, Physical and Health Sciences Ethics Panel, University of Bradford (approval reference E493).

Data sharing statement

Protocol and data collection forms are available upon request from the corresponding author.

Patient Involvement

The design of the study was based on EM's experience as a clinical pharmacist including discussions with patients and their families on accessing medicines. The research question was therefore derived from patients' and family carers' experience on accessing medicines towards the end-of-life. A Hospice Service User Coordinator provided support with the customer survey based on their experience of conducting surveys. Furthermore, patients within a hospice day centre supported the piloting of the customer survey.

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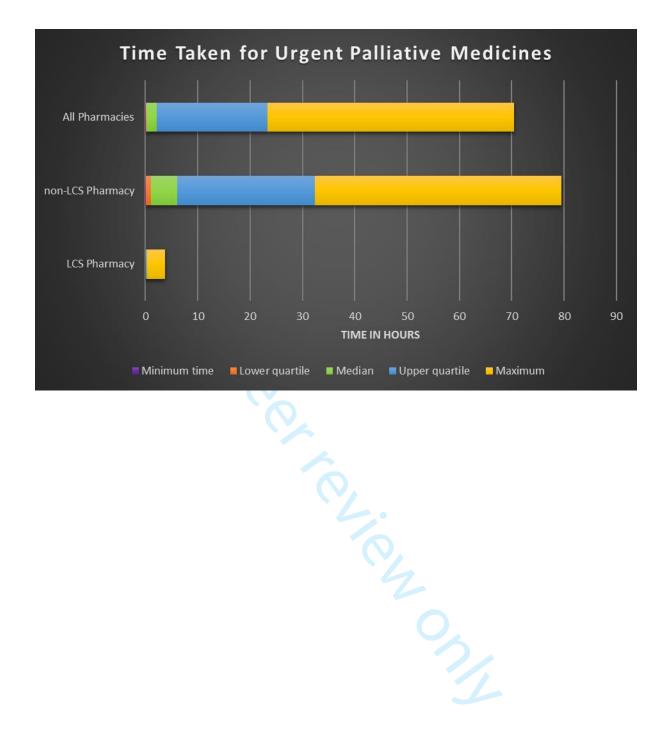
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How timely is access to palliative care medicines in the community? A mixed methods study in a UK city

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SCHOLARONE™ Manuscripts Title: How timely is access to palliative care medicines in the community? A mixed methods study in a UK city

Elizabeth Miller¹, Julie D. Morgan², Alison Blenkinsopp²

- ¹ Sheffield Teaching Hospitals NHS Foundation Trust, Sheffield, UK
- ² University of Bradford, Bradford, UK

Corresponding author: elizabeth.miller@sth.nhs.uk

Pharmacy Department, Royal Hallamshire Hospital, Sheffield S10 2JF

Tel: 0114 2711900

ABSTRACT

Objective: To investigate timely access to palliative medicines/drugs (PMs) from community pharmacies to inform palliative care service delivery.

Design: Mixed methods in two sequential phases: 1. Prospective audit of prescriptions and concurrent survey of patients/representatives collecting PMs from pharmacy; 2. Interviews with community pharmacists (CPs) and other healthcare professionals (HCPs).

Setting: Five community pharmacies in Sheffield, UK and healthcare professionals that deliver palliative care in that community.

Participants: Phase 1: Five CPs: two providing access to PMs within a locally commissioned service (LCS) and three not in the LCS; 55 patients/representatives who completed the survey when accessing PMs; Phase 2: 16 HCPs, including five Phase 1 CPs, were interviewed.

Results: The prescription audit collected information on 75 prescriptions (75 patients) with 271 individual PMs; 55 patients/representatives (73%) completed the survey. Patients/representatives reported 73% of PMs were needed urgently. In 80% of cases patients/representatives received all PMs on the first pharmacy visit. One in five had to travel to more than one pharmacy to access PMs. The range of PMs stocked by pharmacies was the key facilitating factor. CPs reported practical issues causing difficulty keeping PMs in stock and playing a reactive role with palliative prescriptions. Confidentiality concerns were cited by other HCPs who were reluctant to share key patient information proactively with pharmacy teams. Inadequate information transfer, lack of CP integration into the care of palliative patients, and poor HCP knowledge of which pharmacies stock PMs meant patients and their families were not always able to access PMs promptly.

Conclusions: Consistent routine information transfer and integration of pharmacy teams in the care of palliative patients are needed to achieve timely access to PMs. Commissioners of PM access schemes should review and monitor access. HCPs need to be routinely made aware and reminded about the service and its locations.

Key Words: palliative care; community pharmacy services; pharmacists; prescriptions; interprofessional issues

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Article Summary

Strengths and limitations of this study

This is the first published study to identify the relative impact of factors contributing to non-timely access to palliative medicines/drugs.

This paper is the first in the UK to examine perspectives of different healthcare professionals (HCPs) on factors supporting and hindering access to palliative medicines/drugs.

The study is also novel in its examination of customer experience of accessing palliative medicines/drugs and the survey achieved a high response rate.

The study is possibly limited by the low number of sites but adds value to the literature in terms of barriers that need to be considered if more timely access to palliative medicines/drugs is to be more widely implemented.

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INTRODUCTION

Palliative care is a holistic approach that seeks to improve the quality of life of patients with life-limiting or life-threatening illnesses.[1] Population aging together with an increase in those dying with cancer and non-communicable diseases will increase the need for palliative care at the end-of-life[2] with predictions suggesting end-of-life care provision in the community and care homes needs to double by 2040.[3] Whilst the phrase 'end-of-life' is not precisely defined, it is commonly used in United Kingdom (UK) policy and professional guidance to refer to the final year of life [4]. Nevertheless, for many people, end-of-life care will encompass a much shorter timescale and timely access to medicines for pain and symptom management, which are referred to in this study as palliative medicines/drugs (PMs), will be a crucial aspect of palliative care service delivery.[4]

For most patients in primary care the source of medicines is from their community pharmacy (retail pharmacy or "chemist shop"); however previous research and service audits show access to medicines such as injectable medicines used for symptom control in palliative care towards the end-of-life may not be as timely as patients and their families may need and wish.[5–9] Pharmacies cannot stock every possible PM; local formularies which provide lists of preferred medicines to support symptom management towards the end-of-life help address this in the UK.[7,8] However knowledge on which PMs are listed in the formulary and those pharmacies holding stocks may be lacking among prescribers[7,8] which could lead to prescriptions being issued for 'non-formulary' items not on the local list and/or prescriptions being presented to pharmacies that do not routinely hold PMs.[5–12] Delays may also be caused by legal errors on controlled drug (CD) prescriptions that do not comply with UK government legislation necessitating the pharmacist making professional and ethical judgements in supporting patient care especially in the outof-hours period.[5,7,8] There is a suggestion that hand-written prescriptions may be particularly problematic due to higher prescription error rate and out-of-hours presentation[7,13,14] and they are still in use in the UK for home visits.[5,7,14]

Australian research on a proposed core set of PMs found that pharmacies stocked on average three out of the list of 12,[11] while a systems analysis in Ireland found that not stocking PMs in the pharmacy was the most likely factor leading to delays[10] and this has also been found in the UK.[5,7,8,12] Reported contributory

factors include: the unpredictable nature of PM prescription requests; national stock shortages; the prescription of PMs or strengths not on the recommended list; unlicensed medicines; errors on CD prescriptions and the inability to contact the prescriber, for instance outside GP (family doctor) practice opening hours.[5,8,14]

Community pharmacies may take part in local or nationally commissioned services to support access to PMs in the community. In England, a locally commissioned service (LCS) can be provided by the local clinical commissioning group (CCG) or a local enhanced service (LES) can be commissioned by National Health Service (NHS) England Area Teams in response to public need.[15] Such services differ across geographical regions and are not commissioned from all pharmacies, causing confusion for patients and their caregivers who are often involved in prescription collection and medicines management when a patient's condition deteriorates.[9,16–18] Furthermore, a lack of monitoring of PM availability against those prescribed both within the pharmacy and by the commissioning body could mean PMs are not available when needed. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK)

There is little research internationally on community pharmacists' (CPs) involvement in supporting timely access to PMs. Hence this study seeks to answer the question 'What barriers are encountered by community pharmacists in delivering timely access to palliative care medicines.' Due to the dearth of published research particularly in the context of community pharmacy services in England the aim of this study was to evaluate timely access to PMs in the community pharmacy setting and make recommendations to inform the commissioning of services and future practice. The objectives were to:

- determine the timeliness of access to PMs in the community;
- investigate the prevalence and nature of prescribing errors on prescriptions for PMs presented to community pharmacies and determine whether errors impact on access to urgent PMs;
- investigate processes for accessing PMs from pharmacies where a locally commissioned service (LCS) operates including referrals when PMs are not available;

 explore the views and experiences of CPs and other stakeholders on accessing PMs from community pharmacies.

METHODS

This study used mixed methods across two sequential phases (See Table 1 for study overview) conducted in Sheffield, UK. Participants in both phases gave informed consent before taking part. Ethical approval was obtained from the University of Bradford.

Table 1 Overview of study phases

Phase 1:

Audit of palliative prescriptions meeting inclusion criteria in participating pharmacies from May - October 2016

Customer survey for those collecting palliative prescriptions in participating pharmacies from May - October 2016

Phase 2:

Semi-structured face-to-face interviews with pharmacists participating in Phase 1 and other healthcare professionals involved in palliative care in the community from September 2016 - March 2017

Phase 1: Audit of PM supplies over six-month data collection period. Sheffield pharmacies were recruited through e-bulletin sent by the Local Pharmaceutical Committee (LPC), fax invitation to LCS PM pharmacies (19 of the 128 in the city), and verbal invitation at a local pharmacy practice development event. CPs expressing interest in taking part were given an information leaflet and consent form via email providing further information and the study inclusion criteria. Eligible CPs participated in the LCS or usually dispensed thirty or more PM prescriptions in a month based on NHS Digital prescription data for opioid analgesics and midazolam dispensed in pharmacies in the region.[19] Exclusion criteria were: i) pharmacists who had worked in the UK for less than 12 months (to ensure participants were familiar with UK and local community pharmacy services), and ii) if the company or manager did not give permission for participation. None of the interested pharmacies had to be excluded based on these criteria.

A pragmatic approach to sample size was taken; the intention was to recruit up to 15 pharmacies however only five CPs consented to participate, partly due to the unexpectedly low level of PM prescriptions reported. Informed consent was obtained. EM personally visited each participating pharmacy to brief them on the project, data collection forms and answer any questions to enhance consistency of data collection.

Consenting pharmacies collected data on thirty consecutively presented prescriptions which contained medicines likely to be prescribed for palliative care patients using criteria developed by EM. Eligible prescriptions were for adults aged 18 years or over and included one or more of the following: a long acting oral or transdermal strong opioid co-prescribed with a short acting opioid; fast acting fentanyl product; prescription of subcutaneous or syringe pump PMs, specified unlicensed medicines used in palliative care, as well as any prescription issued by the palliative care team. Prescription data was collected for six months between May to October 2016. Pharmacy data collection forms were developed by EM and reviewed by JDM and piloted in one community pharmacy. The form recorded anonymised prescription data including: names of medications on the prescription; whether there was a legal or non-legal error on a controlled drug prescription and further information on how that error was resolved including whether the patient's summary care record (SCR) was accessed to resolve an error. Legal and non-legal prescription errors were identified by the CPs and non-legal errors were classified by EM according to criteria within the PRACtICe study.[20] Further details on non-legal errors were completed on a separate form to allow EM to verify the classification. Previous research suggested that delays may be caused by doctors prescribing products not on the community pharmacy PM list[7,8] (e.g. midazolam 5mg/5ml prescribed when midazolam 10mg/2ml on list) hence where prescribers issued legally correct prescriptions for products not recommended on the LCS PM list these were classified as non-legal errors. Subcutaneous items in the audit were checked by EM against the LCS list to identify non-formulary items in both LCS and non-LCS pharmacies.

Prescriptions were classified as urgent when i) the survey respondent stated it was urgent, ii) they included anticipatory subcutaneous medicines and/or PMs to be given by a syringe pump or iii) they were from an out-of-hours provider. The date/time a

prescription was received by the pharmacy and the date/time when it was ready for collection were recorded by pharmacy staff.

Survey of patients/representatives collecting PM prescriptions. The national pharmacy contracting organisation the Pharmaceutical Services Negotiating Committee (PSNC) community pharmacy patient questionnaire (CPPQ)[21] was used as a basis to develop a short customer survey of experiences of patients and their representatives of collecting PM prescriptions from the community pharmacy. Questions included the perceived urgency of the prescription, the customer's previous use of the pharmacy, whether they were the patient or the patient's representative, whether they were able to access all required PMs, whether they had been referred to the pharmacy (e.g. by another HCP) and whether they had to visit more than one pharmacy to access the PMs on the prescription. When patients/representatives indicated that not all items were available they were given the option of completing a free text section to explain how they intended to get these items. A free text section allowed respondents to record their answer to 'are there any things that could have been improved to make your visit better?' The customer survey was developed by EM with input from JDM, AB, a hospice service user coordinator and risk manager. It was piloted in one pharmacy, further refined and piloted with patients within a hospice day centre. Pharmacy teams were provided with a written briefing on how to introduce the survey to patients/representatives. Individuals collecting prescriptions for PMs were invited to participate by pharmacy support staff or the CP depending on the pharmacy. A unique number was used to match the customer survey to the pharmacy data form to allow verification of the data and assess any discrepancies. Patients/representatives not attending the pharmacy, e.g. home deliveries and care home residents, did not complete the customer survey.

Phase 2: Semi-structured interviews with CPs and other HCPs involved in care of palliative patients. EM conducted interviews with the five CPs participating in phase 1 and with a purposive sample of other HCPs involved in palliative care in the community including GPs, community specialist palliative care team, community nurses, district nurses and intermediate care team members. HCPs were invited to participate via e-bulletin, email and through gatekeepers (practice managers and team leaders). Interviews were audio-recorded with consent and transcribed

verbatim by EM. The interviews explored views and experiences of accessing PMs in the community; factors that supported or hindered access and their knowledge of the LCS. The interview schedule is available on request from the authors.

Data analysis

Prescription data were entered into and analysed using IBM SPSS® V.23 statistical software by EM. Frequencies and percentages were calculated for all categorical variables with mean and standard deviation calculated for time to process prescriptions. Crosstabs was used to check relationship between legal or clinical errors and prescription generation method.

Interview transcripts were read by EM for content familiarisation then annotated and coded manually using a priori themes from the study objectives. Following development of an analytical framework, two over-arching themes were then used to 'chart' the coded data: (1) timely access to PMs, (2) the community pharmacist's role in palliative care, using the Framework Method.[22] The framework was revised and iteratively refined with CW and AB against the coded interview transcripts with emergent themes and subthemes applied across the whole data set. Summaries of data were added within the framework to capture participants' views. Mapping and interpretation of findings compared similarities and contrasts between and across professional groups and was supported through discussion and reflection with AB and CW.

Data from both phases were then triangulated where two or more sources agreed or contrasted with each other to help explain the quantitative results of the study. Triangulation enhanced the validity and reliability of the results and enabled integration of the findings, such that it was possible to make recommendations for practice improvement and identify issues for service commissioners to consider.

Patient Involvement

The study was informed by research priorities in palliative care[23] and through EM's professional experience including discussion with patients and carers experiencing medicines access problems following admission to a hospice. The customer survey tool was developed and piloted with patients in a hospice setting.

RESULTS

Participants in each phase of the study

Phase 1 CP audit: Participating pharmacies were diverse in that they included pharmacies classified as independent (having fewer than five branches) and multiple (having five or more branches); two provided access to PMs under an LCS and three did not. Pharmacy sites were a combination of high street/local parade of shops (3), and suburban (2) with both suburban pharmacies co-located with a GP practice. For pharmacies not consenting to take part, the main reason cited was small numbers of palliative care prescriptions dispensed in the pharmacy.

Customer survey: Customer surveys were completed against 55/75 CP audit forms; response rate 73.3%. Non-completion related primarily to home deliveries and care home prescriptions.

Phase 2 CP and other HCP Interviews: 16 individuals participated: CPs (5), GPs (3), Specialist Palliative Care Team (2), Community Nurses (5), and Intermediate Care Team (1). The five CPs were also involved in phase 1. Median interview durations were 51 minutes for CPs and 18.5 minutes for GPs and other HCPs.

Phase 1: Prescription characteristics

A total of 271 prescription items on 75 prescription forms was recorded (range 2 to 33 per pharmacy, median 14) over the 6-month audit period with a mean number of 3.6 prescription items per form. This included 68.3% (n = 185) of PMs identified as urgent, 49.8% (n = 135) containing subcutaneously administered PMs and 24.7% (n = 67) containing subcutaneously administered CDs. In 91.1% (n = 123) of cases, subcutaneous items were chosen from the LCS formulary list. Non-formulary choices were either different presentations of formulary items (5.9%, n = 8) or items not on the LCS list (3.0%, n = 4)). Varying strengths of midazolam ampoules accounted for 41.7% (n = 5) of non-formulary choices.

Prescriptions were computer-generated (n = 245, 90.4%) or hand-written (n = 22, 8.1%), with no prescriptions delivered electronically via the electronic prescription service (EPS); missing data (n = 4, 1.5%). Most prescriptions were written by NHS GPs providing in-hours services (n = 233, 86%), with out-of-hours GPs (n = 33, 12.2%) or specialist palliative care team (n = 5, 1.8%) writing the remainder. There were no non-medical prescriber prescriptions within the sample. Prescriptions were presented to the pharmacy during GP opening hours (from 9-6pm Monday to Friday) (n = 176, 64.9%) or outside GP hours (evenings and weekends) (n = 77, 28.4%); missing data on 6.6% of forms (n = 18).

Phase 1: Prescription audit

Legal errors were present in 1.1% (n = 3) of prescription items; all of which were computer-generated, not specifying a dose on a CD given via infusion. There were no legal errors on handwritten prescriptions. There was insufficient evidence of a difference between prescription generation method and legal errors (Fisher's Exact 2-sided test, p = 0.052). Other non-legal prescribing errors such as incomplete information, dose/ strength error, generic/ brand error, allergy, and quantity error occurred in 3.0% (n = 8) of items. Table 2 summarises prevalence of different medication problems on prescriptions and table 3 indicates types of prescribing errors using categories as in the PRACtlCe study.[20]

Table 2: Prevalence of medication problems

Type of medication problem	Frequency of problem	Total number of prescription items	Percentage prevalence
Legal errors	3	271	1.1%
Prescribing errors (see table 3)	8	271	3.0%
Out of stock with supplier	15	271	0.4%
Non-formulary LCS item requested	12	135	8.9%

Table 3: Prescribing errors (n=271)

Type of prescribing error	Frequency	Percentage prevalence
Incomplete information on prescription	2	0.7%
Dose / strength error	2	0.7%
Generic / brand error	2	0.7%
Allergy ¹	1	0.4%
Quantity error	1	0.4%

¹ Allergy ascertained from patient or patient's medical record on pharmacy dispensing system

Phase 1: Time to access urgent palliative care medicines

Valid time data was available for 57.8% (n = 107) of 185 urgent items (n = 73 missing data; n = 5 excluded where PMs unavailable and prescription taken elsewhere and recorded as 0 minutes). Median time to process urgent PMs (time of prescription receipt to time of complete supply of PMs) was 2 hours (10 minutes in LCS pharmacies and 5 hours in non-LCS pharmacies). The maximum time to process urgent PMs was 3 hours and 39 minutes within LCS pharmacies, and 47 hours and 15 minutes within non-LCS pharmacies (see figure 1).

The median time taken to access urgent medications (n = 107) between pharmacies participating in the LCS and pharmacies not participating in the service was significantly different (independent samples median test p = 0.002 at 95% confidence level); with pharmacies not participating in the LCS taking significantly longer than pharmacies in the LCS.

Figure 1: Time taken to access urgent palliative medicines from community pharmacies (see separate file)

Legal errors had minimal effect on access as all urgent PMs with legal errors were available within 30 minutes of presentation. Legal errors were resolved by: contacting the nursing home to specify the dose to be given on a prescription for PMs via a syringe pump using a community medicines administration record, using the pharmacy patient medication record to access information on a previously issued prescription, and contacting the prescriber. The patient's summary care record (SCR) was not used to resolve errors in the prescription audit sample.

Phase 1: Customer Survey

Survey responses showed that representatives collected PMs on behalf of the patient (65.5%); for both themselves and the patient (1.8%); and patients collected their own PMs (32.7%); 72.9% of surveys overall indicating the prescription included urgent item(s). All cases for urgent subcutaneous medications were collected by a representative on behalf of a patient. In 42.6% of cases the patient attended their usual pharmacy. Patients/representatives also indicated the pharmacy was:

convenient (14.8%); one of several pharmacies used (20.4%), or that they had been referred to the pharmacy for the medications (21.8%).

In 80% of cases patients/representatives received all medications against the prescription at the first pharmacy they visited. In 20% (11/55) one or more items on the prescription was not available, in five of these the item(s) were urgent. Free text sections were completed for six of the 11 cases of unavailable items. Four indicated they would return to collect the item from the pharmacy and two said they would try another pharmacy to obtain the items.

Thirteen respondents made additional comments on whether their experience could have been better. Comments were mostly positive: six indicated 'no', 'none', 'no fine' or similar phrase; five made comments on the staff or service: 'friendly services under difficult circumstances', 'no - staff really friendly and helpful, service was quick and efficient', 'Nothing – excellent and quick service'; and one explained 'nothing much that would make it better, but I phone in advance to make sure my items are in stock'. One respondent requested to 'keep a stock of all required items'.

Overall one in five patients/representatives had to go to more than one pharmacy to get urgently needed PMs, increasing to one in three for urgent subcutaneous injection prescription items. One in every two patients/representatives referred to the pharmacy by another HCP had to go to more than one pharmacy. Data from the prescription audit and customer survey were triangulated to verify the validity of the information in phase 1.

Phase 2: Interview findings

Findings from the interviews are presented in four sections: timely access; challenges; knowledge of LCS; and communication and collaboration. The findings are illustrated by verbatim quotes from the interviews where appropriate.

Timely access

Anticipating need and forward planning were key themes to ensure timely access to PMs.

... I could go in now and say, 'I need these drugs' (and the CP might say) 'Oh I can get them in for 11 o'clock tomorrow morning' [exasperated laugh] it's like

that's not really very helpful, I need them now (HCP7, Community Healthcare Professional)

Community nurses and palliative care team staff reported strategies to enhance access including conducting an end of week check and balance for those on syringe pumps to ensure sufficient stock for over the weekend when fewer staff were available. Specialist palliative care team staff also described making do with the medicines already available in the house and then ordering medication for the next day. CPs perceived that patients/HCPs phoning ahead for large quantities would be helpful. Insufficient quantities of PMs could adversely impact patient symptom management and had consequences for staff resources, however not all situations could be taken into consideration.

... a GP won't prescribe a syringe driver ahead of time...we are always...having to do it now not in a more considered way (HCP4, Community Healthcare Professional)

... I remember having to go [to the pharmacy] in the middle of doing a [syringe] driver because there weren't enough drugs (HCP8, Community Healthcare Professional)

The local clinical commissioning group (CCG) had implemented a template on the GP prescribing system to provide a 'suite' of PMs according to local last days of life algorithms which included some of the injectable medicines listed on the LCS formulary. Even so, in phase 1 several 'non-formulary' medications not on the local CCG PM list were prescribed and in phase 2 CPs in LCS pharmacies described non-compliance with the local formulary as a reason for a lack of timely access to PMs.

We've got three different strengths of oxycodone injection, and they [GPs] prescribe all three, and you might not have one, you might have the other...it's just so frustrating... (P4, Community Pharmacist)

The big problem is midazolam...so many strengths...volumes of ampoules...the GPs just pick one. (P5, Community Pharmacist)

Challenges

CPs described practical issues in supplying PMs for example: stock ordering processes including: timing of deliveries and the inability to return CD items to suppliers due to legal restrictions; CD cabinet size (to meet UK legal requirements for storage); and quantities on prescriptions.

We don't have an ability to be able to keep a lot [controlled drugs] and so we have a particular issue with the quantities that they write on the prescriptions sometimes which can impact on the next patient (P4, Community Pharmacist)

We've only got very small CD cabinets...the more controlled drugs you keep the more issues you are going to have (P1, Community Pharmacist)

Furthermore, patient records and charts to check opioid dose changes and syringe pumps were often not accessible to CPs.

With regards to changing doses or monitoring, I think that would be difficult for a community pharmacist. We have the summary care [record]... with palliative care the dose can change, you've got the pink card...sometimes we see [the pink card] and sometimes we don't (P2, Community Pharmacist)

... they [CPs] don't get the pink card [syringe pump chart] they simply get the prescription (HCP3, GP)

I might only need...extra diamorphine...whereas I might be using midazolam and haloperidol...but I've still got a supply of those, so they [CPs] don't always know what's in the [syringe] driver (HCP4, Community Healthcare Professional)

Knowledge of LCS

HCPs had little knowledge of either the LCS or the pharmacies commissioned to provide it but knew which pharmacies were likely to keep some PMs in stock.

I don't know who's commissioned we just basically know which ones we go to that are more likely to have it. (HCP4, Community Healthcare Professional)

... relatives who are running right left and centre trying to get hold of these meds...there is a commissioned service...but we don't know who they are. (HCP1, Community Healthcare Professional)

GPs generally thought that all pharmacies kept some injectable PMs in stock but said they might ring in advance to check the medication was available if a supply was needed urgently. Non-LCS pharmacist providers knew of the service and how to refer a patient/carer if they did not have the requested medication available. Usually they would phone ahead to the pharmacy to check the medication was available before making a referral. Being able to make a referral depended on whether the carer had access to a car.

... if they haven't got a car it's sometimes a bit tricky (P1, Community Pharmacist)

Communication and Collaboration

Participants described strategies to enhance access to PMs. Two non-LCS pharmacies had worked with GP practices to discuss and agree to stock a subset of the LCS PMs; they had similar response times to pharmacies in the LCS.

... we went to them [GPs] and said, what are the most common drugs you would prescribe in palliative care...they [GPs] came back with a list...so we would try to keep the stock in for what they specified (P3, Community Pharmacist)

CPs reported that some patients/carers contacted the pharmacy when they ordered a prescription for a CD that might not be stocked. Community and specialist palliative care team staff described how they would suggest the pharmacist kept sufficient stock when they had someone on a syringe pump or large quantities of injectable medications. There appeared to be some examples where excellent communication and collaboration existed between GPs, HCPs and CPs which resulted in more timely access for PMs.

... one GP...rang us and said well what have you got in stock and what can you get, which I found really, really useful because as the prescription came in the stock came in and this thing was completely seamless (P3, Community Pharmacist)

When we were down at [previous community nurse location] ...there was a pharmacy next door so...if we had any quick questions, we would go and talk to them...they were more like part of the team (HCP7, Community Healthcare Professional)

However, concerns around patient confidentiality by GPs and other HCPs meant that more often this information was not shared with the pharmacy team in advance of receiving the prescription.

... but you're limited by what you can tell them [pharmacists] obviously from a confidentiality point of view... (HCP11, Community Healthcare Professional)

We don't communicate with them [community pharmacist] what the problem with the patient is we just prescribe the drugs... sometimes they can obviously work it out. (HCP3, GP)

I do have some slight reservations about them [pharmacists] knowing all those ins and outs...I'm not sure how wide that circle is in there [pharmacy]...I'd prefer it ...on just a case by case basis...to an identified clinician... (HCP10, GP)

DISCUSSION

The sequential use of mixed methods to firstly quantify the "problem" and then qualitative methods to provide context to the barriers to timely supply of PMs generated new insights into a longstanding problem. Timeliness of access was found to primarily relate to a mismatch between medicines stocks held by CPs and the PMs that GPs prescribed. Legal errors on CD prescriptions played a much smaller role and had little impact on access to PMs in this study. Stock availability as a

significant factor to support timely access has also been seen in previous studies.[5,8,10,11]

Study results appear to indicate a low prevalence of legal errors on palliative care prescriptions considering 42% of prescriptions were for CDs. Previous studies suggest legal errors on prescriptions can range from less than 1% up to approximately 1.9% [14,20,24–28] though data does not specifically focus on palliative care prescriptions. Legal errors in the current study arose on computergenerated prescriptions for controlled drugs administered by a syringe pump. The primary care organisation prescribing template for PMs may have impacted positively, minimising the number of legal errors.

Our findings of differential time to access PMs between community pharmacies participating in the LCS for PMs and non-participants, indicates that a local service can enhance access. Those pharmacies working with local GP practices to keep a small agreed range of PMs in stock had similar access times to those within the LCS, suggesting that such collaboration can also support more timely access and improve patient and carer experience. Such wider collaboration has been advocated within national policy drivers and enables greater integration of pharmacy teams in improving patient care. [29-32] Some HCPs considered CPs "part of the team" but others saw them only in their "supply" role. Together with concerns about confidentiality this prevented many HCPs from communicating with CPs about available stock or giving advance warning to allow stock to be obtained in a more timely way. A particular concern seemed to be who else, in addition to the pharmacist, might have access to sensitive information. Commissioners could remind primary care staff of community pharmacy ethical and information governance practices to correct any misunderstandings. They could also encourage HCPs to be proactive in checking stock availability with the patient's usual pharmacy.

Palliative patients often rely on family members and friends to support them with managing their medication especially towards the end-of-life.[9,16-18] Our findings show that some families obtain urgently required medicines from a pharmacy different than the one that usually supplied the patient's medicines. It is unclear what effect these changes in continuity of care between pharmacies might have towards the end-of-life, but this study shows that some CPs enhance access by calling other

pharmacies to ascertain they have PMs in stock. A potential solution could be through CPs having read and write access to the summary care record (SCR) allowing them, whether they are the patient's regular, out-of-hours or LCS pharmacy, to record patient care scenarios to ensure safe, continuity of care of PMs. Variable accessibility and difficulties in use of SCR by CPs[33] may suggest wider access to patient records is required.

One in five patients/representatives accessing PMs had to go to more than one pharmacy. This is the first study to quantify the number of patients/representatives who had to do so, with associated inconvenience, wasted time and stress. This finding could be explained by a lack of awareness of the LCS since this is not advertised to the public and there was also low awareness amongst HCPs and GPs in the interviews. Monitoring of LCS/LES services by commissioners may not always be effective. One audit found only one of nineteen pharmacies held all PMs on the formulary list and some CPs were not aware that the scheme was active. (Aslett, M. and Wall-Hayes, L. 2015. 'Access to palliative drugs – community pharmacy scheme – audit.' Unpublished NHS audit report, Birmingham, UK) There was also evidence in phase 1 of prescriptions being written for items not on the LCS list which would not usually be stocked in the pharmacies. Commissioners' service audits could investigate referral patterns from pharmacies not within the LCS, furthermore monitoring of LCS pharmacies may improve practice and caregivers' experience. Commissioners could also act to improve awareness of LCS among local HCPs.

Some limitations affect the interpretation/generalisability of the findings of this study. The small sample of participating pharmacies, missing data, reliance on CPs to identify prescriptions and confounding factors such as time of day, number and type of staff working in the pharmacy may limit interpretation of the results and introduce a degree of bias. Differences in the commissioning of access to PMs within England also may limit the findings as there is no standard service specification stating the outcomes to be measured. Furthermore, the geographical restriction with data only collected in one city could limit application to other areas including those in remote locations, with different out-of-hours providers, and access to palliative care support in the community. Nevertheless, the study adds value to the literature in terms of barriers that need to be considered if more timely access to PMs is to be more

widely implemented and the methodology has enabled new insights into factors contributing to timely access.

CONCLUSION

The findings of this study suggest that legal prescribing errors may now have a smaller impact on access to urgent PMs from community pharmacies compared to mismatches between stock availability and PMs prescribed. Both participation in a LCS or collaboration with local prescribers are likely to improve access to PMs.

Recommendations for future commissioning and practice:

Commissioners should:

- i) encourage GP practices to work with local pharmacies to keep a small range of PMs available
- ii) remind HCPs of the ethical and information governance requirements of community pharmacies and encourage early contact to check stock availability
- iii) involve patients and the public in designing audits of LCS.

Community pharmacies should improve their communication with HCPs around pharmacy opening times and cut-off times for same-day delivery of medicines.

Moving forward, NHS England will be supporting development and integration of CP services into primary care through its Pharmacy Integration Fund[34] and this may also improve interprofessional communication and access to PMs.

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Author Contributions

EM implemented the study, designed and piloted data collection tools, monitored data collection, wrote the analysis plan, analysed the data, wrote and piloted interview schedule, transcribed interviews, developed the thematic framework and drafted and revised the manuscript. JDM and AB supervised EM in planning and undertaking the study including study design, development of data collection tools, data analysis, drafting and revision of the manuscript.

Competing interests

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Ethics approval

Ethical approval obtained 17th December 2015 from the Chair of the Biomedical, Natural, Physical and Health Sciences Ethics Panel, University of Bradford (approval reference E493).

Data sharing statement

Protocol and data collection forms are available upon request from the corresponding author.

Patient Involvement

The design of the study was based on EM's experience as a clinical pharmacist including discussions with patients and their families on accessing medicines. The research question was therefore derived from patients' and family carers' experience on accessing medicines towards the end-of-life. A Hospice Service User Coordinator provided support with the customer survey based on their experience of conducting surveys. Furthermore, patients within a hospice day centre supported the piloting of the customer survey.

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