

Unannounced Medicines Management Inspection Report 2 May 2017











Millverne

Type of service: Residential Care Home Address: 66 Mill Street, Enniskillen, BT74 6DW

Tel No: 028 6634 6000 Inspector: Paul Nixon

1.0 Summary

An unannounced inspection of Millverne took place on 2 May 2017 from 09:40 to 13:10.

The inspection sought to assess progress with any issues raised during and since the previous inspection and to determine if the home was delivering safe, effective and compassionate care and if the service was well led.

Is care safe?

There was evidence that the management of medicines supported the delivery of safe care and positive outcomes for residents. Staff administering medicines were trained and competent. There were systems in place to ensure the management of medicines was in compliance with legislative requirements and standards. It was evident that the knowledge of the staff enables the systems in place for the management of medicines to be robust. There were no areas for improvement identified.

Is care effective?

The management of medicines supported the delivery of effective care. There were systems in place to ensure residents were receiving their medicines as prescribed. One area of improvement was identified in relation to record keeping for food and fluid thickeners and a recommendation was made.

Is care compassionate?

The management of medicines supported the delivery of compassionate care. Staff interactions were observed to be compassionate, caring and timely which promoted the delivery of positive outcomes for residents. Residents consulted with confirmed that they were administered their medicines appropriately. There were no areas for improvement identified.

Is the service well led?

The service was found to be well led with respect to the management of medicines. Written policies and procedures for the management of medicines were in place which supported the delivery of care. Systems were in place to enable management to identify and cascade learning from any medicine related incidents and medicine audit activity. There were no areas for improvement identified.

This inspection was underpinned by The Residential Care Homes Regulations (Northern Ireland) 2005 and the Department of Health, Social Services and Public Safety (DHSSPS) Residential Care Homes Minimum Standards (2011).

1.1 Inspection outcome

	Requirements	Recommendations
Total number of requirements and	0	1
recommendations made at this inspection	-	

Details of the Quality Improvement Plan (QIP) within this report were discussed with Mrs Julianne Treacy, Registered Manager, as part of the inspection process. The timescales for completion commence from the date of inspection.

Enforcement action did not result from the findings of this inspection.

1.2 Actions/enforcement taken following the most recent premises inspection

There were no further actions required to be taken following the most recent inspection on 19 January 2017.

2.0 Service details

Registered organisation/registered person: Carewell Homes Ltd / Mrs Carol Kelly	Registered manager: Mrs Julianne Treacy
Person in charge of the home at the time of inspection: Mrs Julianne Treacy	Date manager registered: 1 April 2005
Categories of care: RC-I, RC-MP, RC-MP(E), RC-DE	Number of registered places: 45

3.0 Methods/processes

Prior to inspection we analysed the following records:

- recent inspection reports and returned QIPs
- recent correspondence with the home
- the management of medicine related incidents reported to RQIA since the last medicines management inspection.

During the inspection the inspector met with three residents, the registered manager, the clinical governance lead and two senior care assistants.

A poster indicating that the inspection was taking place was displayed in the lobby of the home and invited visitors/relatives to speak with the inspector. No one availed of this opportunity during the inspection.

The following records were examined during the inspection:

- medicines requested and received
- personal medication records
- medicine administration records
- medicines disposed of or transferred
- controlled drug record book

- medicine audits
- policies and procedures
- care plans
- training records
- medicines storage temperatures

4.0 The inspection

4.1 Review of requirements and recommendations from the most recent inspection dated 19 January 2017

The most recent inspection of the home was an announced premises inspection. No requirements or recommendations were made.

4.2 Review of requirements and recommendations from the last medicines management inspection dated 01 September 2014

Last medicines mana	gement inspection statutory requirements	Validation of compliance
Requirement 1 Ref: Regulation 13(4) Stated: First time	The registered manager must ensure that anticoagulant medicines are administered in accordance with the prescriber's instructions. Action taken as confirmed during the inspection: The registered manager advised that the procedure for the management of anticoagulants had been reviewed and discussed with senior care staff. No anticoagulants were currently prescribed.	Met
Last medicines mana	gement inspection recommendations	Validation of compliance
Recommendation 1 Ref: Standard 30 Stated: First time	The registered manager should increase the level of audit of liquid medicines and any further discrepancies must be investigated and reported to RQIA. Action taken as confirmed during the inspection: The level of audit of liquid medicines had been increased. The registered manager and staff stated that there had been no discrepancies observed.	Met
Recommendation 2 Ref: Standard 30 Stated: First time	The registered manager should ensure that nutritional supplements are included in the home's auditing procedures. Action taken as confirmed during the inspection: The registered manager and staff advised and the audit records indicated that nutritional supplements were included in the home's auditing procedures.	Met

Recommendation 3

Ref: Standard 30

Stated: First time

The registered manager should review and revise the arrangements in place for the management of anxiolytic and antipsychotic medicines prescribed on an "as required" basis for the management of distressed reactions.

Action taken as confirmed during the inspection:

When a resident was prescribed a medicine for administration on a "when required" basis for the management of distressed reactions, the dosage instructions were recorded on the personal medication record. Staff knew how to recognise signs, symptoms and triggers which may cause a change in a resident's behaviour and were aware that this change may be associated with pain. The reason for and the outcome of administration were recorded. A care plan was generally maintained.

Met

4.3 Is care safe?

Medicines were managed by staff who have been trained and deemed competent to do so. An induction process was in place for care staff who had been delegated medicine related tasks. The impact of training was monitored through team meetings, supervision and annual appraisal. Competency assessments were completed annually. Refresher training in medicines management was provided in the last year.

Systems were generally in place to manage the ordering of prescribed medicines to ensure adequate supplies were available and to prevent wastage. Staff advised of the procedures to identify and report any potential shortfalls in medicines.

There were satisfactory arrangements in place to manage changes to prescribed medicines. Personal medication records were generally updated by two members of staff.

There were procedures in place to ensure the safe management of medicines during a resident's admission to the home.

Records of the receipt, administration and disposal of controlled drugs subject to record keeping requirements were maintained in controlled drug record books. Checks were performed on controlled drugs which require safe custody, at the end of each shift. Additional checks were also performed on other controlled drugs which is good practice.

Appropriate arrangements were in place for administering antibiotic courses and new medicines.

Discontinued or expired medicines were disposed of appropriately.

Medicines were stored safely and securely and in accordance with the manufacturer's instructions. Medicine storage areas were clean, tidy and well organised. There were systems in place to alert staff of the expiry dates of medicines with a limited shelf life, once opened. The medicine refrigerator was checked at regular intervals.

RQIA ID: 1798 Inspection ID: IN028582

Areas for improvement

No areas for improvement were identified during the inspection.

Number of requirements	0	Number of recommendations	0
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4.4 Is care effective?

The sample of medicines examined had been administered in accordance with the prescriber's instructions.

There was evidence that time critical medicines had been administered at the correct time. There were arrangements in place to alert staff of when doses of weekly and three monthly medicines were due.

The sample of records examined indicated that medicines which were prescribed to manage pain had been administered as prescribed. Staff advised that a pain assessment was completed as part of the admission process. Staff were aware that ongoing monitoring was necessary to ensure that the pain was well controlled and the resident was comfortable. Staff advised that the residents could verbalise any pain.

The management of swallowing difficulty was examined. A care plan and speech and language assessment report were in place. However, the food and fluid thickening agent was not recorded on the personal medication record and administrations were not recorded. A recommendation was made.

Staff confirmed that compliance with prescribed medicine regimes was monitored and any omissions or refusals likely to have an adverse effect on the resident's health were reported to the prescriber.

Medicine records were well maintained and facilitated the audit process.

Practices for the management of medicines were audited throughout the month by the staff and management.

Following discussion with the registered manager and staff, it was evident that, when applicable, other healthcare professionals were contacted in response to the needs of residents.

Areas for improvement

The arrangements for the recording of thickening agents need to be reviewed. A recommendation was made.

Number of requirements	0	Number of recommendations	1

4.5 Is care compassionate?

The administration of medicines to residents was completed in a caring manner, residents were given time to take their medicines and medicines were administered as discreetly as possible.

Residents spoken with advised that they were very satisfied with the care experienced. Residents who could not verbalise their feelings in respect of their care were observed to be relaxed and comfortable in their surroundings and in their interactions with staff.

As part of the inspection process, we issued questionnaires to residents, residents' representatives and staff. Four residents and one resident's representative completed and returned questionnaires within the specified timeframe. Comments received were very positive; the responses were recorded as 'satisfied' or 'very satisfied' with the management of medicines in the home.

One member of staff also completed a questionnaire. The responses were positive and raised no concerns about the management of medicines in the home.

Areas for improvement

No areas for improvement were identified during the inspection.

Number of requirements	0	Number of recommendations	0

4.6 Is the service well led?

Written policies and procedures for the management of medicines were in place. Following discussion with staff it was evident that they were familiar with the policies and procedures and that any updates were highlighted to them.

There were robust arrangements in place for the management of medicine related incidents. Staff confirmed that they knew how to identify and report incidents. Medicine related incidents reported since the last medicines management inspection were discussed. There was evidence of the action taken and learning implemented following incidents.

Staff knew the identity of their adult safeguarding lead. They knew that medicine incidents should be considered under safeguarding procedures and how to report these.

A review of the audit records indicated that satisfactory outcomes had been achieved.

Following discussion with the registered manager and care staff, it was evident that staff were familiar with their roles and responsibilities in relation to medicines management.

Staff confirmed that any concerns in relation to medicines management were raised with management.

Areas for improvement

No areas for improvement were identified during the inspection.

5.0 Quality improvement plan

Any issues identified during this inspection are detailed in the QIP. Details of the QIP were discussed with Mrs Julianne Treacy, Registered Manager, as part of the inspection process. The timescales commence from the date of inspection.

The registered provider/manager should note that failure to comply with regulations may lead to further enforcement action including possible prosecution for offences. It is the responsibility of the registered provider to ensure that all requirements and recommendations contained within the QIP are addressed within the specified timescales.

Matters to be addressed as a result of this inspection are set in the context of the current registration of the residential care home. The registration is not transferable so that in the event of any future application to alter, extend or to sell the premises RQIA would apply standards current at the time of that application.

5.1 Statutory requirements

This section outlines the actions which must be taken so that the registered provider meets legislative requirements based on The Residential Care Homes Regulations (Northern Ireland) 2005.

5.2 Recommendations

This section outlines the recommended actions based on research, recognised sources and The Department of Health, Social Services and Public Safety (DHSSPS) Residential Care Homes Minimum Standards (2011). They promote current good practice and if adopted by the registered provider/manager may enhance service, quality and delivery.

5.3 Actions to be taken by the registered provider

The QIP should be completed and detail the actions taken to meet the legislative requirements and recommendations stated. The registered provider should confirm that these actions have been completed and return the completed QIP to pharmacists@rqia.org.uk for assessment by the inspector.

It should be noted that this inspection report should not be regarded as a comprehensive review of all strengths and areas for improvement that exist in the service. The findings reported on are those which came to the attention of RQIA during the course of this inspection. The findings contained within this report do not exempt the registered provider from their responsibility for maintaining compliance with the regulations and standards. It is expected that the requirements and recommendations outlined in this report will provide the registered provider with the necessary information to assist them to fulfil their responsibilities and enhance practice within the service.

Quality Improvement Plan		
Recommendations		
Recommendation 1	The registered provider should ensure that the recording arrangements for thickening agents are reviewed.	
Ref: Standard 31		
	Response by registered provider detailing the actions taken:	
Stated: First time	We have reviewed our recording arrangements and are now keeping a record of all thickening agent use	
To be completed by: 1 June 2017		

^{*}Please ensure this document is completed in full and returned to pharmacists@rgia.org.ukfrom the authorised email address*





The Regulation and Quality Improvement Authority

9th Floor

Riverside Tower 5 Lanyon Place BELFAST

BT1 3BT

Tel 028 9051 7500

Fax 028 9051 7501

Email info@rqia.org.uk

Web www.rqia.org.uk

@RQIANews