

Unannounced Medicines Management Inspection Report 19 June 2017



Gnangara

Type of Service: Residential Care Home
Address: 163 Sligo Road, Drumawill, Enniskillen, BT74 7JZ
Tel No: 028 66325134
Inspector: Paul Nixon

www.rqia.org.uk

Assurance, Challenge and Improvement in Health and Social Care

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 service from their responsibility for maintaining compliance with legislation, standards and best practice.

1.0 What we look for



2.0 Profile of service

This is a residential care home with 15 beds that provides care for residents living with dementia.

3.0 Service details

Organisation/Registered Provider: Fold Housing Association Responsible Individual: Mrs Fiona McAnespie	Registered Manager: See box below.
Person in charge at the time of inspection: Ms Margaret Irwin	Date manager registered: Ms Margaret Irwin - application received - "registration pending".
Categories of care: Residential Care DE – Dementia	Number of registered places: 15

4.0 Inspection summary

An unannounced inspection took place on 22 June 2017 from 09.25 to 13.10.

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).

The inspection assessed progress with any areas for improvement identified during and since the last medicines management inspection and to determine if the home was delivering safe, effective and compassionate care and if the service was well led.

Evidence of good practice was found in relation to medicine administration, medicine records, storage and the management of controlled drugs.

No areas requiring improvement were identified.

The findings of this report will provide the home with the necessary information to assist them to fulfil their responsibilities, enhance practice and residents' experience.

4.1 Inspection outcome

	Regulations	Standards
Total number of areas for improvement	0	0

This inspection resulted in no areas for improvement being identified. Findings of the inspection were discussed with Ms Margaret Irwin, Manager, as part of the inspection process and can be found in the main body of the report.

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

4.2 Action/enforcement taken following the most recent care inspection

Other than those actions detailed in the QIP no further actions required to be taken following the most recent inspection on 11 May 2017. Enforcement action did not result from the findings of this inspection.

5.0 How we inspect

Prior to the inspection a range of information relevant to the service was reviewed. This included the following:

- 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.

A poster informing visitors to the home that an inspection was being conducted was displayed.

During the inspection the inspector met with the manager and three members of care staff.

A total of 15 questionnaires were provided for distribution to residents, their representatives and staff for completion and return to RQIA.

A sample of the following records was 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
- care plans
- training records
- medicines storage temperatures

Areas for improvements identified at the last medicines management inspection were reviewed and the assessment of compliance recorded as met, partially met, or not met.

The findings of the inspection were provided to the person in charge at the conclusion of the inspection.

6.0 The inspection

6.1 Review of areas for improvement from the most recent inspection dated 11 May 2017

The most recent inspection of the home was an unannounced care inspection. The completed QIP will be reviewed by the care inspector. This QIP will be validated by the care inspector at the next care inspection.

6.2 Review of areas for improvement from the last medicines management inspection dated 09 March 2015

Action required to ensure compliance with the Department of Health, Social Services and Public Safety (DHSSPS) Residential Care Homes Minimum Standards (2011).		Validation of compliance
Area for improvement 1 Ref: Standard 30 Stated: First time	The Registered Provider should ensure that appropriate care plans are in place for the management of any anxiolytic or antipsychotic medicines prescribed on an “as required” basis for the management of distressed reactions.	Met
	Action taken as confirmed during the inspection: Appropriate care plans were in place for the management of any anxiolytic or antipsychotic medicines prescribed on an “as required” basis for the management of distressed reactions.	

6.3 Inspection findings

6.4 Is care safe?

Avoiding and preventing harm to patients and clients from the care, treatment and support that is intended to help them.

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 at least annually. Refresher training in medicines management was provided in the last year.

Systems were 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. Antibiotics and newly prescribed medicines had been received into the home without delay. Satisfactory arrangements were in place for the acquisition and storage of prescriptions.

There were satisfactory arrangements in place to manage changes to prescribed medicines. Personal medication records were updated by two members of staff. This safe practice was acknowledged.

In relation to safeguarding, staff advised that they were aware of the regional procedures and who to report any safeguarding concerns to. Training had been completed in October 2016.

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

Records of the receipt, administration and disposal of controlled drugs subject to record keeping requirements were maintained in a controlled drug record book. 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.

Robust arrangements were observed for the management of high risk medicines e.g. warfarin. The use of separate administration charts was acknowledged.

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.

Areas of good practice

There were examples of good practice in relation to staff, training, competency assessments, the management of medicines on admission and controlled drugs.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Total number of areas for improvement	0	0

6.5 Is care effective?

The right care, at the right time in the right place with the best outcome.

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 or three monthly medicines were due.

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 maintained.

The sample of records examined indicated that medicines which were prescribed to manage pain had been administered as prescribed. Staff were aware that ongoing monitoring was necessary to ensure that the pain was well controlled and the resident was comfortable. Staff advised that most of the residents could verbalise any pain, and a pain assessment tool was used as needed. A care plan was maintained. Staff also advised that a pain assessment is completed as part of the admission process.

The management of swallowing difficulty was examined. For those residents prescribed a thickening agent, this was generally recorded on their personal medication record and included details of the fluid consistency. However, one resident did not have the thickening agent recorded on their personal medication record; the manager gave an assurance that this matter would be rectified without delay. Administrations were recorded and care plans and speech and language assessment reports were in place.

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. Areas of good practice included recording the date of replacement of medicine containers. The manager agreed to ensure that staff promptly remove discontinued personal medication records from the current folder and archive them.

Practices for the management of medicines were audited throughout the month by the staff and management. Care staff performed daily audits on all medicines. In addition, a quarterly audit was completed by the community pharmacist.

Following discussion with the manager and staff, it was evident that, when applicable, other healthcare professionals are contacted in response to the healthcare needs of residents. Staff on duty advised that they had good working relationships with the community pharmacy, GP practices and the Health and Social Care Trust.

Areas of good practice

There were examples of good practice in relation to the standard of record keeping and the administration of medicines.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Total number of areas for improvement	0	0

6.6 Is care compassionate?

Patients and clients are treated with dignity and respect and should be fully involved in decisions affecting their treatment, care and support.

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 were observed to be relaxed and comfortable in their surroundings and in their interactions with staff. Staff were noted to be friendly, courteous and happy in their work; they treated the residents with dignity.

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

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

Areas of good practice

Staff listened to residents and took account of their views.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Total number of areas for improvement	0	0

6.7 Is the service well led?

Effective leadership, management and governance which creates a culture focused on the needs and experience of service users in order to deliver safe, effective and compassionate care.

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. In relation to the regional safeguarding procedures, staff confirmed that they were aware that medicine incidents may need to be reported to the safeguarding lead and safeguarding team.

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

Following discussion with the 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 of good practice

There were examples of good practice in relation to governance arrangements, the management of medicine incidents and quality improvement. There were clearly defined roles and responsibilities for staff.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Total number of areas for improvement	0	0

7.0 Quality improvement plan

There were no areas for improvement identified during this inspection, and a QIP is not required or included, as part of this inspection report.

RQIA will phase out the issue of draft reports via paperlite in the near future. Registered providers should ensure that their services are opted in for the receipt of reports via Web Portal. If you require further information, please visit www.rqia.org.uk/webportal or contact the web portal team in RQIA on 028 9051 7500.



The Regulation and
Quality Improvement
Authority

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