

Unannounced Medicines Management Inspection Report 3 October 2017



Magherafelt Manor

Type of Service: Nursing Home
Address: 22 Pound Road, Magherafelt, BT45 6NR
Tel No: 028 7930 0284
Inspector: Rachel Lloyd

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 nursing home with 34 beds that provides care for patients living with a range of healthcare needs as detailed in section 3.0. The nursing home is on the same site as a residential care home.

3.0 Service details

Organisation/Registered Provider: Runwood Homes Ltd Responsible Individual: Mr Gavin O'Hare-Connolly (registration pending)	Registered Manager: Ms Siobhan Conway
Person in charge at the time of inspection: Ms Siobhan Conway	Date manager registered: 31 March 2015
Categories of care: Nursing Homes (NH): DE - Dementia I - Old age not falling within any other category PH - Physical disability other than sensory impairment PH(E) - Physical disability other than sensory impairment - over 65 years	Number of registered places: 34 comprising: A maximum of 22 patients in category NH-DE accommodated in the ground floor unit. A maximum of 12 patients in categories NH-I, NH-PH, NH-PH(E) accommodated in the first floor unit.

4.0 Inspection summary

An unannounced inspection took place on 3 October 2017 from 10.00 to 14.40.

This inspection was underpinned by The Health and Personal Social Services (Quality, Improvement and Regulation) (Northern Ireland) Order 2003, The Nursing Homes Regulations (Northern Ireland) 2005 and the Department of Health, Social Services and Public Safety (DHSSPS) Care Standards for Nursing Homes, April 2015.

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 the administration and storage of medicines, medicine records, care planning, communication with various healthcare professionals, working relationships within the home and the management of the ordering and supply of medicines.

No areas for improvement were identified during the inspection.

The findings of this report will provide the home with the necessary information to assist them to fulfil their responsibilities, enhance practice and patients' 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 Siobhan Conway, Registered 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

No further actions were required to be taken following the most recent inspection on 20 August 2017.

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.

We met with three patients, one care assistant, two registered nurses, the deputy manager and the registered manager.

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

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

A sample of the following records was examined during the inspection:

- | | |
|--|----------------------------------|
| • medicines requested and received | • medicine audits |
| • personal medication records | • care plans |
| • medicine administration records | • training records |
| • medicines disposed of or transferred | • medicines storage temperatures |
| • controlled drug record book | |

Areas for improvement 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 20 August 2017

The most recent inspection of the home was an unannounced follow up care inspection. There were no areas for improvement made as a result of the inspection.

6.2 Review of areas for improvement from the last medicines management inspection dated 9 May 2016

Areas for improvement from the last medicines management inspection		
Action required to ensure compliance with the Department of Health, Social Services and Public Safety (DHSSPS) Care Standards for Nursing Homes, April 2015		Validation of compliance
Area for improvement 1 Ref: Standard 28 Stated: First time	A risk assessment and care plan should be in place for any patient who has medicines administered in disguised form.	Met
	Action taken as confirmed during the inspection: Satisfactory action had been taken. A risk assessment and care plan were in place in the two examples examined.	
Area for improvement 2 Ref: Standard 18 Stated: First time	The reason for and the outcome of administration of medicines prescribed for administration on a "when required" basis for the management of distressed reactions should be routinely recorded.	Met
	Action taken as confirmed during the inspection: The reason for and outcome of administration of these medicines had been recorded in the examples examined. A supplementary sheet for recording this information was in place and this sheet and/or the patient's daily notes included this information.	

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 registered nurses and 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. In relation to safeguarding, staff advised that they were aware of the regional procedures and who to report any safeguarding concerns to.

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 registered nurses. This safe practice was acknowledged.

There were procedures in place to ensure the safe management of medicines during a patient's admission to the home and discharge from 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. Discontinued controlled drugs were denatured and rendered irretrievable prior to disposal.

Medicines were stored safely and securely and in accordance with the manufacturer's instructions. Medicine storage areas were clean, tidy and organised. There were systems in place to alert staff of the expiry dates of medicines with a limited shelf life, once opened. Medicine refrigerators and oxygen equipment were checked at regular intervals. Staff were reminded that refrigerator thermometers should be reset daily after recording temperatures.

Areas of good practice

There were examples of good practice found throughout the inspection in relation to staff training and competency assessment, the management on medicines on admission/discharge,

the management of controlled drugs, the disposal of medicines and the storage of prescriptions and medicines.

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 majority of 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 as to when doses of weekly, monthly or three monthly medicines were due.

The management of distressed reactions, swallowing difficulty and pain were reviewed. The relevant information was mostly recorded in the patient's care plan, personal medication record and records of administration. The registered manager agreed to add details of the prescribed consistency of thickened fluids to the personal medication record for two identified patients.

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

Medicine records were well maintained and facilitated the audit process. Areas of good practice were acknowledged. These included the use of supplementary sheets for the administration of antibiotics, injections, transdermal opioid patches, warfarin and thickening agents, and the assessment of pain. There were a few gaps in the completion of some of these supplementary records. Staff and management were reminded that when these additional records are in use they should be completed accurately at all times.

Practices for the management of medicines were audited throughout the month by the staff and management. In addition, audits were completed by the community pharmacist.

Following discussion with the registered manager and staff, it was evident that other healthcare professionals are contacted when required to meet the needs of patients.

Areas of good practice

There were examples of good practice found throughout the inspection in relation to record keeping, care planning, the administration of medicines and audit procedures.

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 patients was briefly observed. It was completed in a caring manner, patients were given time to take their medicines and medicines were administered as discreetly as possible.

Patients spoken to appeared content and relaxed in the home. Patients 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. Staff demonstrated a good knowledge of patients' wishes and preferences.

At the time of issuing this report none of the questionnaires issued had been returned from patients, relatives or staff.

Areas of good practice

There was evidence that staff listened to and valued patients and took account of their views. Warm relationships were observed between staff and patients.

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; these were not examined on this occasion. 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. In relation to the regional

safeguarding procedures, staff confirmed that they were aware that medicine incidents may need to be reported to the safeguarding team.

A review of the audit records indicated that largely satisfactory outcomes had been achieved. Where a discrepancy had been identified, there was evidence of the action taken and learning which had resulted in a change of practice.

Following discussion with the registered manager and 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 found throughout the inspection in relation to governance arrangements, the management of medicine incidents and maintaining good working relationships. 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.



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
Twitter @RQIANews