

# Unannounced Medicines Management Inspection Report 9 January 2018











# Laganvale

Type of Service: Nursing Home

Address: Dementia Unit 37 Laganvale Mews,

Lisburn, BT67 0RE Tel No: 028 9261 9899

**Inspector: Catherine Glover** 

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 36 beds that provides care for patients living with dementia.

#### 3.0 Service details

Organisation/Registered Provider: Four Seasons Healthcare	Registered Manager: Mrs Shily Paul
Responsible Individual: Dr Maureen Claire Royston	
Person in charge at the time of inspection: Mrs Shily Paul	Date manager registered: 7 November 2007
Categories of care: Nursing Homes DE – Dementia	Number of registered places: 36

## 4.0 Inspection summary

An unannounced inspection took place on 9 January 2018 from 10.00 to 13.30 hours.

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 DHSSPS Care Standards for Nursing Homes 2015.

The inspection assessed progress with any areas for improvement identified since the last care 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 medicines administration, records, storage and controlled drugs.

No areas requiring improvement were identified.

Patients were observed to be relaxed and comfortable in the home and their interactions with staff.

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 Mrs Shily Paul, 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 premises inspection

Other than those actions detailed in the QIP no further actions were required to be taken following the most recent inspection on 28 November 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.

During the inspection we met with two patients, the registered manager and three registered nurses.

Ten questionnaires were provided for distribution to patients and their representatives for completion and return to RQIA. Staff were invited to share their views by completing an online questionnaire.

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

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 28 November 2017

The most recent inspection of the home was an announced premises inspection. The QIP resulting from this inspection will be validated by the estates inspector at the next premises inspection.

# 6.2 Review of areas for improvement from the last medicines management inspection dated 23 February 2017

There were no areas for improvement identified as a result of the last medicines management inspection.

# 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. The impact of training was monitored through team meetings, supervision and annual appraisal. Competency assessments were completed annually. Refresher training in medicines management, distressed reactions and the management of gastrostomy tubes was provided in the last year. Staff have also completed safeguarding training and 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 and handwritten entries on medication administration 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.

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.

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 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 and oxygen equipment were checked at regular intervals.

#### Areas of good practice

There were examples of good practice in relation to staff training, competency assessment, 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, monthly or three monthly medicines were due.

The management of pain and swallowing difficulties was examined and found to be satisfactory. There were appropriate entries on the personal medication records, medicine administration records and care plans 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 patient's health were reported to the prescriber.

Medicine records were well maintained and facilitated the audit process. Areas of good practice were acknowledged. They included additional records for the administration of transdermal patches and running stock balances of some medicines.

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

Following discussion with the registered manager and staff and examination of care files, 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 in relation to the standard of record keeping, care planning and the administration of medicines.

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## **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 had been completed at the commencement of this inspection and was not observed. However, staff were very knowledgeable regarding patients' needs and wishes. The administration of medicines was discussed with the registered manager at the conclusion of the inspection and demonstrated that the medicines rounds were closely monitored.

At the time of this inspection a number of patients were having an appointment with the attending hairdresser. Staff were assisting the patients and engaging in a sing-song, encouraging everyone to join in. The atmosphere was cheerful and good relationships were evident.

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.

None of the questionnaires that were left in the home were returned within the timeframe for inclusion in this report.

#### Areas of good practice

Staff listened to patients and relatives 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. Management advised that these were reviewed regularly. Following discussion with staff it was evident that they were familiar with the policies and procedures and that any updates were highlighted to staff.

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 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 registered nurses, 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.





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