

# Unannounced Medicines Management Inspection Report 23 February 2017











# Laganvale (Dementia Unit)

Type of Service: Nursing Home

Address: 37 Laganvale Mews, Moira, BT67 0RE

Tel no: 028 9261 9899 Inspector: Cathy Wilkinson

## 1.0 Summary

An unannounced inspection of Laganvale (Dementia Unit) took place on 23 February 2017 from 10.15 to 12.30.

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 patients. 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 working relationship with the community pharmacist, the knowledge of the staff and their proactive action in dealing with any issues 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 patients were receiving their medicines as prescribed. There were no areas for improvement identified.

### 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 patients. Patients 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 Nursing Homes Regulations (Northern Ireland) 2005 and the Department of Health, Social Services and Public Safety (DHSSPS) Care Standards for Nursing Homes, April 2015.

# 1.1 Inspection outcome

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

This inspection resulted in no requirements or recommendations being made. 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.

## 1.2 Actions/enforcement taken following the most recent care inspection

Other than those actions detailed in the Quality Improvement Plan (QIP) there were no further actions required to be taken following the most recent inspection on 3 October 2016.

### 2.0 Service details

Registered organisation/registered person: Four Seasons Healthcare Dr Maureen Claire Royston	Registered manager: Mrs Shily Paul
Person in charge of the home at the time of inspection: Mrs Shily Paul	Date manager registered: 7 November 2007
Categories of care: NH-DE	Number of registered places: 36

### 3.0 Methods/processes

Prior to inspection the following records were analysed:

- 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 the nursing sister and the registered manager.

A sample of the following records was examined:

- 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 3 October 2016

The most recent inspection of the home was an unannounced care inspection. The QIP from this inspection will be validated by the care inspector at their next inspection.

# 4.2 Review of requirements and recommendations from the last medicines management inspection dated 25 September 2014

Last medicines mana	gement inspection statutory requirements	Validation of compliance
Requirement 1  Ref: Regulation 13(4)  Stated: Second time	Further monitoring of the administration of liquid medicines in the Dementia Unit must be undertaken to ensure that they are administered in accordance with prescribed directions.	
	Action taken as confirmed during the inspection: Daily stock balances are recorded for the majority of liquid medicines and those that were examined were found to be accurate. There were no discrepancies in the audits of liquid medicines that were completed during the inspection.	Met
Requirement 2 Ref: Regulation 13(4) Stated: First time	All administrations of Schedule 2 and Schedule 3 controlled drugs subject to safe custody requirements must be recorded in the controlled drug record book.  Action taken as confirmed during the inspection: All administrations of Schedule 2 and 3 controlled drugs had been recorded in the controlled drug record book.	Met

Last medicines mana	validation of compliance	
Recommendation 1 Ref: Standard 37 Stated: First time	The registered person should ensure that the recording system in place for patients who are prescribed 'when required' medicines for the treatment of distressed reactions is reviewed to ensure care plans are developed and the reason and outcome of each administration is recorded on every occasion.	Met
	Action taken as confirmed during the inspection: None of the current patients were prescribed medicines for distressed reactions. Staff were knowledgeable about the use of these medicines and the records that should be maintained.	
Recommendation 2 Ref: Standard 39 Stated: First time	The registered nurses should always cross reference the stock balances specified in the controlled drug record book with the actual stocks during each controlled drug stock reconciliation check.	Mat
	Action taken as confirmed during the inspection: No discrepancies were noted between the controlled drugs record book and the reconciliation checks. Staff confirmed that these records were cross referenced.	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 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. 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.

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.

Appropriate arrangements were in place for administering medicines in disguised form.

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. Medicine refrigerators and oxygen equipment were checked at regular intervals.

### **Areas for improvement**

No areas for improvement were identified during the inspection.

Number of requirements	0	Number of recommendations	0

### 4.4 Is care effective?

The sample of medicines examined had been administered in accordance with the prescriber's instructions. Some small discrepancies were noted in the personal medication record for one patient and a supply of eye drops for a second patient. This was highlighted to the registered manager who agreed to follow up after the inspection. 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.

None of the current patients were prescribed medicines for administration on a "when required" basis for the management of distressed reactions. However, staff knew how to recognise signs, symptoms and triggers which may cause a change in a patient's behaviour and were aware that this change may be associated with pain.

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 patient was comfortable. Staff advised that most of the patients 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.

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 running stock balances for medicines that were not contained in the blister pack system and extra records for the administration of transdermal patches.

Practices for the management of medicines were audited throughout the month by the staff and management. This included weekly audits on creams and thickeners. In addition, a quarterly audit was 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 for improvement**

No areas for improvement were identified during the inspection.

	Number of requirements	0	Number of recommendations	0
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### 4.5 Is care compassionate?

Patients were observed to be relaxed and comfortable in their surroundings and in their interactions with staff. Patients were treated courteously, with dignity and respect. Good relationships were evident.

It was not possible to observe the medicine round during this inspection; however staff were knowledgeable about patients' preferences and confirmed that medicines were administered discreetly.

Three patients completed the questionnaires. All of the responses indicated that they were "very satisfied" with how medicines were managed.

### Areas for improvement

No areas for improvement were identified during the inspection.

Number of requirements 0 Number of recommendations 0
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### 4.6 Is the service well led?

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.

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.

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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

There were no issues identified during this inspection, and a QIP is neither required, nor included, as part of this inspection report.

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.





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