



Unannounced Post-Registration Medicines Management Inspection Report 19 November 2018



Milesian Manor Nursing Home

Type of Service: Nursing Home
Address: 9 Ballyheifer Road, Magherafelt, BT45 5DX
Tel No: 028 7963 1842
Inspector: Catherine Glover

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 provider 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 which provides care for up to 23 patients with a range of needs as detailed in Section 3.0.

3.0 Service details

Organisation/Registered Provider: Macklin Care Homes Ltd Responsible Individual: Mr Brian Macklin	Registered Manager: Miss Caitriona Bridghin Doole
Person in charge at the time of inspection: Miss Caitriona Doole	Date manager registered: 25 April 2018
Categories of care: Nursing Home (NH) 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: 23 Patients to be accommodated on the first floor and seven identified bedrooms (201 - 207) on the second floor. The home is also approved to provide care on a day basis to six persons on the first floor and two persons on the second floor.

4.0 Inspection summary

The previous Milesian Manor nursing home was deregistered and patients were transferred to the newly built home which was registered separately. The building also accommodates Milesian Manor Residential Home.

An unannounced inspection took place on 19 November 2018 from 10.00 to 13.00.

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 since the pre-registration 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, medicine records and the management of controlled drugs.

Areas for improvement were identified in relation to recording the date of opening of medicines to facilitate the audit process and the cold storage of medicines.

Patients were relaxed and comfortable in the home. Good relationships with staff were evident.

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	2

Areas for improvement and details of the Quality Improvement Plan (QIP) were discussed with Miss Caitriona Dooley, 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.

4.2 Action/enforcement taken following the pre-registration inspection

The most recent inspection of the home was an unannounced care inspection undertaken on 12 June 2018. Other than those actions detailed in the QIP no further actions were required to be taken. 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 home registered

During the inspection the inspector met with two patients, one registered nurse and the registered manager.

We provided the registered manager with 10 questionnaires to distribute to patients and their representatives, for completion and return to RQIA. 'Have we missed you?' cards were left in the foyer of the home to inform patients/their representatives of how to contact RQIA, to tell us of their experience of the quality of care provided. Flyers providing details of how to raise any concerns were also left in the home.

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
- personal medication records
- medicine administration records
- medicines disposed of or transferred
- controlled drug record book
- medicine audits
- 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 12 June 2018

The most recent inspection of the home was an unannounced care inspection. The completed QIP was returned and approved 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

This Quality Improvement Plan was issued in respect of the previously registered Milesian Manor Nursing Home for the inspection conducted on 7 December 2017.

Areas for improvement from the last care inspection		
	Action required to ensure compliance with the Care Standards for Nursing Homes (2015)	Validation of compliance
<p>Area for improvement 1</p> <p>Ref: Standard 28</p> <p>Stated: First time</p>	<p>The registered person shall ensure that medicines not contained within the monitored dosage system are marked with the date of opening to facilitate audit.</p> <hr/> <p>Action taken as confirmed during the inspection:</p> <p>The date of opening had not been recorded on a significant number of medicines that were not contained within the monitored dosage system. This means that they could not be audited to ensure that they had been administered as prescribed.</p> <p>This area for improvement was stated for a second time.</p>	<p>Not met</p>

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. Refresher training in medicines management was provided in the last year. In relation to safeguarding, staff advised that they were aware of the regional procedures and who to report any safeguarding concerns to.

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

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 had been received into the home without delay.

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.

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. Staff were reminded that only controlled drugs should be stored in the controlled drugs cabinet and all unnecessary items should be removed.

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

Satisfactory arrangements were in place for the safe disposal of discontinued or expired medicines.

Medicines were generally 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 temperature of the medicine refrigerator should be checked daily. It was noted that there were a significant number of days when it had not been checked. An area for improvement was identified.

Areas of good practice

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

Areas for improvement

An area for improvement was identified in relation to recording the temperature of the medicines refrigerator daily.

	Regulations	Standards
Total number of areas for improvement	0	1

6.5 Is care effective?

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

The majority of medicines were contained in a monitored dosage system and had been administered as prescribed. As stated in Section 6.2, the date of opening had not been recorded on a significant number of medicines not contained within the monitored dosage system and these medicines could therefore not be audited. The area for improvement identified previously was stated for a second time.

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 medicines were due.

When a patient 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 patient’s behaviour and were aware that this change may be associated with pain. The reason for and the outcome of administration were recorded. It was noted that one patient did not have a care plan in place for the use of these medicines. It was agreed with the registered manager that this would be addressed following the inspection.

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 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.

Following discussion with the staff and observation of care notes, 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 the standard of record keeping and care planning.

Areas for improvement

No new 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 not observed during this inspection, however the registered nurses were knowledgeable about the patients' medicines and medical requirements.

It was found that there were good relationships between the staff and the patients. Staff were noted to be friendly and courteous; they treated the patients with dignity. It was clear from discussion and observation of staff, that the staff were familiar with the patients' likes and dislikes.

We spoke to two patients during the inspection. The patients were relaxed and comfortable in the home and said that they were happy living there. They said that the staff were kind and the food was good. We were shown the cinema room where a church service was being shown and the bar area where staff said families came to enjoy celebrations with relatives. Patients were also attending the barber who had organised a silent disco whilst they were waiting for their haircut.

None of the questionnaires that were issued for completion by patients and their representatives were returned within the specified time frame (two weeks).

Any comments from patients and their representatives in questionnaires received after the return date will be shared with the registered manager for information and action as required.

Areas of good practice

Staff listened to patients 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

We discussed arrangements in place in relation to the equality of opportunity for patients and the importance of staff being aware of equality legislation and recognising and responding to the diverse needs of patients. Arrangements are in place to implement the collection of equality data.

Written policies and procedures for the management of medicines were in place. They were not reviewed in detail 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 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. 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.

Audits were completed by staff in the home and comprehensive audit was completed by the deputy regional manager monthly. 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.

One area for improvement identified at the last medicines management inspection had not been addressed effectively. To ensure that this is fully addressed and the improvement sustained, it was suggested that the QIP should be regularly reviewed as part of the quality improvement process.

Staff confirmed that any concerns in relation to medicines management were raised with management. They advised that there were good working relationships within the home. The registered manager said that the senior management team were very supportive.

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 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

Areas for improvement identified during this inspection are detailed in the quality improvement plan (QIP). Details of the QIP were discussed with Miss Caitriona Dooley, Registered Manager, as part of the inspection process. The timescales commence from the date of inspection.

The registered provider/manager should note that if the action outlined in the QIP is not taken to comply with regulations and standards this may lead to further enforcement action including possible prosecution for offences. It is the responsibility of the registered provider to ensure that all areas for improvement identified 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 nursing 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.

7.1 Areas for improvement

Areas for improvement have been identified where action is required to ensure compliance with 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.

7.2 Actions to be taken by the service

The QIP should be completed and detail the actions taken to address the areas for improvement identified. The registered provider should confirm that these actions have been completed and return the completed QIP via the Web Portal for assessment by the inspector.

Quality Improvement Plan	
Action required to ensure compliance with the Department of Health, Social Services and Public Safety (DHSSPS) Care Standards for Nursing Homes, April 2015	
<p>Area for improvement 1</p> <p>Ref: Standard 28</p> <p>Stated: Second time</p> <p>To be completed by: 19 December 2018</p>	<p>The registered person shall ensure that medicines not contained within the monitored dosage system are marked with the date of opening to facilitate audit.</p> <p>Ref: 6.2</p> <p>Response by registered person detailing the actions taken: Staff meeting held and supervisions given out regarding the importance of dating and timing the opening of medications. Staff also attended further training session with Pharmacist.</p>
<p>Area for improvement 2</p> <p>Ref: Standard 30</p> <p>Stated: First time</p> <p>To be completed by: 19 December 2018</p>	<p>The registered person shall ensure that the refrigerator temperature is monitored and recorded daily.</p> <p>Ref: 6.4</p> <p>Response by registered person detailing the actions taken: Staff now bring the refridgerator temperatures to daily meeting with home manager to audit it being monitored and recorded daily. Importance of monitoring and recording covered in training session.</p>

Please ensure this document is completed in full and returned via the Web Portal



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