

Inspection Report

Name of Service: Brooklands Healthcare Magherafelt
Provider: Brooklands Healthcare Ltd
Date of Inspection: 28 November 2024

Information on legislation and standards underpinning inspections can be found on our website <https://www.rqia.org.uk/>

1.0 Service information

Registered Provider:	Brooklands Healthcare Ltd
Responsible Individual:	Mr Jarlath Conway
Registered Manager:	Mrs Deirdre Mary Monaghan
<p>Service Profile: Brooklands Healthcare Magherafelt is a nursing home registered to provide nursing care for up to 47 patients. The home operates over two floors.</p> <p>There is a residential care home which also occupies the ground floor and the registered manager for this home manages both services.</p>	

2.0 Inspection summary

An unannounced inspection took place on 28 November 2024, from 10.00am to 2.00pm. This was completed by a pharmacist inspector and focused on medicines management within the home.

The inspection was undertaken to evidence how medicines are managed in relation to the regulations and standards and to determine if the home is delivering safe, effective and compassionate care and is well led in relation to medicines management. The areas for improvement identified at the last care inspection were carried forward for review at the next inspection.

Mostly satisfactory arrangements were in place for the safe management of medicines. Medicines were stored securely. Medicine records and medicine related care plans were well maintained. There were effective auditing processes in place to ensure that staff were trained and competent to manage medicines and patients were administered their medicines as prescribed. However, an improvement was necessary in relation to the management of medicines requiring cold storage.

Whilst an area for improvement was identified, there was evidence that patients were being administered their medicines as prescribed.

Patients were observed to be relaxed and comfortable in the home and in their interactions with staff. It was evident that staff knew the patients well.

Details of the inspection findings, including areas for improvement carried forward for review at the next inspection, and the new area for improvement identified, can be found in the main body of this report and in the quality improvement plan (QIP) (Section 4.0).

RQIA would like to thank the staff for their assistance throughout the inspection.

3.0 The inspection

3.1 How we inspect

RQIA's inspections form part of our ongoing assessment of the quality of services. Our reports reflect how the home was performing against the regulations and standards, at the time of our inspection, highlighting both good practice and any areas for improvement. It is the responsibility of the service provider to ensure compliance with legislation, standards and best practice, and to address any deficits identified during our inspections.

To prepare for this inspection information held by RQIA about this home was reviewed. This included areas for improvement identified at previous inspections, registration information, and any other written or verbal information received from patients, relatives, staff or the commissioning trust.

Throughout the inspection process, inspectors seek the views of those living, working and visiting the home; and review/examine a sample of records to evidence how the home is performing in relation to the regulations and standards.

Through actively listening to a broad range of service users, RQIA aims to ensure that the lived experience is reflected in our inspection reports and quality improvement plans.

3.2 What people told us about the service and their quality of life

Staff expressed satisfaction with how the home was managed. They also said that they had the appropriate training to look after patients and meet their needs. They said that the team communicated well and the management team were readily available to discuss any issues and concerns should they arise.

Staff advised that they were familiar with how each patient liked to take their medicines and medicines were administered in accordance with individual patient preference. Staff also said that they prioritised patients who required pain relief and time-critical medicines during each medicine round.

RQIA did not receive any completed questionnaires or responses to the staff survey following the inspection.

3.3 Inspection findings

3.3.1 What arrangements are in place to ensure that medicines are appropriately prescribed, monitored and reviewed?

Patients in nursing homes should be registered with a general practitioner (GP) to ensure that they receive appropriate medical care when they need it. At times patients' needs may change and therefore their medicines should be regularly monitored and reviewed. This is usually done by a GP, a pharmacist or during a hospital admission.

Patients in the home were registered with a GP and medicines were dispensed by the community pharmacist.

Personal medication records were in place for each patient. These are records used to list all of the prescribed medicines, with details of how and when they should be administered. It is important that these records accurately reflect the most recent prescription to ensure that medicines are administered as prescribed and because they may be used by other healthcare professionals, for example, at medication reviews or hospital appointments.

The personal medication records reviewed were accurate and up to date. In line with best practice, a second member of staff had checked and signed the personal medication records when they were written and updated to confirm that they were accurate.

Copies of patients' prescriptions/hospital discharge letters were retained so that any entry on the personal medication record could be checked against the prescription.

All patients should have care plans which detail their specific care needs and how the care is to be delivered. In relation to medicines these may include care plans for the management of distressed reactions, pain, modified diets etc.

The management of distressed reactions, pain and thickening agents was reviewed. Care plans contained sufficient detail to direct the required care. Medicine records were well maintained. The audits completed at the inspection indicated that these medicines were administered as prescribed.

The management of warfarin was reviewed. Warfarin is a high risk medicine. Safe systems must be in place for blood monitoring and receipt of dosage directions to ensure that the correct dose is administered. Review of the warfarin administration records and audits completed at the inspection identified satisfactory arrangements were in place for the management of warfarin. However, one obsolete warfarin administration record had not been archived appropriately and remained in the medicines file. Only the current dosage directions should be available on the medicines file to ensure that nurses do not refer to obsolete directions in error and administer the wrong dose to the patient. This was highlighted to nursing staff for immediate correction.

3.3.2 What arrangements are in place to ensure that medicines are supplied on time, stored safely and disposed of appropriately?

Medicine stock levels must be checked on a regular basis and new stock must be ordered on time. This ensures that the patient's medicines are available for administration as prescribed. It is important that they are stored safely and securely so that there is no unauthorised access and disposed of promptly to ensure that a discontinued medicine is not administered in error.

Records reviewed showed that medicines were available for administration when patients required them. Staff advised that they had a good relationship with the community pharmacist and that medicines were supplied in a timely manner.

The medicine storage areas were observed to be securely locked to prevent any unauthorised access. They were tidy and organised so that medicines belonging to each patient could be easily located. Temperatures of medicine storage areas were monitored and recorded to ensure that medicines were stored appropriately.

Medicines which require cold storage must be stored between 2°C and 8°C to maintain their stability and efficacy. In order to ensure that this temperature range is maintained it is necessary to monitor the maximum and minimum temperatures of the medicines refrigerator each day and to then reset the thermometer. Appropriate action must be taken if the temperature recorded is outside the recommended range. The maximum temperature recorded was outside the recommended range for over one month and appropriate action had not been taken. An area for improvement was identified.

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

3.3.3 What arrangements are in place to ensure that medicines are appropriately administered within the home?

It is important to have a clear record of which medicines have been administered to patients to ensure that they are receiving the correct prescribed treatment.

A sample of the medicines administration records was reviewed. Most of the records were found to have been accurately completed. Records were filed once completed and were readily retrievable for audit/review.

Controlled drugs are medicines which are subject to strict legal controls and legislation. They commonly include strong pain killers. The receipt, administration and disposal of controlled drugs should be recorded in the controlled drug record book. There were satisfactory arrangements in place for the management of controlled drugs.

Management and staff audited the management and administration of medicines on a regular basis within the home. There was evidence that the findings of the audits had been discussed with staff and addressed. The date of opening was recorded on medicines to facilitate audit and disposal at expiry.

3.3.4 What arrangements are in place to ensure that medicines are safely managed during transfer of care?

People who use medicines may follow a pathway of care that can involve both health and social care services. It is important that medicines are not considered in isolation, but as an integral part of the pathway, and at each step. Problems with the supply of medicines and how information is transferred put people at increased risk of harm when they change from one healthcare setting to another.

A review of records indicated that satisfactory arrangements were in place to manage medicines at the time of admission or for patients returning from hospital. Written confirmation of prescribed medicines was obtained at or prior to admission and details shared with the GP and community pharmacy. Medicine records had been accurately completed and there was evidence that medicines were administered as prescribed.

3.3.5 What arrangements are in place to ensure that staff can identify, report and learn from adverse incidents?

Occasionally medicines incidents occur within homes. It is important that there are systems in place which quickly identify that an incident has occurred so that action can be taken to prevent a recurrence and that staff can learn from the incident. A robust audit system will help staff to identify medicine related incidents.

Management and staff were familiar with the type of incidents that should be reported. The medicine related incidents which had been reported to RQIA since the last inspection were discussed. There was evidence that the incidents had been reported to the prescriber for guidance, investigated and the learning shared with staff in order to prevent a recurrence.

The audits completed at the inspection indicated that medicines were being administered as prescribed.

3.3.6 What measures are in place to ensure that staff in the home are qualified, competent and sufficiently experienced and supported to manage medicines safely?

To ensure that patients are well looked after and receive their medicines appropriately, staff who administer medicines to patients must be appropriately trained. The registered person has a responsibility to check that they staff are competent in managing medicines and that they are supported.

There were records in place to show that staff responsible for medicines management had been trained and deemed competent.

It was agreed that the findings of this inspection would be discussed with staff to facilitate ongoing improvement.

4.0 Quality Improvement Plan/Areas for Improvement

An area for improvement has been identified where action is required to ensure compliance with Standards.

	Regulations	Standards
Total number of Areas for Improvement	3*	3*

* the total number of areas for improvement includes five which are carried forward for review at the next inspection.

The new area for improvement and details of the Quality Improvement Plan were discussed with Mrs Deirdre Mary Monaghan, Registered Manager, as part of the inspection process. The timescales for completion commence from the date of inspection.

Quality Improvement Plan	
Action required to ensure compliance with The Nursing Homes Regulations (Northern Ireland) 2005	
Area for improvement 1 Ref: Regulation 20 (1) (c) (l) Stated: First time To be completed by: 28 August 2024	The registered person must ensure all appropriate staff are in receipt of training in dysphagia. Action required to ensure compliance with this regulation was not reviewed as part of this inspection and this is carried forward to the next inspection. Ref: 2.0
Area for improvement 2 Ref: Regulation 27 (2) (d) Stated: First time To be completed by: 28 July 2024	The registered person must submit a time bound action plan detailing how and when improvements to the environment will be made in terms of homeliness and warmth. Action required to ensure compliance with this regulation was not reviewed as part of this inspection and this is carried forward to the next inspection. Ref: 2.0

<p>Area for improvement 3</p> <p>Ref: Regulation 27 (2) (t)</p> <p>Stated: First time</p> <p>To be completed by: 28 July 2024</p>	<p>The registered person shall submit a time bound action plan detailing how the one outstanding recommendation in the fire safety risk assessment, dated 16 January 2024, will be dealt with.</p> <hr/> <p>Action required to ensure compliance with this regulation was not reviewed as part of this inspection and this is carried forward to the next inspection.</p> <p>Ref: 2.0</p>
<p>Action required to ensure compliance with the Care Standards for Nursing Homes (December 2022)</p>	
<p>Area for improvement 1</p> <p>Ref: Standard 30</p> <p>Stated: First time</p> <p>To be completed by: Immediate and ongoing (28 November 2024)</p>	<p>The registered person shall ensure that the maximum, minimum and current temperatures of the medicines refrigerator are monitored and recorded daily and that appropriate action is taken if the temperature recorded is outside the recommended range 2-8°C.</p> <p>Ref: 3.3.2</p> <hr/> <p>Response by registered person detailing the actions taken: Treatment room fridges are monitored daily to ensure that medications are held as per manufacturers instructions. Records are maintained and regular audits completed. Staff nurses have completed training and supervision on ensuring fridge temperatures are maintained as required with appropriate action taken when out of recommended temperature range.</p>
<p>Area for improvement 2</p> <p>Ref: Standard 18 (1)</p> <p>Stated: First time</p> <p>To be completed by: 28 July 2024</p>	<p>The registered person shall review the identified bed rail assessment in consultation with the patient's aligned named worker, in accordance with current safety guidance.</p> <hr/> <p>Action required to ensure compliance with this standard was not reviewed as part of this inspection and this is carried forward to the next inspection.</p> <p>Ref: 2.0</p>
<p>Area for improvement 3</p> <p>Ref: Standard 47 (1)</p> <p>Stated: First time</p> <p>To be completed by: 28 July 2024</p>	<p>The registered person shall ensure that the two identified lounge doors close securely in accordance with fire safety.</p> <hr/> <p>Action required to ensure compliance with this standard was not reviewed as part of this inspection and this is carried forward to the next inspection.</p> <p>Ref: 2.0</p>

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