

# Inspection Report

**Name of Service:** Woodgrove Care Home

**Provider:** Beaumont Care Homes Limited

**Date of Inspection:** 9 October 2025

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

## 1.0 Service information

<b>Organisation/Registered Provider:</b>	Beaumont Care Homes Limited
<b>Responsible Individual:</b>	Mrs Ruth Burrows
<b>Registered Manager:</b>	Mrs Zoe Patterson
<b>Service Profile:</b> Woodgrove Care Home is a nursing home registered to provide nursing care for up to 32 patients. The home specialises in caring for people with physical disability and nursing needs relating to old age and terminal illness. Patient bedrooms are located on the ground and first floor. There are a range of communal areas throughout the home and patients have access to an enclosed garden.	

## 2.0 Inspection summary

An unannounced inspection took place on 9 October 2025, from 10.10am to 2.00pm. The inspection 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 inspection also reviewed the area for improvement identified at the last medicines management inspection. The area for improvement identified at the last care inspection was 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, improvement was necessary in relation to cold storage.

The area for improvement in relation to controlled drugs identified at the last medicines management inspection was assessed as met. Details of the inspection findings, including the area 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).

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.

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.

#### **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. They stated medication rounds were tailored to respect each individual's preferences, needs and timing requirements.

No completed questionnaires or responses to the staff survey were received 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. A small number of minor discrepancies were highlighted for immediate corrective action and on-going vigilance.

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.

Patients will sometimes get distressed and will occasionally require medicines to help them manage their distress. It is important that care plans are in place to direct staff when it is appropriate to administer these medicines and that records are kept of when the medicine was given, the reason it was given and what the outcome was. If staff record the reason and outcome of giving the medicine, then they can identify common triggers which may cause the patient's distress and if the prescribed medicine is effective for the patient.

The management of medicines, prescribed on a 'when required' basis for distressed reactions, was reviewed. Directions for use were clearly recorded on the personal medication record and patient-centred care plans were in place for the majority of patients prescribed these medicines. One patient required a care plan to be implemented and the manager provided assurance that this would be implemented immediately. Staff knew how to recognise changes in a patient's behaviour and were aware that this change may be associated with pain and other factors. Records of administration included the reason for and outcome of each administration.

The management of pain was discussed. Staff advised that they were familiar with how each patient expressed their pain and that pain relief was administered when required. Care plans were in place and reviewed regularly.

Some patients may need their diet modified to ensure that they receive adequate nutrition. This may include thickening fluids to aid swallowing and food supplements in addition to meals. Care plans detailing how the patient should be supported with their food and fluid intake should be in place to direct staff. All staff should have the necessary training to ensure that they can meet the needs of the patient.

The management of thickening agents was reviewed. Speech and language assessment reports and care plans were in place. Records of prescribing and administration which included the recommended consistency level were maintained.

Care plans were in place when patients required insulin to manage their diabetes. There was sufficient detail to direct staff if the patient's blood sugar was outside of the recommended range. Staff were reminded that in use insulin pen devices must be individually labelled and the date of opening must be recorded to facilitate audit and disposal at expiry.

### **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 area was observed to be securely locked to prevent any unauthorised access. It was tidy and organised so that medicines belonging to each patient could be easily located. The temperature of the medicine storage area was monitored and recorded to ensure that medicines were stored appropriately. Satisfactory arrangements were in place for the storage of controlled drugs and the safe disposal of medicines.

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. Temperature of the medicine refrigerator were monitored and recorded daily but appropriate action had not been taken when the temperature recorded was outside the recommended range. An area for improvement was identified.

### **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. 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.

Occasionally, patients may require their medicines to be crushed or added to food/drink to assist administration. To ensure the safe administration of these medicines, this should only occur following a review with a pharmacist or GP and should be detailed in the patient's care plan. Written consent and care plans were in place when this practice occurred.

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 inspector signposted staff to the RQIA provider guidance in relation to the statutory notification of medication related incidents available on the RQIA website.

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 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 the necessary improvements.

## 4.0 Quality Improvement Plan/Areas for Improvement

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

	Regulations	Standards
<b>Total number of Areas for Improvement</b>	0	2*

\* the total number of areas for improvement includes one which was carried forward for review at the next inspection.

The new area for improvement and details of the Quality Improvement Plan were discussed with Mrs Zoe Patterson, Registered Manager, as part of the inspection process. The timescale for completion commences from the date of inspection.

<b>Quality Improvement Plan</b>	
<b>Action required to ensure compliance with the Care Standards for Nursing Homes (December 2022)</b>	
<p><b>Area for improvement 1</b></p> <p><b>Ref:</b> Standard 30</p> <p><b>Stated:</b> First time</p> <p><b>To be completed by:</b> Immediate and ongoing (9 October 2025)</p>	<p>The registered person shall ensure that the maximum, minimum and current temperature of the medicines refrigerator is monitored and recorded daily and that appropriate action is taken if the temperature recorded is outside the recommended range of 2-8°C.</p> <p>Ref: 3.3.2</p> <hr/> <p><b>Response by registered person detailing the actions taken:</b> Following the inspection, the fridge was defrosted and cleaned. Staff have been reminded of their responsibilities on ensuring Max/Min temperatures are recorded on a daily basis. Temperatures have been reviewed and are currently recording at correct temperatures on daily checks. Compliance will be checked as part of the internal governance audits and during the completion of the Reg 29 audit.</p>
<p><b>Area for improvement 2</b></p> <p><b>Ref:</b> Standard 44</p> <p><b>Stated:</b> First time</p> <p><b>To be completed by:</b> 9 May 2025</p>	<p>The registered person shall ensure that the source of the malodour in the identified room is detected and effectively managed.</p> <hr/> <p><b>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.</b></p> <p>Ref: 2.0</p>

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



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