

Percentage of long stay home care patients whose primary informal caregiver experienced distress, anger or depression in relation to their caregiving role or were unable to continue in that role (Retired)

Alternate Name*

Percentage of long-stay home care patients with a primary unpaid caregiver whose caregiver is unable to continue in caring activities or expresses feelings of distress, anger or depression

INDICATOR DESCRIPTION

Description*

This indicator describes the percentage of long-stay home care patients whose primary informal caregiver is unable to continue in caring activities and/or expresses feelings of distress, anger or depression in patients who had at least one unpaid caregiver. A lower percentage is better.

HQO Reporting tool/product

Public reporting

Dimension*

Patient-centred

Type*

Outcome

DEFINITION AND SOURCE INFORMATION

Unit of Measurement*

Percentage

Calculation Methods*

Numerator divided by denominator times 100

Numerator (short description i.e. not inclusions/exclusions)*

All clients with a primary caregiver who is unable to continue in caring activities AND/OR All clients with a primary caregiver who expresses feelings of distress, anger or depression

Denominator (short description i.e. not inclusions/exclusions)*

All clients with a completed RAI-HC assessment who have primary caregiver



Adjustment (risk, age/sex standardization)- generalized

None

Data Source

Home Care Reporting System (HCRS)

Data provided to HQO by

Canadian Institute for Health Information (CIHI)

Reported Levels of comparability /stratifications (defined)

Time

Region

OTHER RELEVANT INFORMATION

Caveats and Limitations

The results are derived from RAI-HC assessment information in the HCRS database. Therefore, the results are representative of long-stay home care patients receiving publicly funded home care (i.e., home care patients who require care for more than 60 days represent approximately half of publicly funded home care patients. The remaining population are short-stay home care patients who require short-term care while they recover from injury or surgery.) Distress, anger, or depression and ability to continue providing care are subjective measures. Results may underestimate true rates of distress if some informal caregivers do not want to admit that they feel burdened by having to care for a friend or family member or if they do not exhibit obvious signs of distress. Similarly, cultural differences exist that can affect one's perception of distress caused by caring for a friend or family member since it is simply accepted as something that should be done, regardless of the stress it may cause. Data are not risk adjusted for factors associated with rates of distress, such as cognitive impairment in the patient, hours of informal care provided by the caregiver, or hours of formal care provided to the patient. Includes patients of varying levels of complexity, including palliative patients.

Comments Summary

This indicator was jointly developed by interRAI and the Canadian Institute for Health Information (CIHI). The caregiver may live with the patient or the caregiver may visit the patient regularly or respond to needs that the patient has. A home care patient may have more than one person providing unpaid support. The primary informal caregiver is the person who is most helpful to the patient and the person who the patient can rely most upon. The questions in the RAI-HC asking about the caregiver status are meant to assess the reserve of the informal caregiver support system. The questions are posed to the home care patient and informal caregiver separately. The current situation and the projected future needs are both considered. G2a. A caregiver is unable to continue in caring activities The caregiver, client, or assessor believes that a caregiver(s) is not able to continue in caring activities. This can be for any reason, for example: lack of desire to continue, geographically inaccessible, other competing requirements (child care, work requirements), and personal health issues. G2c. Primary caregiver expresses feelings of distress Primary caregiver expresses, by any means, that he/she is distressed, angry, depressed, or in conflict because of caring for the client.

TAGS

TAGS*

Home Care

Percentage of long stay home care patients whose primary informal caregiver experienced distress, anger or depression in



Outcome Patient Reported Measures Other Patient-centred Home Care Reporting System (HCRS)

PUBLISH

PUBLISH DATETIME*

17/04/2018 14:55:00