

NHS Lanarkshire Care Homes Protocol Group

Guideline for review of cognitive enhancing (anti-dementia) medication for clinicians

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GUIDELINE FOR REVIEW OF COGNITIVE ENHANCING (ANTI-DEMENTIA) MEDICATION FOR CLINICIANS i.e. Donepezil (Aricept), Galantamine (Reminyl), Rivastigmine (Exelon), Memantine (Ebixa)

For patients prescribed anti-dementia medication, it is appropriate to review the prescription when there is a change in the patient's presentation e.g. evidence of increased frequency of agitation, when there are indicators of poor prognosis or when undertaking polypharmacy review.

Changes to medication must be discussed with the patient if they have capacity. **Where capacity is absent and there is an existing legal proxy i.e. welfare attorney or guardian, they must be consulted/informed of any changes.**

If the patient lacks capacity and if there is no formal legal proxy, the principles of the Adults with Incapacity (Scotland) Act 2000 apply and treatment options should be discussed with relevant others, such as next of kin, carer or patient advocate. In either circumstance, an appropriate Section 47 certificate of incapacity is required.

The following guidance will support the review process but further advice can be sought from Old Age Psychiatry services if required.

There is a change in presentation e.g. new/increased symptoms of stress/distress.

Rule out other possible causes of stress/distress e.g. pain, infection. Refer to management of stress and distress guideline.

Titrate the dose down gradually leaving 4 weeks between dose changes.

During the 4 week period between dose changes monitor for any functional or cognitive deterioration.

If there is deterioration the patient should be returned to the previous dose as quickly as possible.

If there is no change, continue dose reduction until the medication is stopped.

If stress/distress symptoms have not resolved, refer back to Old Age Psychiatry.

There is a marked deterioration in cognitive function or there are poor prognostic indicators.

YES

NO

LEAVE UNCHANGED

Titrate the dose down gradually leaving 4 weeks between dose changes.

During the 4 week period monitor for any functional or cognitive deterioration.

If there is deterioration the patient should be returned to the previous dose as quickly as possible.

If there is no change, continue dose reduction

If the patient is approaching the end of life or is receiving palliative care, it is appropriate to reduce or stop cognitive enhancing medication taking into consideration the individual clinical circumstances.

On a patient's admission to hospital, where possible, these medicines should not be withdrawn abruptly.
The Old Age Psychiatry liaison team can be contacted for advice if required.

Decisions to change, reduce or withdraw cognitive enhancing medication should be recorded in the patient's care plan or GP record and a plan for review clearly documented.