Have last-observation-carried-forward analyses caused us to favour more toxic dementia therapies over less toxic alternatives? A systematic review

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ABSTRACT

Background: Intention-to-treat analysis is used in the analysis of randomized controlled trials to preserve trial power in the presence of missing subject data as well as to control for both known and unknown confounding factors. One form of intention-to-treat analysis is last-observation-carried-forward (LOCF). Concerns exist regarding whether it is appropriate to use LOCF in analyses involving progressive conditions or in situations where missing data are nonrandom (e.g., subjects drop out because of treatment side effects or differing disease severity).

Objective: To examine the use of intention-to-treat imputation of missing data techniques, and specifically LOCF, in randomized controlled trials of the use of cholinesterase inhibitors and memantine to treat Alzheimer’s disease, vascular dementia, mixed dementia and mild cognitive impairment.

Methods: We conducted a systematic electronic search of MEDLINE and the Cochrane Central Register of Controlled Trials from 1984 to 2008 for double-blinded, randomized controlled trials of cholinesterase inhibitors or memantine that examined progressive symptoms in Alzheimer’s disease, vascular dementia, mixed dementia and mild cognitive impairment. We collected data on the use of intention-to-treat and non-intention-to-treat analyses and on contraindications to the use of LOCF analysis and we performed quality assessments of included trials.

Results: Of the 57 studies that met the inclusion criteria, 12 did not report intention-to-treat analyses. Of the 34 studies that employed LOCF as the only form of intention-to-treat analysis, 24 reported conditions that could produce biased LOCF analyses favouring the drug under study. The latter finding was more common in cholinesterase inhibitor trials than in memantine studies.

Conclusions: The published results of some randomized controlled trials of dementia drugs may be inaccurate (i.e., drug effectiveness may be exaggerated) or invalid (i.e., there may be false-positive results) because of bias introduced through the inappropriate use of LOCF analyses. This bias favours cholinesterase inhibitors, potentially preventing funding of and patient access to less toxic treatment options such as memantine. Licensing agencies should consider whether to accept LOCF analyses in research on dementias and other chronic progressive conditions.

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Competing interests: None declared.

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Open Medicine 2009 3(2):31-50
It is estimated that 24.3 million people worldwide suffer from dementia and that annual costs for Alzheimer’s disease are as high as $155 billion in the United States (1996 US dollars).\textsuperscript{1,2} One potential way to decrease the negative impact of dementia on people with this condition, on their families and on societies is to optimize the use of dementia medications,\textsuperscript{3} with due consideration of both their effectiveness and their toxicity.

The effectiveness of most medications is tested via randomized controlled trials (RCTs). It is inevitable that some participants drop out of such studies before they are completed. Unfortunately, if analyses include only participants who remain in the trial, then study power is lost and erroneous conclusions may be generated. The principle of intention-to-treat (ITT) analysis, in which all patients are included in the analysis according to the group to which they were assigned at randomization, has become the accepted standard for the analysis of RCTs to try to counteract this problem.\textsuperscript{3} The strength of ITT analysis is that it not only preserves power but also promotes balance between treatment groups for both known and unknown confounders, thereby preserving the benefits of randomization.

Ideally, all possible data are collected on all subjects, including those who drop out of the study; however, this is not always possible. In order for ITT approaches to analyze all patients randomly assigned to a group, several methods to impute missing data have been developed.\textsuperscript{3–10} Unfortunately, no statistical strategy can deal fully with all the different combinations of reasons for dropping out, dropout rates and different disease courses. At best, these techniques to impute missing data are educated estimates. One commonly employed technique to impute missing data is last-observation-carried-forward (LOCF), also known as end-point analysis.

LOCF substitutes subjects’ missing outcomes with the last measurement taken before they dropped out. It requires that 2 basic assumptions be met: the subjects’ responses would have been constant from the last observed value (i.e., the point at which they dropped out) to the end point of the trial; and, missing values are missing completely at random (i.e., dropout is not related to variables such as drug side effects, group assignment, disease severity or symptoms).\textsuperscript{5,7} Authors have highlighted 3 factors that cause the second condition to be breached in a manner that intro-

![Figure 1: Differential last observation carried forward (LOCF) bias when there are more or earlier dropouts in the treatment group than in the control group. (Effect measured by LOCF \(c-d > a-b\), resulting in an exaggerated positive effect, biased in favour of treatment.)](image)
roduces bias that will exaggerate the effectiveness of treatments as estimated by LOCF analyses; these include earlier dropouts or greater dropout rates in the treatment group and more rapid disease progression in subjects who drop out of the treatment group. These factors result in more subjects who drop out of the treatment group having their decline artificially frozen at an earlier stage of disease, thereby potentially biasing results in favour of the drug under study (i.e., overestimating effectiveness relative to the placebo). By extension, study results may also be biased against the drug under study (i.e., underestimating effectiveness) if there are earlier dropouts or greater dropout rates in the control group or if there are subjects whose disease progresses more rapidly among those who drop out of the control group (Figures 1 and 2).

Since 1998, researchers have expressed concern that the use of LOCF in dementia drug trials contravenes the assumption of disease stability and the assumption of random missing data and hence risks generating biased results. To better understand the significance of these concerns in dementia research we systematically reviewed the use of ITT and LOCF analyses, contraindications to the use of LOCF analysis, and the use of non-ITT analyses in RCTs of drugs approved for the treatment of Alzheimer’s disease, vascular dementia, mixed dementia and mild cognitive impairment in Canada (i.e., cholinesterase inhibitors such as donepezil, rivastigmine and galantamine, and the N-methyl-D-aspartate (NMDA) receptor antagonist memantine).

Methods

We performed an electronic literature search of MEDLINE and the Cochrane Central Register of Controlled Trials from January 1984 (the year of publication of the McKhann criteria for Alzheimer’s disease) to February 2008 using the OVID search interface. The search strategy included the following terms: randomized controlled trials, dementia, Alzheimer, vascular dementia, mixed dementia, donepezil, Aricept, rivastigmine, Exelon, galantamine, Reminyl, memantine, Ebixa and cholinesterase inhibitor.

The principal investigator reviewed titles and abstracts to select an overly inclusive list of potential articles to be subjected to a full review of text and reference sections to identify relevant RCTs. The full

**Figure 2:** Differential last observation carried forward (LOCF) bias when there are more or earlier dropouts in the control group than in the treatment group. (Effect measured by LOCF [c–d] < true effect [a–b], resulting in an underestimate of effectiveness, biased against treatment.)
text of selected RCT reports was then independently re-
viewed by 2 certified specialists in geriatric medicine
with clinical expertise in dementia, formal research
methodological training and recognized expertise in the
review of dementia drug trials to determine which RCT
reports met the inclusion criteria for the systematic re-
view.

**Inclusion criteria.** We included double-blinded, ran-
domized controlled trials of cholinesterase inhibitors or
memantine that examined progressive symptoms (e.g.,
cognition, function) in Alzheimer’s disease, vascular de-
mentia, mixed dementia or mild cognitive impairment
and that employed DSM-IV (Diagnostic and Statistical
Manual of Mental Disorders) or NINCDS–ADRDA
(National Institute of Neurological and Communicative
Disorders and Stroke – Alzheimer’s Disease and Re-
lated Disorders Association) criteria for Alzheimer’s
disease or NINDS–AIREN (National Institute of Neuro-
logical Disorders and Stroke – Association Interna-
tionale pour la Recherche et l’Enseignement en
Neurosciences) criteria for vascular dementia. Trials of
cholinesterase inhibitors not currently licensed in
Canada (tacrine, metrifonate) were not reviewed. The
systematic review was restricted to studies with full trial
reports published in English-language peer-reviewed
journals. The diagnostic criteria for mild cognitive impairment were not specified, as they were in develop-
ment when the relevant studies were published.

Although the reference sections of open-label stud-
ies, reviews, meta-analyses, commentaries, editorials,
studies of pooled data from previous studies and toler-
ability and safety studies were searched for relevant
RCTs, the articles themselves were not included in the
systematic review. Subgroup analyses and secondary or
retrospective analyses were also excluded.

**Figure 3: Selection of studies for review.**
Data collection. Data collected included publication details, investigative site locations, funding, drug comparators, drug doses, diagnostic criteria employed, type(s) of analysis employed, discussion of the limitations of the forms of analysis employed, dropout characteristics (e.g., number, timing, patient characteristics, reasons for dropout), contraindications to the use of LOCF and the results of each study’s primary and secondary outcome measures.

The 2 previously mentioned reviewers independently extracted data from all included studies and then met to review their findings and discuss discrepancies. When consensus could not be achieved, discrepancies were forwarded to a third party for independent review.

Results

Of the 1146 articles identified by the search strategies, 191 papers (including RCT reports, reports of nonrandomized trials, commentaries, systematic reviews and meta-analyses) were selected for full text and reference section review. Of these, 57 RCT reports met the eligibility criteria for systematic review (Fig. 3).5-12,14-20,22,26-79

Reviewer agreement. Although agreement on abstracted items was not formally measured, the methods employed resulted in consensus on almost all abstracted items. After the reasons for different ratings were explained (differences were mostly a result of difficulty finding the relevant data in the studies reviewed), the reviewers agreed on all but 5 final ratings. These were arbitrated by a third party. The kappa score, if it had been measured, would have been unusually high.

Trial characteristics. Details of the 57 included trials are provided in Tables 1 and 2. Forty-five studies enrolled patients with Alzheimer’s disease (21 involved donepezil, 11 rivastigmine [1 of these studies was a donepezil–rivastigmine comparison study], 7 galantamine and 6 memantine), 8 studies enrolled patients with vascular dementia or mixed dementia (3 involved donepezil, 3 galantamine and 2 memantine) and 4 studies enrolled patients with mild cognitive impairment (2 involved donepezil, 1 rivastigmine and 1 galantamine).

In 40 trials there was an explicit statement of pharmaceutical industry funding. In 6 trials industry funding was implied (the authors were pharmaceutical industry employees but the source of funding was not explicitly stated). Three studies were funded by industry in partnership with public funders, and 4 studies were entirely publicly funded (Table 1). The source of funding for 4 studies could not be determined. All 57 study reports were rated as demonstrating high-quality methodology with a Jadad–Schultz score greater than or equal to 3 (Table 1).

Reporting of dropouts. Data on dropouts are provided in Tables 3 and 4. Dropouts were described in 94% of cholinesterase inhibitor studies and 100% of memantine trials. Seven of the 49 cholinesterase inhibitor trials (14%) and none of the memantine trials reported data on the timing of dropouts. The reasons for dropout were often difficult to discern, as many were described as adverse events that might have been due to drug side effects but were not reported as such. Cholinesterase inhibitor studies were more likely than memantine studies to demonstrate a higher dropout rate in the treatment group than in the control group (73% of cholinesterase inhibitor studies v. 25% of memantine studies). When cholinesterase inhibitor studies were combined there was a higher dropout rate in the treatment group than in the control group (23.2% in the treatment group v. 16.8% in the control group) (Table 4). When memantine trials were combined the opposite pattern was noted: there were fewer dropouts in the treatment group than in the control group (14.6% in the treatment group v. 18.5% in the control group) (Table 4). Ten studies (18%) discussed potential bias associated with dropouts.

Types of non-ITT analyses conducted. The most common non-ITT analysis (employed in 35 trials) was observed case analysis (Table 2). Other forms of non-ITT analysis included fully evaluable population analysis (5 RCTs), treatment per protocol analysis (5 RCTs) and completer analysis (3 RCTs) (Table 2).

Types of ITT analyses conducted. Twelve (21%) of the 57 studies did not identify the type of analysis performed (5 studies) or performed only non-ITT analysis (7 studies) (Table 2). Of the 45 studies in which an identifiable form of ITT analysis was performed, 42 (93%) employed LOCF (Table 2). Thirty-four of the trials in which ITT analysis was performed (76%) relied on LOCF as the only form of ITT analysis (Table 2).

Ten of the 57 studies (17.5%) reported employing ITT techniques other than LOCF (Table 2); 6 of 49 cholinesterase inhibitor studies (12%) and 4 of 8 memantine studies (50%) employed ITT techniques other than LOCF. The 6 alternative approaches for ITT imputation of missing data included the following: replacement of missing values with the mean changes observed in the placebo group;42,58 time-response relationship for change in ADAS-cog/11 (the Alzheimer Disease Assessment Scale – Cognitive Subscale 11-item) score analyzed using generalized linear modelling;50,52 mixed-effects modelling;56,63 mixed-models repeated measures;76,79 replacement of missing data with worst ranks57 and sensitivity analyses consisting of a number of simulations.58

Of the 42 studies employing LOCF, only 8 reported performing another type of ITT analysis to confirm the
| Study                          | Funding source(s) | Medication(s) studied* | Indication | Sample size(s), no. of patients | Jadad-Schulz quality score |
|-------------------------------|-------------------|------------------------|------------|--------------------------------|-----------------------------|
| Rogers et al. (1996)          | Industry          | Donepezil (1, 3, 5 mg) | AD         | Controls 40, Active comparators 121 | 4                           |
| Rogers et al. (1998)          | Industry          | Donepezil (5, 10 mg)   | AD         | Controls 153, Active comparators 315 | 4                           |
| Rogers et al. (1998)          | Industry          | Donepezil (5, 10 mg)   | AD         | Controls 162, Active comparators 311 | 4                           |
| Agid et al. (1998)            | Industry          | Rivastigmine (4, 6 mg) | AD         | Controls 133, Active comparators 269 | 3                           |
| Corey-Bloom et al. (1998)     | Industry          | Rivastigmine (1-4, 6-12 mg) | AD         | Controls 235, Active comparators 464 | 5                           |
| Burns et al. (1999)           | Industry          | Donepezil (5, 10 mg)   | AD         | Controls 274, Active comparators 544 | 4                           |
| Forette et al. (1999)         | Industry          | Rivastigmine (6-9 mg; BID or TID) | AD         | Controls 24, Active comparators 90 | 4                           |
| Rösler et al. (1999)          | Industry          | Rivastigmine (1-4, 6-12 mg) | AD         | Controls 239, Active comparators 486 | 5                           |
| Winblad et al. (1999)         | Not reported      | Memantine (10 mg)      | AD, VD     | Controls 84, Active comparators 82 | 3                           |
| Greenberg et al. (2000)       | Public            | Donepezil (5 mg) cross-over | AD         | Controls 30, Active comparators 30 | 5                           |
| Homma et al. (2000)           | Industry          | Donepezil (5 mg)       | AD         | Controls 129, Active comparators 134 | 3                           |
| Kumar et al. (2000)           | Industry (authors employed) | Rivastigmine (1-4, 6-12 mg) | AD         | Controls 103, Active comparators 216 | 3                           |
| Raskind et al. (2000)         | Industry          | Galantamine (24, 32 mg) | AD         | Controls 213, Active comparators 423 | 5                           |
| Tariot et al. (2000)          | Industry          | Galantamine (16, 24 mg) | AD         | Controls 286, Active comparators 552 | 5                           |
| Wilcock et al. (2000)         | Industry          | Galantamine, (24, 32 mg) | AD         | Controls 215, Active comparators 438 | 5                           |
| Feldman et al. (2001)         | Industry          | Donepezil (5-10 mg)    | AD         | Controls 146, Active comparators 144 | 5                           |
| Mohs et al. (2001)            | Industry          | Donepezil (5-10 mg)    | AD         | Controls 217, Active comparators 214 | 4                           |
| Tariot et al. (2001)          | Industry          | Donepezil (5-10 mg)    | AD         | Controls 105, Active comparators 103 | 4                           |
| Thomas et al. (2001)          | Not reported      | Donepezil, vitamin E   | AD         | Controls 20, Active comparators 20 | 4                           |
| Winblad et al. (2001)         | Industry          | Donepezil (5-10 mg)    | AD         | Controls 144, Active comparators 142 | 5                           |
| Rockwood et al. (2001)        | Industry          | Galantamine (24-32 mg) | AD         | Controls 125, Active comparators 261 | 5                           |
| Study                          | Funding source(s)      | Medication(s) studied* | Indication | Sample size(s), no. of patients | Jadad-Schultz quality score |
|-------------------------------|------------------------|------------------------|------------|---------------------------------|-----------------------------|
| Wilkinson et al. (2001)†4     | Industry               | Galantamine (18, 24, 36 mg) | AD         | 87                              | 198                         | 5                           |
| Doraiswamy et al. (2002)†8     | Industry               | Rivastigmine (1-4, 6-12 mg) | AD         | Not reported                    | Not reported                | 3                           |
| Pratt et al. (2002)60          | Industry (authors employed) | Donepezil (5, 10 mg)  | VD         | 290                             | 603                         | 5                           |
| Erkinjuntti et al. (2002)63    | Industry               | Galantamine (24 mg)    | MC, VD     | 196                             | 396                         | 4                           |
| Orgogozo et al. (2002)65       | Industry               | Memantine (20 mg)      | VD         | 141                             | 147                         | 3                           |
| Wilcock et al. (2002)66        | Industry (authors employed) | Memantine (20 mg)      | VD         | 271                             | 277                         | 4                           |
| Krishnan et al. (2003)68       | Industry               | Donepezil (10 mg)      | AD         | 33                              | 34                          | 5                           |
| Tune et al. (2003)72           | Industry               | Donepezil (10 mg)      | AD         | 14                              | 14                          | 3                           |
| Reisberg et al. (2003)18       | Industry, public       | Memantine (20 mg)      | AD         | 126                             | 126                         | 5                           |
| Black et al. (2003)61          | Industry               | Donepezil (5, 10 mg)   | VD         | 199                             | 404                         | 5                           |
| Wilkinson et al. (2003)19      | Industry               | Donepezil (5, 10 mg)   | VD         | 193                             | 423                         | 5                           |
| AD 2000 Collaborative Group (2004)8 | Public               | Donepezil (5-10 mg)    | Phase 1    | 244                             | 242                         | 4                           |
| Holmes et al. (2004)38         | Industry               | Donepezil (10 mg)      | AD         | 55                              | 41                          | 5                           |
| Seltzer et al. (2004)40        | Industry               | Donepezil (10 mg)      | AD         | 57                              | 96                          | 4                           |
| Tariot et al. (2004)29         | Industry               | Memantine (20 mg)      | AD†        | 201                             | 202                         | 5                           |
| Bullock et al. (2004)19        | Industry (authors employed) | Galantamine (24 mg)    | MD         | 97                              | 188                         | 4                           |
| Salloway et al. (2004)67       | Industry               | Donepezil (10 mg)      | MCI        | 137                             | 133                         | 4                           |
| Karaman et al. (2005)19        | Not reported           | Rivastigmine (6-12 mg) | AD         | 20                              | 24                          | 4                           |
| Bullock et al. (2005)19        | Industry               | Rivastigmine (3-12 mg), donepezil (5-10 mg) | AD | 499                           | (donepezil) | 495 | 5 |
| Brodaty et al. (2005)55        | Industry (authors employed) | Placebo, immediate-release galantamine (16-24 mg), prolonged-release galantamine (16-24 mg) | AD | 324 | 647 | 5 |
| Study                                      | Funding source(s)     | Medication(s) studied* | Indication | Sample size(s), no. of patients | Jadad-Schultz quality score |
|--------------------------------------------|-----------------------|------------------------|------------|-------------------------------|-----------------------------|
| Petersen et al. (2005)†                    | Industry, public      | Donepezil (10 mg)      | MCI        | Controls 259, Active comparators 253 | 4                           |
| Koontz and Baskys (2005)†                  | Industry              | Galantamine (24 mg)    | MCI        | Controls 11, Active comparators 8  | 4                           |
| Dos Santos Moraes et al. (2006)†           | Public                | Donepezil (10 mg)      | AD         | Controls 18, Active comparators 17 | 4                           |
| Johanssen et al. (2006)††                  | Industry              | Donepezil (10 mg)      | AD         | Controls 103, Active comparators 99 | 4                           |
| Winblad et al. (2006)†                     | Industry              | Donepezil (5-10 mg)    | AD         | Controls 120, Active comparators 128 | 5                           |
| Rockwood et al. (2006)††                   | Industry, public      | Galantamine (16-24 mg) | AD         | Controls 66, Active comparators 64 | 5                           |
| Mazza et al. (2006)†                       | Not reported          | Donezepil (5 mg), ginko | AD        | Controls 26, Active comparators 25 | 5                           |
| Peskind et al. (2006)†                     | Industry              | Memantine (20 mg)      | AD         | Controls 202, Active comparators 201 | 5                           |
| Auchus et al. (2007)†                      | Industry (authors employed) | Galantamine (flexible dose) | VD | Controls 391, Active comparators 397 | 4                           |
| Bakchine and Loft (2007)††                 | Industry              | Memantine (20 mg)      | AD         | Controls 152, Active comparators 318 | 4                           |
| Black et al. (2007)†                       | Industry              | Donepezil (10 mg)      | AD         | Controls 167, Active comparators 176 | 5                           |
| Feldman et al. (2007)†                     | Industry              | Rivastigmine (3-12 mg) | MCI        | Controls 509, Active comparators 508 | 5                           |
| Feldman et al. (2007)†                     | Industry              | Rivastigmine (2-12 mg; BID or TID) | AD | Controls 222, Active comparators 229 (BID group), 227 (TID group) | 5                           |
| Mowla et al. (2007)†                       | Public                | Placebo, rivastigmine (6-12 mg), rivastigmine (6-12 mg) + fluoxetine | AD | Controls 40, Active comparators 41 rivastigmine alone | 4                           |
| Van Dyck et al. (2007)††                   | Industry              | Memantine (20 mg)      | AD         | Controls 172, Active comparators 178 | 5                           |
| Winblad et al. (2007)†                     | Industry              | Placebo, rivastigmine patch (10 cm² or 20 cm²), rivastigmine tablet (12 mg) | AD | Controls 302, Active comparators 293 (10 cm² patch), 303 (20 cm² patch), 297 (12 mg tablet) | 5                           |

AD = Alzheimer’s dementia; VD = vascular dementia; MD = mixed dementia; MCI = mild cognitive impairment. BID = 2 times per day; TID = 3 times per day. *Studies used a placebo control unless otherwise indicated. † Patients already on donepezil.
Table 2: Types of ITT and non-ITT analyses employed and number of contraindications to LOCF analysis

| Study                        | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|------------------------------|------------------------------------------|--------------------------------------|---------------------------------------------------|----------------------------------------------------------------------------------|---------------------|--------------------------|----------------------------------------------------------------------------------|
| DONEPEZIL IN ALZHEIMER’S DEMENTIA                                      |                                          |                                      |                                                   |                                                                                 |                     |                          |                                                                                  |
| Rogers et al. (1996)²⁶       | No                                       | Unknown                              | Unknown                                          | 0-2                                               | LOCF described but term not used                               | None                 |                                                                                     |
| Rogers et al. (1998)¹²       | Yes                                      | Placebo 7%                           | Unknown                                          | 1-3                                               | LOCF                                                            | FEP                  | +                                                                                   |
| Rogers et al. (1998)²⁷       | Yes                                      | Placebo 20%                          | Unknown                                          | 1-3                                               | LOCF                                                            | FEP                  | +                                                                                   |
| Burns et al. (1999)¹⁸        | Yes                                      | Placebo 20%                          | Unknown                                          | 1-3                                               | LOCF                                                            | FEP, OCA, retrieved dropout | +                                                                                   |
| Greenberg et al. (2000)²⁹    | Yes                                      | Placebo 5%                           | Unknown                                          | 1-3                                               | Not specified                                                   | Not specified        | +                                                                                   |
| Homma et al. (2000)³⁰        | No                                       | Unknown                              | Unknown                                          | 0-2                                               | Not specified                                                   | TPP                  | +                                                                                   |
| Feldman et al. (2001)³¹      | No                                       | No                                   | Unknown                                          | 0-1                                               | LOCF                                                            | OCA                  |                                                                                     |
| Mohs et al. (2001)³²         | No                                       | Unknown                              | Unknown                                          | 0-2                                               | LOCF                                                            | OCA                  |                                                                                     |
| Tariot et al. (2001)³³       | No                                       | Unknown                              | Unknown                                          | 0-2                                               | LOCF                                                            | OCA                  |                                                                                     |
| Thomas et al. (2001)³⁴       | No                                       | No                                   | No                                               | 0                                                 | No ITT analysis performed                                        | CA                   | +                                                                                   |
| Winblad et al. (2001)³⁵      | No                                       | No                                   | Unknown                                          | 0-1                                               | LOCF                                                            | OCA                  |                                                                                     |
| Krishnan et al. (2003)³⁶     | No                                       | Unknown                              | Unknown                                          | 0-2                                               | LOCF                                                            | OCA                  |                                                                                     |
| Tune et al. (2003)³⁷         | No                                       | Unknown                              | Unknown                                          | 0-2                                               | Not specified                                                   | Not specified        | +                                                                                   |
| Study                                    | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-----------------------------------------|------------------------------------------|--------------------------------------|-------------------------------------------|--------------------------------------------------------------------------------------|----------------------|--------------------------|-------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------|
| AD2000 Collaborative Group (2004)        | No                                       | Unknown                              | Yes                                       | 1-2                                                                                  | Most recent previous score was used (similar to LOCF) if one existed; if not, next subsequent valid score was substituted | None                     |                                                                                       | +                                                                                                                                  |
| Holmes et al. (2004)                     | No                                       | Unknown                              | Yes                                       | 1-2                                                                                  | LOCF                 | OCA                      |                                                          |                                                                                                                                  |
| Hopkins et al. (2000)                    | No                                       | No                                   | No                                        | 0                                                                                     | Not specified        | Not specified            |                                                          |                                                                                                                                  |
| Seltzer et al. (2004)                    | Yes                                      | Unknown                              | Unknown                                   | 1-3                                                                                  | LOCF                 | FEP                      |                                                          |                                                                                                                                  |
| Johansson et al. (2006)                  | No                                       | Unknown                              | Unknown                                   | 0-2                                                                                  | LOCF                 | OCA                      |                                                          |                                                                                                                                  |
| Winblad et al. (2006)                    | Yes                                      | Unknown                              | Unknown                                   | 1-3                                                                                  | LOCF and modelling; missing data were replaced with mean of observed values for change from baseline to month 6 in placebo group (LOCF and modelling provided similar point estimates in SIB, ADCS-ADL-severe, CGI-I, MMSE and NPI) | CA                      |                                                          |                                                                                                                                  |
| Mazza et al. (2006)                      | No                                       | Unknown                              | Unknown                                   | 0-2                                                                                  | Not specified        | Not specified            |                                                          |                                                                                                                                  |
| Black et al. (2007)                      | Yes                                      | Unknown                              | Unknown                                   | 1-3                                                                                  | LOCF                 | OCA                      |                                                          |                                                                                                                                  |

**RIVASTIGMINE IN ALZHEIMER’S DEMENTIA**

| Study                                    | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-----------------------------------------|------------------------------------------|--------------------------------------|-------------------------------------------|--------------------------------------------------------------------------------------|----------------------|--------------------------|-------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------|
| Study                        | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-----------------------------|-----------------------------------------|--------------------------------------|--------------------------------------------------|-----------------------------------------------------------------|---------------------|--------------------------|------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------|
| Corey-Bloom et al. (1998)⁴⁸ | Yes Placebo 7% 1-4 mg 15% 6-12 mg 35% | Yes Unknown 2-3 | LOCF                                             | OCA                                                             | +                   |
| Forette et al. (1999)⁴⁹     | Yes Placebo 21% 6-9 mg via BID dosing 50% 6-9 mg via TID dosing 38% | Unknown Unknown 1-3 | No ITT analysis performed                       | FEP                                                             | +                   |
| Rösler et al. (1999)⁵⁰      | Yes Placebo 13% 1-4 mg 14% 6-12 mg 32% | Yes Unknown 2-3 | LOCF                                             | OCA                                                             | +                   |
| Kumar et al. (2000)⁷⁷       | Yes Placebo 16% 1-4 mg 14% 6-12 mg 33% | Unknown Unknown 1-3 | No ITT analysis performed                       | OCA                                                             | +                   |
| Dora’swamy et al. (2002)⁵⁸  | Not ruled out                           | Unknown Unknown 0-3 | LOCF                                             | OCA                                                             |                     |
| Karaman et al. (2005)⁹⁵     | Yes Placebo 0% 6-12 mg 13%               | No No 1 | No ITT analysis performed                       | OCA                                                             | +                   |
| Bullock et al. (2005)¹⁰⁵     | Yes Donepezil 5-10 mg 36% Rivastigmine 3-12 mg 47% | Yes Unknown 2-3 | LOCF                                             | OCA, FEP                                                       | +                   |
| Feldman et al. (2007)¹³⁶     | Yes Placebo 15% Rivastigmine BID 24% Rivastigmine TID 17% | Unknown Unknown 1-3 | LOCF                                             | OCA. Retrieved dropout + LOCF                                   | +                   |
| Mowla et al. (2007)⁷³       | No Unknown Unknown 0-2                   | Not specified Not specified                   | Not specified                                                    | +                   |
| Study                          | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-------------------------------|------------------------------------------|--------------------------------------|--------------------------------------------------------------------------------------|----------------------|---------------------------|------------------------------------------------------------------------|--------------------------------------------------------------------------------|
| Winblad et al. (2007)        | Yes Placebo 12% 10 cm² 22% 20 cm² 21% 12 mg 22% | Unknown Unknown                      | 1-3                                                                                  | LOCF                 | OCA Retrieved dropout    |                                                                         | +                                                                               |
| Raskind et al. (2000)        | Yes Placebo 9% 24 mg 32% 32 mg 42%        | Unknown Unknown                      | 1-3                                                                                  | LOCF; time response relationship for change in ADAScog/11 analyzed using generalized linear interactive modelling (results of modelling not provided) | OCA                     |                                                                         |                                                                                   |
| Tariot et al. (2000)         | Yes Placebo 16% 8 mg 23% 6mg 22% 24 mg 22% | Unknown Unknown                      | 1-3                                                                                  | LOCF                 | OCA                       |                                                                         | +                                                                               |
| Wilcock et al. (2000)        | Yes Placebo 13% 24mg 20% 32mg 25%         | Unknown Unknown                      | 1-3                                                                                  | The term LOCF was not employed but the technique was described in the paper; the time response relationship for change in ADAScog/11 was analyzed using generalized linear mixed modelling (results of modelling not provided) | OCA                     |                                                                         |                                                                                   |
| Rockwood et al. (2001)       | Yes Placebo 11% 24 or 32mg 33%            | Unknown Unknown                      | 1-3                                                                                  | LOCF                 | OCA                       |                                                                         | +                                                                               |
### No. of contraindications to the use of LOCF analysis

| Study | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-------|-----------------------------------------|--------------------------------------|------------------------------------------|---------------------------------------------|----------------------|------------------------|-----------------------------------------------|----------------------------------------------------------------------------------|
| Wilkinson et al. (2001) | Yes | Placebo 16% 18 mg 28% 24 mg 28% 36 mg 48% | Unknown | Unknown | 1-3 | LOCF | TPP | + |
| Brodaty et al. (2005) | Yes | Placebo 23% Galantamine (immediate release) 31% Galantamine (prolonged release) 25% | Unknown | Unknown | 1-3 | LOCF | OCA | + |
| Rockwood et al. (2006) | No | Unknown | Unknown | 0-2 | LOCF, mixed-effects modelling (point estimates of outcomes based on modelling not provided) | OCA | | |

### MEmantine in Alzheimer’s Dementia

| Study | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-------|-----------------------------------------|--------------------------------------|------------------------------------------|---------------------------------------------|----------------------|------------------------|-----------------------------------------------|----------------------------------------------------------------------------------|
| Winblad et al. (1999) | No | Unknown | Unknown | 0-2 | Missing end-point data were replaced by worst ranks | TPP | | |
| Reisberg et al. (2003) | No | Unknown | Unknown | 0-2 | LOCF; missing values replaced with mean observed value for decline in placebo group (point estimates of outcomes based on modelling not given) | OCA | | |
| Tariot et al. (2004) | No | Unknown | Unknown | 0-2 | LOCF | OCA | | |
| Peskind and Loft (2006) | No | Unknown | Unknown | 0-2 | LOCF, MMRM (point estimates of outcomes based on modelling not given) | OCA | | |
| Study                           | Greater dropout rate in treatment group?* | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout group? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|--------------------------------|------------------------------------------|--------------------------------------|-----------------------|-----------------------------------------------|----------------------|--------------------------|--------------------------------------------------------------------------------|--------------------------------------------------------------------------------|
| Bakchiche et al. (2007)†³      | Yes                                      | Unknown                              | Unknown               | 1-3                                           | LOCF mentioned but results were not provided | CA, OCA                  |                                                                                   |                                                                                   |
| Van Dyck (2007)²⁸              | No                                       | Unknown                              | Unknown               | 0-2                                           | LOCF, MMRM (point estimates of outcomes based on modelling not given) | OCA                     |                                                                                   |                                                                                   |
| **DONEPEZIL IN VASCULAR DEMENTIA AND MIXED DEMENTIA** |                                                                                   |                                      |                                      |                                                                                   |                                                                                   |                                                                                   |                                                                                   |
| Pratt et al. (2002)²⁰          | Yes                                      | Unknown                              | Unknown               | 1-3                                           | LOCF                               | OCA                      | +                                                                                 |                                                                                   |
| Black et al. (2003)³¹          | Yes                                      | Unknown                              | Unknown               | 1-3                                           | LOCF                               | OCA                      | +                                                                                 |                                                                                   |
| Wilkinson et al. (2003)³⁵      | Yes                                      | Unknown                              | Unknown               | 2-3                                           | LOCF                               | OCA                      | +                                                                                 |                                                                                   |
| **GALANTAMINE IN VASCULAR DEMENTIA AND MIXED DEMENTIA** |                                                                                   |                                      |                                      |                                                                                   |                                                                                   |                                                                                   |                                                                                   |
| Erkinjuntti et al. (2002)³⁵    | Yes                                      | Unknown                              | Unknown               | 2-3                                           | LOCF, mixed-effects modelling (results of modelling not provided)             | OCA                      | +                                                                                 |                                                                                   |
| Bullock et al. (2004)³⁴        | Yes                                      | Unknown                              | Unknown               | 1-3                                           | Used term “observed case analysis” but described LOCF | None                     | +                                                                                 |                                                                                   |
| Auchus et al. (2007)³⁶         | Yes                                      | Unknown                              | Unknown               | 1-3                                           | LOCF                               | OCA                      | +                                                                                 |                                                                                   |
| Study                              | Greater dropout rate in treatment group? | Earlier dropouts in treatment group? | More rapid progressors in treatment dropout? | Total no. of contraindications (demonstrated to potential maximum) to LOCF analysis† | Type of ITT analysis | Type of non-ITT analyses | Only non-ITT analysis performed or type of ITT analysis not specified | LOCF was only ITT analysis and study demonstrated factors that can introduce bias in favour of study drug in LOCF analysis |
|-----------------------------------|----------------------------------------|--------------------------------------|--------------------------------------------|---------------------------------------------------------------------------------|---------------------|--------------------------|--------------------------------------------------------------|------------------------------------------------------------------------------------------------------------------|
| **MEMANTINE IN VASCULAR DEMENTIA AND MIXED DEMENTIA** |                                        |                                      |                                            |                                                                                |                     |                          |                                                              |                                                                                                                  |
| Orgogozo et al. (2002)§§         | Yes                                    | Placebo 16% 20 mg 21%                | Unknown                                    | 1-3                                                                             | LOCF                | OCA, TPP                 | +                                                            |                                                                                                                  |
| Wilcock et al. (2002)§§          | No                                     | Unknown                              | Unknown                                    | 0-2                                                                             | LOCF                | TPP                      |                                                              |                                                                                                                  |
| **DONEPEZIL IN MILD COGNITIVE IMPAIRMENT** |                                        |                                      |                                            |                                                                                |                     |                          |                                                              |                                                                                                                  |
| Salloway et al. (2004)§§         | Yes                                    | Placebo 17% 10 mg 32%                | Unknown                                    | 1-3                                                                             | LOCF                | OCA, FEP                 | +                                                            |                                                                                                                  |
| Petersen et al. (2005)§§         | Yes                                    | Placebo 25% 10 mg 36%                | Yes                                        | 3                                                                               | Employed a sensitivity analysis consisting of a number of simulations (modeling) | For secondary outcomes “missing values were imputed with the use of a projection method appropriate for assessing responses among subjects with neurodegenerative diseases.” |                                                              |                                                                                                                  |
| **GALANTAMINE IN MILD COGNITIVE IMPAIRMENT** |                                        |                                      |                                            |                                                                                |                     |                          |                                                              |                                                                                                                  |
| Koontz and Baskys (2005)§§       | Yes                                    | Placebo 36% 24 mg 50%                | Unknown                                    | 1-3                                                                             | Not specified       | Description suggests OCA | +                                                            |                                                                                                                  |
| **RIVASTIGMINE IN MILD COGNITIVE IMPAIRMENT** |                                        |                                      |                                            |                                                                                |                     |                          |                                                              |                                                                                                                  |
| Feldman et al. (2007)§§          | No                                     | Unknown                              | Unknown                                    | 0-3                                                                             | LOCF described, but the term was not used | OCA                      |                                                              |                                                                                                                  |

ITT= intention to treat; LOCF = last observation carried forward; FEP = fully evaluable population; OCA = observed-case analysis; TPP = treatment per protocol; SIB = severe impairment battery; ADCS-ADL-severe = the Modified Alzheimer’s Disease Cooperative Study activities of daily living inventory for severe Alzheimer’s disease; CGI-I = clinical global impression of improvement; MMSE = mini-mental state examination; NPI = neuropsychiatric inventory; CA = completer analysis; BID = 2 times per day; TID = 3 times per day; MWM = mixed-models repeated measures.

§ If there was a greater dropout rate in the treatment group, the dropout rates are provided.

† Total number of contraindications to the use of LOCF: the lower number represents the number of contraindications explicitly demonstrated whereas the higher number represents the potential maximum number of contraindications (the difference between the higher and lower numbers represents potential contraindications that could not be ruled out owing to lack of information on dropouts in the publication reviewed).

§§ In this study the use of donepezil in both Alzheimer’s dementia and mixed dementia was investigated.

$ This study compared treatment with rivastigmine and donepezil.

¶ In this study the use of memantine in both Alzheimer’s dementia and vascular dementia was investigated.
results. In 3 of these 8 studies the authors did not comment on the results of the alternative non-LOCF ITT analysis. In 4 of the 5 studies in which the authors commented on the results of the alternative ITT analysis, they did not report the values calculated by this analysis but they did indicate that the direction of the results was unchanged. It is uncertain whether the point estimates of the outcomes were similar when the alternative ITT analyses were performed.

In only 1 study were the point estimates of outcomes measuring drug efficacy generated by LOCF verified with point estimates generated by an alternative form of ITT analysis.32 The values of 3 positive outcomes were verified in this study.

**Contraindications to the use of LOCF as the only form of ITT analysis.** Of the 34 studies employing LOCF as the only form of ITT analysis, 24 (71%) explicitly demonstrated contraindications (factors that could introduce bias) to its use. It was unclear whether the remaining 10 studies were free of contraindications, because most studies failed to report adequate data regarding the timing of dropouts and the severity of disease of the participants who dropped out. Consequently, Table 2 provides a range of potential contraindications to the use of LOCF for each study (the lower number representing the number of explicitly identified contraindications). Seven of the 57 trials in this review (12%) discussed the limitations of LOCF or non-ITT approaches.

### Discussion

Despite previously published cautions that LOCF analysis may introduce bias into dementia research, LOCF remains the most widely employed analytic technique in this research area; its results are rarely verified by other forms of ITT analysis. Further, the majority of the publications reviewed in the present study did not report the results of an ITT analysis, did not verify the results of LOCF with alternative ITT analyses when conditions that could introduce LOCF analytic bias in favour of the study drug existed, or did not comment on the results of alternative ITT analyses that were performed.

These problems were particularly evident in cholinesterase inhibitor trials. In the majority of these trials, either no ITT results were provided or LOCF ITT analysis was performed in the presence of contraindicating factors. For example, a higher dropout rate in the treatment group than in the control group was more common in cholinesterase inhibitor studies than in memantine studies (73% v. 25%), potentially biasing study results in favour of cholinesterase inhibitors and against memantine. Owing to a lack of data on the timing of dropouts and on the severity of disease in study participants who dropped out, our results may in fact underestimate the true prevalence of conditions promoting bias.

The concern that LOCF analysis introduces bias can be explored via ITT sensitivity analyses. If similar out-

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**Table 3: Account of dropouts by drug class**

| Description of dropouts                                      | Studies of cholinesterase inhibitors, no. (%) (n = 49) | Studies of memantine, no. (%) (n = 8) |
|--------------------------------------------------------------|-------------------------------------------------------|-------------------------------------|
| Studies describing total no. of study dropouts               | 46 (94)                                               | 8 (100)                             |
| Studies with a greater dropout rate in the experimental group| 36 (73)                                               | 2 (25)                              |
| Studies with a greater dropout rate in the control group     | 8 (16)                                                | 2 (25)                              |
| Studies with similar dropout rates between groups            | 2 (4)                                                 | 4 (50)                              |

| Timing of dropouts                                          |                                                      |                                      |
|--------------------------------------------------------------|-------------------------------------------------------|-------------------------------------|
| Studies describing dropout timing                            | 7 (14)                                                | 0 (0)                               |

**Table 4: Combined data for cholinesterase inhibitor and memantine trials**

|                                | Studies of cholinesterase inhibitors (n = 49) | Studies of memantine (n = 8) |
|--------------------------------|----------------------------------------------|-------------------------------|
|                                | Control group | Experimental groups | Control group | Experimental groups |
| Total no. of study participants | 7275           | 11969              | 1349           | 1539                          |
| No. of study participants completing study                     | 6050           | 9198              | 1099           | 1315                          |
| Dropout rate (%)                                               | 16.8           | 23.2              | 18.5           | 14.6                          |
comes are generated when other forms of ITT analysis are employed, this provides some reassurance (but does not guarantee) that LOCF analytic bias does not alter results. Only 1 study verified the point estimates of efficacy calculated by LOCF analysis with an alternative ITT analysis. The 3 positive point estimates verified by alternative ITT analyses in this study are the only ones out of the hundreds of positive outcomes reported for LOCF analyses in dementia trials to have been verified in this way.

Some may erroneously argue that results of previous studies have been adequately confirmed by non-ITT analyses (i.e., techniques that exclude subjects without data from analysis), such as observed case analysis, completer analysis, fully evaluable population analysis or treated-per-protocol analysis. Like LOCF analysis, these non-ITT techniques may be systematically biased in favour of the group with greater, earlier or more severely affected dropouts and, consequently, they are not reliable, valid sensitivity analyses. The biases inherent in these non-ITT techniques have been highlighted by the International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use,\(^{42}\) the European Agency for the Evaluation of Medicinal Products\(^{14}\) and by a number of authors.\(^{15,53,67,81}\)

Furthermore, the use of such non-ITT techniques and of LOCF analysis is completely unnecessary: other forms of ITT analysis that do not treat dropouts artificially by freezing values at the point of dropout but rather model for expected natural decline in dropouts could easily be performed either as the primary analysis or as a sensitivity analysis. The available approaches range from techniques that simply apply the rate of decline noted in the control group to all dropouts to more complex modelling procedures that are available in standard statistical programs. More appropriate forms of analysis have been employed in dementia research.\(^{15,50,52,58,68,78,79}\)

Petersen’s study of mild cognitive impairment\(^{68}\) may serve as a model for future research, as it employs both modelling for dropouts and sensitivity analyses of the effect of various modelling assumptions and approaches.

The present study cannot quantify the magnitude of the effect of the use of LOCF analysis on trial results; it is restricted to highlighting the high prevalence of conditions promoting bias in favour of more toxic therapies and against less toxic alternatives, such as memantine. As verification of results obtained using non-ITT and LOCF analyses requires individual patient data that are not publicly available, the onus is on the investigators who publish these trials to disprove the possibility that these analyses have introduced bias by performing ITT sensitivity analyses as performed in Petersen’s study of mild cognitive impairment.\(^{68}\) This is particularly true for those studies demonstrating higher dropout rates in treatment groups.

These results are meaningful in day-to-day clinical care. Because this bias has likely exaggerated results in favour of more toxic therapies (e.g., cholinesterase inhibitors), this may have created inappropriate barriers to the funding and prescription of less toxic treatment options for dementia (e.g., memantine). Without accurate analyses, physicians cannot optimally counsel patients and families regarding appropriate therapies, and patients and families cannot provide truly informed consent when making treatment decisions. In addition, meta-analyses and pharmaco-economic studies cannot be performed accurately and we cannot make reliable statements regarding whether trial results truly cross thresholds of clinical significance. These concerns, as well as the fact that LOCF analytic bias may prevent the funding and use of future less toxic treatments, should be of great concern to patient advocacy groups, such as the Alzheimer Society of Canada and the US Alzheimer’s Association.

In summary, it is highly unlikely, given the high prevalence of conditions promoting LOCF analytic bias in this study, that point estimates of some of the hundreds of positive outcomes generated in trials have not been affected in some way. The question is likely not whether bias has been introduced, but rather the number of outcomes that have been biased and the degree to which they have been biased. As such, the present results provide empirical support for recent recommendations to researchers, licensing bodies and research guidelines bodies\(^8\) regarding their use of LOCF analysis. One of these recommendations is that the CONSORT group (www.consort-statement.org) consider incorporating guidelines regarding appropriate analyses for studies of medications used to treat chronic progressive disorders into the CONSORT Statement so that journal editors, funding agencies, ethics review boards and drug formulary committees can request that these recommendations be followed in future studies of dementia and other chronic progressive disorders. In the meantime, researchers should ensure that analyses promoting bias are avoided or scrutinized using alternative ITT sensitivity analyses. Further, licensing agencies (e.g., the US Food and Drug Administration, the European Agency for the Evaluation of Medicinal Products, and Health Canada) should review this situation immediately to determine whether they will continue to accept LOCF analyses in research on dementia and other chronic progressive conditions.
34. Thomas A, Iacono D, Bonanni L, D’Andreamattee G, Onofri M. Donepezil, rivastigmine, and vitamin E in Alzheimer disease: A combined P200 event-related potentials/neuropsychologic evaluation over 6 months. Clin Neuropsychol. 2001;24(1):31–42.

35. Winblad B, Engedal K, Soininen H, Verhey F, Waldemar G, Wimo A, et al. A 1-year, randomized, placebo-controlled study of donepezil in patients with mild to moderate AD. Neurology. 2001;57(3):489–495.

36. Krishnan KRR, Charles HC, Doraiswamy PM, Mintzer J, Weisler R, Yu X, et al. Randomized, placebo-controlled trial of the effects of donepezil on neuronal markers and hippocampal volumes in Alzheimer’s disease. Am J Psychiatry. 2003;160(11):2003–2011.

37. Tune I, Tiseo PJ, Ieni J, Perdomo C, Pratt RD, Votaw JR, et al. Donepezil HCI (E2020) maintains functional brain activity in patients with Alzheimer disease: Results of a 24-week, double-blind, placebo-controlled study. Am J Geriatr Psychiatry. 2003;11(2):169–177.

38. Holmes C, Wilkinson D, Dean C, Vethanayagam S, Oliveri S, Langley A, et al. The efficacy of donepezil in the treatment of neuropsychiatric symptoms in Alzheimer disease. Neurology. 2004;63(2):214–9.

39. Moraes WD, Poyares DR, Guillemínault C, Ramos LR, Bertolucci PHF, Tufik S. The effect of donepezil on sleep and REM sleep EEG in patients with Alzheimer disease: a double-blind placebo-controlled study. Sleep. 2006;29(2):199–205.

40. Selzer B, Zolnoumi P, Nunez M, Goldman R, Kumar D, Ieni J, et al. Efficacy of donepezil in early-stage Alzheimer disease: A randomized placebo-controlled trial. Arch Neurol. 2004;61(12):1852–1856.

41. Johanssen P, Salmon E, Hampel H, Xu Y, Richardson S, Qvitzau S, et al. Assessing therapeutic efficacy in a progressive disease: A study of donepezil in Alzheimer’s disease. CNS Drugs. 2006;20(4):311–325.

42. Winblad B, Kilander L, Eriksson S, Minthon L, Båtsman S, Westerholm A, et al. Donepezil in patients with severe Alzheimer’s disease: Double-blind, parallel-group, placebo-controlled study. Lancet. 2006;367(9516):1057–1065.

43. Agid Y, Dubois B, Anand R, Gharabawi G; International Rivastigmine Investigators. Efficacy and tolerability of rivastigmine in patients with dementia of the Alzheimer type. Curr Ther Res. 1998;59(12):837–845.

44. Corey-Bloom J, Anand R, Veach J; ENA 713 B352 Study. A randomized trial evaluating the efficacy and safety of ENA 713 (rivastigmine tartrate), a new acetylcholinesterase inhibitor, in patients with mild to moderately severe Alzheimer’s disease. Int J Geriatr Psychopharmacol. 1998;1:55–65.

45. Forette F, Anand R, Gharabawi G. A phase II study in patients with Alzheimer’s disease to assess the preliminary efficacy and maximum tolerated dose of rivastigmine (Exelon). Eur J Neurol. 1999;6(4):423–429.

46. Rösmler M, Anand R, Cicin-Sain A, Gauthier S, Agid Y, Dal-Bianco P, et al. Efficacy and safety of rivastigmine in patients with Alzheimer’s disease: International randomised controlled trial. BMJ. 1999;318(7184):693–698.

47. Kumar V, Anand R, Messina J, Hartman R, Veach J. An efficacy and safety analysis of Exelon in Alzheimer’s disease patients with concurrent vascular risk factors. Eur J Neurol. 2000;7(2):159–169.

48. Doraiswamy PM, Krishnan KRR, Anand R, Sohn H, Danyuk J, Hartman RD, et al. Long-term effects of rivastigmine in moderately severe Alzheimer’s disease: does early initiation of therapy offer sustained benefits. Prog Neuropsychopharmacol Biol Psychiatry. 2002;26(4):705–712.

49. Karaman Y, Erdoğan F, Köseöglu E, Turan T, Ersoy ÖA. A 12-month study of the efficacy of rivastigmine in patients with advanced moderate Alzheimer’s disease. Dement Geriatr Cogn Disord. 2005;19(1):51–56.

50. Raskind MA, Peskind ER, Wessel T, Yuan W; Galantamine USA-1 Study Group. Galantamine in AD: A 6-month randomized, placebo-controlled trial with a 6-month extension. Neurology. 2000;54(12):2261–2268.

51. Tariot PN, Solomon PR, Morris JC, Kershaw P, Lilienfeld S, Ding C; The Galantamine USA-10 Study Group. A 5-month, randomized, placebo-controlled trial of galantamine in AD. Neurology. 2000;54(12):2269–2276.

52. Wilcock GK, Lilienfeld S, Gaens E; Galantamine International-1 Study Group. Efficacy and safety of galantamine in patients with mild to moderate Alzheimer’s disease: Multicentre randomised controlled trial. BMJ. 2000;321(7274):1445–1449.

53. Rockwood K, Mintzer J, Truyen L, Wessel T, Wilkinson D. Effects of a flexible galantamine dose in Alzheimer’s disease: A randomized, controlled trial. J Neurol Neurosurg Psychiatry. 2001;71(5):589–595.

54. Wilkinson D, Murray J; The Galantamine Research Group. Galantamine: A randomized, double-blind, dose comparison in patients with Alzheimer’s disease. Int J Geriatr Psychiatry. 2001;16(9):852–857.

55. Brodaty H, Corey-Bloom J, Potocnik FCV, Truyen L, Gold M, Damaraju CRV. Galantamine prolonged-release formulation in the treatment of mild to moderate Alzheimer’s disease. Dement Geriatr Cogn Disord. 2005;20(2-3):120–132.

56. Rockwood K, Fay S, Song X, MacKnight C, Gorman M; Video-Imaging Synthesis of Treating Alzheimer’s Disease (VISTA) Investigators. Attainment of treatment goals by people with Alzheimer’s disease receiving galantamine: A randomized controlled trial. CMAJ. 2006;174(8):1099–1105.

57. Winblad B, Portis N. Memantine in severe dementia: Results of the 9M-Best Study (Benefit and efficacy in severely demented patients during treatment with memantine. Int J Geriatr Psychiatry. 1999;14(2):135–146.

58. Reisberg B, Doody R, Stöffler A, Schmitt F, Ferris S, Möbius HJ; Memantine Study Group. Memantine in moderate-to-severe Alzheimer’s disease. N Engl J Med. 2003;348(14):1333–1341.

59. Tariot PN, Farlow MR, Grossberg GT, Graham SM, McDonald S, Gergel I; Memantine Study Group. Memantine treatment in patients with moderate to severe Alzheimer disease already receiving donepezil: a randomized controlled trial. JAMA. 2004;291(3):317–324.

60. Pratt RD, Perdomo CA. Donepezil-treated patients with probable vascular dementia demonstrate cognitive benefits. Ann N Y Acad Sci. 2002;977:513–522.

61. Black S, Román GC, Geldmacher DS, Salloway S, Hecker J, Burns A, et al. Efficacy and tolerability of donepezil in vascular dementia: Positive results of a 24-week, multicenter, international, randomized, placebo-controlled clinical trial. Stroke. 2003;34(16):2233–2230.

62. Wilkinson D, Doody R, Helme R, Taukman K, Mintzer J, Kertesz A, et al. Donepezil in vascular dementia: A randomised, placebo-controlled study. Neurology. 2003;61(4):479–486.

63. Erkinjuntti T, Kurz A, Gauthier S, Bullock R, Lilienfeld S, Damaraju CV. Efficacy of galantamine in probable vascular dementia
and Alzheimer’s disease combined with cerebrovascular disease: A randomised trial. *Lancet.* 2002;359(9314):1283–1290.

64. Bullock R, Erkinjuntti T, Liliienfeld S; GAL-INT-6 Study Group. Management of patients with Alzheimer’s disease plus cerebrovascular disease: 12-Month treatment with galantamine. *Dement Geriatr Cogn Disord.* 2004;17(1-2):29–34.

65. Orgogozo J, Rigaud A, Stößler A, Möbius H, Forette F. Efficacy and safety of memantine in patients with mild to moderate vascular dementia: A randomized, placebo-controlled trial (MMM 300). *Stroke.* 2002;33(7):1834–1839.

66. Wilcock G, Möbius HJ, Stößler A; MMM 500 group. A double-blind, placebo-controlled multicentre study of memantine in mild to moderate vascular dementia (MMM500). *Int Clin Psychopharmacol.* 2002;17(6):297–305.

67. Salloway S, Ferris S, Kluger A, Goldman R, Griesing T, Kumar D, et al. Efficacy of donepezil in mild cognitive impairment: A randomized placebo-controlled trial. *Neurology.* 2004;63(4):651–657.

68. Petersen RC, Thomas RG, Grundman M, Bennett D, Doody R, Ferris S, et al. Vitamin E and donepezil for the treatment of mild cognitive impairment. *N Engl J Med.* 2005;352(23):2379–2388.

69. Koontz J, Basks J. Effects of galantamine on working memory and global functioning in patients with mild cognitive impairment: A double-blind placebo-controlled study. *Am J Alzheimers Dis Other Demen.* 2005;20(5):395–302.

70. Mazza M, Capuano A, Bria P, Mazza S. Ginkgo biloba and donepezil: A comparison in the treatment of Alzheimer’s dementia in a randomized placebo-controlled double-blind study. *Eur J Neurol.* 2006;13(9):981–985.

71. Black SE, Doody R, Li H, McRae T, Jambor KM, Xu Y, et al. Donepezil preserves cognition and global function in patients with severe Alzheimer disease. *Neurology.* 2007;69(3):459–469.

72. Feldman HH, Lane R; Study 304 Group. Rivastigmine: A placebo controlled trial of twice daily and three times daily regimens in patients with Alzheimer’s disease. *J Neurol Neurosurg Psychiatry.* 2007;78(10):1056–1063.

73. Mowla A, Mosavinasab M, Haghsenas H, Haghighi AB. Does serotonin augmentation have any effect on cognition and activities of daily living in Alzheimer’s dementia? A double-blind, placebo-controlled clinical trial. *J Clin Psychopharmacol.* 2007;27(5):484–487.

74. Winblad B, Cummings J, Andreasen N, Grossberg G, Onofri M, Sadowsky C, et al. A six-month double-blind, randomized, placebo-controlled study of a transdermal patch in Alzheimer’s disease – rivastigmine patch versus capsule. *Int J Geriatr Psychiatry.* 2007;22(5):456–467.

75. Bakchine S, Loft H. Memantine treatment in patients with mild to moderate Alzheimer’s disease: Results of a randomised, double-blind, placebo-controlled 6-month study. *J Alzheimer Dis.* 2007;11(4):471–479.

76. Auchus AP, Brashear HR, Salloway S, Korczyn AD, De Deyn PP, Gassmann-Mayer C; GAL-INT-26 Study Group. Galantamine treatment of vascular dementia: A randomized trial. *Neurology.* 2007;69(5):448–458.

77. Feldman HH, Ferris S, Winblad B, Sfikas N, Mancione L, He Y, et al. Effect of rivastigmine on delay to diagnosis of Alzheimer’s disease from mild cognitive impairment: The InDDEx study. *Lancet Neurol.* 2007;6(6):501–512.

78. van Dyck CH, Tariot PN, Meyers B, Malca RE; Memantine MEM-MD-01 Study Group. A 24-week randomized, controlled trial of memantine in patients with moderate-to-severe Alzheimer disease. *Alzheimer Dis Assoc Disord.* 2007;21(2):136–143.

79. Peskind ER, Potkin SG, Pomara N, Ott BR, Graham SM, Olin JT, et al. Memantine treatment in mild to moderate Alzheimer disease: A 24-week randomized, controlled trial. *Am J Geriatr Psychiatry.* 2006;14(8):704–715.

80. Structure and content of clinical study reports, E3: International conference on harmonisation of technical requirements for registration of pharmaceuticals for human use. Geneva: The Conference: 1995.

81. Ritchie CW, Ames D, Clayton T, Lai R. Metaanalysis of randomized trials of the efficacy and safety of donepezil, galantamine, and rivastigmine for the treatment of Alzheimer disease. *Am J Geriatr Psychiatry.* 2004;12(4):358–369.

82. Molnar FJ, Hutton B, Fergusson D. Does analysis using “last observation carried forward” introduce bias in dementia research. *CMAJ.* 2008;179(8):751–753.

**Citation:** Molnar FJ, Man-Son-Hing M, Hutton B, Fergusson DA. Have last-observation-carried-forward analyses caused us to favour more toxic dementia therapies over less toxic alternatives? A systematic review *Open Med* 2009;3(2):31-50

**Published:** 12 May 2009

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