Limitations of real-world treatment with atorvastatin monotherapy for lowering LDL-C in high-risk cardiovascular patients in the US

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Background: Guidelines endorse statin therapy for lowering low-density lipoprotein cholesterol (LDL-C) to recommended levels, in patients with cardiovascular disease (CVD) risk, if needed, after lifestyle changes. Atorvastatin is a common statin with greater LDL-C lowering efficacy than most other statins; its availability in generic form will likely increase its use. This study assessed attainment of guideline-recommended LDL-C levels in high-risk CVD patients treated with atorvastatin monotherapy.

Methods: Analyses of two retrospective US cohorts of patients who received a prescription for atorvastatin monotherapy between January 1, 2008 and December 31, 2010 (index date defined as first prescription date) in the GE Centricity Electronic Medical Record (EMR) (N = 10,693) and Humana Medicare (N = 16,798) databases. Eligible patients were ≥18 years, diagnosed with coronary heart disease or atherosclerotic vascular disease, with ≥1 LDL-C measurement between 3 months and 1 year postindex date, and continuously enrolled for 1 year prior to and following the index date.

Results: Of the eligible patients, 21.8%, 29.6%, 29.9%, and 18.7% (GE Centricity EMR) and 25.4%, 32.9%, 27.8%, and 14.0% (Humana Medicare) received 10, 20, 40, and 80 mg doses of atorvastatin, respectively. The mean ± standard deviation (SD) follow-up LDL-C levels were 2.1 ± 0.8 mmol/L (83 ± 30 mg/dL) and 2.3 ± 0.8 mmol/L (88 ± 31 mg/dL) for the GE Centricity EMR and Humana Medicare cohorts, respectively. Regardless of dose, only 28.3%–34.8% of patients had LDL-C levels ≤1.8 mmol/L (<70 mg/dL), and 72.0%–78.0% achieved LDL-C ≤2.6 mmol/L (<100 mg/dL) in both cohorts. As many as 41% and 13% of patients had LDL-C levels >0.5 mmol/L (>20 mg/dL) above LDL-C ≤1.8 mmol/L (70 mg/dL) and 2.6 mmol/L (100 mg/dL), respectively, in both cohorts; these percentages were generally similar across atorvastatin doses.

Conclusion: In this real-world US setting, a large number of high-risk CVD patients did not attain guideline-recommended LDL-C levels with atorvastatin monotherapy. More than 65% of the patients had LDL-C levels >1.8 mmol/L (>70 mg/dL), and of these, 30%–40% had LDL-C levels ≤0.5 mmol/L (≥20 mg/dL) above this, regardless of dose. This suggests that more effective lipid-lowering strategies, such as statin uptitration, switching to a higher efficacy statin, and/or combination therapy, may be required to achieve optimal LDL-C lowering in high-risk patients.

Keywords: statin therapy, managed-care, lipid-lowering therapy

Introduction

Lowering of low-density lipoprotein cholesterol (LDL-C) levels by statin therapy has been shown to be highly related to cardiovascular disease (CVD) risk reduction...
in a meta-analysis of 26 randomized-controlled statin clinical studies. As such, statin therapy is endorsed by various guidelines as the first-line pharmacotherapeutic approach for reducing LDL-C levels and CVD risk when lifestyle changes, including diet (eg, reduced saturated fats and cholesterol, plant stanols/sterols, and increased fiber), weight control, and exercise do not adequately control LDL-C levels.

Earlier guidelines targeted LDL-C <2.6 mmol/L (<100 mg/dL) as the optimal goal for primary prevention, in patients at high CVD risk with multiple coronary heart disease (CHD) risk factors. An optional treatment target of LDL-C <1.8 mmol/L (<70 mg/dL) was further recommended in 2004, based on clinical trial evidence showing that LDL-C 2.6 mmol/L (100 mg/dL) did not appear to be a threshold below which no further CV risk reduction could be achieved. The latter target is also considered reasonable for high-risk patients with atherosclerotic vascular disease, and similar guidelines have also been adopted internationally. Although LDL-C <1.8 mmol/L (<70 mg/dL) is recommended as a therapeutic option for all high CVD-risk patients, those in the category of very high risk (ie, patients with pre-existing CHD, plus major multiple risk factors [eg, diabetes], severe and poorly controlled risk factors, multiple risk factors of the metabolic syndrome, or acute coronary syndrome) should be most highly considered for this target. It should also be noted that recent evidence-based guidelines have recommended treatment of moderate-risk patients with moderate-intensity statins to reduce LDL-C levels from baseline by 30%–50% and of high-risk patients with high-intensity statins to reduce LDL-C levels by ≥50%, rather than to reach specific LDL-C targets.

While lowering of LDL-C to guideline-recommended levels has improved over time due to an increased awareness and use of statin therapy in clinical practice, many higher-risk patients still do not attain recommended LDL-C levels when prescribed statin monotherapy. Although ~2/3 of high-risk patients attain LDL-C levels <2.6 mmol/L (<100 mg/dL), only ~1/3 achieve levels <1.8 mmol/L (<70 mg/dL). Several observational studies and national surveys have indicated that various barriers, including sub-optimal lipid-lowering therapy, noncompliance, tolerability, and cost factors, have contributed to this treatment gap. Cholesterol guidelines have indicated that more effective lipid-lowering therapy, including up titration to higher statin doses, switching to higher potency statins, and/or combination statin therapy with other lipid-lowering agents, may be necessary to further improve LDL-C levels in these patients.

Atorvastatin is a common statin used in clinical practice that has been shown to effectively reduce LDL-C and CVD risk, in randomized, controlled clinical trials. Even though atorvastatin is more potent in terms of LDL-C lowering efficacy than most other statins, it has been less widely used in comparison with generic statins, particularly simvastatin, likely attributable to cost and other factors. However, the availability of generic atorvastatin will likely increase its use; thus it is of interest to examine LDL-C-lowering in high-risk patients treated with this statin in a real-world setting. This analysis assessed the attainment of LDL-C levels of <1.8 mmol/L (<70 mg/dL) and <2.6 mmol/L (<100 mg/dL) in two retrospective cohort studies of high-risk CVD patients treated with atorvastatin in the US, as well as the distance from these specified cut points in those patients who did not reach these levels.

**Study methods**

**Data sources**

Patient records from the GE Centricity Electronic Medical Record (EMR) database (GE Healthcare, Little Chalfont, UK) and administrative claims from Humana Medicare (Humana Inc., Louisville, KY, USA) were assessed between the dates of January 1, 2007 and December 31, 2011. The GE Centricity EMR database captures patient-level clinical data, including drug prescription and laboratory results, from over 25,000 physicians (~66% primary care), representing ~30 million US patients. We also analyzed administrative claims data from the Humana Medicare database. Humana Medicare claims data represent 8.1 million covered lives in 50 states, Washington DC, and Puerto Rico, among whom 4.3 million are Medicare members and 1.9 million are Advantage prescription drug plan beneficiaries. For the purpose of the present analysis, pharmacy data were merged with medical, lab, and patient data for Humana Medicare beneficiaries, using unique patient IDs. In both analyses, data were de-identified, and patient privacy was protected.

**Study design and patient eligibility**

Included in the studies were patients at high CVD risk who received a prescription for atorvastatin monotherapy (the date of the first prescription was defined as the index date) between January 1, 2008 and December 31, 2010 (index period). Patients were required to be ≥18 years of age at the index date and to have at least one recorded LDL-C measure ≥3 months and ≤12 months postindex date. For the Medicare analysis, there was an additional requirement for one atorvastatin prescription within 45 days of the LDL-C measure.
Limitations of atorvastatin monotherapy

Patients were required to be continuously enrolled in the respective system for 1 year prior to the index date (baseline period), and for 1 year following the index date (follow-up period) in the Humana Medicare analyses. Having at least one office visit in both the baseline and follow-up periods was used as a proxy for continuous enrollment over the study period, in the GE Centricity EMR analyses. High-risk patients were identified as those with evidence of CHD or atherosclerotic vascular disease (AVD), including ischemic heart disease, ischemic stroke, peripheral arterial disease, or atherosclerosis. The qualifying disease was identified using the International Classification of Diseases (ninth revision), Clinical Modification (ICD-9-CM) diagnosis/procedure or Current Procedural Terminology (CPT®) procedure codes during the baseline period and up until the follow-up LDL-C measure (Table S1). Excluded were patients who received any lipid-modifying treatment regimen, other than atorvastatin monotherapy, between the index date and the qualified LDL-C measure.

Outcomes of interest

Various guidelines recommend treatment of LDL-C to levels <2.6 mmol/L (<100 mg/dL) for all patients with CHD and/or AVD but also suggest treatment to LDL-C <1.8 mmol/L (<70 mg/dL) as reasonable or optional in such patients.2,4 The aim of this analysis was to assess mean LDL-C levels and the proportions of patients with levels <1.8 mmol/L (<70 mg/dL) and <2.6 mmol/L (<100 mg/dL), in high-risk patients with CHD and/or AVD who were treated with atorvastatin monotherapy. In this analysis, all patients were assessed against each specified LDL-C cut point, by atorvastatin dose and across doses. In addition, for patients above the specified LDL-C cut-points, we also assessed the proportions who were ≤0.3, 0.3–0.5, or >0.5 mmol/L (≤10, 11–20, or >20 mg/dL) away from the cut point overall, across atorvastatin doses and by daily atorvastatin dose.

Statistical analysis

The GE Centricity EMR and Humana Medicare analyses were purely descriptive, and there was no hypothesis testing or statistical comparison either within or between the two studies. They are presented side-by-side only to provide a broad view of atorvastatin monotherapy usage and associated LDL-C levels in two US populations.

Descriptive statistics were used to assess baseline patient characteristics, atorvastatin dose, and LDL-C levels. For continuous variables, means and standard deviations were calculated. Categorical variables were expressed as frequencies and proportions, with 95% confidence intervals calculated for the outcome measures of interest. Statistical analyses were performed using SAS versions 9.1 and 9.2 (SAS Institute, Inc., Cary, NC, USA) for the analysis of the GE Centricity EMR and the Humana Medicare databases, respectively.

Results

Patient characteristics

Of 438,403 patients on atorvastatin therapy in the GE Centricity EMR database during the index period, a total of 10,693 patients satisfied the inclusion criteria (Figure 2). Among these patients, the mean age was 68.0 years, and the majority were male (58.7%). The mean baseline LDL-C level was 2.1 mmol/L (82.2 mg/dL), with 34.1% of patients receiving some type of lipid-lowering medication in the 12 months prior to the index date (Table 1). CHD was the most common qualifying diagnosis (73.7%), followed by atherosclerosis (21.9%). In addition, 20.3% of patients had diabetes. The most commonly prescribed doses of atorvastatin were 20 (29.7%) and 40 (29.9%) mg; atorvastatin 80 mg was less-often prescribed (18.7%).

Of the 695,639 patients on atorvastatin in the Humana Medicare database during the index period, 16,798 patients met the inclusion criteria (Figure 2). Among these patients, the mean age was 73.7 years, and 52.1% were male. The mean LDL-C at baseline was 2.5 mmol/L (98.2 mg/dL), with 80.8% receiving some type of lipid-lowering therapy in the 12 months prior to the index date (Table 1). Similar to the findings for patients in the GE database, CHD was the most common qualifying diagnosis (69.6%), and atherosclerosis was identified in 23.7% of patients. Overall, 18.8% of this
high-risk population also had diabetes. The most commonly prescribed dose of atorvastatin was 20 mg (32.9%), and the least prescribed was 80 mg (14.0%).

**LDL-C levels at follow-up**

In each study, the mean LDL-C levels in the follow-up period were similar across atorvastatin doses. The mean LDL-C in the GE Centricity EMR analysis ranged from 2.2 mmol/L (83.5 mg/dL) to 2.1 mmol/L (82.4 mg/dL) for patients taking atorvastatin 10 mg and 80 mg, respectively (Table 2).

**Table 1** Patient characteristics

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>GE Centricity EMR</th>
<th>Humana Medicare</th>
</tr>
</thead>
<tbody>
<tr>
<td>N=10,693</td>
<td>N=16,798</td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>68.0±10.2</td>
<td>73.7±8.4</td>
</tr>
<tr>
<td>Male</td>
<td>6,275 (58.7)</td>
<td>8,750 (52.1)</td>
</tr>
<tr>
<td>Diabetes</td>
<td>2,171 (20.3)</td>
<td>3,159 (18.8)</td>
</tr>
<tr>
<td>Baseline LDL-C mmol/L ± SD</td>
<td>2.1±0.8 (82.2±29.7)</td>
<td>2.5±0.9 (98.2±35.3)</td>
</tr>
<tr>
<td>AWD or CHD diagnosis</td>
<td>7,879 (73.7)</td>
<td>11,690 (69.6)</td>
</tr>
<tr>
<td>Ischemic stroke</td>
<td>1,033 (9.7)</td>
<td>864 (5.1)</td>
</tr>
<tr>
<td>PAD</td>
<td>521 (4.9)</td>
<td>617 (3.7)</td>
</tr>
<tr>
<td>Atherosclerosis</td>
<td>2,338 (21.9)</td>
<td>3,976 (23.7)</td>
</tr>
<tr>
<td>Atorvastatin dose before LDL-C measure**</td>
<td>2.326 (21.8)</td>
<td>4,262 (25.3)</td>
</tr>
<tr>
<td>10 mg</td>
<td>2,326 (21.8)</td>
<td>4,262 (25.3)</td>
</tr>
<tr>
<td>20 mg</td>
<td>3,170 (29.7)</td>
<td>5,520 (32.9)</td>
</tr>
<tr>
<td>80 mg</td>
<td>3,198 (29.9)</td>
<td>4,672 (27.8)</td>
</tr>
<tr>
<td>10 mg</td>
<td>1,999 (18.7)</td>
<td>2,344 (14.0)</td>
</tr>
</tbody>
</table>

**Notes:** Values are expressed as mean ± SD or N (%). *The last LDL-C measure taken in 12 month baseline period closest to the index date; ‡Dose based on last prescribed before the follow-up LDL-C measure; §LDL-C level more than 0.5 mmol/L (20 mg/dL) above the 1.8 mmol/L (70 mg/dL) cut point. Of note, 32.4% of GE Centricity EMR patients and 40.7% of Humana Medicare patients had LDL-C levels more than 0.5 mmol/L (20 mg/dL) above the 1.8 mmol/L (70 mg/dL) cut point (Figure 3). These percentages were similar regardless of atorvastatin dose in the GE Centricity EMR analysis (33.2%, 33.0%, 31.5%, and 32.1% at 10, 20, 40, and 80 mg, respectively) and Humana Medicare databases.

For Humana Medicare patients, mean LDL-C levels ranged from 2.3 mmol/L (88.7 mg/dL) to 2.2 mmol/L (85.9 mg/dL) for 80 mg doses.

**Table 2** Follow-up LDL-C levels by atorvastatin dose

<table>
<thead>
<tr>
<th>Atorvastatin dose</th>
<th>GE Centricity EMR</th>
<th>Humana Medicare</th>
</tr>
</thead>
<tbody>
<tr>
<td>N=10,693</td>
<td>N=16,798</td>
<td></td>
</tr>
<tr>
<td>LDL-C mmol/L (mg/dL)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>10 mg</td>
<td>2.2±0.8 (83.5±29.6)</td>
<td>2.3±0.8 (88.7±29.4)</td>
</tr>
<tr>
<td>20 mg</td>
<td>2.1±0.8 (82.7±29.6)</td>
<td>2.3±0.8 (88.8±31.1)</td>
</tr>
<tr>
<td>40 mg</td>
<td>2.2±0.8 (83.1±30.6)</td>
<td>2.3±0.8 (86.9±31.3)</td>
</tr>
<tr>
<td>80 mg</td>
<td>2.1±0.8 (82.4±31.6)</td>
<td>2.2±0.9 (85.9±33.9)</td>
</tr>
<tr>
<td>All doses combined</td>
<td>2.2±0.8 (82.4±30.3)</td>
<td>2.3±0.8 (87.9±31.2)</td>
</tr>
</tbody>
</table>

**Notes:** Values are expressed as mean ± SD. **Abbreviations:** EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol; SD, standard deviation.
(42.4%, 42.3%, 38.6%, and 37.4% at 10, 20, 40, and 80 mg respectively) (Tables S2–S5).

**Discussion**

In this real-world study, a substantial number of high-risk CVD patients did not achieve guideline-recommended LDL-C levels on atorvastatin monotherapy. Although a large proportion of these patients (71%−78%) attained the recommended treatment levels of LDL-C <100 mg/dL, more than 65% did not achieve the optional level of <1.8 mmol/L (<70 mg/dL), even when on therapy at the highest atorvastatin dose of 80 mg. Moreover, 30%−40% of these patients had LDL-C levels more than 0.5 mmol/L (20 mg/dL) above 1.8 mmol/L (70 mg/dL), regardless of atorvastatin dose. Overall, these results indicate a need for more effective LDL-C-lowering therapy in high CVD-risk patients.

Cholesterol-lowering management has improved over time due to increased awareness and better use of lipid-lowering therapy. However, several observational studies and national surveys have shown that many high-risk CHD patients still do not meet guideline-recommended LDL-C levels on statin monotherapy, attributed to various reasons, including less than optimal use of lipid-lowering therapy, noncompliance, and cost factors. The attainment rates for specified LDL-C levels in our study were similar to those previously reported during 1998–2008 in the US, in which 58%−81% of high-risk CHD patients attained LDL-C levels <2.6 mmol/L (<100 mg/dL) and 26%−43% attained LDL-C levels of <1.8 mmol/L (<70 mg/dL).

For those high-risk patients who do not achieve recommended LDL-C levels on statin monotherapy, more intensive treatment, by statin up titration, switching to a higher efficacy statin, and/or combination therapy can be considered. In our study of high-risk CHD and AVD patients, the majority were treated with medium efficacy-level, LDL-C-lowering doses of atorvastatin (20 and 40 mg doses). Nonetheless, the proportion of patients who attained LDL-C <1.8 mmol/L (<70 mg/dL) improved little at any atorvastatin dose, in both databases. It should be noted that while our study included patients who were both newly initiated and on existing atorvastatin therapy, as many as 34%−81% of the patients in both databases had been on some type of lipid-lowering therapy during the baseline period, including 22%−70% who were on prior atorvastatin therapy, and this may have

![Figure 3 Distribution of LDL-C levels around <1.8 mmol/L (<70 mg/dL) and <2.6 mmol/L (<100 mg/dL) cut points.](https://www.dovepress.com/)

**Abbreviations:** EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.
expected to be completed in 2014 will provide additional
information on the role of ezetimibe in combination with
simvastatin with respect to clinical outcomes.39

Although the reasons for therapy choice were not evaluated
in our analysis, taken together, our study and these studies
suggest that many high-risk patients would likely benefit
from uptitration of their atorvastatin dose, where possible,
and/or switching to a higher potency statin (rosuvastatin) or
combination therapy. Doubling the statin dose can provide an
additional 6% reduction in LDL-C levels,16,28,40 and switching
from atorvastatin doses to a higher potency statin (rosuvas-
tatin), can provide further increases in LDL-C-lowering,
by as much as 10%, depending on the dose. For example,
the LDL-C-lowering efficacy of atorvastatin ranges from
38% to 55% for 10, 20, 40, and 80 mg doses, whereas that of
rosuvastatin ranges from 45% to 60% for 5, 10, 20, and
40 mg doses.23,41 In randomized controlled clinical trials,
combination therapy with ezetimibe has been shown to
increase LDL-C lowering by as much as 21.3%–27.0%
compared with statin monotherapy, when coadministered
in statin-naïve and/or patients on ongoing statin therapy.42,45
In a recent study, ezetimibe added to atorvastatin reduced LDL-C
levels significantly more than doubling the atorvastatin dose
or switching to (or doubling) higher potency rosuvastatin.46

As noted in these studies, the real-world treatment differences
did not achieve the expected results.

A strength of our study is that it provides information on attainment of guideline-recommended LDL-C levels
with atorvastatin monotherapy use in a real-world clinical
practice setting. However, as with most retrospective,
observational studies, there are inherent limitations. We
were unable to determine whether LDL-C values were
accounted for the minimal change observed in LDL-C
levels. Atorvastatin uptitration for individual patients was not assessed in this study, but our results are consistent with
the percent patient attainment of LDL-C levels <1.8 mmol/L
(<70 mg/dL) observed overall with moderate to high doses
of statin in other studies.24,25

Several observational and randomized controlled clinical
trials have reported that statin uptitration and/or combination
therapy with higher LDL-C-lowering efficacy are more
effective than moderate-potency therapy for reducing LDL-C
and improving goal attainment in high-risk patients.24,25,29–35
However, these studies showed that >50% of these patients
still did not achieve the more stringent LDL-C levels of
<1.8 mmol/L (<70 mg/dL) and that high-potency statin
therapy as well as the combination of statin with ezetimibe
or niacin therapies were used less frequently compared with
moderate-potency statins in these patients. Lower use of
such alternatives was attributed to patient noncompliance/
tolerance, physician nonadherence to guidelines, and/or
cost factors. A lack of definitive data showing the benefit of
combination therapy on cardiovascular outcomes compared
with statin monotherapy may limit its use.36–38 Both fibrates
and niacin failed to show further reductions of events when
combined with statins compared with statins alone.36–38

Furthermore, the coadministration of niacin and fibrates with
statins increases the risk of myopathy and rhabdomyolysis. In
a recent study, niacin plus the antiflushing agent laropiprant
in combination with simvastatin increased the incidence of
some types of nonfatal, serious side effects, thus bringing
into question the overall safety of niacin.37 An ongoing trial
expected to be completed in 2014 will provide additional

\[
\begin{array}{c|c|c|c|c}
\text{GE Centricity EMR} & \text{Humana Medicare} \\
\hline
\text{Atorvastatin 10 mg} & \text{Atorvastatin 20 mg} & \text{Atorvastatin 40 mg} & \text{Atorvastatin 80 mg} \\
\hline
\end{array}
\]

Figure 4 Proportion of patients with LDL-C <1.8 mmol/L (<70 mg/dL) and <2.6 mmol/L (<100 mg/dL) by atorvastatin dose.
Abbreviations: EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.
based on direct methods or were estimated based on the Friedewald equation. With regard to the cohort selection process, it is possible that the CHD and AVD patients within the databases may not have been fully identified due to a lack of specific information regarding all CHD and AVD risk factors. Although we were able to identify atorvastatin prescriptions written by the physician (in the GE Centricity EMR) and claims for atorvastatin prescriptions filled (in the Humana Medicare database), we were unable to verify patient adherence/compliance to their prescribed medication. Furthermore, because we were unable to confirm if and when atorvastatin prescriptions were filled in the GE Centricity EMR database, we were unable to require as inclusion criteria one atorvastatin prescription fill within 45 days of the follow-up LDL-C measure, in this database. While this study provides an assessment of LDL-C-reduction in atorvastatin users identified who were both newly initiated and on existing therapy during a 3-year time period, LDL-C-reduction was not evaluated separately in these patients groups. Those patients identified as new users and who were not at LDL-C goal may not have yet had the opportunity for further up titration of lipid-lowering therapy. Inclusion of both new and previous users of lipid-lowering therapy also precluded assessment of LDL-C-reductions from pretreatment lipid levels. Furthermore, the majority of patients in the Humana Medicare analysis were on prior atorvastatin therapy. Although our data suggest that a high percentage of patients were on newly initiated atorvastatin monotherapy in the GE Centricity EMR database, the minimal reduction observed in LDL-C levels suggests that many of these patients may have been on prior lipid-lowering therapy that was not captured in the EMR. We also did not have information on intended patient lipid goals or patient therapy choices, including whether patients underwent atorvastatin up titration, in either database. We also had no information on side effects or other patient-level factors that may have precluded treatment to lower LDL-C targets.

In conclusion, in this real-world setting, a large percentage of high-risk CVD patients treated with atorvastatin monotherapy were not at LDL-C goal, regardless of dose. These data suggest that more effective lipid-lowering strategies, such as statin up titration, use of higher potency statins, and/or combination therapy, may be needed to achieve optimal LDL-C-lowering in high-risk patients. It may be prudent for physicians to carefully weigh considerations of safety, tolerability, and anticipated efficacy when making therapeutic decisions, based on individual patient risk. Additionally, improved patient and provider education is still needed to reinforce use of effective therapies and medication adherence.

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Author contributions

All authors jointly developed the manuscript content and were involved in at least one of the following: conception, design, data acquisition, analysis, statistical analysis, interpretation of data, drafting the manuscript, and/or revising the manuscript for important intellectual content. All authors provided final approval of the version to be published.

Disclosure

NJ Zhang received research funding from Merck & Co., Inc. to conduct the Humana Medicare analysis; C Zhao, D Ramey, JE Tomassini, AM Tershakovec, and DR Neff are current employees, and E Marrett and Q Zhang are former employees of Merck Sharp & Dohme Corp., a subsidiary of Merck & Co., Inc. and may hold stock/stock options in the company.

References


Supplementary materials

Table S1 Diagnosis and procedure codes used to identify patients

<table>
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<tr>
<th>Comorbid disease</th>
<th>ICD9 diagnosis codes</th>
<th>CPT codes</th>
<th>ICD-9 procedure codes</th>
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<td>Diabetes</td>
<td>250.xx</td>
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<td>CHD/AVD</td>
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<td>Acute myocardial infarction</td>
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<td>Previous myocardial infarction</td>
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<td>Ischemic stroke</td>
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<td>Peripheral arterial disease</td>
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<td>Angina pectoris</td>
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<tr>
<td>Other acute, subacute ischemic heart disease (eg, unstable angina, acute coronary syndrome)</td>
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</tr>
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<td>Revascularization</td>
<td>33140–33141, 33510, 33511, 33512, 33513, 33514, 33515, 33516, 33517, 33518, 33519, 33521, 33522, 33523, 33533, 33534, 33535–33536, 33537, 33572, 92973, 92980, 92981, 92982, 92984, 92995, 92996, 93540</td>
<td>06.66, 36.01–36.02, 36.05, 36.1x–36.3x</td>
<td></td>
</tr>
<tr>
<td>Acute carotid artery procedures</td>
<td>34001, 35001, 35002, 35201, 35231, 35261, 35286, 35301, 35390, 35501, 35506, 35508, 35509, 35510, 35526, 35601, 35606, 35626, 35642, 35691, 35694, 35701, 35901</td>
<td>00.61–00.65, 38.01–38.02, 38.11–38.12, 38.31–38.32, 38.41–38.42, 39.28, 39.7</td>
<td></td>
</tr>
<tr>
<td>Other forms of chronic ischemic heart disease</td>
<td>414.xx, v45.8, v45.81, v45.82</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Atherosclerosis</td>
<td>429.2x, 433.x0, 434.x0, 437.0x, 437.1x, 440.xx, 444.0x, 444.1x, 445.8x</td>
<td>36.0x–36.3x, 38.12–38.16, 38.18, 38.34, 38.44, 39.22–39.26, 39.28, 39.29, 39.50, 39.71, 39.73, 39.90, 99.10</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: AVD, atherosclerotic vascular disease; CHD, coronary heart disease; CPT, current procedural terminology; ICD-9, International Classification of Diseases, 9th revision.

Table S2 GE Centricity EMR: proportion of patients with LDL-C < 1.8 mmol/L (<70 mg/dL) and proportion with levels >1.8 mmol/L (>70 mg/dL), categorized by distance from goal

<table>
<thead>
<tr>
<th>Atorvastatin dose</th>
<th>% (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LDL-C &lt; 1.8 mmol/L (≤70 mg/dL)</td>
<td>LDL-C ≥1.8 to &lt;2.1 mmol/L (≥70 to &lt;80 mg/dL)</td>
</tr>
<tr>
<td>10 mg</td>
<td>32.5 (31.6–33.4)</td>
</tr>
<tr>
<td>20 mg</td>
<td>35.3 (34.4–36.2)</td>
</tr>
<tr>
<td>40 mg</td>
<td>34.6 (33.7–35.5)</td>
</tr>
<tr>
<td>80 mg</td>
<td>37.3 (36.4–38.2)</td>
</tr>
<tr>
<td>All doses combined</td>
<td>34.8 (33.9–35.7)</td>
</tr>
</tbody>
</table>

Abbreviations: CI, confidence interval; EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.
### Table S3 GE Centricity EMR: proportion of patients with LDL-C <2.6 mmol/L (<100 mg/dL) and proportion with levels ≥2.6 mmol/L (>100 mg/dL), categorized by distance from goal

<table>
<thead>
<tr>
<th>Atorvastatin dose</th>
<th>% (95% CI)</th>
<th>LDL-C &lt;2.6 mmol/L (&lt;100 mg/dL)</th>
<th>LDL-C ≥2.6 to &lt;2.9 mmol/L (≥100 to &lt;110 mg/dL)</th>
<th>LDL-C ≥2.9 to &lt;3.1 mmol/L (≥110 to &lt;120 mg/dL)</th>
<th>LDL-C ≥3.1 mmol/L (≥120 mg/dL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 mg</td>
<td>77.9 (77.1–78.7)</td>
<td>7.6 (7.1–8.1)</td>
<td>4.1 (3.7–4.5)</td>
<td>10.3 (9.7–10.9)</td>
<td></td>
</tr>
<tr>
<td>20 mg</td>
<td>77.7 (76.9–78.5)</td>
<td>8.0 (7.5–8.5)</td>
<td>4.8 (4.4–5.2)</td>
<td>9.5 (8.9–10.1)</td>
<td></td>
</tr>
<tr>
<td>40 mg</td>
<td>78.4 (77.6–79.2)</td>
<td>7.0 (6.5–7.5)</td>
<td>4.4 (4.0–4.8)</td>
<td>10.2 (9.6–10.8)</td>
<td></td>
</tr>
<tr>
<td>80 mg</td>
<td>77.9 (77.1–78.7)</td>
<td>6.4 (5.9–6.9)</td>
<td>5.4 (5.0–5.8)</td>
<td>10.3 (9.7–10.9)</td>
<td></td>
</tr>
<tr>
<td>All doses combined</td>
<td>78.0 (77.2–78.8)</td>
<td>7.3 (6.8–7.8)</td>
<td>4.6 (4.2–5.0)</td>
<td>10.0 (9.4–10.6)</td>
<td></td>
</tr>
</tbody>
</table>

**Abbreviations:** CI, confidence interval; EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.

### Table S4 Humana Medicare: proportion of patients with LDL-C <1.8 mmol/L (<70 100 mg/dL) and proportion with levels ≥1.8 mmol/L (>70 mg/dL) categorized by distance from goal

<table>
<thead>
<tr>
<th>Atorvastatin dose</th>
<th>% (95% CI)</th>
<th>LDL-C &lt;1.8 mmol/L (&lt;70 mg/dL)</th>
<th>LDL-C ≥1.8 to &lt;2.1 mmol/L (≥70 to &lt;80 mg/dL)</th>
<th>LDL-C ≥2.1 to &lt;2.3 mmol/L (≥80 to &lt;90 mg/dL)</th>
<th>LDL-C ≥2.3 mmol/L (≥90 mg/dL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 mg</td>
<td>25.7 (25.0–26.4)</td>
<td>15.4 (14.9–16.0)</td>
<td>16.5 (15.9–17.1)</td>
<td>42.4 (41.7–43.2)</td>
<td></td>
</tr>
<tr>
<td>20 mg</td>
<td>27.3 (26.6–27.8)</td>
<td>15.3 (14.8–15.8)</td>
<td>15.1 (14.6–15.6)</td>
<td>42.3 (41.6–43.1)</td>
<td></td>
</tr>
<tr>
<td>40 mg</td>
<td>29.7 (29.0–30.9)</td>
<td>15.1 (14.6–15.7)</td>
<td>15.6 (15.1–16.2)</td>
<td>38.6 (37.9–39.3)</td>
<td></td>
</tr>
<tr>
<td>80 mg</td>
<td>33.0 (32.3–33.7)</td>
<td>14.9 (14.4–15.4)</td>
<td>14.7 (14.2–15.2)</td>
<td>37.4 (36.7–38.1)</td>
<td></td>
</tr>
<tr>
<td>All doses combined</td>
<td>28.3 (27.6–29.0)</td>
<td>15.5 (15.0–16.1)</td>
<td>15.5 (15.0–16.1)</td>
<td>40.7 (40.0–41.4)</td>
<td></td>
</tr>
</tbody>
</table>

**Abbreviations:** CI, confidence interval; EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.

### Table S5 Humana Medicare: proportion of patients with LDL-C <2.6 mmol/L (<100 mg/dL) and proportion with levels >2.6 mmol/L (>100 mg/dL), categorized by distance from goal

<table>
<thead>
<tr>
<th>Atorvastatin dose</th>
<th>% (95% CI)</th>
<th>LDL-C &lt;2.6 mmol/L (&lt;100 mg/dL)</th>
<th>LDL-C ≥2.6 to &lt;2.9 mmol/L (≥100 to &lt;110 mg/dL)</th>
<th>LDL-C ≥2.9 to &lt;3.1 mmol/L (≥110 to &lt;120 mg/dL)</th>
<th>LDL-C ≥3.1 mmol/L (≥120 mg/dL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 mg</td>
<td>71.5 (70.8–72.2)</td>
<td>9.1 (8.7–9.5)</td>
<td>6.8 (6.4–7.2)</td>
<td>12.6 (12.2–13.1)</td>
<td></td>
</tr>
<tr>
<td>20 mg</td>
<td>70.6 (69.9–71.3)</td>
<td>9.2 (8.8–9.6)</td>
<td>6.2 (5.8–6.6)</td>
<td>14.0 (13.5–14.5)</td>
<td></td>
</tr>
<tr>
<td>40 mg</td>
<td>73.3 (72.6–74.0)</td>
<td>8.5 (8.1–8.9)</td>
<td>5.6 (5.3–6.0)</td>
<td>12.6 (12.1–13.1)</td>
<td></td>
</tr>
<tr>
<td>80 mg</td>
<td>73.8 (73.1–74.5)</td>
<td>8.5 (8.1–8.9)</td>
<td>5.3 (5.0–5.6)</td>
<td>12.4 (11.9–12.9)</td>
<td></td>
</tr>
<tr>
<td>All doses combined</td>
<td>72.0 (71.3–72.7)</td>
<td>8.8 (8.4–9.2)</td>
<td>6.0 (5.6–6.4)</td>
<td>13.2 (12.7–13.7)</td>
<td></td>
</tr>
</tbody>
</table>

**Abbreviations:** CI, confidence interval; EMR, electronic medical record; LDL-C, low-density lipoprotein cholesterol.