Geographic variations in lipid-lowering therapy utilization, LDL-C levels, and proportion retrospectively meeting the ACC/AHA very high-risk criteria in a real-world population of patients with major atherosclerotic cardiovascular disease events in the United States

Seth J. Baum, Pallavi B. Rane, Sasikiran Nunn, Mohdhar Habib, Kiran Philip, Kainan Sun, Xin Wang, Rolin L. Wade

Department of Integrated Medical Sciences, Charles E. Schmidt College of Medicine, Florida Atlantic University, Boca Raton, FL USA
IQVIA, Plymouth Meeting, PA USA

Key words: Atherosclerotic cardiovascular disease
Very high-risk
LDL-C
Lipid-lowering therapy

Objective: We assessed national- and state-level geographic variations among patients with a history of ≥1 major atherosclerotic cardiovascular disease (ASCVD) event in: (1) the proportion of patients with retrospectively identified 2018 American College of Cardiology/American Heart Association guideline very high-risk (VHR) ASCVD criteria; (2) utilization of guideline-directed lipid-lowering therapy (LLT); and (3) the proportion of patients with persistent low-density lipoprotein cholesterol (LDL-C) elevations despite statin and/or ezetimibe use.

Methods: A retrospective cohort study using the Prognos LDL-C database linked to IQVIA longitudinal medical and prescription claims databases. The study period was from January 01, 2011, to November 30, 2019 and the index period was from January 01, 2016, to November 30, 2019; the index date was defined as the most recent LDL-C test during the index period. The study included patients aged ≥18 years at index who had a measured LDL-C level during the index period and had ≥1 inpatient/outpatient claim for ASCVD during the 5-year pre-index period.

Results: Of patients with any ASCVD (N=4652,468), 1537,514 (33.1%) patients had ≥1 major ASCVD event. Among patients with ≥1 major ASCVD event, the VHR ASCVD criteria were retrospectively identified in 1139,018 (74.1%) patients; Hawaii had the highest (81.7%) and Colorado the lowest (65.0%) proportion of these patients. Nationally, 48.8% and 50.2% of patients with ≥1 major ASCVD event and retrospectively identified VHR ASCVD criteria, respectively, had current LLT use; Massachusetts and Colorado had the highest and lowest proportions, respectively. After standardizing for age and sex, 57.3% and 58.8% of patients with ≥1 major ASCVD event and retrospectively identified VHR ASCVD criteria, respectively, had LDL-C ≥70 mg/dL (≥1.8 mmol/L) despite statin and/or ezetimibe use, with substantial state-level variations observed.

Conclusions: The study highlights high rates of elevated LDL-C and pervasive underuse of LLT in health-insured patients with a history of major ASCVD events treated in the United States, with state-level geographic variations observed.

1. Introduction

Cardiovascular disease remains the leading cause of death in the United States (US) [1], with a combined annual direct and indirect cost burden of $555 billion, which is predicted to rise to $1.1 trillion from 2015 to 2035 [2]. Low-density lipoprotein cholesterol (LDL-C) is a well-established, causal risk factor for atherosclerotic cardiovascular disease (ASCVD) [3,4]. Accordingly, lowering of LDL-C with lipid-lowering therapy (LLT) significantly reduces the risk of cardiovascular events [5]. Recent updates in the 2018 American College of Cardiology/American Heart Association (ACC/AHA) Multisociety blood cholesterol guidelines introduced the very high-risk (VHR) ASCVD classification [6]. Patients in this VHR group have multiple major ASCVD events (i.e. recent acute coronary syndrome [ACS], history of myocardial infarction [MI] other than ACS, history of ischemic stroke [IS], or symptomatic peripheral arterial disease [PAD]), or a single major ASCVD event and a history of multiple high-risk conditions [6]. In patients with VHR ASCVD and
LDL-C $\geq 70$ mg/dL ($\geq 1.8$ mmol/L) despite high-intensity or maximally tolerated statin therapy, the addition of non-statin therapy (ezetimibe and/or a proprotein convertase subtilisin/kexin type 9 inhibitor [PCSK9i]) is recommended [6].

Previous studies of US clinical practice have reported suboptimal utilization of guideline-directed LLT for patients with hypercholesterolemia, with poor statin adherence, underdosing and/or discontinuation frequently observed [7-15]. However, the clinical and treatment characteristics of patients at the highest cardiovascular risk (e.g. those with a history of $\geq 1$ major ASCVD event) remain to be fully elucidated in a real-world setting. Additionally, given the recent introduction of the VHR ASCVD risk stratification in the 2018 ACC/AHA blood cholesterol guidelines, [6] the proportion of patients meeting these criteria in real-world, US clinical practice remains largely unknown. Importantly, the provision of guideline-directed LLT to reduce cardiovascular risk in patients with major ASCVD events should be uniform across the US; however, to date, VHR ASCVD status, LLT utilization, and LDL-C levels in patients with major ASCVD events have not been assessed at the state level in the US. This is an important gap in the literature, with assessment at the state level required to identify potential heterogeneity in the treatment patterns, clinical outcomes, and cardiovascular risk status of patients with major ASCVD events. Moreover, state-level analysis allows for the generalizability of national trends to be assessed, and the uncovering of disparities can inform subsequent public health interventions and policies [16].

Therefore, the objectives of the current study were to describe US national- and state-level geographic variations among patients with a history of $\geq 1$ major ASCVD event with regard to the proportion of patients with retrospectively identified 2018 ACC/AHA guideline VHR ASCVD criteria, [6] utilization of guideline-directed LLT, and the proportion of patients with persistent LDL-C elevations despite statin and/or ezetimibe use.

2. Methods

2.1. Study design and databases

This was a retrospective cohort study using the nationally representative Prognos LDL-C database (Prognos, New York, NY, USA) [17] linked to anonymized longitudinal medical and prescription claims data across IQVIA (Plymouth Meeting, PA, USA) data sources [18]. Prognos data aggregates LDL-C values from large national and regional laboratory providers in the US, and the Prognos Registry is the largest source of clinical diagnostics information in 35 disease areas, with over 8 billion medical records for 150 million patients. The IQVIA Longitudinal Prescription claims (LRx) database captures information on adjudicated dispensed prescriptions sourced from retail, mail, long-term care, and specialty pharmacies, and represents 86% of prescriptions dispensed in retail pharmacies, 55% of prescriptions dispensed by standard mail service, along with 40%-70% of specialty pharmacy volume. Being a phar-
macy claims database, the IQVIA LRx database does not contain clinical and diagnostic information. Thus, to capture clinical characteristics, the IQVIA LRx database was linked to the IQVIA PharMetrics® Plus health plan claims database and the IQVIA Professional Fee Claims database (Dx). The aggregated IQVIA PharMetrics Plus database comprises adjudicated claims for more than 130 million unique patients across the US and is sourced directly from insurance companies. IQVIA PharMetrics Plus data have a diverse representation of geography, employers, payers, providers, and therapy areas. The IQVIA Dx database contains unadjudicated medical claims from office-based physicians, ambulatory facilities, and hospital-based physicians, and is sourced from clearing houses (also referred to as switches) involved in claims processing. The study utilized de-identified health claims data and thus was exempt from institutional review board review.

The overall study period was from January 01, 2011, to November 30, 2019. The index period was from January 01, 2016, to November 30, 2019, to allow for at least a 5-year pre-index period, with the index date defined as the most recent LDL-C test during the index period (Supplementary Fig. 1). No minimum post-index follow-up was required. The 5-year pre-index period was used to capture the patient’s ASCVD status, history of risk factors, and major ASCVD events among all patients, to identify those who retrospectively met the VHR ASCVD criteria per the updated ACC/AHA guidelines published in November 2018 during the index period of the current study [6] (Supplementary Fig. 1; Supplementary Table 1).

Patients across the different IQVIA databases and the Prognos LDL-C dataset were linked to create a final cohort using a Health Insurance Portability and Accountability Act (HIPAA)-compliant encrypted ID. The study complied with all applicable laws regarding patient privacy, using HIPAA-compliant de-identified retrospective data sources. No direct patient contact or primary collection of individual patient data occurred. Study results were in tabular form and aggregate analyses that omitted patient identification.

2.2. Patients and outcomes

The study included patients aged ≥18 years at index, who had a measured LDL-C level during the index period and had ≥1 inpatient/outpatient claim for ASCVD during the 5-year pre-index period identified using International Classification of Diseases (ICD)-9, ICD-10, and/or Current Procedural Terminology codes.

From all patients identified with ASCVD, the proportion of patients with a history of ≥1 major ASCVD event(s) and the proportion of patients with retrospectively identified 2018 ACC/AHA criteria for VHR ASCVD in the 5-year pre-index period were estimated at the national and state levels [6]. While state-level data were calculated and reported for all states, while ranking, only states with a sample size in the top 90th percentile were considered (states with a sample size in the bottom 10th percentile were not ranked).

Per the 2018 ACC/AHA guidelines [6], major ASCVD events were defined as the presence of recent ACS (within the past 12 months), history of MI (other than a recent ACS event), history of IS, or symptomatic PAD, identified using ICD-9/ICD-10 codes. Per these guidelines [6], VHR ASCVD was retrospectively defined in the current study as a history of multiple major ASCVD events, or 1 major ASCVD event and a history of multiple high-risk conditions during the 5-year pre-index period. The detailed criteria and associated operational definitions, including diagnostic codes, for major ASCVD events and high-risk conditions used in the current study are reported in Supplementary Table 1.

LDL-C distribution of the patients was assessed using the most recent LDL-C value measured on the index date. Current LTT use was estimated at the national and state levels, with patients classed as in receipt of current LTT if they received any statin, ezetimibe, or PCSK9i therapy on the index date or during the 3 months prior to the index date. National-level estimates for LTT use on the index date or during the 12 months prior to the index date were also calculated for all patient groups. The following were the categories of LTT use: “PCSks9 use (monotherapy or in combination with statins)”; “statin only”; “statin plus ezetimibe”; and “ezetimibe only”.

Age- and sex-standardized rates of patients with a history of ≥1 major ASCVD event, patients with retrospectively identified VHR ASCVD criteria per the 2018 ACC/AHA guidelines [6], current LTT use, and patients with elevated LDL-C (≥70 mg/dL [≥1.8 mmol/L] and ≥100 mg/dL [≥2.6 mmol/L]) despite current statin and/or ezetimibe use in the 2 ASCVD subgroups were reported. These were calculated by using the age and sex distribution of a standard population of patients with ASCVD in the US. A standard population of patients with ASCVD was used for age and sex standardization as opposed to census population estimates because the age distribution of patients with ASCVD is different from the age distribution of the general population in the US; thus, the census population cannot be considered a standard population [19]. Prevalence estimates of the standard ASCVD population by sex and across each age group and sex category in the US were computed in a separate sample of patients who were enrolled in the IQVIA PharMetrics Plus database during 2017. These prevalence rates across each age group and sex category were then applied to the most recent census data (2017) available at the time of the analysis to obtain the final standard population to be used for the purpose of this study [19,20]. Directly standardized rates were calculated using the following formula: directly standardized rate = \( \frac{r_1N_1 + r_2N_2 + r_3N_3 + \ldots + r_kN_k}{N_1 + N_2 + N_3 + \ldots + N_k} \), where \( r_k \) equals the rate in k-th stratum of study sample (stratum refers to each age group category by sex) and \( N_k \) equals the number of persons in k-th stratum of the standard population.

2.3. Statistical analysis

Analyses were conducted using SAS version 9.3 (SAS Institute, Cary, NC, USA). The study was descriptive in nature and formal statistical tests were not conducted. Mean, median, and standard deviation (SD) were generated as measures of central tendency and variance for continuous variables. Frequencies and percentages were calculated for categorical variables.

3. Results

3.1. Baseline characteristics

From a total population of 33,910,626 patients with a measured LDL-C level in the linked database during the index period, the study included 4652,468 patients with a history of any ASCVD in the 5-year pre-index period. The mean (SD) age of the ASCVD patients was 70.0 (11.8) years and 52.6% were male; their mean (SD) baseline LDL-C was 90.8 (35.3) mg/dL (2.4 [0.9] mmol/L). The majority of patients (56.5%) were insured by a commercial payer, and the Southern region of the US had the highest proportion (51.5%) of patients with ASCVD (Table 1). Of patients with any ASCVD, 1537,514 (33.1%) patients had a history of ≥1 major ASCVD event in the 5-year pre-index period (Table 1). The mean (SD) LDL-C in patients with a history of ≥1 major ASCVD event was 87.2 (35.6) mg/dL (2.3 [0.9] mmol/L); additional patient clinical characteristics are reported in Table 2.

3.2. Prevalence of VHR ASCVD criteria

Table 3 reports geographic variations in study outcomes in the states with the 10 highest and 10 lowest proportions of patients for each respective outcome; geographic variations in study outcomes across all states is reported in Supplementary Table 2. Among patients with a history of ≥1 major ASCVD event, the 2018 ACC/AHA guideline VHR ASCVD criteria were retrospectively identified in 1139,018 (74.1%) patients during the 5-year pre-index period. The most common VHR ASCVD qualifying event identified during the 5-year pre-index period was...
Table 1
Baseline demographic characteristics.

| Demographics | Patients with a history of ASCVD (N=4652,468) | Patients with a history of ≥1 major ASCVD event (N=1537,514) | Patients with retrospectively identified VHR ASCVD criteria (N=1139,018) |
|--------------|-----------------------------------------------|---------------------------------------------------------------|------------------------------------------------------------------|
| Age, years   | Mean ± SD                                     | 70.0 ± 11.8                                                   | 71.0 ± 11.3                                                      |
|              | Median (min, max)                             | 71 (18, 119)                                                  | 72 (18, 119)                                                    |
| Age group, years, N (%) | 18–34                                     | 42,481 (0.9)                                               | 14,068 (0.9)                                                   | 5513 (0.5)                                        |
|              | 35–44                                        | 96,669 (2.1)                                              | 34,453 (2.2)                                                   | 18,135 (1.6)                                      |
|              | 45–54                                        | 332,144 (7.1)                                             | 119,093 (7.8)                                                  | 72,745 (6.4)                                      |
|              | 55–64                                        | 890,908 (19.2)                                           | 305,829 (19.9)                                                 | 199,896 (17.6)                                    |
|              | 65+                                           | 3290,266 (70.7)                                          | 1064,071 (69.2)                                               | 842,729 (74.0)                                     |
| Sex, N (%)   | Male                                         | 2446,291 (52.6)                                          | 846,922 (55.1)                                                  | 622,203 (54.6)                                    |
|              | Female                                        | 221,767 (47.4)                                          | 687,548 (45.0)                                                  | 516,815 (45.4)                                    |
| Geographic region, N (%) | Northeast                     | 963,084 (20.7)                                           | 297,772 (19.4)                                                 | 215,266 (18.9)                                    |
|              | Midwest                                      | 489,360 (10.5)                                          | 172,428 (11.2)                                                 | 131,773 (11.6)                                    |
|              | South                                        | 2393,943 (51.5)                                         | 793,764 (51.6)                                                 | 590,291 (51.8)                                    |
|              | West                                         | 806,068 (17.2)                                          | 273,545 (17.8)                                                 | 201,685 (17.7)                                    |
|              | Unknown                                      | 13 (0.0)                                                  | 5 (0.0)                                                       | 3 (0.0)                                           |
| Payer type, N (%) | Commercial                  | 2630,016 (56.5)                                         | 877,517 (57.1)                                                 | 618,654 (54.3)                                    |
|              | Medicare                                     | 1984,505 (42.7)                                         | 644,005 (41.9)                                                 | 509,195 (44.7)                                    |
|              | Other                                        | 37,947 (0.8)                                             | 15,992 (1.0)                                                  | 11,169 (1.0)                                      |

ASCVD, atherosclerotic cardiovascular disease; SD, standard deviation; VHR, very high-risk.

![Fig. 1](image-url) Heatmap of US geographic variations in proportions of (A) patients with a history of ≥1 major ASCVD event and (B) patients with retrospectively identified VHR ASCVD criteria.

Data are age- and sex-standardized percentages. Due to low sample sizes, values for Alaska, Montana, North Dakota, South Dakota, Vermont, and Wyoming are masked. ASCVD, atherosclerotic cardiovascular disease; NA, not available; VHR, very high-risk.

a history of MI (Table 2). In this analysis, Hawaii had the highest proportion (81.7%), while Colorado had the lowest proportion (65.0%) of patients with retrospectively identified VHR ASCVD criteria (Table 3; Supplementary Table 2; Fig. 1).

3.3. Geographic variations in LLT use

We assessed LLT use both in the 12-month and 3-month period prior to the index LDL-C date. Nationally, 64.5% of patients with a history of ≥1 major ASCVD event and 66.6% of patients with retrospectively identified VHR ASCVD criteria had ≥1 claim of any LLT use in the 12-month pre-index period (Table 2). In the 3-month pre-index period, 48.8% of patients nationally with a history of ≥1 major ASCVD event had ≥1 claim of current LLT use and, of these, 93.4% of patients received statin monotherapy with 42.2%, 44.4%, and 6.7% of them receiving high-, medium-, and low-intensity statins, respectively. Only 3.5% of patients received statin plus ezetimibe combination therapy and only 0.9% of patients received PCSK9i therapy (monotherapy or in combination with statins) (Table 2).

Nationally, 50.2% of patients with retrospectively identified VHR ASCVD criteria were in receipt of current LLT in the 3-month pre-index period. Of these, 42.2% of patients received high-intensity statins, and 0.9% of patients received PCSK9i (Table 2). Fig. 2 displays a heatmap of US geographic variations in age- and sex-standardized current LLT use among patients with a history of ≥1 major ASCVD event and patients with retrospectively identified VHR ASCVD criteria. The state with the highest current LLT use among patients with a history of ≥1 major ASCVD event and among patients with retrospectively identified VHR ASCVD criteria was Massachusetts, with 55.4% and 57.3% of patients, respectively, treated in the 3-month pre-index period (Fig. 2). The state with the lowest current LLT utilization among patients with a history of ≥1 major ASCVD event and among patients with retrospectively identified VHR ASCVD criteria was Colorado, with only 39.6% and 42.2% of patients, respectively, treated in the 3-month pre-index period (Fig. 2).
3.4. Geographic variations in persistent LDL-C elevations despite statin and/or ezetimibe use

Given that only 0.9% of both patients with ≥1 major ASCVD event and patients with retrospectively identified VHR ASCVD criteria used PCSK9i therapy, we excluded these patients while assessing persistent LDL-C elevations despite current LLT use. At the national level, after standardizing for age and sex, 57.3% and 20.3% of patients had a history of ≥1 major ASCVD event had LDL-C ≥70 mg/dL (≥1.8 mmol/L) and LDL-C ≥100 mg/dL (≥2.6 mmol/L), respectively, despite current LLT with statins and/or ezetimibe. Similarly, nationally, 58.8% and 23.7% of patients with retrospectively identified VHR ASCVD criteria had LDL-C ≥70 mg/dL (≥1.8 mmol/L) and LDL-C ≥100 mg/dL (≥2.6 mmol/L), respectively, despite current LLT with statins and/or ezetimibe.

The Central Illustration displays a heatmap of US geographic variations in the proportions of patients with a history of ≥1 major ASCVD event and patients with retrospectively identified VHR ASCVD criteria with elevated LDL-C ≥70 mg/dL (≥1.8 mmol/L) and LDL-C ≥100 mg/dL (≥2.6 mmol/L), despite current LLT with statins and/or ezetimibe. Substantial state-level variations in persistent LDL-C elevations despite current LLT with statins and/or ezetimibe were observed: Maine and Hawaii had the highest (68.5%) and lowest (46.9%) proportions, respectively, of patients with a history of ≥1 major AS-

Table 2
Clinical characteristics.

| Index LDL-C, mg/dL* | Patients with a history of ASCVD (N=4652,468) | Patients with a history of ≥1 major ASCVD event (N=1537,514) | Patients with retrospectively identified VHR ASCVD criteria (N=1139,018) |
|---------------------|---------------------------------|-------------------------------------------------|-------------------------------------------------|
| Mean ± SD           | 9.0 ± 3.5                        | 8.7 ± 3.6                                       | 8.7 ± 3.6                                       |
| Median (min, max)   | 85 (10,495)                     | 81 (10,492)                                     | 81 (10,492)                                     |

* LDL-C was assessed among all patients with or without current LLT. ACS, acute coronary syndrome; ASCVD, atherosclerotic cardiovascular disease; CABG, coronary artery bypass graft; CHF, congestive heart failure; eGFR, estimated glomerular filtration rate; HeFH, heterozygous familial hypercholesterolemia; IS, ischemic stroke; LDL-C, low-density lipoprotein cholesterol; LLT, lipid-lowering therapy; MI, myocardial infarction; PAD, peripheral arterial disease; PCI, percutaneous coronary intervention; PCSK9i, proprotein convertase subtilisin/kexin type 9 inhibitor; SD, standard deviation; VHR, very high-risk.
Table 3
Geographic variation in major ASCVD events, retrospectively identified VHR ASCVD criteria, current LLT patterns, and LDL-C ≥70 mg/dL despite current LLT with statins and/or ezetimibe (age and sex standardized).

| Patient subgroup | National average | State-level: 10 highest proportions | State-level: 10 lowest proportions |
|------------------|------------------|-----------------------------------|-----------------------------------|
|                  |                  | ≥1 major ASCVD event (%) | VHR ASCVD criteria (%) | ≥1 major ASCVD event on statins and/or ezetimibe (%) | VHR ASCVD criteria (% of major ASCVD events on statins and/or ezetimibe) | ≥1 major ASCVD event with LDL-C ≥70 mg/dL (≥1.8 mmol/L) despite statins and/or ezetimibe (%) | VHR ASCVD criteria with LDL-C ≥70 mg/dL (≥1.8 mmol/L) despite statins and/or ezetimibe (%) |
|                  |                  | (%)                      | (%)                          | (%)                                          | (%)                                                                 | (%)                                                                 | (%)                                                                 |
|                  |                  | 33.6                     | 72.1                        | 48.3                                        | 49.8                                                                 | 57.3                                                                 | 58.8                                                                 |
| Maine            |                  | (40.7)                   | (81.7)                      | (55.1)                                      | Massachusetts (57.0)                                          | Maine (68.5)                                                     | Maine (70.6)                                                     |
| Oregon           |                  | (40.0)                   | (80.1)                      | (54.2)                                      | Pennsylvania (55.5)                                          | Michigan (63.9)                                                  | Wisconsin (65.8)                                                  |
| Kentucky         |                  | (39.7)                   | (77.3)                      | (54.0)                                      | Louisiana (55.3)                                             | Delaware (55.5)                                                 | Wisconsin (63.9)                                                  |
| Ohio             |                  | (38.9)                   | (76.5)                      | Connecticut (53.8)                           | Louisiana (55.3)                                             | Connecticut (53.0)                                              | Michigan (64.7)                                                   |
| Michigan         |                  | (38.5)                   | (76.4)                      | Kentucky                                    | Delaware (53.7)                                              | Connecticut (55.0)                                              | DC (62.6)                                                        |
| Hawaii           |                  | (37.9)                   | (76.4)                      | Washington                                   | Missouri (52.8)                                              | Indiana (60.9)                                                  | Arkansas (62.5)                                                   |
| Iowa             |                  | (37.7)                   | (76.3)                      | Arkansas                                     | Illinois (51.8)                                              | Missouri (54.0)                                                 | Florida (61.8)                                                    |
| New Hampshire (37.6) |            |                         |                             | New Hampshire (51.5)                         | Rhode Island (51.2)                                         | New Hampshire (53.6)                                            | Georgia (61.7)                                                    |
| Nevada           |                  | (37.6)                   |                             | (75.7)                                      | Michigan (52.3)                                              | Illinois (46.9)                                                 | Maine (66.1)                                                      |
| Minnesota (37.3) |                  | (75.4)                   |                             |                                           | Ohio (51.2)                                                  | Arizona (47.9)                                                  | Illinois (61.5)                                                   |
| New York (27.0)  |                  | (65.0)                   |                             |                                           | Colorado (41.7)                                              | New Mexico (42.8)                                               | DC (54.5)                                                        |
| New Jersey (30.9)|                  | (68.1)                   |                             |                                           | Arizona (43.1)                                               | New Mexico (44.1)                                               | Florida (51.4)                                                    |
| Kansas           |                  | (31.9)                   |                             |                                           | Colorado (42.1)                                              | Delware (50.6)                                                  | Hawaii (47.9)                                                     |
| Oklahoma (32.2)  |                  | (68.8)                   |                             |                                           | Oklahoma (43.4)                                              | Nevada (45.1)                                                   | Kentucky (56.6)                                                   |
| Florida          |                  | (32.3)                   |                             |                                           | Virginia (68.9)                                              | Oklahoma (45.4)                                                 | Washington (52.7)                                                 |
| Colorado (32.9)  |                  | (69.7)                   |                             |                                           | Florida (46.7)                                               | Florida (47.0)                                                  | North Carolina (54.0)                                             |
| Texas (33.0)     |                  | (70.3)                   |                             |                                           | Massachusetts (45.3)                                         | Oregon (45.5)                                                   | North Carolina (55.5)                                             |
| South Carolina (33.2) |            |                         |                             |                                           | North Carolina (48.1)                                        | Utah (53.8)                                                     | Arizona (55.3)                                                   |
| Rhode Island (33.4) |            |                         |                             |                                           | New Jersey (46.8)                                           | New York (48.1)                                                 | North Carolina (56.6)                                             |
| DC              |                  | (33.6)                   |                             |                                           | New Jersey (70.5)                                           | New York (48.1)                                                 | Arizona (54.5)                                                    |

*DC is not a state; however it met the criteria for inclusion in the state-level analysis. ASCVD, atherosclerotic cardiovascular disease; DC, District of Columbia; LDL-C, low-density lipoprotein cholesterol; LLT, lipid-lowering therapy; VHR, very high-risk.
CVD event with LDL-C $\geq$70 mg/dL (≥1.8 mmol/L) despite current LLT with statins and/or ezetimibe (Table 3; Supplementary Table 2; Central Illustration). Maine and Colorado had the highest and lowest proportions, respectively, of patients with a history of $\geq$1 major ASCVD event and patients with retrospectively identified VHR ASCVD criteria, where Maine and Colorado had the highest and lowest proportions, respectively, of patients with LDL-C $\geq$100 mg/dL (≥2.6 mmol/L) despite current LLT with statins and/or ezetimibe (Central Illustration). A similar trend was seen among patients with retrospectively identified VHR ASCVD criteria, where Maine and Colorado had the highest and lowest proportions, respectively, of patients with LDL-C $\geq$100 mg/dL (≥2.6 mmol/L) despite current LLT with statins and/or ezetimibe (Central Illustration). Maine and Hawaii had the highest (70.6%) and lowest (47.9%) proportions, respectively, of patients with retrospectively identified VHR ASCVD criteria with LDL-C $\geq$70 mg/dL (≥1.8 mmol/L) despite current LLT with statins and/or ezetimibe (Table 3; Supplementary Table 2; Central Illustration).

4. Discussion

The current study provides novel real-world data on US geographic variations in LLT utilization and LDL-C levels, with this being the first study to report these outcomes at both the national and subnational level, in patients treated across US clinical practices with a history of $\geq$1 major ASCVD event and among patients with retrospectively identified 2018 ACC/AHA guideline VHR ASCVD criteria during a 5-year pre-index period. Nationally, the results indicate a pervasive unmet treatment need for effective lowering of LDL-C in patients with the highest-risk ASCVD. For example, 58.8% of patients with retrospectively identified VHR ASCVD criteria had LDL-C $\geq$70 mg/dL (≥1.8 mmol/L) despite current LLT with statins and/or ezetimibe. The current results support existing real-world studies conducted in the US that report unmet treatment needs [7–15]. The factors contributing to the elevated LDL-C levels observed in the current study are unclear; however, they may have included patients not taking their medications as prescribed, clinicians not regularly assessing follow-up LDL-C levels, along with a reduced focus on LDL-C levels as a secondary prevention performance measure during the study time period [21]. Broadly, the current results indicate that lipid-modifying management strategies need to be intensified; in patients with VHR ASCVD, the addition of a PCSK9i to maximally tolerated LLT was recommended in the 2018 ACC/AHA guidelines to achieve LDL-C lowering [6], with it now recognized that patients with higher cardiovascular risk derive greater benefit from PCSK9i treatment versus those with lower cardiovascular risk [22–24].

At the state level, we observed substantial geographic variations in the proportion of patients with retrospectively identified 2018 ACC/AHA guideline VHR ASCVD criteria, utilization of guideline-directed LLT, and the proportion of patients with elevated LDL-C despite current statin and/or ezetimibe use. Thus, patients’ cardiovascular risk status and likelihood to receive guideline-directed LLT varied by the state in which they were treated. For example, Massachusetts and Colorado had the highest and lowest proportions, respectively, of patients with a history of $\geq$1 major ASCVD event and patients with retrospectively identified VHR ASCVD criteria with current LLT use. Notably, among patients with a history of $\geq$1 major ASCVD event, Hawaii had the highest proportion of patients with retrospectively identified VHR ASCVD criteria, but these patients were also the “best treated”, with Hawaii having the lowest proportion of patients with retrospectively identified VHR ASCVD criteria with LDL-C $\geq$70 mg/dL (≥1.8 mmol/L) while treated with statins and/or ezetimibe.

The reasons for the state-level variations observed in the current study are likely to be multifactorial and complex, and it should be noted that the assessment of causal variables driving the heterogeneity was beyond the scope of the current study. However, a potential explanation for the clinical inertia observed in the states with the lowest levels of LLT utilization may include unmet educational needs for healthcare providers and patients [25–27]. For example, it has been reported that approximately 50% of healthcare providers in US clinical practices did not read the 2013 ACC/AHA blood cholesterol guidelines [25], and US clinical practices with the highest levels of statin utilization were much more likely to adopt the 2013 ACC/AHA blood cholesterol guidelines versus practices with an underutilization of statins [27]. Additionally, physician factors (e.g., beliefs about statins and cholesterol) and patient factors (e.g., statin intolerance) are also likely to play a role in explaining the state-level variations observed in the current study. For instance, a recent study demonstrated that patients treated by physicians with beliefs in statin benefit were more likely to receive 2013 ACC/AHA guideline-recommended statin intensity, whereas patients treated by clinicians expressing statin safety concerns were less likely to receive statins at a guideline-recommended intensity [28].

The current study had a number of strengths, including the very large sample size and being the first study to examine geographic variations across the US, and reporting metrics related to patients with a history of major ASCVD events and patients with retrospectively identified VHR ASCVD criteria at a subnational level. Additionally, the use of the IQVIA LRs database to supplement claims from the IQVIA PharMetrics Plus database enabled the capture of statin prescriptions that were paid out...
of pocket; otherwise these prescriptions may not have been captured in the traditionally used payer-sourced databases alone [29].

However, the study had a number of limitations. The study was subject to the common limitations arising from the use of insurance claims databases for clinical research such as limited generalizability to populations without health insurance, along with potential errors in coding and recording of data, and the potential for selection bias. To improve health inequity in the US and inform clinical practice and policy, future studies that assess US geographical variations in LLT patterns and LDL-C levels in ASCVD patients without health insurance are needed. Also, other than age and sex standardization, we did not adjust the data, and the potential influence of confounding variables on LLT patterns and LDL-C levels was not assessed. Lastly, the study contains limited data from the time period following the publication of the updated 2018 ACC/AHA blood cholesterol guidelines and it is likely that the results are more reflective of US clinical practice resulting from the 2013 ACC/AHA blood cholesterol guidelines published during the pre-index period of the study [30]. Consequently, the results of the study, including the low utilization rates of non-statin therapies, should be interpreted with the study time period in mind.

In conclusion, the current retrospective cohort study is the first to describe LLT patterns and LDL-C distributions in health-insured patients with major ASCVD events as retrospectively defined by the 2018 ACC/AHA guideline criteria [6] at both the national and subnational level in the US. The study highlights high rates of elevated LDL-C and pervasive underuse of LLT in patients with a history of major ASCVD events, including in the highest-risk patients with retrospectively identified VHR ASCVD criteria, with substantial state-level geographic variation. Consequently, there is an unmet treatment need for improved lipid management—through the adoption and implementation of clinical guidelines and intensification of LLT with non-statin therapy, where appropriate—in order to homogenize treatment and improve cardiovascular risk reduction across the US.

5. Data sharing

Qualified researchers may request data from Amgen clinical studies. Complete details are available at the following: http://www.amgen.com/datasharing.

6. Author contributions

Conception or design of the study; or the acquisition, analysis, or interpretation of data for the work: SJB, PBR, SN, KS, XV, RLW

Drafting the manuscript or revising it critically for important intellectual content: SJB, PBR, SN, MH, KP, KS, XV, RLW

All authors had full access to the study data, gave final approval of the version to be published and agree to be accountable for all aspects of the work.

Declaration of Competing Interest

Seth J. Baum has served on scientific advisory boards, provided consulting, and performed clinical research for Amgen, Sanofi/Regeneron, Esperion, Akcea, AstraZeneca, Boehringer Ingelheim/Lilly, Novo Nordisk, and Gilead; has served as a speaker for Amgen, Boehringer Ingelheim/Lilly, Novo Nordisk, and Arazel; and serves as president of Excel Medical Clinical Trials, LLC and Preventive Cardiology, Inc; spouse owns VitalRemedyMD. Kainan Sun, Xin Wang, and Rolin L. Wade are employees of IQVIA, which received consulting fees from Amgen to conduct this study. Sasikiran Nunna was an employee of IQVIA when the study was conducted and is a current employee of Bristol Myers Squibb. Pallavi B. Rane, Kiran Philip, and Mohdhar Habib are employees of Amgen Inc. and hold Amgen stock.

Acknowledgments

Medical writing support, which was in accordance with Good Publication Practice (GPP3) guidelines, was provided by Liam Gillies, PhD, CMPP, of Cactus Life Sciences and was funded by Amgen Inc. The authors thank Matt Hill of IQVIA for generating the heatmap figures for this manuscript.

Funding

The study was funded by Amgen Inc. The sponsor was involved in the study design; in the analysis and interpretation of data; in writing the report; and in the decision to submit the article for publication.

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.ajpc.2021.100177.

References

[1] Benjamin EJ, Blaha MJ, Chiuve SE, et al. Heart disease and stroke statistics—2017 update: a report from the American Heart Association. Circulation. 2017;135:e146–603. doi:10.1161/CIR.0000000000000485.
[2] American Heart Association & American Stroke Association. Cardiovascular Disease: a continuing burden for America — projections through 2035. https://healthmetrics.heart.org/wp-content/uploads/2017/10/Cardiovascular-Disease-A-Costly-Burden.pdf. Accessed November 2019.
[3] Verschure WM, Jacobs DR, Bloomberg BP, et al. Serum total cholesterol and long-term coronary heart disease mortality in different cultures. Twenty-five-year follow-up of the seven countries study. JAMA 1995;274(2):131–6. doi:10.1001/jama.1995.03530020049031.
[4] Pekkanen J, Linn S, Helis G, et al. Ten-year mortality from cardiovascular disease in relation to cholesterol level among men with and without preexisting cardiovascular disease. N Engl J Med 1990;322(24):1700–7. doi:10.1056/NEJM1990061413224030.
[5] Baigent C, Blackwell L, Emberson J, et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26 randomised trials. Lancet 2010;376:1670–81. doi:10.1016/S0140-6736(10)63550-5.
[6] Grundy SM, Stone NJ, Bailey AL, et al. 2018 AHA/ACC/AACVPR/AAPA/ABC/ACPM/ADA/AGS/APA/ASCPCJA/PCNA guideline on the management of blood cholesterol: executive summary: a report of the American College of Cardiology/American Heart Association Task Force on clinical practice guidelines. J Am Coll Cardiol 2019;73(24):3168–209. doi:10.1016/j.jacc.2018.11.002.
[7] Pittman DG, Chen W, Bowlin SJ, Foody JM. Adherence to statins, subsequent healthcare costs, and cardiovascular hospitalizations. Am J Cardiol 2013;110(1):1162–6. doi:10.1016/j.amjcard.2011.01.052.
[8] Serban M-C, Colantonio LD, Manthirpragada AD, et al. Statin intolerance and risk of future coronary heart events and all-cause mortality following myocardial infarction. J Am Coll Cardiol 2017;69:1386–95. doi:10.1016/j.jacc.2016.12.036.
[9] Colantonio LD, Huang L, Moeda RL, et al. Adherence to high-intensity statins following a myocardial infarction hospitalization among Medicare beneficiaries. JAMA Cardiol 2017;2(8):890–5. doi:10.1001/jamacardio.2017.0911.
[10] Superko H, Williams PT, Dansinger M, Scharfer E. Trends in low-density lipoprotein-cholesterol blood values between 2012 and 2017 suggest sluggish adoption of the recent 2013 treatment guidelines. Clin Cardiol 2019;42(1):101–10. doi:10.1002/clc.22119.
[11] Gaskin FS, Iyengar R, Eatherly M, et al. Impact of new ACC/AHA cholesterol treatment guidelines on statin utilization patterns in the United States. J Am Coll Cardiol 2016;67(13 suppl):S162. doi:10.1016/S0735-1097(16)31965-5.
[12] Ramsaran E, Prespe P, Sundaresan D, et al. Adherence to blood cholesterol treatment guidelines among physicians managing patients with atherosclerotic cardiovascular disease. Am J Cardiol 2019;124(2):169–75. doi:10.1016/j.amjcard.2019.04.017.
[13] Gamoza CM, Safford MM, Levitin EB, et al. Statin underuse and low prevalence of LDL-C control among U.S. adults at high risk of coronary heart disease. Am J Med Sci 2014;348(2):108–14. doi:10.1097/MAJ.0000000000000292.
[14] Kern DM, Balu S, Tunceli O, Anzalone D. Statin treatment patterns and clinical profile of patients with risk factors for coronary heart disease defined by National Cholesterol Education Program Adult Treatment Panel III. Curr Med Res Opin 2014;30(12):2443–51. doi:10.1185/03007995.2014.971151.
[15] Pokharel Y, Tang F, Jones PG, et al. Adoption of the 2013 American College of Cardiology/American Heart Association cholesterol management guideline in cardiology practices nationwide. JAMA Cardiol 2017;2(4):361–9. doi:10.1001/jamacardio.2016.5922.
[16] Barnes JE, Briaz PA, Goodman RA, Bowman BA. Prevention of chronic disease in the 21st century: elimination of the leading preventable causes of premature death and disability in the USA. Lancet 2014;384(9937):45–52. doi:10.1016/S0140-6736(14)60486-8.
