

A standardized framework for risk-based assessment of treatment effect heterogeneity in observational healthcare databases

Alexandros Rekkas, MSc¹ David van Klaveren, PhD^{2,3}, Patrick B. Ryan, PhD⁴

Ewout W. Steyerberg, PhD^{3,5} David M. Kent, PhD² Peter R. Rijnbeek, PhD¹

¹ Department of Medical Informatics, Erasmus University Medical Center, Rotterdam, Netherlands

² Predictive Analytics and Comparative Effectiveness (PACE) Center, Institute for Clinical Research and Health Policy Studies (ICRHPS), Tufts Medical Center, Boston, MA, USA

³ Department of Public Health, Erasmus University Medical Center, Rotterdam, Netherlands

⁴ Janssen Research and Development, 125 Trenton Harbourton Rd, Titusville, NJ 08560, USA

⁵ Department of Biomedical Data Sciences, Leiden University Medical Center, Leiden, The Netherlands

Corresponding author

Alexandros Rekkas, MSc
Department of Medical Informatics
Erasmus University Medical Center
3000 CA Rotterdam, P.O. Box 2040
Email: a.rekkas@erasmusmc.nl

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Abstract

Background: Treatment effects are often anticipated to vary across groups of patients with different baseline risk. Implementation of a risk-based approach in the RCT setting has shown evidence of increase in statistical power for the evaluation of treatment effect heterogeneity. The aim of this study was to extend this approach to the observational setting using a standardized scalable framework.

Methods: The proposed framework consists of five steps: 1) definition of the problem, i.e. the population, the treatment, the comparator and the outcome(s) of interest; 2) identification of relevant databases; 3) development of a prediction model for the outcome(s) of interest; 4) estimation of relative and absolute treatment effect within strata of predicted risk, after adjusting for observed confounding; 5) presentation of the results. We demonstrate our framework by evaluating heterogeneity of the effect of angiotensin-converting enzyme (ACE) inhibitors versus beta blockers on three efficacy and six safety outcomes across three observational databases.

Findings: In our demonstration, patients at low risk of acute myocardial infarction (MI) received negligible absolute benefits for all three efficacy outcomes, though they were more pronounced in the highest risk quarter, especially for hospitalization with heart failure. However, failing diagnostics showed evidence of residual imbalances even after propensity score adjustment.

Interpretation: Application of our framework allows for observing differential risk of outcomes across strata, which offers the opportunity to consider the benefit-harm tradeoff between alternative treatments within targeted subpopulations. Further insights may arise by application to safety and effectiveness questions on a large scale.

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Keywords: observational data, heterogeneity of treatment effect, risk stratification, subgroup analysis

1 Introduction

2 Treatment effects can often vary substantially across individual patients, causing overall effect estimates to be
3 inaccurate for a significant proportion of the patients at hand^{1,2}. Understanding heterogeneity of treatment effects
4 (HTE) has been crucial for both personalized (or precision) medicine and comparative effectiveness research, giving
5 rise to a wide range of approaches for its discovery, evaluation and application in clinical practice. A common
6 approach to evaluating HTE in clinical trials is through subgroup analyses, which are rarely adequately powered
7 and can lead to false conclusions of absence of HTE or exaggerate its presence^{3,4}. In addition, patients differ with
8 regard to multiple characteristics simultaneously, resulting in much richer HTE compared to the one explored with
9 regular on-variable-at-a-time subgroup analyses [Kent, BMJ 2018].

10 Baseline risk is a summary score inherently related to treatment effect that can represent more closely the
11 variability in patient characteristics^{3,5–8}. For example, an invasive coronary procedure—in comparison with medical
12 treatment—improves survival in patients with myocardial infarction at high (predicted) baseline risk but not in
13 those at low baseline risk⁹. It has also been shown that high-risk patients with pre-diabetes benefit substantially
14 more from a lifestyle modification program than low-risk patients¹⁰.

15 Recently, systematic guidance on the application of risk-based methods for the assessment of HTE has been
16 developed for RCT data^{11,12}. After risk-stratifying patients using an existing or an internally derived prediction
17 model, risk stratum-specific estimates of relative and absolute treatment effect are evaluated. Several methods
18 for predictive HTE analysis have been adapted for use in observational data, but risk-based methods are still not
19 readily available and have been highlighted as an important future research need¹².

20 The Observational Health Data Science and Informatics (OHDSI) collaborative has established a global network
21 of data partners and researchers that aim to bring out the value of health data through large-scale analytics by
22 mapping local databases to the Observational Medical Outcomes Partnership (OMOP) Common Data Model
23 (CDM)^{13,14}. A standardized framework applying current best practices for comparative effectiveness studies within
24 the OHDSI setting has been proposed¹⁵. This framework was successfully implemented on a large scale for
25 estimation of average effects of all first-line hypertension treatment classes on a total of 52 outcomes of interest
26 across a global network of nine observational databases¹⁶.

27 We aimed to develop a framework for risk-based assessment of treatment effect heterogeneity in observational
28 healthcare databases, extending the existing methodology developed for the RCT setting. We implemented the
29 framework in a publicly available package providing an out-of-the-box solution for implementing such analyses at
30 scale within any observational database mapped to OMOP-CDM. In a case study we analyzed heterogeneity of the
31 effects of first-line hypertension treatment.

¹ 2 Methods

² The proposed framework defines 5 distinct steps: 1) definition of the research aim; 2) identification of the
³ databases within which the analyses will be performed; 3) prediction of outcomes of interest; 4) estimation
⁴ of absolute and relative treatment effects within risk strata; 5) presentation of the results. We developed
⁵ an open-source R-package for the implementation of the proposed framework and made it publicly available
⁶ (<https://github.com/OHDSI/RiskStratifiedEstimation>). An overview of the entire framework can be found in
⁷ Figure 1.

⁸ 2.1 Step 1: General definition of the research aim

⁹ The typical research aim is: “to compare the effect of treatment to a comparator treatment in patients with disease
¹⁰ with respect to outcomes O_1, \dots, O_n ”.

¹¹ We use a comparative cohort design. This means that at least three cohorts of patients need to be defined at this
¹² stage of the framework:

- ¹³ ■ A single treatment cohort (T) which includes patients with disease receiving the target treatment of interest.
- ¹⁴ ■ A single comparator cohort (C) which includes patients with disease receiving the comparator treatment.
- ¹⁵ ■ One or more outcome cohorts (O_1, \dots, O_n) that contain patients developing the outcomes of interest

¹⁶ 2.2 Step 2: Identification of the databases

¹⁷ Including in our analyses multiple databases representing the population of interest potentially increases the
¹⁸ generalizability of results. Furthermore, the cohorts should preferably have adequate sample size with adequate
¹⁹ follow-up time to ensure precise effect estimation, even within smaller risk strata. Other relevant issues such as the
²⁰ depth of data capture (the precision at which measurements, lab tests, conditions are recorded) and the reliability
²¹ of data entry should also be considered.

²² 2.3 Step 3: Prediction

²³ Our method relies on adequately separating patients into subgroups based on their baseline risk for the outcomes
²⁴ of interest. Therefore, a model—either an existing external model adequately validated on an internally developed
²⁵ one—assigning patient-level risk is required. For internally developing a risk prediction model we adopt the
²⁶ standardized framework focused on observational data that ensures adherence to existing guidelines^{17–19}.

1 We first need to define a target cohort of patients, i.e. the set of patients on whom the prediction model will be
2 developed. In our case, the target cohort is generated by pooling the already defined treatment and comparator
3 cohorts. We develop the prediction model on the propensity score-matched (1:1) subset of the pooled sample to
4 avoid differentially fitting between treatment arms, thus introducing spurious interactions with treatment^{20,21}. We
5 also need to define a set of patients that experience the outcome of interest, i.e. the outcome cohort. Finally,
6 we need to decide the time frame within which the predictions will be carried out, i.e. the patients' time at risk.
7 Subsequently, we can develop the prediction model.

8 It is important that the prediction models display good discriminative ability to ensure that risk-based subgroups
9 are accurately defined. A performance overview of the derived prediction models including discrimination and
10 calibration both in the propensity score matched subset, the entire sample and separately for treated and comparator
11 patients should also be reported.

12 **2.4 Step 4: Estimation**

13 We estimate treatment effects (both on the relative and the absolute scale) within risk strata defined using
14 the prediction model of step 3. We often consider four risk strata, but fewer or more strata can be considered
15 depending on the available power for accurately estimating stratum-specific treatment effects. Effect estimation
16 may be focused on the difference in outcomes for a randomly selected person from the risk stratum (average
17 treatment effect) or for a randomly selected person from the treatment cohort within the risk stratum receiving
18 the treatment under study (average treatment effect on the treated).

19 Any appropriate method for the analysis of relative and absolute treatment effects can be considered, as long
20 as the this is done consistently in all risk strata. Common statistical metrics are odds ratios or hazard ratios
21 for relative scale estimates and differences in observed proportions or differences in Kaplan-Meier estimates for
22 absolute scale estimates, depending on the problem at hand. We estimate propensity scores within risk strata
23 which we then use to match patients from different treatment cohorts or to stratify them into groups with similar
24 propensity scores or to weigh each patient's contribution to the estimation process²².

25 Prior to analyzing results, it is crucial to ensure that all diagnostics are passed in all risk strata. The standard
26 diagnostics we carry out include analysis of the overlap of propensity score distributions and calculation of
27 standardized mean differences of the covariates before and after propensity score adjustment. Finally, we use effect
28 estimates for a large set of negative control outcomes (i.e. outcomes known to not be related with any of the
29 exposures under study) to evaluate the presence of residual confounding not accounted for by propensity score
30 adjustment^{23–25}.

¹ **2.5 Step 5: Presentation of results**

² In the presence of a positive treatment effect and a well-discriminating prediction model we expect an increasing
³ pattern of the differences in the absolute scale, even if treatment effects remain constant on the relative scale
⁴ across risk strata. Due to this scale-dependence of treatment effect heterogeneity, results should be assessed both
⁵ on the relative and the absolute scale.

⁶ **2.6 Case study**

⁷ As a demonstration, we evaluated if our proposed method was able to identify treatment effect heterogeneity
⁸ of ACE inhibitors compared to beta blockers using acute myocardial infarction (MI) risk quarter specific effect
⁹ estimates, both on the relative and on the absolute scale. We focused on three efficacy outcomes (acute MI,
¹⁰ hospitalization with heart failure and ischemic or hemorrhagic stroke) and six safety outcomes (hypokalemia,
¹¹ hyperkalemia, hypotension, angioedema, cough and abnormal weight gain). We used data from three US-based
¹² claims databases. The analysis plan was the framework outlined in steps 1 through 5.

¹³ **3 Results**

¹⁴ **3.1 Step 1: General definition of the research aim**

¹⁵ We considered the following research aim: “compare the effect of ACE inhibitors (T) to the effect of beta blockers
¹⁶ (C) in patients with established hypertension with respect to nine outcomes (O_1, \dots, O_9)”. The cohorts are:

- ¹⁷ ▪ Treatment cohort: Patients receiving any drug within the ACE inhibitor class with at least one year of
¹⁸ follow-up before treatment initiation and a recorded hypertension diagnosis within that year.
- ¹⁹ ▪ Comparator cohort: Patients receiving any drug within the beta blocker class with at least one year of
²⁰ follow-up before treatment initiation and a recorded hypertension diagnosis within that year.
- ²¹ ▪ Outcome cohorts: We considered three efficacy and six safety outcome cohorts. These were patients in the
²² database with a diagnosis of: acute MI; hospitalization with heart failure; ischemic or hemorrhagic stroke
²³ (efficacy outcomes); hypokalemia; hyperkalemia; hypotension; angioedema; cough; abnormal weight gain
²⁴ (safety outcomes). Among the safety outcomes we focus on angioedema and cough which are two known
²⁵ adverse events linked to treatment with ACE inhibitors²⁶. Results on the rest of the safety outcomes are
²⁶ included in the supplement.

¹ All cohort definitions were identical to the ones used in the multinational study that provided overall treatment
² effect estimates comparing all anti-hypertensive drug classes with each other¹⁶. More information can be found in
³ the supplementary material.

⁴ 3.2 Step 2: Identification of the databases

⁵ For our demonstration we used data from three US claims databases, namely IBM MarketScan Commercial Claims
⁶ and Encounters (CCAE), IBM MarketScan Medicaid (MDCD), and IBM MarketScan Medicare Supplemental
⁷ Beneficiaries (MDCR). Our analyses included a total of 924 459, 107 046, and 106 905 patients initiating treatment
⁸ with ACE inhibitors and 465 763, 76 546, and 73 213 patients initiating treatment with beta blockers in CCAE,
⁹ MDCD and MDCR respectively (Table 1). Adequate numbers of patients were included in all strata of predicted
¹⁰ acute MI risk (Supplement: Table XX).

¹¹ 3.3 Step 3: Prediction

¹² We internally developed separate prediction models for acute MI in all three databases. The prediction models were
¹³ estimated on the propensity score matched (1:1) subset of the sample, using caliper of 0 · 2 and after excluding
¹⁴ patients having the outcome any time prior to treatment initiation. We chose a 2-year time at risk for patients and
¹⁵ developed the prediction models using LASSO logistic regression with 3-fold cross validation for hyper-parameter
¹⁶ selection.

¹⁷ The models had moderate discriminative performance (internally validated) with no major issues of overfitting to
¹⁸ any cohort except for the case of CCAE, where the derived prediction model performed better in the comparator
¹⁹ cohort (Table 2). We also observed lower performance of the prediction model developed in MDCR compared to
²⁰ the other 2 databases.

²¹ 3.4 Step 4: Estimation

²² We used patient-level predictions to stratify the sample into four acute MI risk quarters. Within risk quarters,
²³ relative effects were estimated using Cox regression and absolute effects were derived from the Kaplan-Meier
²⁴ estimate differences at two years after treatment initiation. To adjust for observed confounding within each
²⁵ risk quarter, we estimated propensity scores using the same approach as step 3 and stratified patients into five
²⁶ propensity score strata. The risk quarter-specific effect estimates were derived by averaging over the estimates
²⁷ within the propensity score fifths.

1 In the lowest acute MI risk quarter of CCAE and MDCC we observed strong separation of the propensity
2 score distributions, therefore, effect estimates derived in these strata are not well-supported (Figure 2). This
3 problematic behavior is also visible in the covariate balance plots comparing standardized mean differences of
4 patient characteristics before and after PS adjustment, where in many cases the commonly accepted bound of
5 0 · 1 is violated (Figure 3). This is more pronounced in the lowest acute MI risk quarter of CCAE, but remains an
6 issue for a small number of covariates in all CCAE risk strata. This diagnostic also fails for the two lower acute
7 MI risk quarters of MDCC. Often the persisting imbalances were linked to pregnancy outcomes, which can be
8 explained by the contraindication of ACE inhibitors in this condition. Analyses in MDCR passed all diagnostics.

9 Finally, the distribution of the estimated relative risks with regard to 30 negative control outcomes indicated
10 unresolved confounding within the lowest acute MI risk quarter of CCAE (Figure 4). Hazard ratios significantly
11 different than 1 (true effect size) were concentrated in the lower right part of Figure 4: panel Q1. This suggests
12 significant negative effects of ACE inhibitors compared to beta blockers on causally unrelated outcomes, pointing
13 at unresolved differences between the two treatment arms. This was not the case in the other risk quarters of
14 CCAE, or in any risk quarter of MDCC and MDCR (Supplement, Figure XX).

15 **3.5 Step 5: Presentation of results**

16 The overall estimated hazard ratios for the main outcomes are presented in Table 3. For hospitalization with acute
17 MI there was an increasing trend in favor ACE inhibitors compared to beta blockers on the relative scale (hazard
18 ratios decreased) with increasing acute MI risk. More specifically, hazard ratios decreased from 1 · 29 (1 · 00 to
19 1 · 68; 95% CI) and 1 · 58 (0 · 78 to 3 · 28; 95% CI) to 0 · 77 (0 · 71 to 0 · 83; 95% CI), 0 · 84 (0 · 76 to 0 · 94;
20 95% CI) in CCAE and MDCC respectively (Figure 4). In MDCR hazard ratios increased from 0 · 93 (0 · 75 to
21 1 · 17; 95% CI) in the lowest MI risk quarter to 1 · 03 (0 · 92 to 1 · 16; 95% CI). Relative treatment effect estimates
22 for hospitalization with heart failure favored ACE inhibitors across all risk strata in all databases. In the case of
23 stroke in CCAE we found quite constant hazard ratios which became weaker in the highest risk quarter patients
24 (0 · 88 with 95% CI from 0 · 80 to 0 · 96). In the other two databases no significant relative treatment effects
25 were observed for stroke. In terms of the safety outcomes, we found an increased ACE inhibitor risk of cough and
26 angioedema on the relative scale across all risk strata. In the case of cough, this effect decreased with increasing
27 risk of acute MI—from 1 · 41 (1 · 37 to 1 · 46; 95% CI), 1 · 28 (1 · 18 to 1 · 38; 95% CI), and 1 · 38 (1 · 29 to
28 1 · 48; 95% CI) to 1 · 30 (1 · 26 to 1 · 34; 95% CI), 1 · 06 (1 · 00 to 1 · 12; 95% CI), and 1 · 11 (1 · 04 to 1 · 18;
29 95% CI) in CCAE, MDCC, and MDCR, respectively.

30 We observed an increasing trend of treatment effect on the absolute scale with increasing acute MI risk in favor of

¹ ACE inhibitors in terms of acute MI in all databases except for MDCR—from -0 · 05% (-0 · 10% to -0 · 005%;
² 95% CI), -0 · 04% (-0 · 14% to 0 · 05%; 95% CI), and 0 · 08% (-0 · 19% to 0 · 34%; 95% CI) in the lowest acute
³ MI risk quarter to 0 · 47% (0 · 31% to 0 · 63%; 95% CI), 0 · 93% (0 · 35% to 1 · 50%; 95% CI), and -0 · 39%
⁴ (-0 · 96% to 0 · 18%; 95% CI) in the highest acute MI risk quarter in CCAE, MDCCD, and MDCR, respectively
⁵ (Figure 5). We found no difference on the absolute scale for stroke across risk strata. Absolute risk differences did
⁶ not favor ACE inhibitors compared to beta blockers in terms of cough, even though this effect again diminished
⁷ with increasing acute MI risk—from -3 · 97% (-4 · 40% to -3 · 54%; 95% CI), -4 · 54% (-6 · 97% to -2 · 12%; 95%
⁸ CI), and -3 · 64% (-4 · 60% to -2 · 68%; 95% CI) in the lowest acute MI risk quarter to -2 · 57% (-3 · 02% to
⁹ -2 · 13%; 95% CI), -0 · 20% (-1 · 58% to 1 · 17%; 95% CI), and -1 · 08% (-2 · 25% to 0 · 08%; 95% CI) in the
¹⁰ highest acute MI risk quarter in CCAE, MDCCD, and MDCR, respectively. In terms of angioedema absolute risk
¹¹ differences were very small due to the rarity of the outcome.
¹² The results of all the analyses performed can be accessed and assessed through a publicly available web application
¹³ (<https://data.ohdsi.org/AceBeta9Outcomes>).

¹⁴ 3.6 Interpretation

¹⁵ The overall benefits of ACE inhibitors compared to beta blockers for acute MI and hospitalization with heart
¹⁶ failure are driven mainly by the higher acute MI risk patients in CCAE and MDCCD, hence the observed increasing
¹⁷ patterns of the absolute treatment benefits. In MDCR we found no significant overall difference on the relative
¹⁸ scale for acute MI and, consequently, no differences in acute MI risk strata were observed on any scale. For heart
¹⁹ failure, MDCR patients at the lower half of acute MI risk had lower absolute benefits compared to the patients
²⁰ at the upper half. Finally, the small overall relative effect for stroke resulted in smaller absolute benefits of ACE
²¹ inhibitors across acute MI risk strata in all databases.
²² For patients at lower acute MI risk, the cough and angioedema risk increase related to treatment with ACE
²³ inhibitors may be important factors to consider for medical decision making, given the small benefits observed for
²⁴ the main outcomes. However, diagnostics failed in lower risk patients within CCAE and MDCR which renders
²⁵ these conclusions less dependable.
²⁶ Note that any conclusions drawn are for demonstration purposes only and should be interpreted under this very
²⁷ limited setting.

4 Discussion

1 The major contribution of our work is the development of a risk-based framework for the assessment of treatment
2 effect heterogeneity in large observational databases. This fills a gap identified in the literature after the development
3 of guidelines for performing such analyses in the RCT setting^{11,12}. As an additional contribution we developed
4 the software for implementing this framework in practice and made it publicly available. We made our software
5 compatible to databases mapped to OMOP-CDM which allows researchers to easily implement our framework
6 in a global network of healthcare databases. In our case study we demonstrated the use of our framework for
7 the evaluation of treatment effect heterogeneity ACE inhibitors compared to beta blockers on three efficacy and
8 six safety outcomes. We propose that this framework is implemented any time treatment effect estimation in
9 high-dimensional observational data is undertaken.

10 In recent years several methods for the analysis of treatment effect heterogeneity have been developed in the RCT
11 setting²⁷. However, low power and restricted prior knowledge on the mechanisms of variation in treatment effect
12 are often inherent in RCTs, which are usually adequately powered only for the analysis of the primary outcome.
13 Observational databases contain a large amount of information on treatment assignment and outcomes of interest,
14 while also capturing key patient characteristics. They contain readily available data on patient subpopulations of
15 interest for which no RCT has focused before either due to logistical or ethical reasons. However, observational
16 databases can be susceptible to biases, poorly measured outcomes and missingness, which may obscure true HTE or
17 falsely introduce it when there is none²⁸. Therefore, inferences on both overall treatment effect estimates and HTE
18 need to rely on strong, often unverifiable, assumptions, despite the advancements and guidance on best practices.
19 However, well-designed observational studies on average replicate RCT results, even though often differences
20 in magnitude may occur³³. Our framework is in line with the recently suggested paradigm of high-throughput
21 observational studies using consistent and standardized methods for improving reproducibility in observational
22 research²⁵.

23 Our framework highlights the scale dependency of HTE and how it relates to baseline risk. Treatment effect is
24 mathematically determined by baseline risk, if we assume a constant non-zero effect size³⁴. Patients with low
25 baseline risk can only experience minimal benefits, before their risk is reduced to zero. In contrast, high risk patients
26 are capable of displaying much higher absolute benefits. This becomes evident when evaluating the effects of ACE
27 inhibitors on cough and angioedema, compared to treatment with beta blockers. Despite the small relative cough
28 risk increase of ACE inhibitors, the large baseline cough risk resulted in larger absolute risk differences, compared
29 to the other considered outcomes. Conversely, in the case of angioedema, the substantial relative risk increase
30 with ACE inhibitors only translated in a small absolute risk increase due to the quite low baseline angioedema risk.

1 The application of our framework in the case study is for demonstration purposes and there are several limitations
2 to its conclusions. First, death could be a competing risk. We could expand our framework in the future to
3 potentially support subdistribution hazard ratios and cumulative incidence reductions. Second, we only used the
4 databases readily available to us and not all the available databases mapped to OMOP-CDM. Therefore, the
5 generalizability of our results still needs to be explored in future studies. These studies should also address the
6 particular aspects of the databases at hand, such as their sampling frame, the completeness of the data they
7 capture and many others that were not assessed in our demonstration. Third, we did not correct for multiplicity
8 when presenting the results. We are interested in presenting trends in the data and not detecting the specific
9 subgroups within which a non-null treatment effect is detected. The implementation of our framework, however,
10 generates all the relevant output required for a researcher to correct for multiple testing, if that is required.

11 In conclusion, the case study demonstrates the feasibility of our framework for risk-based assessment of treatment
12 effect heterogeneity in large observational data. It is easily applicable and highly informative whenever treatment
13 effect estimation in high-dimensional observational data is of interest.

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6 Tables and figures

Table 1: Number of patients, person years and events within quarters of predicted risk for acute MI for the three efficacy outcomes of the study (acute MI, hospitalization with heart failure and ischemic or hemorrhagic stroke) across the three databases.

Outcome	ACE inhibitors			Beta blockers		
	Patients	Person years	Outcomes	Patients	Person years	Outcomes
CCAE						
acute myocardial infarction	924 196	1 327 973	4 102	457 375	648 612	2 492
hospitalization with heart failure	924 459	1 328 430	3 764	465 763	660 580	3 711
stroke	917 501	1 319 236	3 741	464 989	659 472	2 454
MDCD						
acute myocardial infarction	107 046	162 590	1 448	76 307	112 767	1 361
hospitalization with heart failure	105 544	160 237	2 819	74 649	110 455	3 005
stroke	104 953	159 344	1 799	76 546	113 048	1 623
MDCR						
acute myocardial infarction	106 905	163 260	1 764	72 733	110 821	1 480
hospitalization with heart failure	106 191	162 258	3 004	73 182	111 710	3 592
stroke	103 531	158 369	2 323	73 213	111 613	2 241

Table 2: Discriminative ability (c-statistic) of the derived prediction models for acute myocardial infarction in the matched set (development set), the treatment cohort, the comparator cohort and the entire population in CCAE, MDCC and MDCR.

Population	CCAE	MDCD	MDCR
Matched	0 · 73 (0 · 72, 0 · 74)	0 · 78 (0 · 77, 0 · 79)	0 · 66 (0 · 65, 0 · 68)
Treatment	0 · 69 (0 · 68, 0 · 70)	0 · 76 (0 · 75, 0 · 77)	0 · 65 (0 · 63, 0 · 66)
Comparator	0 · 77 (0 · 76, 0 · 78)	0 · 82 (0 · 81, 0 · 82)	0 · 64 (0 · 63, 0 · 66)
Entire population	0 · 72 (0 · 71, 0 · 73)	0 · 79 (0 · 78, 0 · 80)	0 · 65 (0 · 64, 0 · 66)

Table 3: Overall hazard ratios.

Outcome	CCAE	MDCD	MDCR
acute myocardial infarction	0 · 83 (0 · 79, 0 · 88)	0 · 87 (0 · 80, 0 · 96)	1 · 02 (0 · 94, 1 · 10)
hospitalization with heart failure	0 · 66 (0 · 62, 0 · 69)	0 · 86 (0 · 81, 0 · 92)	0 · 85 (0 · 80, 0 · 90)
stroke	0 · 88 (0 · 83, 0 · 93)	0 · 97 (0 · 90, 1 · 05)	0 · 91 (0 · 85, 0 · 97)

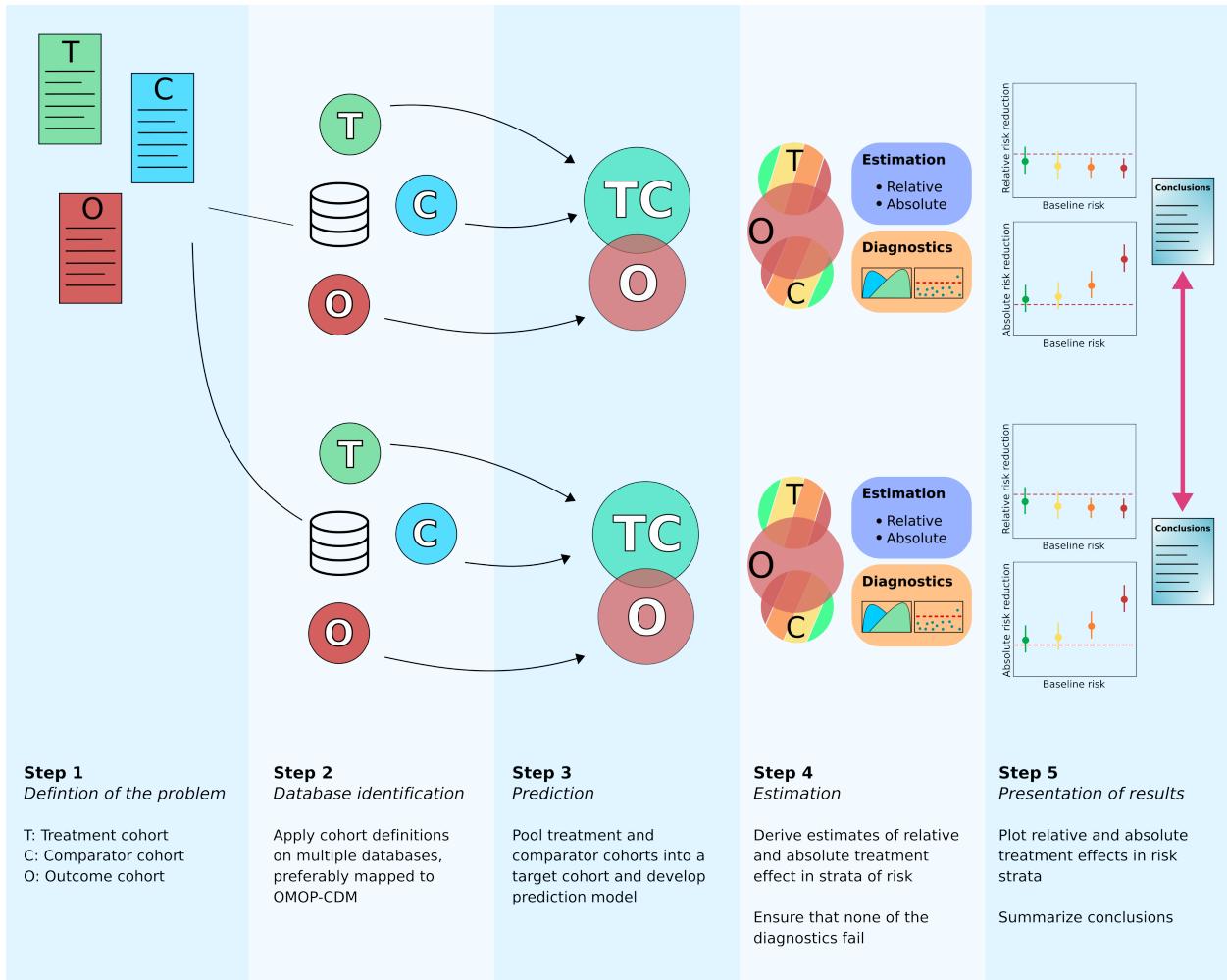


Figure 1: Illustration of how the framework is applied on two observational databases, preferably mapped to OMOP-CDM.

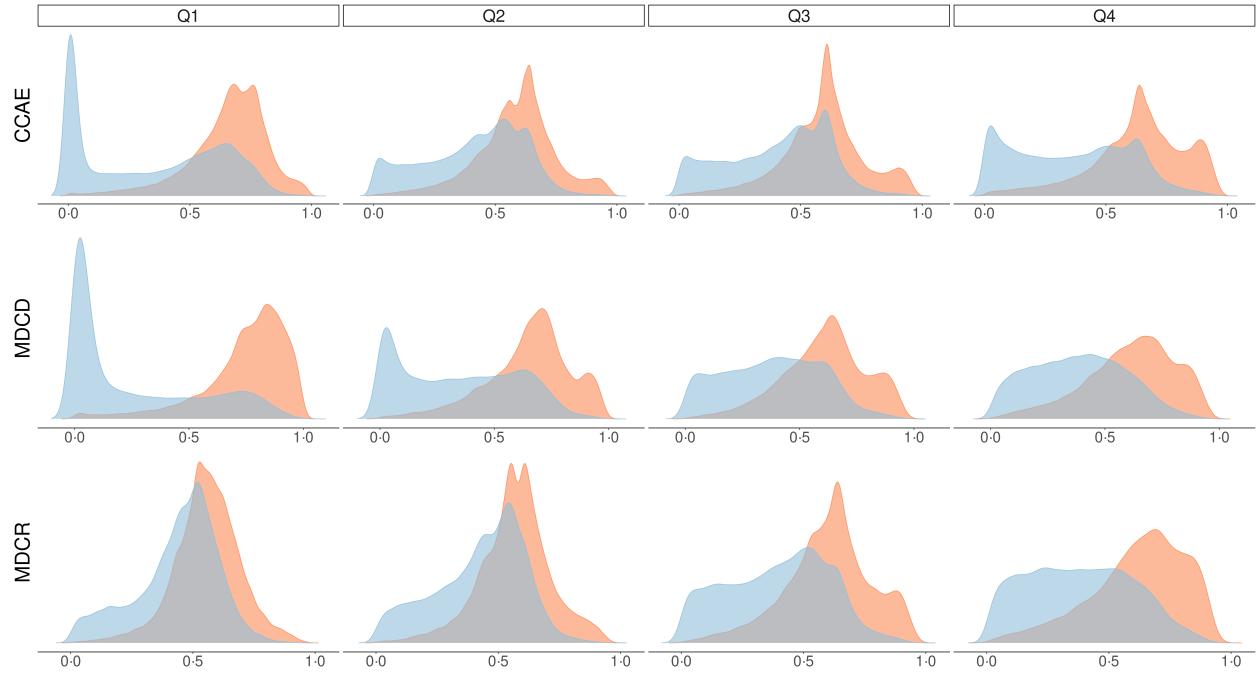


Figure 2: Preference score distributions for the evaluation of heterogeneity of the effect of ACE inhibitors compared to beta blockers on acute MI based on quarters of predicted acute MI risk. The preference score is a transformation of the propensity score that adjusts for prevalence differences between populations.

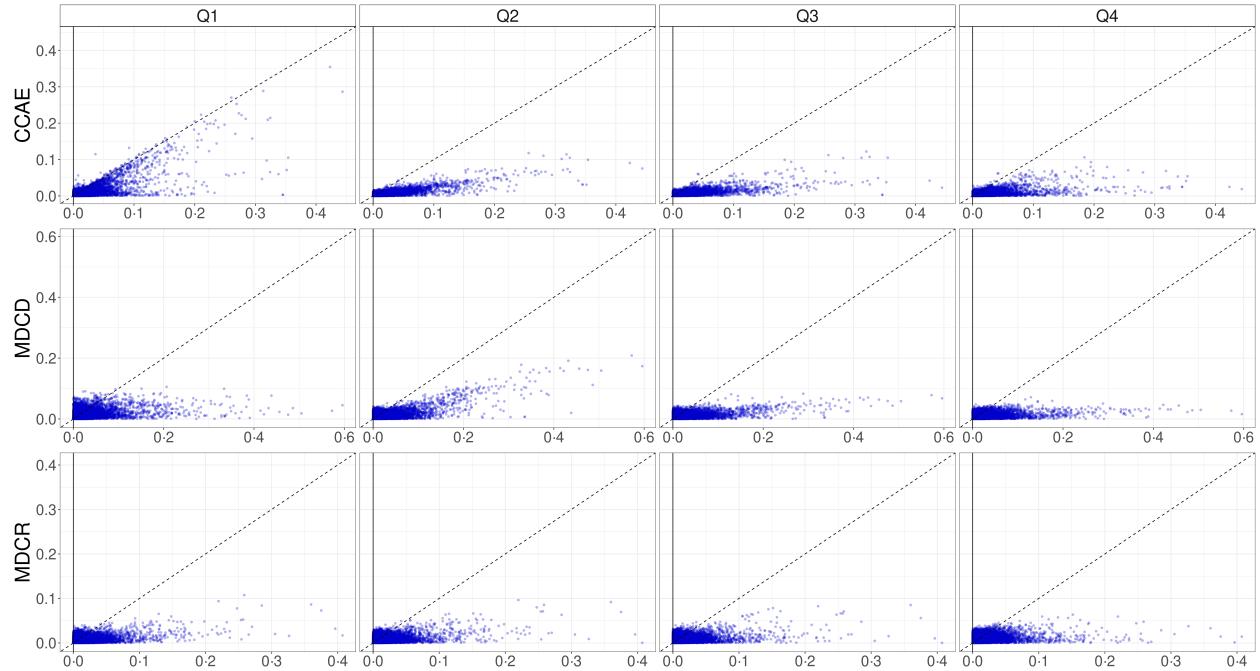


Figure 3: Patient characteristic balance for ACE inhibitors and beta blockers before and after stratification on the propensity scores. Each dot represents the standardized difference of means for a single covariate before (x-axis) and after (y-axis) stratification.

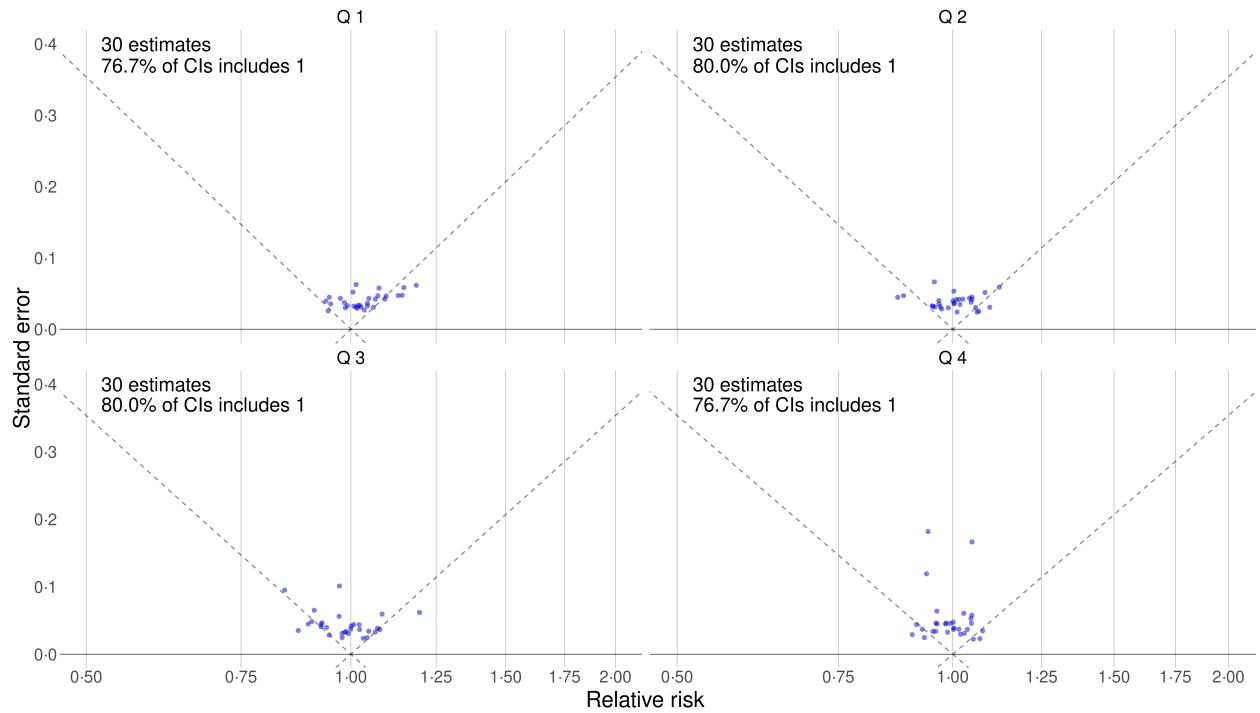


Figure 4: Systematic error. Effect size estimates for the negative controls (true hazard ratio = 1). Estimates below the diagonal dashed lines are statistically significant ($\alpha = 0.05$) different from the true effect size. A well-calibrated estimator should have the true effect size within the 95 percent confidence interval 95 percent of times.

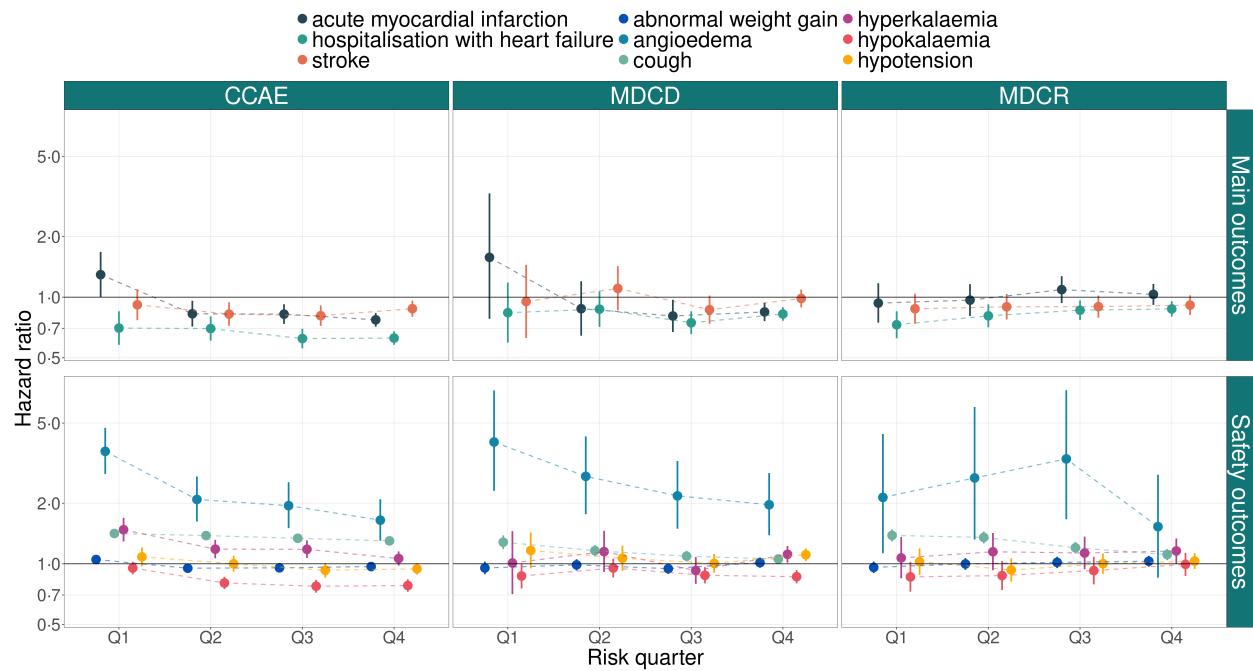


Figure 5: Overview of heterogeneity of ACE inhibitors treatment on the relative scale (hazard ratios) within strata of predicted risk of acute MI. Values below 1 favor ACE inhibitors, while values above 1 favor beta blockers.

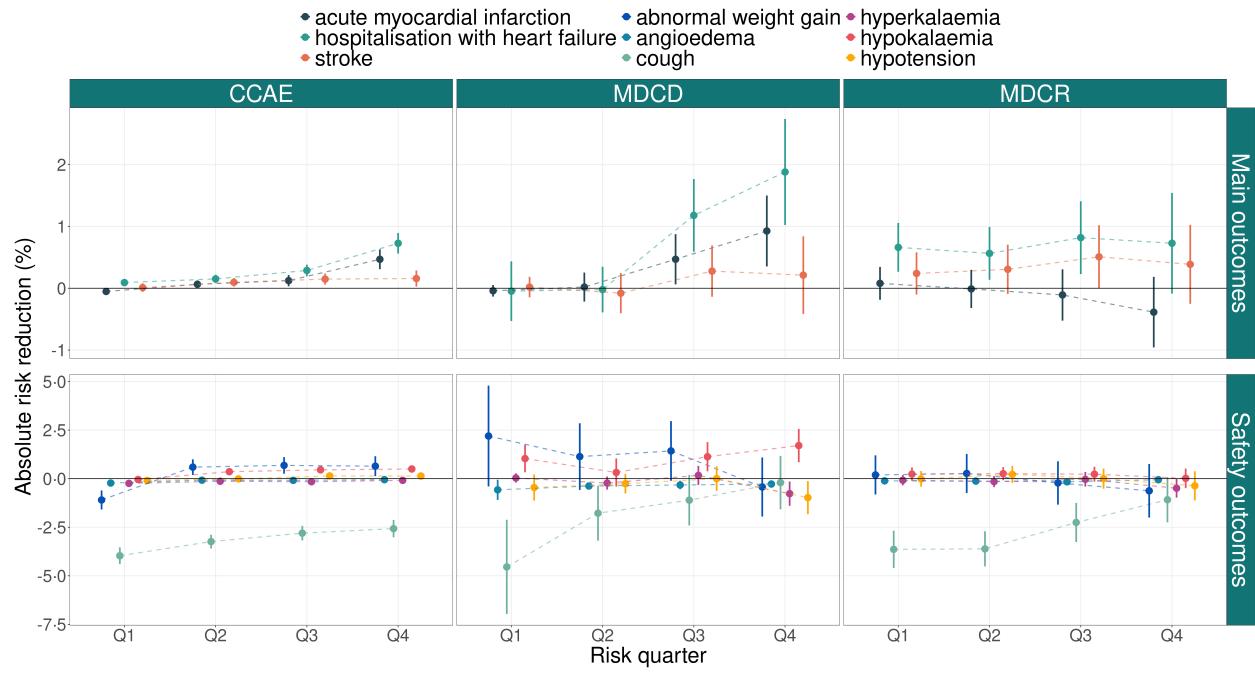


Figure 6: Overview of heterogeneity of ACE inhibitors treatment on the absolute scale within strata of predicted risk of acute MI. Estimates of absolute treatment effect are derived as the difference in Kaplan-Meier estimates at two years after inclusion. Values above 0 favor ACE inhibitors, while values below 0 favor beta blockers.