
IMPROVEMENT-FOCUSED CAUSAL RECOURSE (ICR)

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ABSTRACT

Algorithmic recourse recommendations, such as Karimi et al.’s (2021) causal recourse (CR), inform stakeholders of how to act to revert unfavorable decisions. However, there are actions that lead to acceptance (i.e., revert the model’s decision) but do not lead to improvement (i.e., may not revert the underlying real-world state). To recommend such actions is to recommend fooling the predictor. We introduce a novel method, Improvement-Focused Causal Recourse (ICR), which involves a conceptual shift: Firstly, we require ICR recommendations to guide towards improvement. Secondly, we do not tailor the recommendations to be accepted by a specific predictor. Instead, we leverage causal knowledge to design decision systems that predict accurately pre- and post-recourse. As a result, improvement guarantees translate into acceptance guarantees. We demonstrate that given correct causal knowledge ICRguides towards both acceptance and improvement.

Keywords algorithmic recourse · gaming · causal inference · interpretable machine learning · robustness

1 Introduction

Predictive systems are increasingly deployed for high-stakes decisions, for instance in hiring [Raghavan et al., 2020], judicial systems [Zeng et al., 2017], or when distributing medical resources [Obermeyer and Mullainathan, 2019]. A range of work [Wachter et al., 2017, Ustun et al., 2019, Karimi et al., 2021] develops tools that offer individuals possibilities for so-called algorithmic recourse (i.e. actions that revert unfavorable decisions). Joining previous work in the field, we distinguish between reverting the model’s prediction \hat{Y} (acceptance) and reverting the underlying real-world state Y (improvement) and argue that recourse should lead to acceptance *and* improvement [Ustun et al., 2019, Barocas et al., 2020]. Existing methods, such as counterfactual explanations (CE; Wachter et al. [2017]) or causal recourse (CR; Karimi et al. [2021]), ignore the underlying real-world state and only optimize for acceptance. Since ML models are not designed to predict accurately in interventional environments (i.e. environments where actions have changed the data distribution), acceptance does not necessarily imply improvement.

Let us consider a simple motivational example. The goal is to predict whether hospital visitors without recent test certificate are infected with Covid in order to restrict access to tested and low-risk individuals. In the example, the model’s *prediction* \hat{Y} represents whether someone is classified to be infected, whereas the *prediction target* Y represents whether someone is actually infected. Target and prediction differ in how they are affected by actions. E.g., intervening on the *symptoms* may change the diagnosis \hat{Y} , but will not affect whether someone is infected (Y).

Both counterfactual explanations (CE) and causal recourse (CR) only target \hat{Y} (Figure 1). Therefore, CE and CR may suggest to alter the *symptoms* (e.g., by taking cough drops) and thereby may recommend to *game* the predictor: Although the intervention leads to acceptance the actual Covid risk Y is not improved.¹

One may argue that this is an issue of the prediction model and may adapt the predictor strategically to make gaming less lucrative than improvement [Miller et al., 2020]. In our example, the model’s reliance on the symptom state would need to be reduced. However, such strategic adaptations may come at the cost of predictive performance since gameable variables, like the symptom state, can be highly predictive [Shavit et al., 2020]. Thus, we tackle the problem by adjusting the explanation.

¹In E.1, the case is formally demonstrated.

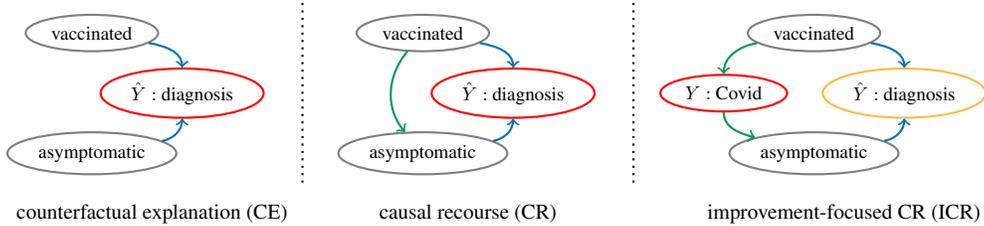


Figure 1: Directed Acyclic Graph (DAG) illustrating the perspective on model and data taken by counterfactual explanations (CE, left) and causal recourse (CR, center) in contrast to improvement-focused recourse (ICR, right). Blue edges represent the causal links induced by the prediction model, green edges the real-world causal links, gray nodes the covariates, and the red (yellow) node the primary (secondary) recourse target. CR respects the causal relationships but only between input features. ICR is the only approach that takes the target Y into account. While CE and CR aim to revert the prediction \hat{Y} , ICR aims to revert the target Y .

Contributions We present improvement-focused causal recourse (ICR), the first recourse method that targets improvement instead of acceptance. Since estimating the effects of actions is a causal problem, causal knowledge is required. More specifically, we show how to exploit either knowledge of the structural causal model (SCMs) or the causal graph to guide towards improvement (Section 5). On a conceptual level we argue that the individual’s improvement options should not be limited by an acceptance constraint (Section 4). In order to nevertheless yield acceptance, we show how to exploit said causal knowledge to design post-recourse decision systems that in expectation recognize improvement (Section 6), such that improvement guarantees translate into acceptance guarantees (Section 7). On synthetic and semi-synthetic data, we demonstrate that ICR, in contrast to existing approaches, leads to improvement and acceptance (Section 8).

2 Related Work

Contrastive Explanations Contrastive explanations explain decisions by contrasting them with alternative decision scenarios [Karimi et al., 2020a, Stepin et al., 2021]; a well known example are counterfactual explanations (CE) that highlight the minimal feature changes required to revert the decision of a predictor $\hat{f}(x)$ [Wachter et al., 2017, Dandl et al., 2020]. However, CEs are ignorant of causal dependencies in the data and therefore in general fail to guide action [Karimi et al., 2021]. In contrast, the causal recourse (CR) framework by Karimi et al. [2022] takes the causal dependencies between covariates into account: More specifically, Karimi et al. [2022] use structural causal models or causal graphs to guide individuals towards acceptance.² The importance of improvement was discussed before [Ustun et al., 2019, Barocas et al., 2020], but as of now no improvement-focused recourse method was proposed.

Strategic Classification The related field of strategic modeling investigates how the prediction mechanism incentivizes rational agents [Hardt et al., 2016, Tsirtsis and Gomez Rodriguez, 2020]. A range of work [Bechavod et al., 2020, Chen et al., 2020, Miller et al., 2020] thereby distinguishes models that incentivize *gaming* (i.e., interventions that affect the prediction \hat{Y} but not the underlying target Y in the desired way) and *improvement* (i.e., actions that also yield the desired change in Y). Strategic modeling is concerned with adapting the model, where except for special cases the following three goals are in conflict: incentivizing improvement, predictive accuracy, and retrieving the true underlying mechanism [Shavit et al., 2020].

Robust algorithmic recourse The robustness of CEs and CR has been investigated before [Rawal et al., 2021, Pawelczyk et al., 2020, Upadhyay et al., 2021, Dominguez-Olmedo et al., 2021, Pawelczyk et al., 2022], yet only with respect to generic shifts of model and data. Only Pawelczyk et al. [2020] investigate the robustness regarding refits on the same data. They find that on-the-manifold CEs are more robust than standard CEs. In contrast, we empirically compare the robustness of CE, CR and ICR with respect to refits on the same data.

3 Background and Notation

Prediction model We assume binary probabilistic predictors and cross-entropy loss, such that the optimal score function $h^*(x)$ models the conditional probability $P(\hat{Y} = 1|X = x)$, which we abbreviate as $p(y|x)$. We denote the

²For the interested reader, we formally introduce CR in our notation in A.4.

estimated score function as $\hat{h}(x)$, which can be transformed into the binary decision function $\hat{f}(x) := [\hat{h}(x) \geq t]$ via the decision threshold t .

Causal data model We model the data generating process using a structural causal model (SCM) $\mathcal{M} \in \Pi$ [Pearl, 2009, Peters et al., 2017]. The model $\mathcal{M} = \langle X, U, \mathbb{F} \rangle$ consists of the endogenous variables $X \in \mathcal{X}$, the mutually independent exogenous variables $U \in \mathcal{U}$, and structural equations $\mathbb{F} : \mathcal{U} \rightarrow \mathcal{X}$. Each structural equation f_j specifies how X_j is determined by its endogenous causes and the corresponding exogenous variable U_j . The SCM entails a directed graph \mathcal{G} , where variables are connected to their direct effects via a directed edge.

The index set of endogenous variables is denoted as D . The parent indexes of node j are referred to as $pa(j)$ and the children indexes as $ch(j)$. We refer to the respective variables as $X_{pa(j)}$. We write $X_{pa(j)}$ to denote all parents excluding Y and $(X, Y)_{pa(j)}$ to denote all parents including Y . All ascendant indexes of a set S are denoted as $asc(S)$, its complement as $nasc(S)$, all descendant indexes as $d(S)$, and its complement as $nd(S)$.

SCMs allow to answer causal questions. This means that they cannot only be used to describe (conditional) distributions (observation, rung 1 on Pearl’s ladder of causation [Pearl, 2009]), but can also be used to predict the (average) effect of actions $do(x)$ (intervention, rung 2) and imagine the results of alternative actions in light of factual observation $(x, y)^F$ (counterfactuals, rung 3).

As such, we model actions as structural interventions $a : \Pi \rightarrow \Pi$, which can be constructed as $do(a) = do(\{X_i := \theta_i\}_{i \in I})$, where I is the index set of features to be intervened upon. A model of the interventional distribution can be obtained by fixing the intervened upon values to θ_I (e.g. by replacing the structural equation $f_I := \theta_I$). Counterfactuals can be computed in three steps [Pearl, 2009]: First, the factual distribution of exogenous variables U given the factual observation of the endogenous variables x^F is inferred (*abduction*) (i.e., $P(U_j | X^F)$). Second, the structural interventions corresponding to $do(a)$ are performed (*action*). Finally, we can sample from the counterfactual distribution $P(X^{SCF} | X = x^F, do(a))$ using the abducted noise and the intervened-upon structural equations (*prediction*).

4 The Two Tales of Contrastive Explanations

In the introduction we have demonstrated that CE and CR may suggest to game the predictor (i.e. guide towards acceptance without improvement). To tackle the issue, we will introduce a new explanation technique called improvement-focused causal recourse (ICR) in Section 5.

In this section we lay the conceptual justification for our method. More specifically, we argue that for recourse the acceptance constraint of CR should be *replaced* by an improvement constraint. Therefore, we first recall that a multitude of goals may be pursued with contrastive explanations [Wachter et al., 2017] and separate two purposes of contrastive explanations: *contestability of algorithmic decisions* and *actionable recourse*. We then argue that improvement is an essential requirement for recourse and that the individual’s options for improvement should not be limited by acceptance constraints.

Contestability and recourse are distinct goals. *Contestability* is concerned with the question of whether the algorithmic decision is correct according to common sense, moral or legal standards. Explanations may help model authorities to detect violations of such standards or enable explainees to contest unfavorable decisions [Wachter et al., 2017, Freiesleben, 2021]. Explanations that aim to enable contestability must reflect the model’s rationale for an algorithmic decision. *Recourse recommendations* on the other hand need to satisfy various constraints unrelated to the model, such as causal links between variables [Karimi et al., 2021] or their actionability [Ustun et al., 2019]. Consequently, explanations geared to contest are more complete and true to the model while recourse recommendations are more selective and true to the underlying process.³ We believe that the selectivity and reliance of recourse recommendations on factors besides the model itself is not a limitation but an indispensable condition for making explanations more relevant to the explainee.

In the context of recourse, improvement is desirable for model authority and explainee. We consider improvement to be an important normative requirement for recourse, both with respect to explainee and model authority. Valuable recourse recommendations enable explainees to plan and act; thus, such recommendations must either provide indefinite validity or a clear expiration date [Wachter et al., 2017, Barocas et al., 2020, Venkatasubramanian and Alfano, 2020]. Problematically, when model authorities give guarantees for non-improving recourse, this constitutes a binding commitment to misclassification. However, if model authorities do not provide recourse guarantees over time, this diminishes the value of recourse recommendations to explainees. They might invest effort into non-improving actions

³We do not claim that recourse and contestability always diverge, we only describe a difference in focus. If contesting is successful it may even provide an alternative route towards recourse.

that ultimately do not even lead to acceptance because the classifier changed.⁴ In contrast, improvement-focused recourse is honored by any accurate classifier. We conclude that, given these advantages for both model authority and explainee, recourse recommendations should help to improve the underlying target Y .⁵

Improvement should come first, acceptance second. Taken that we constrain the optimization on improvement, how to guarantee acceptance remains an open question. One approach would be to constrain the optimization on both improvement and acceptance. However, a restriction on acceptance is either redundant or, from our moral standpoint, questionable: If improvement already implies acceptance, the constraint is redundant. In the remaining cases, we can predict improvement with the available causal knowledge but would withhold these (potentially less costly) improvement options because of the limitations of the observational predictor. To ensure that acceptance ensues improvement, we instead suggest to exploit the assumed causal knowledge for accurate post-recourse prediction (Section 6), such that acceptance guarantees can be made (Section 7).

5 Improvement-Focused Causal Recourse (ICR)

We continue with the formal introduction of ICR, an explanation technique that targets improvement ($Y = 1$) instead of acceptance ($\hat{Y} = 1$). Therefore we first define the improvement confidence γ , which can be optimized to yield ICR. Like previous work in the field [Karimi et al., 2020b], we distinguish two settings: In the first setting, knowledge of the SCM can be assumed, such that we can leverage structural counterfactuals (rung 3 on Pearl’s ladder of causation) to introduce the individualized improvement confidence γ^{ind} . In the second setting only the causal graph is known, which we exploit to propose the subpopulation-based improvement confidence γ^{sub} (rung 2).

Individualized improvement confidence For the individualized improvement confidence γ^{ind} we exploit knowledge of a SCM. SCMs can be used to answer counterfactual questions (rung 3). In contrast to rung-2-predictions, counterfactuals are tailored to the individual and their situation [Pearl, 2009]: They ask what would have been if one had acted differently and thereby exploit the individual’s factual observation. Given unchanged circumstances, counterfactuals can be seen as individualized causal effect predictions.

In contrast to existing SCM-based recourse techniques [Karimi et al., 2022] we include both the prediction \hat{Y} and the target variable Y as separate variables in the SCM. As a result, the SCM can be used not only to model the individualized probability of acceptance, but also the individualized probability of improvement.

Definition 1 (Individualized improvement confidence). *For pre-recourse observation x^{pre} and action a we define the individualized improvement confidence as*

$$\gamma^{ind}(a) = \gamma(a, x^{pre}) := P(Y^{post} = 1 | do(a), x^{pre}).$$

Since the pre-recourse (factual) target Y cannot be observed, standard counterfactual prediction cannot be applied directly. However, we can regard the distribution as a mixture with two components, one for each possible state of Y . We can estimate the mixing weights using h^* and each component using standard counterfactual prediction. Details including pseudocode are provided in B.1.

Subpopulation-based improvement confidence For the estimation of the individualized improvement confidence γ^{ind} knowledge of the SCM is required. If the SCM is not specified, but the causal graph is known instead and there are no unobserved confounders (causal sufficiency), we can still estimate the effect of interventions (rung 2).

In contrast to counterfactual distributions (rung 3), interventional distributions describe the whole population and therefore provide limited insight into the effects of actions on specific individuals. Building on Karimi et al. [2020b], we thus narrow the population down to a subpopulation of similar individuals, for which we then estimate the subpopulation-based causal effect. More specifically, we consider individuals to belong to the same subgroup if the variables that are not affected by the intervention take the same values. For action a , we define the subgroup characteristics as $G_a := nd(I_a)$ (i.e., the non-descendants of the intervened-upon variables in the causal graph).⁶ More formally, we define the subpopulation-based improvement confidence γ^{sub} as the probability of Y taking the favorable outcome in the subgroup of similar individuals (Definition 2).

⁴For instance, in the introductory example, an intervention on the symptom state would only be honored by a refit of the model on pre- and post-recourse data for the small percentage of individuals who were already vaccinated, as documented in more detail in E.1. Also, gaming actions may not be robust concerning model multiplicity, as seen in the experiments (Section 8).

⁵We do not claim that gaming is necessarily bad; it may be justified when predictors perform morally questionable tasks.

⁶The estimand resembles the conditional treatment effect with G_a being effect modifiers [Hernán MA, 2020].

Definition 2 (Subpopulation-based improvement confidence). *Let a be an action that potentially affects Y , i.e. $I_a \cap asc(Y) \neq \emptyset$.⁷ Then we define the subpopulation-based improvement confidence as*

$$\gamma^{sub}(a) = \gamma(a, x_{G_a}^{pre}) := P(Y^{post} = 1 | do(a), x_{G_a}^{pre}).$$

The set G_a is chosen for practical reasons. In order to make the estimation more accurate, we would like to condition on as many characteristics as possible. However, without access to the SCM, one can only identify interventional distributions for subgroups of the population by conditioning on their (unobserved) post-intervention characteristics (but not by conditioning on their pre-intervention characteristics) [Pearl, 2009, Glymour et al., 2016]. If we were to select a subgroup from a post-recourse distribution by conditioning on pre-recourse characteristics that are affected by a (e.g. strong pre-recourse symptoms), we yield a group that the individual may not be part of (e.g. people with strong post-recourse symptoms). In contrast, for X_{G_a} pre- and post-intervention values coincide, such that we can estimate γ^{sub} : Assuming causal sufficiency, the standard procedure to sample interventional distributions can be applied, only that additionally $X_{G_a}^{post} := x_{G_a}^{pre}$. Based on the sample γ^{sub} can be estimated (as detailed in B.3).

The estimation of γ^{sub} does not require knowledge of the SCM, but is less accurate than γ^{ind} . In the introductory example, for the action *get vaccinated* the set of subgroup-characteristics G_a is empty. As such, γ^{sub} is concerned with the effect of a vaccination over the whole population. If we were to observe *zip code*, a variable that is not affected by *vaccination*, γ^{sub} would indicate the effect of vaccination for subjects that share the explainee’s *zip code*. In contrast, γ^{ind} also takes the explainee’s *symptom state* into account.

Optimization problem To generate ICR recommendations, we can optimize Equation 1. We aim to find actions that meet a user-specified improvement target confidence $\bar{\gamma}$ with minimal cost for the recourse seeking individual. The cost function $cost(a, x^{pre})$ captures the effort the individual requires to perform action a [Karimi et al., 2020b]. As for CE or CR, the optimization problem for ICR is computationally challenging (B.4). It can be seen as a two-level problem, where on the first level the intervention targets I_a , and on the second level the corresponding intervention values θ_a are optimized [Karimi et al., 2020b]. Since we target improvement, we can restrict I_a to causes of Y . Following Dandl et al. [2020], we use the genetic algorithm NSGA-II [Deb et al., 2002] for optimization.

$$\operatorname{argmin}_{a=do(X_I=\theta)} \quad cost(a, x^{pre}) \quad \text{s.t.} \quad \gamma(a) \geq \bar{\gamma}. \quad (1)$$

6 Accurate Post-Recourse Prediction

Recourse recommendations should not only lead to improvement Y but also revert the decision \hat{Y} . Whether acceptance guarantees naturally ensue from γ depends on the ability of the predictor to recognize improvements. As follows, we demonstrate how the assumed causal knowledge can be exploited to design accurate post-recourse predictors. We find that an individualized post-recourse predictor is required to translate γ^{ind} into an individualized acceptance guarantee, but curiously that the observational predictor is sufficient in supopulation-based settings.

Individualized post-recourse prediction If we were to use the optimal pre-recourse observational predictor h^* for post-recourse prediction, there would be an imbalance in predictive capability between ML model and individualized ICR: ICR individualizes its predictions using x^{pre} and the SCM. This knowledge is not accessible by the predictor h^* , which only makes use of x^{post} . As such, improvement that was accurately predicted by ICR is not necessarily recognized by h^* and γ^{ind} cannot be directly translated into an acceptance bound. We demonstrate the issue at an Example in E.3.⁸

In order to settle the imbalance between ICR and the predictor, we suggest to leverage the SCM not only when generating individualized ICR recommendations but also when predicting post-recourse, such that the predictor is at least as accurate as γ^{ind} . More formally, we suggest to estimate the post-recourse distribution of Y conditional on x^{pre} , $do(a)$, and the post-recourse observation $x^{post,a}$ (Definition 3). This post-recourse prediction resembles the counterfactual distribution, except that we additionally take the factual post-recourse observation of the covariates into account.

⁷If a cannot affect Y , we can predict $P(Y|x^{pre}, do(a)) = P(Y|x^{pre})$ using the optimal observational predictor h^* .

⁸One may also argue that standard predictive models are not suitable since optimality of the predictor in the pre-recourse distribution does not necessarily imply optimality in interventional environments (as Example 1, E.1 demonstrates). We can refute this criticism using Proposition 3, where we learn that \hat{h}^* is stable with respect to ICR actions.

Definition 3 (Individualized post-recourse predictor). *We define the individualized post-recourse predictor as*

$$h^{*,ind}(x^{post}) = P(Y^{post} = 1 | x^{post}, x^{pre}, do(a))$$

For SCMs with invertible equations, $h^{*,ind}$ can be estimated using a closed form solution. Otherwise we can sample from the counterfactual post-recourse distribution $p(y^{post}, x^{post} | x^{pre}, do(a))$ (as we did for the estimation of γ^{ind}), select the samples that conform with x^{post} and compute the proportion of favorable outcomes (details in B.2).

For the individualized post-recourse predictor, improvement probability and prediction are closely linked (Proposition 1). More specifically, the expected post-recourse prediction $h^{*,ind}$ is equal to the individualized improvement probability $\gamma(x^{pre}, a)$. We will exploit Proposition 1 in Section 7, where we derive acceptance guarantees for ICR.

Proposition 1. *The expected individualized post-recourse score is equal to the individualized improvement probability $\gamma^{ind}(x^{pre}, a) := P(Y^{post} = 1 | x^{pre}, do(a))$, i.e.*

$$E[\hat{h}^{*,ind}(x^{post}) | x^{pre}, do(a)] = \gamma^{ind}(a).$$

Subpopulation-based post-recourse prediction Curiously we find that for ICR actions a the optimal observational pre-recourse predictor h^* remains accurate: in the subpopulation of similar individuals the expected post-recourse prediction corresponds to the improvement probability $\gamma^{sub}(a)$ (Proposition 3). This allows us to derive acceptance guarantees for h^* in Section 7.

This result is in contrast to the negative results for CR, where actions may not affect prediction and the underlying target coherently, such that the predictive performance deteriorates (as demonstrated in the introduction, and more formally in E.1). The key difference to CR is that ICR actions exclusively intervene on causes of Y : Interventions on non-causal variables may lead to a shift in the conditional distribution $P(Y | X_S)$ (where $S \subseteq D$ is any set of variables that allows for optimal prediction). In contrast, given causal sufficiency, the conditional $P(Y | X_S)$ is stable to interventions on causes of Y .

Proposition 2. *Given nonzero cost for all interventions, ICR exclusively suggests actions on causes of Y . Assuming causal sufficiency, for optimal models the conditional distribution of Y given the variables X_S that the model uses (i.e. $P(Y | X_S)$) is stable w.r.t interventions on causes. Therefore, optimal predictors are intervention stable w.r.t ICR actions.*

Proposition 3. *Given causal sufficiency and positivity⁹, for interventions on causes the expected subgroup-wide optimal score h^* is equal to the subgroup-wide improvement probability $\gamma^{sub}(a) := P(Y^{post} = 1 | do(a), x_{G_a}^{pre})$, i.e.*

$$E[\hat{h}^*(x^{post}) | x_{G_a}^{pre}, do(a)] = \gamma^{sub}(a).$$

Link between CR and ICR: Proposition 2 has further interesting consequences. For CR actions a that only intervene on causes of Y and that are guaranteed to yield a predicted score ζ in the subpopulation, we can infer that $\gamma^{sub}(a) \geq \zeta$. For instance, if acceptance with respect to a 0.5 decision threshold can be guaranteed, that implies improvement with at least 50% probability. As such, in subpopulation-based settings (1) improvement guarantees can be made for CR if only interventions on causes are lucrative, and (2) CR can be adapted to also guide towards improvement by a restricting actions to intervene on causes.

7 Acceptance Guarantees

For the presented accurate post-recourse predictors, improvement guarantees translate into acceptance guarantees (Proposition 4). The reason is that the post-recourse prediction is linked to γ (Propositions 1 and 3).

Proposition 4. *Let g be a predictor with $E[g(x^{post}) | x_S^{pre}, do(a)] = \gamma(x_S^{pre}, a)$. Then for a decision threshold t the post-recourse acceptance probability $\eta(t; x_S^{pre}, a) := P(g(x^{post}) > t | x_S^{pre}, do(a))$ is lower bounded by the respective improvement probability:*

$$\eta(t; x_S^{pre}, a, g) \geq \frac{\gamma(x_S^{pre}, a) - t}{1 - t}.$$

Proof (sketch): We decompose the expected prediction (γ) into true positive rate (TPR), false negative rate (FNR) and acceptance rate. By bounding TPR and FNR we yield the presented acceptance bound. The proof is provided in D.4.

⁹Positivity ensures that the post-recourse observation lies within the observational support [Neal, 2020], where the model was trained (i.e., $p^{pre}(x^{post}) > 0$).

Using Proposition 4, we can tune confidence γ and the model’s decision threshold to yield a desired acceptance rate. For instance, we can guarantee acceptance with (subgroup-wide) probability $\eta \geq 0.9$ given $\gamma = 0.95$ and a global decision threshold $t = 0.5$.

Furthermore we can leverage the sampling procedures that we use to compute γ to estimate the individualized or subpopulation-based acceptance rate $\eta(t; x_S^{pre}, a, g)$ (as detailed in B.1 and B.3). To guarantee acceptance with certainty, the decision threshold can be set to $t = 0$.

For the explainee, it is vital that the acceptance guarantee is presented in a human-intelligible fashion. In contrast to previous work in the field, we suggest to communicate the acceptance guarantee in terms of a probability.¹⁰ Furthermore, for subpopulation-based recourse, the set of subgroup characteristics should be transparent. In the hospital admission example, the subpopulation-based acceptance guarantee could be communicated as follows: *Within a group of individuals that share your zip code, a vaccination leads to acceptance with at least probability η .*

8 Experiments

In the experiments we evaluate the following questions, assuming correct causal knowledge and accurate models of the conditional distributions in the data:

Q1: Do CE, CR and ICR lead to improvement?

Q2: Do CE, CR and ICR lead to acceptance (by pre- and post- post-recourse predictor)?

Q3: Do CE, CR and ICR lead to acceptance by other predictors with comparable test error?¹¹

Q4: How costly are CE, CR and ICR recommendations?

Setup We evaluate CE, individualized and subpopulation-based CR and ICR with various confidence levels, over multiple runs, and on multiple synthetic and semi-synthetic datasets with known ground-truth (listed below).¹² Random forests were used for prediction, except in the *3var* settings where logistic regression models were used. Following Dandl et al. [2020], we use NSGA-II [Deb et al., 2002] for optimization. For a full specification of the SCMs including the linear cost functions we refer to C.2. Details on the implementation and access to the code are provided in C.1.

3var-causal: A linear gaussian SCM with binary target Y , where all features are causes of Y .

3var-noncausal: The same setup as *3var-causal*, except that one of the features is an effect of Y .

5var-skill: A categorical semi-synthetic SCM where programming skill-level is predicted from causes (e.g. *university degree*) and non-causal indicators extracted from GitHub (e.g. *commit count*).

7var-covid: A semi-synthetic dataset inspired by a real-world covid screening model [Jehi et al., 2020, Wynants et al., 2020].¹³ The model includes typical causes like *covid vaccination* or *population density* and symptoms like *fever* and *fatigue*. The variables are mixed categorical and continuous with various noise distributions. Their relationships include nonlinear structural equations.

Results The results are visualized in Figure 2 and provided in tabular form in C.3.

Q1 (Figure 2a): In scenarios where gaming is possible and lucrative (*3var-noncausal*, *5var-skill* and *7var-covid*) ICR reliably guides towards improvement, but CE and CR game the predictor and yield improvement rates close to zero. For instance, on *5var-skill* CE and CR exclusively suggest to tune the GitHub profile (e.g. by adding more commits). Since the employer offered recourse it should be honored although the applicants remain unqualified. In contrast, ICR suggests to get a degree or to gain experience, such that recourse implementing individuals are suited for the job.

On *3var-causal*, where gaming is not possible, CR also achieves improvement. However, since acceptance w.r.t to a decision threshold $t = 0.5$ is targeted, only improvement rates close to 50% are achieved (the expected predicted score translates into γ^{sub} (Proposition 3)).

For subp. ICR, γ^{obs} is below $\bar{\gamma}$, because the subpopulation may include individuals that were already accepted pre-recourse, such that γ^{sub} and γ^{obs} may not coincide.

Q2 (Figure 2d): All methods yield the desired acceptance rates w.r.t. to the pre-recourse predictor.¹⁴ For CE and CR η^{obs} is higher than for ICR, and for ind. recourse higher than for subp. recourse. Curiously, although no acceptance

¹⁰For CR, the acceptance confidence is encoded in a hyperparameter, as explained in E.2.

¹¹The problem that refits on the same data with similar performance have different mechanism is known as the Rashomon problem or model multiplicity [Breiman, 2001, Pawelczyk et al., 2020, Marx et al., 2020].

¹²For ground-truth counterfactuals, simulations are necessary [Holland, 1986].

¹³The real-world screening model is used to decide whether individuals need a test certificate to enter a hospital. It can be accessed via <https://riskcalc.org/COVID19/>.

¹⁴ICR holds the acceptance rates from Proposition 4, as analyzed in more detail in C.3.

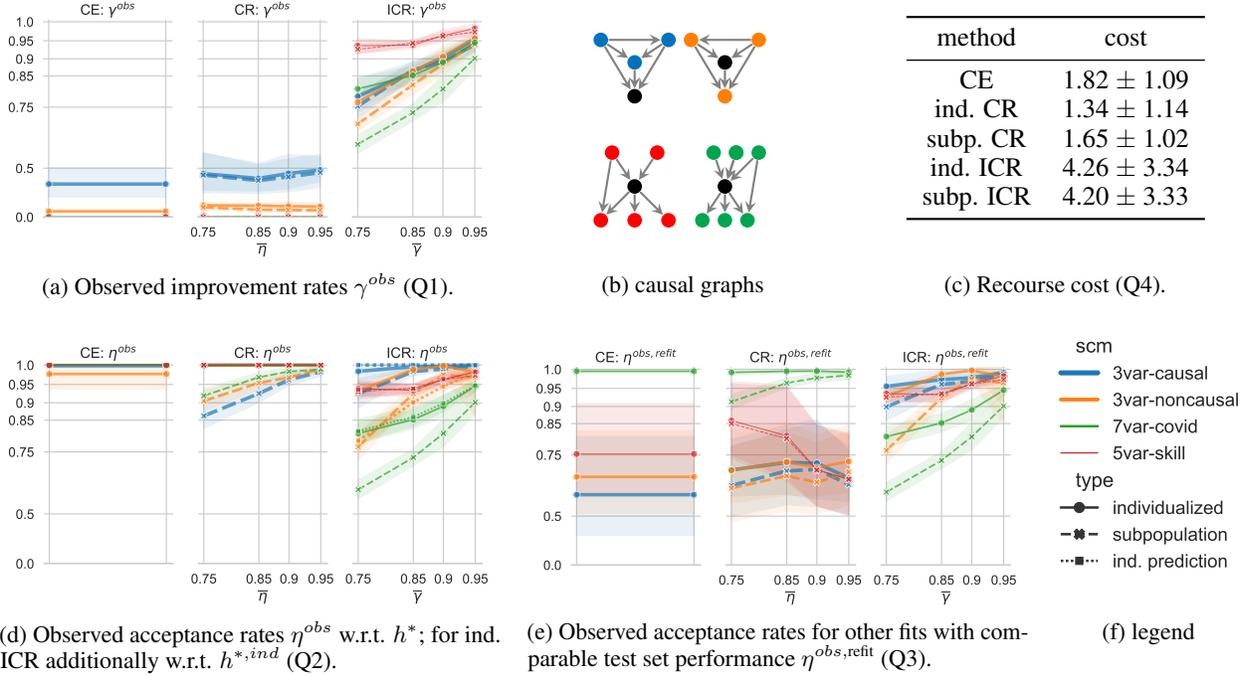


Figure 2: Experimental results for CE, CR and ICR on four datasets over 10 runs on 200 individuals each. For the probabilistic methods the confidences 0.75, 0.85, 0.9, 0.95 were targeted (for CR: $\bar{\eta}$, for ICR: $\bar{\gamma}$). For CE no slack is allowed, such that the results correspond to a confidence level of 1.0. Values are reported on a quadratic scale.

guarantees could be derived for the pre-recourse predictor and ind. ICR, we find that both pre- and ind. post-recourse predictor reliably lead to acceptance.¹⁵

Q3 (Figure 2e): We observe that CE and CR actions are unlikely to be honored by other model fits with similar performance on the same data. This result is highly relevant to practitioners, since models deployed in real-world scenarios are regularly refitted. As such, individuals that implemented acceptance-focused recourse may not be accepted after all, since the decision model was refitted in the meantime. In contrast, ICR acceptance rates are nearly unaffected by refits. The result confirms our argument that improvement-focused recourse may be more desirable for explainees (Section 4).

Q4 (Table 2c): CR actions are cheaper than ICR actions, since improvement may require more effort than gaming. As such, CR has benefits for the explainee: For instance, on *5var-skill*, CR suggests to tune the GitHub profile (e.g. by adding more commits), which requires less effort than earning a degree or gaining job experience. Detailed results on cost are reported in C.3.

In conclusion, ICR actions require more effort than CR, but lead to improvement and acceptance while being more robust to refits of the model.

9 Limitations and Discussion

Causal knowledge and assumptions Individualized ICR requires a fully specified SCM; Subpopulation-based ICR is less demanding but still requires the causal graph and causal sufficiency. SCMs and causal graphs are rarely readily available in practice [Peters et al., 2017] and causal sufficiency is difficult to test [Janzing et al., 2012]. Research on causal inference gives reason for cautious optimism that the difficulties in constructing SCMs and causal graphs can eventually be overcome [Spirtes and Zhang, 2016, Peters et al., 2017, Heinze-Deml et al., 2018, Malinsky and Danks, 2018, Glymour et al., 2019].

There are further foundational problems linked to causality that affect our approach: causal cycles, an ontologically vague target Y (e.g. in hiring), disparities in our data, or causal model misspecification [Barocas and Selbst, 2016,

¹⁵Given that the ind. post-recourse predictor is much more difficult to estimate, the pre-recourse predictor in combination with individualized acceptance guarantees (B.1) may cautiously be used as fallback.

Barocas et al., 2017, Bongers et al., 2021]. All of these factors are considered difficult open problems and may have detrimental impact on our, as well as on any other, recourse framework.

Guiding action without causal knowledge is impossible; when causal knowledge is available, our work provides a normative framework for improvement-focused recourse recommendations. Thus, we join a range of work in explainability [Frye et al., 2020, Heskes et al., 2020, Wang et al., 2021, Zhao and Hastie, 2021] and fairness [Kilbertus et al., 2017, Kusner et al., 2017, Zhang and Bareinboim, 2018, Makhlouf et al., 2020] that highlights the importance of causal knowledge.

Contestability Improvement-focused recourse guides individuals towards actions that help them to improve, e.g., it recommends a vaccination to lower the risk to get infected with Covid. If, however, a explainee is more interested in contesting the algorithmic decision, (improvement-focused) recourse recommendations are not sufficient. Think of an individual who is denied entrance to an event because of their high Covid risk prediction, which is based on a non-causal, spurious association with their country of origin¹⁶. In such situations, we suggest to additionally show explainees diverse explanations, which enable to contest the decision. For example, such an explanation could be: if your country of origin would be different, your predicted Covid risk would have been lower.

10 Conclusion

In the present paper, we took a causal perspective and investigated the effect of recourse recommendations on the underlying target variable. We demonstrated that acceptance-focused recourse recommendations like counterfactual explanations or causal recourse may not improve the underlying prediction but game the predictor instead. The problem stems from predictive, but non-causal relationships, which are abundant in machine learning applications.¹⁷

We tackled the problem in the explanation domain and introduced Improvement-Focused Causal Recourse (ICR), an explanation technique that guides towards improvement of the prediction target and demonstrated how to design post-recourse predictors such that improvement leads to acceptance. We confirm the theoretical results in experiments. With ICR we hope to inspire a shift from acceptance- to improvement-focused recourse.

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¹⁶E.g., due to a spurious association with the causal variable *type of vaccine*.

¹⁷For instance, in hiring, certain keywords in the CV may be associated with qualification, but adding them to the CV does not improve aptitude [Strong, 2022].

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A Extended Background

As follows, we recapitulate well-known definitions in our notation, provide more detailed background on related work and recapitulate results that we use in the proofs. Readers who are already familiar with recourse terminology and d -separation (A.1 and A.2), and who are not interested in more detailed introductions of intervention stability (A.3, only required for the proof of Proposition 2) or causal recourse (A.4), may skip this section.

A.1 Overview of important terms

An overview of important terms is provided in Table 1.

A.2 d -separation

Two variable sets X, Y are called d -separated [Geiger et al., 1990, Spirtes et al., 2000] by the variable set Z in a graph \mathcal{G} (denoted as $X \perp_{\mathcal{G}} Y|Z$), if, and only if, for every path p it either holds that (i) p contains a chain $i \rightarrow m \rightarrow j$ or a fork $i \leftarrow m \rightarrow j$ where $m \in Z$ or (ii) p contains a collider $i \rightarrow m \leftarrow j$ such that m and for all of its descendants n it holds that $m, n \notin Z$. Given the causal Markov property, d -separation in a causal graph implies (conditional) independence in the data [Peters et al., 2017].

A.3 Generalizability and intervention stability

For Proposition 2, we leverage necessary conditions for invariant conditional distributions as derived in [Pfister et al., 2021]. The authors introduce a d -separation based intervention stability criterion that is applied to a modified version of \mathcal{G} . For every intervened upon variable X_l an auxiliary intervention variable, denoted as I_l , is added as direct cause of X_l , yielding \mathcal{G}^* . The intervention variable can be seen as a switch between different mechanisms. A set $S \subseteq \{1, \dots, d\}$ is called *intervention stable* regarding a set of actions if for all intervened upon variables X_l (where $l \in I^{\text{total}}$) the d -separation $I^l \perp_{\mathcal{G}^*} Y|X_S$ holds in \mathcal{G}^* . The authors show that intervention stability implies an invariant conditional distribution, i.e., for all actions $a, b \in \mathbb{A}$ with $I^a, I^b \subseteq I^{\text{total}}$ it holds that $p(y^a|x_S) = p(y^b|x_S)$ (Pfister et al. [2021], Appendix A).

A.4 Causal recourse

ICR is closely related to the CR framework [Karimi et al., 2020b, 2021], but differs substantially in its motivation and target. In order to allow for a direct comparison we briefly sketch the main ideas and the central CR definitions in our notation. Like ICR, CR aims to guide individuals to revert unfavorable algorithmic decisions (recourse). Therefore, they suggest to search for cost-efficient actions that lead to acceptance by the prediction model. Actions are modeled as structural interventions $a : \Pi \rightarrow \Pi$, which can be constructed as $a = do(\{X_i := \theta_i\}_{i \in I})$, where I is the index set of features to be intervened upon [Karimi et al., 2021]. The conservativeness of the suggested actions can be adjusted using the hyperparameter γ_{LCB} , that determines the adaptive threshold $\text{thresh}(a)$ and thereby how many standard deviations the expected prediction shall be away from the model’s decision threshold t . In order to accommodate different levels of causal knowledge, two probabilistic versions of CR were introduced [Karimi et al., 2020b]: While individualized recourse assumes knowledge of the SCM, subpopulation-based CR only assumes knowledge of the causal graph.

Table 1: Overview of important terms and their meanings.

term	meaning
explainee	individual for whom the explanation is generated, e.g. loan applicant
model authority	decision-making entity, e.g. credit institute
recourse	action of the explainee that reverts unfavorable decision
acceptance	desirable model prediction ($\hat{Y} = 1$)
improvement	(yield) desirable state of the underlying target ($Y = 1$)
gaming	yield acceptance without improvement, e.g. treating the symptoms
pre-/post-recourse	before/after implementing recourse recommendation
contestability	the explainee’s ability to contest an algorithmic decision
robustness of recourse	probability that recourse is accepted despite model/data shifts

Individualized recourse Individualized recourse predicts the effect of actions using structural counterfactuals [Karimi et al., 2021], which require a full specification of the SCM.

Given a function that evaluates the cost of actions ($\text{cost}(a, x^{pre})$), the optimization goal for individualized causal recourse is given below. The adaptive threshold thresh bounds the prediction away from the decision threshold.¹⁸

$$a^* \in \underset{a \in \mathbb{A}}{\text{argmin}} \quad \text{cost}(a, x^{pre}) \quad \text{s.t.} \quad \mathbb{E}[\hat{h}(x^{post}) | do(a), x^{pre}] \geq \text{thresh}(a)$$

$$\text{with } \text{thresh}(a) := 0.5 + \gamma_{LCB} \sqrt{\text{Var}[\hat{h}(x^{post,a})]}$$

Subpopulation-based recourse: If no knowledge of the SCM is given, counterfactual distributions cannot be estimated and consequently individualized recourse recommendations cannot be computed. Subpopulation-based CR is based on the average treatment effect within a subgroup of similar individuals [Karimi et al., 2020b]. More specifically individuals belong to the same group if the non-descendants $nd(I)$ of intervention variables (which ceteris paribus remain constant despite the intervention) take the same value. The subpopulation-based objective is given below.

$$a^* \in \underset{a \in \mathbb{A}}{\text{argmin}} \quad \text{cost}(a, x^{pre}) \quad \text{s.t.} \quad \mathbb{E}_{X_{d(I)} | do(X_I = \theta), x_{nd(I)}^{pre}} [\hat{h}(x_{nd(I)}^{pre}, \theta, X_{d(I)})] \geq \text{thresh}(a).$$

¹⁸Further constraints have been suggested, e.g., $x^{post,a} \in \mathcal{P}$ lausible or $a \in \mathcal{F}$ easible [Laugel et al., 2019, Ustun et al., 2019, Mahajan et al., 2020, Dandl et al., 2020, Karimi et al., 2021].

B Estimation and Optimization

As follows we provide detailed explanations of the proposed estimation procedures. First, we explain how to sample from the individualized post-recourse distribution, which allows us to estimate the individualized improvement and acceptance rates (γ^{ind} and η^{ind} , B.1). Based on the same sampling mechanism we can also estimate the individualized post-recourse prediction $h^{*,ind}$ (B.2). Then we explain how to sample from the subpopulation-based post-recourse distribution, which allows us to estimate the subpopulation-based improvement and acceptance rates (γ^{sub} and η^{sub} , B.3). Furthermore, we provide details on optimization (B.4) and demonstrate that the optimal observational predictor h^* can also be estimated using the SCM (B.5).

B.1 Estimation of the individualized improvement confidence γ^{ind} and individualized acceptance rate η^{ind}

We recall that γ^{ind} is the counterfactual probability of the underlying target Y taking the favorable outcome, and η^{ind} the counterfactual probability of the prediction \hat{Y} taking the favorable outcome. In order to estimate γ^{ind} and η^{ind} we first sample covariates and target from the counterfactual post-recourse distribution and then compute the proportion of favorable outcomes for Y and \hat{Y} in the sample.

In general, sampling from counterfactual distributions based on a SCM is performed in three steps (Section 3, [Pearl, 2009]).

1. *Abduction*: The exogenous noise variables are reconstructed from the observations, i.e., $p(u_{Y,D}|x^{pre})$ is estimated.
2. *Intervention*: The intervention $do(a)$ on the SCM \mathcal{M} is performed by replacing the respective structural equations $f_{I_a} := \theta_{I_a}$, yielding $\mathcal{M}_{do(a)}$.
3. *Prediction*: The abducted noise variables are sampled from $p(u_{Y,D}|x^{pre})$ and passed through the model $\mathcal{M}_{do(a)}$ to sample from the counterfactual distribution $P(Y^{post}, X^{post}|x^{pre}, do(a))$.

Given knowledge of the SCM, the challenge is to sample the exogenous variables from $p(u_{Y,D}|x^{pre})$ (abduction). As follows we explain the abduction in two steps. First, we explain how we can abduct u_j for variables for which both the node x_j and all parents $(x, y)_{pa(j)}$ are observed, which we refer to as the standard abduction case. Then we factorize the abduction of the joint $p(u_{Y,D}|x^{pre})$ into several components which can be reduced to said standard abduction case. The sampling procedure is summarized in Algorithm 1.

B.1.1 Recap: Standard abduction

If for a node u_j both the node $(x, y)_j$ and the parents $(x, y)_{pa(j)}$ are observed, we can apply standard abduction. The standard abduction procedure depends on the type of structural equation and exogenous noise distribution.

Given invertible structural equations, observation of $x_j, x_{pa(j)}$ determines u_j . More specifically, u_j can be reconstructed using

$$u_j = f^{-1}(x_j; x_{pa(j)}).$$

For instance, for additive structural equations $f_j(u_j; x_{pa(j)}) = g(x_{pa(j)}) + u_j$, the inversion is given by $f_j^{-1}(x_j; x_{pa(j)}) = x_j - g(x_{pa(j)})$.

In our experiments we also included binomial variables with a sigmoidal (non-invertible) structural equation. More specifically, the structural equations are defined as $x_j = [\sigma(l(x_{pa(j)})) \leq u_j]$ with $U_j \sim Unif(0, 1)$. Here σ refers to the sigmoid function and l to some linear combination. $[cond]$ evaluates to 1 when the condition is true and otherwise to 0. Intuitively, $\sigma(l(x_{pa(j)}))$ can be seen as a nonlinear activation function which determines the probability of the node being activated ($x_j = 1$). u_j acts as a dice, where values $\leq \sigma(l(x_{pa(j)}))$ imply $x_j = 1$ and vice versa.

For those variables, if $x_j = 1$, we know that $u_j \leq \sigma(l(x_{pa(j)}))$ and vice versa, such that we can abduct U_j as follows (and can therefore sample u_j):

$$P(U_j|x_j; x_{pa(j)}) = \begin{cases} Unif(0, \sigma(l(x_{pa(j)}))), & \text{for } x_j = 1 \\ Unif(\sigma(l(x_{pa(j)})), 1), & \text{for } x_j = 0 \end{cases}$$

As we will see in the next section, our estimation procedure can be flexibly extended to SCMs with different types of structural equations, as long as a procedure to sample from the abducted exogenous noise variable for the standard case (where parents and the node itself are observed) is available.

Algorithm 1: Sampling from the individualized post-recourse distribution

Data: pre-recourse observation x^{pre} , action a (where $do(a) := do(X_{I_a} := \theta)$), sample size M , structural causal model \mathcal{M} with structural equations f_j , observational predictor h

Result: sample from $p(y^{post}, x^{post} | x^{pre}, do(a))$

get $\mathcal{M}_{do(a)}$ by updating $f_i(x_{pa(i)}; u_i) := \theta_i$ for $i \in I_a$;

for m **in** $(0, \dots, M - 1)$ **do**

 sample y' from $Binomial(h(x^{pre}))$;

for j **in** D **do**

 sample $u_j^{(m)}$ from $p(u_j | (x, y')_j, (x, y')_{pa(j)})$ \triangleright comment: leveraging standard abduction;

end

 sample $u_Y^{(m)}$ from $p(u_Y | y', x_{pa(Y)})$;

 compute $(x^{post}, y^{post})^{(m)} = f_{\mathcal{M}_{do(a)}}(u^{(m)})$;

end

B.1.2 Factorization of $p(u|x)$

We have demonstrated how to abduct individual nodes in the standard setting where the corresponding endogenous variable and its parents are observed.

As follows we demonstrate how to sample from the joint distribution of the exogenous variables given an observation of X (and without observing Y). Therefore, we show that $p(u|x)$ can be seen as a mixture of two distributions, one for each possible state y' of Y . In order to sample from it, we (1) need to sample y' from the mixing distribution $p(y|x)$ and (2) given y' , sample from the respective abducted noise variable $p(u|y', x)$.

$$p(u|x) \stackrel{\text{law tot. prob.}}{=} \sum_{y' \in \{0,1\}} p(u, y'|x) \stackrel{\text{cond. prob.}}{=} \sum_{y' \in \{0,1\}} p(u|y', x)p(y'|x) \quad (2)$$

The binomial mixing distribution $p(y|x)$ can be obtained and sampled from by leveraging the cross-entropy optimal predictor h^* (which can for instance be derived from the SCM, see B.5). In order to sample from $p(u|y', x)$ we leverage the Markov factorization, which allows us to sample each component independently using the standard abduction procedure described above.

$$p(u|x, y') \stackrel{\text{d-sep.}}{=} P(u_Y | x_{pa(Y)}, y') \prod_{k \in ch(Y)} P(u_k | x_k, x_{pa(k)}, y') \prod_{k \notin ch(Y)} P(u_k | x_k, x_{pa(k)}). \quad (3)$$

The overall procedure is summarized in Algorithm 1.

B.1.3 Estimation of γ^{ind} and η^{ind}

Given the procedure to sample from the individualized post-recourse distribution we can estimate γ^{ind} by taking the mean over the samples taken for Y^{post} . Similarly, for each sample for X^{post} we can compute the prediction \hat{y}^{post} using either $h \geq t$ or $h^{ind} \geq t$. By taking the mean over all sampled predictions \hat{y}^{post} we can estimate the respective acceptance probability $\eta(t; x^{pre}, a, h)$ or $\eta(t; x^{pre}, a, h^{ind})$.

B.2 Estimation of the individualized post-recourse prediction

We continue to show how the individualized post-recourse prediction can be estimated. We recall that $h^{*,ind}$ is

$$h^{*,ind}(x^{post}; x^{pre}, a) = P(Y^{post} = 1 | x^{post}, x^{pre}, do(a)).$$

We can estimate $h^{*,ind}$ by leveraging the procedure to sample from the post-recourse covariate distribution (Algorithm 1). More specifically, we draw samples (y', x') from $P(Y^{post}, X^{post} | do(a), x^{pre})$ and keep those that conform with x^{post} (i.e., $x' = x^{post}$). Within the subsample, we compute the proportion of samples for which $y' = 1$ to estimate $p(y^{post} | x^{pre}, x^{post}, do(a))$. In more formal terms, we approximate Eq. 4 using rejection sampling and Monte Carlo integration [Koller and Friedman, 2009].

If the structural equations are invertible¹⁹ or the nodes are categorical the procedure is tractable, since many or all

¹⁹Meaning that the abducted joint distribution has point mass probability for two configurations, one for each possible state of Y .

Algorithm 2: Estimating $h^{*,ind}$

Data: pre-recourse observation x^{pre} , action a , sample size M , structural causal model \mathcal{M} , observational predictor h , $m = 0$

Result: $\hat{h}^{ind}(x^{post}; x^{pre}, do(a))$

while $m < M$ **do**

 sample (x', y') using Alg. 1 and $x^{pre}, a, \mathcal{M}, h$;

if $x' = x^{post}$ **then**

$m = m + 1$; store y' as $y^{(m)}$;

end

end

$$\hat{h}^{ind}(x^{post}) = \frac{1}{M} \sum_{m=1}^M y^{(m)}$$

samples conform with x^{post} . Otherwise the estimation may become intractable. We see the application of likelihood weighting or MCMC as promising directions and refer interested readers to Koller and Friedman [2009]. In addition to the sampling-based procedure we also derive a closed-form solution for settings with invertible structural equations, which is provided in Proposition 5, Eq. 5.

Proposition 5. *In general, the individualized post-recourse predictor can be estimated as*

$$\begin{aligned} & p(y^{post} | x^{pre}, x^{post}, do(a)) \\ &= \frac{\int_{\mathcal{U}} p(y^{post}, x^{post} | u, do(a)) p(u | x^{pre}) du}{\sum_{y' \in \{0,1\}} (\int_{\mathcal{U}} p(y', x^{post} | u, do(a)) p(u | x^{pre}) du)} \end{aligned} \quad (4)$$

Given invertible structural equations, the individualized post-recourse prediction function reduces to

$$\begin{aligned} & p(y^{post} | x^{post}, x^{pre}, do(a)) \\ &= \frac{p(U_{-I} = f_{do(a)}^{-1}(y^{post}, x^{post}) | x^{pre}, do(a))}{\sum_{y' \in \{0,1\}} p(U_{-I} = f_{do(a)}^{-1}(y', x^{post}) | x^{pre}, do(a))}. \end{aligned} \quad (5)$$

B.3 Estimation of the subpopulation-based improvement confidence γ^{sub} and the subpopulation-based acceptance rate η^{sub}

As follows we detail how to estimate γ^{sub} and η^{sub} . We focus on actions a that potentially affect Y , meaning that they intervene on causes of Y .²⁰

In order to estimate γ^{sub} and η^{sub} we sample (x', y') from the subpopulation-based post-recourse distribution. Given a sample from the subpopulation-based post-recourse distribution we can estimate γ^{sub} and η^{sub} by taking the respective sample means.

We explain the sampling procedure in two steps: We first recall how causal graphs can be leveraged to sample interventional distributions, and then explain why we can apply the procedure to sample from the subpopulation-based post-recourse distribution.

Recap: Sampling interventional distributions leveraging a causally sufficient causal graph \mathcal{G} Given a causal graph \mathcal{G} (that fulfills the global Markov property), the joint distribution $P(X, Y)$ can be reformulated using the Markov factorization, which makes use of the d -separations in the graph.

$$p(x, y) = p(y | x_{pa(y)}) \prod_{j \in D} p(x_j | (x, y)_{pa(j)})$$

As a consequence, we can sample from the joint distribution by sampling each component given its respective parents. In order to ensure that the parents for each node have been sampled already, the graph is traversed in topological order, starting with the root node and ending with the sink nodes [Koller and Friedman, 2009].

Given that causal sufficiency (no unobserved confounders) and the principle of independent mechanisms hold, the same

²⁰Actions that do not affect Y trivially do not lead to improvement. The respective probability of $Y = 1$ can be estimated using the optimal observational predictor.

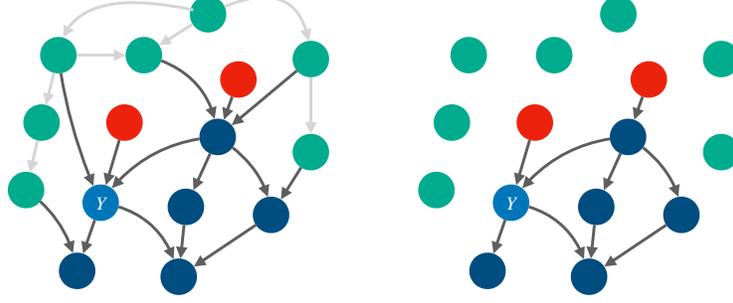


Figure 3: Causal graph $\mathcal{G}_{\bar{I}_a}$ visualizing the subpopulation-based post-recourse setting, including the prediction target Y (light blue), intervened-upon variables I_a (red), the subgroup characteristics G_a (cyan) and the descendants Γ that shall be resampled (dark blue). \bar{I}_a indicates that incoming edges to I_a were removed. Right: Causal graph $\mathcal{G}_{\bar{I}_a G_a}$ where incoming edges to I_a and outgoing edges from G_a were removed. We observe that in this manipulated graph G_a is d -separated from Γ . Thus, according to the second rule of do -calculus, for G_a intervention and conditioning coincide.

Algorithm 3: Sampling from the subpopulation-based post-recourse distribution

Data: pre-recourse observation x^{pre} , action a with $I_a \cap asc(Y) \neq \emptyset$ ($do(a) := do(X_{I_a} := \theta)$), sample size M , causal graph \mathcal{G} , conditional distributions $P(X_j | X_{pa(j)})$ for $j \in \Gamma$ with $\Gamma := \{r : r \in asc(Y) \wedge r \in d(I)\}$

Result: sample from $p(y, x_\Gamma | do(a), x_{G_a})$

for $m \leftarrow 0$ **to** M **do**

- $\Gamma^{sorted} \leftarrow \text{topologicalsort}(\Gamma; \mathcal{G}_{do(a)})$ ▷ sort such that causes precede effects ;
- for** j **in** Γ^{sorted} **do**
- sample $(x, y)_j^{post, (m)} \sim P((X, Y)_j | (X, Y)_{pa(j)} = (x, y)_{pa(j)}^{post})$;
- end**

end

procedure can also be applied when sampling from interventional distributions of the form $p(x, y | do(a))$ by leveraging the so-called truncated factorization. The intervened upon nodes are not sampled from their parents, but fixed to the values θ_a . The remaining nodes Γ are sampled as before:

$$p((x, y)_\Gamma | do(a)) = \prod_{j \in \Gamma} p((x, y)_j | (x, y)_{pa(j) \cap \Gamma}, \theta_{pa(j) \cap I_a})$$

with $\Gamma := D \setminus I_a$

Sampling from the subpopulation-based post-recourse distribution using \mathcal{G} We recall that for actions a that potentially affect Y the subpopulation-based post-recourse distribution is defined as

$$P(Y^{post}, X^{post} | do(a), X_{G_a}^{post} = x_{G_a}^{pre}). \quad (6)$$

As we will see, the previously described sampling procedure can be applied. Therefore we apply the second rule of do -calculus to show that in Equation 6 conditioning on x_{G_a} is equal to intervening $do(X_{G_a} = x_{G_a})$. More specifically, if we remove all outgoing edges from X_{G_a} and all incoming edges to I_a , then X_{G_a} and X_Γ with $\Gamma := D \setminus I_a \cap G_a = d(I_a)$ are d -separated, meaning that conditioning and intervention are equivalent (Figure 3).

$$\begin{aligned} & P((Y, X)_\Gamma^{post} | do(a), X_{G_a}^{post} = x_{G_a}^{pre}) \\ &= P((Y, X)_\Gamma^{post} | do(a), do(X_{G_a}^{post} = x_{G_a}^{pre})) \end{aligned}$$

As follows we can leverage the procedure to sample interventional distributions to sample from the subpopulation-based post-recourse distribution. The procedure is illustrated in Algorithm 3.

B.3.1 Learning the conditional distributions $P(X_j | x_{pa(j)})$

In this work we assume that we have prior knowledge that allows us to sample from the components of the factorization ($P(X_j | x_{pa(j)})$), e.g. available if we know the SCM).

If the conditional distributions are not known, they can be learned from observational data; depending on which assumptions about distribution and functional can be made, different techniques may be employed. For categorical variables the problem reduces to standard supervised learning with cross-entropy loss. For linear Gaussian data, the conditional distribution can be estimated analytically from the covariance matrix [Page Jr, 1984]. A variety of estimation techniques exist for continuous settings with nonlinearities [Bishop, 1994, Bash-tannyk and Hyndman, 2001, Sohn et al., 2015, Trippe and Turner, 2018, Winkler et al., 2019, Hothorn and Zeileis, 2021].

B.4 Optimization

Like the optimization problems for CE [Wachter et al., 2017, Tsirtsis and Gomez Rodriguez, 2020] or CR [Karimi et al., 2020b], the optimization problem for ICR is computationally challenging. It can be seen as a two-stage problem, where in the first stage the intervention targets I_a , and in the second stage the corresponding intervention values θ_a are optimized [Karimi et al., 2020b]. For the selection of intervention targets I_a alone $2^{d'}$ combinations exist, with $d' \leq d$ being the number of causes of Y . We jointly optimize the intervention targets and the intervention values using a genetic algorithm called NSGA-II [Deb et al., 2002]. For mixed categorical and continuous data, previous work in the field [Dandl et al., 2020] suggests to use NSGA-II in combination with *mixed integer evaluation strategies* [Li et al., 2013]. The exact hyperparameter configurations are reported in C.3.

B.5 Estimation of the optimal observational predictor h^* using the SCM

Instead of leveraging supervised learning with cross-entropy loss, we can factorize the optimal observational predictor as shown in Proposition 6 and then leverage the SCM for the estimation.

Proposition 6. *The optimal observational predictor can be factorized into conditional distributions of nodes given their parents (using the Markov factorization). More specifically, we yield*

$$p(y|x) = \frac{p(x, y)}{p(x)} = \frac{p(x, y)}{\sum_{y' \in \{0,1\}} p(x, y')} \quad (7)$$

$$\stackrel{\text{M.f.}}{=} \frac{p(y|x_{pa(j)}) \prod_{j \in D} p(x_j|(x, y)_{pa(j)})}{\sum_{y' \in \{0,1\}} p(y'|x_{pa(j)}) \prod_{j \in D} p(x_j|(x, y')_{pa(j)})} \quad (8)$$

$$= \frac{p(y|x_{pa(j)}) \prod_{j \in ch(y)} p(x_j|x_{pa(j)}, y)}{\sum_{y' \in \{0,1\}} p(y'|x_{pa(j)}) \prod_{j \in ch(y)} p(x_j|x_{pa(j)}, y')} \quad (9)$$

It remains to show how the conditional distribution $p(x_j|x_{pa(j)})$ of a node given its parents can be estimated. Generally it holds that

$$p(x_j|x_{pa(j)}) \quad (10)$$

$$\stackrel{\text{law tot. prob.}}{=} \int_{\mathcal{U}_j} p(x_j|x_{pa(j)}, u_j) p(u_j|x_{pa(j)}) du \quad (11)$$

$$\stackrel{\text{SCM, } u_j \perp x_{pa(j)}}{=} \int_{\mathcal{U}_j} [f(x_{pa(j)}, u_j) = x_j] p(u_j) du. \quad (12)$$

The integral can be approximated using Monte Carlo integration: we can sample from $p(u_j)$, compute the respective $\tilde{x}_j = f_j(x_{pa(j)}, \tilde{u}_j)$ and compute the proportion of cases where $x_j = \tilde{x}_j$. If X_j and U_j are continuous, this may require huge sample sizes to converge.

Furthermore, we may be able to leverage assumptions about f_j to derive a closed form solution. If f_j is invertible, the integral reduces to $p(x_j|x_{pa(j)}) = p(U_j = f_j^{-1}(x_j, x_{pa(j)}))$. For binary nodes with $x_j := [\sigma(l(x_{pa(j)})) \leq u_j]$ and $U_j \sim \text{Unif}(0, 1)$, we directly see that $p(x_j|x_{pa(j)}) = \sigma(l(x_{pa(j)}))$.

C Details on Experiments

In this section we provide additional details on the experiments. More specifically, we explain which open-source libraries we use, how to access our code and how to reproduce the results in C.1. We formally introduce the synthetic and semi-synthetic datasets that we used in our experiments in C.2 and the corresponding figures. Details on hyperparameters, models as well as detailed results are reported in C.3 and the corresponding tables.

C.1 Implementation

The code relies of efficient tensor calculations with `numpy` [Harris et al., 2020], `pytorch` [Paszke et al., 2019] and `jax` [Bradbury et al., 2018]. For named dataframes we use `pandas` [pandas development team, 2020]. For plotting we rely on `matplotlib` [Hunter, 2007] and `seaborn` [Waskom, 2021]. We use the evolutionary optimization library `deap` [Fortin et al., 2012] and `NSGA-II` [Deb et al., 2002] to solve the combinatorial optimization problem.²¹ In order to speed up the computation, we cache queries and results for the improvement confidence using `functools.cache`. For continuous variables the intervention can be rounded to a specified number of digits to increase the probability of reusing a cached result (with neglectable loss of precision).²²

All code is publicly available via <https://github.com/gcskoenig/icr>. The repository contains the user-friendly python package `icr`, which we use in our experiments to generate and evaluate recourse. Furthermore, the scripts for the experiments, the scripts for the visualization of the results as well as a `README.md` with instructions for the installation of all dependencies are contained in the repository, such that the experiments are reproducible.

C.2 Synthetic and Semi-Synthetic Datasets

3var-causal and *3var-noncausal* are abstract, synthetic settings. *5var-skill* is inspired by Montandon et al. [2021], who use GitHub profiles to detect the role of a developer. In our SCM we model *senior-level skill* as a binary variable which is caused by *programming experience* and the education *degree*. The skill is causal for GitHub metrics such as the number of *commits*, the number of programming *languages* and the number of *stars*. The *7var-covid* dataset is inspired by Jehi et al. [2020]. The following variables are introduced: population density D , flu vaccination V_I , number of covid vaccination shots V_C , deviation from average BMI B , whether someone is free of covid disease C , whether the individual has influence I , appetite loss S_A , fever S_{Fe} and fatigue S_{Fa} . The corresponding structural equations, noise distributions and causal graphs are provided in Figure 4 (*3var-causal*), 5 (*3var-noncausal*), 6 (*5var-skill*) and 7 (*7var-covid*). A pairplot for each dataset is presented in Figure 8. In our notation σ is the sigmoid function, N the Gaussian distribution, Cat a categorical distribution, $Unif$ the uniform distribution, $Bern$ a Bernoulli distribution and GaP a Gamma-Poisson mixture. $[cond]$ is 1 when the condition is met and 0 if not. As a consequence variables with $[Z \leq U]$ and $U \sim Unif(0, 1)$ are bernoulli distributed with $Bern(Z)$.

C.3 Detailed Results

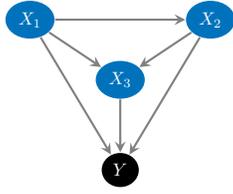
In this section we report all experimental results in tabular form. More specifically, the results for *3var-causal* are reported in Table 2, for *3var-noncausal* in Table 3, for *5var-skill* in Table 4 and for *7var-covid* in Table 5. For each experiment we report the specified confidence γ (or η for CR), as well as the observed improvement rate γ_{obs} , the observed acceptance rate η_{obs} , the observed acceptance rate by the individualized post-recourse predictor η_{obs}^{indiv} , the observed acceptance rate on refits η_{obs}^{refit} and the average recourse cost for individuals who were rejected and whom were provided with a recourse recommendation. A visual summary of the results is provided in Section 8.

In order to enable a more direct comparison of the CR and ICR targets, we equalize the optimization thresholds for ICR and CR. More specifically, for CR we require the (individualized or subpopulation-based) acceptance probability to be $\geq \eta$, and for ICR we require the (individualized or subpopulation-based) improvement probability to be $\geq \bar{\gamma}$, where $\bar{\gamma} = \bar{\eta}$.²³ Furthermore, in order to be able to estimate the effects of recourse actions, CR assumes causal sufficiency, meaning that there are no two endogeneous variables that share an unobserved cause. If the target variable Y is exogeneous then any causal model with more than one endogeneous direct effect of Y violates the assumptions. In order to enable an application of CR on datasets with more than one effect variable we assume knowledge of the SCM

²¹We also implemented abduction based on probabilistic inference. Thereby we rely on `pyro` [Bingham et al., 2018] for discrete inference and `numpyro` [Phan et al., 2019] for MCMC inference of continuous variables. For our experiments we used the analytical formulas presented in B

²²All packages are open source. For detailed license information we refer to the respective package websites.

²³A short comment on the choice of a non-adaptive threshold can be found in E.2.

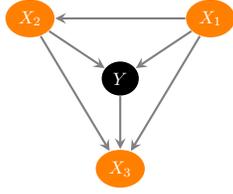


(a) Causal graph

$$\begin{aligned}
 X_1 &:= U_1, & U_1 &\sim N(0, 1) \\
 X_2 &:= X_1 + U_2, & U_2 &\sim N(0, 1) \\
 X_3 &:= X_1 + X_2 + U_3, & U_3 &\sim N(0, 1) \\
 Y &\sim [\sigma(X_1 + X_2 + X_3) \leq U_Y], & U_Y &\sim Unif(0, 1)
 \end{aligned}$$

(b) Structural Equations

Figure 4: SCM for *3var-causal*. The cost function is given as $cost(a) = \delta_1 + \delta_2 + \delta_3$, where δ is the vector of absolute changes to the intervened upon variables. E.g., for $do(a) = do(X_1 = x'_1)$, $\delta_1 = |x'_1 - x_1|$ and $\delta_2 = \delta_3 = 0$

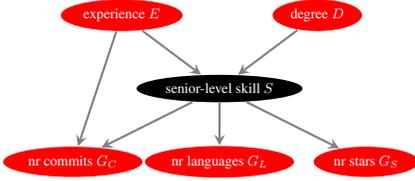


(a) Causal graph

$$\begin{aligned}
 X_1 &:= U_1, & U_1 &\sim N(0, 1) \\
 X_2 &:= X_1 + U_1, & U_1 &\sim N(0, 1) \\
 Y &:= [\sigma(X_1 + X_2) \leq U_Y], & U_Y &\sim Unif(0, 1) \\
 X_3 &:= X_1 + X_2 + Y + U_3, & U_3 &\sim N(0, 0.1)
 \end{aligned}$$

(b) Structural Equations

Figure 5: SCM for *3var-noncausal* with $cost(a) = \delta_1 + \delta_2 + \delta_3$.

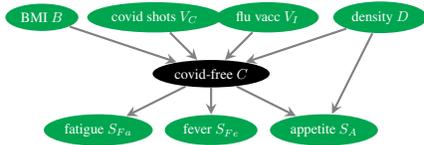


(a) Causal graph

$$\begin{aligned}
 E &:= U_E; U_E \sim GaP(8, 8/3) \\
 D &:= U_D; U_D \sim Cat(0.4, 0.2, 0.3, 0.1) \\
 S &:= [\sigma(-10 + 3E + 4D) \leq U_S]; U_S \sim Unif(0, 1) \\
 G_C &:= 10E(11 + 100D) + U_{G_C}; U_{G_C} \sim GaP(40, 40/4) \\
 G_L &:= \sigma(10S) + U_{G_L}; U_{G_L} \sim GaP(2, 2/4) \\
 G_S &:= 10S + U_{G_S}; U_{G_S} \sim GaP(5, 5/4)
 \end{aligned}$$

(b) Structural Equations

Figure 6: SCM for *5var-skill* with $cost(a) = 5\delta_E + 5\delta_D + 0.0001\delta_{G_C} + 0.01\delta_{G_L} + 0.1\delta_{G_S}$.

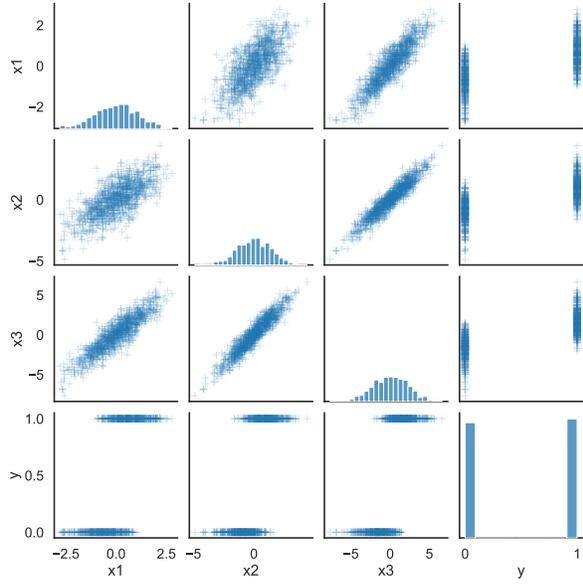


(a) Causal graph

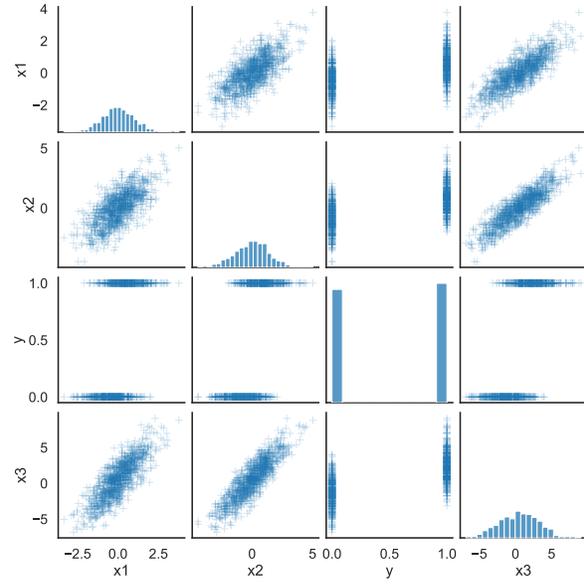
$$\begin{aligned}
 D &:= U_D; U_D \sim \Gamma(4, 4/3) \\
 V_I &:= U_{V_I}; U_{V_I} \sim Bern(0.39) \\
 V_C &:= U_{V_C}; U_{V_C} \sim Cat(0.24, 0.02, 0.15, 0.59) \\
 B &:= U_B; U_B \sim N(0, 1) \\
 C &:= [\sigma(-(D - 3 - V_I - 2.5V_C + 0.2B^2)) \leq U_C]; \\
 U_C &\sim Unif(0, 1) \\
 S_A &:= [\sigma(-2C) \leq U_{S_A}]; U_{S_A} \sim Unif(0, 1) \\
 S_{F_e} &:= [\sigma(5 - 9C) \leq U_{S_{F_e}}]; U_{S_{F_e}} \sim Unif(0, 1) \\
 S_{F_a} &:= [\sigma(-1 + B^2 - 2C) \leq U_{S_{F_a}}]; \\
 U_{S_{F_a}} &\sim Unif(0, 1)
 \end{aligned}$$

(b) Structural Equations

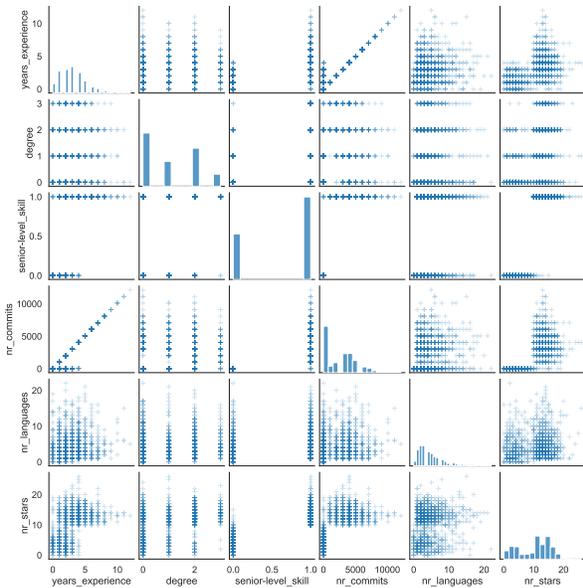
Figure 7: SCM for *7var-covid* with cost function $cost(a) = \delta_D + \delta_{V_I} + \delta_{V_C} + \delta_B + \delta_{S_A} + \delta_{S_{F_e}} + \delta_{S_{F_a}}$.



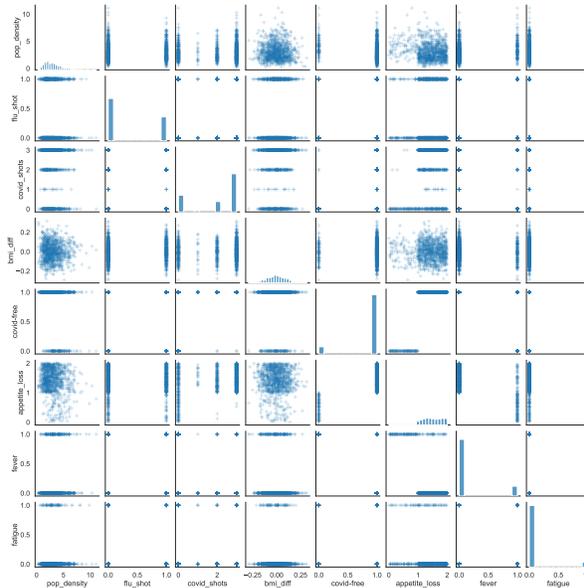
(a) Pairplot for *3var-causal*.



(b) Pairplot for *3var-noncausal*.



(c) Pairplot for *5var-skill*.



(d) Pairplot for *7var-covid*.

Figure 8: Pairplots for the SCMs.

Table 2: Results for 3var-causal.

3var-causal	$\bar{\gamma} / \bar{\eta}$	$\gamma_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}^{\text{individ.}}$	\pm	$\eta_{\text{obs.}}^{\text{refit}}$	\pm	\emptyset cost	\pm
CE	-	0.41	0.09	1.00	0.00	-	-	0.60	0.20	3.08	0.41
ind. CR	0.75	0.47	0.10	1.00	0.00	-	-	0.70	0.10	2.46	0.37
ind. CR	0.85	0.44	0.08	1.00	0.00	-	-	0.72	0.12	2.39	0.25
ind. CR	0.90	0.47	0.09	1.00	0.00	-	-	0.72	0.14	2.36	0.35
ind. CR	0.95	0.49	0.07	1.00	0.00	-	-	0.67	0.10	2.44	0.31
subp. CR	0.75	0.46	0.11	0.86	0.04	-	-	0.64	0.14	2.66	0.41
subp. CR	0.85	0.43	0.08	0.93	0.02	-	-	0.69	0.14	2.64	0.32
subp. CR	0.90	0.45	0.09	0.96	0.02	-	-	0.70	0.15	2.73	0.42
subp. CR	0.95	0.48	0.09	0.98	0.01	-	-	0.64	0.14	2.86	0.41
ind. ICR	0.75	0.79	0.06	0.98	0.02	1.0	0.0	0.96	0.03	3.27	0.50
ind. ICR	0.85	0.86	0.03	1.00	0.01	1.0	0.0	0.97	0.02	3.82	0.30
ind. ICR	0.90	0.90	0.02	1.00	0.01	1.0	0.0	0.98	0.03	3.70	0.31
ind. ICR	0.95	0.95	0.01	1.00	0.00	1.0	0.0	0.99	0.01	4.08	0.24
subp. ICR	0.75	0.75	0.04	0.93	0.04	-	-	0.90	0.04	3.34	0.49
subp. ICR	0.85	0.87	0.03	0.98	0.01	-	-	0.96	0.02	4.05	0.29
subp. ICR	0.90	0.89	0.02	0.99	0.01	-	-	0.97	0.02	3.87	0.25
subp. ICR	0.95	0.94	0.02	1.00	0.00	-	-	0.99	0.01	4.22	0.28

Table 3: Results for 3var-noncausal

3var-noncausal	$\bar{\gamma} / \bar{\eta}$	$\gamma_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}^{\text{individ.}}$	\pm	$\eta_{\text{obs.}}^{\text{refit}}$	\pm	\emptyset cost	\pm
CE	-	0.17	0.03	0.98	0.04	-	-	0.67	0.15	2.28	0.26
ind. CR	0.75	0.25	0.03	1.00	0.00	-	-	0.70	0.13	2.28	0.21
ind. CR	0.85	0.24	0.02	1.00	0.00	-	-	0.73	0.13	2.29	0.17
ind. CR	0.90	0.24	0.04	1.00	0.00	-	-	0.71	0.11	2.24	0.16
ind. CR	0.95	0.23	0.04	1.00	0.00	-	-	0.73	0.12	2.18	0.32
subp. CR	0.75	0.22	0.03	0.91	0.03	-	-	0.63	0.15	2.18	0.12
subp. CR	0.85	0.19	0.03	0.95	0.02	-	-	0.67	0.15	2.33	0.21
subp. CR	0.90	0.19	0.03	0.97	0.01	-	-	0.65	0.14	2.42	0.19
subp. CR	0.95	0.19	0.03	0.99	0.01	-	-	0.69	0.14	2.26	0.32
ind. ICR	0.75	0.77	0.03	0.93	0.02	0.79	0.03	0.93	0.02	2.16	0.11
ind. ICR	0.85	0.86	0.02	0.99	0.01	0.90	0.02	0.99	0.01	2.51	0.08
ind. ICR	0.90	0.91	0.03	1.00	0.00	0.94	0.01	1.00	0.00	3.00	0.08
ind. ICR	0.95	0.96	0.02	0.98	0.07	0.98	0.01	0.98	0.08	3.32	0.16
subp. ICR	0.75	0.69	0.03	0.77	0.05	-	-	0.76	0.05	2.11	0.20
subp. ICR	0.85	0.82	0.03	0.93	0.02	-	-	0.92	0.02	2.42	0.11
subp. ICR	0.90	0.89	0.03	0.98	0.01	-	-	0.97	0.01	2.86	0.13
subp. ICR	0.95	0.94	0.02	0.97	0.10	-	-	0.96	0.12	3.19	0.15

Table 4: Results for 5var-skill

5var-skill	$\bar{\gamma} / \bar{\eta}$	$\gamma_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}^{\text{individ.}}$	\pm	$\eta_{\text{obs.}}^{\text{refit}}$	\pm	\emptyset cost	\pm
CE	-	0.00	0.00	1.00	0.00	-	-	0.76	0.14	1.34	1.28
ind. CR	0.75	0.00	0.00	1.00	0.00	-	-	0.86	0.11	0.27	0.28
ind. CR	0.85	0.00	0.00	1.00	0.00	-	-	0.81	0.14	0.24	0.20
ind. CR	0.90	0.00	0.01	1.00	0.00	-	-	0.70	0.15	0.10	0.00
ind. CR	0.95	0.00	0.00	1.00	0.00	-	-	0.66	0.16	0.11	0.03
subp. CR	0.75	0.00	0.00	1.00	0.00	-	-	0.85	0.11	4.06	4.97
subp. CR	0.85	0.00	0.00	1.00	0.00	-	-	0.80	0.15	0.24	0.19
subp. CR	0.90	0.00	0.01	1.00	0.00	-	-	0.70	0.15	0.10	0.01
subp. CR	0.95	0.00	0.00	1.00	0.00	-	-	0.66	0.15	0.12	0.04
ind. ICR	0.75	0.94	0.02	0.94	0.02	0.94	0.02	0.94	0.02	4.95	5.32
ind. ICR	0.85	0.94	0.01	0.93	0.02	0.94	0.01	0.93	0.02	9.80	0.27
ind. ICR	0.90	0.96	0.02	0.96	0.02	0.96	0.02	0.96	0.02	10.38	0.23
ind. ICR	0.95	0.98	0.01	0.98	0.01	0.98	0.01	0.98	0.01	11.23	0.21
subp. ICR	0.75	0.93	0.01	0.93	0.02	-	-	0.93	0.01	4.72	5.08
subp. ICR	0.85	0.94	0.01	0.94	0.01	-	-	0.94	0.02	9.74	0.17
subp. ICR	0.90	0.96	0.01	0.96	0.01	-	-	0.96	0.01	10.46	0.53
subp. ICR	0.95	0.97	0.01	0.97	0.01	-	-	0.97	0.01	10.88	0.21

Table 5: Results for 7var-covid

7var-covid	$\bar{\gamma} / \bar{\eta}$	$\gamma_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}$	\pm	$\eta_{\text{obs.}}^{\text{individ.}}$	\pm	$\eta_{\text{obs.}}^{\text{refit}}$	\pm	\emptyset cost	\pm
CE	-	0.00	0.00	1.00	0.00	-	-	1.00	0.00	0.60	0.12
ind. CR	0.75	0.01	0.00	1.00	0.00	-	-	0.99	0.01	0.56	0.02
ind. CR	0.85	0.00	0.00	1.00	0.00	-	-	0.99	0.00	0.55	0.02
ind. CR	0.90	0.00	0.00	1.00	0.00	-	-	1.00	0.00	0.55	0.03
ind. CR	0.95	0.00	0.00	1.00	0.00	-	-	0.99	0.01	0.54	0.07
subp. CR	0.75	0.01	0.01	0.92	0.02	-	-	0.91	0.02	0.52	0.03
subp. CR	0.85	0.00	0.01	0.97	0.01	-	-	0.96	0.01	0.75	0.40
subp. CR	0.90	0.00	0.00	0.98	0.01	-	-	0.98	0.01	0.55	0.03
subp. CR	0.95	0.00	0.00	0.99	0.01	-	-	0.98	0.01	0.51	0.07
ind. ICR	0.75	0.81	0.03	0.81	0.03	0.82	0.04	0.81	0.03	1.26	0.02
ind. ICR	0.85	0.85	0.03	0.85	0.03	0.86	0.03	0.85	0.03	1.14	0.44
ind. ICR	0.90	0.89	0.03	0.89	0.03	0.90	0.02	0.89	0.03	1.61	0.02
ind. ICR	0.95	0.95	0.01	0.95	0.01	0.95	0.01	0.95	0.01	1.97	0.06
subp. ICR	0.75	0.61	0.04	0.61	0.04	-	-	0.61	0.04	1.06	0.03
subp. ICR	0.85	0.73	0.03	0.73	0.03	-	-	0.73	0.03	1.09	0.34
subp. ICR	0.90	0.81	0.04	0.81	0.04	-	-	0.81	0.04	1.42	0.05
subp. ICR	0.95	0.90	0.03	0.90	0.03	-	-	0.90	0.03	1.73	0.06

including Y for CR as well and draw ground-truth interventional samples from the SCM instead of identifying the interventional distribution from observational data.

For *3var-causal* and *3var-noncausal* we configured NSGA-II to optimize over 600 generations with a population size of 300, for *5var-skill* and *7var-covid* 1000 generations with 500 individuals were used. For all experiments the crossover probability was 0.3 and the mutation probability 0.05. For all settings continuous variables were rounded to 1 decimal point. For the 3 variable settings a standard `sklearn LogisticRegression` was used, for the refits without penalty. For the nonlinear dataset a `RandomForestClassifier` with max depth 30, 50 estimators and balanced subsampling was applied. The experimental results were computed on a Quad core Intel Core i7-7700 Kaby Lake processor. For each setting, the experiments took between 24 to 48 hours.

D Proofs

As follows we provide the full proofs for Propositions 1 - 5.

D.1 Linking individualized prediction with γ^{ind} , Proof of Proposition 1

Proposition 1. *The expected individualized post-recourse score is equal to the individualized improvement probability $\gamma^{ind}(x^{pre}, a) := P(Y^{post} = 1|x^{pre}, do(a))$, i.e.*

$$E[\hat{h}^{*,ind}(x^{post})|x^{pre}, do(a)] = \gamma^{ind}(a).$$

Proof: It holds that

$$\begin{aligned} & E[\hat{h}^{*,ind}(x^{post})|x^{pre}, do(a)] \\ &= E[E[Y|x^{pre}, x^{post}]|x^{pre}, do(a)] \\ \text{total exp.} &= E[Y|x^{pre}, do(a)] \\ &= \gamma^{ind}(a). \end{aligned}$$

D.2 Intervention stability w.r.t. ICR actions, Proposition 2

Proposition 2. *Given nonzero cost for all interventions, ICR exclusively suggests actions on causes of Y . Assuming causal sufficiency, for any optimal predictor the conditional distribution of Y given the variables that the model uses X_S (i.e. $P(Y|X_S)$) is stable w.r.t interventions on causes. Therefore, optimal predictors are intervention stable w.r.t. ICR actions.*

Proof: We prove the statement in six steps.

ICR only intervenes on causes: The goal of meaningful recourse is to improve Y with minimal cost. Only interventions on causes alter Y . Consequently, actions on non-causes of Y would not be suggested by meaningful recourse.

Given causal sufficiency, a graph \mathcal{G} and an endogenous Y , the set of endogeneous direct parents, direct effects and direct parents of effects are the minimal d -separating set $S_{\mathcal{G}}$: Standard result, see e.g. Peters et al. [2017], Proposition 6.27.

The set $S_{\mathcal{G}^}$ in the augmented graph \mathcal{G}^* coincides with $S_{\mathcal{G}}$:* The minimal d -separating set contains direct causes, direct effects and direct parents of direct effects. I_l is never a direct cause of X_l . Also, since I_l has no endogenous causes, it cannot be a direct effect. Furthermore, since we restrict interventions to be performed on causes, I_l cannot be a direct parent of a direct effect.

$S_{\mathcal{G}}$ is intervention stable: As follows, all intervention variables are d -separated from Y in \mathcal{G}^* by $S_{\mathcal{G}}$. Therefore $S_{\mathcal{G}}$ is intervention stable. An example is given in Figure 9.

Then also the markov blanket is intervention stable: Since d -separation implies independence $MB(Y) \subseteq S_{\mathcal{G}}$. Therefore, if $X_T \perp Y|X_{MB(Y)}$ then also $X_T \perp Y|S_{\mathcal{G}}$. If any element $s \in S_{\mathcal{G}}$ it holds that $s \notin MB(Y)$, then it must hold that $X_s \perp Y|X_{MB(Y)}$. Therefore, if $X_T \perp Y|X_{MB(Y)}$, X_s then also $X_T \perp Y|X_{MB(Y)}$ and therefore any independence entailed by $S_{\mathcal{G}}$ also holds for $MB(Y)$. Since Pfister et al. [2021] only require the independence that is implied by d -separation in their invariant conditional proof, the same implication holds for the $MB(Y)$. As follows, $P(Y|X_{MB(Y)})$ is invariant with respect to interventions on any set of endogenous causes.

Then any superset of the markov blanket is intervention stable: We prove the statement by contradiction. The markov blanket d -separates the target variable Y from any other set of variables. If adding a set of variables S_1 to the markov blanket would open a path to any other set of variables S_2 , then it would hold that $S := S_1 \cup S_2$ is not d -separated from Y ($P(Y|MB(Y)) = P(Y|MB(Y), S_1, S_2) \neq P(Y|MB(Y), S_1) = P(Y|MB(Y))$)

D.3 Linking observational prediction and γ^{sub} , Proposition 3

Proposition 3. *Given causal sufficiency and positivity²⁴, for interventions on causes the expected subgroup-wide optimal score h^* is equal to the subgroup-wide improvement probability $\gamma^{sub}(a) := P(Y^{post} = 1|do(a), x_{G_a}^{pre})$, i.e.*

$$E[\hat{h}^*(x^{post})|x_{G_a}^{pre}, do(a)] = \gamma^{sub}(a).$$

²⁴Positivity ensures that the post-recourse observation lies within the observational support, where the model was trained (i.e., $p^{pre}(x^{post}) > 0$), [Neal, 2020]).

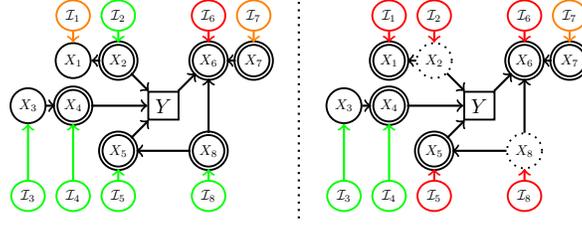


Figure 9: A schematic drawing illustrating under which interventions I_1, \dots, I_8 the Markov blanket (double circle) is intervention stable. In this setting, we consider the intervention variables to be independent treatment variables: We would like to know how the different actions influence the conditional distribution, irrespective of how likely they are to be applied. Therefore, they are modeled as parent-less variables. Green indicates no intervention stability, red indicates intervention stability. Orange indicates intervention stability of non-causal variables. Dotted variables are not observed. *Left*: Since all endogenous variables are observed, $MB_O(Y)$ is stable w.r.t. interventions on every endogenous cause of Y (Proposition 3). *Right*: Unobserved variables (X_2, X_8) open paths between interventions on causes and Y .

Proof: The proposition follows from Proposition 2. More specifically

$$E[h^*(x^{post,a})|x_G^{pre}, a] = E[E[Y|x^{post,a}]|x_G^{pre}, a] \stackrel{\text{total exp.}}{=} E[Y|x_G^{pre}, a] \stackrel{\text{def.}}{=} \gamma^{sub}(a). \quad (13)$$

D.4 Acceptance Bound, Proof of Proposition 4

Proposition 4. *Let g be a predictor with $E[g(x^{post})|x_S^{pre}, do(a)] = \gamma(x_S^{pre}, a)$. Then for a decision threshold t the post-recourse acceptance probability $\eta(t; x_S^{pre}, a) := P(g(x^{post}) > t | x_S^{pre}, do(a))$ is lower bounded:*

$$\eta(t; x_S^{pre}, a) \geq \frac{\gamma(x_S^{pre}, a) - t}{1 - t}.$$

Proof: Positivity ($p^{pre}(x^{post}) > 0$) is necessary for subpopulation-based ICR since only then we can assume that the model is actually optimal for any input that it receives. The problem is discussed in more detail in Hernan MA [2020], Neal [2020].

As follows we denote \hat{h}^* as the random variable indicating the predictions of the post-recourse predictors described in Section 5.

From Propositions 1 and 3, for both individualized and subpopulation-based post-recourse predictors we know that

$$E[\hat{h}(x^{post,a})^* | x_S^{pre}, do(a)] = \gamma(x_S^{pre}, a).$$

We decompose the expected prediction

$$\gamma(x_S^{pre}, a) = E[\hat{h}^* | x_S^{pre}, a] \quad (14)$$

$$= E[\hat{h}^* | \hat{h}^* > t]P(\hat{h}^* > t) + E[\hat{h}^* | \hat{h}^* \leq t]P(\hat{h}^* \leq t) \Big|_{x_S^{pre}, a} \quad (15)$$

$$= E[\hat{h}^* | \hat{h}^* > t]P(\hat{h}^* > t) + E[\hat{h}^* | \hat{h}^* \leq t](1 - P(\hat{h}^* > t)) \Big|_{x_S^{pre}, a} \quad (16)$$

$$= E[\hat{h}^* | \hat{h}^* > t]P(\hat{h}^* > t) + E[\hat{h}^* | \hat{h}^* \leq t] - P(\hat{h}^* > t)E[\hat{h}^* | \hat{h}^* \leq t] \Big|_{x_S^{pre}, a} \quad (17)$$

$$= E[\hat{h}^* | \hat{h}^* \leq t] + P(\hat{h}^* > t) \left(E[\hat{h}^* | \hat{h}^* > t] - E[\hat{h}^* | \hat{h}^* \leq t] \right) \Big|_{x_S^{pre}, a} \quad (18)$$

which can be reformulated to yield the acceptance rate η :

$$\frac{\gamma - E[\hat{h}^* | \hat{h}^* \leq t]}{E[\hat{h}^* | \hat{h}^* > t] - E[\hat{h}^* | \hat{h}^* \leq t]} \Big|_{x_S^{pre}, a} = P(\hat{h}^* > t | x_S^{pre}, a) = \eta(x_S^{pre}, a). \quad (19)$$

It holds that $E[\hat{h}^{*,ind}|\hat{h}^* \leq t] = FNR(t)$ and $E[\hat{h}^*|\hat{h}^* > t] = TPR(t)$.

We can show that $E[\hat{h}^*|\hat{h}^* \leq t] \leq t$:

$$0 \leq FNR(t|x_S^{pre}, a) \quad (20)$$

$$= P(Y^{a,post} = 1|h^* \leq t, x_S^{pre}, a) \quad (21)$$

$$= E[Y^{a,post}|h^* \leq t, x_S^{pre}, a] \quad (22)$$

$$= E[E[Y^{a,post}|x^{post,a}]|h^* \leq t, x_S^{pre}, a] \quad (23)$$

$$= E[h^*|h^* \leq t, x_S^{pre}, a] \quad (24)$$

$$\leq t \quad (25)$$

and analog that $1 \geq TPR(t) \geq t$. Therefore

$$\eta(t, x_S^{pre}, a) = \frac{\gamma - FNR(t)}{TPR(t) - FNR(t)} \Big|_{x_S^{pre}, a} \geq \frac{\gamma(x_S^{pre}, a) - FNR(t)}{1 - FNR(t)} \geq \frac{\gamma(x_S^{pre}, a) - t}{1 - t}. \quad (26)$$

D.5 Individualized post-recourse prediction, proof of Proposition 5

Proposition 5. *In general, the individualized post-recourse predictor can be estimated as*

$$p(y^{post}|x^{pre}, x^{post}, do(a)) \quad (27)$$

$$= \frac{\int_{\mathcal{U}} p(y^{post}, x^{post}|u, do(a))p(u|x^{pre})du}{\sum_{y' \in \{0,1\}} (\int_{\mathcal{U}} p(y', x^{post}|u, do(a))p(u|x^{pre})du)} \quad (28)$$

Given binary decision problems with invertible structural equations, the individualized post-recourse prediction function reduces to

$$p(y^{post}|x^{post}, x^{pre}, do(a)) \quad (29)$$

$$= \frac{p(U_{-I} = f_{do(a)}^{-1}(y^{post}, x^{post})|x^{pre}, do(a))}{\sum_{y' \in \{0,1\}} p(U_{-I} = f_{do(a)}^{-1}(y', x^{post})|x^{pre}, do(a))}. \quad (30)$$

Proof: It holds that

$$p(y^{post}|x^{pre}, x^{post}, do(a)) \stackrel{\text{def. cond.}}{=} \frac{p(y^{post}, x^{post}|x^{pre}, do(a))}{p(x^{post}|x^{pre}, do(a))} \quad (31)$$

$$(32)$$

We can reformulate the conditional distribution $p(y^{post}, x^{post}|x^{pre}, do(a))$ as two parts, one that describes the probability of a state of the context given x^{pre} , and one that describes the probability of a post-recourse state x^{post}, y^{post} given a certain noise state u and $do(a)$.

$$p(y^{post}, x^{post}|x^{pre}, do(a)) \quad (33)$$

$$\stackrel{\text{marginal.}}{=} \int_{\mathcal{U}} p(y^{post}, x^{post}, u|x^{pre}, do(a))du \quad (34)$$

$$\stackrel{\text{chain rule}}{=} \int_{\mathcal{U}} p(y^{post}, x^{post}|u, x^{pre}, do(a))p(u|x^{pre})du \quad (35)$$

$$(y, x)^{post} \perp x^{pre} | u \int_{\mathcal{U}} p(y^{post}, x^{post}|u, do(a))p(u|x^{pre})du. \quad (36)$$

In combination we yield

$$p(y^{post}|x^{pre}, x^{post}, do(a)) \quad (37)$$

$$= \frac{\int_{\mathcal{U}} p(y^{post}, x^{post}|u, do(a))p(u|x^{pre})du}{\int_{\mathcal{Y}} (\int_{\mathcal{U}} p(y', x^{post}|u, do(a))p(u|x^{pre})du) dy'} \quad (38)$$

$$= \frac{\int_{\mathcal{U}} p(y^{post}, x^{post}|u, do(a))p(u|x^{pre})du}{\sum_{y' \in \{0,1\}} (\int_{\mathcal{U}} p(y', x^{post}|u, do(a))p(u|x^{pre})du)} \quad (39)$$

For a setting with invertible structural equations this reduces to

$$p(y^{post}|x^{post}, x^{pre}, do(a)) \tag{40}$$

$$= \frac{p(y^{post}, x^{post}|x^{pre}, do(a))}{p(x^{post}|x^{pre}, do(a))} \tag{41}$$

$$= \frac{p(U_{-I} = f^{-1}(y^{post}, x^{post})|x^{pre}, do(a))}{\sum_{y' \in \{0,1\}} p(U_{-I} = f^{-1}(y^{post}, x^{post})|x^{pre}, do(a))}. \tag{42}$$

where $-I$ is the index set for variables that have not been intervened on (since the noise terms for the intervened upon variables are isolated variables in the interventional graph).

E Misc

E.1 Negative Result: Algorithmic recourse is neither meaningful nor robust

In the introduction we claimed that CR recommendations [Karimi et al., 2020b, 2021] may not lead to improvement. Now, we formally demonstrate the case on the Covid hospital admission example (Figure 1) which we extend with the full structural causal model (Example 1). Furthermore, we show that CR is not robust to refits of the model on mixed pre- and post-recourse data. All code is publicly available via <https://anonymous.4open.science/r/icr-aaai/README.md>.

Example 1. Let V indicate whether someone is fully vaccinated, Y indicate whether someone is free of Covid and S whether someone is asymptomatic. The data is generated by the following structural causal model (SCM) entailing the causal graph depicted in Figure 1:

$$V := U_V, \quad U_V \sim \text{Bern}(0.5) \quad (43)$$

$$Y := V + U_Y \text{ mod } 2, \quad U_Y \sim \text{Bern}(0.09) \quad (44)$$

$$S := Y + U_S \text{ mod } 2, \quad U_S \sim \text{Bern}(0.05) \quad (45)$$

For prediction, a `sklearn` logistic regression model is fit on 2000 samples, yielding \hat{h} with $\beta_v \approx 3.7$, $\beta_s \approx 5.1$, $\beta_0 \approx -4.3$. Visitors are allowed to enter the hospital if $\hat{h} < 0.5$. Intervening on (flipping) V and S costs 0.5 and 0.1 respectively.

Lack of improvement: Given a decision threshold of 0.5, the model admits everyone without symptoms ($S = 1$), irrespective of their vaccination status V . Therefore, in order to revert rejections ($S = 0$), both individualized and subpopulation-based CR suggest removing the symptoms S (*do*($S = 1$), for instance by taking cough drops). However, since they only treat the symptoms S , the actual Covid risk Y is unaffected: none of the recourse-implementing individuals actually improve. We say the predictor is *gamed*.

Lack of robustness: For individuals who implement recourse the association between symptom state S and Covid risk Y is broken. Thus, the predictive power of the model for recourse-seeking individual drops from ≈ 95 percent pre-recourse to ≈ 5 percent post-recourse.²⁵ A refit of the model on a mix pre- and post-recourse data (2000 samples each) yields \hat{h} with $\beta_V \approx 4.1$, $\beta_S \approx 3.3$, $\beta_0 \approx -4.8$. Since the association between symptom state and disease status is broken post-recourse, the new model rejects individuals if they are not vaccinated, irrespective of their symptom state. For that reason, recourse recommendations that were designed for the original model only lead to acceptance by the refitted model for those individuals who happened to be vaccinated anyway.

The example demonstrates that CR recommendations are prone to gaming the predictor and therefore may neither lead to improvement nor be robust to model refits.

E.2 Interpretability of improvement confidence γ

Counterfactuals are concerned with changing the inputs to the model such that the model prediction changes in the desired way. Since the prediction function is deterministic and accessible, the post-recourse prediction can be determined exactly.

In contrast CR and ICR deal with the effects of real-world interventions on real-world variables. As such, the effects of recourse actions on the covariates (and the underlying prediction target) cannot be determined exactly. Therefore both CR and ICR have to deal with uncertainty.

CR deals with this uncertainty by phrasing the optimization objective for CR in terms of an expectation over the prediction distribution and by using an action-adaptive confidence threshold. This threshold bounds the expected prediction away from the model’s decision threshold (e.g. $t = 0.5$). Using the conservativeness parameters, the user can roughly steer how far the expected prediction shall be away from the decision boundary.

In contrast, ICR deals with the uncertainty by letting the user specify the confidence γ , which can be intuitively interpreted as improvement probability (whereas the expected prediction cannot be interpreted as acceptance probability). A lower-bound on the acceptance probability for a combination of γ and t is given in Proposition 4. Furthermore, we can estimate the individualized and subpopulation-based acceptance rates for a specific situation (a, x^{pre}) as detailed in B.1 and B.3. The human-interpretable improvement and acceptance confidences are vital for the explainee to make an informed decision.

In order to allow a direct comparison of the methods, we rephrase the CR objective to optimize the acceptance probability η in our experiments.

²⁵The previously wrongly-rejected individuals are correctly classified after implementing recourse.

E.3 Imbalance between standard predictors and individualized ICR recommendations

In Section 6 we argued that there is an imbalance in predictive capability between (optimal) observational predictors and the pre-recourse SCM (which used to predict γ^{ind}). We illustrate the problem on a simple example.

Example 2. *Let there be a three variable chain $X_1 \rightarrow Y \rightarrow X_2$ where at every step the value is incremented by one with 50% chance and the maximum value is set to 2 ($X_1 := U_1, Y := X_1 + U_Y, X_2 := \min(2, Y + U_2)$ where $U_1, U_2, U_Y \sim \text{Bern}(0.5)$). Let us assume a factual observation $x^{pre} = (0, 2)$ and action $a = do(X_1 = 1)$ yielding $x^{post} = (1, 2)$. For the observation $x^{pre} = (0, 2)$ we can infer that U_Y must have been 1, since two increments are needed to get from 0 to 2. However, from the post-intervention observation $x^{post} = (1, 2)$ we cannot infer where the increment happened (U_Y or U_2). As a consequence, an optimal predictive model that only has access to x^{post} would predict that y^{post} for $x^{post} = (1, 2)$ could be 1 or 2 with equal likelihood. In contrast, with access to x^{pre} and the SCM we can infer that $y^{post} = 2$ since $U_Y = 1$.*

In the above example, given knowledge of the SCM, the pre-intervention observation x^{pre} and the performed action a we can already abduct U_Y perfectly and therefore correctly determine the post-intervention state of Y (even without access to the post-intervention observation x^{post}). In contrast, with the post-recourse observation alone it is impossible to reconstruct U_Y and therefore impossible to determine the post-intervention state of Y .²⁶ In the context of ICR this means that the observational predictor's post-recourse predictions are not directly linked with γ : they may not honor the implementation of actions with $\gamma^{ind} = 1$. As a consequence, we suggested to use the SCM for post-recourse prediction in Section 6.

²⁶The optimal pre-recourse predictor $\hat{h}^*(x^{post})$ predicts 0.5 for both $y = 1$ and $y = 2$.