

# Mortality prediction models, causal effects, and end-of-life decision making in the intensive care unit

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End-of-life (EOL) decision making in the intensive care unit (ICU) is challenging for both families and clinicians. This decision-making process is ideally framed around a shared understanding of a patient's values and goals, all taken in the context of their critical illness and prognosis. However, clinicians commonly face uncertainty regarding prognosis and may have difficulty offering families an accurate assessment of the likely outcomes of treatment decisions. Adding to the complexity of these scenarios, clinicians, patients and families are each susceptible to unconscious but influential cognitive biases when making decisions under stress. Given these challenges, and a rapidly growing interest in data science to inform care in the ICU, investigators have explored the use of prediction models (eg, machine learning or ML algorithms) to assist with prognostication.<sup>1-3</sup> Prediction models describe an outcome distribution among individuals with a particular set of characteristics, such as risk of acute kidney injury among individuals with particular laboratory values and clinical characteristics in a population. However, they do not compare how that outcome distribution would change were different treatment decisions made in that population—this requires causal effect estimation, rather than prediction modelling. Herein, we explain why prediction modelling alone is not sufficient to inform many ICU treatment decisions, including EOL decision making, and describe why causal effect estimation is necessary.

Consider the following case in which a prediction model is used, rather than causal effect estimation: a 68-year-old man is admitted to the ICU with severe pneumonia and requires mechanical ventilation. After 5 days, he requires continued full support from the ventilator and has developed delirium. His family is concerned about prolonging intensive care but worries about transitioning to

comfort measures prematurely if continued intensive care could, in fact, achieve their goals. His family and clinicians would like to use the best available evidence to inform their decision. Given an increasing interest in mortality prediction models, the clinical team explores this as a tool for decision support. Their chosen algorithm is a prediction model—trained on available data containing measurements of treatments, outcomes and other characteristics of previously ventilated patients—which returns a 70% probability of death within the next 30 days.

The first question we must ask is: what is the precise interpretation of this prediction? This is an estimate of the probability of death for a population of patients 'like this patient'—patients mechanically ventilated for 5 days with similar baseline characteristics and clinical risk factors up to that moment in their ICU course. Importantly, this probability is contingent on the treatment decisions, after day 5, that were made in the population from which the training data came. For example, if all patients similar to this one in the training data transitioned after day 5 to comfort measures only, then the algorithm would predict a 100% risk of 30-day mortality. Alternatively, if most of the patients similar to this index case in the training data frequently pursued tracheostomy, the model may predict a low 30-day mortality. These two extreme conditions demonstrate that the interpretation is highly dependent on the distribution of treatment decisions among patients 'like this patient' in the training data.

Second, and more importantly, we consider the following question: how is this probability useful (or not useful) to both the family and the clinicians? We must first clearly articulate the question that clinicians and family members are truly interested in answering for this patient. They are not simply concerned about predicting if survival is possible. Instead,

they are interested in knowing: what would the outcome be if one treatment strategy was chosen compared with the outcome if a different treatment strategy was chosen. To be more concrete, they may want to know if continued mechanical ventilation and attempts at ventilator liberation for another week would result in a different 30-day survival than a more limited trial of 48 hours of ventilation. They are concerned with the balance between unnecessarily prolonging intensive care versus a missed opportunity for survival if they transition to comfort measures prematurely. More importantly, in the context of the patient's values and goals, they likely want to understand the effect of these treatment strategies on long-term quality-of-life.

### CONTRASTING CAUSAL EFFECTS WITH PREDICTION MODELS

We refer to the outcome that would have been observed had, perhaps contrary to fact, a particular treatment been given, as the *counterfactual outcome* under that treatment. Using our earlier clinical scenario as an example, if our critically ill patient had decided to undergo tracheostomy on day 7, we might wonder 'what would have happened if he did not undergo tracheostomy and instead remained intubated'. The outcome under our 'what if' scenario is contrary to what actually happened, that is, the counterfactual outcome. *Causal effects* compare counterfactual outcomes for a person (or a population) under different treatment strategies, asking 'what would be the outcome if we choose treatment A compared to the outcome if we choose treatment B'. Clinicians, patients and families intuitively think in counterfactuals when weighing the risks and benefits of different decisions, including EOL decision making. Thus, causal estimates would seem to be the natural approach to support decision making. Yet prediction models, rather than causal estimates, have received rapidly growing attention in the literature while their limitations are often overlooked.

In contrast to causal estimates, mortality prediction models are mapping inputs (or 'features') to a chosen outcome, such as mortality. They might help estimate if a patient is at a higher risk of death, but they offer little help in making the best decision in that scenario. However, as we noted previously, these estimates depend on the distribution of treatments that were given to patients like ours in the training data. As such, if historic treatment distributions differ from those in current practice, then the prediction will be inaccurate.

The appeal of predictive approaches, from a data science perspective, is that they can be readily applied with existing healthcare data and established machine learning algorithms. However, these models ignore assumptions about causal structure—the relationship between the variables that can only be informed by expert knowledge. For example, users of a prediction model may claim that a high fraction of inspired oxygen is associated with higher ICU mortality but that prediction could not justify a claim that an intervention to reduce the fraction

of inspired oxygen administered would reduce mortality without first reasoning about how these variables are connected to one another.<sup>4</sup> Specifically, they must defend assumptions about how the treatment and outcome are related, causally or by associational pathways. Because prediction models omit this step, they cannot provide an estimate of expected outcomes when divergent treatment decisions are chosen.

### APPLYING CAUSAL INFERENCE TO ICU DATA

Having established the need for causal effects of different ICU decisions, rather than predictions of mortality, we will describe how they can be estimated. An intuitive and effective approach to designing observational analyses for estimation of causal effects is to specify a hypothetical pragmatic randomised trial (ie, a 'target trial'), one that would answer the question of interest but may be impossible or impractical to conduct in practice.<sup>5</sup> This hypothetical trial helps us be explicit about the important aspects of our analysis, including the causal question it aims to answer and avoid biases introduced by the study design (eg, immortal time bias).<sup>6</sup> Specifically, we need to define the eligibility criteria, the treatment strategies of interest, the follow-up period (including a clear definition of 'time zero', the start of follow-up, eg, mechanical ventilation day 5 in the above example), the outcomes of interest and the statistical analysis plan. This also requires expert knowledge of the clinical context.

We describe one such trial, for example, in [table 1](#). This trial is ethically and logistically infeasible; therefore, an analysis of observational data, designed with identical features as the trial, is the next best approach. In particular, to emulate the trial described in [table 1](#), we would, after obtaining appropriate observational data: (1) restrict our data to individuals meeting the eligibility criteria, (2) classify those who immediately discontinue mechanical ventilation as adherent to strategy one and those continuing mechanical ventilation on day 6 as adherent to strategy two and (3) compare estimates of the risk of 30-day mortality among those adherent to strategy one versus strategy two, adjusted for measured prebaseline prognostic factors (ie, *measured confounders*). Adjustment is required because treatment is not randomly assigned in observational data (in other words, treatment is related to the outcome via associational pathways). If all the relevant confounders are measured and adjusted for, then the same effect estimates will be obtained from the observational data analysis as from the trial had it been conducted (except for random variation).

While this may be a useful example for the technical aspects of this process, the trial described in [table 1](#) is not the one of interest to decision makers. For example, treatment strategies such as 'continue mechanical ventilation for another week, unless liberation from the ventilator is achieved' versus 'continue mechanical ventilation for another 48 hours, unless liberation from the ventilator is achieved' address questions around time-limited trials





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