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Prompt admission to the ICU: an instrument to improve mortality for deteriorating ward patients

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Introduction
In a recent article in Intensive Care Medicine, Harris and co-authors present a study on the importance of prompt transfer of deteriorating ward patients to the ICU [1]. Between 2010 and 2011, they conducted a large observational study involving 49 hospitals in the UK considering the problem of ongoing bed block causing unwanted delays in the admission of patients. We would like to congratulate the authors on conducting such challenging and groundbreaking study designed to answer a valid but also somewhat delicate research question. A key feature of this study is the applied instrumental variable (IV) analysis, which we believe has a non-harvested potential within critical care research.

Do the efforts for a fast ICU transfer make a difference?
The assessment of deteriorating ward patients is a key part of the daily work for every intensivist. This involves many complicated questions: “Will this patient benefit from ICU admission?”, or “How many free beds do we have in the ICU now and later?”. Indeed, triage decisions are embedded in the complex environment of each hospital, where the ICU capacity and the efficiency on admitting or discharging patients without delays are crucial [1–3]. When the decision for an ICU transfer is made, the ward, rapid response and ICU teams usually feel relieved when the process is fast. However, in real life, there are multiple (expected and unexpected) bottlenecks that need to be overcome [4–6].

To randomize to prompt or “watchful waiting” ICU admission for deteriorating ward patients is likely to lack clinical equipoise and therefore difficult to study with a randomized design [7]. Thus, Harris and co-authors settled on an intelligent cohort study, starting from ward patients referred for assessment by critical care teams. They followed these patients prospectively, including those recommended and those not recommended for ICU admission, and were able to capture outcomes for reviewed patients by linking data specifically collected in this study to the Intensive Care National Audit & Research Centre (ICNARC) database.

In the analysis, the authors applied IV methodology to estimate the causal effect of prompt transfers (< 4 h of assessment) compared to what they defined as a “watchful waiting” cohort. The authors observed that prompt admissions reduced 90-day mortality in the whole cohort by 7.4% (95% CI 1.7, 18.5%) and by 16.2% (95% CI 1.1, 31.3%) in those recommended for immediate ICU admission.

Although the study and the IV method have some limitations as discussed in the manuscript, the (SPOT)light results highlight the need for optimization of the patient’s flow and ICU efficiency. The authors observed that transfer delays are frequent, related to ICU strain and associated with worse outcome. The solutions are not easy and require efforts from several players within the hospital. The current study, especially if replicated in other settings, highlights the need for more research on ICU triage and discharge decisions. Solutions could include incorporating concepts from other fields, such as engineering and logistics [8].
**IV method: what it is and which potentials?**

A randomized controlled trial (RCT) is the gold standard for drawing a causal conclusion because its design ensures that treatment allocation does not depend on any known or unknown patient characteristics. In contrast, observational studies do not have randomization to ensure treatment groups are comparable, increasing the possibility of confounding. Accordingly, an attempt is made to collect information on as many potential confounders as possible with adjustment for imbalances using statistical methods. However, it will always be a matter of faith if we trust that enough confounders have been collected. Presented like this, it appears that the two study designs are completely distinct, but the truth is that they are just in the opposite ends of a continuum of study designs where we have more or less control of how treatment is allocated (100% control in RCT; 0% in the other end). Re-examining the RCT reveals that the important part is not that we have 100% control of the treatment allocation mechanism, but instead it is that we by design introduce a component in the treatment allocation which is unrelated to characteristics of the included patient. This is sometimes called an exogenous push (i.e. the instrument).

The key idea in IV analyses is that this exogenous push could also come from sources other than randomization. Harris et al. use the fact that overall bed occupancy in the ICU (the instrument) will affect the probability of prompt admission to ICU irrespective of the specific clinical condition of that patient. Or phrased differently: there are patients who had they been presented on a day with many open beds in the ICU would have been given prompt admission, but had the same patients been presented on a day with few free beds in the ICU they would not be given prompt admission. The link between randomization and IV appear clear in this setting. Methods for handling non-compliance in RCTs [9–11] are also a case of IV analyses.

The precise details on IV estimation depend on outcome and instrument type; see Greenland [12] for an introduction, but the basic idea is always the same: (1) estimate the causal effect of the instrument on the outcome. This effect can be thought of as the multiplication of the causal effect of the instrument on the treatment and the causal effect of the treatment on outcome. (2) Estimate the causal effect of the instrument on the exposure. (3) Divide the estimate in one by the one from two to produce an estimate of the causal effect of treatment on exposure. The method only works under the (untestable) assumption that the IV only affects the considered outcomes through its effect on treatment. With bed occupancy as the instrument, this assumption is
not trivial as one could fear that treatment quality in the ICU is reduced in conditions of high bed occupancy; this potential problem is, however, addressed in the paper. No knowledge of confounding factors between prompt admission and survival is required. We only need to know that there are no confounding factors between instrument and exposure and instrument and outcome; see Fig. 1 for a graphical illustration of the assumptions.

The statistical methods for conducting IV analysis are still developing (see e.g. Tchetgen Tchetgen [13] for survival outcomes) and they are a bit harder to implement than standard regression. However, these are just technicalities which should not distract us from the essential point: the analysis by Harris illustrates how IV methods can replace the need for detailed clinical information on each patient with simple administrative data on hospital utilization. The latter data is much “easier” to come by than the first (without in anyway belittling the work done by the authors). The new requirement is that researchers must think outside the usual suspects of variables influencing treatment allocation. We believe that such variables are common (e.g. occupancy, administrative changes, changes in funding of wards, changes in uptake areas, etc.) and we encourage our colleagues to “go look for that exogenous pushing”.

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Compliance with ethical standards

Conflicts of interest
The authors declare that they have no conflicts of interest.

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