Tailoring treatments to older people in intensive care. A way forward

Márlon Juliano Romero Aliberti1,2*, Sébastien Bailly3 and Matthew Anstey4,5,6

The number of older adults requiring critical care has increased worldwide as the population ages. People aged 80 years and older represent the fastest growing subgroup in intensive care unit (ICU) admissions. In high-income countries, for example, they already constitute 15% and 20% of all ICU admissions [1–3]. Increasing evidence suggests that outcomes of older patients in ICU are not as poor as one might initially expect. Nonetheless, the appropriateness of admission to ICU remains context-specific. Where there are resource limitations and competing priorities for those beds, it would be better if decision-making incorporated objective parameters to answer core questions, like “Will this older patient benefit from ICU resources?” or “How to best care for this older patient with a critical illness?” Although we do not have the perfect answers to these questions, recent advances regarding older ICU patients are notable [4].

In older people, age-related conditions—frailty, cognitive impairment, functional disability, sensory deficits, comorbidities, polypharmacy—capture an overall baseline vulnerability, pivotal in estimating the patient’s ability to cope with acute insults (e.g., emergency surgery, septic shock) [5–7]. Such conditions share similar pathophysiology mechanisms, including neuromuscular weakness, decreased oxygen utilization, increased inflammation, and immunosenescence, all important in the ICU context. Thus, unlike robust older patients, those with high baseline vulnerability are susceptible to multi-system organ failure and adverse outcomes when facing an acute insult [7–9]. That is why ‘biological age’ differs from ‘chronological age’ and is essential for decision-making in older patients [4–6].

Heterogeneity is a hallmark of the older population, making it challenging to identify early—among older patients of similar age—those more likely to benefit from ICU resources and those in whom escalating acute care would be useless [5–7]. Physiological changes that affect vital signs (e.g., blunted heart rate response to stress) and laboratory findings (e.g., lower serum creatinine) predispose to atypical presentations of acute diseases as people age, particularly in those accumulating frailty and other age-related conditions [10]. Consequently, traditional prognostic tools [the Sequential Organ Failure Assessment (SOFA) score and Simplified Acute Physiology Score (SAPS) II] alone can also misclassify illness severity in older patients [5].

In this issue of Intensive Care Medicine, Mousai and colleagues offer cutting-edge knowledge on this topic [11]. These authors examined the heterogeneity of people aged 80 years and older in critical care by clustering analysis of bedside clinical variables assessed on admission to ICUs in 22 European countries, integrating the VIP2 cohort study [5]. Besides demographic factors, reasons for ICU admission, and SOFA scores, the authors considered information on age-related conditions, which allowed them to identify seven clusters with distinct phenotypes (Fig. 1) [11]. Notably, Mousai and colleagues confirmed the validity of the clusters by providing external validation in another dataset comprising critical care patients affected by coronavirus disease 2019 (COVID-19) (the COVIP study) and by incorporating initially excluded patients with limitations of life-sustaining treatment into sensitivity analyses [7].
For many years, ICU admissions in people aged 80 years and older were a synonym for suffering and moral distress in patients and families [4]. However, this view has proven to be superficial, as Mousai and colleagues showed while reporting divergent outcomes across the clusters (Fig. 1) [11]. While 30-day mortality reached 57% in the VIP2 patients with age-related conditions and high SOFA scores, it was only 7% among the oldest old patients presenting very mild age-related impairments and low SOFA scores. Other specific profiles denoting respiratory, cardiac, and renal failure had a 30-days mortality rate of around 10%, with only the phenotype presenting age-related conditions approaching nearly 20% of mortality risk, despite low SOFA scores [11].

Such findings corroborate practical implications. Experienced intensive care clinicians understand that age can attract frailty and other age-related conditions, and those latter factors are responsible for limited reserves to survive critical illness. It would not surprise that a frail patient with a high SOFA score has a very high risk of dying (cluster G). A frail patient in ICU without such a high SOFA score still has a considerable risk of dying (cluster F). On the contrary, an otherwise robust octogenarian (cluster A) or nonagenarian (cluster C), without an illness with high SOFA scores, is highly likely to survive an ICU admission. Mousai and colleagues innovated by presenting this whole picture as distinct phenotypes (Fig. 1), which could aid clinicians in conversations at the bedside on goals of care with patients and families (i.e., advanced care planning or decisions on withdrawing or withholding ICU treatments) [11–13].

Personalized medicine implies considering each patient individually to be able to manage specifically their health care. A midpoint between generalizing all ICU older patients and viewing each patient individually is to identify homogenous sub-groups constituted by patients who share similar clinical characteristics at a time point. In this way, cluster approaches aim to minimize the differences between similar individuals and maximize the differences between individuals with distinct features [14]. However, clustering is not an exact science as results depend on data and methodological approaches. For example, clusters F and G comprised 10% and 30% of patients living in nursing homes, respectively, subgroups that would often not be appropriate for ICU admissions. Although observed clusters identified by Mousai and colleagues can be considered solid sub-groups of older patients in the ICU, their findings in Europe lack generalisability to other ICU

Fig. 1 Distinct clinical and prognostic profiles of older patients from the VIP2 cohort
populations and contexts [11]. Furthermore, their findings were based on ICU and 30-day mortality, arguably very short-term outcomes for this cohort of patients. Future research should focus on patient-centred outcomes, as quality of life and functional measures are the most relevant health aspects for older ICU patients [12].

In conclusion, Mousai and colleagues leveraged routine and vulnerability measures that were obtained at the bedside to describe clusters of older patients sharing similar clinical and prognostic profiles on ICU admission [11]. The next step is looking at which interventions might be appropriate for the different clusters and which interventions might not be. This may involve not admitting older patients with high frailty and SOFA scores to ICU. It may involve providing short-term vasopressor support but not mechanical ventilation or dialysis to other patients. In the meantime, we should acknowledge that age is insufficient reasoning [15]—it is time to integrate geriatric principles into critical care medicine.

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Author details
1 Laboratorio de Investigigacao Medica em Envelhecimento (LIM-66), Servico de Geriatria, Faculdade de Medicina, Hospital das Clínicas HCFCMUSPUniversidade de Sao PauloClínicas Médica, Av. Dr. Eneas de Carvalho Aguiar 155, 8º Andar, Sao Paulo, SP 05403-000, Brazil. 2 Research Institute, Hospital Sírio-Libanês, Sao Paulo, Brazil. 3 Grenoble Alpes University, Inserm, U1300, Grenoble Alpes University Hospital, Grenoble, France. 4 Department of Intensive Care, Sir Charles Gardiner Hospital, Perth, Australia. 5 School of Medicine, University of Western Australia, Perth, Australia. 6 School of Public Health, Curtin University, Perth, Australia.

Author contributions
MJRA, SB, and IMA were substantially involved in drafting the manuscript. All authors critically revised the manuscript and approved the final version of the article.

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