

Editorial Commentary: Community-acquired pneumonia, comparison of three mortality prediction scores in the emergency department

Comentario editorial: Neumonía adquirida en la comunidad, comparación de tres puntajes de predicción de mortalidad en el departamento de emergencias

Lena Barrera^{1,2} 
lena.i.barrera@correounivalle.edu.co



OPEN ACCESS

Citation: Barrera L. Editorial Commentary: Community-acquired pneumonia, comparison of three mortality prediction scores in the emergency department. *Colomb Méd (Cali)*, 2022; 53(3):e1015377
<http://doi.org/10.25100/cm.v53i3.5377>

Received: 08 Aug 2022

Revised: 11 Aug 2022

Accepted: 19 Aug 2022

Published: 20 Sep 2022

Copyright: © 2022 Universidad del Valle



Conflict of interest:

None

Corresponding author:

Lena Barrera MD. Msc. PhD.
Profesora Asociada. Departamento de Medicina Interna. Universidad del Valle-Cali (Colombia). **E-mail:** lena.i.barrera@correounivalle.edu.co

1 Universidad del Valle, Facultad de Salud, Escuela de Medicina, Departamento de Medicina Interna. Cali, Colombia. **2** Editor asociado, Revista Colombia Medica. Cali, Colombia,

Related article: <https://colombiamedica.univalle.edu.co/index.php/comedica/article/view/4287/5097>

Successive measurements of global disease burden have documented that lower respiratory tract infections, including pneumonia, are among the top 10 causes disability adjusted life-years and in 2019 pneumonia was the fourth cause of mortality for all ages¹. In Colombia, acute respiratory infections are the leading cause of mortality within the group of infectious diseases, 52.3% of the total reported between 2005 and 2019². Noteworthy, the COVID-19 epidemic increased the impact of respiratory tract infections on the global disease burden, with estimates of 18 million excess deaths during the period January 2020 to December 2021 worldwide³.

The assessment of an adult with pneumonia or suspected pneumonia demands the identification of the likelihood of death and/or hospitalization. Several scales have been constructed to estimate this probability to improve the predictive capacity of clinical evaluation. Among these scales, the CRB-65 and the CURB-65 stand out; being the first one recommended for use with clinical criteria, and the second one when laboratory data such as urea nitrogen are available⁴. Additionally, for individual with sepsis there have been developed to predict mortality such as SOFA (Sequential Organ Failure Assessment) and more recently the qSOFA (quick SOFA) which has an accurate prediction of mortality in this population^{5,6}.

Hincapié C et al assessed the CURB-65, CRB-65 and SOFA scales to predict mortality and/or admission to the intensive care unit in adults with pneumonia in three cohorts of patients admitted in three medium- and high-complexity hospitals in the city of Medellín-Colombia⁷. The study included 1,110 patients with suspected pneumonia who were identified in the emergency department and followed up until discharge and/or death. The authors found that the highest discrimination capacity, measured by the ROC curve, for the outcome hospitalization in an intensive care unit was 0.61, 0.58 and 0.59 for the CURB-65, CRB-65 and SOFA, respectively. In relation to mortality, the ROC found was 0.66, 0.63, and 0.63 for CURB-65, CRB65, and SOFA, respectively. The calibration was appropriate, that is, the ability to predict mortality and admission to the intensive care unit e for the three scales. Some readers have expressed their disagreement with the possible limited use of the scales, particularly the CURB-65 and the CRB-65 in the evaluation of an adult patient with pneumonia expressed by the authors.

The diagnosis of pneumonia both in the context of emergency care and in outpatient services is a challenge due to the heterogeneity of the clinical picture. The sensitivity of the clinical diagnosis has been reported between 45% and 69%⁸⁻¹⁰. Age, the immune status of the person, and the type of germ are the conditions most related to variability in clinical presentation¹¹. Hincapié et al.⁶, report that the diagnosis was based on the data obtained from the clinical records and their verification by the researchers. In this analysis, it would be useful to know the concordance between the diagnosis assigned by the researchers and the one established by the treating group, as well as the diagnostic capacity of the criteria used in the study⁷. A discrepancy in favor of a greater diagnostic capacity used by the researchers would suggest that the treating group did not identify the diagnosis early and therefore delays in the care received, such as the start of antibiotics. These delays would lead to a potentially greater probability of complications¹² and consequently leave in the sample a population composed mainly of people in the highest levels of severity, which could explain the low discriminative capacity of the scales found in the study.

In a complementary way, the validation of a score requires a sample that meets at least two characteristics: the occurrence of the outcome close to the real value and the representation of groups with different levels of risk within the cohort^{13,14}. Hincapié et al.⁶, found the mortality rate between 17% and 33% in the three cohorts⁷. This mortality is similar to that identified by Narvaéz P et al.¹⁵, who found a mortality rate of 20% in patients hospitalized with pneumonia secondary to pneumococcus in the city of Bogotá but it is higher than that observed in the cohorts from which CURB-65 and CRB-65 were derived, 7%⁴. Again, this difference could indicate that the population included in the study would be mostly in the groups with the highest severity and therefore the small number of patients with low scores did not allow to assess correctly their discriminative capacity but it did allow for the calibration^{4,16}. The authors could clarify this comment by sharing the distribution of the population included based on the categories established in the CURB-65 and CRB-65 scales.

The limited discriminatory capacity of the CURB-65 and CRB-65 scores has also been reported by other authors¹⁶. Aujesky et al.¹⁴, found that the ability to identify patients at low risk of mortality was greater using the PSI (Pneumonia Severity Index) scale compared to the CURB-65. Consequently, the guideline for the diagnosis and treatment of reported acquired pneumonia prepared by the American Society of Infectious Diseases recommends the use of the PSI to guide management. However, the NICE (National Institute for Health and Care Excellence) guide of the United Kingdom continues to recommend the use of the CURB-65, which was derived from cohorts identified in the health services of that country¹⁶. This corroborate the well-known need to validate the performance of scales in each population when using them in populations different from which they were derived. The adequate representation of the entire spectrum of the disease in the sample studied, as well as the influence of the interventions, determines their utility for a particular population.

The study by Hincapié et al.⁶, illustrates the variability in the predictive performance of the CURB-65 and CRB-65 scales and suggests the need to improve the early diagnosis of community-acquired pneumonia⁷. The potential limitation in the discriminative capacity of these scales could be clarified in a cohort assembled with a population treated at different levels of care in such a way that all the spectrum of the pneumonia condition can be included.

References

1. Roth GA, Mensah GA, Johnson CO, Addolorato G, Ammirati E, Baddour LM, et al. Global burden of cardiovascular diseases and risk factors, 1990-2019: update from the GBD 2019 Study. *J Am Coll Cardiol*. 2020; 76(25): 2982-3021. doi: 10.1016/j.jacc.2020.11.010.
2. Ministerio de Salud y Protección Social. Análisis de Situación de Salud, Colombia 2019. Bogotá, Colombia: Ministerio de Salud y Protección Social, Dirección de Epidemiología y Demografía; 2019. Disponible en: <https://www.minsalud.gov.co/sites/rid/Lists/BibliotecaDigital/RIDE/VS/ED/PSP/asis-2019-colombia.pdf>

3. Wang H, Paulson KR, Pease SA, Watson S, Comfort H, Zheng P, et al. Estimating excess mortality due to the COVID-19 pandemic: a systematic analysis of COVID-19-related mortality, 2020-21. *Lancet*. 2022; 399(10334): 1513-36. Doi: 10.1016/S0140-6736(21)02796-3
4. Lim WS, van der Eerden MM, Laing R, Boersma WG, Karalus N, Town GI, et al. Defining community acquired pneumonia severity on presentation to hospital: an international derivation and validation study. *Thorax*. 2003;58(5):377-82. doi: 10.1136/thorax.58.5.377
5. Vincent JL, Moreno R, Takala J, Willatts S, De Mendonça A, Bruining H, et al. The SOFA (Sepsis-related Organ Failure Assessment) score to describe organ dysfunction/failure. On behalf of the Working Group on Sepsis-Related Problems of the European Society of Intensive Care Medicine. *Intensive Care Med*. 1996;22(7):707-10. doi: 10.1007/BF01709751.
6. Hincapié C, Ascuntar J, León A, Jaimes F. Community-acquired pneumonia: comparison of three mortality prediction scores in the emergency department. *Colomb Med (Cali)*. 2021; 52(4): e2044287. Doi: 10.25100/cm.v52i4.4287
7. Wipf JE, Lipsky BA, Hirschmann JV, Boyko EJ, Takasugi J, Peugeot RL, et al. Diagnosing pneumonia by physical examination: relevant or relic? *Arch Intern Med*. 1999;159(10):1082-7. doi: 10.1001/archinte.159.10.1082.
8. Metlay JP, Kapoor WN, Fine MJ. Does this patient have community-acquired pneumonia? Diagnosing pneumonia by history and physical examination. *JAMA*. 1997; 278(17): 1440-5.
9. Atamna A, Shiber S, Yassin M, Drescher MJ, Bishara J. The accuracy of a diagnosis of pneumonia in the emergency department. *Internat J Infect Dis*. 2019; 89: 62-5. Doi: 10.1016/j.ijid.2019.08.027
10. Ticona JH, Zaccone VM, McFarlane IM. Community-Acquired Pneumonia: A Focused Review. *Am J Med Case Rep*. 2021; 9(1): 45-52. doi: 10.12691/ajmcr-9-1-12.
11. Alba AC, Agoritsas T, Walsh M, Hanna S, Iorio A, Devereaux PJ, et al. Discrimination and calibration of clinical prediction models: users' guides to the medical literature. *JAMA*. 2017; 318(14): 1377-84. doi: 10.1001/jama.2017.12126.
12. Kent P, Cancelliere C, Boyle E, Cassidy JD, Kongsted A. A conceptual framework for prognostic research. *BMC Med Res Methodol*. 2020; 20(1): 172. doi: 10.1186/s12874-020-01050-7
13. Narváez PO, Gomez-Duque S, Alarcon JE, Ramirez-Valbuena PC, Serrano-Mayorga CC, Lozada-Arcinegas J, et al. Invasive pneumococcal disease burden in hospitalized adults in Bogota, Colombia. *BMC Infect Dis*. 2021; 21(1): 1059. doi: 10.1186/s12879-021-06769-2
14. Aujesky D, Auble TE, Yealy DM, Stone RA, Obrosky DS, Meehan TP, et al. Prospective comparison of three validated prediction rules for prognosis in community-acquired pneumonia. *Am J Med*. 2005; 118(4): 384-92. doi: 10.1016/j.amjmed.2005.01.006
15. Metlay JP, Waterer GW, Long AC, Anzueto A, Brozek J, Crothers K, et al. Diagnosis and treatment of adults with community-acquired pneumonia. an official clinical practice guideline of the American Thoracic Society and Infectious Diseases Society of America. *Am J Respir Crit Care Med*. 2019; 200(7): e45-e67. doi: 10.1164/rccm.201908-1581ST.
16. National Institute for Health and Care Excellence. Pneumonia (community-acquired): antimicrobial prescribing. NICE guideline 138. England: NICE; 2022.