



Adjunctive treatment with oral dexamethasone in non-ICU patients hospitalised with community-acquired pneumonia: a randomised clinical trial

SHAREABLE PDF

Esther Wittermans¹, Stefan M.T. Vestjens¹, Simone M.C. Spoorenberg¹, Willem L. Blok², Jan C. Grutters^{3,4}, Rob Janssen⁵, Ger T. Rijkers⁶, Frank W.J.M. Smeenk⁷, G. Paul Voorn⁸, Ewoudt M.W. van de Garde^{9,10}, Willem Jan W. Bos^{1,11} and the Santeon-CAP Study Group¹²

¹Dept of Internal Medicine, St Antonius Hospital, Nieuwegein, The Netherlands. ²Dept of Internal Medicine, OLVG, Amsterdam, The Netherlands. ³Dept of Pulmonology, St Antonius Hospital, Nieuwegein, The Netherlands. ⁴Division of Heart and Lungs, University Medical Center Utrecht, Utrecht, The Netherlands. ⁵Dept of Pulmonology, Canisius Wilhelmina Hospital, Nijmegen, The Netherlands. ⁶Dept of Science, University College Roosevelt, Middelburg, The Netherlands. ⁷Dept of Pulmonology, Catharina Hospital, Eindhoven, The Netherlands. ⁸Dept of Medical Microbiology and Immunology, St Antonius Hospital, Nieuwegein, The Netherlands. ⁹Dept of Clinical Pharmacology, St Antonius Hospital, Nieuwegein, The Netherlands. ¹⁰Division of Pharmacoepidemiology and Clinical Pharmacology, Faculty of Science, Utrecht University, Utrecht, The Netherlands. ¹¹Division of Nephrology, Dept of Internal Medicine, Leiden University Medical Center, Leiden, The Netherlands. ¹²The complete membership of the Santeon-CAP Study Group is provided in the Acknowledgements.

Corresponding author: Esther Wittermans (e.wittermans@antoniusziekenhuis.nl)



Shareable abstract (@ERSpublications)

Adjunctive treatment with oral dexamethasone in adults hospitalised with community-acquired pneumonia (CAP) reduced length of stay and ICU admission rate. However, it remains unclear for which CAP subgroup the risk-benefit ratio is optimal. <https://bit.ly/35tXfPX>

Cite this article as: Wittermans E, Vestjens SMT, Spoorenberg SMC, *et al.* Adjunctive treatment with oral dexamethasone in non-ICU patients hospitalised with community-acquired pneumonia: a randomised clinical trial. *Eur Respir J* 2021; 58: 2002535 [DOI: 10.1183/13993003.02535-2020].

This single-page version can be shared freely online.

Copyright ©The authors 2021. For reproduction rights and permissions contact permissions@ersnet.org

This article has supplementary material available from erj.ersjournals.com

Received: 3 July 2020
Accepted: 27 Dec 2020

Abstract

Background Adjunctive intravenous corticosteroid treatment has been shown to reduce length of stay (LOS) in adults hospitalised with community-acquired pneumonia (CAP). We aimed to assess the effect of oral dexamethasone on LOS and whether this effect is disease severity dependent.

Methods In this multicentre, stratified randomised, double-blind, placebo-controlled trial, immunocompetent adults with CAP were randomly assigned (1:1 ratio) to receive oral dexamethasone (6 mg once daily) or placebo for 4 days in four teaching hospitals in the Netherlands. Randomisation (blocks of four) was stratified by CAP severity (pneumonia severity index class I–III and IV–V). The primary outcome was LOS.

Results Between December 2012 and November 2018, 401 patients were randomised to receive dexamethasone (n=203) or placebo (n=198). Median LOS was shorter in the dexamethasone group (4.5 days, 95% CI 4.0–5.0 days) than in the placebo group (5.0 days, 95% CI 4.6–5.4 days; p=0.033). Within both CAP severity subgroups, differences in LOS between treatment groups were not statistically significant. The secondary ICU admission rate was lower in the dexamethasone arm (5 (3%) *versus* 14 (7%); p=0.030); 30-day mortality did not differ between groups. In the dexamethasone group the rate of hospital readmission tended to be higher (20 (10%) *versus* 9 (5%); p=0.051) and hyperglycaemia (14 (7%) *versus* 1 (1%); p=0.001) was more prevalent.

Conclusion Oral dexamethasone reduced LOS and ICU admission rate in adults hospitalised with CAP. It remains unclear for which patients the risk-benefit ratio is optimal.