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- 3 National Institute for Health and Care Excellence. Venous thromboembolic diseases: diagnosis, management and thrombophilia testing. London: National Institute for Health and Care Excellence (UK), 2020.
- 4 Konstantinides SV, Meyer G, Becattini C, et al. 2019 ESC Guidelines for the diagnosis and management of acute pulmonary embolism developed in collaboration with the European Respiratory Society (ERS). Eur Heart J 2020; 41: 543-603.
- 5 Roncon L, Zuin M, Zonzin P. Age-adjusted D-dimer cut-off levels to rule out venous thromboembolism in COVID-19 patients. Thromb Res 2020; 190: 102.

## **Authors' reply**

We appreciate the issues related to normal age-related D-dimer concentrations raised by Marco Zuin and colleagues. The ACTION study<sup>1</sup> is a randomised trial designed pragmatically to answer the question of whether patients hospitalised with COVID-19 and elevated D-dimer concentration, without another indication for anticoaqulation, should routinely receive therapeutic anticoagulation with the factor Xa inhibitor, rivaroxaban. We found that rivaroxaban 20 mg once daily for 30 days had no benefit and significantly increased bleeding compared with inhospital prophylactic heparin.

Early in the COVID-19 pandemic, observational data suggested that thrombotic events were high in patients with COVID-19 and even higher among patients with an elevated D-dimer concentration.2 In light of the available information at the time, coupled with the fact that the D-dimer concentration of many patients with COVID-19 was measured when they were hospitalised, we included an elevated D-dimer concentration, defined as above the assay (not age-adjusted) upper limit of normal in each site, as an inclusion criterion for the ACTION trial. D-dimer concentrations are not routinely measured to guide therapeutic anticoagulation decision making in patients without COVID-19,3 and the purpose of enrolling patients with an elevated D-dimer concentration in ACTION was to increase the trial population's risk of thrombotic

events and not to establish D-dimer concentrations as a diagnostic tool to guide therapeutic anticoagulation in patients hospitalised with COVID-19.

A prespecified subgroup analysis of ACTION showed that the main results were consistent, irrespective of D-dimer concentrations.1 Similar results have been shown in other randomised trials investigating anticoagulation in patients with COVID-19, in which D-dimer concentrations at presentation (elevated vs normal vs not collected) did not influence the main results.<sup>4,5</sup> Furthermore, when we used ageadjusted D-dimer upper limits of normal, as proposed by Zuin and colleagues, we found that most (>90%) patients over the age of 50 years still had an elevated ageadjusted D-dimer concentration at study entry. Not surprisingly, when we excluded the fewer than 10% of patients who did not have an elevated age-adjusted D-dimer concentration, our main results remained consistent with those in the overall population (win ratio 0.87 [95% CI 0.59-1.26]). Therefore, the results from the ACTION trial are relevant, robust, and provide highquality evidence to avoid the routine use of therapeutic rivaroxaban—in the absence of another evidence-based indication for oral anticoagulation—in patients hospitalised with COVID-19, irrespective of D-dimer concentration.

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- 1 Lopes RD, de Barros e Silva PGM, Furtado RHM, et al. Therapeutic versus prophylactic anticoagulation for patients admitted to hospital with COVID-19 and elevated D-dimer concentration (ACTION): an open-label, multicentre, randomised, controlled trial. Lancet 2021: 397: 2253-63.
- 2 Tang N, Li D, Wang X, Sun Z. Abnormal coagulation parameters are associated with poor prognosis in patients with novel coronavirus pneumonia. J Thromb Haemost 2020: 18: 844–47.
- Ortel TL, Neumann I, Ageno W, et al. American Society of Hematology 2020 guidelines for management of venous thromboembolism: treatment of deep vein thrombosis and pulmonary embolism. *Blood Adv* 2020; 4: 4693-738.
- 4 The REMAP-CAP, ACTIV-4a, and ATTACC Investigators. Therapeutic anticoagulation with heparin in critically ill patients with COVID-19. N Engl J Med 2021; published online Aug 4. https://doi.org/10.1056/NEJMoa2103417.
- The ATTACC, ACTIV-4a, and REMAP-CAP Investigators. Therapeutic anticoagulation with heparin in noncritically ill patients with COVID-19. N Engl J Med 2021; published online Aug 4. https://doi.org/10.1056/ NEIMoa2105911.

## **Department of Error**

Shikino K, Sato R, Hanazawa N, Ikusaka M.
Chronic clicking tinnitus due to palatal tremor:
essential or secondary? Lancet 2021; 397: e16—
In this Clinical Picture, Manato Yasuda has
been added as an author. In the second
sentence of the fifth paragraph, treatment
dose has been corrected to 3 mg clonazepam
orally per day. These corrections have been
made to the online version as of Oct 7, 2021.