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* You should state whether an appropriate sample size was computed when the study was being designed
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* If no explicit power analysis was used, you should describe how you decided what sample (replicate) size (number) to use

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* Report exact p-values wherever possible alongside the summary statistics and 95% confidence intervals. These should be reported for all key questions and not only when the p-value is less than 0.05.

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* Indicate how samples were allocated into experimental groups (in the case of clinical studies, please specify allocation to treatment method); if randomization was used, please also state if restricted randomization was applied
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* Where provided, these should be in the most useful format, and they can be uploaded as “Source data” files linked to a main figure or table
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All data required to reproduce the reported analyses appears in the article text, tables, and figures. Data of this submission are anticipated to be part of a submission package to the European Medicines Agency for the request of approval of anakinra for the management of COVID-19 guided by the biomarker suPAR. Once this is finalized, the data may become available to your readers. This can be the case after signing contract with the sponsor of the study which is the Hellenic Institute for the Study of Sepsis. The responsible official DPO responsible for GDPR (headed@sepsis.gr).