

عنوان مقاله:

In silico analysis of the substitution mutations and evolutionary trends of the SARS-CoV-2 structural proteins in Asia

محل انتشار:

مجله علوم پایه پزشکی ایران، دوره 25، شماره 11 (سال: 1401)

تعداد صفحات اصل مقاله: 9

نویسندگان:

Mohammad Abavisani - *Student Research Committee, Mashhad University of Medical Sciences, Mashhad, Iran*

Karim Rahimian - *Bioinformatics and Computational Omics Lab (BioCOOL), Department of Biophysics, Faculty of Biological Sciences, Tarbiat Modares University, Tehran, Iran*

Mansoor Kodori - *Non communicable Diseases Research Center, Bam University of Medical sciences, Bam, Iran*

Reza Khayami - *Department of Medical Genetics and Molecular Medicine, Faculty of Medicine, Mashhad University of Medical Sciences, Mashhad, Iran*

Mahsa Mollapour Sisakht - *Department of Biochemistry, Erasmus University Medical Center, P.O. Box ۲۰۴۰, ۳۰۰۰ CA Rotterdam, The Netherlands*

Mohammadamin Mahmanzar - *Department of Bioinformatics, Kish International Campus University of Tehran, Kish, Iran*

Zahra Meshkat - *Department of Microbiology and Virology, Faculty of Medicine, Mashhad University of Medical Sciences, Mashhad, Iran*

خلاصه مقاله:

Objective(s): To address a highly mutable pathogen, mutations must be evaluated. SARS-CoV-2 involves changing infectivity, mortality, and treatment and vaccination susceptibility resulting from mutations. **Materials and Methods:** We investigated the Asian and worldwide samples of amino-acid sequences (AASs) for envelope (E), membrane (M), nucleocapsid (N), and spike (S) proteins from the announcement of the new coronavirus ۲۰۱۹ (COVID-۱۹) up to January ۲۰۲۲. Sequence alignment to the Wuhan-۲۰۱۹ virus permits tracking mutations in Asian and global samples. Furthermore, we explored the evolutionary tendencies of structural protein mutations and compared the results between Asia and the globe. **Results:** The mutation analyses indicated that ۵.۸۱%, ۷۰.۶۳%, ۲۶.۵۹%, and ۳.۳۶% of Asian S, E, M, and N samples did not display any mutation. Additionally, the most relative mutations among the S, E, M, and N AASs occurred in the regions of ۵۰۸ to ۶۳۵ AA, ۷ to ۱۴ AA, ۶۶ to ۸۸ AA, and ۱۶۴ to ۲۰۵ AA in both Asian and total samples. D۶۱۴G, T۹۱, I۸۲T, and R۲۰۳M were inferred as the most frequent mutations in S, E, M, and N AASs. Timeline research showed that substitution mutation in the location of ۶۱۴ among Asian and total S AASs was detected from January ۲۰۲۰. **Conclusion:** N protein was the most non-conserved protein, and the most prevalent mutations in S, E, M, and N AASs were D۶۱۴G, T۹۱, I۸۲T, and R۲۰۳M. Screening structural protein mutations is a robust approach for developing drugs, vaccines, and more specific diagnostic tools.

لینک ثابت مقاله در پایگاه سیویلیکا:

<https://civilica.com/doc/1540935>

