

Novel Corona Virus

Puca Edmond*

Department of Endocrinology, University Hospital Center of Mother Teresa, Tirana, Albania

Short Communication

Received: 01/07/2021

Accepted: 15/07/2021

Published: 22/07/2021

***For correspondence:**

Puca Edmond, Department of
Agricultural Engineering, RIMT
University, Mandi Gobindgarh,
Punjab, India

E-mail: pucaedmon@gmail.com

Keywords: SARS-CoV-2;
Nucleocapsids; Polyprotein

ABSTRACT

Corona virus have an unsegmented, single-stranded, positive-sense RNA genome of around 30 kb, encased by a 5'-cap and 3'-poly(A) tail. The genome of SARS-CoV-2 is 29,891 bp long, with a G+C substance of 38%. These infections are surrounded with an envelope containing viral nucleocapsid. The nucleocapsids in CoVs are orchestrated in helical balance, which mirrors an atypical quality in certain sense RNA infections. The electron micrographs of SARS-CoV-2 uncovered a wandering round layout with some level of pleomorphism, virion breadths fluctuating from 60 to 140 nm, and unmistakable spikes of 9 to 12 nm, giving the infection the presence of a sun oriented crown. The CoV genome is orchestrated directly as 5'-pioneer UTR-replicase-auxiliary qualities (S-E-M-N)- 3' UTR-poly(A). Frill qualities, for example, 3a/b, 4a/b, and the hemagglutinin-esterase quality (HE), are likewise observed blended with the basic qualities. SARS-CoV-2 has additionally been discovered to be masterminded comparably and encodes a few extra proteins, despite the fact that it comes up short on the HE, which is normal for some betacoronaviruses. The positive-sense genome of CoVs fills in as the mRNA and is meant polyprotein 1a/1ab (pp1a/1ab). A replication-record complex (RTC) is shaped in twofold film vesicles (DMVs) by nonstructural proteins (nsps), encoded by the polyprotein gene. Accordingly, the RTC incorporates a settled arrangement of subgenomic RNAs (sgRNAs) by means of discontinuous transcription.

INTRODUCTION

SARS-CoV-2 is viewed as another Betacoronavirus having a place with the subgenus Sarbecovirus. A couple of other basic zoonotic infections (MERS-related CoV and SARS-related CoV) have a place with similar class. In any case, SARS-CoV-2 was distinguished as a particular infection dependent on the percent personality with different Betacoronavirus; preserved open perusing outline 1a/b (ORF1a/b) is beneath 90% identity. A general 80% nucleotide identity was seen between SARS-CoV-2 and the first SARS-CoV, alongside 89% identity with ZC45 and ZXC21 SARS-related CoVs of bats [1]. What's more, 82% identity has been seen between SARS-CoV-2 and human SARS-CoV Tor2 and human SARS-CoV. A succession personality of just 51.8% was seen between MERS-related CoV

and the as of late arose SARS-CoV-2. Phylogenetic investigation of the auxiliary qualities additionally uncovered that SARS-CoV-2 is nearer to bat SARS-related CoV. In this way, SARS-CoV-2 may have started from bats, while other enhancer hosts may have assumed a part in infection transmission to people. Of note, the other two zoonotic CoVs (MERS-related CoV and SARS-related CoV) likewise began from bats [2].

DESCRIPTION

Coronavirus disease (COVID-19) is an infectious disease caused by a newly discovered coronavirus. Corona virus have an unsegmented, single-stranded, positive-sense RNA genome of around 30 kb, encased by a 5'-cap and 3'-poly(A) tail. The genome of SARS-CoV-2 is 29,891 bp long, with a G+C substance of 38%. These infections are surrounded with an envelope containing viral nucleocapsid. The nucleocapsids in CoVs are orchestrated in helical balance, which mirrors an atypical quality in certain sense RNA infections. The electron micrographs of SARS-CoV-2 uncovered a wandering round layout with some level of pleomorphism, virion breadths fluctuating from 60 to 140 nm, and unmistakable spikes of 9 to 12 nm, giving the infection the presence of a sun oriented crown. The CoV genome is orchestrated directly as 5'-pioneer UTR-replicase-auxiliary qualities (S-E-M-N)- 3' UTR-poly(A) [3]. Frill qualities, for example, 3a/b, 4a/b, and the hemagglutinin-esterase quality (HE), are likewise observed blended with the basic qualities. SARS-CoV-2 has additionally been discovered to be masterminded comparably and encodes a few extra proteins, despite the fact that it comes up short on the HE, which is normal for some betacoronaviruses. The positive-sense genome of CoVs fills in as the mRNA and is meant polyprotein 1a/1ab (pp1a/1ab). A replication-record complex (RTC) is shaped in twofold film vesicles (DMVs) by nonstructural proteins (nsps), encoded by the polyprotein gene. Accordingly, the RTC incorporates a settled arrangement of subgenomic RNAs (sgRNAs) by means of discontinuous transcription [4].

SARS-CoV-2 is viewed as another Betacoronavirus having a place with the subgenus Sarbecovirus. A couple of other basic zoonotic infections (MERS-related CoV and SARS-related CoV) have a place with similar class. In any case, SARS-CoV-2 was distinguished as a particular infection dependent on the percent personality with different Betacoronavirus; preserved open perusing outline 1a/b (ORF1a/b) is beneath 90% identity. A general 80% nucleotide identity was seen between SARS-CoV-2 and the first SARS-CoV, alongside 89% identity with ZC45 and ZXC21 SARS-related CoVs of bats. What's more, 82% identity has been seen between SARS-CoV-2 and human SARS-CoV Tor2 and human SARS-CoV. A succession personality of just 51.8% was seen between MERS-related CoV and the as of late arose SARS-CoV-2. Phylogenetic investigation of the auxiliary qualities additionally uncovered that SARS-CoV-2 is nearer to bat SARS-related CoV. In this way, SARS-CoV-2 may have started from bats, while other enhancer hosts may have assumed a part in infection transmission to people. Of note, the other two zoonotic CoVs (MERS-related CoV and SARS-related CoV) likewise began from bats [5].

CONCLUSION

Advances in novel microbiological techniques have to be implemented to combat the treatment procedures in SARS-CoV-2. Research and reviews in microbiology and biotechnology journal encourages novel research towards a new microbial world.

Most people infected with the COVID-19 virus will experience mild to moderate respiratory illness and recover without requiring special treatment. Older people, and those with underlying medical problems like cardiovascular disease, diabetes, chronic respiratory disease, and cancer are more likely to develop serious illness.

The best way to prevent and slow down transmission is to be well informed about the COVID-19 virus, the disease it causes and how it spreads. Protect yourself and others from infection by washing your hands or using an alcohol based rub frequently and not touching your face.

REFERNCES

1. Liang H, et al. Novel corona virus disease (COVID-19) in pregnancy: What clinical recommendations to follow? AOGS. 2020;123:2-4.
2. Kumar D, et al. Corona virus: a review of COVID-19. EJMO. 2020;4:8-25.
3. Peng QY, et al. Findings of lung ultrasonography of novel corona virus pneumonia during the 2019–2020 epidemic. Intensive Care Med. 2020;46:845-850.
4. Ikhlaq A, et al. Awareness and attitude of undergraduate medical students towards 2019-novel corona virus. Pak J Med Sci. 2020;36:32-36.
5. Zheng F, et al. Clinical characteristics of 161 cases of corona virus disease 2019 (COVID-19) in Changsha. Eur Rev Med Pharmacol Sci. 2020;24:3404-3410.