

# Identification of Natural Chemical Inhibitors against SARS-Associated Coronavirus

Chelladurai G

Department of Zoology, G.Venkataswamy Naidu College,  
Kovilpatti, Tamil Nadu, India

Corresponding author: marinesafety20@gmail.com

Received on: 03/05/2023

Accepted on: 15/09/2023

Published on: 27/09/2023

## ABSTRACT

**Aim:** This study was aimed to investigate the available natural drugs show in vitro anti SARS- CoV activity.

**Method and Materials:** Properties of some natural drugs like saikosaponins, amentoflavone and L-Isoleucine –Essential amino acid were used for the investigations.

**Results:** Results of drugs may act preventing viral replication by inhibiting viral DNA polymerase and capacity of preventing the invasion of pathogens *via* the increase of immunity

**Conclusion:** It was concluded that some natural drugs from plant extract and the compound of L- Isoleucine may be considered for further researches of anti-SARS-CoV drugs.

**Keywords:** SARAS-COV, natural drugs, amino acids and antiviral mask.

**How to cite this article:** Chelladurai G (2023). Identification of Natural Chemical Inhibitors against SARS-Associated Coronavirus. J. Chem. Res. Adv., 04(02): 26-28.

## Introduction

The Severe acute respiratory syndrome (SARS) is an infectious disease caused by a newly identified human coronavirus (SARS-CoV). The disease can produce severe pneumonia with a reported fatal outcome of 15% to 20%. Currently, no effective drug exists to treat SARS-CoV infection. The urgency of the outbreak has led to the empiric use of broad- spectrum antibiotics and antiviral agents in affected patients in several countries. Intensive efforts are under way to gain more insight into the mechanisms of viral replication, in order to develop targeted antiviral therapies and vaccines (Emily et al., 2004). The some natural drugs can act as antiviral strategies are generally found direct antiviral effects, inhibition of viral entry and replication at the cellular level by targeting virus-related processes, and enhancement of host immune response (Driscoll, 2002).

This natural molecules extracted from living organism can play an important role in the development of new drugs as they may have advantages over conformist chemical compound-based medications, such as fewer side effects, less long-term toxicity, variable bioavailability, and unidentified chemical structures and biological activities. Hence the present study to identification of some natural chemical inhibitors against coronavirus.

### Natural drugs

#### 1. Saikosaponins (Isolation from Plant Source)

It is more effective against COVID- 19 and used in the treatment of Virus diseases and it may act include preventing viral replication by inhibiting viral DNA polymerase; binding to specific cell-surface receptors and inhibiting viral penetration or un-coating; inhibiting viral protein synthesis; or blocking late stages of virus assembly and also suppress immune function by one of several mechanisms of action. The others may act through activation of T-Cells (Al-Jabri et al., 1996). This compound might enter in to the COVID-19 virus and will inhibit the viral replication and may prevent the further development. So far, this drug is more useful and active to development of viral infected peoples (Hao et al., 2012). It is purely natural drug and obtained from only plant

**Copyright:** Chelladurai. Open Access. This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (<http://creativecommons.org/licenses/by/4.0/>), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

materials. There is no any side effected when some of chemical based drugs.

### 2. Amentoflavone (Isolation from Plant Source)

It is more effective against COVID- 19 and Amentoflavone or Biflavonoid with anti-inflammatory, anti-viral and cancer chemopreventive activity. It inhibits vascularization of tumors by blocking the activity of angiogenic VEGFs. Blocks the induction of COX-2 and up-regulates PPAR- $\gamma$ . It is a negative modulator of the GABAA receptor at the benzodiazepine binding site (Chang et al., 2007). These compounds have inhibited the SARS protease inhibitor, which is identified and inhibit the peptide DNA replication of this virus.

#### New Developing Product against COVID-19

This kind of mask will be prepared with natural drugs like Saikosaponins or Amentoflavone. It is a SARS-CoV Protease inhibitor. It more resistance to microbes and also SARS Virus and eco-friendly to human body. The following pharmacological activity has been reported of this compound.

- Antimicrobial properties
- It can protect against neuro-inflammation
- Anti-Oxidant and anti-Inflammatory Properties
- May prevent cancer formation
- Helps With Depression and Anxiety
- Protects against skin Aging, Inflammation, vascular system
- Protects the liver and metabolism, Promotes bone growth and prevent the hair loss

#### Properties of Natural drugs

- It occurs in the divided solid state
- Melting range of these compounds is more than 400°F
- It is partially miscible in water. Its solubility is measured by gram per liter
- It is pale yellow powder, does not mix well with water

**Note:** The above mentioned information is more useful to preparation of mask against COVID-19. It is not soluble easily in water and melting point is also high, so we can prepare the mask and use it for long time.

### 3. L-Isoleucine -Essential amino acid (Vaccine for COVID-19)

Isoleucine is a class of branched chain amino acid, which is essential for some physiological functions of humans and other vertebrates. The

L- isoleucine will specifically be built into proteins of immune cells like lymphocytes [Zakaryan et al., 2017, Chuang et al., 1990]. They plays a critical role for immune functions, including maintaining the development of immune organs include cells and stimulating the secretion of immune molecules substances in human and other vertebrates. In recent clinical study, dietary l-isoleucine supplementation can relieve the acute diarrhoea and cardiovascular diseases and other malfunction organs. Which is related to the production of host defense peptides induced by isoleucine [Hale et al., 2004, Rivas-Santiago et al., 2001]. Thus, isoleucine has the capacity of preventing the invasion of pathogens *via* the increase of immunity. The L isoleucine is a essential peptide to break the surface protein of COVID -19 virus and also inhibit the DNA replication hence it may act as vaccine against CoV -virus.

### Conclusion

It was concluded that some natural drugs from plant extract and the compound of L- Isoleucine may be considered for further researches of anti-SARS-CoV drugs. Moreover, further research is needed on these molecules to exhibit the antiviral drug against COVID -19.

### Reference

- Al-Jabri AA, Wigg MD and Oxford JS (1996). Initial in vitro screening of drug candidates for their potential antiviral activities. In: Mahy, B.W.J., Kangro, H.O., Editors. Virology methods manual. London: Academic Press Ltd. p. 293-356.
- Chang JS, Wang KC, Liu HW, Chen MC, Chiang LC and Lin CC (2007). Sho-saiko-to (Xiao-Chai-Hu-Tang) and crude saikosaponins inhibit hepatitis B virus in a stable HBV-producing cell line. *Am. J. Chin. Med.*, 35: 341-51.
- Chuang JC, Yu CL and Wang SR (1990). Modulation of human lymphocyte proliferation by amino acids. *Clin. Exp. Immunol.*, 81(1): 173-6.  
10.1111/j.1365-2249.1990.tb05310.x.
- Driscoll JS (2002). Antiviral drugs. Aldershot, UK: Ashgate Publishing Ltd.
- Emily LC, Chin-Yo Lin, Hwee Cheng Tan, Ai Ee Ling and Bing Lim (2004). Inhibition of SARS Coronavirus Infection in Vitro with Clinically Approved Antiviral Drugs, *Emerg Infect Dis.*, 10(4): 581-586.

- Hale LL, Pharr GT, Burgess SC, Corzo A and Kidd MT (2004). Isoleucine needs of thirty- to forty-day-old female chickens: immunity. *Poult. Sci.*, 83(12): 1979-85. 10.1093/ps/83.12.1979
- Hao BJ, Wu YH, Wang JG, Hu SQ, Keil DJ and Hu HJ (2012). Hepatoprotective and antiviral properties of isochlorogenic acid A from *Laggeraalata* against hepatitis B virus infection. *J Ethnopharmacol*, 144: 190-4.
- Rivas-Santiago CE, Rivas-Santiago B, León DA, Castañeda-Delgado J and Hernández PR (2011). Induction of  $\beta$ -defensins by l-isoleucine as novel immunotherapy in experimental murine tuberculosis. *Clin Exp Immunol*, 164(1): 80-9.
- Zakaryan E, Arabyan A and Zandi (2017). "Flavonoids: promising natural compounds against viral infections," *Arch. Virol.*, 162(9): 2539-2551.

\*\*\*\*\*