

# Beware the Snake Oil Salesmen

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*"The snake venom theory by Dr. Bryan Ardis is built upon the interpretation of the unpurified fraudulent "SARS-COV-2" genome which is itself built upon references to other fraudulent genomes of human and animal "coronaviruses" created in the very same way. Attempting to claim any connections between the random A,C,T,G's in a computer database is a useless and pointless exercise as the RNA that was fabricated into the genome of a "virus" was never purified, isolated, and proven to physically exist in the first place. Thus any connections between the protein codes said to belong to a "virus" which are then said to be closely related to supposed snake "coronaviruses" is immediately invalid.*

*Using this invalid premise to then claim that people have been poisoned by snake venom in the vaccines, the drugs, and the water supply is nothing but unsubstantiated science fiction that seems designed to have a few purposes:*

- 1. To keep people engaged in the lie that a new disease known as "Covid-19" exists and that there is a singular cause.*
- 2. To restore faith in monoclonal antibodies and other toxic alternative treatments.*

3. *To use the theory to promote and sell anti-venom supplements.*
4. *To divide and distract those questioning the official narrative.*
5. *To make the “Truther” community look foolish by falling for loosely tied-together circumstantial evidence that is easily debunked.”*

~ *Mike Stone*

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## Beware the Snake Oil Salesmen

by [Mike Stone](#), [Viroliegy](#)

April 18, 2022

**“My story has never been to create fear, panic, and anxiety about water.”** He said he told Peters that he believes **“there’s actually a snake venom connection to all of COVID-19, and I think that’s the weapon.”** – Dr. Bryan Ardis

<https://www.thedailybeast.com>

Summarizing his theory, Dr. Ardis said, **“They are using Krait venom and Cobra venom, calling it Covid-19, you’re drinking it, it’s getting into your brainstem and it’s paralyzing your diaphragm’s ability to breathe.”**

<https://www.google.com/amp/s/miamistandard.news/>

I really didn’t want to write this article. I was hopeful that people would easily see right through the unsubstantiated claims of Dr. Bryan Ardis that snake venom is the cause of

“Covid.” I was hopeful that people would take the time to research the information presented in support of the snake venom theory to see if it held any merit at all. I thought his whirlwind alternative media tour on the who’s who of questionable sources (including the likes of Stew Peters, Mike Adams, and Infowars) would have people questioning why this theory was allowed to be so heavily promoted so quickly. I thought that the fact that the man who created the “Covid” snake venom theory was actually selling his own anti-venom line of supplements would be enough grounds to be skeptical of his motive and his claims.

☆ thedrardisshow.com ↻

Dr. Ardis

The Dr Ardis Show is Hosted By Dr Bryan Ardis, D.C.

## The Anti-V Collection - Quercetin, Vitamin C, N-acetyl L-cysteine (NAC), Zinc

ArdisLabs

★★★★★ (No reviews yet)

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**MSRP: \$122.96**

**\$109.00**

**(You save \$13.96)**



The Anti-V Collection is a special formulation created by Dr. Bryan Ardis, D.C.

The Anti-V Collection is a four bottle supplement kit that includes 4 separate bottles of Quercetin, Immune-C, N-acetyl L-cysteine (NAC), and Zinc. Each kit is a 30 day supply of premium ingredients that are made in our facility in Plano, TX.

Dr. Ardis, D.C. recommends this supplement collection because:

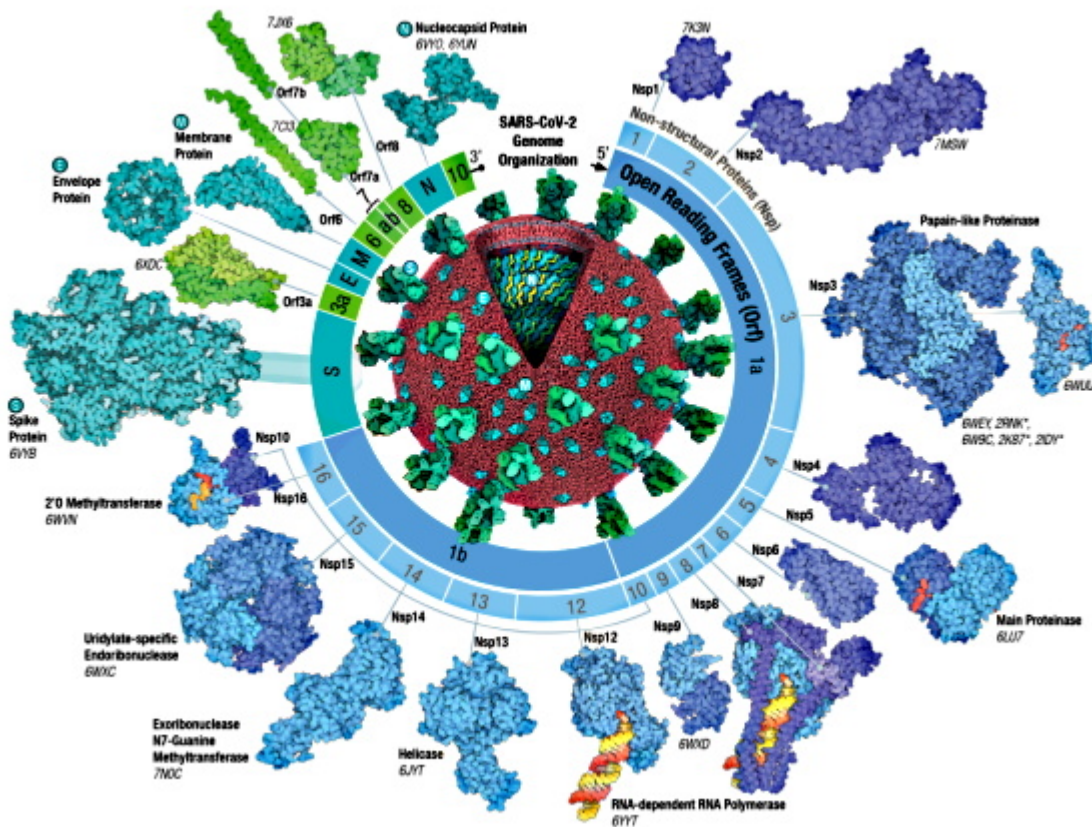
1. Vitamin C is found in numerous studies to inhibit destructive and deadly impact of V.
2. NAC (N-acetyl L-cysteine) helps the liver make Glutathione, is proven to be completely inhibitory to things such as S. V. Phosphodiesterase.
3. Zinc ion receptors are the second most targeted receptors of all cells by V.
4. Quercetin which is a natural substance that behaves like a zinc ionophore and should complement the zinc receptor protections against V.

It seems I was wrong. Just like the baseless vaccine shedding and [gain of function/bioweapons](#) narratives, this new snake venom theory has sadly spread through the “Truther” community like wildfire, with many who rightfully challenge the existence of “viruses” clinging to the idea of a new invisible enemy to defeat. They believe that it must be a new toxin. It can’t possibly be the same factors we have seen each and every year leading to disease. This toxin must be hiding in the [vaccines](#), the drugs, and/or even the very water we drink. What these “Truthers” do not realize is that this very line of thinking gives credibility to the idea of a new disease which requires new treatments in order to combat it. This is exactly what the pharmaceutical companies want you to believe.

However, there is NO NEW DISEASE. There is no need for any new or even existing pharmaceutical interventions to treat the same symptoms of detoxification people go through each and every year. In fact, the current treatments can easily be shown to have led to numerous unnecessary deaths. There is no new threat known as “Covid-19” which is being caused by any one factor. The factors leading to the symptoms of disease people are experiencing are multi-causal as they are every year.

Now this is not to say that the vaccines, the drugs, or even the water supply are free of toxins. These are all sources of toxicity and should be investigated as to their composition and effects on our health. However, the theory that there is one factor in all of these sources, i.e. snake venom, and this one factor is leading to the symptoms of disease people are experiencing is, at present time, completely baseless. And it all begins at the very foundation of the fraudulent genome.

## **The Fraudulent Genome**



You take that snake or that serpent and **you figure out how to isolate genes from that serpent and get those genes of that serpent to insert itself into your God-given created DNA.** I think this is the plan all along, was to get the serpents', the evil one's DNA, into your God-created DNA."

<https://www.thedailybeast.com/covid-conspiracy-theorists-are-at-war-over-snake-venom>

He also said **genetic sequence** testing done on sick patients in Wuhan **found their genetic sequence matched two snakes,** the Chinese Krait and King Cobra, not bats."

<https://www.google.com/amp/s/miamistandard.news/>

From Dr. Ardis' interview with Mike Adams, he supplied the article "**Snakes could be the source of the Wuhan coronavirus outbreak**" from CNN as his starting point for the "Covid"/snake connection. Within the article, you can see that this claim originates from the fraudulent genomes:

**“The researchers used an analysis of the protein codes favored by the new coronavirus and compared it to the protein codes from coronaviruses found in different animal hosts, like birds, snakes, marmots, hedgehogs, manis, bats and humans. Surprisingly, they found that the protein codes in the 2019-nCoV are most similar to those used in snakes.”**  
<https://www.google.com/amp/s/amp.cnn.com/>

To anyone who actually researched the [creation of the original “SARS-COV-2” genome](#), it is readily apparent that it is a fraudulent computer-generated creation stemming from the unpurified lung fluid of a single patient. The sequenced material could have come from multiple sources, including host DNA/RNA, bacteria, and microbes/microorganisms. It could have even come from outside contamination. There is no way to tell what the origin of the RNA is or even if it was a single source as no particles assumed to be “SARS-COV-2” were ever properly [purified and isolated](#) directly from the fluids of the sick patient before being sequenced. Thus, any relation this fabricated sequence has to any other sequence is invalid as the source was never identified to exist as a physical entity to begin with. Considering that the [bat](#) and snake “coronavirus” sequences for which the “SARS-COV-2” sequence [was then compared to](#) also come from unpurified sources, it is easy to see that any claims as to the origins of the sequenced material is a horrible foundation to build upon for an origin theory of a nonexistent “virus” and/or disease.

Even if this snake-venom connection was valid, the enzyme phospholipase A2 group IIA or sPLA2-IIA, which Dr. Ardis bases much of his claims on, only has *similarities* to rattlesnake venom. These peptides are “almost identical” to the venoms of animals and yet they are regularly found in healthy humans and other mammals. From his own source:

## **Like Venom Coursing Through the Body: Researchers Identify Mechanism Driving COVID-19 Mortality**

“Researchers from the University of Arizona, in collaboration with Stony Brook University and Wake Forest School of Medicine, analyzed blood samples from two COVID-19 patient cohorts and **found that circulation of the enzyme – secreted phospholipase A2 group IIA, or sPLA2-IIA, –** may be the most important factor in predicting which patients with severe COVID-19 eventually succumb to the virus.

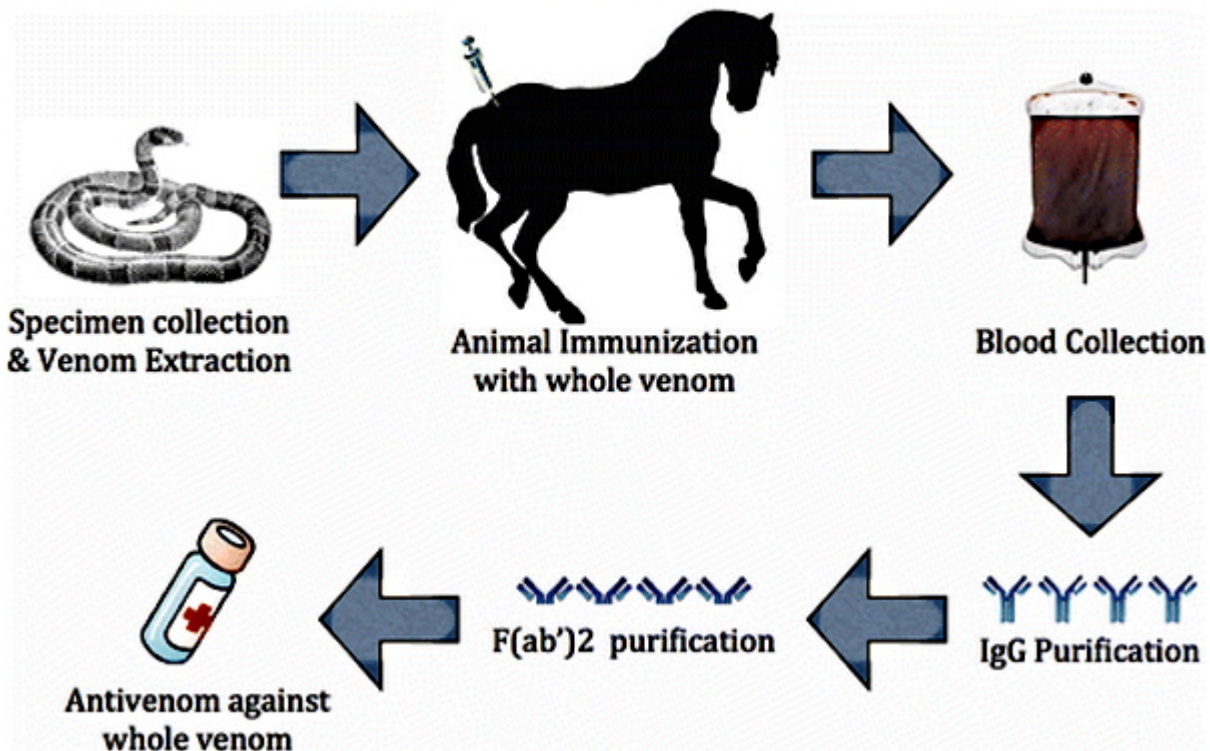
The sPLA2-IIA enzyme, which has **similarities** to an active enzyme in rattlesnake venom, **is found in low concentrations in healthy individuals** and has long been known to play a critical role in defense against bacterial infections, destroying microbial cell membranes.”

Thus, the snake enzymes are in fact normal human enzymes that are regularly found in healthy individuals. There is no mystery as to why these would be present in a sample. We should be able to put this “Covid” snake venom nonsense to bed right here. However, let’s press on a see what else we can uncover.

### **Antivenom = Monoclonal Antibodies**

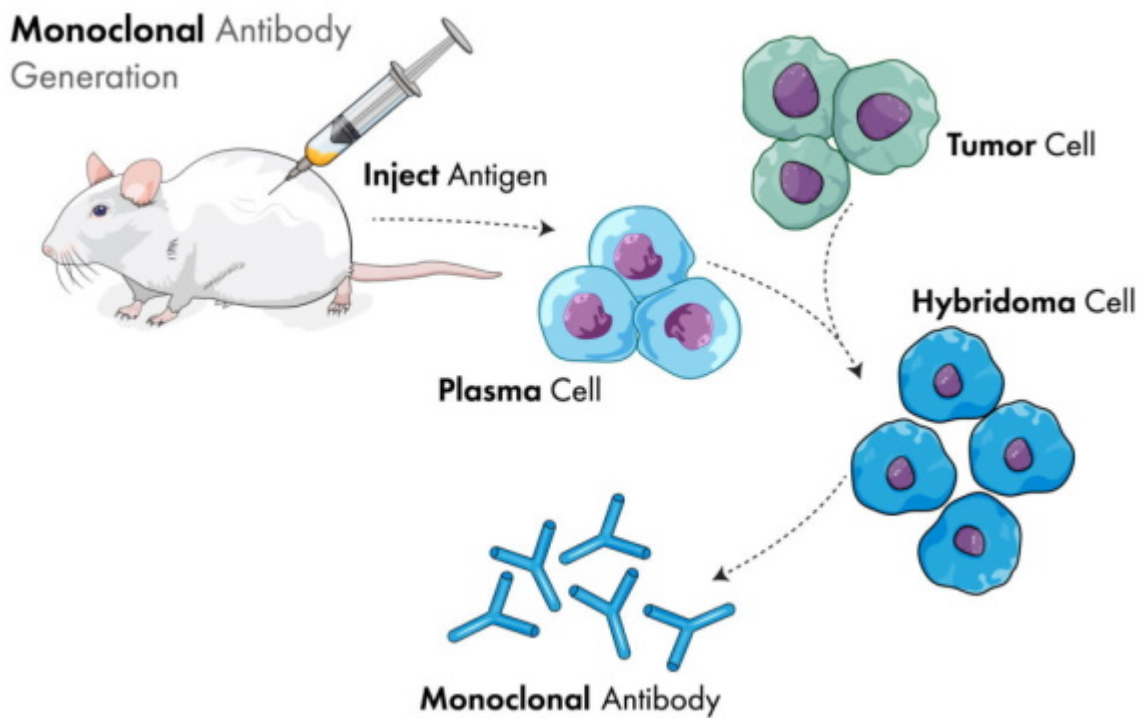
One thing I will give Dr. Ardis credit for is spotlighting the connection between the creation of antivenoms with the creation of monoclonal antibodies. The processes for both are very similar and the desired outcome is the exact same: the creation of [theoretical antibodies](#). In the case of snake antivenom, it is normally created by a series of injections of the venom of a snake into an animal and then collecting the blood after a period of time. This is usually done through horses but other animals can be used as the host as well. Thus, the antivenom used for a snakebite victim is typically an injection of horse blood.

## Traditional Antivenom Production



Monoclonal antibodies, on the other hand, are created by injecting mice with an antigen, extracting the resulting blood which contains the theoretical antibodies, and culturing it in myeloma (i.e. cancer) cells. For the creation of “SARS-COV-2” therapies, it is said that they are typically created either from the B cells of recovered “Covid” patients or by [immunizing mice genetically modified to have a humanized immune system and harvesting the “effective” antibodies from them.](#)





Both of these therapies have their basis in animal blood and the creation of the theoretical antibodies. Both are associated with toxic side effects. Sadly, while he was originally right about the fact that monoclonal antibodies are toxic and should not be used to treat the symptoms now collectively known as "Covid," Dr. Ardis changed his tune when another doctor texted him asking if he would use antivenom for a snake bite:

**"Last December, Dr Bryan Ardis received a text message from an Emergency Room physician friend of his that sent him down an unexpected and bizarre rabbit hole that may explain the adverse events from the vaccines that we've been reporting. The text read: "Hey Dr Ardis...If you got bit by a rattlesnake, would you go to a hospital and get anti-venom?"**

**"He says, "I realized, all of a sudden, monoclonal antibodies ARE anti-venom. The Federal Government doesn't want us using anti-venom. Why are they fighting anti-venom and why are we finding anti-venom works against COVID? Is**

it not a virus? Is it a venom? This is what I want to know: **Is COVID a venom and is this why they don't want you using monoclonal antibodies?"**

<https://www.google.com/amp/s/cairnsnews.org/>

Do you see the trick? They want you to equate monoclonal antibodies with antivenom. This is supposed to be an "aha" moment where you realize that there is no way that you would not inject antivenom (i.e. horse blood) into yourself if bitten by a snake. It's a no-brainer, right? We have all seen the movies where a person is bitten by a venomous snake and quickly dies if not given the antivenom.

If you are willing to accept the injection of horse blood into your body to survive a snake bite, why wouldn't you also inject the cancer-cell cultured blood of genetically altered mice in order to combat "Covid?"

As Dr. Ardis points out, monoclonal antibodies are essentially antivenom. However, he wrongly states that monoclonal antibodies are an effective therapy. According to a September 2021 Cochrane review of the available studies, they found insufficient evidence to claim that monoclonal antibodies are an effective treatment for "SARS-COV-2:"

### **Are laboratory-made, COVID-19-specific monoclonal antibodies an effective treatment for COVID-19?**

"The evidence for each comparison is based on single studies. None of these measured quality of life. Our **certainty in the evidence** for all non-hospitalised individuals **is low**, and for hospitalised individuals is **very low to moderate**. We consider the current evidence **insufficient to draw meaningful conclusions regarding treatment with SARS-CoV-2-neutralising mAbs.**"

<https://www.cochrane.org/CD013825/>

In other words, the evidence for the usefulness of monoclonal antibodies is non-existent. Unfortunately, the Cochrane Review failed to point out that there are various risks and adverse reactions associated with their use:

### **Do mAbs have risks?**

**“Therapeutic mAbs, typically administered by intravenous (IV) infusion, have been a valuable and generally safe treatment option for a variety of conditions for many years. However, they are also known to cause a range of side effects and reactions, which can be immediate or delayed. Serious adverse events associated with mAbs include infusion reactions, acute anaphylaxis, and serum sickness, as well as longer-term complications such as infections, cancer, autoimmune disease, and cardiotoxicity.”**

<https://www.ecri.org/components/>

In January 2022, the FDA restricted the use of some monoclonal therapies (Bamlanivimab and Etesevimab) that are authorized against “Covid-19” as they were shown to be ineffective:

### **Coronavirus (COVID-19) Update: FDA Limits Use of Certain Monoclonal Antibodies to Treat COVID-19 Due to the Omicron Variant**

**“In light of the most recent information and data available, today, the FDA revised the authorizations for two monoclonal antibody treatments – bamlanivimab and etesevimab (administered together) and REGEN-COV (casirivimab and imdevimab) – to limit their use to only when the patient is likely to have been infected with or exposed to a variant that is susceptible to these treatments.**

**Because data show these treatments are highly unlikely to be active against the omicron variant, which is circulating at a very high frequency throughout the United States,**

**these treatments are not authorized for use in any U.S. states, territories, and jurisdictions at this time.** In the future, if patients in certain geographic regions are likely to be infected or exposed to a variant that is susceptible to these treatments, then use of these treatments may be authorized in these regions.

**Monoclonal antibodies are laboratory-made proteins that mimic the immune system's ability to fight off harmful pathogens such as viruses, like SARS-CoV-2.** And like other infectious organisms, SARS-CoV-2 can mutate over time, resulting in certain treatments not working against certain variants such as omicron. This is the case with these two treatments for which we're making changes today."

<https://www.fda.gov/news-events/>

On April 16th, 2022, the FDA revoked the use of Bamlanivimab alone as it's benefits were shown not to outweigh its risks. Somehow despite this evidence, the FDA still allows for it to be used in combination with Etesevimab, even though they previously revoked their use together in January 2022:

### **Coronavirus (COVID-19) Update: FDA Revokes Emergency Use Authorization for Monoclonal Antibody Bamlanivimab**

"Today, the U.S. Food and Drug Administration **revoked the emergency use authorization (EUA) that allowed for the investigational monoclonal antibody therapy bamlanivimab,** when administered alone, to be used for the treatment of mild-to-moderate COVID-19 in adults and certain pediatric patients. Based on its ongoing analysis of emerging scientific data, specifically the sustained increase of SARS-CoV-2 viral variants that are resistant to bamlanivimab alone **resulting in the increased risk for treatment failure,** the FDA has determined that the known and potential benefits of bamlanivimab, when administered alone, no longer outweigh the known and potential risks for

**its authorized use.** Therefore, the agency determined that the criteria for issuance of an authorization are no longer met and has revoked the EUA.

On Nov. 9, 2020, based on the totality of scientific evidence available at the time, the FDA issued an EUA to Eli Lilly and Co. authorizing the emergency use of bamlanivimab alone for the treatment of mild to moderate COVID-19 in adults and pediatric patients (12 years of age and older weighing at least 40 kg) with positive results of direct SARS-CoV-2 viral testing, and who are at high risk for progressing to severe COVID-19 and/or hospitalization. **Importantly, although the FDA is now revoking this EUA, alternative monoclonal antibody therapies remain available under EUA, including REGEN-COV (casirivimab and imdevimab, administered together), and bamlanivimab and etesevimab, administered together, for the same uses as previously authorized for bamlanivimab alone.** The FDA believes that these alternative monoclonal antibody therapies remain appropriate to treat patients with COVID-19 when used in accordance with the authorized labeling based on information available at this time.”

<https://www.fda.gov/news-events/press-announcements/>

If the FDA's confusing revoking of the EUA's of these monoclonal antibodies has you concerned that you will not be able to use them against an imaginary "virus," don't worry. The FDA authorized the use of a new "Omicron-specific" monoclonal antibody called Bebtelovimab on February 11th, 2022. Granted, it still carries the same risks, adverse side effects, and uncertainty over clinical worsening listed for the previously ineffective antibody therapies. From the FDA fact sheet:

**Coronavirus (COVID-19) Update: FDA Authorizes New Monoclonal Antibody for Treatment of COVID-19 that**

## **Retains Activity Against Omicron Variant**

“Possible side effects of bebtelovimab include **itching, rash, infusion-related reactions, nausea and vomiting**. Serious and unexpected adverse events including **hypersensitivity, anaphylaxis and infusion-related reactions** have been observed with other SARS-CoV2 monoclonal antibodies and could occur with bebtelovimab. In addition, **clinical worsening following administration of other SARS-CoV-2 monoclonal antibody treatment** has been reported and therefore is possible with bebtelovimab. **It is not known if these events were related to SARS-CoV-2 monoclonal antibody use or were due to progression of COVID-19.**”

<https://www.fda.gov/news-events/press-announcements/>

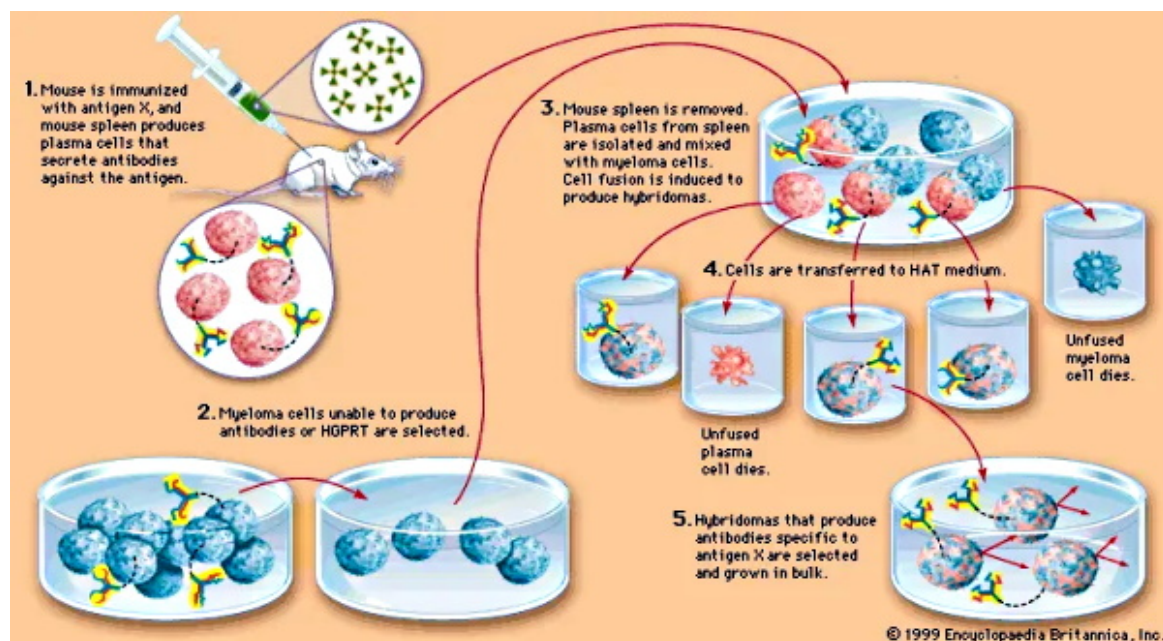
## **Coronavirus (COVID-19) Update: FDA Authorizes New Monoclonal Antibody for Treatment of COVID-19 that Retains Activity Against Omicron Variant**

- **Hypersensitivity Including Anaphylaxis and Infusion-Related Reactions:** Serious hypersensitivity reactions, **including anaphylaxis**, have been observed with administration of other SARS-CoV-2 monoclonal antibodies and could occur with administration of bebtelovimab. If clinically significant hypersensitivity reactions occur, discontinue and initiate appropriate supportive care. Infusion-related reactions may occur up to 24 hours post injection. These reactions may be severe or life threatening. (5.1)
- **Clinical Worsening After SARS-CoV-2 Monoclonal Antibody Administration:** Clinical worsening of COVID-19 after administration of SARS-CoV-2 monoclonal antibody treatment has been reported and may include **signs or symptoms of fever, hypoxia or increased respiratory difficulty, arrhythmia (e.g., atrial fibrillation, sinus tachycardia, bradycardia), fatigue, and altered mental**

**status.** Some of these events required hospitalization. It is not known if these events were related to SARS-CoV-2 monoclonal antibody use or were due to progression of COVID-19. (5.2)

- **Limitations of Benefit and Potential for Risk in Patients with Severe COVID-19:** Treatment with bebtelovimab has **not been studied** in patients hospitalized due to COVID-19. Monoclonal antibodies, such as bebtelovimab, **may be associated with worse clinical outcomes** when administered to hospitalized patients with COVID-19 requiring high flow oxygen or mechanical ventilation. (5.3)

<http://www.fda.gov/media/156152/download>



Mixed with cancer cells. Sounds healthy...

It should be fairly clear that, unlike Dr. Ardis' claims, monoclonal antibodies are not effective, carry numerous risky side effects, and can actually worsen the disease they are supposed to treat. Interestingly, this same risk of dangerous side effects and worsening disease outcomes is associated with snake antivenom as well. From the fact sheet of a commonly used antivenom for rattlesnake bites, we find these admitted side effects:

## **Rattlesnake Antivenin Side Effects Center**

“Rattlesnake Antivenin (antivenin crotalidae polyvalent) is an antivenin product used only to treat envenomation caused by bites of crotalids (pit vipers) including rattlesnakes, copperhead and cottonmouth moccasins, and others. **Common side effects of Rattlesnake Antivenin include allergic reactions such as flushing, itching, hives, swelling of the face/tongue/throat, cough, shortness of breath, blue color to the skin, vomiting, and anaphylaxis (severe allergic reaction).**”

“**Immediate systemic reactions (allergic reactions or [anaphylaxis](#)) can occur whenever a horse-serum-containing product is administered.** An immediate reaction (e.g. shock, anaphylaxis) usually occurs within 30 minutes. Symptoms and signs may develop before the needle is withdrawn and may include **apprehension, flushing, [itching](#), [urticaria](#); [edema](#) of the face, [tongue](#), and [throat](#); [cough](#), [dyspnea](#), [cyanosis](#), vomiting, and collapse.** There have been isolated reports of **[cardiac arrest](#)** and death associated with Antivenin (Crotalidae) Polyvalent (equine origin) use.”

“**Serum sickness usually occurs 5 to 24 days after administration and its frequency may be related to the number of Antivenin vials administered.**<sup>30</sup> The incubation period may be less than 5 days, especially in those who have received horse-serum-containing preparations in the past. The usual symptoms and signs are **[malaise](#), [fever](#), urticaria, [lymphadenopathy](#), edema, [arthralgia](#), [nausea](#), and vomiting.** Occasionally, neurological manifestations develop, such as meningismus or peripheral neuritis. Peripheral neuritis usually involves the shoulders and **[arms](#)**. **[Pain](#)** and muscle weakness are frequently present, and permanent atrophy may develop.”

<https://www.rxlist.com/rattlesnake-antivenin-side-effects-d>



Maybe the use of antivenom to treat a snakebite isn't the super cure it has been sold to be? Is it possible that, as with many pharmaceutical products and interventions, the antivenom itself is creating the very symptoms it is said to treat? For some further insight, let's look at a few highlights from an paper from September 2019, right before this "crisis," which reviewed the use of antivenom and had a few revealing claims about the "anti" toxin. You will see it reiterated that the injection of antivenom created from either horse, sheep, goats, and/or rabbits can cause immediate hypersensitivity and anaphylaxis or a delayed "serum sickness" which can occur weeks after the treatment. It is stated that the antivenom has limited efficacy and can be entirely ineffective based on the geographic location. Improper use of antivenom contributes to increased severe outcomes and the production of antibodies in animals leads to a large number (70%) of immunoglobulins that do not react to snake venom:

## **Perspective on the Therapeutics of Anti-Snake Venom**

### **3. Current Information in the Design of New Antivenoms**

**"Currently, the only accepted treatment for snakebite envenomation involves intravenous administration of conventional antivenoms comprising antibodies or antibody fragments derived from the plasma of large mammals (generally horses, but also sheep, goats, or rabbits) that have been previously immunized with non-lethal venomous doses [14,15]. Hyperimmunized animals produce antibodies against the venom proteins and serum is extracted from their blood for the treatment of envenomation [6,16]. Conventional serum therapy aims to bind and neutralize the snake venom proteins [17]. It is a fact that the antivenom allows the body to try to reverse the damage caused by the venom. However, it is known that such therapy can cause problems related to different antivenom characteristics,**

such as:

- **Immediate hypersensitivity reaction to the alien immunoglobulins, including anaphylactic and pyrogenic reactions** such as chills, rigor, headache, and tachycardia. **Delayed antivenom reactions or serum sickness is observed after 8 to 12 days of treatment;** these are characterized by cutaneous eruptions, fever, and allergies, among other effects [18];
- **Limited efficacy of antivenom therapy** to protect the affected organ/s against immediate local tissue damage and low stability;
- **Ineffectiveness of the antivenom** due to significant geographic variation in the composition of the venom;
- **Antigenic reactivity** due to the taxonomic diversity of the snakes;
- **Improper use of the antivenom** due to incorrect medical management, **which contributes to a high incidence of adverse reactions, a low toxin neutralizing potency, or both.**

**“Current antibody production faces challenges during the immunization of the animal (equine or ovine), leading to the production of a huge number of antibodies that are not related to the snake venom. Around 70% of the immunoglobulins obtained do not act directly against venom toxins [26]. Despite the abovementioned facts, this is the only FDA approved therapy to treat snake venom.”**

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6767026/>

A few other studies also point out the severe reactions regularly attributed to the use of antivenom. The first is a study from 2016 which points out that not only are adverse reactions common, they occur at a high rate. It is stated that this is due to poor quality control and manufacturing

problems:

## **Adverse reactions to snake antivenom, and their prevention and treatment**

“Antivenom is the mainstay of treatment of snakebite envenoming. **However, adverse reactions to snake antivenom that is available are common in many parts of the world where snakebite is prevalent.** Both acute (anaphylactic or pyrogenic) and delayed (serum sickness type) reactions occur. Acute reactions are usually mild but severe systemic anaphylaxis may develop, often within an hour or so of exposure to antivenom. Serum sickness after antivenom has a delayed onset between 5 and 14 days after its administration. Ultimately, the prevention reactions will depend mainly on improving the quality of antivenom.”

“**The high rate of acute adverse reactions to antivenom is an example of how poor manufacturing and quality control by antivenom producers cause problems for patients and their doctors.** This highlights the importance of addressing issues related to poor quality and potentially unsafe antivenom. **Ultimately, the prevention of reactions will depend mainly on improving the quality of antivenom.** Until these improvements take place, doctors will have to depend on pharmacological prophylaxis as well as careful observation of patients receiving antivenom in preparation for prompt management of acute as well as delayed reactions when they occur.”

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4767202/>

This next source is from 2018 and it points out that early antivenoms were unsafe and caused severe life-threatening events. While they now have “acceptable” safety profiles, antivenoms still have varying quality and range from 10% adverse reactions to greater than 50%. This same variation in quality is seen in the production of monoclonal antibodies:

## **Antivenom therapy: efficacy of premedication for the prevention of adverse reactions**

**“However, in their initial applications, antivenoms did not exhibit good safety results and could even cause life-threatening side effects [8]. The main reason was that first antivenoms were poorly purified preparations or crude sera. Over the years, for many of the original applications, heterologous serums were replaced by other drugs with better safety profiles, such as antibiotics, vaccines and homologous serums. However, in cases of envenomation by snakes, scorpions or arachnids, antivenoms remain the only effective treatment [4]. Currently, after many improvements, antivenoms exhibit acceptable safety profiles [1, 9, 10]. Nevertheless, antivenom quality still varies widely depending on the producer, while some antivenoms exhibit adverse reaction rates of less than 10%, others have values of greater than 50% [11, 12].”**

<https://jvat.biomedcentral.com/articles/10.1186/s40409-018-0144-0>

In is interesting to note that there are many factors that are said to influence the severity of a venomous snakebite including the age, sex, and health of the person bitten as well as the type of snake, the geographical location of the snake, the season the bite occurred in, what the snake ate, and how recently the snake released its venom. Antivenoms themselves have been shown to have varying effects in quality due to the geographical location of the snake which somehow renders the antivenom ineffective and even dangerous in different countries and continents, even against the same type of snake. It is said that this has kept locals from seeking out medical care and sticking to traditional healers:

“Snake venoms are highly complicated. At least 26 separate enzymes have been identified with 10 of these enzymes common to all snake venoms (though in different

concentrations). All snake bites are not equal. The quality of venom depends not only on the type of snake but on the season, the geographical region, the age of the snake, and how recently it has released venom previously.”

<https://www.vin.com/apputil/content/>

## Antivenom's fatal flaw

“A study led by Dr Fry has found that antivenoms produced using snakes from one region **may perform poorly or fail completely against the same species of snakes from other regions.**

Researchers tested the effectiveness of two African and two Indian saw-scaled viper antivenoms against saw-scaled vipers from 10 regions.

**The results showed that the two African antivenoms were only effective against snakes from restricted ranges.**

**One antivenom performed well against West African saw-scaled vipers and the other performed best against the East African saw-scaled vipers.**

**The Indian antivenom only worked against saw-scaled vipers from the region where the antidote was produced and failed against toxins from other Indian regions. It failed completely against African saw-scaled vipers.**

“These antivenoms are being sold and used interchangeably to treat all saw-scaled viper bites, **and in many cases they are not working,**” Dr Fry says.

“In Kenya, snakebite deaths have increased dramatically after hospitals switched supplies of a very effective African antivenom with a cheaper Indian variety.”

“This creates a knock-on effect in these communities. It's hard enough to convince people living in these regions not

to go to traditional healers to treat snakebite. **And if someone does seek proper medical care but dies because of ineffective antivenom, it will be even harder to convince the next victim to seek out antivenom.**"

## **Viper venom's lethal evolution**

It's the variety of the saw-scaled viper's prey, from rodents to insects, that researchers say could be the reason why antivenom from one region might not work in another.

**"Antivenom is effective and reliable when venom composition does not vary greatly between individual snakes,"** UQ PhD candidate in Toxinology Bianca op den Brouw wrote in an article for The Conversation.

"Unfortunately, the venom composition from saw-scaled vipers **varies between populations** and is **thought to be partly due** to an evolutionary adaptation linked to their diet.

"Different saw-scaled viper populations feed on different prey. The physiology of these prey animals differs, and this dictates what makes a toxin effective.

**"From a medical perspective, this means that the antibodies in an antivenom may not be able to adequately recognise and fight all the harmful toxins in the venom."**

<http://uq.edu.au/research/impact/stories/antivenoms-fatal-law/>

Maybe the proceeding information on how snakebite antivenoms are created as well as the high rate of adverse events from the antibodies used for antivenom now has you questioning that initial "no-brainer" thought: *"Of course I would use antivenom if bit by a snake."* If so, you are on the right track as, based on information from the African Snakebite Institute, in most snake bite cases, antivenom is not used and many snake bites are often unattended and/or unreported. In fact, it is

apparently a well-known “myth” (i.e. truth in this case) that the antivenom kills more people than the snake venom itself. Most people (over 80%) never receive antivenom as, like the previous sources stated, it can have disastrous side-effects. Most snake bites do not cause symptoms warranting the use of something so toxic. In fact, snake bite victims are not immediately injected with antivenom and typically are sent home after observation:

**“Yet people often have a poor understanding of how it works and there are endless myths about antivenom killing more people than the snake venom itself.”**

**“Few snakebite victims are treated with antivenom (less than 20 % of those hospitalised after a snakebite) as most victims are not severely envenomated or the bite may be from a snake that is not considered potentially deadly or is not covered by the antivenom (Rhombic Night Adder, Berg Adder and Stiletto Snake). Antivenom is relatively scarce, expensive and can have disastrous side-effects.** The biggest danger is an acute allergic reaction (anaphylaxis) or, to a lesser degree, serum sickness that can affect the immune system several days after treatment.”

**“Snakebite victims are not automatically injected with antivenom as most of them never experience symptoms severe enough to justify its use.** The majority of snakes have control over their venom glands and are quite reluctant to waste their venom on humans. **They very often give ‘dry’ bites with no subsequent symptoms of envenomation or the snake might inject a little bit of venom that will cause discomfort or some symptoms but nothing serious.** Such patients are usually hospitalised for a day, carefully monitored and then sent home.”

**“As already mentioned, some snakebite victims quickly have an allergic reaction to antivenom and this happens in more than 40% of all cases where antivenom is used.** Some of

those victims go into anaphylactic shock which is a life-threatening medical condition and must be treated with adrenaline. **This has to do with the fact that our antivenom is made from horse blood and the allergy is basically an allergy to horse proteins.**"



Bill Haast – repeated snake bite victim from the world's deadliest snakes tragically died at the young age of 100 from natural causes. ?

If snake bites regularly do not cause symptoms and do not require the use of antivenom, are snake bites really as toxic and harmful as we previously thought? Are the dangerous side effects linked to snake bites really just the reactions to having horse blood injected into the body as treatment? Is this another case where the treatment causes the symptoms of disease it was supposed to prevent? If the examples of these next few individuals are taken into consideration, it's entirely plausible to conclude that we have been misled about the dangers stemming from snakebites in order to cover for the toxic effects of the treatment:



## **Repeated snake bite for recreation: Mechanisms and implications**

“There is a debate in the fatality/immunity due to repeated snake bites in human beings either accidentally or incidentally. **Haast and Winer[11] reported complete recovery of a patient without any specific therapy even after bitten by a deadly snake Bangarus Caeruleus[11]** and the authors attributed it to cross protection of existing antibody between species of Bangarus and Indian, African and Egyptian cobras, as he had a history of bites from these snakes earlier.”

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3883202/>

**This snake-man got himself bitten over 200 times to become immune to venom**

“Bill Haast, a scientist turned snake-man from America, **was bitten at least 173 times by poisonous snakes** in his life till mid-2008 of which **he was fatally injured about 20 times.**”

“In the 1950s, **he had few ill-effects and didnt need any anti-venom** in spite of the fact that he was **bitten by the cobras about 20 times** as per the report published in **Today I Found Out.**”

<https://www.google.com/amp/s/www.indiatoday.in/>

**Man makes deadly snakes bite him 160 times in hunt for human antidote**

“An amateur scientist has **deliberately endured more than 160 self-inflicted snake bites** in a bid to become immune to venom.

Tim Friede is obsessed by finding a human antidote to poisonous snake bites, which kill an estimated 100,000 people every year.

Mr Friede was recently **bitten by a taipan and a black mamba, two deadly snakes** he keeps at his home in Wisconsin, USA, in addition to his two rattlesnakes and water cobra.

He said he experienced a “real throbbing sensation” **but he “felt great” after the bites.**

“It really hurts and it swells **but that’s it,**” he said.”

<https://www.google.com/amp/s/www.independent.co.uk/>

### **Poison pass: the man who became immune to snake venom**

“A lot has been written about [Steve Ludwin](#), **widely known as the man who injects snake venom**, and lately his life has turned into a non-stop frenzy of international journalists and film crews revelling in the seeming sheer insanity of it.”

“**He’s been shooting, swallowing and scratching venom into his skin from some of the world’s deadliest snakes for 30 years.** “Snakes are fucking everywhere. The symbol for medicine is two snakes. They’re ingrained in our brain and DNA,” he tells me, **proudly insisting that he hasn’t been ill for decades and has developed “a superhuman immune system”**. And it’s tempting to believe him. He does look undeniably fit.”

<https://www.google.com/amp/s/amp.theguardian.com/>

### **The Photographer Who Was Bitten by a Black Mamba... and Got the Shot**

“**After several minutes and then hours passed and Laita was still feeling fine** – experts recommend heading straight for a hospital, by the way – the crew concluded that Laita didn’t have any venom in his system. The photographer believes that it was either a “dry bite,” when a snake doesn’t release any venom, or that his heavy flow of blood pushed out the venom.”

[\*The Photographer Who Was Bitten By A Black Mamba... And Got The\*](#)

## Shot

As can be seen, there are numerous examples of people being deliberately and accidentally bitten by the world's deadliest snakes who are completely fine and do not require treatment from antivenom whatsoever. Are we to conclude that these people are the lucky few who somehow have amazing super-human "immune" systems that render snake venom ineffective? Or have snake bites and the associated symptoms of venom toxicity been blown out of proportion? Could this be a case where some have had bad reactions to a snake bite just as there are those who have severe allergic reactions to bee stings while the majority of snake bite and bee sting victims come away unscathed? Could this be similar to the supposed [rabies cases](#) where the majority of those who were bitten by "rabid" animals actually went on to be just fine without getting the rabies vaccination?

## **The Treatments Are Worse Than the Disease**

It's very apparent that in the case of monoclonal antibodies and anivenom, the adverse effects of the drugs are actually worse than the supposed diseases they are meant to treat. Could this be due to the fact that, like "viruses," so-called antibodies have never been properly purified, isolated, and proven to exist? The results of studies using antibodies are regularly unreproducible and irreproducible. It is well-known that antibodies are in fact not as specific as they are claimed to be and are said to regularly bind to the wrong proteins. Perhaps it is difficult to produce safe and effective products when the entities that are supposed to be produced and supplied in the animal blood are entirely theoretical? Maybe the ridiculous snake venom theory should be viewed in the context that it is a bad idea to be injecting anything, let alone animal blood, into our bodies in an attempt to make ourselves feel better when trusting the body and allowing it to heal is often times the best course of

action we can take.

### **In Summary:**

- Dr. Bryan Ardis put forth a theory that snake venom is the cause of “Covid-19” primarily **based on fraudulent genomic data**
- The snake connection stems from research linking proteins from the fabricated “SARS-COV-2” genome to **bat and snake “coronavirus” proteins**
- The enzyme phospholipase A2 group IIA or sPLA2-IIA, which Dr. Ardis bases much of his claims on, only has **similarities** to rattlesnake venom
- These peptides are “almost identical” to the venoms of animals and **are regularly found in healthy humans and other mammals**
- Dr. Ardis pointed out that, based on a text, he uncovered the connection between antivenom and monoclonal antibodies and stated that **they are the same thing**
- He **wrongly** concluded that monoclonal antibodies **are an effective treatment** for snake poisons that could be in the vaccines, Remdesivir, and water
- According to a Sept 2021 Cochrane Review, **their certainty in the evidence for the use of monoclonal antibodies** in the treatment of “Covid” for all non-hospitalised individuals **was low**, and for hospitalised individuals was **very low to moderate**
- They considered the current evidence **insufficient to draw meaningful conclusions** regarding treatment with “SARS-CoV-2-neutralising” mAbs
- Monoclonal antibodies are known to cause a range of side effects and reactions, **which can be immediate or delayed**
- Serious adverse events associated with mAbs include **infusion reactions, acute anaphylaxis, and serum sickness**, as well as longer-term complications such as **infections, cancer, autoimmune disease, and**

## **cardiotoxicity**

- In February 2022, the FDA revised the authorizations for two monoclonal antibody treatments – bamlanivimab and etesevimab (administered together) and REGEN-COV (casirivimab and imdevimab) – **to limit their use to only when the patient is likely to have been infected with or exposed to a variant that is susceptible to these treatments**
- The data showed these treatments are **highly unlikely to be active against the omicron variant** which is circulating at a very high frequency throughout the United States
- These treatments are **not authorized** for use in any U.S. states, territories, and jurisdictions at this time
- Monoclonal antibodies are **laboratory-made proteins that mimic** the immune system's ability to fight off harmful pathogens
- In April 2022, the U.S. Food and Drug Administration **revoked the emergency use authorization (EUA) that allowed for the investigational monoclonal antibody therapy bamlanivimab**, when administered alone, to be used for the treatment of mild-to-moderate "COVID-19" in adults and certain pediatric patients
- Based on its ongoing analysis of emerging scientific data, specifically the sustained increase of "SARS-CoV-2 viral" variants that are resistant to bamlanivimab alone **resulting in the increased risk for treatment failure**, the FDA determined that the known and potential benefits of bamlanivimab, when administered alone, **no longer outweigh the known and potential risks for its authorized use**
- Importantly, although the FDA revoked this EUA, alternative monoclonal antibody therapies remain available under EUA, including REGEN-COV (casirivimab and imdevimab, administered together), **and bamlanivimab and etesevimab, administered together**, for the same uses

as previously authorized for bamlanivimab alone

- *In other words, the use of Bamlanivimab and Etesevimab was revoked as well as the use of Bamlanivimab but they can still be used together as an alternative to Bamlanivimab alone...*
- For the Omicron-specific Bebtelovimab authorized by the FDA in February 2022, possible side effects include
  1. Itching
  2. Rash
  3. Infusion-related reactions
  4. Nausea
  5. Vomiting
- Serious and unexpected adverse events including **hypersensitivity, anaphylaxis and infusion-related reactions** have been observed with other “SARS-CoV2” monoclonal antibodies and could occur with bebtelovimab
- In addition, **clinical worsening following administration of other “SARS-CoV-2” monoclonal antibody treatment has been reported** and therefore is possible with bebtelovimab
- The FDA claims that **it is not known** if these events were **related to “SARS-CoV-2” monoclonal antibody use** or were due to progression of “COVID-19”
- Signs or symptoms of worsening outcomes include:
  1. Fever
  2. Hypoxia or increased respiratory difficulty
  3. Arrhythmia (e.g., atrial fibrillation, sinus tachycardia, bradycardia)
  4. Fatigue
  5. Altered mental status
- Treatment with Bebtelovimab **has not been studied** in patients hospitalized due to “COVID-19”
- Monoclonal antibodies, such as Bebtelovimab, **may be associated with worse clinical outcomes** when administered to hospitalized patients with “COVID-19” requiring high flow oxygen or mechanical ventilation

- Antivenom carries the **same risks of severe side effects and worsening condition** as monoclonal antibodies
- The listing for common side effects of Rattlesnake Antivenin include allergic reactions such as:
  1. Flushing
  2. Itching
  3. Hives
  4. Swelling of the face/tongue/throat
  5. Cough
  6. Shortness of breath
  7. Blue color to the skin
  8. Vomiting, and anaphylaxis (severe allergic reaction)
- Immediate systemic reactions (allergic reactions or anaphylaxis) can occur **whenever a horse-serum-containing product is administered**
- There have been isolated reports of **cardiac arrest and death** associated with Antivenin (Crotalidae) Polyvalent (equine origin) use
- **Serum sickness usually occurs 5 to 24 days after administration** and its frequency may be related to the number of Antivenin vials administered
- The usual symptoms and signs are:
  1. Malaise
  2. Fever
  3. Urticaria
  4. Lymphadenopathy
  5. Edema
  6. Arthralgia
  7. Nausea
  8. Vomiting
- Occasionally, **neurological manifestations develop**, such as meningismus or peripheral neuritis
- **Peripheral neuritis** usually involves the shoulders and arms and pain and muscle weakness are frequently present, **and permanent atrophy may develop**
- A 2019 review on antivenom stated that currently, the

only accepted treatment for snakebite envenomation involves intravenous administration of conventional antivenoms comprising antibodies or antibody fragments **derived from the plasma of large mammals (generally horses, but also sheep, goats, or rabbits) that have been previously immunized with non-lethal venomous doses**

- It is known that such therapy can cause problems related to different antivenom characteristics, such as:
  1. **Immediate hypersensitivity reaction to the alien immunoglobulins, including anaphylactic and pyrogenic reactions** such as chills, rigor, headache, and tachycardia.
  2. **Delayed antivenom reactions or serum sickness is observed after 8 to 12 days of treatment;** these are characterized by cutaneous eruptions, fever, and allergies, among other effects
  3. **Limited efficacy of antivenom therapy** to protect the affected organ/s against immediate local tissue damage and low stability
  4. **Ineffectiveness of the antivenom** due to significant geographic variation in the composition of the venom;
  5. **Antigenic reactivity** due to the taxonomic diversity of the snakes
  6. **Improper use of the antivenom due to incorrect medical management, which contributes to a high incidence of adverse reactions, a low toxin neutralizing potency, or both**
- Current antibody production faces challenges during the immunization of the animal (equine or ovine), **leading to the production of a huge number of antibodies that are not related to the snake venom**
- Around 70% of the immunoglobulins obtained **do not act directly against venom toxins**
- According to a 2016 study, **adverse reactions to snake antivenom that is available are common** in many parts of



the world where snakebite is prevalent

- The **high rate of acute adverse reactions to antivenom** is an example of how poor manufacturing and quality control by antivenom producers cause problems for patients and their doctors
- The prevention of reactions **will depend mainly on improving the quality of antivenom**
- According to their initial applications, antivenoms **did not exhibit good safety results and could even cause life-threatening side effects**
- Currently, after many improvements, antivenoms **exhibit “acceptable” safety profiles** yet antivenom **quality still varies widely** depending on the producer, while some antivenoms exhibit adverse reaction rates of less than 10%, **others have values of greater than 50%**
- All snake bites **are not equal** and the quality of venom depends not only on the **type of snake** but on the **season**, the **geographical region**, the **age** of the snake, and how **recently it has released venom previously**
- A study led by Dr. Fry found that antivenoms produced using snakes from one region **may perform poorly or fail completely against the same species of snakes from other regions**
- The results showed that the two African antivenoms **were only effective against snakes from restricted ranges**
- One antivenom performed well against West African saw-scaled vipers and the other performed best against the East African saw-scaled vipers
- The Indian antivenom only worked against saw-scaled vipers from the region where the antidote was produced and **failed against toxins from other Indian region and it failed completely against African saw-scaled vipers**
- “These antivenoms are being sold and used interchangeably to treat all saw-scaled viper bites, and **in many cases they are not working,**” Dr Fry said
- If someone does seek proper medical care **but dies because of ineffective antivenom, it will be even harder**

**to convince the next victim to seek out antivenom**

- **Antivenom is effective and reliable when venom composition does not vary greatly between individual snakes**
- **Unfortunately, the venom composition from saw-scaled vipers varies between populations and is thought to be partly due to an evolutionary adaptation linked to their diet**
- **From a medical perspective, this means that the antibodies in an antivenom may not be able to adequately recognise and fight all the harmful toxins in the venom**
- **There are endless myths about antivenom killing more people than the snake venom itself**
- **Few snakebite victims are treated with antivenom (less than 20 % of those hospitalised after a snakebite)**
- **Antivenom is relatively scarce, expensive and can have disastrous side-effects**
- **Snakebite victims are not automatically injected with antivenom as most of them never experience symptoms severe enough to justify its use**
- **Snakes very often give 'dry' bites with no subsequent symptoms of envenomation or the snake might inject a little bit of venom that will cause discomfort or some symptoms but nothing serious**
- **Such patients are usually hospitalised for a day, carefully monitored and then sent home**
- **Some snakebite victims quickly have an allergic reaction to antivenom and this happens in more than 40% of all cases where antivenom is used**
- **This has to do with the fact that antivenom is made from horse blood and the allergy is basically an allergy to horse proteins**
- **Haast and Winer reported complete recovery of a patient without any specific therapy even after bitten by a deadly snake *Bangarus Caeruleus* and the authors attributed it to cross protection of existing antibody between species of *Bangarus* and Indian, African and**

Egyptian cobras, as he had a history of bites from these snakes earlier

- Bill Haast, a scientist turned snake-man from America, was **bitten at least 173 times by poisonous snakes** in his life till mid-2008 of which he was seriously injured **about 20 times**
- In the 1950s, he had **few ill-effects and didnt need any anti-venom** in spite of the fact that he was **bitten by the cobras about 20 times**
- An amateur scientist named Tim Friede **deliberately endured more than 160 self-inflicted snake bites** in a bid to become immune to venom
- Mr Friede was recently **bitten by a taipan and a black mamba**, two deadly snakes he keeps at his home in Wisconsin, USA, in addition to his two rattlesnakes and water cobra
- He said he experienced a “real throbbing sensation” **but he “felt great” after the bites**
- Steve Ludwin, widely known as the man who injects snake venom, **has been shooting, swallowing and scratching venom into his skin from some of the world’s deadliest snakes for 30 years**
- He **hasn’t been ill for decades** and has developed “a superhuman immune system”
- A photographer was bit by the deadliest snake, a Black Mamba, and after hours passed, **he was still feeling fine** and needed no treatment

The snake venom theory by Dr. Bryan Ardis is built upon the interpretation of the unpurified fraudulent “SARS-COV-2” genome which is itself built upon references to other fraudulent genomes of human and animal “coronaviruses” created in the very same way. Attempting to claim any connections between the random A,C,T,G’s in a computer database is a useless and pointless exercise as the RNA that was fabricated into the genome of a “virus” was never purified, isolated, and proven to physically exist in the first place. Thus any

connections between the protein codes said to belong to a "virus" which are then said to be closely related to supposed snake "coronaviruses" is immediately invalid.

Using this invalid premise to then claim that people have been poisoned by snake venom in the vaccines, the drugs, and the water supply is nothing but unsubstantiated science fiction that seems designed to have a few purposes:

1. To keep people engaged in the lie that a new disease known as "Covid-19" exists and that there is a singular cause.
2. To restore faith in monoclonal antibodies and other toxic alternative treatments.
3. To use the theory to promote and sell anti-venom supplements.
4. To divide and distract those questioning the official narrative.
5. To make the "Truther" community look foolish by falling for loosely tied-together circumstantial evidence that is easily debunked.

If we are to take the claims of Dr. Ardis seriously that the symptoms associated with snake venom is the true cause of a disease known as "Covid-19," how does his theory explain for the fact that the antivenom and monoclonal antibody treatments cause the exact same symptoms of the disease they are supposed to treat? How would it be determined that the worsening clinical outcomes after injection are from the snake bites/venom rather than the antivenom/monoclonal antibodies given as treatment? How does his theory account for the numerous instances where people have been deliberately bitten by snakes, injected with the venom of snakes, and drank of the venom of the snakes with little to no harmful effects whatsoever? How does his theory account for the fact that the vast majority of "Covid" cases are asymptomatic and the vast majority of snake bite cases need no treatment at all? There are many holes in this theory which will easily be picked

apart to make those who follow it look foolish for having done so.

There is no "SARS-COV-2." There is no "Covid-19." There is no new disease nor any new symptoms of disease requiring treatment from vaccines, monoclonal antibodies, Remdesivir, Hydroxychloroquine, Ivermectin, NAC, nor any other treatment. There is no need for any anti-venom supplements.

Beware those who will sell you the cause of the disease and the solution.

**[Connect with Mike Stone](#)**

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