

## Pfizer COVID-19 mRNA Vaccine Blood Clots in the Heart Up to 40cm Long Clots

Surviving a Pfizer clot shot heart attack. Pfizer's Eliquis is now world's 6th best selling drug!

By [Dr. William Makis](#)

Theme: [Science and Medicine](#)

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[COVID Intel](#)

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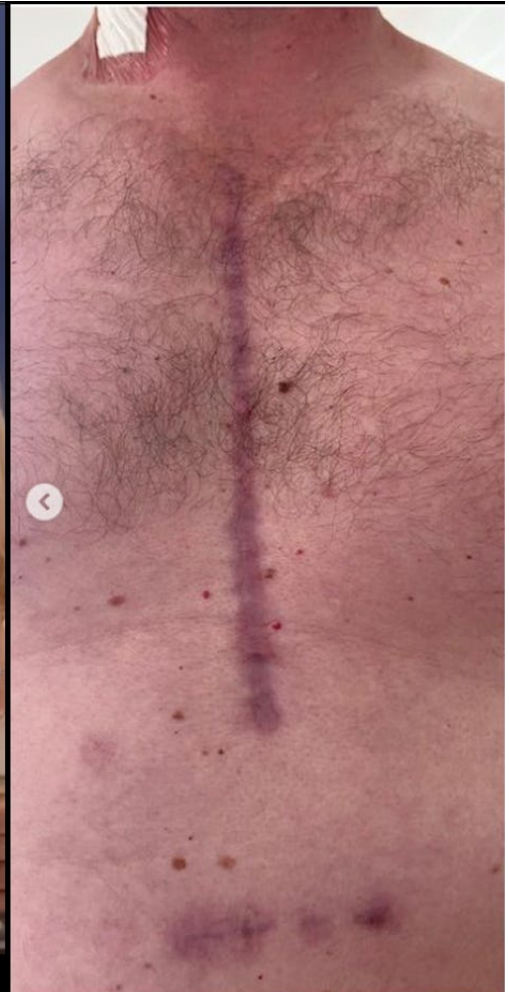
*Feb. 2023 - Sydney, NSW, Australia - 42 year old Darius works 5-7 days a week abseiling buildings in construction. He was bodybuilding and training 6 times a week. He had two Pfizer vaccines and barely survived a heart attack 2 weeks later - needed 5 stents put in*

- Darius was an extremely fit 42 year old
- 2 weeks after his 2nd Pfizer mRNA, he had a heart attack, 99.3% blockage and had to have 5 stents put in
- he still has chest pains up to 10/10 to this day
- he has been on anti-depressants and has been suicidal

<p>My name is Darius. I'm 42 years old and live in Kogarah, Sydney. I was working 5-7 days a week obsselling buildings in the construction industry. You need to be extremely fit to do this work because it's very physically demanding. Prior to the 🏠, I was body building and training 6 times a week at the gym, rode a push bike for 12km twice a week, and walked 9-12km a week.</p> <p>I had no underlying heart conditions. In fact, in 2020 before undergoing surgery after a car accident, I had a pre-op heart test where a dye was injected to ensure I had no artery blockages. I got the all clear. There's also no family history of heart disease.</p> <p>In 2021, construction workers were mandated to get the 🦠, so I got my first 🦠 on the 19th of July 2021, 2:02pm (left arm) at RPA Mallet Centre (Newtown Hospital). Six hours later, I started to feel very light headed and I couldn't put one foot in front of the other without losing my balance. The next day was the same. On the 3rd day, I got my balance back but I lost my voice, had a high temperature and sore joints. I felt like a 90 year old and like I'd been hit by a truck. I could only perform light duties at work and it took about 2 weeks to feel like myself again.</p>	<p>I had my second 🦠 on Monday, 9th of August (left arm) at 3:12pm. When I got home a few hours later, I felt light headed and unsteady on my feet again. My brain felt foggy and confused, and I was so fatigued I went straight to bed. I had the sweats and nightmares throughout the night, and woke the next day to find I'd lost my voice completely. My voice didn't return for 8 days.</p> <p>After the first 🦠, I was short of breath just climbing a few flights of stairs. I had sore joints and sweated profusely. These symptoms all worsened after the second 🦠. About 2 weeks after the second 🦠, I had 4-5 days where I felt almost back to normal.</p> <p>My boss told me on the 25th of August, I was speaking really fast, mumbling a lot, and jumping from one topic to another. He said I was incoherent.</p> <p>The next day, on the 26th of August, the worksite was closed due to rain so I had the day off and planned a 12km walk with my boss at around 7:30am. Before my walk, I did a set of 4 bicep curls but after my 8th rep, I felt a sharp pain in my chest. The pain was so bad I had to lay down and I felt as if the world was caving in on me. I had a shooting pain going up and down my veins between my wrist and elbow on my left arm, and my chest felt like a screwdriver was twisting from side to side into my heart. The pain was excruciating and lasted about 15 minutes.</p>	<p>I stepped outside for some fresh air and ended up vomiting up blood, phlegm and foam. The pain then left me. My initial thought was that what I threw up didn't seem normal which concerned me, but because I was no longer feeling pain, I thought I should be okay. I went ahead with my initial plan to go for a walk with my boss. I told him, "I think I just had a heart attack" but didn't think more of it as I was no longer feeling any pain. That same night from 12pm onwards, I was waking throughout the night with chest pains. When I woke on the morning on the 27th of August, I was sweating and very fatigued, but despite the chest pains (I have a high pain threshold), I thought I'd be okay to drive to work. Five minutes into the drive, I pulled over at a service station because the chest pains became excruciating. I called St George Hospital to book an appointment for after work but after hearing my symptoms, the nurse told me to turn off my car immediately and I was not to drive because everything I'd told her indicated I was having a heart attack. She called me an ambulance and the initial tests done by the paramedics showed all my vital signs to be fine. When I arrived at the hospital, the nurse took one look at me and said "he's having a heart attack" and immediately ordered I be given blood thinners. The blood tests came back showing my heart was bleeding blood and proteins — this indicates a heart attack.</p>
<p>I was told my artery had narrowed 99.3%. I was taken directly from emergency to the operating theatre. On the way, they asked for permission to cut my clothes off me and I was injected with a blood dye to assist in manoeuvring a camera through my veins to show which artery needed to be opened first with a stent. I was awake for the procedure to ensure my heartbeat remained as stable as possible, and my heart rate and pain were closely monitored.</p> <p>I was taken to the cardiac ward afterwards and I was told they'd be reporting my case as an adverse reaction to the 🦠 database. The doctors said because I had no underlying conditions, they had reason to believe my symptoms were a result of the 🦠 and this was put in writing. They likened my heart to that of a 97 year old with heart issues. I was told 4 other arteries had narrowed and I'd need further surgery ASAP.</p> <p>On Monday, the 30th of August at 9am, I underwent the same procedure and was in surgery for 4-4.5 hours. They went through my groin area as my wrist was still healing from the procedure I had a few days earlier. The pain medication was wearing off and I was growing impatient, not realising they were inserting stents to open up all 4 arteries at the same time. I was told the damage was very bad, and I was in hospital for 6 days post operation.</p>	<p>After the procedures to insert the 5 stents, I couldn't work for 3 weeks because I wasn't to lift anything during that time. On the first day back, I was on the ropes when I felt a really sharp pain in my chest. I thought I'd moved one of the stents when I reached out my arm to point something. After persevering with the chest pain for 3 days at work, I went back to the hospital because I was told to go back if I had any pain.</p> <p>I was seen to immediately and a camera was inserted through my right wrist to investigate. They found a 6th artery had narrowed since my last operation to insert the 5 stents 4 weeks prior. This most recent narrowing artery sits in-between two major arteries which already had stents. I was told they couldn't open this 6th artery because the stents in the two major arteries on either side would likely collapse and kill me. I was advised to find tools and coping strategies to live with the pain. I was in hospital for 3 days (discharged 29th of September). On a positive note, they said the other stents all looked good.</p> <p>I wasn't to lift more than 300g with that arm after the most recent surgery so I was off work for 3 weeks. Before I was due to go back, I woke at 6am one morning and couldn't breathe! I went to the mirror to find my tonsils were the size of 50 cents coins and blocking my airways.</p>	<p>I went straight to hospital and was immediately given steroids. The swelling went down after 6 hours. The doctors and nurses said they'd never seen tonsils swell so suddenly, particularly as I hadn't been unwell and didn't have a sore throat prior. This was about 2.5 months after the 2nd 🦠. I went back to work 3-4 days later.</p> <p>I'd arranged with my cousin to drive to Melbourne to see my daughter on the 30th of November. On the day of the trip, I felt very sick when I woke up. I felt light headed and nauseous. At 9am, whilst on the ropes at work, I was still feeling quite unwell when I vomited blood and foam (similar to when I had my initial heart attack). It was decided I'd come down to supervise instead of going on the ropes. It was a hot day, so I was sitting in the ute with the air conditioner on, but I felt myself drifting in and out of consciousness and had to try keep myself awake throughout the day. I called my mum at 11-12pm and told her, "I think I'm bleeding inside".</p> <p>Because everything was booked and paid for, and I was really looking forward to meeting my 4 month old granddaughter for the first time, we made the decision to go ahead with our trip. We left Sydney in two separate cars at 2pm and after driving 2.5 hours we stopped at the Yass service station.</p>
<p>The minute I got out of the car, my head was spinning like crazy. I felt hot and cold, nauseous and extremely sick. My cousin was parked about 20 metre in-front of me but I struggled to walk to him. When I reached his car, I barely managed to open the car door before collapsing forward onto the back seat. I couldn't physically hold myself up another second. We made the decision to leave my car at Yass and my cousin would drive, with me travelling in the back seat.</p> <p>We stopped at a servo at 1am and as soon as I stood up, I collapsed, falling backwards onto the ground twice. My cousin helped me up but I fell forward a third time, this time hitting the front passenger side door and mirror with my head, slumping to the ground like a sloth! My cousin thought I was having a heart attack because I was incoherent and mumbling. He drove me to a nearby country hospital where they ran blood tests that showed I was bleeding internally. With no doctors available, an ambulance was called to take me to a Melbourne hospital. During the 2 hour trip, my heart nearly stopped 6 times and the paramedics had to keep waking me to keep me conscious. Along the way, a second ambulance was called as they needed senior paramedics to assist on the way to the hospital.</p>	<p>I was taken straight to the operating theatre upon arrival at The Northern Hospital and I gave consent to be given a blood transfusion. I was given a whole bag of blood in 15 minutes — this is normally done over an hour but I needed blood urgently. It was very painful. A camera was inserted down my throat and they found the internal bleeding was due to my stomach being slashed and burst ulcers from the blood thinners. I was given a further 5 bags of blood over the next 4 days, and treated with laser burning to seal up the damage to my stomach. I was in hospital for 7 days and I stayed with my daughter for 2 weeks following surgery. I wasn't able to work for 2 months.</p> <p>In mid January 2022, I had internal bleeding again from the blood thinners and was admitted to St George hospital. They used laser surgery to stop the bleeding. My heart surgeon said to stop the really strong blood thinners and put me on Nurofen, but I was put back on the strong blood thinners 4 months later.</p> <p>I have to take blood thinners for the rest of my life. I need medication to coat my stomach to reduce the risk of internal bleeding from the thinners, and I'm on blood pressure tablets. My condition also makes me sweat excessively, so I have to change my shirt at least 3-4 times a day. I sweat just sitting on the couch whereas prior to the 🦠, I could do a workout without breaking a sweat.</p>	<p>I'm an anti depressants because I was so depressed I wanted to commit suicide. But when I thought of my daughter and granddaughter, and the surgeons who worked so hard to save my life, I decided not to.</p> <p>My cardiologist said to me on 3-4 occasions, "We'll try make you live to 60." I was absolutely dumbfounded and in a complete state of shock. It's taken me a long time to process his words — that I have about 15 years to live.</p> <p>Every single day, I live with chest pains that never fully go away. Some days the pain stops me in my tracks and I can't even hold a conversation. My pain can range from a 5 (on a 10 scale) to a 9 or 10 which feels like I'm having a heart attack. I had noticed when I was at my mum's place, being around her pets helped with my pain so in December 2022 I decided to get two cats of my own. They are an addition to my arsenal of coping strategies for my chronic pain — keeping me busy and helping distract me from my chest pain. Animals are a blessing with their undeniable healing influence.</p> <p>If the pains feel different or more intense than what I already experience daily, particularly if I get the feeling of the world caving in on me like when I had my first heart attack, I will go back to hospital to be checked. The most recent hospital visit was only in December 2022 — thankfully the tests came back okay.</p>
<p>My life has been forever changed. I will never be 100% of my old self ever again. Prior to the 🦠, I was making a comfortable living doing work I love, going to the gym and coffee shops, and just enjoying life. I was forced to take the 🦠 so I could work to be able to eat, pay the rent and bills. Since the heart attacks and health issues, I fatigue easily and have constant chest pains — I can barely work enough to support myself.</p> <p>I have been denied compensation despite my application rejection letter clearly stating, "The reporting practitioner has provided opinion that the most likely cause of the diagnosed injury(s) is 🦠. Other causes have been considered and appear less likely to have contributed to this individual's diagnosis." Their reason is that my condition of myocardial infarction is not listed as a claimable clinical condition in relation to the 🦠.</p> <p>My hope is to find enough injured victims who are courageous enough to take a stand to bring about accountability and compensation for their ongoing mental and physical hardships and suffering.</p> <p>I also hope my story raises awareness of the common signs of a heart attack, so if anyone reading this experiences similar symptoms, they take it seriously and seek medical attention.</p> <p>I am just a number, but I am still hurting and all I ask is to be treated with dignity.</p>		

Nov. 25, 2022 - Margaret River, WA, Australia - 51 year old Mark was in peak health and fitness, worked on an Oil Rig as a Supervisor. Had three Pfizer mRNA jabs (Pfizer #1 FH3221, #2 FK6268, #3 1F1056A).

- Got the Pfizer mRNA Booster on March 29, 2022
- started having shortness of breath
- April 26, 2022 hospitalized, April 27 open heart surgery to remove 30-40cm clot that was putting pressure on the heart
- His Vaccine Injury compensation application was denied by Australian govt.





<p>I'm 51, a husband to my lovely wife Carissa, and father of two children. Prior to the 🦠👤, I was in peak health and fitness and had no medical issues at all. I work FIFO on an Oil Rig as a Maintenance Supervisor. It is common procedure to have regular medical check-ups, and these have always come back good. Before this job, I served in the navy for 8 years as a Sub-mariner, so fitness was always a priority of mine. I enjoyed mountain bike riding and doing regular gym work outs.</p> <p>I got the job because of the work mandate, I held out as long as I could in hope that the mandates would drop. I am not an anti-👤 as I had taken other 🦠 in the Navy, but this new one seemed rushed through and it concerned me. Reluctantly, I had to get the 🦠👤 on the 1st of December 2021 or lose my job. I felt nothing, 3 weeks went by and I went ahead to get the second 🦠👤 on the 23rd of December 2021. The next morning, I felt fatigued and hung over. I hoped I would feel right for Christmas. I thought it would pass as I had reassurance from others on the rig that they had similar experiences.</p>	<p>I pushed through and got the 🦠👤 on the 29th of March 2022. The next day I developed flu like symptoms and a persistent cough. I pushed through and thought this too would pass. I noticed when I was doing the regular task of cutting up firewood that I struggled, I suddenly had shortness of breath and had to rest on the back of the ute. It occurred to me that maybe I was having a reaction to the 🦠👤 as I had never felt this before. Since I had heard many stories about dismissive behaviour from medical staff, I didn't go to the GP. Though I managed to have good sleep, I felt aches in my body particularly around the rib cage and my cough lingered. I thought it too would pass and I prepared to go back to work on the Monday 4th of April 2022. By then even though the cough was still there, I was feeling less achy, so I drove 3 hrs to Perth, caught a 2 hr Qantas flight to Karratha then a 30-minute helicopter ride to the Oil rig. My role as a supervisor involves lots of office work, some site visits and addressing the team in meetings.</p>	<p>On 8th of April 2022, I worked out doing a gym circuit on the helideck and towards the end, my calf muscle was sore, but as it was towards the end of the session, I pushed through thinking it was a strain. It caused a limp, but I persevered with it for 3 days before the pain stopped. By this time, I was more concerned about my cough as it appeared to be getting worse, I thought I could have asthma and shared my concern with my wife, but I had no prior history of it. A few times I also struggled to address my crew, in the meetings as standing and reading was making me short of breath. It started to take its toll on me. I could not easily ascend stairs or climb ladders anymore.</p> <p>On Friday 22nd of April, I went to the gym and started with the row machine. I aimed to complete 2000m but only managed to do 500m before really needing to stop as I was so short of breath. It all appeared to be getting worse. The following week on the 26th of April 2022, I went to the medic on the rig and expressed my concerns, relating the symptoms back to the 🦠👤 when it all started.</p>
<p>Regular covid testing (RAT) had already been done prior to arriving and had already been ruled out due to negative test results. The doctor did an Oxygen Saturation Test and found that my oxygen levels were dropping to 92% well below normal range. He relayed the results to a doctor in Karratha which was protocol. But it was left to me to make the call. My manager approached me and asked, "Do you want to get off the rig?" He had noticed me struggling at the meetings. As I had 8 days to go and sitting behind the computer at work was manageable, so I thought I could stick it out. That same night my cough started to worsen and kept me awake. I called the medic late and asked for cold and flu tablets. I propped myself up on pillows and this helped a bit. The following day, I was encouraged to leave the rig and a helicopter made a diversion to pick me up from the rig to take me to Karratha Airport. I had to carry my bag from the helipad to the airport building about 800m away and it took me 5 rest stops to get my breath back each time.</p>	<p>My plan was to see my local doctor who had my medical history, so I took a flight to Perth airport and met my wife there. I felt awful at this stage. I had chest pains, felt clammy and struggled to breathe. It was a difficult decision to make deciding on going to Perth hospital which had been in the news for lengthy wait times in ER, so we headed to Bunbury Hospital 1.5 hrs away and went straight to emergency. We arrived there at 10pm and I was taken away in a wheelchair, but the triage nurses would not allow my wife to enter the hospital due to her non 🦠👤 status. Horribly, she was sent to wait in the car park for further updates.</p> <p>After relaying the current symptoms, the medical staff did an Oxygen Saturation Test and it was again seriously low. Even with the application of an oxygen mask the levels did not improve, so they fitted oxygen tubes into my nose. Next a CT scan was done to get a better picture of what was happening. Very quickly a diagnosis was given, that the condition was a Pulmonary embolism in my lungs. I was told that I was very sick as there were blood clots in my lungs and calf and they highly recommended that I be airlifted by the Royal Flying Doctors Service, to Sir Charles Gardener Hospital in Perth the next day. I started on blood thinners that night.</p>	<p>A nurse met Carissa in the car park, to give her my update and suggested going home to Margaret River, as there was nothing she could do there.</p> <p>On the 27th of April 2022, I arrived at Sir Charles Gardener Hospital at 8am. The hospital was a Cardiothoracic Specialist Hospital and there were many people waiting in the ER. My serious case was taken to four surgeons who discussed the options and gave me three courses of action. The first was to just take blood thinners but they mentioned just doing that would leave a 1 in 6 chance of dying. The second was to take medication usually given to stroke victims to dissipate clots but the side effects were mainly the chance of a brain bleed, the last option was to clear the blood clots surgically by removing the heart and cleaning out the clots. The deeper ones in the lungs would be mopped out by the blood thinners. It was a difficult decision to make, however the best course of action was the third, and it was even harder to tell my wife and family that over the phone. During discussions with the surgeons, I had related symptoms to all starting with the 🦠👤. One of the surgeons did firmly say to me that my condition was not due to the 🦠👤 as I had had 🦠 and not 🦠. I wasn't going to argue!</p>
<p>I was in theatre at 9am the same day and the open-heart surgery took 5-6 hrs. The surgeons removed a 30-40cm blood clot which was placing pressure on the heart and inserted an IVC filter in the veins to stop clots travelling from the calf to the heart. It would have to remain for 6 months and then be removed. The severity of blood clots was a real mystery to the surgeons as usual causes could be a flight from long flight or presence of cancer, so a lymph node test was done which came back negative 2 weeks later. Overall, the surgery was a success.</p> <p>Whilst I was in an induced coma for two days and spent 3 days in ICU and 5 days in the High Dependency Unit, my wife travelled to Perth. She was refused entry once again and left to rely on my daughter to be given updates. One ICU nurse tried to arrange entry, but it was met with backlash from other medical staff. After an 8-day hospital stay, I was moved to an Airbnb for post-surgery treatment. A nurse visited every day to monitor my progress and change the dressings.</p>	<p>For the next 5 months, I focused on my recovery from open heart surgery. I was advised to rest and do light walking only. I used up all my sick leave. I had regular check-ups. A month later, I did a follow up CT scan at Busselton Hospital. There was still concern over some clots in the lower half of the lungs, but I was discharged as I was still on blood thinner medication. I returned to work again on the 29th of September 2022.</p> <p>On the 11th of November 2022, I went in for a day procedure to remove the IVC filter at the 6-month mark. I have continued to take the blood thinner called 'Apixaban' and after following up with the Haematologist and Immunologist, they have recommended to take this for the rest of my life. Both specialists recognised that the 🦠👤 was what provoked the pulmonary embolism and gave a written medical report stating this. Disappointedly, the application for the Adverse Reaction Compensation Scheme was denied.</p>	<p>At present, I feel back to my normal self, though still have a slight pain at the base of my sternum. The GP thinks it is just tenderness from the surgery. I can do the regular activities like mountain bike riding and cutting timber once again without the shortness of breath, though I must be mindful of what I do, due to being on blood thinners.</p> <p>I share my story because I want to bring the truth and awareness of what is happening to others. Where there is risk there must be choice and I stand by this 100%. Hopefully my story can help make a difference.</p>



Nov. 25, 2021: Myrtle Beach, SC, Tim - "4 weeks after my last Pfizer vax, these passed through my heart, and I was subsequently life flighted to a Trauma Care Unit where my chest plate was knocked open and these were vacuumed out of my chest."



**Tim** ✓  
@TimSurvivedThis

...

4 weeks after my last Pfizer vax, these passed through my heart, and I was subsequently life flighted to a Trauma Care Unit where my chest plate was knocked open and these were vacuumed out of my chest....The CLOT SHOT is real.

**November 25, 2021**

12:45 PM

Ec



**Tim** ✓

@TimSurvivedThis

True Conservative....Not What These People Posing as Conservatives call themselves Today....I mean a Reagan Conservative....2 Genders....Read people very Well

📍 Myrtle Beach, SC 📅 Joined February 2023

363 Following 955 Followers

Oct. 21, 2021 - Gosford, NSW, Australia - 19 year old Cienna Knowles was a professional equestrian. She got 2 Pfizer mRNA Vaccines (#1: FG3712, #2FG6431) and developed blood clots in the lungs and the heart

- Cienna had the 2nd Pfizer mRNA on Oct. 21, 2021
- Oct. 24, 2021 she was told she has pulmonary embolism
- Nov. 11, 2021 she was told 3 of 5 arteries in her heart were 100% blocked
- She was put on Eliquis (Apixaban), Pfizer's anti-coagulant drug





I was 19 when I had the 🍌. I am 20 now. I was a professional elite equestrian athlete prior to the 🍌. I'd won champion multiple times and I'm obsessed with horses. As well as riding, I have always been insanely into fitness, running, going to the gym and motorbikes. I had also played touch football for the 'Newcastle Knights' and had represented my school in many different sports. I've always had an excellent immune system, had never been sick before and had no pre-existing conditions. The only medication I'd been on for about 12 months prior to taking the 🍌.

I started an office job in June 2021 and was verbally told by my manager that my role was menial, so it had come under review. I didn't receive anything formal in writing from the company stating this, only verbally and via text from my manager that I needed to be fully 🍌 by the 1st October 2021 to remain employed. Everyone at work knew I did not want to have this 🍌. I'd never had an issue with any other 🍌 and had the flu 🍌 a couple of years back when it was a severe flu season. At school I would always volunteer to go first as I wasn't scared of them and everyone else was.

I had no immediate reaction but woke up at midnight in my own sweat with pain in my joints and muscles, like I had been running a marathon and struggled to move - I couldn't walk or straighten my arms. I had a fever, was vomiting, had sore eyes, an insane headache, all over weakness and a severe pain in the left side of my chest - my heart was racing and I could feel my pulse through my entire body. It was painful to breathe, and I could only take short gasps of air. I was in so much pain I was crying, and it felt like I was dying. I told Mum and Dad called my Dad in the morning. Dad said to go to the doctors and get checked out and that he'd meet me there (I also called my boss to let her know). The doctor was super concerned as I was visibly not well and my blood pressure was really high.

I had my bloods taken immediately after the consult and they were sent express to Newcastle Lab. After the bloods were taken I burst into tears which is not like me so Dad was really worried. We got Nurofen and electrolytes from the chemist afterwards, then as I was driving home the GP called and told me I needed to go straight to the hospital due to abnormalities in my bloods, he told me my D-dimer level was off the charts (I didn't know what that meant so googled it on the way and found it was the test for blood clots).

My parents were with me the following morning (20th October) when I was seen by the Respiratory Physician. He informed me that I had a pulmonary embolism. Mum asked him to explain what that was, and said it meant I had blood clots. He then said words to the effect of "On the balance of probabilities, I think the clots are due to the contraceptive pill" and told me that most clots develop in the legs before moving to the heart and lungs. He told me it's a lifelong issue and if I was to have kids in future, I'd need to be medicated throughout the entire process to keep the baby and myself safe. He advised me not to do any form of exercise at all - no horse riding, motorbikes, flying, and I was to be on complete rest for 6-12 months as my body had just fought for its life.

I had no words, and felt like I did not want to be alive at that point - What kind of life is it for a 19 year old to be told you can't do anything you love? I asked my parents to leave the room, I wanted to cry and scream by myself (as I'm not one to show emotion in front of people).

They discharged me around 7am that evening saying they couldn't do anything to help me. They provided me a medical certificate for work and stated I was unable to drive for the next month due to my 🍌 injury. We were given no advice. Mum knew something was really wrong so she was asking questions about what could happen. It was explained that I was at high risk of heart attack if a clot was to pass through my heart, or otherwise I could have a clot go to my brain and kill me instantly. Mum and Dad demanded the hospital report my adverse event before I left. I was then left to find my own specialist.

On the 1st November, my parents and I had an appointment with the doctor who administered both my 🍌, as we wanted to ensure he reported my adverse reaction and to seek further advice around my 🍌 injury. He said he'd report it to NSW Health but encouraged me to get another GP as he was retiring in a month. My parents both received confirmations from Central Coast Health about the investigation into my 🍌 injury in the days following my discharge from hospital and we attended an online consultation on the 8th November with someone from the NSW 🍌 safety panel, along with another doctor who informed me that they'd done testing on my blood samples.

I bled heavily for 3 weeks from when I began taking Apixiban - it was so heavy that I was wearing a maternity pad and a super tampon and would need to change them after 20 minutes. Blood would also pour from me in the shower. I'd never seen anything like it before. It then stopped for a week then went through two cycles of being heavy again for 2 weeks and stopping for 2 weeks, then it completely stopped for around 6 months. I've had it return to normal (light flow and 3-4 days in duration) in the last few months.

Now that I am on extreme clotting risk, I can't have any contraception. I stopped taking the blood thinners in May as my clots had completely dissolved. I'd also been going to a naturopath the entire time and was taking vitamins and supplements. I still experience extreme fatigue, and my arteries are permanently scarred. After I shared what happened on my social media last year, my story blew up and I felt like I couldn't go anywhere without people knowing who I was - I'd become a famous sick kid overnight. Even going to the bakery to get bread, I'd be approached by strangers who knew my name and what had happened to me. I developed really bad social anxiety over this time because of it (which I'd never had before) and didn't want to leave the house.

I applied for Work Cover but my work had only processed it a few months ago. We heard back last week that it was declined, so we are appealing it. They state I wasn't mandated, but communications I have from my manager say otherwise - I was stood down on leave without pay and was told not to return to work or enter the premises until I provided proof that I was fully 🍌. I haven't been able to go back to my job as they've wanted me to go back full time (8am-5pm, 6 days per week), and I'm not able to do that. They have since stated last week they can accommodate a gradual return to work to aid the transition to full time. I'd also applied for the Government Compensation scheme and had heard nothing - it's still in review, 6 months on from my application.

The 🍌 was new, still being trialled and long term safety was unknown - my worry was that it would affect fertility. I remember saying to my mum "I was old and had all my kids, I wouldn't be near as hesitant or phased about that". Around the time, a good friend got Myocarditis after their 🍌, then another friend (a surfer) ended up with Pericarditis - I could see that things were happening to people. My Dad is a Police Officer and was hesitant himself. He told me he had written on his consent form that he doesn't consent to taking this 🍌 and doesn't agree with the mandate.

I went ahead and had the first 🍌 on Thursday 30th September 2021 (left arm) to keep my job. After the 🍌 I had a fever, headache, body aches and felt sick, but the doctor had warned this was normal, so I didn't worry too much. On Friday 1st October I received a call from my boss to say I couldn't come back to work until I was fully 🍌. I was put on leave without pay from the 4th October until I could provide my 🍌 passport. I didn't look for another job as the 🍌 was a requirement everywhere so I felt like I had no other choice but to risk getting the second 🍌. So I went and had the second 🍌 on Thursday 21st October (left arm).

I rang my parents crying as I drove myself to Gosford Hospital. I arrived at 1:30pm on the 22nd October. Both my 🍌 parents were told they couldn't come into the hospital due to 🍌 restrictions. They did tests and scans and even was on the phone asking me what they were doing, but I couldn't understand what was happening or comprehend what they were telling me as I was in a lot of pain. I was so emotional and scared that the hospital had to speak on my behalf. I was informed I had blood clots in my lungs, had minimal white blood cells left in my body and that I had an extremely low immune system. They discharged me at 7am with limited information, no medical plan and no discharge summary. I was given blood thinning medication (Apixiban) and advised I could drive home and that I should rest. They told me I did not need to be hospitalised and that I should continue seeing my GP.

The next evening (23rd October) around 6pm, the pain in my chest became stronger. I called my Dad and he arranged for my brother to pick me up and said that he'd meet me at the hospital. Soon after I arrived I was admitted to the respiratory ward and the doctor asked if I wanted to say goodbye to my family and I became quite upset because I took it as I was dying and may not see them again.

They'd taken multiple blood tests, drawing blood from both arms and my veins were bruised and extremely sore. It also felt quite a bit of blood - the side effects from the blood thinning medication I'd started 2 days prior had given me internal bleeding, nose bleeds and my period had come on after starting it and was extremely heavy. I struggled to breathe and was in excruciating pain (which they gave me Endone for). They informed me that with the number of clots on my lungs, it's just like trying to breathe with broken ribs. I was never given oxygen or any kind of medical drip over my time in there.

Just after 12pm I was taken for an ultrasound, which found no signs of blood clots in my legs. The nurse who scanned my legs apologised to me about how I was being treated - I didn't know why she was apologising, but I think she'd seen my notes and read what got said about what happened to me. She said there was no way she'd let her 12 year old daughter get the 🍌 for this reason, and mentioned she only got 🍌 to keep her job.

They ruled out any genetic dispositions that might cause blood clotting, and noted that my D-dimer results were 7 times higher than normal and were alarming. I was referred for an emergency haematologist appointment, which I had on the 11th of November. He advised that 3 of 5 arteries in my heart were totally blocked and stated I would be on blood thinners for 6-12 months as the clots were significantly large and it would take that long for my body to break them down. He also confirmed I didn't have clots anywhere other than my heart and lungs. He'd said my body had gone into attack mode against the 🍌 and that was why I was so sick. He was shocked I wasn't admitted into hospital the first time I went due to how serious the clotting was. He said I should've had a 2 week minimum in intensive care and been given oxygen, as when Pulmonary embolism is not monitored in the initial 2 week period, the fatality rate is around 80%. He noted that I was lucky to be here - all this was new information to me and it scared and shocked me. He explained that when I'm older and want to conceive, I'll need injections in my stomach every 3 months prior to trying, during the entire pregnancy, and 3 months post-partum, and that it's not possible for me to be able to have kids naturally anymore.

AFL lawyers heard my story after it gained a bit of traction. They contacted my mum and felt that my case was strong as it's gone worldwide and had wanted me to lead the class action against the mandate. We agreed, and my mum and dad signed the paperwork that I was to lose, I would not be held accountable for any legal fees. I understood that funding would come from donation or had already been raised.

I did my statement on the 19th December 2021 and provided all my medical records and everything to them. We lost the case, and although the news article states I owe money, that's not true - I don't owe anything. I didn't attend court. I wasn't required to and my lawyers went for me. I didn't really want to hear about it so I didn't pay much attention to it. I was trying to recover and was not in the mindset to listen to anything about it. It dragged on for 10 months - I just wanted to move on. My lawyers said that although there were other stories read out, the judge listened intently to my story and had looked saddened about it. On the 31st August, I woke up to my Instagram blowing up because of articles the media had written about me and the case. I felt angry they'd slandered my name and said things about me that weren't true - it felt like a slap in the face.

### One Year

Well, this week might've been the 1 year anniversary of my death, for my friends and family. The date I had my second 🍌 (the 21st of October), will forever hold a special place in my heart because it became my second chance at life. I will never forget the pain, being too scared to close my eyes at night and knowing I'd had an 80% chance they wouldn't open for me the next morning. From never having a single health issue prior, I'd never thought I'd become a 'sick kid' being wheeled around a hospital. Everything I loved doing, I was told I was no longer allowed to do; Drive my car, ride my horses and motorbike, fly on planes and exercise, even the ability to conceive a child naturally. I took all these things for granted. Although this chapter of hardship changed me, I didn't let it define me. It shaped me into the better person I am today. I will never make a 'full recovery' but I've learnt this is the new me and there are things I have to do differently in my future, and that's okay. If someone tells you that you can't do something, take it, but don't take it onboard - prove them wrong. Life's too short, do what you want, say what you want and tell people you love them. Tomorrow is never promised. How bloody great is it to be alive! Cienna - October 2022.

## My Take...

These situations are usually lethal.

According to a Dec.30, 2022 [USA Today Fact check](#), there is NO LINK between Pfizer COVID-19 mRNA Vaccines and blood clots.

FDA spokesperson Abigail Capobianco also told USA Today Fact Checkers that there is no link.

The COVID Cartel will simply NOT ALLOW a link to be established between Pfizer COVID-19 mRNA Vaccines and blood clots, nor will it allow any research papers to be published. [South Korean researchers](#) tried, and they were forced to return with the “[correct answer](#)” a few months later.

That’s why the Australian government is denying all vaccine injury claims involving blood clots and mRNA vaccines.

Eliquis is a blood thinner developed by Bristol-Myers Squibb and Pfizer to treat blood clots.

- Eliquis sales in [2019](#) were \$12.1 billion
- Eliquis sales in [2020](#) were \$14.1 billion
- Eliquis sales in [2021](#) were \$16.7 billion
- Eliquis sales in [2022](#) were \$18.3 billion

AstraZeneca was taken off the market in March 2021 and very few J&J were given out. Both were taken off the market for causing “rare” blood clots.

That raises an interesting question.

What is driving the rise in Eliquis sales to the point where it is the 6th best selling drug in the world? Climate change? Long COVID?



## The 50 best-selling pharmaceuticals of 2022: COVID-19 vaccines poised to take a step back

By Brian Buntz | April 18, 2023

Drug name	Manufacturer(s)	2022 Sales	Indication(s)
<b>1. Comirnaty COVID-19 vaccine</b>	Pfizer/BioNTech	\$55,918,791,640	The COVID-19 vaccine topped the list of 2022's 50 best-selling pharmaceuticals. It sold slightly more last year.
<b>2. Humira (adalimumab)</b>	AbbVie	\$21,237,000,000	rheumatoid and psoriatic arthritis, ankylosing spondylitis, Crohn's disease, ulcerative colitis
<b>3. Keytruda (pembrolizumab)</b>	Merck	\$20,937,000,000	various cancers
<b>4. Paxlovid</b>	Pfizer	\$18,933,000,000	prevention of severe COVID-19
<b>5. Spikevax COVID-19 vaccine</b>	Moderna	\$18,435,000,000	reduce risk of COVID-19 infection
<b>6. Eliquis (apixaban)</b>	Bristol Myers Squibb and Pfizer	\$18,269,000,000	blood clots
<b>7. Eylea (afibercept)</b>	Regeneron Pharmaceuticals, Bayer	\$12,721,221,200	age-related macular degeneration, macular edema and diabetic retinopathy
<b>8. Biktarvy (bictegravir, emtricitabine, and tenofovir alafenamide)</b>	Gilead Sciences	\$10,390,000,000	HIV
<b>9. Revlimid (lenalidomide)</b>	Bristol Myers Squibb (Celgene)	\$9,978,000,000	myelodysplastic syndrome, multiple myeloma, and mantle cell lymphoma
<b>10. Stelara (ustekinumab)</b>	Janssen (Johnson & Johnson)	\$9,723,000,000	plaque psoriasis, psoriatic arthritis, ulcerative colitis and Crohn's disease

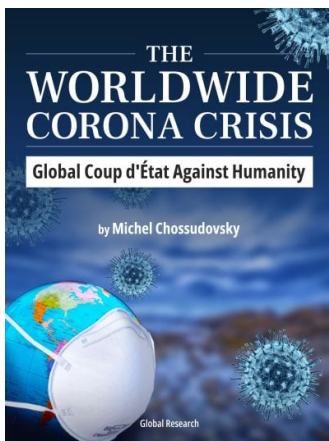
I believe that Pfizer and Moderna COVID-19 mRNA Vaccines cause blood clots and just like what happened with myocarditis, the true risks of clotting from these mRNA vaccines have been greatly downplayed and suppressed by the authorities.

On [October 3, 2023](#), Biden's White House announced that the makers of Eliquis will participate in Biden's "Inflation Reduction Act" program to "lower prescription drug costs for Americans.

\*

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*Dr. William Makis is a Canadian physician with expertise in Radiology, Oncology and Immunology. Governor General's Medal, University of Toronto Scholar. Author of 100+ peer-reviewed medical publications.*



## The Worldwide Corona Crisis, Global Coup d'Etat Against Humanity

by Michel Chossudovsky

Michel Chossudovsky reviews in detail how this insidious project “destroys people’s lives”. He provides a comprehensive analysis of everything you need to know about the “pandemic” — from the medical dimensions to the economic and social repercussions, political underpinnings, and mental and psychological impacts.

*“My objective as an author is to inform people worldwide and refute the official narrative which has been used as a justification to destabilize the economic and social fabric of entire countries, followed by the imposition of the “deadly” COVID-19 “vaccine”. This crisis affects humanity in its entirety: almost 8 billion people. We stand in solidarity with our fellow human beings and our children worldwide. Truth is a powerful instrument.”*

### Reviews

*This is an in-depth resource of great interest if it is the wider perspective you are motivated to understand a little better, the author is very knowledgeable about geopolitics and this comes out in the way Covid is contextualized. —Dr. Mike Yeadon*

*In this war against humanity in which we find ourselves, in this singular, irregular and massive assault against liberty and the goodness of people, Chossudovsky’s book is a rock upon which to sustain our fight. –Dr. Emanuel Garcia*

*In fifteen concise science-based chapters, Michel traces the false covid pandemic, explaining how a PCR test, producing up to 97% proven false positives, combined with a relentless 24/7 fear campaign, was able to create a worldwide panic-laden “plandemic”; that this plandemic would never have been possible without the infamous DNA-modifying Polymerase Chain Reaction test – which to this day is being pushed on a majority of innocent people who have no clue. His conclusions are evidenced by renown scientists. —Peter Koenig*

*Professor Chossudovsky exposes the truth that “there is no causal relationship between the virus and economic variables.” In other words, it was not COVID-19 but, rather, the deliberate implementation of the illogical, scientifically baseless lockdowns that caused the shutdown of the global economy. –David Skripac*

*A reading of Chossudovsky’s book provides a comprehensive lesson in how there is a global*

*coup d'état under way called "The Great Reset" that if not resisted and defeated by freedom loving people everywhere will result in a dystopian future not yet imagined. Pass on this free gift from Professor Chossudovsky before it's too late. You will not find so much valuable information and analysis in one place. -Edward Curtin*

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