

Most Painful Thing In The World? CRPS! The Neuroscience Center Relieves Pain w Combination Therapy

T

The Neuroscience Center

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Most Painful Thing In The World? Complex Regional Pain Syndrome! The Neuroscience Center relieves pain with cutting-edge technology! Dr. Steve Best has invented, and patented, a Combination Therapy, which includes Transcranial Magnetic Stimulation with Simultaneous IV Ketamine. The results are remarkable!
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D

Commenting publicly as Duncan MacKinnon

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- 

Gracie Bagosy

1 year ago

I am not a patient at this clinic any longer.


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Kristin Shadrick

1 month ago

Would you be so kind to talk to me? Kristinshadrick@gmail.com either of you?

Reply •
- 

Gracie Gean Chronic Pain Advocacy & Consulting

1 month ago

Gracie Bagosy Feel free to email me
www.GGPainAdvocacy.com

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D

G

GARTHBOOTS 9 months ago

I have fullbody rsd and im in hell !!

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E

Ern 1 month ago

God Bless. You have my prayers.

Reply • 1

G

GARTHBOOTS 1 month ago

Thankyou Ern Im sorry you have this horrific disease.I try to stay positive its hard at times but Im helping myself as much as i can.GOD BLESS TO YOU ERN AND OTHERS WHO SUFFER!!

Reply •

d

dlny911 1 year ago

Just called after suffering over 30 years with RSD/CRPS. They take no insurances whatsoever no matter what- so cash only which means if you are alone and on disability with Medicaid and Medicare you are fucked. They DO NOT CARE UNLESS you have MONEY. I just called- nope - they do not care. They want to end your suffering ONLY if

Reply • 1

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E

Ern 2 months ago

my prayers Angela. tip: don't use your full name.

Reply •

**Wes Northcutt** 2 weeks ago

you are 100% right...all that matters in medicine \$\$\$

Reply •

**Phoenix Chastaine** 1 month ago

Dr. Kirkpatrick understands CRPS like no other. He comes right and says there are still way, way to many of these poor souls being totutred by doctors who are ignorant of how horrific the daily life experience of theses patientst are. Too often, they barely even assessed physically, they don't believe the patients details of the excruciating and how

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**Phoenix Chastaine** 1 month ago

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Reply •

L

Lisa Jacobs 2 months ago

I have had C.R.P.S. THAT STARTED WHEN I CALLED FOR A LOAN TO PUT AN ADDITION ON MY HOUSE. I HAVE A SISTER THAT IS 16 YEARS OLDER THEN ME. SHE HAS DRAINED MY PARENTS BANK ACCOUNTS 6 TIMES! she is not a drug addict! she is a sociopath!! When I called to get the loan I found out she rold my mother who was 82 at the time that she was just signing a paper in my name to help me! once again my mother believed her! she knows I won't put my mom in jail so thats why she did it! my sister. only sister) stole every dime I had 199,000 dollors. I was at work. so my mind was not on my work and I fell. I got right back up. but I had a nervous breakdown and fell to the floor. She is not my sister to me because a sister wouldn't do that! my mom and I had nothing to eat for 2 weeks. We were so hungry i called her and asked her to buy us some food? sne came over 2 weeks later and gave us left iver spaghetti. I believe that I got this from my nervous system shock when I fell. I truly believe if my nerves didn't go into shock I wouldn't of got C.R.P.S.!!! my mom is 91 and has to work . god bless her she is a strong woman!! my sister made up lies to my family and told them I was a drug addict and stayed in bed all day and ate pills. the truth is I did stay in bed all day and still do because of the pain!! I want nothing to do woth her! because she gave me C.R.P.S. IT WAS MY NERVES NOT THE FALL!! my nerves had to go somewhere! so now we are in our own. I dont have any money to take care of my mom. but my SISTER couldn't care

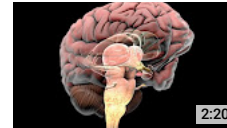


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14:48



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Complex Regional

nothing so I can go there!! my point is! I think I got it because of what she has done to my nerves!!

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Reply •

E

Ern

1 month ago

It's possible you are. Emotional provocation or stress will make it worse. The best thing I use is Lorazepam. Try methadone and Fentanyl if you can get it. The methadone is a proven narcotic for RSD, works better than others and is very cheap and easier to get because the doctors are sure that you'r not chasing a

[Read more](#)

Reply •


L

LadyBuggy05

2 months ago

I just got diagnosed with it. I'm 11. I've also been diagnosed with Lupus, Rheumatic Fever, and Rheumatoid Arthritis since I was 4.

Reply •




Tony Beer

2 months ago

PLEASE READ THE POST BELOW THIS ONE IS VERY INTERESTING THANK YOU

Reply •



Tony Beer

2 months ago

I shared this about a year ago and will share it again now so people can understand. Warning it's a long post ♥

Hi my name is Complex Regional Pain Syndrome... some friends who have known me for a very long time call me RSD or CRPS..... I'm an invisible inflammatory disease that attacks your sympathetic nervous system.

I am now velcroed to you for life. If you have CRPS you hope for remission but there is no cure.

I'm so sneaky--I don't show up in your blood work, in x-rays, MRI's can't detect me, basically there is no test to prove you have me. There are only test to rule out other things.

Others around you can't see me or hear me, but YOUR body feels me.

I can attack you anywhere and anyway I please. And, I will. Constantly.

I can cause severe pain or, if I'm in a good mood, I can just cause you to ache all over. Remember when you and energy ran around together and had fun?

I took energy from you, and gave you exhaustion. Try to have fun now.

I can take good sleep from you and in its place, give you brain fog and lack of concentration.

I can make you want to sleep 24/7, and I can also cause insomnia.

I can make you tremble internally or make you feel cold or hot when everyone else feels normal.

I can cause one limb to change color, look bruised, feel super sensitive randomly for seemingly no reason.

I can also give you swollen hands and feet, swollen face and eyelids, swollen everything. OH, and just because I started off in one part of your body, don't think I can't travel and effect other limbs or any other part of your body I so choose to torment. I can, and likely I will.

I can make you feel very anxious with panic attacks or very depressed. I can also cause other mental health problems. You know crazy mood swings? That's me. Crying for no reason? Angry for no reason? That's probably me too. It is hard to not feel hopeless when you have me beating your body up constantly.

I can make you scream out loud, anytime of day or night, anywhere you are because I can create pain that makes you sure someone just stabbed you with a knife. Making you look crazy is fun for me.

I can make your hair fall out, your nails become dry and brittle, cause acne, cause dry skin, the sky's the limit with me.


I can make you gain weight and no matter what you eat or how much you exercise, I can keep that weight on you. I can also make you lose weight. I don't discriminate.


Some of my other inflammatory disease friends often join me, giving you even more to deal with.

If you have something planned, or are looking forward to a great day, I can take that away from you. You didn't ask for me. I chose you for various reasons: That sports injury, needle stick,, or that car accident, or maybe it was the surgery to correct a bone problem.. Whatever the cause, I'm here to stay.

I hear you're going to see a doctor to try and get rid of me. That makes me laugh. Just try. You will have to go to many, many doctors until you find one who can even try to help you effectively. Most of them will make you feel like you are to blame, or worse, it is all in your head. I'll convince them that you are crazy because normal people know that you can't have all those symptoms all over your body and still walk around looking normal.








CanadaQbank

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living with complex regional pain syndrome (CRPS)

Kirsten Evans

41,206 views

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There are so many other ways I can make you sick and miserable, the list is endless - if your body is all of a sudden dealing with things that were never issues before...yep.... that's probably me.
[Read more](#)

Reply •



captain0ldy0da 1 month ago

My life written out here

Reply •



Tony Beer 1 month ago

sorry to hear that my friend CRPS is pure evil

Reply •

E

Ern 2 months ago (edited)

Please recognize that this is a sales pitch with one pure objective - to sell the idea that something can be done. But for how much and where? Will your insurance cover it? I haven't left the house without an ambulance for two years and before that I was laying down in a van. This sounds like one clinic and my doctors aren't telling me about it and
[Read more](#)

Reply •

h

hotlonnie78 5 months ago

<http://www.gofundme.com/help-lonnie-fight-like-a-girl>

Reply •



Another Awkward Cosplayer 9 months ago

I have CRPS in my legs, although I don't feel boring anymore the pain is still there, and very bad. The worst pain overall though, is the fact that my family doesn't believe me.

Reply •



A.V. 6 months ago

Another Awkward Cosplayer antonisanto

Reply •



Ern 2 months ago

I understand Cosplayer. I have had it for 17 years and have lived with the same circumstance. Fortunately my wife has struggled with understanding and has been an immense help. Most of my family and friends are gone. But for my wife I would be very lonely. I send my prayers.

Reply •



Kara Alvarez 10 months ago

I didn't go to that clinic, but I had CRPS for about 5 years. It had been completely gone for 12 years now 😊😊

Reply •



Rollin Shultz 11 months ago

The way they describe the pain levels, it sounds as bad as Diabetic neuropathy. I have been taking tramadol for three years for my nerve pain and it works great on only nerve pain, no other pains such as muscular. Unlike what I have heard I get NO withdrawal symptoms from it. In fact if I don't use a countdown timer to regulate my intake, I forget to take my pill until the pain sneaks up on me and then I must wait an additional hour or two for the new pill to take effect resulting in approx. three hours of needless pain. My neuropathy is in my entire body and the tramadol eliminates the pain in all but my feet and toes, which are reduced to about a level 2.

I received NO offers of any cure or even treatment to improve my condition from the medical establishment, especially my neurologist who is supposed to be an expert, although they were glad to take my money for countless tests and visits, while offering no benefit for the money they received. I almost died as my condition worsened and begged God many times to kill me and set me free.

I began to see improvement after some months of being on a neuropathy supplement formula (do your own search) that contains B1, B12, and the most important (alleged nerve re-generator) R-Alpha Lipoic. I took it on faith based on so many testimonials they could not have been faked. I examined the testimonials for patterns and repetition and I found them to be similar but personally influenced, and no one is that creative to make so many similar but different accounts. There is always some kind of pattern to faked testimonials, it goes with the territory.

regeneration, so again dig deep do your own research and make your own decision. And always avoid cheap low quality supplements and pay extra for high quality products, you will thank [Read more](#)

Reply •

C

[Celtic Contemporary](#) 11 months ago

I have CRPS in both of my hands

Reply •



[Tony Beer](#) 1 year ago

the pain is very real .i have had CRPS for about 3 years but after 3 spinal operation and ALIF that's a fusion i have had intractable pain for almost 20 years due to spinal pain .its hell .don't let anyone tell you its all in your head ..ITS REAL

Reply •

E

[Ern](#) 2 months ago

Of course it is Tony. It's real for all of us. Don't waste much energy trying to convince anyone. Most people - even compassionate people, don't understand. Even if they do understand pain they don't understand relentless pain and living in the loss of your life. They have done brain scans of people with their foot in a vise

Reply • 1

K

[Kade Larson](#) 1 year ago

I have CRPS in my left foot for just over 7 years now, I haven't had any thing that has truly helped.

Reply •

A.

[A. Peerson](#) 2 years ago

Ketamine, via NMDA antagonism, is an effective antidepressant.

Reply •

E

[Ern](#) 2 months ago

Ketaminie is a horse tranquilizer. It is very dangerous to work with. Should be administered at a hospital.

Reply •



[Angela Siegfried](#) 1 month ago

It is not very dangerous to work with. On the contrary the only reason it isn't being used more extensively and or researched more is because Katimine is very inexpensive and there's not enough money in it for pharmaceuticals to want to research it further. It has been used in both humans and pets as an anesthesia.

Reply •



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