Graeme Parker of Brampton, Huntingdon residing in motorhome reg J243 ENN Tel: 07783 631527

Mr John Woods 1 Little End Rd Eaton Socon Saint Neots PE19 8JH

21st July 2023

Dear John Woods

I am writing to you from Newquay Cornwall where life as a Surfer revolves around the ocean swells coming and going. My family residing in Cambridgeshire since 1973 following the posting of my father (deceased 2016) to RAF Wyton and then Brampton. Currently aged 53 last month I was in Huntingdon to attend Priory Fields Surgery for my annual medical and learnt of your health condition from the 21st June edition of the Hunts Post newspaper. Journalist Debbie Davis informing that funds had recently been raised for the Arthur Rank Hospice Charity and the Motor Neuron Disease Foundation by cycling around Grafham Water Reservoir.

My mother (Wendy Mary Parker) died in Hinchingbrooke Hospital Huntingdon from a paralysis disease in November 2002, I having been living in Brazil since 1999 employed as a Design Engineer in their aerospace industry. My mother's illness commenced in September 2000 with her left foot and ankle becoming lame only nine months into her retirement following 20 years employment within Hinchingbrooke Hospital. My mother's condition spread up her left leg and by the spring of 2002 she had lost use of both her legs. Consultant neurologists including those practising in London over this almost two year period were unable to provide a diagnosis fear being that it was Motor Neuron Disease, however my mother's illness did not show 'classic symptoms' of losing the ability to smile. In the spring of 2002 Wendy Mary Parker received an experimental treatment involving transfer of blood plasma, following this her condition worsened, the paralysis accelerating up through the torso eventually causing her death. Consultant neurologist only confirming Motor Neuron Disease after having administered experimental treatment.

During August 2001 via the internet in Brazil I surfed across the American Mr Donald Byrn who had spent many years battling against his diagnosis of ALS (Amyotrophic Lateral Sclerosis the American term for MND). Travelling to his home in Phoenix Arizona I met Donald and two of his assistants that were Mormons running his cause named 'New Hope Appeals'. Donald offering treatment to restore the central nervous system derived from peanut shell extract, the active substance being **Luteolin** which can also be found in celery sticks. Questioning Donald face to face he told me via an assistant, as he was surviving via a mechanical respirator, that he believed his condition had been caused via drinking from a hosepipe on farmland.

Our Brampton family home garden bordering a cereal crop field via a drainage ditch, my mother's death affected me very much. Having lost engineering employment due to the 911 felling of New York's twin towers, aged 32 I committed myself to childhood dream of surfing and based myself in Ilha Santa Catarina off the south coast of Brazil. Returning to this location after a successful surf trip to South Africa August - September 2004 I had been befriended and encouraged by local professionals and continued to train fulltime. However, life took an alternate path in October 2004 following the rupture of my left side vas deferens and semen entering blood stream, 15 months later I began to experience onset of a paralysis disease that commenced in the outer two fingers of my left hand.

As the disease spread up my arm I observed 'growth' occurred during sleep, as life became a Nightmare on Elm Street fighting off my arm going lame I practised guitar, press-ups and surfing. Trying not to go to sleep

I would experience tremor of my left wrist as the 'infection' would create nodal spores up my arm that would then connect and similar to a flicking of an electrical switch the limb would be paralysed and I would shake it back to life. I learnt that the infection was linked to my metabolism and noticed coffee (caffeine) would accelerate growth, the toxins of smoking a cigarette would quell growth and occasional smoking of marijuana would lower my metabolism and encourage infection's growth.

Paralysis attacks culminated 12 months after first attacks to left hand outer fingers with the infection synapsing across the nape of my neck and highly efficiently leaving me with both arms and hands paralysed, shaking them both back to life - days followed of trying to not go to sleep by doing press-ups until I dropped.

This was February 2007, housed by a hotel owning family that had been witness to my activity since infection onset my father was invited to Brazil and following him back to his hotel in downtown Florianopolis capital city of Santa Catarina, I stopped at a market stall selling medicinal tea managed by a tall well-built Indian gentleman lacking sight in one eye. Explaining my condition he prescribed a traditional remedy for parasites named 'Flor da Amazonia' below is a scan of a packet providing this medicinal compound.



Translating the paragraph 'Indicacao' = 'Recommendation' provides 'Sistema Nervoso' = Central Nervous System and 'Vermes em Geral' = Worms in General.

Since returning to the UK in March 2012 (I was deported by Brazilian Federal Police) I have been regularly taking medicinal tea 'Ipe Roxo' principal ingredient of 'Flor da Amazonia' which is sold in Europe under alternate name 'Pau D'Arco', it's the inner bark of a tree native to South America. Living in a motorhome the morning chill causes the outer two fingers of both hands to be slightly lame and wiggled back to life and if I'm lazy or sunbathing exposing my back to the Sun I experience an involuntary spasm to my lower back where I believe a spore of 'Neurol Worms' (worms the size of a neuron) are slowly attacking my spinal column.

My attempts to present my experiences and discoveries to the NHS have so far been unsuccessful and hope you don't mind that this presentation to you John Woods of St Neots in the Ouse Valley be a part of my next presentation.

In November 2016 I attended a class of 1986 Hinchingbrooke School reunion and learnt that two of my class mates and residents of Brampton had recently lost their mother's to a paralysis disease diagnosed as MND and attending Jehovah Witness reunion in Huntingdon I learnt that the grandmother of another classmate had died from MND.

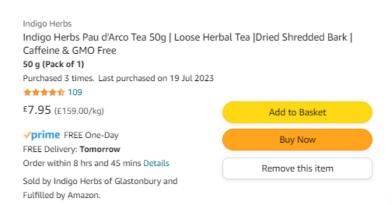
Fully understanding the 'hope' that terminally ill patients cling too I could not but write this letter to you and provide you with a packet of the same medicinal tea I am taking.

Pau D'Arco and Luteolin available from Amazon

Also recommend Mebendazole - Threadworm tablets available from Boots the Chemist.

Avoid Caffeine.







Capsules 500mg, Maxim...

Deal

-20% £14.90 RRP: £18.62
(£0.25/count)

✓prime FREE One-Day Get it Tomorrow, Jul 22

Add to Basket

As a screenwriting storyteller I have incorporated my mother's tale into story 'The Golden Dolphin and the Wormwood' which can be viewed on YouTube channel Apollobath TV link:https://www.youtube.com/watch?v=2LFtovygmtk&t=1s

Am also providing you with 'Chapter 5 – Paralysis Disease The Wormwood' of a document presented to Cambridgshire Mental Health and invite your study of chapter 11 of dossier 'Camaialot A' available at website www.apollobath.org link 'Crown Office and Embassy'.

Best Wishes

fundraiser Lee 'Celebration of life' for

Tributes have been paid to St Neots man Lee Taylor after a celebration of life event was held on Friday, June 16.

May just days after cheques were Neurone Disease (MND), died in presented at his final fundraiser for the Arthur Rank Hospice Lee, aged 46 who had Motor Charity.

The event involved friends and children Mya-Mai and Hudson cycling around Grafham Water. family - including his young

Together, with their family and ball at The Priory Centre, in St and the Motor Neurone Disease raised £20,800 which has now Arthur Rank Hospice Charity friends, they hosted a charity been donated equally to the Neots, in March. The event (MND) Association.

MND in 2019, he set himself a When Lee, of St Neots, was diagnosed with bulbar onset

DEBBIE DAVIES

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and campaigned on social media series of fundraising challenges to raise awareness of "living During his illness, Lee positively"

benefited from care and charities and from the Addenbrooke's support from both nosted at the MND Clinic hospice.

that he was determined to leave are comforted by the fact heartbroken but we most amazing father. We are "Lee was the husband and Lee's wife Tracey said:

Huntingdon, St Neots and across raised helps us provide holistic their support. The money they "We are grateful to Lee and John and their families for all care for more people in condition. "It was a privilege to Sharon Allen, said: positive attitude to living engage with Lee. His chief executive,

Windows, said: "When I was first Mr Woods, owner of St Neots Cambridgeshire."

well for as long as possible will

diagnosed with a life-limiting continue to help other people

achieve so much to help others." us with so many memories and

diagnosed it was Lee's positivity thinking about what the future holds for someone with MND. that helped me cope with

bond. I miss him terribly, but his legacy will live on forever just as "After a while we supported each other and had a special he wanted it to. He was a legend."

