

GcMAF has worked for my family

THESE have been several letters and articles about Immuno Biotech's GcMAF in the Guernsey Press this year. I would like to offer some personal experience that your readers might find useful in making their own minds up about the efficacy of GcMAF.

I have a daughter with a rare genetic disorder, LMBBS – Laurence Moon Bardet Biedl Syndrome – which means, among other things, that her immune system does not work at full strength. As a result, she is often ill with colds, stomach bugs, etc. and always has low energy, sometimes sleeping 12 hours a day, so rather like CFS – Chronic Fatigue Syndrome. She also has had acne, cysts and dyslexia. The first piece on GcMAF in the Press a few months ago prompted me to look into it further as, in a nutshell, it restores the body's natural immune

system to full strength. Her symptoms overlap with those of autism and chronic fatigue for which GcMAF has documented success in relieving. Possibly because if you are continually fighting infections, you have low energy levels and little energy left over for other things or because they are actually caused by

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LMBBS sufferer who has been taking GcMAF

infections. My teenage daughter is currently in the UK doing studies and there was no problem in having GcMAF shipped to her there. She has been having weekly injections for just over three months and the results are remarkable. Acne has virtually gone. A cyst that would normally

take a course of antibiotics to get rid of has gone – she has been on and off antibiotics for cysts for the past five years and not needed antibiotics since starting GcMAF. More importantly, she is bounding around full of energy, talking nineteen to the dozen and says she is able to think so much more clearly. In her words, 'It has

changed my life. I feel and think like a normal person – no longer wading through treacle and fog.' And now to the key thing. The worst part of this genetic disorder is that the retina of the eye slowly dies so that sufferers usually become blind by their late teens – a form of macular degeneration. An

ophthalmologist recently checked her eyes and said 'she has surprisingly good visual function'. It seems that GcMAF is keeping the expected retinal decline at bay. You cannot imagine what that feels like for her to learn that. On the strength of all these positive tangible results, my teenage son, who has the same genetic disorder, started on GcMAF. Again, a transformation in his energy levels has been dramatic, resulting in improvements in behaviour and attitude. He is so much happier as he feels normal. And what about his eyes? A few days ago, his retinas were checked. Amazingly, they are actually improving. According to the experts in this disorder, that cannot happen. Yet I have the pictures of the retinas to prove that it can and the only possible cause is GcMAF. The exact mechanism at work is unclear mainly because the cause of the retinal

degeneration in this genetic condition is not known in the first place but the evidence-based results that it works is indisputable. The best evidence for something working is the evidence of one's own experience. GcMAF lived up to its claim of restoring the immune system to work at full strength for my children. In case you are wondering, I tried GcMAF on myself first to check for side-effects. No problems and it cleared up a low-level infection that had been sapping my energy levels for several days and swelling my lymph glands. Fortunately, I am not in a position to comment on its effects on cancer from first-hand experience, but, based on its positive effect on my children's immune systems, I would urge anyone with cancer to read the research papers on GcMAF. Similarly for autism and chronic fatigue syndrome on the basis that GcMAF-treatable infection could well be the cause in both cases.

Name and address withheld.

Lingering questions over Guernsey Water