

Miraculous drug for MS

Sir, — I am one of many multiple sclerosis sufferers in this country but I am luckier than some.

Shortly after my diagnosis, I found the drug LDN, which helps all autoimmune diseases by regulating the autoimmune system and stopping it from attacking the body's own cells.

I found this drug before my disability became unbearable, before I was in a wheelchair and before I was a burden to my family and needed round-the-clock care that many people with autoimmune diseases require. Such people are often in pain, often depressed and helpless.

Finding LDN was amazing and it opened up a world I never knew existed.

It made me realise that a cheap, safe little drug is not available on the NHS, which should be delighted to help patients and save money.

It made me aware that patients often know a lot more about their diseases than doctors do, which is fairly understandable, but I had just never thought about it.

It made me realise that most doctors haven't even heard of LDN until a patient tells them about it.

I am proud that my article became front-page material for our village paper and that my story is included in the e-books people have been writing to make fellow sufferers aware of this drug, which can improve and even save lives.

There is 25 years of patients' evidence to prove that this drug works, that it is the best thing since Penicillin and until there is a cure it's the best thing indeed.

But people don't know about it, they aren't told about it, unless they find out for themselves on the internet or because they might read this letter.

Last week was the first international LDN awareness week. Users worldwide are working at getting the word out about LDN. Here in Britain, we are running a petition calling on the Government to look into LDN, so it can be made available on the NHS.
— Yours faithfully,

Silvia Lane

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