

# Pills For Every Ill

September 14, 2015 By [Nicole Lemelle](#)

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“Medicine is a science of uncertainty and an art of probability.” ~ William Osler

For the past few weeks, I have been feeling run down. I even had moments when my left hand stopped working. It just didn't respond to anything I wanted it to do. I thought the problem would pass but it never did. So I finally made an appointment with my neurologist. I told him about the difficulty I was having with my fine motor skills and how I was more tired than usual. He gave me a routine examination. Then he prescribed the standard MS go-to treatment of steroids.

Being a veteran at receiving Solu-Medrol, I opted for the in-home 5 day IV infusion. That way I could stay in the comforts of my living room instead of traveling to a clinic or hospital.

When the nurse arrived at my home, he had a pile of papers for me to sign. The last thing he needed to know, before starting the treatment, was all of the medications I was taking. As I read from my drug list I keep on my phone, I noticed he ran out of room on the form he was using to document my answers. Then he said, “That's never happened before. I've never run out of room”.

So I asked him, “Do you think I take a large amount of medicine?”

He very diplomatically said, “If that's what you need then it's not a lot.”

That encounter got me thinking. Do I take a lot of prescription drugs? I know I ingest a whole heap of pills. But is it more than others with MS? Or am I in harmony with the norm? It just seems I take a pill for every ill.

Below is my list of medications that help me manage my MS and it's symptoms.

Rituxan® - MS Treatment

Clonazepam® - 2mg

Mybetriq® - 50mg

Baclofen® - 20mg

Nudexta® - 20mg

Alendronate® - 70mg

Reglan® - 10mg

Effexor® - 75mg

Trazodone - 50 mg

Ultram® - 50mg

Ritalin® - 20mg

Cambigan® - Eye Drops

(This list excludes my vitamins)

So I want to know, do you come close to the amount of medicine I consume? Do you take more? Do you take less? Or do you think I should not compare myself to others because everyone's MS is different?

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