



Pamela

Homocystinuria | Nevada | Annual costs: \$3600

It would mean so much to have the medical nutrition equity act passed. I have a diagnosis of homocystinuria which was discovered very late in life. I have suffered from strokes and blood clots in my lungs because of it. Although I currently have coverage through our insurance, it is always a fight to get them to pay. It requires many frustrating hours of appeals and mountains of documentation from my doctors. When I turn 65 in a few years and am eligible for Medicare, I will lose coverage for medical nutrition. I will then be forced to pay \$800 - \$1000 a month to receive these necessities to stay healthy. This is of great concern to me. Please pass this legislation. I am a person, not just a diagnosis to be forgotten!

Pamela depends on medical nutrition to stay healthy. Pamela and thousands of others across the country are depending on you to co-sponsor the Medical Nutrition Equity Act. Please contact the offices of Sen. Casey, Rep. McGovern or Rep. Herrera Beutler.