Due to the burdens of participating in a clinical registry, your provider is not participating in one and neither are any other local providers. You are unable to participate in a patient registry due to the burdens of consistently reporting clinical information.

CMS will only cover the treatment if you are participating in a prospective clinical study – like a clinical registry – which could be a provider-based clinical data registry, a patient registry or something else CMS will require.

Unfortunately, there aren’t any in your community, and you don’t have the time or flexibility to attend study visits required as part of the trial.

Because you have a health condition or a co-morbidity listed in the exclusion criteria, you are not eligible to participate in the trial.

If it is a blinded study design, you may receive a placebo instead of the medicine.

While biopharmaceutical companies have made tremendous progress in scientific and medical innovations for neurodegenerative diseases like Alzheimer’s over the past decade, efforts are in vain if patients are prevented from accessing these life-altering treatments.