

Study: Medical marijuana associated with fewer hospitalizations for individuals with sickle cell disease

13 August 2020

Individuals with sickle cell disease (SCD) who receive medical marijuana to treat pain may require fewer visits to the hospital, according to a new study in *Blood Advances*. Adults with SCD who requested and obtained medical marijuana were admitted to the hospital less frequently than those who did not obtain it.

SCD is the most common inherited red blood cell disorder in the United States, affecting an estimated 100,000 people. According to the Centers for Disease Control and Prevention, SCD affects one out of every 365 Black or African American births and one out of every 16,300 Hispanic American births. SCD is characterized by abnormal, sickle-shaped [red blood cells](#) that can adhere to and block blood vessels, preventing oxygen from reaching the tissues. When this occurs, individuals living with SCD experience severe pain events which may drive them to seek emergency care. It is estimated that there are more than 100,000 SCD-related hospital stays in the United States each year.

There is a need for other options for management of pain so that individuals with SCD do not have to go through the time, hardship, and expense of hospitalization and can manage their symptoms at home. Previous studies have shown that cannabis and cannabinoid products can effectively treat chronic pain associated other conditions such as cancer. More controlled studies of marijuana for the treatment of pain in SCD are still needed.

"When we offered [medical marijuana](#) as an option to our patients with sickle cell disease, we found the majority of people who were interested were already using illicit marijuana to treat pain," said the study's lead author Susanna Curtis, MD, of the Yale School of Medicine and Yale Cancer Center. "Illicit marijuana is not regulated, so its quality and

contents are not standardized. And particularly for people with [sickle cell disease](#), many of whom identify as Black, we know that while Black and white people use marijuana at similar rates, Black people are four times more likely to be arrested for possession. We didn't want our patients using unsafe products or being arrested for trying to control the pain of their condition."

Dr. Curtis and her team examined data from 50 individuals with SCD at the Adult Sickle Cell Program at Yale New Haven Hospital who were certified for medical marijuana use. Of those certified, 29 obtained medical marijuana and 21 did not. Those who obtained medical marijuana visited the hospital less frequently on average over the following six months. Several patients even reduced their hospital admission rates by three, four, or five visits. Receiving medical marijuana was not associated with a change in emergency room or infusion center visits, total health care utilization, or opioid use. Researchers did not observe any change to hospitalization rates in individuals who did not obtain medical marijuana.

Dr. Curtis suggested a possible reason for the reduced hospitalization rates could be that medical marijuana helps individuals better tolerate their pain at home.

The researchers also sought to understand why many patients were unable to access medical marijuana, and why some continued to occasionally use illicit cannabis despite obtaining medical marijuana. In a survey, individuals who obtained medical marijuana reported that they felt it was safer than illicit marijuana and they felt it was effective at controlling their pain; however, they did also report barriers such as greater expense and difficulty of access.

Race and socioeconomic status may also be barriers for patients with SCD. "About 80% of our clinic population identifies as Black, and another 15% as Latinx, and unfortunately people of color who visit the hospital with pain are often not believed or accused of being drug-seeking. Medical marijuana is associated with significant stigma, and stigma is already a big part of the life of a person with sickle cell," said Dr. Curtis.

Cannabis and cannabinoid products also present legal and medical challenges. While 33 states have medical marijuana laws, only five of those states list SCD as a qualifying condition. Medical marijuana products vary greatly in their chemical content and forms of administration. In the study, patients who obtained medical [marijuana](#) were more likely to use edible products as opposed to inhaled products. Previous research suggests the pain-relieving effect of edible products has a slower onset but a longer duration than that of inhaled products. This variability between different products can cause difficulty for patients trying to manage their symptoms.

Dr. Curtis highlighted the need for further research to understand the efficacy, side effects, and drug interactions of various cannabis products on SCD treatment. "My patients are living with a very difficult disease that causes them a lot of [pain](#). We need controlled trials to look at each product, and the effects of how it is taken, so that we can offer regulated, pharmaceutical-grade treatment options."

Provided by American Society of Hematology

APA citation: Study: Medical marijuana associated with fewer hospitalizations for individuals with sickle cell disease (2020, August 13) retrieved 11 October 2022 from <https://medicalxpress.com/news/2020-08-medical-marijuana-hospitalizations-individuals-sickle.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.