

Table 1. Overview of diseases for which CBD may have therapeutic benefits taken from Pisanti et al (2017) [69]

Disease	Effects
<b>Alzheimer's Disease</b>	Anti-inflammatory antioxidant, antiapoptotic <i>in vitro</i> and <i>in vivo</i> AB-evoked neuroinflammatory and neurodegenerative responses.
<b>Parkinson's Disease</b>	Attenuation of the dopaminergic impairment <i>in vivo</i> ; neuroprotection; improvement of psychiatric rating and reduction of agitation, nightmare and aggressive behavior in patients.
<b>Multiple Sclerosis</b>	Improved signs of EAE in mice, anti-inflammatory and immunomodulatory properties.
<b>Huntington's Disease</b>	Neuroprotective and antioxidant in mice transgenic models; no significant clinically important differences in patients.
<b>Hypoxemia Ischemia Injury</b>	Short term neuroprotective effects; inhibition of excitotoxicity, oxidative stress and inflammation <i>in vitro</i> and in rodent models.
<b>Pain</b>	Analgesic effect in patients with neuropathic pain resistant to other treatments.
<b>Psychosis</b>	Attenuation of the behavioral and glial changes in animal models of schizophrenia; anti-psychotic properties on ketamine-induced symptoms.
<b>Anxiety</b>	Reduction of muscular tension, restlessness, fatigue, problems in concentration, improvement of social interaction in rodent models of anxiety and stress; reduced social anxiety in patients.
<b>Depression</b>	Anti-depressant effect in genetic rodent model of depression.
<b>Cancer</b>	Antiproliferative and anti-invasive actions in a large range of cancer types; induction of autophagy-mediated cancer cell death; chemo preventive effects.
<b>Nausea</b>	Suppression of nausea and conditioned gaping in rats.
<b>Inflammatory Disease</b>	Anti-inflammatory properties in several <i>in vitro</i> and <i>in vivo</i> models; inhibition of inflammatory cytokines and pathways.
<b>Rheumatoid Arthritis</b>	Inhibition of TNF- $\alpha$ in an animal model.
<b>Infection</b>	Activity against methicillin-resistant staphylococcus aureus.
<b>Inflammatory Bowel and Crohn's Disease</b>	Inhibition of macrophage recruitment and TNF- $\alpha$ secretion <i>in vivo</i> and <i>ex vivo</i> ; reduction in disease activity index in Crohn's patients.
<b>Cardiovascular Disease</b>	Reduced infarct size through anti-oxidant and anti-inflammatory properties <i>in vitro</i> and <i>in vivo</i> .
<b>Diabetic Complications</b>	Attenuation of fibrosis and myocardial dysfunction.