The Impact of Caring
FOR A LOVED ONE WITH DRAVET SYNDROME

Caregivers and families are unsung heroes who play a vital role in the health and well-being of children and young adults with Dravet syndrome. The disease affects not only those with the condition, but also the entire family unit—parents, siblings, caregivers, and loved ones. This commitment can often result in a substantial financial, physical, psychosocial, and emotional burden.

WHAT IS DRAVET SYNDROME?
Dravet syndrome is a rare form of intractable (treatment-resistant) epilepsy that begins in infancy and is associated with potentially life-threatening frequent and severe seizures, developmental delay, cognitive impairment, and an elevated risk of sudden unexplained death in epilepsy (SUDEP).

Many Parents Experience the Impact at Work
81% stopped working because of their caregiving responsibilities

48 eight-hour work days missed per year

77 eight-hour work days of lost productivity at work per year

The Responsibility of Care Consumes Significant Amounts of Time
1 hr. or less per day is the only time 77% of caregivers report having completely for themselves

256 eight-hour work days of lost leisure time per year

The Burden Ripples Through the Entire Family
79% of young siblings (ages 9-12) expressed fear that their sibling might die

58% of siblings (ages 9-12) reported feeling worried or scared when their sibling has a seizure

66% reported having suffered from depression

33% have moderate to high levels of anxiety or depression due to the devastating nature of Dravet syndrome

It's important that people who care for children and young adults with Dravet syndrome learn ways to cope. Many advocacy groups offer tips and parent support groups to connect with other Dravet families. Visit the Dravet Syndrome Foundation at dravetfoundation.org to learn more about ways to manage caring for a loved one.

References: