# Childhood Tumour Trust 10 year timeline

## 2016

Childhood Tumour Trust (CTT) is registered as a charity

## 2017

CTT wins a National Diversity Award.

## 2018

First order for CTT's Body Map for the 'Personal Child Health Record'.

## 2019

Over 300 people attend CTT family days out to Alton Towers and Chessington.

The launch of CTT's CPD module on neurofibromatosis type 1.

## 2020

The launch of CTT's Zoom sessions to keep young people connected through Covid-19 and beyond.

## 2021

The unveiling of the CTT mascot, Patches.

'Patches and the Very Special Diagnosis' book is launched and is now available in seven languages, braille, British Sign Language and has been made into an animation.

## 2022

CTT take a group of young adults to a neurofibromatosis summit in New York.

CTT hold the first young adult camp.

## 2023

CTT fund a PhD into breast cancer awareness and neurofibromatosis type 1.

## 2024

CTT launch awareness campaigns for breast cancer and neurofibromatosis type 1 .

CTT's first cross-border family gathering at Dublin Zoo with NF Ireland.

## 2025

BMJ paper is published, 'Alone on our NF1 island': a patient-led mixed-method survey study to understand the care pathway for neurofibromatosis type 1 patients in the UK.

RARE REVOLUTION MAGAZINE

This graphic was created in collaboration with Childhood Tumour Trust and RARE Revolution Magazine