

*"I have more hope knowing that there is somebody who is really prepared to help me to confront the hard steps that I have to face and the fights I have to have, in order to get heard where it matters, when caring with someone with this disease!"*

# Impact report 2021/2022



Huntington's  
Disease  
Association

The Huntington Disease Association wants to give a better life for people with Huntington's disease. We do this in a number of ways, this report shows the difference we make. This is all possible thanks to your support.

## Advisory Service



We have 19 advisers based across England and Wales. They are there to support anyone affected by Huntington's disease

1

Youth Engagement Coordinator



4060

Individuals Supported



1026

Family Sessions



317

Individuals supported by Youth Engagement Coordinator

## Professionals



We held two courses for professionals, attended by **84 people**



We delivered 103 training sessions attended by **891 people**



We held 102 information sessions attended by **967 people**



# Branches, Support Groups, and HD Voice

39

Branches and support groups



5

Leader sessions delivered



13

Families with Huntington's disease contributed to HD Voice



## Webinars



We held 26 webinars attended by **612 people**



We held 57 carers groups attended by **301 people**



4

Sessions were attended by Young Adults



**18,740**

Conversations with families



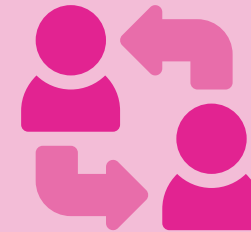
## Website and Social Media



**268,766**

Web visits

## Supporting people with Juvenile Huntington's disease



**9**

Peer support sessions

## Working with Professionals

### Followers

 15,286

 2,216

 3,840

 1,312



**41**

People supported



**484**

Multi-Disciplinary meetings attended

### Engagement

(number of interactions)

 117,393

 78,469

 8,812

 5310



**64**

Clinics attended



**31**

Welfare grants awarded

