



## Crohn's & Colitis Top Ten Impacts Survey

### What's the aim of this Survey?

The goal of this survey is to understand the ways that Crohn's and Colitis affects the lives of people living with these conditions, allowing us to identify key areas for improvement.

The survey is based on our 'Top Ten Impacts project, which identified the ten areas of people's lives most impacted by living with Crohn's and Colitis in the UK.

[You can read more about the project here](#)

### Who is running the survey?

The project is being run by Crohn's & Colitis UK. The data is being collected by Common Collective, a team of researchers, who then provide it to us.

### What does the survey involve?

The survey should take around 10 minutes to complete.

You will be asked a series of questions about your experiences related to the top ten impacts.

### Who can take part??

In order to take part you must:

- Be diagnosed with Crohn's, Colitis or another form of IBD
- Be 18 years+
- Live in the UK

### How will taking part make a difference?

By sharing your experiences, Crohn's & Colitis UK will deepen their understanding of how your condition impacts your quality of life. We will share key findings from the survey publicly to enable the charity to help drive meaningful change.

### What happens to your data?

All data collected from the survey will be anonymised and made available on a digital hub, ensuring that individual responses remain confidential:

- When you answer questions in this survey you will be sharing data about yourself and the ways you are affected by your condition with Common Collective and Crohn's & Colitis UK.
- The data will be stored securely electronically by Common Collective Ltd on an encrypted secure EU-based cloud server.
- The Common Collective research team will also review anonymised data to analyse responses.
- The anonymised data also will be securely transferred to Crohn's & Colitis UK and be used at an aggregate level to illustrate the impact these conditions have on quality of life. By aggregate level we mean that the data that Crohn's & Colitis UK shares publicly to illustrate the findings about the impacts for all those living with the condition who completed the survey. Findings will represent group level responses and never individual responses.