

Patient Data Journey

Step 1

- Panthera works with your local Health Care Professional (HCP) to identify patients with particular diagnoses in compliance with the General Data Protection Regulation law.
- The HCP sends this information directly to Docmail, an NHS-approved mailing company, which dispatches the letters out. Please note – Panthera has no access to any patient data at this point.



Step 2

- If you feel a trial invitation letter is something you'd like to explore in more detail, you can contact Panthera's call centre. We will ask you a few basic questions about you and your condition.
- You might then be eligible to undergo a more detailed examination at one of Panthera's clinics, which will be explained to you over the phone. All calls are recorded for quality assurance purposes and you will be made aware of this.



Step 3

- If your call handler informs you that you are suitable to proceed to an initial clinic visit with one of our Research Physicians, we will ask for your consent to be put into our database and will book you in for an appointment. We will also supply certain information regarding the local Panthera clinic.
- If you consent but choose not to participate in the specific trial, we won't send you any further information but will check with you whether we can contact you in future for other potentially relevant clinical trials. If you opt out at this point, we will record your preferences on our database and you won't hear from us again.

