You are being invited to take part in a research project. Before you decide if you are happy to take part, and if you would like your child to take part, it is important for you to understand why the project is being done and what taking part will involve. Please take time to read this information sheet and decide whether or not you would like to take part and discuss it with others if you wish. Ask us if there is anything that is not clear or if you would like more information. Thank you for taking the time to read this.

Who is doing this project?
The project is being led by Professor Lucy Bray at Edge Hill University who is working with an international group of nurses, doctors, play specialists, psychologists, youth workers, children and young people and parents to develop the rights based standards. You can find out more about the team, project and standards on our webpage. https://edgehill.ac.uk/health/research/rights-based-standards-for-children-undergoing-clinical-procedures

What is the purpose of the project?
We are inviting you to share your views about the standards to help us work out if they are useful for children, parents and professionals and if there is anything we need to change. We are keen to hear from children aged 7-15 years old, parents/carers and professionals. Your child does not need to have had a procedure to be able to take part.

Why have we been invited to take part?
We would like to make sure the standards are useful for children, parents and professionals and so we are asking for feedback to help us see what you think of them and if anything needs changing.

Do I or my child have to take part?
No, it is up to you to decide whether or not you or your child take part. You and your child can take part separately from each other. If your child would like to share their views, we will just ask you to tick a box at the beginning of their survey to say you have given them permission before they start answering the questions.
What will happen if I take part?
You will be asked to read the standards and tell us what you think in a short online survey. The survey will take about 10-15 minutes to fill out, depending on how much you want to tell us. You will not have to answer any questions you do not want to, and you can end the survey at any time. We will not be asking for any information which will identify who you are. Please do not add any identifiable information to the survey. If you click ‘done’ at the end of the survey then this means you are happy for us to use your answers as part of the project. As it is anonymous, once you have clicked done or submit then you cannot take your answers back.

What will happen if my child takes part?
If your child decides they would also like to take part and share their views then please read the children's information sheet about the study with them. There is a link to the children’s survey at the end of the parent survey. There is also a box at the beginning of their survey which we will ask you to tick to say you are happy for them to take part. The survey for children will take about 10-15 minutes to fill out. They can share their views by ticking the boxes or typing in what they think. We will not be asking for any information which will identify who they are. Your child will not be able to go back and see your answers.

Are there any possible disadvantages or risks from taking part?
We hope that taking part will not involve any risks for you or your child. Some of the questions may be upsetting as they may prompt you or your child to think back to procedure they have had in the past. You do not have to answer any questions that you do not want to and at the end of the survey there are some suggestions for places for support, you can skip straight to these if you like. As we cannot identify who has filled out a survey, we will not be able to offer individual help or support.

What are the possible benefits of taking part?
We hope that what we find out from this project will help us further develop the Rights- based standards' to make a difference to children having a clinical procedure.

What will happen to the results of the project?
The research team will share the results in articles, presentations and we will also work with parents and children to write a summary of the key findings which will be posted on our webpage. The report from the project will be shared with key organisations across the world.
Who has reviewed this study?
This study has been reviewed and given ethics clearance through the Edge Hill University Health Research Ethics Committee (HREC) Reference: ETH2021-0261

What if I have a question, concern or something goes wrong?
If you have a question or concern about any part of this project, please contact the lead researcher, Professor Lucy Bray (brayl@edgehill.ac.uk).
If you wish to make a complaint about the project please contact Philip Bentley, URESC Secretary in the research office of the University at research@edgehill.ac.uk who is independent to the project team.

Data Protection Notice
Edge Hill University is the organiser for this study based in the United Kingdom. We will be using information from you in order to undertake this project and will act as the data controller for this project. This means that we are responsible for looking after your information and using it properly. Edge Hill University will process your data for the purpose of research. Research is a task that we perform in the public interest. Your rights to access, change or withdraw your information is limited, as you have taken part in the survey anonymously. We are not collecting any information from you which is identifiable.

You can find out more about how we use your information by referring to the University privacy policy: edgehill.ac.uk/about/legal/privacy

Thank you for reading this information sheet and thinking about taking part in this project