

# National Registry of Rare Kidney Diseases (RaDaR)

## Children's Information Sheet

Thank you for reading this information sheet.

The problem with your kidneys is rare - that means not many people have it. Doctors want to find out more about it. They have set up a project called **RaDaR** to collect information on people like you who have rare kidney problems.

If you would like to take part, information about your kidney problem will be put on a computer database for the researchers to look at.

You and your family can also see this information with a special password on a website called Patient View if your hospital have this.

Researchers may sometimes write to you and your family with information about other research projects. It is up to you if you want to take part in these.

If you have any questions about RaDaR you can talk to your doctor when you next visit the hospital.