



Are you or someone you know living with Alport Syndrome?

Do you want to find out more?

**Saturday
21st November
2015, 10am-4pm**

**Castlefield Rooms
18-20 Castle Street
Manchester
M3 4LZ**



Travel bursaries available if your journey costs more than £20 (with receipts)

Lunch will be provided as well as activities for children. There will be a separate workshop for teens and young adults

An informal event created by those living with Alport Syndrome and supported by:



The 4th National Alport Information Day

New content and format:

- Hear about the latest work on new treatments for Alport Syndrome from The 2015 International workshop on Alport Syndrome in Germany
- Put your questions to the medical team - Kidney, Genetics, Organ transplantation, Hearing and Sight experts
- Hear about the experiences of others living with Alport Syndrome and our new research project to improve the quality of life

Special focus: the Manchester team will run specific breakouts for teenagers and young adults

What to do next...

Registration now open. Simply email your details (name, address, phone number and email) to: info@alportuk.org

By registering you will receive updates and information

a brighter future for people living with alport syndrome

Registered charity number 1154774

shaping futures

Royal
Manchester
Children's
Hospital



University of Manchester