



Are you or someone you know living with Alport Syndrome?

**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:



a brighter future for people living with alport syndrome

Registered charity number 1154774

The 4th National Alport Information Day

- Introductions – Susie Gear**
 - Introduce yourself: our name and where you are from
 - Tell us about something fun that you have done in the last year
- Manchester research – Rachel Lennon**
- Genetics - Frances Flinter**

Breakout for teenagers and young adults – Future of Alport Syndrome research – Rachel Lennon and Salmon xxxx

- Kidneys and hearing – Neil Turner**
- Sharing experiences of living with Alport Syndrome - Melissa Stepney**
- Diet – Julie xxxxxx (need to check surname in email)**

Lunch

- Eyes - Omar Mahroo**
- Registries and Athena study – Neil Turner**
- Breakout reports back on Alport Syndrome research – Rachel Lennon with teens/young adults**
- alport uk – Susie Gear**

Make sure we have your email...

Email us at info@alportuk.org, so you can receive updates and information



University of Manchester