



# Are you or your family living with Alport Syndrome?

## Do you want to find out more?

**Saturday  
1 December  
2012, 10am-4pm**

**Where:** Central London venue to be confirmed, close to good transport links.

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, family event created by those living with Alport Syndrome and supported by:



## The 1st National Alport Family Support Day

- Learn more about Alport Syndrome with Kidney, Genetics, Organ transplant, Hearing and Sight experts - put your questions to the medical team
- Find out about the experiences of others living with Alport Syndrome
- Meet families in similar situations and share your experiences

### What to do next...

**Registration now open.** Simply email your details (name, address, phone number and email) to: [meetings@alport.info](mailto:meetings@alport.info)

We are also planning an **International Science workshop** in Autumn 2013

By registering you will receive updates and information



RARE DISEASE

UK

The National Alliance for people with rare diseases & all who support them