

# Families for the Treatment of Hereditary MND (FATHoM)



18<sup>th</sup> April 2017

Saïd Business School, Oxford University  
Park End Street, Oxford  
Hosted by the Oxford MND Research and Care Centre

<b>10:00 – 10:30</b>	<b>WELCOME, TEA &amp; COFFEE</b>
<b>10:30 – 11:15</b>	<b>THE GENETICS OF MND</b>
<b>11:15 – 11:30</b>	<b>REFRESHMENTS</b>
<b>11:30 – 12:30</b>	<b>GENETIC TESTING</b>
<b>12:30 – 14:00</b>	<b>LUNCH</b>
<b>14:00 – 15:00</b>	<b>RESEARCH</b>
<b>15:15 – 15:30</b>	<b>REFRESHMENTS</b>
<b>15:15 – 16:00</b>	<b>EMERGING AND FUTURE TREATMENTS</b>
<b>16:00 – 17:00</b>	<b>Q&amp;A SESSION</b>

Families for the Treatment of Hereditary MND (FATHoM) is an initiative to bring together the community of families affected by inherited (genetic) forms of MND. This first event is a meeting led by Professor Martin Turner and Professor Kevin Talbot consisting of expert talks on key issues affecting such families. It is open to any person living with an inherited form of MND, and the close relatives of those affected by inherited forms of MND. Registration, lunch and refreshments are provided free of charge. There is charged parking at the nearby railway station car park (accessed via Osney Lane). To register or for more information, visit [this link](#) or email [lynn.ossheer@ndcn.ox.ac.uk](mailto:lynn.ossheer@ndcn.ox.ac.uk)