

Appendix 1: Survey 1**Patients' and Families' Experiences Following Treatment for a Brain Tumour in Childhood**

1. What is your connection to brain tumours?
 - I have/had a brain tumour as a child
 - I have/had a brain tumour as a young adult
 - I am a parent/carer of a child with a brain tumour
 - Other (please specify)
2. Was the person provided with paediatric or adult services to recover after tumour treatment?
 - Paediatric services
 - Adult services
 - Don't know
 - Other (please specify)
3. What treatment, if any, was given for the brain tumour?
4. At which hospital(s) was neurosurgical treatment of the tumour done?
5. What help did you receive from nurses, therapists, and doctors in the health service?
6. What help did you receive from psychologists and teachers in health or educational services?
7. What are your experiences with help outlined in Q5 and Q6, and could they be improved upon?
8. Which aspects of help during recovery do you think is most important for children and young adults to receive after treatment for a brain tumour?
9. What questions relating to recovery after treatment for a brain tumour would you like to be answered by future research?

Please add any further information that you think would be helpful to us from your experiences (or lack) of rehabilitation services, including how we could improve on this, if at all.