

*Drug resistant Epilepsy*

*Number (percentage, of total unless otherwise specified)*

<i>Total</i>	131
<i>Underwent epilepsy surgery</i>	97 (74%)
<i>Resective surgery</i>	82 (62.6%)
<i>Disconnection</i>	15 (11.5%)
<i>Surgery recommended but parents declined</i>	11 (8.4%)
<i>Evaluated but not felt to be a surgical candidate</i>	6 (4.6%)
<i>Awaiting discussion or surgery</i>	4 (3.1%)
<i>Not referred for surgical assessment</i>	6 (4.6%)
<i>PET scan performed</i>	29 (22.1%)
<i>Invasive monitoring performed</i>	30 (22.9%)
<i>&gt;1 year post surgery follow up</i>	92 (70.2%)
<i>Engel Scale 1 at 1 year follow up</i>	55 – (59.8% of those > 1 year post surgery)
<i>Engel 1-3 at 1 year follow up</i>	76 – (82.6% of those > 1 year post surgery)
<i>Engel Scale 4 at 1 year follow up</i>	16 – (17.4% of those > 1 year post surgery)
<i>Required further epilepsy surgery</i>	13 – (14.1% of those > 1-year post surgery). 1 underwent vagal nerve stimulator implantation
<i>Histology confirmed FCD</i>	68 (70% of surgical cohort); Resection 71.4%, Disconnection 60%, p=0.3762
<i>Histology nondiagnostic</i>	13 (13.4% of surgical cohort)
<i>Engel Scale 1 in those with confirmed FCD</i>	41 (64% of FCD 1 year post surgery)
<i>Engel Scale 1 in those with non-FCD histology</i>	14 (50% of those 1 year post surgery)
<i>FCD Subtypes (% total FCD)</i>	FCD Type 2a - 12 (17.6%) FCD Type 2b - 47 (69.2%) FCD – subtype unknown - 9 (13.2%)
<i>Other Histology (n=29)</i>	Non-diagnostic - 19 Gliosis – 3 Polymicrogyria – 2 Normal – 1 Glioneuronal tumour – 1 Low grade ganglioglioma (NOS) – 1 Hippocampal sclerosis (no FCD) – 1 Pleomorphic xanthoastrocytoma (PXA) – 1

Table 2. Pharmacoresistant and epilepsy surgery patients. *Abbreviations: Focal cortical dysplasia (FCD), Positron emission tomography (PET), Multidisciplinary Team (MDT)*