

Stefanie Frahssek¹, Rene Schmidt⁵, Andreas Faldum⁵, Johannes Wolff⁶, Gudrun Fleischhack⁷, Monika Warmuth-Metz⁸, Jürgen Krauss⁹, Rolf-Dieter Kortmann¹⁰, Niels Galley¹¹, Joachim Kühl¹, and Stefan Rutkowski³; ¹University Children's Hospital, Würzburg, Germany, ²Department of Psychology, University of Bochum, Bochum, Germany, ³Department for Pediatric Hematology and Oncology, University Medical Center Hamburg-Eppendorf, Hamburg, Germany, ⁴Department for Pediatric Oncology, Charité University-Medicine, Berlin, Germany, ⁵Institute of Biometry, University of Münster, Münster, Germany, ⁶Abbview, Oncology Development, Chicago, IL, USA, ⁷Pediatrics III, University Children's Hospital, Essen, Germany, ⁸Department of Neuroradiology, University Medical Center Würzburg, Würzburg, Germany, ⁹Department of Neurosurgery, University Medical Center Würzburg, Würzburg, Germany, ¹⁰Department of Radiotherapy, University of Leipzig, Leipzig, Germany, ¹¹Institute of Psychology University of Cologne, Cologne, Germany

Neurocognitive deficits are frequent in childhood brain tumor survivors and affect mental intelligence, psychomotor and executive abilities. The differential impact of factors such as disease (location, histology) or treatment (local (LI) vs. craniospinal irradiation (CSI)) on these parameters is not fully understood. Between 2007–2011 and 2013–2017 300 testings were performed on-site by one neuropsychologist. Of these, 274 tests from n=208 children with medulloblastoma (MB), ependymoma (EP) and supratentorial embryonal tumors (SET) <4 years at diagnosis are currently included into the analysis. Applied tests included the Bayley II, WUEP-KD, K-ABC, tapping speed (TS), CPT_Hits/CPT_DT, and, as a new score, CPT_Power which integrates the latter. Treatment consisted of surgery and chemotherapy ± LI/CSI. All children receiving CSI and MB children with LI showed substantial deficits in general intelligence scores. In contrast, children with MB or SET without CSI/LI and those with EP receiving LI performed surprisingly well after 2 and 5 years follow-up. Motor function (TS) was reduced in all children except those with SET without irradiation. Of note, mental processing speed (as measured in CPT_Power) was not essentially reduced in MB and EP patients, indicating that mental processing is less affected than motor speed (TS) in children with infratentorial tumors. In conclusion, our data show that besides the established detrimental effects of CSI on general intelligence, infratentorial tumor location is a main risk factor for motor dysfunction irrespective of irradiation. Appropriate sensitive testing tools are warranted to assess cognitive function without the interfering influence of motor dysfunction.

QOL-25. MICROSTRUCTURAL BRAIN CHANGES ASSOCIATED WITH NEUROCOGNITIVE AND FUNCTIONAL OUTCOMES OF INTRACRANIAL GERM CELL TUMOUR SURVIVORS – A DIFFUSIONAL KURTOSIS IMAGING STUDY

Wan-Yee Winnie Tso¹, Sai Kam Hui², Tatia Mei Chun Lee³, Anthony Pak Yin Liu¹, Patrick Ip¹, Kevin Cheng⁴, Daniel Fong⁵, Dorita Chang³, Frederick KW Ho⁶, Ka Man Yip¹, Dennis Ku⁷, Daniel KL Cheuk⁷, Chung Wing Luk⁷, Ming Kong Shing⁷, LK Leung¹, Pek Lan Khong², and Godfrey Chi Fung Chan¹; ¹Department of Paediatrics & Adolescent Medicine, LKS Faculty of Medicine, University of Hong Kong, Hong Kong, Hong Kong, ²Department of Diagnostic Radiology, LKS Faculty of Medicine, University of Hong Kong, Hong Kong, Hong Kong, ³State Key Laboratory of Brain and Cognitive Sciences, The University of Hong Kong, Hong Kong, Hong Kong, ⁴Department of Neurosurgery, Queen Mary Hospital, Hong Kong, Hong Kong, ⁵Department of Nursing, LKS Faculty of Medicine, University of Hong Kong, Hong Kong, Hong Kong, ⁶Institute of Health and Wellbeing, University of Glasgow, Glasgow, United Kingdom, ⁷Hong Kong Children's Hospital, Hong Kong, Hong Kong

BACKGROUND: Childhood intracranial germ cell tumour (iGCT) survivors are prone to radiotherapy-related neurotoxicity which can lead to neurocognitive dysfunction. Diffusion kurtosis imaging (DKI) is a MRI technique that quantifies microstructural changes in the grey and white matter of the brain. This study aims to investigate the associations between MR-DKI metrics, the cognitive and functional outcomes of childhood iGCT survivors. **METHOD:** 20 childhood iGCT survivors who had received cranial radiotherapy were recruited. DKI parameters were determined for iGCT survivors and 18 control subjects. Neurocognitive assessment using the Hong Kong Wechsler Intelligence Scales for Children (HKWISC)/ Wechsler Adult Intelligence Scale – Revised (WAIS-R) and functional assessment using the Lansky/ Karnofsky performance scales were performed for GCT survivors. **RESULTS:** There were significant negative correlation between the IQ scores and the mean diffusivity (MD) in multiple white matter regions of iGCT survivors including: anterior limb of internal capsule, superior fronto-occipital fasciculus, anterior corona radiata, uncinate fasciculus, cingulum and hippocampus. Mean kurtosis (MK) values of the superior fronto-occipital fasciculus were positively correlated with IQ scores. For grey matter, the MD of the olfactory, insula, caudate, heschl gyrus, parahippocampal gyrus, hippocampus, anterior cingulum, frontal inferior operculum, middle and superior temporal gyrus, middle and superior frontal orbital gyri, cuneus and

precentral gyrus were negatively correlated with IQ scores. Most of the microstructural changes with associated functional impairment were white matter regions. **CONCLUSION:** Our study identified vulnerable brain regions with significant white and grey matter microstructural changes that were associated with impaired cognitive function or deficits in physical functioning.

QOL-26. I'VE GOT FRIENDS NOW: PAEDIATRIC PATIENTS' EXPERIENCES OF METHYLPHENIDATE TREATMENT FOR NEUROCOGNITIVE LATE-EFFECTS ASSOCIATED WITH BRAIN TUMOUR

Sarah Verity^{1,2}, Rebecca Hill^{1,2}, Gail Halliday¹, Jade Ryles¹, and Simon Bailey^{1,2}; ¹Newcastle Upon Tyne Hospitals NHS Foundation Trust, Newcastle Upon Tyne, United Kingdom, ²Northern Institute of Cancer Research, Newcastle University, Newcastle Upon Tyne, United Kingdom

BACKGROUND: Whilst rates of survival following paediatric brain tumour have increased, quality of survival continues to present a significant challenge to children and their families. The neurocognitive impact of cranial radiotherapy (CRT) in childhood upon future intellectual development is well established. Both CRT and chemotherapy are associated with medium-term slowed speed of cognitive processing, attention, and memory impairment, and with longer-term failure to achieve pre-morbid intellectual potential and low Health-Related Quality of Life (HRQoL). Methylphenidate is a psychostimulant drug shown to be effective in alleviating some of the neurocognitive symptoms of cancer treatment, however the subjective experience of paediatric participants is not reported. **AIM:** The current study aimed to explore the subjective experience of HRQoL in paediatric neuro-oncology patients currently receiving methylphenidate. **METHODS:** A retrospective audit was conducted on 12 paediatric neuro-oncology patients in receipt of methylphenidate. Both standardised and novel measures were used to assess aspects of HRQoL, specifically; social life, perceived independence, mood, confidence, school life, self-esteem, interpersonal relationships and fatigue levels. Data collected were analysed using Thematic Analysis. **RESULTS:** Five key themes were identified; physical, emotional, social, academic and neuropsychological impact. **CONCLUSION:** The current findings evidence the perception of patients that methylphenidate supported them to regain previously lost functionality. Methylphenidate has the potential to increase HRQoL in this population and to provide children with the opportunity to regain a sense of normality in their lives.

QOL-27. SWALLOWING ASSESSMENT IN PEDIATRIC PATIENTS WITH BRAIN TUMOR

Natalia Oliveira Machado, Ana Paula Duarte, Aline Azevedo dos Santos, Bruna Minniti Mançano, and Carlos Almeida Jr; Barretos' Children and Young Adults Cancer Hospital, Barretos, Sao Paulo, Brazil

BACKGROUND: Neurosurgical intervention is the initial modality of treatment for the vast majority of pediatric brain tumors. However, studies on the swallowing process in pediatric patients with brain tumors are scarce, especially comparing changes that can be identified before and after surgery. In clinical practice, it is possible to observe that these patients may present modifications in the swallowing phases both before and after surgery. Therefore, we conducted a longitudinal study with a cohort of 20 patients ranging in age from 0 to 17 years, in order to characterize the swallowing disorders. **RESULTS:** 30% of the patients presented some change in orofacial motricity in the organs related to initiation, coordination, and maintenance of swallowing at the time of hospital admission, and 65% of the patients exhibited these changes after surgery. Due to worsening in swallowing after surgery, 40% of the patients required modification of the consistency of oral diet or required the use of an alternative route of feeding. **CONCLUSIONS:** There is a high prevalence of swallowing disorders in pediatric patients with brain tumors, mainly regarding the proper functioning of organs related to initiation, coordination, and maintenance of swallowing even before the surgical intervention, and these changes increase after surgery - especially in patients with posterior fossa tumors. The role of the speech/language pathologist is of paramount importance, given their role in the assessment and adequacy of the feeding route, identifying patients at risk of pulmonary aspirations, minimizing swallowing complications, and also facilitating communication with patients and their families.

QOL-28. NEUROCOGNITIVE PROFILE OF PEDIATRIC MEDULLOBLASTOMA PATIENTS PRIOR TO RADIATION THERAPY

Muhammad Baig, Ineke Osthorn, Susan McGovern, David Grosshans, Mary McAleer, Kristina Woodhouse, Arnold Paulino, Grace Yang, Peter Stavinoha, and Wafik Zaky; MD Anderson Cancer Center, Houston, TX, USA

Neurocognitive late effects are unfortunately common following treatment for pediatric medulloblastoma. While radiation therapy is an essen-