# **Objectives**

Typical hemolytic Qualitative analysis revealed lesions on diffusion-(HUS) uremic syndrome İS characterised by the triad of acute renal failure, weighted imaging (DWI) in basal ganglia and/or thalami thrombocytopenia and hemolytic anemia and in with corresponding T2 hyperintensities in subgroup A, Europe is mostly caused by Shiga toxin-producing whereas subgroup B did not show qualitative DWI Escherichia coli (STEC). Neurological involvement is alterations even if T2 hyperintense lesions were found found in between 17 – 52%. Magnetic resonance in these regions (Table 1). Quantitative analysis demonstrated abnormal ADC values in all HUS patients imaging (MRI) may reveal abnormalities especially affecting the basal ganglia and / or thalami, but also with a trend to more regions being affected in the deep white matter. Early MRI may offer such subgroup A (16 versus 10.5 respectively, median options, as previously postulated by Donnerstag et al. number of regions, p=0.18 (Table 2)). [1].

We aimed to evaluate microstructural cerebral changes in children suffering from hemolytic uremic syndrome (HUS) based on apparent diffusion coefficient (ADC)

# Viethods

In 9 pediatric HUS patients with neurological symptoms with a median age 3.0 y (0.8 - 14.6), conventional magnetic resonance imaging (cMRI) at 1.5T was retrospectively analysed. All examinations included axial diffusion weighted, single-shot, spin-echo echo planar sequences. ADC values were measured in 35 different brain regions and compared with age-related ADC reference values from previously published pediatric controls. Depending on clinical outcome our **Table 1** Qualitative imaging findings in basal ganglia/ thalami in cohort was divided into 2 subgroups. Subgroup A children with HUS: basal ganglia/thalamic lesions with T2 hyperintense showed devastating neurological outcome whereas corresponding DWI alterations were only found in subgroup A subgroup B showed improvement without lasting with devastating neurological outcome. neurological deficit.

Cerebral microstructural changes in children suffering from hemolytic uremic syndrome Bültmann E<sup>1</sup>, Zapf A<sup>3</sup>, Mussgnug HJ<sup>1</sup>, Lanfermann H<sup>1</sup>, Kanzelmeyer N<sup>2</sup> Hartmann H<sup>2</sup>

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### Results

Pat ID	Age (y)	Subgroup	Basal ganglia and/or thalamic alterations		
			Hyperintense T2 lesions	DWI lesions	
1	1.4	А	+	+ (DWI	
4	2.3	А	+	+ (DWI 압, ADC =)	
6	7.3	А	+	+ (DWI	
8	3	А	+	+ (DWI 압, ADC 圦)	
9	0.8	А	_	_	
9 FU	0.8	А	+	+ (DWI 압, ADC - ()	
2	3.5	В	+	_	
3	1.8	В	_	_	
5	10	В	_	_	
7	14.6	В	+	-	



Subgroup	Number	of regions w	Number of	
		ADC values	hyperintense T2	
	Below	Above	Total	regions
A	51 (29%)	21 (12%)	72 (41%)	28
B	17 (12%)	20 (15%)	37 (27%)	5
Total A + B	68 (22%)	41 (13%)	109 (35%)	33

**Table 2** Frequencies of ADC values below and above the agedependent reference range per HUS subgroup

# Conclusion

Using DWI images qualitative and quantitative differences were found between HUS patients with devastating neurological outcome and those without. ADC values may help to detect more extensive cerebral changes than conventional qualitative findings. Both modalities together may permit prognostic statements in pediatric HUS patients.

#### **References:**

1. Donnerstag F, Ding X, Pape L, Bultmann E, Lucke T, Zajaczek J, et al. Patterns in early diffusion-weighted MRI in children with haemolytic uraemic syndrome and CNS involvement. Eur Radiol. 2012;22:506-13.