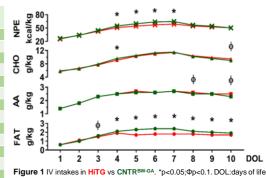


Hypertriglyceridemia in Very Low Birth Weight Infants on Routine Parenteral Nutrition: A Case-Control Study



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Background and Objectives		Table	1	HiTG	CNTR ^{BW-GA}	Diff.	р
Hypertriglyceridemia (HiTG) often occurs in infants on parenteral nutrition (PN), especially those with low birth weight (BW). In case of HiTG, the ESPGHAN/ESPEN/ESPR 2018 guidelines recommend an intravenous (IV) lipid (FAT) titration. The consequences of IV FAT	Г	Birth	weight (BW) - g	875±173	877±170	-2±41	0.5
		Gest	ational age (GA) – days	193±15	193±14	-0±1	0.9
			s, no. (%)	69 (51%)	64 (47%)	+5 (+4%)	0.5
titration in small infants are largely unknown.	_ T	SGA	^{2SDS} , n (%)	15 (11%)	13 (10%)	+2 (+1%)	0.5
To investigate the modifications of IV FAT, amino acids (AA),			ır5min – no	8 7 8	8 7 8	0	0.4
carbohydrates (CHO) and non-protein energy (NPE) intakes in infants	፟ .	Surfa	actant therapy- no. (%)	97 (71%)	85 (63%)	+12 (+9%)	0.1
with a BW less than 1250g on routine PN who developed HiTG (>250 mg/dL).	L	Perir	atal steroids-no. (%)	115 (88%)	117 (90%)	-2 (-2%)	0.7
	Г	- LOS		30 (22%)	36 (26%)	-6 (-4%)	0.4
Methods	ES	NEC	≥ grade II	8 (6%)	6 (4%)	+2 (+1%)	0.6
	ဗ	Chol	estasis	13 (10%)	15 (11%)	-2 (-1%)	0.7
We retrospectively reviewed nutrition, growth and neurodevelopment		RDS	and HMD	124 (91%)	123 (90%)	+1 (+1%)	0.8
of a cohort of infants with a BW<1250 g consecutively admitted to the "G. Salesi" Children's Hospital between Jan-2004 and Dec-2016 who		BPD		34 (25%)	42 (31%)	-8 (-6%)	0.3
received routine PN. Patients with and without HiTG were match- paired for BW and gestational age (GA). Paired t-test, Mann-Whitney test, Wilcoxon test or McNemar test were used for the statistical analysis.	<u> </u>	Asph	yxia	8 (6%)	7 (5%)	+1 (+1%)	0.8
	S.	PDA		89 (65%)	95 (70%)	-6 (-4%)	0.4
	ž	PVL	II-IV	3 (2%)	6 (4%)	-3 (-2%)	0.3
	ÌΙ	IVH 2	≥ grade III	12 (9%)	10 (7%)	+2 (+1%)	0.7
Results	L	ROP	≥ grade III	2 (1%)	0 (0%)	+2 (+1%)	0.3
	Г	— Weig	jht 36W – g	1861±354	1895±343	-34±283	0.2
Six hundred and fifty-eight infants of the cohort were analyzed and 196 (30%) had at least one HiTG episode in the first 10 days of life (DOL). One hundred and thirty-six HiTG patients were compared with 136 matched-controls (CNTR, Table 1). In the first 10 DOL, IV FAT and NPE, but not IV AA, were significantly	0 0	Weig	ht SDS-36W - no.	-1.91±0.88	-1.81±0.86	-0.10±0.71	0.1
)W-UP n=120	Tota	Lenght 36W - cm	42.6±2.5	42.7±2.5	-0.1±2.2	0.7
	<u>``</u>	Tota	Lenght SDS-36W- no.	-1.87±0.99	-1.84±0.96	-0.04±0.85	0.7
	q		d Circ 36W – cm	30.5±1.5	30.6±1.5	-0.1±1.9	0.5
lower in HiTG infants (Figure 1). The incidence of hyperglycaemia	d NPE, but not IV AA, were significantly 1). The incidence of hyperglycaemia HC SDS-36W - no1.63±1.02 -1.53±1.08 -0.10±1.	-0.10±1.36	0.4				
episodes associated with HiTG, was not significantly higher than in		Weig	ht at 24Mo – g	11627±1759	11562±2184	+65±2564	0.9
CNTR (12% vs 7%, p=0.2). We found no differences between groups	p=0.2). We found no differences between groups WTSDS-24Mo - no -0.29±1.30	-0.24±1.69	-0.05±1.90	0.9			
in the incidence of the main complications of prematurity.	§ ₹	TL at	24Mo – g	87.5±4.2	86.1±4.6	+1.4±6.0	0.1
Anthropometry at 36 weeks (W) and at 24 months corrected age (CA), and neurodevelopment at 24Mo CA (HiTG vs CNTR - Cognitive	Ж, <u>Г</u>	TL S	DS-24Mo - no	0.72±1.36	0.44±1.46	+0.28±1.82	0.3
composite score, n=39: 94±13 vs 95±14, p=0.6, Motor composite	Weight at 24Mo - g 11627±1759 11562±2184 +65±2564 0.9 pups H WT SDS-24Mo - no -0.29±1.30 -0.24±1.69 -0.05±1.90 0.9 rifty. TL at 24Mo - g 87.5±4.2 86.1±4.6 +1.4±6.0 0.1 CA), ititive B TL SDS-24Mo - no 0.72±1.36 0.44±1.46 +0.28±1.82 0.3						
score, n=22: 100±14 vs 101±11, p=0.8) were also not different.	L	_ HC S	SDS-24Mo - no	-1.05±1.56	-0.72±1.24	-0.33±1.78	0.2



Conclusion

titration resulted in a significantly lower IV FAT and NPE intakes. HiTG

preterm infants receiving parenteral lipid emulsions. BMC Pediatr

At our institution, HiTG occurred in 30% of VLBW infants and IV FAT

infants on routine PN had similar growth and neurodevelopment than controls

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Disclosure & Author Affiliation

No conflict of interest to disclose.

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