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Comparison of Two Fat Emulsions on Interleukin-1β, Interleukin-8 and Fatty Acid Composition in Infants Post Gastrointestinal Surgery: A Randomized Trial Protocol V.2

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Abstract

Surgical intervention in infants is associated with postoperative sepsis and severe outcomes due to immature immune system¹. Gastrointestinal surgery induces excessive cytokine secretion, which may lead to increased postoperative mortality and morbidity².

Parenteral nutrition plays a crucial role in pediatric patients who undergo gastrointestinal surgery³. Intravenous lipid emulsion is an integral part of parenteral nutrition because it contains high energy density and low osmolarity, hence becoming the main source of energy and essential fatty acids^{4,5}. Fatty acid determines structural integrity and fluidity of cell membrane, and it has been proven that fatty acid helps to regulate expression of various genes and modulate cell-signaling pathway, which occurs during inflammation^{6,7}. The current standard type of IVFE is a 50:50 mixture of medium chain triglyceride (MCT) and long chain triglyceride (LCT)⁸. This type of emulsion is rich in ω -6 and contains high levels of linoleic acid (LA, C18:2 ω -6) and alpha-linolenic acid (ALA, C18:3 ω -3). According to several studies, ω -6 is associated with impaired cell-mediated immunity and higher potential risk of elevated proinflammatory markers and severe inflammatory response. This mechanisms may lead to the increase in mortality, morbidity, duration of treatment, and recovery time in patients who undergo gastrointestinal surgery⁹⁻¹¹.

Calder (2010) showed that the structure modification of fatty acids may alter their functions7. Some studies have shown that the addition of ω -3 in soy oil-based fat emulsion may improve patients' outcome by modulating inflammatory response^{3,4,12}. ω -3, particularly eicosapentaenoic acid (EPA) and doxohexanoic acid (DHA), is a competitive enzyme inhibitor of arachidonic acid (AA). ω -3 has potential anti-inflammatory properties by inhibiting AA pathway and generating inflammatory eicosanoids, such as prostaglandine E3, thromboxane A3 and leukotriene B5, which are considered less inflammatory. Up to this time, the effect of ω -3-enriched intravenous fat emulsion compared to standard intravenous fat emulsion on the IL-1b ,IL-8 levels and plasma fatty acid composition in infants who undergo gastrointestinal surgery has yet to be elucidated.

The purpose of this study is aimed to investigate the effect of ω -3-enriched intravenous fat emulsion compared to standard intravenous fat emulsion on the IL-1b ,IL-8 levels and plasma fatty acid composition in infants who undergo gastrointestinal surgery

Trial Design: Type: Parallel randomized controlled trial Allocation ratio:1:1 Framework: Superiority

Aim of the study:

The purpose of this study is to investigate the effect of ω -3-enriched intravenous fat emulsion compared to standard MCT/LCT intravenous fat emulsion on the IL-1b ,IL-8 levels and plasma fatty acid composition in infants who undergo gastrointestinal surgery.

PICO approach:

- P: Patients underwent gastrointestinal surgery.
- I: $\omega\mbox{-}3\mbox{-}enriched$ intravenous fat emulsion
- C: MCT/LCT standard intravenous fat emulsion

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	Outc ome meas ure	Tool for meas urem ent	Unit of meas urem ent
Prima ry	Infla mmat ory respo nse IL-1	Blood	pg/ml
	Infla mmat ory respo nse IL-8	Blood	Pg/ml
Seco ndary	Fatty Acid comp ositio n	Blood	(%of Total fatty acids)

 $S {\rightarrow} \text{ Randomized control study}$

Research Question

Will the Intravenous omega-3 enriched-fat emulsion make difference in IL-1β, IL-8 and fatty acid composition when compared to standard intravenous MCT/LCT fat emulsion?

Materials

MATERIALS

X Interleukin-1beta R&D Systems

X Interleukin-8 R&D Systems

Primary Outcome (p): IL-1β, IL-8

IL-1β, IL-8 levels will be measured by Quantikine HS ELISA R& D System before surgery and 3 days after surgery.

Secondary outcome (Fatty Acid Composition)

The profiling of fatty acid composition was analyzed using gas chromatography tandem mass spectrometry (GC-MSMS) before surgery and 3 days after surgery.

Safety warnings

Allergy reaction: rash

Before start

A-The experimental group (ω -3-enriched intravenous fat emulsion)

1-The parents or the legal guardian of patients who fulfill the inclusion and exclusion criteria will have to sign informed consent after the investigator explains the study.

2-Before surgery, blood samples of patients will be drawn (3-4 cc) in order to examine their IL-1β, IL-8 levels and plasma fatty acid composition.

3-After surgery, patients will get ω-3-enriched intravenous fat emulsion (SMOFlipid) for three consecutive days in 1-4 gram/kilogram/day dosing.

4-On day three after surgery, blood samples of patients will be drawn (3-4 cc) in order to examine their IL-1β, IL-8 levels and plasma fatty acid composition post-treatment.

5-All of blood samples will be sent to Prodia laboratory.

B-The control group (MCT/LCT standard intravenous fat emulsion)

1- The parents or the legal guardian of patients who fulfill the inclusion and exclusion criteria will have to sign informed consent after the investigator explains the study.

2- Before surgery, blood samples of patients will be drawn (3-4 cc) in order to examine their IL-1 β , IL-8 levels and plasma fatty acid composition.

3- After surgery, patients will get ω -3-enriched intravenous fat emulsion

(SMOFlipid) for three consecutive days in 1-4 gram/kilogram/day dosing.

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