

# Compatibility Reference Guide

## TESTING METHODS

- Admixture stability with SMOf lipid was tested by visual inspection, pH, large-diameter lipid globule size distribution (PFAT5), and mean lipid droplet diameter in compliance with USP <729> standards. No microbiological or chemical tests were conducted.
- Results are only valid for the branded products listed at the time of testing.

SMOf lipid® Lipid Injectable Emulsion, USP 20%		Aminosyn® II 15% Amino Acid Injection	Plenaminate™ 15% Amino Acids Injection	Clinisol 15% Amino Acid Injection	ProSol 20% Amino Acid Injection	Travasol 10% Amino Acid Injection
Macronutrients <sup>a</sup>	Amino Acid Solution (g/L)	32-91	32-91	33-91	35-111	28-67
	SMOf lipid 20% (g/L)	12-67	12-67	12-67	14-74	9-56
	Dextrose 70% (g/L)	88-342	88-342	88-343	103-371	68- 297
Trace Elements & Adult Multivitamins	Infuvite Adult Injection (Baxter)	0 to 10 mL	0 to 10 mL	0 to 10 mL	0 to 10 mL	0 to 10 mL
	Addamel N <sup>b</sup> (Fresenius Kabi)	0 to 10 mL	0 to 10 mL	0 to 10 mL	0 to 10 mL	0 to 10 mL
Electrolytes <sup>c</sup>	Sodium	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L
	Potassium	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L
	Calcium	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L
	Magnesium	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L	0 to 15 mEq/L
	Chloride	0 to 300 mEq/L	0 to 300 mEq/L	0 to 300 mEq/L	0 to 300 mEq/L	0 to 300 mEq/L
	Acetate	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L	0 to 150 mEq/L
	(Inorganic) Phosphate <sup>d</sup>	0 to 15 mmol/L	0 to 15 mmol/L	0 to 15 mmol/L	0 to 15 mmol/L	0 to 15 mmol/L
Extended Stability		9 days total storage: 7 days storage at 2°-8°C, then 2 days at 20°-25°C.	9 days total storage: 7 days storage at 2°-8°C, then 2 days at 20°-25°C.	8 days total storage: 7 days storage at 2°-8°C, then 1 day at 20°-25°C.	8 days total storage: 7 days storage at 2°-8°C, then 1 day at 20°-25°C.	8 days total storage: 7 days storage at 2°-8°C, then 1 day at 20°-25°C.

<sup>a</sup>The range does not necessarily reflect the minimum concentration of each macronutrient to ensure stability.

<sup>b</sup>Addamel N (for adults) is not FDA-approved for use in the U.S. Per 10 mL, it contains: iron (1.1 mg), zinc (6.5 mg), copper (1.26 mg), chromium (0.01 mg), fluoride (0.95 mg), and molybdenum (0.019 mg).

<sup>c</sup>The electrolyte salts used were sodium chloride, potassium acetate, calcium chloride, magnesium sulfate, and sodium phosphates. Any significant change in additions to the admixture compared to what has been evaluated in this study may affect stability/compatibility.

<sup>d</sup>The same limits are valid when additions of organic phosphate, sodium glycerophosphate (Glycophos) are used. Glycophos is not FDA-approved for use in the U.S.

## ADDITIVES & STORAGE

- Additions to the PN admixtures should be evaluated by a pharmacist for compatibility. If it is deemed advisable to introduce additives, use strict aseptic techniques to avoid microbial contamination.
- Infuse admixtures containing SMOf lipid immediately. If not used immediately, store admixtures under refrigeration at 2°C to 8°C (36°F to 46°F) for no longer than 24 hours. Infusion must be complete within 24 hours after removal from refrigeration. Discard any remaining admixture.

**If you have any questions, please contact medical information at 1.800.551.7176 (option 4) or email [Nutrition.MedInfo.USA@fresenius-kabi.com](mailto:Nutrition.MedInfo.USA@fresenius-kabi.com).**

## SMOFLIPID (lipid injectable emulsion), for intravenous use

### BRIEF SUMMARY OF PRESCRIBING INFORMATION

This brief summary does not include all the information needed to use SMOFLIPID safely and effectively. Please see full prescribing information for intravenous use at [www.freseniuskabinutrition.com](http://www.freseniuskabinutrition.com).

### INDICATIONS AND USAGE

SMOFLIPID is indicated in adult and pediatric patients, including term and preterm neonates, as a source of calories and essential fatty acids for parenteral nutrition (PN) when oral or enteral nutrition is not possible, insufficient, or contraindicated.

### DOSAGE AND ADMINISTRATION

The recommended daily dosage in adults is 1 to 2 grams/kg per day and should not exceed 2.5 grams/kg per day. Pediatric dosage is in Table 1, and do not exceed an infusion rate of 0.15 g/kg/hour.

SMOFLIPID 1000 mL is supplied as a Pharmacy Bulk Package for admixing only and is not for direct infusion. Prior to administration, transfer to a separate PN container for individual patient use. Use a non-vented, non-DEHP 1.2 micron in-line filter during administration. Protect the admixed PN solution from light.

Table 1: Recommended Pediatric Dosage

Pediatric Age group	Initial Dose	Maximum Dose	Duration of infusion
Birth to 2 years of age (including preterm and term neonates*)	0.5 to 1 g/kg/day Increase the dose by 0.5 to 1 g/kg/day	3 g/kg/day	20 to 24 hours for preterm and term neonates 12 to 24 hours for patients 1 month to 2 years
2 to <12 years of age	1 to 2 g/kg/day Increase the dose by 0.5 to 1 g/kg/day	3 g/kg/day	12 to 24 hours
12 to 17 years of age	1 to 2 g/kg/day	2.5 g/kg/day	12 to 24 hours

\* The neonatal period is defined as including term, post-term, and preterm newborn infants. The neonatal period for term and post-term infants is the day of birth plus 27 days. For preterm infants, the neonatal period is defined as the day of birth through the expected age of delivery plus 27 days (i.e., 44 weeks post-menstrual age).

### CONTRAINDICATIONS

- Known hypersensitivity to fish, egg, soybean, or peanut protein, or to any of the active ingredients or inactive ingredients in SMOFLIPID.
- Severe disorders of lipid metabolism characterized by hypertriglyceridemia (serum triglycerides > 1,000 mg/dL).

### WARNINGS AND PRECAUTIONS

Risk of Parenteral Nutrition-Associated Liver Disease (PNALD) and Other Hepatobiliary Disorders: PNALD, or Intestinal failure associated liver disease (IFALD) can present as cholestasis or hepatic stenosis, and may progress to steatohepatitis with fibrosis and cirrhosis (possibly leading to chronic hepatic failure). The etiology of PNALD is multifactorial; however, intravenously administered phytosterols (plant sterols) contained in plant-derived lipid emulsions, including SMOFLIPID, have been associated with development of PNALD.

In a randomized study of neonates and infants expected to be treated with PN for at least 28 days, parenteral nutrition-associated cholestasis (PNAC), a precursor to PNALD, developed less frequently in SMOFLIPID-treated patients than in 100% soybean oil lipid emulsion-treated patients.

Monitor liver tests in patients treated with SMOFLIPID and consider discontinuation or dosage reduction if abnormalities occur.

#### Other Hepatobiliary Disorders

Hepatobiliary disorders including cholecystitis and cholelithiasis have developed in some parenteral nutrition-treated patients without preexisting liver disease. Monitor liver tests when administering SMOFLIPID. Patients developing signs of hepatobiliary disorders should be assessed early to determine whether these conditions are related to SMOFLIPID use.

- Death in Preterm Neonates: Deaths in preterm neonates after infusion of lipid injectable emulsions containing only soybean oil have been reported in the medical literature. Autopsy findings in these preterm neonates included intravascular lipid accumulation in the lungs. Preterm and small-for-gestational-age neonates have poor clearance of intravenous lipid emulsion and increased free fatty acid plasma levels following lipid emulsion infusion. This risk due to poor lipid clearance should be considered when administering intravenous lipid emulsions. Monitor patients receiving SMOFLIPID for signs and symptoms of pleural or pericardial effusion.
- Hypersensitivity Reactions: SMOFLIPID contains soybean oil, fish oil, and egg phospholipids, which may cause hypersensitivity reactions. Cross reactions have been observed between soybean and peanut. SMOFLIPID is contraindicated in patients with known hypersensitivity to fish, egg, soybean, peanut protein, or to any of the active or inactive ingredients in SMOFLIPID. If a hypersensitivity reaction occurs, stop infusion of SMOFLIPID immediately and initiate appropriate treatment and supportive measures.
- Infections: Lipid emulsions, such as SMOFLIPID, can support microbial growth and are an independent risk factor for the development of catheter-related bloodstream infections. To decrease the risk of infectious complications, ensure aseptic techniques are used for catheter placement, catheter maintenance, and preparation and administration of SMOFLIPID. Monitor for signs and symptoms of infection including fever and chills, as well as laboratory test results that might indicate infection (including leukocytosis and hyperglycemia). Perform frequent checks of the intravenous catheter insertion site for edema, redness, and discharge.
- Fat Overload Syndrome: This is a rare condition that has been reported with intravenous lipid emulsions, and is characterized by a sudden deterioration in the patient's condition (e.g., fever, anemia, leukopenia, thrombocytopenia, coagulation disorders, hyperlipidemia, hepatomegaly, deteriorating liver function, and central nervous system manifestations such as coma). A reduced or limited ability to metabolize lipids, accompanied by prolonged plasma clearance (resulting in higher lipid levels), may result in this syndrome. Although fat overload syndrome has been most frequently observed when the recommended lipid dose or infusion rate was exceeded, cases have also been described when the lipid formulation was administered according to instructions.  
If signs or symptoms of fat overload syndrome occur, stop SMOFLIPID. The syndrome is usually reversible when the infusion of the lipid emulsion is stopped.
- Refeeding Syndrome: Administering PN to severely malnourished patients may result in refeeding syndrome, which is characterized by the intracellular shift of potassium, phosphorus, and magnesium as patients become anabolic. Thiamine deficiency and fluid retention may also develop. To prevent these complications, closely monitor severely malnourished patients and slowly increase their nutrient intake.
- Hypertriglyceridemia: The use of SMOFLIPID is contraindicated in patients with hypertriglyceridemia with serum triglyceride concentrations >1,000 mg/dL.  
Patients with conditions such as inherited lipid disorders, obesity, diabetes mellitus, or metabolic syndromes have a higher risk of developing hypertriglyceridemia with the use of SMOFLIPID. In addition, patients with hypertriglyceridemia may have worsening of their hypertriglyceridemia with administration of SMOFLIPID. Excessive dextrose administration may further increase such risk.

Evaluate patients' capacity to metabolize and eliminate the infused lipid emulsion by measuring serum triglycerides before the start of infusion (baseline value) and regularly throughout treatment. If triglyceride levels are above 400 mg/dL in adults, stop the SMOFLIPID infusion and monitor serum triglyceride levels to avoid clinical consequences of hypertriglyceridemia such as pancreatitis. In pediatric patients with hypertriglyceridemia, lower triglyceride levels (i.e., below 400 mg/dL) may be associated with adverse reactions. Monitor serum triglyceride levels to avoid potential complications with hypertriglyceridemia such as pancreatitis, lipid pneumonia, and neurologic changes, including kernicterus.

To minimize the risk of new or worsening of hypertriglyceridemia, assess high-risk patients for their overall energy intake including other sources of lipids and dextrose, as well as concomitant drugs that may affect lipid and dextrose metabolism.

- Aluminum Toxicity: SMOFLIPID contains no more than 25 mcg/L of aluminum. Prolonged PN administration in patients with renal impairment may result in aluminum reaching toxic levels. Preterm infants are at greater risk because their kidneys are immature, and they require large amounts of calcium and phosphate solutions, which contain aluminum. Patients with impaired kidney function, including preterm infants, who receive parenteral levels of aluminum at greater than 4 to 5 mcg/kg/day can accumulate aluminum at levels associated with central nervous system and bone toxicity. Tissue loading may occur at even lower rates of administration.
- Essential Fatty Acid Deficiency: Treatment-emergent cases of moderate or severe essential fatty acid deficiency (EFAD) (defined as the triene [Mead acid] to tetraene [arachidonic acid] ratio >0.2 and >0.4, respectively) were not observed in pediatric clinical trials of SMOFLIPID up to 28 days. However, cases of EFAD have been reported in adults and pediatric patients in the postmarketing period with the use of SMOFLIPID. The median time to onset was greater than 28 days among cases that reported time to onset. Monitor patients for laboratory evidence (e.g., abnormal fatty acid levels) and clinical symptoms of EFAD (e.g., skin manifestations and poor growth) because these signs may emerge before laboratory evidence of EFAD is confirmed. Laboratory testing using the triene to tetraene ratio may not be adequate to diagnose EFAD, and assessment of individual fatty acid levels may be needed. Ensure patients are receiving recommended dosages of SMOFLIPID to prevent EFAD.
- Monitoring/Laboratory Tests: Throughout treatment monitor serum triglycerides, fluid and electrolyte status, blood glucose, liver and kidney function, coagulation parameters, and complete blood count including platelets.

The lipids contained in SMOFLIPID may interfere with some laboratory blood tests (e.g., hemoglobin, lactate dehydrogenase [LDH], bilirubin, and oxygen saturation) if blood is sampled before lipids have cleared from the bloodstream. Conduct these blood tests at least 6 hours after stopping the infusion. SMOFLIPID contains vitamin K that may counteract anticoagulant activity.

### ADVERSE REACTIONS

Most common adverse drug reactions >1% of adult patients who received SMOFLIPID from clinical trials were nausea, vomiting, hyperglycemia, flatulence, pyrexia, abdominal pain, increased blood triglycerides, hypertension, sepsis, dyspepsia, urinary tract infection, anemia and device-related infection.

Less common adverse drug reactions in ≤ 1% of adult patients who received SMOFLIPID were dyspnea, leukocytosis, diarrhea, pneumonia, cholestasis, dyspepsia, increased blood alkaline phosphatase, increased gamma-glutamyltransferase, increased C-reactive protein, tachycardia, liver function test abnormalities, headache, pruritis, dizziness, rash and thrombophlebitis.

The most common adverse drug reactions in >1% of pediatric patients who received SMOFLIPID anemia, vomiting, gamma-glutamyltransferase increased, nosocomial infection, cholestasis, pyrexia, C-reactive protein increased, hyperbilirubinemia, abdominal pain, bilirubin conjugated increase, diarrhea, tachycardia, thrombocytopenia, hyperglycemia, sepsis.

Less common adverse reactions in ≤ 1% of pediatric patients who received SMOFLIPID were decreased hematocrit, metabolic acidosis, increased blood triglycerides, infection, increased blood alkaline phosphatase, increased alanine aminotransferase, fluid overload, hypertension, hypertriglyceridemia, and rash.

The following adverse reactions have been identified during post-approval use of SMOFLIPID in countries where it is registered. Cardiac disorders: palpitations; General disorders and administration site conditions: chills, chest pain, malaise; Hepatobiliary disorders: cholestasis; Infections and Infestations: infection; Metabolism and nutrition disorders: fatty acid deficiency; Respiratory, Thoracic and Mediastinal Disorders: dyspnea; Skin and subcutaneous tissue disorders: hyperhidrosis; Vascular disorders: phlebitis.

To report SUSPECTED ADVERSE REACTIONS, contact Fresenius Kabi USA, LLC at 1-800-551-7176, option 5, or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).

### DRUG INTERACTIONS

Soybean and olive oils in SMOFLIPID contain vitamin K, which may counteract the anticoagulant activity of vitamin K antagonists such as warfarin. In patients who receive concomitant SMOFLIPID and warfarin, increase monitoring of laboratory parameters for anticoagulant activity.

### USE IN SPECIFIC POPULATIONS

- Pregnancy and Lactation: Administration of the recommended dose of SMOFLIPID is not expected to cause major birth defects, miscarriage, or other adverse maternal or fetal outcomes. No animal reproduction studies have been conducted with SMOFLIPID. Administration of the recommended dose of SMOFLIPID is not expected to cause harm to a breastfed infant. There are no data on the presence of SMOFLIPID in human or animal milk or its effects on milk production.
- Pediatric Use: The safety and effectiveness of SMOFLIPID have been established as a source of calories and essential fatty acids for parenteral nutrition when oral or enteral nutrition is not possible, insufficient, or contraindicated in pediatric patients, including term and preterm neonates. Use of SMOFLIPID in neonates is supported by evidence from short-term (i.e., 1- to 4- week) studies, and one study following neonates beyond 4 weeks [see *Clinical Studies* (14.2)]. Use of SMOFLIPID in older pediatric patients is supported by evidence from a short-term (i.e., <28 days) study in pediatric patients 28 days to 12 years of age and additional evidence from studies in adults [see *Clinical Studies* (14)]. The most common adverse reactions in SMOFLIPID-treated pediatric patients were anemia, vomiting, gamma-glutamyltransferase increased, and nosocomial infection [see *Adverse Reactions* (6.1)]. PNALD, also referred to as IFALD, has been reported in pediatric patients who received SMOFLIPID for more than 2 weeks. PNAC (a precursor to PNALD) was reported less frequently in SMOFLIPID-treated patients compared to soybean oil lipid emulsion-treated patients in Pediatric Study 1 [see *Warnings and Precautions* (5.1) and *Adverse Reactions* (6.1)]. Although clinically significant cases of EFAD were not observed during short-term use in pediatric clinical studies, cases of EFAD have been reported with the use of SMOFLIPID in the postmarketing setting [see *Warnings and Precautions* (5.9), *Adverse Reactions* (6.1)]. Monitor pediatric patients for laboratory evidence of EFAD because they may be particularly vulnerable to neurologic complications if adequate amounts of essential fatty acids are not provided [see *Warnings and Precautions* (5.9)]. Deaths in preterm infants after infusion of lipid injectable emulsions containing only soybean oil have been reported in medical literature [see *Warnings and Precautions* (5.2)]. Because of immature renal function, preterm infants receiving prolonged treatment with SMOFLIPID may be at risk for aluminum toxicity [see *Warnings and Precautions* (5.8)].

### OVERDOSAGE

In the event of an overdose, fat overload syndrome may occur. Stop the SMOFLIPID infusion until triglyceride levels have normalized and symptoms have abated. The effects are usually reversible by stopping the lipid infusion. If medically appropriate, further intervention may be indicated. Lipids are not dialyzable from plasma.