## ANNUAL REVIEW

## Y-Site Compatibility of Medications with Parenteral Nutrition

Christine A. Robinson, PharmD<sup>1</sup> and Jaclyn E. Sawyer, PharmD<sup>2</sup>

<sup>1</sup>Clinical Associate Professor, Department of Pharmacy Practice and Administration, Ernest Mario School of Pharmacy, Rutgers University, Piscataway, New Jersey and Department of Pharmacy, Morristown Memorial Hospital, Morristown, New Jersey; <sup>2</sup>Pharmacy Clinical Specialist, Cardiology/CICU Cincinnati Children's Hospital Medical Center, Cincinnati, Ohio

KEYWORDS medication, compatibility, parenteral nutrition, y-site

J Pediatr Pharmacol Ther 2009;14:48-56

The most discussed change in parenteral nutrition compatibility within the last two years was with ceftriaxone. In the summer of 2007, Roche Laboratories updated their prescribing information for Rocephin® (ceftriaxone sodium) to include a contraindication for the co-administration of ceftriaxone with calcium-containing intravenous solutions in neonates due to reported fatal cases of pulmonary and renal precipitates in this patient population.1 The additional warnings to avoid a potential interaction are the most controversial. It is now recommended that ceftriaxone and calcium-containing intravenous solutions, such as parenteral nutrition, not be administered within 48 hours of each other regardless of patient age or remote administration sites. Institutions have struggled with the theoretical expansion of this interaction to older children and adults. At this time, no published case reports could be found in the literature.

Obtaining and maintaining venous access in pediatric patients can be complicated. Many patients require multiple treatment modalities to be administered intravenously including medications, fluids, blood products and nutrition. Clinicians must optimize available access

to ensure appropriate and timely administration of all products prior to establishing additional access. Separate administration of intravenous products, if feasible, is always preferred, however, many times simultaneous administration of medications and parenteral nutrition will be required making compatibility considerations essential. It is important to recognize that compatibility only reflects the physical interactions such as formation of a precipitate and does not necessarily address stability or pharmacologic activity of the products. Published data may report both compatibility and stability, however most evaluate visual compatibility alone. Currently there are multiple resources to use when answering the question of compatibility with parenteral nutrition. We strove to evaluate and present the available published data as a comprehensive and practical reference. Primary literature regarding y-site compatibility of medications with three different parenteral nutrition formulas, 3-in-1, 2-in-1 and lipids alone was reviewed. When conflicting results were encountered the clinical strength was considered. When published data was not accessible Trissel's Handbook on Injectable Drugs<sup>2</sup> was used.

Address correspondence to: Christine A. Robinson, PharmD, Department of Pharmacy, Morristown Memorial Hospital, 100 Madison Ave, Morristown, NJ 07962, email: crobins1@rci.rutgers.edu

© 2009 Pediatric Pharmacy Advocacy Group

## **ABBREVIATIONS**

- C Compatibility has been demonstrated. When Y-site compatibility was not available, medications compatible in solution for 24 hours were assumed to be Y-site compatible. Medications compatible with 3-in-1 admixtures were assumed to be compatible with lipids alone.<sup>2</sup>
- I Incompatibility has been demonstrated
- Compatibility data not available
- **C/I** Conflicting compatibility has been demonstrated and strength of the evidence supports compatible
- I/C Conflicting compatibility has been demonstrated and strength of the evidence supports incompatible

Medication	Admixture Type				References	
	2-in-1	lipids	3-in-1	Comments	С	ı
Acetazolamide	ı	_	_	White precipitate forms immediately		3
Acyclovir sodium	I	ı	ı	White precipitate forms immediately		3,4,5
Albumin	c	I	I	Although albumin appears visually compatible with 2:1 solutions, the potential of increased growth of fungi and bacteria warrants separate administration; visually apparent emulsion disruption	2,6	7
Aldesleukin	C	C	_		8	
Alprostadil	C	_	_		9	
Amikacin sulfate	c	C/I	C/I	Visual breaking of emulsion within 1 hour in select formulations	3,4,5,10,11,12	13
Aminophylline	C/I	C	C		4,5,14,15,16	3
Amphotericin B	1	ı	ı	Yellow precipitate forms immediately		4,5
Ampicillin sodium	C/I	c	c		4,5,17	3,10,12, 18
Ampicillin sodium - Sulbactam sodium	c	c	c		4,5	
Argatroban	c	_	_		19	
Ascorbic acid	C	_	_		20	
Atracurium besylate	c	_	_		20	
Aztreonam	C	c	C		4,5	
Bumetanide	c	c	c		4,5	
Buprenorphine HCI	C	C	C		4,5	
Butorphanol tartrate	c	C	c		4,5	
Caffeine citrate	C	_	_		21	
Carboplatin	c	C	c		4,5	
Cefamandole nafate	C	c	C		10,12,17	



Cefazolin sodium	C/I	c	c	Incompatible at a dextrose concentration of 25%	4,5,10,17,21	4
Cefepime HCI	C	_	_		22	
Cefoperazone sodium	C	C	C		4,5,10	
Cefotaxime sodium	C	C	C		3,4,5,10	
Cefotetan disodium	C	c	C		4,5	
Cefoxitin sodium	C	C	C		4,5,10,17	
Ceftazidime sodium	C	c	C		3,4,5,23	
Ceftizoxime sodium	C	C	C		4,5	
Ceftriaxone sodium	C/I	c	C/I	Per manufacturing labeling, ceftriaxone should not be administered with calcium- containing IV solutions within 48 hours of each other	4,5,20	1
Cefuroxime sodium	C	C	C		4,5	
Cephalothin sodium	C	_	_		10,12,18	
Chloramphenicol sodium succinate	С	c	_		2,10	
Chlorpromazine HCI	C	c	c		4,5	
Cimetidine HCI	C	C	C		4,5,24	
Ciprofloxacin lactate	I	c	c	Amber discoloration in 1 to 4 hours	5	4
Cisplatin	ı	c	c	Amber discoloration in 1 to 4 hours	5	4
Clindamycin phosphate	C	C	c		4,5,10,17	
Cyclophosphamide	C	C	C		4,5	
Cyclosporine	C/I	C/I	C/I	For 2:1, found to be compatible with Dextrose 5%/ Amino Acids 4.25%, but not compatible with Dextrose 25%/Amino Acid 3.5%	4,5,25	4,5
Cytarabine	ı	С	С	Substantial loss of natural turbidity occurs immediately	5	4
Dexamethasone sodium phosphate	c	c	C		3,4,5	
Diazepam	c	_	_		20	
Digoxin	C	C	C		4,5,26	
Diphenhydramine HCI	c	C	c		4,5	
Dobutamine HCI	C	C	c		3,4,5,11	
Dopamine HCl	C	C/I	C/I		3,4,5,26	5
Doxorubicin HCl	ı	ı	ı	Substantial loss of natural turbidity occurs immediately; emulsion disruption occurs immediately		4,5
Doxycycline hyclate	c	I	ı	Emulsion disruption occurs immediately	4,10	5

Droperidol	c	I	ı	Emulsion disruption occurs in 1 to 4 hours	4	5
Enalaprilat	c	C	c		4,5	
Epinephrine HCI	c		_		20	
Epoetin alfa	C	_	_		27	
Erythromycin lactobionate	C	C	c		10,17,20	
Famotidine	c	c	c		4,5,24,28, 29,30,31,32	
Fentanyl citrate	C	C	C		3,4,5,33	
Fluconazole	C	c	C		4,5,34	
Fluorouracil	ı	C/I	C/I	Slight haze, small crystals and amber discoloration form in 1 to 4 hours; turbidity forms immediately; very small amount of white precipitate forms immediately in select 3:1 formulations	5	4,5
Folic acid	C	_	_		20,35,36,37	
Foscarnet	C	_	_		38	
Furosemide	C/I	c	c	Small amount of precipitate forms in 4 hours in select formulations	3,5,20,26	4
Gallium nitrate	c	c	C		4,5	
Ganciclovir sodium	I/C	I	ı	Concentrations of ≥ 10mg/mL result in precipitation within 0 to 30 mins	39,40	4,5,39
Gentamicin sulfate	c	c	c		3,4,5,10,11, 12,13,17,18,20	
Granisetron HCI	c	C	c		4,5	
Haloperidol lactate	c	ı	1	Emulsion disruption occurs immediately	4,20	5
Heparin sodium	c	ı	ı	Emulsion disruption occurs immediately with heparin 100 units/mL	4,20	5
Hydrochloric acid	C	_	_		41	
Hydrocortisone sodium / phosphate / succinate	c	c	c		4,5,20	
Hydromorphone HCI	c	I/C	I/C	Emulsion disruption occurs immediately in select formulations	4,5	5
Hydroxyzine HCl	C	C	C		4,5	
Ibuprofen lysine	ı	_	_			42
Idarubicin HCI	C	_	_		43	
Ifosfamide	C	C	C		4,5	
Imipenem-Cilastatin sodium	c	c	c		4,5	



lmmune Globulin	—/C	_	_	Only supportive of Gammagard* 2.5%; not recommended to infuse with other drugs or solutions	44	
Indomethacin sodium trihydrate	ı	_	_			45
Insulin, regular human	C	C	c		4,5,20	
Iron dextran	C/I	_	I/C	For 2:1, found to be compatible in solution at amino acid concentrations of 2% or greater	46,47,48	47,49
Isoproterenol HCI	c	c	c	For 2:1, compatible with dextrose 25%/amino acids 4.25% (electrolytes were not added)	26,50	
Kanamycin sulfate	C	C	C		17,18,50,51	
Leucovorin calcium	C	C	C		4,5	
Levorphanol tartrate	c	ı	ı	Emulsion disruption occurs immediately	4	5
Lidocaine HCl	c	c	c	For 2:1, compatible with dextrose 25%/amino acids 4.25% (electrolytes were not added)	26,50	
Linezolid	c	-	_	Compatible with dextrose 20%/amino acids 4.9%; electrolytes were not added	52	
Lorazepam	C	ı	I	Partial emulsion disruption occurs in 1 hour	4	5
Magnesium sulfate	C	C	C		4,5	
Mannitol	C	C	c		4,5	
Meperidine HCI	C	C	C		4,5,53	
Meropenem		C	c		5	
Mesna	C	C	C		4,5	
Methotrexate sodium	ı	C	c	For 2:1, hazy precipitate forms in 0 to 1 hour	5	4
Methyldopate HCl	c	C/I	C/I	For 2:1, compatible with dextrose 25%/amino acids 4.25% (electrolytes were not added); cracked the lipid emulsion in select formulations	26,50	26
Methylprednisolone sodium succinate	c	c	c		4,5	
Metoclopramide HCI	I/C	c	c	Substantial loss of natural turbidity occurs immediately in select formulations	2,5	4
Metronidazole	c	C	C		3,4,5,20	
Mezlocillin sodium	C	C	C		4,5	
Miconazole	C	C	c		4,5,10	

				war.		
Midazolam HCI	I/C	1	ı	White precipitate forms immediately in select formulations	54	4,5,20
Milrinone lactate	C	_	_		55,56	
Minocycline HCl	ı	ı	ı	Bright yellow discoloration forms immediately; emulsion disrupts immediately		4,5
Mitoxantrone HCI	I	c	c	Substantial loss of turbidity occurs immediately	5	4
Morphine sulfate	C	C/I	C/I	For 3:1, morphine 1 mg/mL compatible, but 15 mg/mL was not compatible; emulsion disruption occurs immediately in select formulations	3,4,5,20,53	5
Nafcillin sodium	C	C	C		4,5,10,12	
Nalbuphine HCl	c	ı	ı	Emulsion disruption occurs immediately	4	5
Netilmicin sulfate	C	C	c		4,5	
Nitroglycerin	C	C	c		4,5	
Norepinephrine bitartrate	C	C	C		4,26	
Octreotide acetate	C	C	C		4,5	
Ondansetron HCI	c	ı	ı	Emulsion disruption occurs immediately	4	5
Oxacillin sodium	C	C	C		10,12,17	
Paclitaxel	C	C	c		4,5	
Penicillin G potassium	C	C	C		3,10,12,17,20	
Penicillin G sodium	C	_	_		10,12	
Pentobarbitol sodium	c	ı	ı	Emulsion disruption occurs immediately	4	5
Phenobarbitol sodium	c	I	I	Emulsion disruption occurs immediately	4	5
Phenytoin sodium	ı	I	_	Heavy white precipitate forms immediately; incompatible with dextrose		2,20
Piperacillin sodium	C	c	c		4,5,10,12	
Piperacillin sodium / Tazobactam sodium	c	c	c		4,5	
Potassium chloride	c	c	c		4,5	
Potassium phosphate	ı	I	ı	Increased turbidity occurs immediately; emulsion disruption occurs immediately		4,5
Prochlorperazine edisylate	c	c	c		4,5	
Promethazine HCI	C/I	c	c	Amber discoloration in 4 hours in select formulations	4,5	4
Propofol	c	_	_	Propofol injection contains approximately 10 gm fat / 100 mL	57	

Ranitidine HCl	c	c	C		3,4,5,20,24	
Sargramostim	c	_	_		58	
Sodium bicarbonate	I/C	c	c	Small amount of precipitate forms in 1 hour in select formulations	4,5	4
Sodium nitroprusside	C	C	c		4,5	
Sodium phosphate	ı	ı	I	Increased turbidity occurs immediately; emulsion disruption occurs immediately		4,5
Tacrolimus	C	C	C		4,5	
Ticarcillin disodium	C	C	C		4,5,10,12,17	
Ticarcillin disodium- Clavulanate potassium	c	c	c		4,5,20	
Tobramycin sulfate	c	c	c		3,4,5,10, 11,12,13,17	
Trimethoprim- Sulfamethoxazole	c	c	c		4,5	
Urokinase	C	_	_		2	
Vancomycin HCI	c	c	c		3,4,5,10,11,20	
Vecuronium bromide	C	_	_		20	
vitamin K1 - phytonadione	c	c	_		18,59	
Zidovudine	C	C	C		3,4,5	

## REFERENCES

- 1. Rocephin [package insert]. Nutley, NJ: Roche Laboratories Inc; August 2008.
- 2. Trissel LA. Handbook on Injectable Drugs, 14<sup>th</sup> ed, Bethesda, MD: American Society of Health-System Pharmacists, Inc; 2007.
- 3. Veltri M, Lee CKK. Compatibility of neonatal parenteral nutrient solutions with selected intravenous drugs. Am J Health-Syst Pharm 1996;53:2611-2613.
- 4. Trissel LA, Gilbert DL, Martinez JF, et al. Compatibility of parenteral nutrient solutions with selected drugs during simulated Y-site administration. Am J Health-Syst Pharm 1997;54:1295-1300.
- 5. Trissel LA, Gilbert DL, Martinez JF, et al. Compatibility of medications with 3-in-1 parenteral nutrition admixtures. J Parenter Enteral Nut 1999;23:67-74.
- Mirtallo JM, Caryer K, Schneider PJ, et al. Growth of bacteria and fungi in parenteral nutrition solutions containing albumin. Am J Hosp Pharm 1981;38:1907-1910.

- 7. Ambados F. Destabilization of fat emulsion in total nutrient admixtures by concentrated albumin 20% infusion. Aust J Hosp Pharm 1999;29:210-212.
- 8. Anderson PM, Rogosheske JR, Ramsey NKC, et al. Biological activity of recombinant interleukin-2 in intravenous admixtures containing antibiotic, morphine sulfate or total parenteral nutrition solution. Am J Hosp Pharm 1992;49:608-612.
- 9. Dice JE. Physical compatibility of alprostadil with commonly used IV solutions and medications in the neonatal intensive care unit. J Pediatr Pharmacol Ther 2006;11:233-236.
- 10. Watson D. Piggyback compatibility of antibiotics with pediatric parenteral nutrition solutions. J Parenter Enteral Nutr 1985;9:220-224.
- 11. Schilling CG. Compatibility of drugs with a heparin-containing neonatal total parenteral nutrient solution. Am J Hosp Pharm 1988;45:313-314.

- 12. Kamen BA, Gunther N, Sowinsky N, et al. Analysis of antibiotic stability in a parenteral nutrition solution. Pediatr Infect Dis 1985;4:387-389.
- 13. Bullock L, Clark JH, Fitzgerald JF, et al. The stability of amikacin, gentamicin, and tobramycin in total nutrient admixtures. J Parenter Enteral Nutr 1989;13:505-509.
- 14. Andreu A, Cardona D, Pastor C, et al. Intravenous aminophylline: in vitro stability of fat-containing TPN. Ann Pharmacother 1992;26:127-128.
- Niemiec PW Jr, Vanderveen TW, Hohenwarter MW, et al. Stability of aminophylline injection in three parenteral nutrient solutions. Am J Hosp Pharm 1983;40: 428-432.
- Sykes R, McPherson C, Foulks K, et al. Aminophylline compatibility with neonatal total parenteral nutrition. J Pediatr Pharmacol Ther 2008;13:76-79.
- 17. Baptista RJ, Lawrence RW. Compatibility of total nutrient admixtures and secondary antibiotic infusions. Am J Hosp Pharm 1985;42:362-363.
- 18. Schuetz DH, King JC. Compatibility and stability of electrolytes, vitamins and antibiotics in combination with 8% amino acids solutions. Am J Hosp Pharm 1978;35:33-44.
- 19. Honisko ME, Fink JM, Militello MA, et al. Compatibility of argatroban with selected cardiovascular agents. Am J Health-Syst Pharm 2004;61:2415-2418.
- 20. Gilbar PJ, Groves CF. Visual compatibility of total parenteral nutrition solution (Synthamin 17 premix) with selected drugs during simulated Y-site injection. Aust J Hosp Pharm 1994;24:167-170.
- 21. Nahata MC, Zingarelli J, Durrell DE. Stability of caffeine citrate injection in intravenous admixtures and parenteral nutrition solutions. J Clin Pharm Ther 1989;14:53-55.
- Maxipime [package insert]. Princeton, New Jersey: Bristol Meyers Squibb Company; December 2003.
- 23. Wade CS, Lampasona V, Mullins RE, Parks RB. Stability of ceftazidime and amino acids in parenteral nutrient solutions. Am J Hosp Pharm 1991;48:1515-1519.
- 24. Hatton J, Luer M, Hirsch J, et al. Histamine receptor antagonists and lipid stability in total nutrient admixtures. J Parenter Enteral Nutr 1994;18:308-312.

- Jacobson PA, Maksym CJ, Landvay A, Weiner N, Whitmore R. Compatibility of cyclosporine with fat emulsion. Am J Hosp Pharm 1993;50:687-690.
- Baptista RJ, Dumas GJ, Bistrian BR, et al. Compatibility of total nutrient admixtures and secondary cardiovascular medications. Am J Hosp Pharm 1985;42:777-778.
- 27. Ohls RK, Christensen RD. Stability of human recombinant epoetin alfa in commonly used neonatal intravenous solutions. Ann Pharmacother 1996;30:466-468.
- DiStefano JE, Mitrano JE, Baptista FP, et al. Long-term stability of famotidine 20 mg/ mL in a total parenteral nutrient solution. Am J Hosp Pharm 1989;46:2333-2335.
- 29. Bullock L, Fitzgerald JF, Glick MR, et al. Stability of famotidine 20 and 40 mg/L and amino acids in total parenteral nutrient solutions. Am J Hosp Pharm 1989;46:2321-2325.
- 30. Bullock L, Fitzgerald JF, Glick MR. Stability of famotidine 20 and 50 mg/L in total nutrient admixtures. Am J Hosp Pharm 1989;46:2326-2329.
- 31. Montoro JB, Pou L, Salvador P, et al. Stability of famotidine 20 and 40 mg/L in total nutrient admixtures. Am J Hosp Pharm 1989;46:2329-2332.
- Shea BF, Souney PF. Stability of famotidine in a 3-in-1 total nutrient admixture. DCIP 1990;24:232-235.
- Moshfeghi M, Ciuffo J. Visual compatibility of fentanyl citrate with parenteral nutrient solutions [Letters]. Am J Health-Sys Pharm 1998;55:1194-1197.
- 34. Couch P, Jacobson P, Johnson CE. Stability of fluconazole and amino acids in parenteral nutrient solutions. Am J Hosp Pharm 1992;49:1459-1462.
- 35. Chen MF, Boyce HW, Triplett L. Stability of the B vitamins in mixed parenteral nutrition solution. J Parenter Enteral Nutr 1983;7:462-464.
- 36. Barker A, Hebron BS, Beck PR, Ellis B. Folic acid and total parenteral nutrition. J Parenter Enteral Nutr 1984;8:3-8.
- Louie N, Stennett DJ. Stability of folic acid in 25% dextrose, 3.5% amino acids, and multivitamin solution. J Parenter Enteral Nutr 1984;8:421-426.

- 38. Baltz JK, Kennedy P, Minor JR, Gallelli J. Visual compatibility of foscarnet with other injectable drugs during simulated Y-site administration. Am J Hosp Pharm 1990;47:2075-2077.
- Outman WR, Mitrano FP, Baptista RJ. Visual compatibility of ganciclovir sodium and parenteral nutrient solution during simulated Y-site injection. Am J Hosp Pharm 1991;48:1538-1539.
- 40. Johnson CE, Jacobson PA, Chan E. Stability of ganciclovir sodium and amino acids in parenteral nutrient solutions. Am J Hosp Pharm 1994;51:503-508.
- 41. Mirtallo JM, Rogers KR, Johnson JA, et al. Stability of amino acids and the availability of acid in total parenteral nutrition solutions containing hydrochloric acid. Am J Hosp Pharm 1981;38:1729-1731.
- 42. Holt RJ, Siegert SWK, Krishna A. Physical compatibility of ibuprofen lysine injected with selected drugs during simulated Y-site injection. J Pediatr Pharmacol Ther 2008;13:156-161.
- 43. Turowski RC, Durthaler JM. Visual compatibility of idarubicin hydrochloride with selected drugs during simulated Y-site injection. Am J Hosp Pharm 1991;48:2181-2184.
- 44. Lindsay CA, Dang K, Adams JM, Ou CN, Baker CJ. Stability and activity of intravenous immunoglobulin with neonatal dextrose and total parenteral nutrient solutions. Ann Pharmacother 1994;28:1014-1017.
- 45. Ishisaka DY, VanVleet J, Marquardt E. Visual compatibility of indomethacin sodium trihydrate with drugs given to neonates by continuous infusion. Am J Hosp Pharm 1991;48:2442-2443.
- Wan KK, Tsallas G. Dilute iron dextran formulation for addition to parenteral nutrient solutions. Am J Hosp Pharm 1980;37:206-210
- 47. Mayhew SL, Quick MW. Compatibility of iron dextran with neonatal parenteral nutrient solutions. Am J Health-Syst Pharm 1997;54:570-571.
- 48. Tu YH, Knox NL, Biringer JM, et al. Compatibility of iron dextran with total nutrient admixtures. Am J Hosp Pharm 1992;49:2233-2235.

- 49. Vaughan LM, Small C, Plunkett V. Incompatibility of iron dextran and a total nutrient admixture. Am J Hosp Pharm 1990;47:1745-1746.
- 50. Athanikar N, Boyer B, Deamer R, et al. Visual compatibility of 30 additives with a parenteral nutrient solution. Am J Hosp Pharm 1979;36:511-513.
- 51. Feigin RD, Moss KS, Shackelford PG. Antibiotic stability in solutions used for intravenous nutrition and fluid therapy. Pediatrics 1973;51:1016-1026.
- Trissel LA, Williams KY, Gilbert DL. Compatibility screening of linezolid injection during simulated Y-site administration with other drugs and infusion solutions. J Am Pharm Assoc 2000;40:515-519.
- 53. Pugh CB, Pabis DJ, Rodriguez C. Visual compatibility of morphine sulfate and meperidine hydrochloride with other injectable drugs during simulated Y-site injection. Am J Hosp Pharm 1991;48:123-125.
- 54. Bhatt-Mehta V, Rosen DA, King RS, Maksym CJ. Stability of midazolam hydrochloride in parenteral nutrient solutions. Am J Hosp Pharm 1993;50:285-288.
- 55. Akkerman SR, Zhang H, Mullins RE, Yaughn K. Stability of milrinone lactate in the presence of 29 critical care drugs and 4 i.v. solutions. Am J Health-Syst Pharm 1999;56:63-68.
- 56. Veltri MA, Conner KG. Physical compatibility of milrinone lactate injection with intravenous drugs commonly used in the pediatric intensive care unit. Am J Health-Syst Pharm 2002;59:452-454.
- 57. Bhatt-Mehta V, Paglia RE, Rosen DA. Stability of propofol with parenteral nutrient solutions during simulated Y-site injection. Am J Health-Syst Pharm 1995;52:192-196.
- 58. Trissel LA, Bready BB, Kwan JW, Santiago NM. Visual compatibility of sargramostim with selected antineoplastic agents, anti-infectives, or other drugs during simulated Y-site injection. Am J Hosp Pharm 1992;49:402-406.
- Dahl GB, Svensson L, Kinnander NJ, Zander M, Bergstrom UK. Stability of multivitamins in soybean oil fat emulsion under conditions simulating intravenous feeding of neonates and children. J Parenter Enteral Nutr 1994;18:234-239.