

Liquid Dosage Forms

Gautami.j*

Department of pharmaceutics, SRM College of pharmacy, Potheri, Kattankulathur, Tamil Nadu 603203, India

***Corresponding author:** Gautami.j, Department of pharmaceutics, SRM College of pharmacy, Potheri, Kattankulathur, Tamil Nadu 603203, India, Tel: +998-712418594; E-mail: gautami_j@gmail.com

Received: July 25, 2016; Accepted: August 20, 2016; Published: August 24, 2016

Abstract

Liquid state forms are meant for internal, parental or external use. They are available in monophasic and biphasic forms. Monophasic liquid dosage forms are true or colloidal solution. Water is mainly used as a solvent for majority of monophasic liquid dosage forms. The liquid which consists of two phases are known as biphasic liquids.

Keywords: Liquid dosage forms, Suspensions, Emulsions, Ointments, Internal use, External use

Introduction

Liquid form of a dose of a drug used as a drug or medication intended for administration or consumption [1-15]. Advantages

- Faster absorption than solids
- Palatable pleasant to taste
- Best choice for children and old persons

Monophasic liquid dosage forms: It contains only one phase.

Classification [16-30]

A. Liquids for internal use: Drops, Elixirs, Linctus, Syrups, and draughts.

B. Liquids for external use:

Liquid to be applied to the skin: Liniments and lotions.

Liquids meant for body cavity: Gargles, throat paints, mouth washes, throat paints, mouth washes, eye drops, eye lotions, ear drops, nasal drops, sprays and inhalations

Syrups: Saturated solution of sucrose in purified water, sweet viscous preparations. Concentration of sugar is 66% (w/w). Syrups containing medicinal substances are called medicated syrups and those containing aromatic or flavored substances are known as flavored syrups.

Preparation: Add sucrose to purified water and heat it to dissolve sucrose with occasional stirring. Cool it and add more of purified water to make the required weight. Syrups used in formulation of antibiotics, saline drugs, vitamins, antitussives, sedatives [31-35].

Elixirs: Sweet aromatic colored preparations. Main Ingredients of elixir are ethyl alcohol, water, glycerin, propylene glycol, flavoring agent, syrup and preservatives. Medicated elixir contains very potent drug such as antibiotics, antihistamines, sedatives. Flavoring elixirs used as flavours and vehicles [36-38].

Linctuses: Viscous liquid and oral preparations that are generally prescribed for the relief of cough. They contain medicament which have demulcent, sedative or expectorant action. linctuses should be taken in a small doses sipped and swallowed slowly without diluting it with water in order to have maximum and prolonged effect of medications. Simple syrup is used a vehicle for most of the linctuses. Tolu syrup is preferred in certain cases because of its aromatic odour and flavor [39-41].

Drops: Liquid preparations meant for oral administration. The oil soluble vitamins such as vitamin A and D concentrations in fish liver oil are presented as drops for administration. Since these preparations contain potent medications the dose must be measured accurately [42-45].

Liniments: Liquid and semi liquid preparations meant for application to the skin. Liniments are usually applied to the skin with friction and rubbing of the skin. Liniments may be alcoholic or oily solutions or emulsions. Alcohol helps in penetration of medicament in to the skin and also increases its counterirritant or rubefacient action. Arachis oil is used in some liniments which spread more easily on the skin. Soap is also included as ingredients in some of the liniments which helps in easy application of liniment on the skin.

Liniments contain medicaments possessing analgesic, rubefacient, soothing, counter irritant or stimulating properties. Liniment should not be applied to broken skin it may cause excess irritation [46-48].

Lotions: Liquid preparations meant for external application without friction. They are applied direct to the skin with the help of some absorbent material such as cotton, wool or gauze soaked in it. Lotions may be used for local action as cooling, soothing or protective purpose. They are generally prescribed for antiseptic action ex: Calamine lotion [49-50].

Gargles: Aqueous solutions used to prevent or treat throat infections. They are usually available in concentrated for with direction for dilution with warm water before use. They are brought in to contact with mucous membrane of the throat and are allowed to remain in contact with it for a few seconds [51-53].

Mouth washes: Aqueous solutions with a pleasant taste and odour used to make clean and deodorize the buccal cavity. Generally they contain antibacterial agents, alcohol, glycerin, sweetening agents, flavouring agents and coloring agents [54-55].

Throat paints: Viscous liquid preparations used for mouth and throat infections. Glycerin is commonly used as a base it

adheres to mucous membrane for a long period and it possesses a sweet taste [56].

Nasal drops; solutions of drugs that are instilled in to the nose with a dropper. They are usually aqueous and not oily drops. Nasal drops should be isotonic having neutral pH and viscosity similar to nasal secretions by using methyl alcohol [57-60].

Ear drops: Sterile solution or suspensions of drugs that are instilled in to the eye with a dropper. The eye drops are usually made in aqueous vehicle. It should be sterile isotonic with lacrymal secretions, buffered and free from foreign particles to avoid irritation to the eye [61-64].

Eye lotions: Aqueous solutions used for washing the eyes. The eye lotions are supplied in concentrated form and are required to be diluted with warm water immediately before use. It should be isotonic and free from foreign particles to avoid irritation to the eye [65-70].

Ear drops: solutions of drugs that are instilled in to the ear with a dropper. These are generally used for cleaning the ear, softening the wax and for treating the mild infections [71-72].

Biphasic liquid dosage forms: It contains two phases [73-80].

Ex: Suspension and emulsion

Suspensions: Biphasic liquid dosage form of medicament in which finely divided solid particles are dispersed in a liquid or semisolid vehicle. The solid particles act as disperse phase whereas liquid vehicle acts as the continuous phase. Suspensions are generally taken orally or by parental route. They are also used for external application [81-85].

Many suspensions are supplied as dry powders which are converted in to suspensions by adding the specified amount of vehicle before use. This is done to ensure the stability of suspension

Ex: Ampicillin for oral suspensions, Barium sulphate suspensions, Insulin zinc suspension

Emulsion: Biphasic liquid preparation containing two immiscible liquids, one of which is dispersed as minute globules in to the other. The liquid which is converted in to minute globules is called the disperse phase and the liquid in which the globules are dispersed is called the continuous phase. Normally two immiscible liquids cannot be dispersed for a long period. So an emulsifying agent is added to the system. It forms the film around the globules in order to scatter them indefinitely in the continuous phase, So that a stable emulsion is formed [86-90].

Emulsions are of two types [91-100]

I. Oil in water type (O/W): Emulsion in which oil is I the dispersed phase whereas water is in the continuous phase. The O/W type emulsions are preferred for internal use. In these emulsions gum acacia, tragacanth, methyl cellulose, saponins synthetic substances and soaps formed from monovalent bases like sodium, potassium are used as an emulsifying agent.

II. Water in oil type (W/O): Emulsion in which water is in the dispersed phase whereas oil is in continuous phase. Wool

wax, resins, beeswax and soaps formed from divalent bases like calcium, magnesium and zinc are used as an emulsifying agent. The W/O emulsions are mainly used externally as lotions or creams.

III. Intravenous emulsion: The oil soluble hormones vitamin A,D and K are administered as intravenous injection. The emulsified oils are also injected as diagnostic aids. The emulsion should have small globule size and must be sterile.

IV. Emulsion for external use: The emulsions for external application may be both O/W or W/O type but O/W type emulsion is preferred. When a drug is emulsified its rate of penetration through the skin may get reduced. It helps to prolong the action of a drug. Generally the emulsions for application to the skin are semisolid at room temperature and are considered to be an excellent vehicle.

Conclusion:

Liquid dosage forms are formulated to release the active principle immediately after oral administration to obtain rapid and complete systemic drug absorption when compared to oral route. Liquid state forms are meant for internal, parental or external use.

REFERENCES

- 1. Peter ASA, Hymavathi TV, Yasoda DP. A Study on the Different Methods of Preparation of Lutein from Supercritical Fluid Processed Lutein Esters. J Nutr Food Sci. 2012;2:154.
- 2. Figueiras A, Cardoso O, Veiga F. Preparation and characterization of Trimethoprim inclusion complex with Methyl-β-Cyclodextrin and determination of its antimicrobial activity. Pharm Anal Acta. 2015;6:405.
- 3. Zhang L, Zhao H, Zhou G, Niu T, Yang J. Simulation Database System of the Active Ingredients in Compound Decoction of Chinese Medicine. J Bioequiv Availab.2010;2:131-34.
- Naveed S, Sajid S. Degradation in Pharmaceutical Creams: Ascorbic Acid Demonstrating Phenomenon: A Review. J Bioequiv Availab. 2016;8:080-83.
- Khokhlov AL, Shitov LN, Ryska M, et al. The Pharmacokinetic Properties and Bioequivalence of Methyldopa Formulations: Results of an Open-label, Randomized, Two-period, Crossover, Single-dose Study. J Bioequiv Availab. 2016;8:185-90.
- Delsin SD, Mercurio DG, Fossa MM, Maia Campos PMBG. Clinical Efficacy of Dermocosmetic Formulations Containing Spirulina Extract on Young and Mature Skin: Effects on the Skin Hydrolipidic Barrier and Structural Properties. Clin Pharmacol Biopharm. 2015;4:144.
- Galicia-QC, Valle LCF, Soto MH, et al. Adverse Events Reactions Reported With the Use of a Fixed-Dose Combination of Nor- Pseudoephedrine, Triiodothyronine, Atropine, Aloin and Diazepam in Obese Mexican Patients. J Pharmacovigil. 2015;3:185.
- Karwa M, Arora S, Agrawal SG. Recent Regulatory Amendment in Schedule Y: Impact on Bioequivalence Studies Conducted In India. J Bioequiv Availab. 2013;5:174-76.
- 9. Xie PS, Sun S, Xu S, Guo L. Value the Unique Merit of HPTLC Image Analysis and Extending its Performance by Digitalization for Herbal Medicines Quality Control. J Chromatograph Separat Techniq. 2014;5: 249.
- Singh A, Tandon S, Sand NK. Active Ingredient Estimation of Clopyralid Formulation by Reversed Phase HPLC. J Chromatogr Sep Tech. 2014;6:257.

- 11. James KC, Rangoonwala R, Reshetnykov M. Non-Equivalence of Antibiotic Generic Drugs and Risk for Intensive Care Patients. Pharmaceut Reg Affairs. 2013;2:109.
- 12. Iordaconiu L, Malaescu I, Chirigiu L. Polymeric Membranes: Effects of Catalyst Volume Fraction on Dielectric Relaxation Time and Crystallites Dimensions. Ind Chem. 2016;2:117.
- Gopi S, Amalraj A, Thomas S. Effective Drug Delivery System of Biopolymers Based On Nanomaterials and Hydrogels
 A Review. Drug Des. 2016;5:129.
- 14. Devi AN, Vijendar C, Goud AK, et al. Preparation and Evaluation of Floating Microspheres of Cefdinir in Treatment of Otitis Media and Respiratory Tract Infections. J Pharmacovigilance. 2016; 4: 209.
- 15. Enose AA, Dasan P, Sivaramakrishanan H. Formulation, Characterization and Pharmacokinetic Evaluation of Telmisartan Solid Dispersions. J Mol Pharm Org Process Res. 2016;4:131.
- Newton AMJ, Rani SM, Sukhjinder K. Fabrication and Evaluation of Fast Disintegrating Oral Hybrid Films of Propranolol Hydrochloride by Using Pectin and Synthetic Polymers. J Dev Drugs. 2015;5:157.
- 17. Anumolu PD, Krishna VL, Rajesh CH, et al. Gas Chromatographic Assessment of Residual Solvents Present in Excipient-Benzyl Alcohol. J Chromatogr Sep Tech. 2015;7:321.
- Mohd AB, Vemula SK. Formulation and Pharmacokinetics of Vitamin E TPGS Melt Dispersion Granules: An Approach to Improve Oral Delivery of Flurbiprofen. J Bioequiv Availab. 2016;8:089-94.
- 19. Chauhan MK, Bhatt N. A Simple and Modified Method Development of Vancomycin Using High Performance Liquid Chromatography. J Chromatogr Sep Tech. 2015;6:296.
- 20. Gajbhiye KR, Gajbhiye V, Soni V. Targeted Brain Delivery of Bioactive Molecules Using Nanocarriers. J Bioequiv Availab. 2015;7:112-22.
- Shintani H. Sterilization Validation of Gas Plasma Exposure Based on ISO Documents (Mainly ISO TC 198 And 194 Documents). Pharmaceutical regulatory affaris. 2015;4:137.
- 22. Trivedi MK, Nayak G, Patil S, Tallapragada RM, Mishra R. Influence of Biofield Treatment on Physicochemical Properties of Hydroxyethyl Cellulose and Hydroxypropyl Cellulose. J Mol Pharm Org Process Res. 2015;3:126.
- Sivasakthi M, Sangeetha N. A Comparative Study of the Physico- Chemical Characteristics of the Ready-to Eat Coconut Based Snack. J Food Process Technol. 2015;6:489.
- 24. Figueiras A, Cardoso O, Veiga F, et al. Preparation and characterization of Trimethoprim inclusion complex with Methyl-β-Cyclodextrin and determination of its antimicrobial activity. Pharm Anal Acta. 2015;6:405.
- 25. Hattab MA, Ghaly A. Microalgae Oil Extraction Pretreatment Methods: Critical Review and Comparative Analysis. J Fundam Renewable Energy Appl. 2015;5:172.
- 26. Chen Q, Bruyneel A, Clarke K, Carr C, et al. Collagen- Based Scaffolds for Potential Application of Heart Valve Tissue Engineering. J Tissue Sci Eng. 2012;S11: 003.
- 27. Dzung NT, Dzung NQ, Dzung TV, et al. Application of Multi- Objective Optimization by S and R* Optimal Combination Criteria to Determine the Freeze Drying Mode of Penaeus monodon. J Chem Eng Process Technol. 2011;2:107.
- Gopi S, Amalraj A, Thomas S. Effective Drug Delivery System of Biopolymers Based On Nanomaterials and Hydrogels
 A Review. Drug Des. 2016;5:129.
- 29. Medioni J, Tournou F C, De Y. A New Adaptive Simon- Based Design Focusing on Subpopulation Heterogeneity. Drug Des. 2016;5:128.

- Lopes CM, Soares C. Transdermal Drug Delivery Systems Activated by Physical stimuli: Techniques and Applications. Drug Des.2015;4:e129.
- White AR. The Success of Solanezumab Should Drive Renewed Efforts to Develop Small Molecule Anti-Amyloid Agents for Alzheimer's disease Therapy. Drug Des. 2015;4:e128.
- 32. Gomase VS, Kale KV. Information of Surface Accessibility of the Peptide Fragments of Coat Protein from Alfalfa mosaic virus (AMV) at the Physicochemical and Immunochemical Levels. Drug Des. 2015;4:119.
- 33. Bule MH, Haymete A, Kefale B. Synthesis and In-Vivo Pharmacological Evaluation of Some Novel 4(3H)-Quinazolinone Derivatives as Potential Anti-malarial Agents. Drug Des. 2015;4:121.
- 34. Yuh-Jenn Wu, Chi-Tian Chen, Hsiao-Hui Tsou, Chin-Fu Hsiao. Evaluating the Relative Cost of a Targeted Design versus an Untargeted Design for Randomized Clinical Trials. Drug Des. 2015;4:120.
- 35. Yuh-Jenn Wu, Chi-Tian Chen, Hsiao-Hui Tsou, Chin-Fu Hsiao. Evaluating the Relative Cost of a Targeted Design versus an Untargeted Design for Randomized Clinical Trials. Drug Des. 2015;4:120.
- 36. Chow SC, Pong A. Statistical Designs for Pharmaceutical/Clinical Development. Drug Des. 2014;3:112.
- 37. Anil Vaidya. Drug Designing and Development: Emerging Role of Health Technology Assessment. Drug Des. 2014;3:111.
- 38. Toshihiko Tashima. Multifunctionality of Drug's Pharmacophores within an Organism. Drug Des. 2014;3: 116.
- 39. Chow SC. On Assessment of Analytical Similarity in Biosimilar Studies. Drug Des. 2014;3:e124.
- 40. Lopes CM. Therapeutics Delivery: Innovations Technology Approaches. Lopes, Drug Des. 2014;3:e123.
- 41. Karaman R. The Future of Prodrugs Designed by Computational Chemistry. Drug Des. 2012;1:e103.
- 42. Chow SC, Chiu ST. A Note on Design and Analysis of Clinical Trials. Drug Des. 2013:2:102.
- 43. Nayak UY. Role of Nano-Particles in Drug Therapy-Drug Delivery Approach. Drug Design. 2013;S5: e001.
- 44. Nirmala MJ, Nagarajan R. Microemulsions as Potent Drug Delivery Systems. J Nanomed Nanotechnol. 2016;7:e139.
- 45. AbouAitah KEA, Farghali AA, Swiderska-Sroda A, Lojkowski W, Razin AMF, et al. pH-controlled Release System for Curcumin based on Functionalized Dendritic Mesoporous Silica Nanoparticles. J Nanomed Nanotechnol. 2016;7:351.
- Patil JS. Significance of Particulate Drug Delivery System in Antimicrobial Therapy. Adv Pharmacoepidemiol Drug Saf. 2016;5:139.
- 47. Dudhipala N, Narala A, Janga KY, et al. Amoxycillin Trihydrate Floating-Bioadhesive Drug Delivery System for Eradication of Helicobacter pylori: Preparation, In Vitro and Ex Vivo Evaluation. J Bioequiv Availab. 2016:8:118-124.
- 48. Naydenov T, Stoimenova A, Kassarova M, et al. Opinion of Bulgarian Pharmacists on Drug Delivery Systems, Orodispersible and Pediatric Dosage Forms. J App Pharm. 2015;8:211.
- 49. Shanmugan P, Bandameedi R. Chronotherapeutic Drug Delivery Systems. J Drug Metab Toxicol. 2015;6: 194.
- Kumar P, Bose PP. Targeted Delivery of Paromomycin to Leishmania Infected Macrophage by Hemoglobin Tagged Nanocarrier. J App Pharm. 2015;8:212.
- 51. The Pharmaceutical Codex Principles and practice of Pharmaceutics. 1994;12:189.
- 52. Coelho M. Fate of Vitamins in Premixes and Feeds: Vitamin Stability. FeedManagement. 1991; 42(10):24.
- 53. Akhtar MJ, Khan MA, Ahmad I. Photodegradation of folic acid in aqueous solution. J Pharma Biomed Anal.1999; 19(3):269-75.
- 54. ManzurUl, Haque H. Assay of Vitamins in Pharmaceutical Preparations.1972; 7(10): 213-26.
- 55. Howard CA, Loyd VA, Nicholas GP. Pharmaceutical Dosage Forms and Drug Delivery Systems. 2000; 7Edn: 38-64.

- 56. Ekinci R, Kadakal C. Determination of seven water-soluble vitamins in Tarhana, a traditional Turkish cereal food, by High-Performance Liquid Chromatography. ActaChromatographica.2005;15:289-97.
- 57. Deshpande SW, Nilesh G. Drugs and Cosmetics Act 1940 and Rules 1945. 2000; 2Edn:538-539.
- Nelson DH, Samuels LT. A Method for Determination of 17-Hydroxycorticosteroids in Blood: 17-Hydroxycorticosterone in the Peripheral Circulation. J Clin Endocrinol & Metab. 1952;12:519.
- Glenn EM, Nelson DH. Chemical Method for the Determination of 17-Hydroxycorticosteroids and 17-Ketosteroids in Urine Following Hydrolysis With β-Glucuronidase. J Clin Endocrinol & Metab. 1953;13:911.
- 60. Bayliss RES, Steinbeck AW. A Modified Method for Estimting 17-Hydroxycorticosteroids in Plasma. Biochem J. 1953;54:523.
- Burton RB, Zaffaroni A, Keutmann EH. Paper Chromatography of Steroids. II. Corticosteroids and Related Compounds. J Biol Chem. 1951;188:763.
- 62. Nelson DH, Sandberg AA, Palmer JG, Tyler FH. Blood Levels of 17-Hydroxycorticosteroids Following the Administration of Adrenal Steroids and Their Relation to Levels of Circulating Leukocytes. J Clin Invest 1952;31: 843.
- 63. Stewart PJ, Tucker IG. A survey of current extemporaneously prepared paediatric formulations". Aust J Hosp Pharm 1982;12(3):64-8.
- 64. Tan E, Cranswick NE, Rayner CR, et al. Dosing information for paediatric patients: are they really 'therapeutic orphans'? Med J Aust 2003;179(4);195–98.
- 65. Pal, VandNahata M. Need for extemporaneousformulations inpediatric patients. J Pediatr Pharmacol Ther 2001;6:107-19.
- 66. Joseph FS, Catherine Tuleu. Paediatric formulations-Getting to the heart of the problem, International Journal of Pharmaceutics. 2005;300:56-66.
- 67. Standing JF, Khaki Z Fand, Wong I. Poor formulation information inpublished pediatric drug trials. Pediatrics. 2005;116:559-62.
- 68. Alexander KS, Haribhakti RP, Parker GA. Stability of acetazolamide in suspension compounded from tablets", Am J Hosp Pharm 1991;48:1241-44.
- 69. Fawcett JP, Stark G, Tucker IG, Woodd DJ. Stability of dantrolene oral suspension prepared from capsules. J Clin Pharm Ther. 1994;19:349-53.
- 70. Anonymous. Boots pharmacist and trainee cleared of baby"s manslaughter. Pharm J 2000;264:390-92.
- 71. Joseph F. Paediatric formulations-Getting to the heart of the problem", Int J Pharm 2000;300:56–66.
- 72. Teng J, Song CK, Williams RL, Polli JE. Lack of medication dose uniformity in commonly split tablets. J Am Pharm Assoc 2002;42:195–99.
- 73. Breitkreutz J, Wessel, Tand Boos J. Dosage forms for oral drug administration to children. Paediatr Perinatal Drug Ther 1999;3:25–33.
- 74. Milap C Nahata, Loyd V Allen. Extemporaneous Drug Formulations. Clinical Therapeutics. 2014; 30(11): 2112-20.
- 75. Rafiee TM, Mehramizi A. In vitro release studies of piroxicam from oil-in-water creams and hydroalcoholic gel topical formulations. Drug Dev IndPharm 2000;64:409-14.
- Voudrie MA, Allen DB. Stability of Oseltamivir Phosphate in SyrSpend SF, Cherry Syrup, and SyrSpend SF (For Reconstitution). IJPC 2010;14(1):82-85.
- 77. Winiarski AP, Infeld MH, Tscherne R, et al. Preparation and stability of extemporaneous oral liquid formulations of

oseltamivir using commercially available capsules. J Am Pharm Assoc 2003; 47(6):747-55.

- 78. Glass BD, Haywood A. Stability considerations in liquid dosage forms extemporaneously prepared from commercially available products. Journal of Pharmacy and Pharmaceutical Sciences. 2009;9(3):398-426.
- 79. Donnelly RF, Pascuet E, Ma C, et al. Stability of celecoxib oral suspension. Can J Hosp Pharm. 2009;62(6):464-68.
- Skillman KL, Caruthers RL, Johnson CE. Stability of an extemporaneously prepared clopidogrel oral suspension. Am J Health Syst Pharm 2010: 67(7):559-61.
- Mihaila B, Ellis D, Rozek T, Milne R. Chiral Stability Study of Oral Liquid Clopidogrel Formulations for Infants. J Pharm Prac Res 2012:42(2):106-10.
- Walker SE, Baker D, Law S. Stability of clozapine stored in oral suspension vehicles at room temperature. Can J Hosp Pharm 2005;58(5):279-84.
- Walker SE, Baker D, Law S. Stability of clozapine stored in oral suspension vehicles at room temperature. Can J Hosp Pharm 2005;58(5):279-84.
- Chou J, Decarie D, Dumont RJ, et al. Stability of dexamethasone in extemporaneously prepared oral suspensions. Can J Hosp Pharm 2001;54(2):97-103.
- Cober MP, Johnson CE, Sudekum D, Penprase K. Stability of extemporaneously prepared glycopyrrolate oral suspensions. Am J Health Syst Pharm. 2001;68(9):843-45.
- Ensom MH, Decarie D, Sheppard I. Stability of lansoprazole in extemporaneously compounded suspensions for nasogastric or oral administration. Can J Hosp Pharm. 2007;60(3):184-91.
- 87. Melkoumov A, Soukrati A, Elkin I, Forest JM, et al. Quality evaluation of extemporaneousdelayed-release liquid formulations of lansoprazole. Am J Health Syst Pharm 2011;68(21):2069-74.
- Ensom MH, Decarie D, Rudolph S. Stability of levetiracetam in extemporaneously compounded suspensions. Can J Hosp Pharm 2011;64(3):207-11.
- Johnson CE, Cober MP, Thome T, Rouse E. Stability of an extemporaneous alcohol-free melatonin suspension. Am J Health Syst Pharm. 2011;68(5):420-23.
- Aliabadi HM, Romanick M, Somayaji V, Mahdipoor P, Lavasanifar A. Stability of compounded thioguanine oral suspensions. Am J Health Syst Pharm. 2011;68(10):900-08.
- Donnelly RF, Wong K, Goddard R, Johanson C. Stability of Venlafaxine Immediate-Release Suspensions. International Journal of Pharmaceutical Compounding. 2011;15(1):81-4.
- Abobo CV, Wei B, Liang D. Stability of zonisamide in extemporaneously compounded oral suspensions. Am J Health Syst Pharm. 2009;66(12):1105-09.
- 93. DiGiacinto JL, Olsen KM, Bergman KL, et al. Stability of suspension formulations of lansoprazole and omeprazole stored in amber-colored plastic oral syringes. Ann Pharmacother 2000;34(5):600-05.
- 94. Ghulam A, Keen K, Tuleu C, et al. Poor preservation efficacy versus quality and safety of pediatric extemporaneous liquids. Ann Pharmacother. 2007;41(5):857-60.
- Lim LY, Tan LL, Chan EW, et al. Stability of phenoxybenzamine hydrochloride in various vehicles. Am J Health Syst Pharm. 1997;54(18):2073-78.
- 96. Nahata MC, Morosco RS, Leguire LE. Development of two stable oral suspensions of levodopa-carbidopa for children with amblyopia. J Pediatr Ophthalmol Strabismus. 2000;37(6):333-37.
- 97. Caruthers RL, Johnson CE. Stability of extemporaneously prepared sodium phenylbutyrate oral suspensions. Am J

Health Syst Pharm 2007;64(14):1513-15.

- 98. Cober MP, Johnson CE. Stability of an extemporaneously prepared alcohol-free phenobarbital suspension. Am J Health Syst Pharm 2007;64(6):644-46.
- 99. Burnett JE, Balkin ER. Stability and viscosity of a flavored omeprazole oral suspension for pediatric use. Am J Health Syst Pharm 2006;63(22):2240-47.
- 100. Vaishnani R. Formulation and evaluation of immediate release tablets of paroxetine HCl using different superdisintegrants. Int J Res Pharma Biomedical Sci 2011; 2(3):1095-99.