

Preparation and Evaluation of Lipid Emulsion Containing 13 Vitamins for Injection Without Anaphylactoid Reactions

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Objective: Multivitamins containing Tween 80 can cause anaphylactoid reactions. The objective of this study was to develop a new lipid emulsion containing 13 fat- and water-soluble vitamins for injection (13V-LE) that were simultaneously dissolved in one bottle and to evaluate the stability of and anaphylactoid reactions to 13V-LE.

Methods: Particle size, ζ -potential, and polydispersity of 13V-LE were assayed with a Zetasizer Nano ZS. Entrapment efficiency of 13V-LE was determined with HPLC. Behavior, histamine, and blood pressure of beagle dogs were investigated by observation, fluorospectrophotometry, and sphygmomanometry.

Results: The 13V-LE with the smallest particles and highest entrapment efficiency with stable ζ -potential was composed of soybean oil, glycerin (2.25%, w:v), egg lecithin (1.2%, w:v), and purified water. There was no obvious change in characteristics of the 13V-LE samples in terms of appearance, size distribution, ζ -potential, pH value, or concentration over 6 months. In anaphylactoid reactions tests, when being administered with the multivitamin Infuvite Adult containing Tween 80, six beagles showed grade IV symptoms ($P < 0.01$ vs control), low blood pressure, and high plasma-histamine concentrations ($P < 0.05$ or $P < 0.01$). However, there were no significant differences in behavior, blood pressure, or histamine concentration in the dogs before and after administration in the 13V-LE group.

Conclusion: The 13V-LE formulation is a suitable intravenous lipid emulsion without anaphylactoid reactions.

Keywords: anaphylactoid reaction, multivitamins, lipid emulsion, beagle dog

Introduction

Adverse drug reactions cause increased patient morbidity and mortality, are a considerable source of financial burden to the health-care system, and remain a huge challenge in modern health care.¹ Adverse drug reactions occur in approximately 17% of patients.^{2,3} Hypersensitivity reactions comprise approximately a third of adverse drug reactions. Anaphylactoid reactions represent >60% of hypersensitivity reactions.⁴ Avoiding adverse drug reactions, especially anaphylactoid reactions, is essential from both an economic and ethical standpoint.

Nutrition is the source of healthy growth and organism activity, and indispensable for the recovery of patients. The prevalence of malnutrition in patients can be as high as 50%.⁵⁻⁸ Malnutrition has become an independent risk factor of mortality, morbidity, and hospitalization in patients.⁹⁻¹⁴ Multivitamins are conducive to the

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ingestion of essential vitamins to maintain the body's resistance and repair functions. Multivitamins are the most common supplement, and their use is increasing.¹⁵

Thirteen vitamins needed by the human body are mainly used in multivitamin preparations for parenteral nutrition support.¹⁶

The multivitamin Infuvite Adult is an injection containing 13 vitamins that has been widely used for years. The product comprises two bottles. Bottle 1 contains vitamins A, D, E, K, C, B₁, B₂, B₆, nicotinamide, and dexpantenol. Tween 80 (1.4%) is used to solubilize the fat-soluble vitamins A, D, E, and K, in bottle 1. Bottle 2 contains vitamin B₁₂, biotin, folic acid, using 30% propylene glycol as solvent. The two bottles are mixed before clinical injection. However, a problem occurs with the use of Tween 80. There is increasing evidence that it can cause anaphylactoid reactions.^{17–19} Not surprisingly, multivitamins can produce severe and fatal anaphylactoid reactions.^{20–23} Tween 80 is not suitable as a solubilizing agent for intravenous preparations. In addition to Tween 80–induced anaphylactoid reactions, there are some difficulties in the study of multivitamin preparation. Fat-soluble vitamins are insoluble in water. It is difficult to dissolve fat- and water-soluble vitamins in the same system.

To solve these problems, a new lipid emulsion containing 13 fat- and water-soluble vitamins in one bottle (13V-LE) was developed in this study. Intravenous lipid emulsions were used to supply essential fatty-acids to patients with intestinal failure who needed parenteral nutrition in the US in the 1970s.²⁴ Over the last 50 years, new lipid emulsions have been continuously produced with the goal of improving security and efficacy and realizing physiologically optimal formulations.²⁴ In the present day, lipid emulsions have become a tool for systemic delivery of poorly soluble and cytotoxic drugs.^{25,26} 13V-LE combines water-soluble and fat-soluble vitamins without Tween 80 and forms a cosoluble single freeze-dried compound, which could improve vitamins stability and avert hypersensitivity. The objective of this study was to evaluate the stability and the anaphylactoid reactions of 13V-LE.

Methods

Materials

Vitamins were purchased from Disiman Vitamin (Shanghai, China). Poloxamer 188 (Pluronic F68) was purchased from Shanghai Qingyuanxing Chemical Technology (Shanghai, China). Glycerol, oleate, soybean oil, and egg lecithin (PC 98%) was supplied by Xi'an

Libang Pharmaceutical (Shaanxi, China) and α -Tocopherol purchased from Sigma-Aldrich (MO, USA).

Preparation

Formulation and optimized composition of 13V-LE were begun with filtration sterilization and ultrasonic treatment. Briefly, in the case of 13V-LE, fat-soluble vitamins and soybean oil were considered as oil phase. Egg lecithin and oleate acid were chosen as emulsifier. Water-soluble vitamins, glycerin, and purified water were considered as a water phase.

The new multivitamin 13V-LE contains nine water-soluble vitamins (B₁, B₂, B₆, B₁₂, C, nicotinamide, dexpantenol, biotin, and folic acid) and four kinds of liposoluble vitamins (including vitamins A, D₃, E, and K₁). These are shown in Table 1. In order to solve the problem of the cosolubility of water-soluble and liposoluble vitamins in the same system, 16 emulsifier formulations were designed, as presented in Table 2. Various proportions of lecithin, F68, oleate, and Tween 80 were used as emulsifiers for the preparation of 13V-LE. Soybean oil was considered as a solvent of the oil phase. Glycerin (2.25%, w:v) and purified water were considered as a solvent of the water phase in the preparation of the different formulations. Based on physical appearance, particle size, ζ -potential, and entrapment efficiency (EE), an optimized formulation of emulsifiers was chosen.

To prepare 13V-LE, egg lecithin was dissolved into dehydrated alcohol and agitated until dissolved with the addition of fat-soluble vitamins. Then, soybean oil was added in the dissolved solution and heated at 70°C until the mixture had become a clear and stable oil phase containing emulsifier. Glycerin and oleate acid were dissolved in water and heated at 70°C to form a water phase. After that, the water-phase

Table 1 Composition of 13V-LE

Components	Quantity
Vitamin A	3,300 IU
Vitamin D ₃	200 IU
Vitamin E	10 IU
Vitamin K ₁	0.15 mg
Vitamin B ₁	6 mg
Vitamin B ₂	3.6 mg
Vitamin B ₆	6 mg
Vitamin B ₁₂	0.005 mg
Vitamin C	200 mg
Biotin	0.06 mg
Niacinamide	40 mg
Dexpantenol	15 mg
Folic acid	0.6 mg

Table 2 Emulsifiers for 13V-LE (w:v, %)

Formulations	Lecithin	F68	Oleate	Tween 80
F1	0.6	0	0	0
F2	0.6	0.2	0.04	0.2
F3	0.6	0.4	0.06	0
F4	0.6	0.6	0.02	0.2
F5	1.2	0	0	0
F6	1.2	0	0.06	0.2
F7	1.2	0.6	0.04	0
F8	1.2	0.4	0.02	0
F9	2.0	0.4	0.04	0.2
F10	2.0	0.6	0	0
F11	2.0	0	0.02	0.2
F12	2.0	0.2	0.06	0
F13	3.0	0.6	0.06	0.2
F14	3.0	0.4	0.02	0
F15	3.0	0.2	0	0.2
F16	3.0	0	0.04	0

solution was added slowly to the oil-phase solution containing the emulsifier. The coarse emulsion was prepared by high-speed shearing for 15 minutes at 70°C under the protection of nitrogen. The coarse emulsion was cooled to room temperature and adjusted to pH 5–7 with NaOH. The final emulsion was prepared with high-pressure homogenization (Niro Soavi NS1001L) at 1,000 bar for eight cycles at 40°C. After homogenization, the water-soluble vitamins were added to form 13V-LE, then freeze-dried after sterilization through a 0.22 µm bacteria-retentive filter.

Characterization of Formulations

Particle size, polydispersity, and ζ-potential of the 13V-LE were assayed with a Zetasizer Nano ZS (Malvern Instruments). The samples of 13V-LE were redissolved in 5 mL water, then diluted to 1:1,000 with water before measurement. The pH value of the 13V-LE was tested with a pH meter (PP-50 Professional Meter; Sartorius).

Entrapment Efficiency

The EE of the emulsions was assayed by vitamin concentration in the dispersion phase. 13V-LE was centrifuged at 50,000 rpm for 1.5 hours at 4°C by ultracentrifugation (OPTIMA L-100XP; Beckman Coulter), then collected in the water phase. Vitamin concentrations in the water phase were measured with HPLC. EE was calculated:

$$EE(\%) = \frac{C_{total} V_{total} - C_{water} V_{water}}{C_{total} V_{total}} \times 100$$

where C_{total} is the concentration of vitamin in the emulsion, V_{total} the volume of the emulsion, C_{water} the concentration of vitamin in the water phase after ultracentrifugation, and V_{water} the volume of water phase collected after ultracentrifugation.

Vitamin A was the most abundant and unstable in the oil phase among the vitamins of 13V-LE, so the stability of the formulation was determined by examining the encapsulation rate of vitamin A. The concentration of fat-soluble vitamin A was measured with HPLC using a Waters 2996 UV detector. The column was an Agilent XDB-C₁₈ (4.6×150 mm, 5 µm), and the mobile phase was a mixture (methanol : isopropanol 95:5 v:v). The wavelength was 326 nm and the injection volume 10 µL. The flow rate of the mobile phase was 1.2 mL/min. The column temperature was 25°C. All measurements were repeated in triplicate.

Stability

13V-LE was stored at 40±2°C and at room temperature for 6 months. After that, the physical and chemical stability of these samples were evaluated with assays of appearance, polydispersity, particle-size distribution, ζ-potential, pH value, and drug remaining. All measurements were repeated three times. For appearance evaluation, if a lipid emulsion is stratified, aggregated, flocculated, produces small oil droplets on the surface, or shows other anomalous changes, it can be defined as “bad”. If a lipid emulsion is homogeneous and shows no anomalous changes, it can be defined as “good”.

Compatibility

Compatibility of the 13 vitamins with pharmaceutical excipients and that of 13V-LE with different infusion solutions were investigated. Vitamins were mixed with different pharmaceutical excipients, as shown in [Supplementary Table 1](#). These mixtures were exposed to high temperature (60°C), damp (90%±5%), light of 4,500 ±500 lx for 10 days. The content of vitamins and related substances were tested with HPLC. To test compatibility with different infusion solutions, samples of 13V-LE were redissolved in 5 mL water for injection, then diluted to 500 mL with 0.9% sodium chloride solution, 5% glucose injection, and fatemulsion injection. The osmotic pressure of these mixtures was measured with an Osmomat-3000D (Gonotec).

Anaphylactoid Reaction Tests

Animals and Treatments

Six beagle dogs (three male, three female) weighing 8–12 kg were purchased from Xi'an Dilepu Biology Resources (Shaanxi, China). They were housed in separate cages and allowed access to food and water ad libitum in an environment at room temperature with 40%–70% humidity, and a day–night cycle of 12–12 hours. The dogs were handled in accordance with the criteria outlined in the *Guide for Care and Use of Laboratory Animals*. All experimental protocols were approved by the Animal Ethics Committee at Xi'an Jiaotong University (2017–0045).

13V-LE and Infuvite Adult are compound vitamin injections. The content of each vitamin in the injections is different and the units of measurement inconsistent. For the convenience of research and calculation, vitamin C was used as a marker of 13V-LE and Infuvite Adult to calculate the dose. The dose of vitamin C in 13V-LE and Infuvite Adult for an adult is 200 mg/day. When human body weight is set at 70 kg, the dose of vitamin C is 2.9 mg/kg. The dose-converse factor of dog:person is 1.85. Therefore, the equivalent dose of vitamin C for a dog is 5.3 mg/kg. In this study, 6 mg/kg vitamin C was selected as the dose for the dogs.

There were three groups in the anaphylactoid reaction tests: control (normal saline), 13V-LE, and Infuvite Adult. On the first day of the dosing regimen, all six beagles received intravenous saline injections as the control group. Then, all dogs were divided into two groups of three animals each. Two groups of dogs were cross-injected with Infuvite Adult and 13V-LE on the second and fourth days. On the second day, dogs 1–3 were intravenously injected with Infuvite Adult and dogs 4–6 intravenously injected 13V-LE. On the fourth day, dogs 1–3 were intravenously injected with 13V-LE dogs 4–6 dogs intravenously injected with Infuvite Adult (Table 3). Blood samples were collected in tubes with heparin before and 10 minutes after administration. Samples were centrifuged for 10 minutes, then plasma collected and stored at –20°C until analysis.

Table 3 Modes of administration in anaphylactoid reaction tests

Dogs	Time of administration		
	Day 1	Day 2	Day 4
1–3	Normal saline	Infuvite Adult	13V-LE
4–6	Normal saline	13V-LE	Infuvite Adult

Behavioral Observation

Before and after administration, dogs' eyes, mouth, neck, abdomen, limbs, skin, behavior, gait, and body coordination were observed for 30 minutes for anaphylactoid reactions. Levels of reaction were assessed using the criterion in [Supplementary Table 2](#).

Plasma-Histamine Concentration

Concentrations of plasma histamine were measured with fluorescence spectroscopy. Ethyl alcohol (300 µL) was added to plasma samples of 100 µL for deproteinization. The supernatant was tested after vortex mixing and centrifugation at 10,000 g. Sample supernatants (100 µL) were added to 0.5 M NaOH solution (40 µL) and 2.5 µg/mL *o*-phthalaldehyde (20 µL) and incubated at 37°C for 30 minutes. Subsequently, 10 µL 3 M HCl solution was added to terminate the reaction. The excitation wavelength was 365 nm and the emission wavelength 465 nm. The fluorescence intensity of samples was detected with a fluorescence plate reader, and the standard curve was drawn, and then histamine concentrations of samples were calculated.

Blood-Pressure Detection

Blood pressure was measured with an OMRON HEM-8611 at 0, 10, 20, and 30 minutes after administration of drugs. Dogs were put in the decubitus position without anesthesia and the blood-pressure monitor put on the arm. Measurements were repeated in triplicate at each time point.

Statistical Analysis

SPSS 11.5 was used for statistical analyses. All data are shown as means ± SE. $P < 0.05$ was considered statistically significant. ANOVA with least significant difference was used to analyze quantitative data in multiple groups, including blood pressure at different time points. Student's *t*-test was used to analyze quantitative data between two groups, such as concentrations of histamine. The rank-sum test was used to analyze ordinal data, such as grades of behavior, in the anaphylactoid-reaction tests.

Results and Discussion

Composition and Stability of 13V-LE HPLC

Representative chromatograms were shown in [Supplementary Figure 1](#). The peak of vitamin A was eluted within about 33 minutes under the analytical conditions described. The reproducibility of the detection was studied with repeated injections in triplicate samples containing

vitamin A at three concentrations (2, 3, and 4 µg/mL). Relative standard deviation was <0.3%. The relative recovery of vitamin A at 2, 3, and 4 µg/mL was 101.8%±0.0%, 102.3%±0.0%, and 103.0%±0.0%, respectively (n=3). The standard curve of vitamin A concentration was obtained: $y=3.401x+73.986$, $R^2=0.9994$.

Composition of 13V-LE

In this study, a new formulation of intravenous lipid emulsions was developed to solve the problem of cosolvability of water-soluble vitamins (B₁, B₂, B₆, B₁₂, C, nicotinamide, dexpantenol, biotin, and folic acid) and liposoluble vitamins (A, D₃, E, and K₁) in the same system. The appearance, ζ-potential, particle-size distribution, and EE of 13V-LE in the different emulsifiers (lecithin, F68, oleate, and Tween 80) were assessed to select the optimal formulation of 13V-LE. In the emulsified state, 13V-LE was in an oil-in-water state, and the four liposoluble vitamins (A, D₃, E, and K₁) were encased in oil in the lipid emulsion. Commonly, only the EE of encased vitamins needs to be evaluated. Vitamin A was the most abundant and unstable vitamin in the oil phase in 13V-LE. Therefore, it was chosen as a representative to evaluate the stability of the formulation by examining its EE. As shown in Table 4, F1, F4, F6, F9, F10, F13, and F14 were inferior formulations with poor appearance, such as oil drops, stratification, or flocculation. 13V-LE particle size increased with increasing lecithin content. When emulsifiers contained 0.6% lecithin, F68, oleate, and Tween 80 in F2 and

F3, this contributed to emulsion formation, but with low EE. Particles of F7, F8, F11, F12, F13 and F14 containing F68 or oleate were larger than those of F5 containing lecithin. When an emulsifier contained only 1.2% lecithin, the emulsion had the smallest particles and highest EE of vitamin A with stable ζ-potential (F5). Therefore, egg lecithin 1.2% (w:v) was chosen as an emulsifier.

Before freeze-drying, 13V-LE is an oil-in-water lipid emulsion. The new multivitamin 13V-LE contains 3,300 IU (990 µg) vitamin A, 200 IU vitamin D₃, 10 IU (9.1 mg) vitamin E, 0.15 mg vitamin K₁, 200 mg vitamin C, 6 mg vitamin B₁, 3.6 mg vitamin B₂, 6 mg vitamin B₆, 40 mg nicotinamide, 15 mg dexpantenol, 0.005 mg vitamin B₁₂, 0.06 mg biotin, 0.6 mg folic acid, soybean oil, egg lecithin (1.2%, w:v), glycerin (2.25%, w:v), NaOH, and water. Fat-soluble vitamins and soybean oil were considered as oil phase, egg lecithin was chosen as an emulsifier, and water-soluble vitamins, glycerin, and water were considered as water phase in 13V-LE.

Characterization of Lipid Emulsions

The 13V-LE dispersion, composed of soybean oil, glycerin, egg lecithin, 13 vitamins, and purified water, was analyzed in terms of size and polydispersity. As shown in Figure 1A, the mean diameter of 13V-LE was <200 nm, which is suitable for parenteral administration. Polydispersity is a dimensionless parameter of size distribution for broadness. The lower the polydispersity, the

Table 4 Effects of different emulsions on characterization of 13V-LE

Formulations	Characterization of 13V-LE			
	Physical appearance	Particle size (nm)	ζ-potential (mV)	Entrapment efficiency of vitamin A (%)
F1	Oil drops	—	—	—
F2	Good	152.4±8.7	-20.2±1.5	70.4±3.1
F3	Good	176.3±11.7	-26.3±1.9	85.5±2.0
F4	Stratification	—	—	—
F5	Good	121.1±6.0	-27.7±2.1	99.7±0.3
F6	Oil drops	—	—	—
F7	Good	205.1±3.2	-29.6±3.4	94.9±1.4
F8	Good	186.8±5.5	-24.0±2.0	98.2±0.4
F9	Flocculation	—	—	—
F10	Oil drops	—	—	—
F11	Good	194.1±5.0	-25.4±2.0	95.3±1.2
F12	Good	215.8±9.0	-30.7±1.6	92.4±3.2
F13	Oil drops	—	—	—
F14	Oil drops	—	—	—
F15	Good	217.3±3.8	-25.8±1.4	94.6±2.1
F16	Good	226.8±6.6	-29.5±1.9	93.8±2.5

Note: n=3.

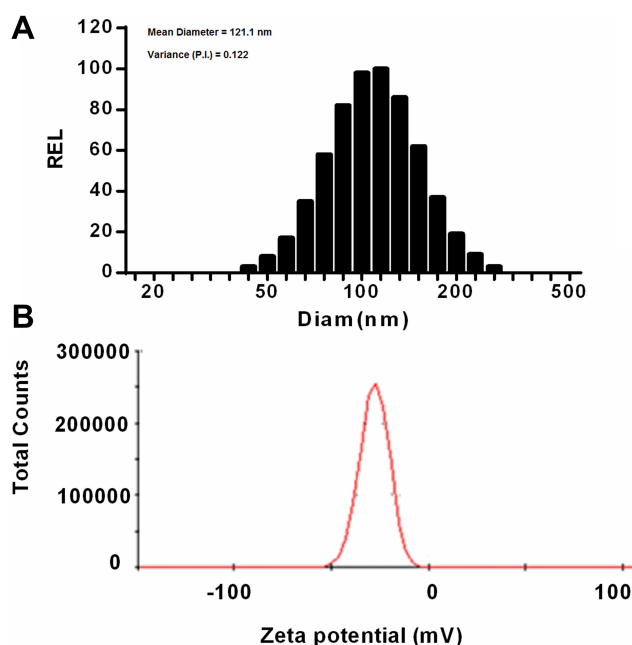


Figure 1 Particle-size and the ζ -potential distribution of 13V. (A) Particle-size distribution; (B) ζ -potential distribution. 13V-LE samples were diluted 1:1,000 with purified water. Particle size and polydispersity were measured with a Zetasizer Nano ZS. Measurements were performed in triplicate.

Abbreviation: PI, polydispersity index.

narrower the distribution of lipid emulsion. The mean polydispersity of 13V-LE was <0.2 , showing that 13V-LE had proper delivery.

ζ -potential is a crucial parameter of emulsion stability. It determines the size of electrostatic repulsion among particles, which affects the aggregation of particles and the stability of a lipid emulsion. Generally, ζ -potential ranges from -20 mV to -45 mV, while that of 13V-LE was -27.7 ± 2.1 mV, suggesting that the emulsions were stable (Figure 1B).

Stability of 13V-LE

13V-LE was stored at 25°C and 40°C for 6 months, then its stability was studied by monitoring size and polydispersity. Table 5 shows that the appearance, particle-size distribution, pH value, ζ -potential, and vitamin A content of the 13V-LE samples underwent no significant change in 6 months, suggesting that 13V-LE had favorable stability. Severe test, accelerated, and long-term storage tests also showed that 13V-LE had favorable stability (Supplementary Table 3–5).

Multivitamins/minerals refers to any supplement containing three or more vitamins or minerals, but no herbs, hormones, or drugs, with each component at a dose less than the tolerable upper level determined by the US Food and Nutrition Board, ie, the maximum daily intake likely to pose

Table 5 Physical stability of 13V-LE during 6 months ($n=3$)

Characterizations of 13V-LE	0 Month	6 Months at 40°C	6 Months at 25°C
Physical appearance	Good	Good	Good
pH value	6.35 ± 0.13	6.17 ± 0.18	6.26 ± 0.2
Average particle size (nm)	121.1 ± 6.0	132.5 ± 11.7	125.4 ± 9.0
Average ζ -potential (mV)	-27.7 ± 2.1	-28.1 ± 3.1	-26.2 ± 3.0
Content of vitamin A (%)	99.7 ± 0.2	96.0 ± 1.9	98.1 ± 1.4

no risk of adverse health effects.²⁷ The content of the vitamins in 13V-LE is the same as that in Infuvite Adult. The differences between 13V-LE and Infuvite Adult are their form and pharmaceutical adjuvants. Tween 80 is used to solubilize the fat-soluble vitamins in Infuvite Adult. In order to avoid Tween 80-induced anaphylactoid reactions, a lipid emulsion was chosen to be the drug-carrier system for 13V-LE. Lipid emulsions have been a significant component of parenteral nutrition and play a crucial role in supplying calories, providing necessary fatty acids that cannot be endogenously synthesized, and more importantly improving safety and efficacy and achieving physiologically optimal formulations.²⁴ Soybean oil, which has high phytosterol content, was the triglyceride source of the first lipid emulsion manufactured and successfully administered intravenously.²⁸ Egg lecithin, a natural phospholipid, is used in oral, dermal, and parenteral products, including oil-in-water emulsions as an emulsifier.²⁹ Soybean oil, egg lecithin, and glycerin formed a stable and homogeneous drug-carrier system for 13V-LE, with high ζ -potential and low polydispersity. Many vitamins, such as vitamin C, are sensitive and easily lost during handling, processing, and storage.³⁰ Loss of vitamin C during storage is lower in freeze-dried powder than vacuum, tunnel-dried, or fresh powder.^{31,32} Cernevit is also a cosoluble single freeze-dried compound containing nine water-soluble vitamins and three liposoluble vitamins without vitamin K. Freeze-dried technology is conducive to the stability of vitamins. Our testing suggested that 13V-LE had good stability.

Compatibility of 13V-LE

The compatibility of vitamins with the pharmaceutical excipients soybean oil, egg lecithin, glycerin, F68, Tween 80, and oleate was investigated. The content of each vitamin and

the related substances did not change significantly and was within the range of standard requirements, suggesting that the vitamins had good compatibility with these excipients (Supplementary Table 6). The 13V-LE was mixed with 0.9% sodium chloride solution, 5% glucose injection, and fat-emulsion injection. None of these mixtures showed anomalous changes in appearance, such as stratification, aggregation, and flocculation. The osmotic pressure of these mixtures were 290.3 ± 0.9 Osm/kg, 292.7 ± 0.9 Osm/kg, and 301.7 ± 1.8 Osm/kg, respectively (Table 6). The osmotic pressure of human blood is 285–310 Osm/kg. The range of osmotic pressure required for intravenous preparations should be the same as the osmotic pressure of blood in the body. The results of osmotic pressure suggested that 13V-LE had good compatibility with different infusion solutions. The drug-carrier system of 13V-LE consisting of soybean oil, egg lecithin, and glycerin is similar to that of propofol, which is an intravenous anesthetic and Vitalipid, which is a fat-soluble injection containing vitamins A, D, E, and K₁.³³ These products have good stability and compatibility with parenteral nutrition admixtures.

Anaphylactoid Reactions

Alterations in Behavior

The behavior of the dogs was observed in anaphylactoid reaction tests. The six beagles of the saline group did not show any abnormal behavior within 30 minutes after administration (Table 7 and Figure 2). In the 13V-LE group, there was no abnormal behavior after intravenous infusion of 13V-LE. Within 30 minutes after intravenous administration of Infuvite Adult, the six dogs displayed grade IV symptoms, such as nose and head scratching and ear flicking (>3 minutes), multiple skin and mucous rubeosis, skin rash, tachypnea, instability of gait, diarrhea, pawing the ground, rubbing the ground, rolling over, somnolence, hypodynamia, and wheezing. Figure 2 show the influence of 13V-LE on the skin of the dogs. There was multiple skin and mucous rubeosis and skin rash in dogs who had been administered Infuvite Adult. Anaphylactoid reactions were grade IV with a mean

Table 6 Osmotic pressure of 13V-LE mixed with different infusion solutions (n=3)

	Compatibility volume (mL)	Osmotic pressure (Osm/kg)
Normal saline	500	290.3 ± 0.9
5% Glucose	500	292.7 ± 0.9
Fat emulsion	500	301.7 ± 1.8

Table 7 Anaphylactoid reaction tests

	Dogs (n)							Score	Grade
	0	I	II	III	IV	V	VI		
Control	6	0	0	0	0	0	0	0	0
13V-LE	6	0	0	0	0	0	0	0	0
Infuvite Adult	0	0	0	0	6	0	0	$25.5 \pm 3.1^{**}$	$4 \pm 0^{**}$

Notes: Behavior of beagles observed for 30 minutes after the intravenous administration of drugs. Values are means \pm SE, n=6. $^{**}P < 0.01$ vs control.

score of 25.5 ± 3.1 , which was obviously higher than the control and 13V-LE groups ($P < 0.01$, Table 7). These results indicated that 13V-LE did not induce any anaphylactoid reaction, but Infuvite Adult did induce anaphylactoid reactions triggered by Tween 80.

Alterations in Blood Pressure

Before administration of Infuvite Adult, systolic and diastolic blood pressure of the dogs were 122 ± 3 mmHg and 69 ± 3 mmHg, respectively. Systolic pressure decreased to 93 ± 4 mmHg at 10 minutes after drug administration, which was significantly lower than systolic pressure before administration ($P < 0.01$). Diastolic pressure was 58 ± 5 mmHg, and there was a declining trend. Systolic pressure at 20 and 30 minutes after administration were 86 ± 7 mmHg and 81 ± 8 mmHg, respectively, which significantly decreased ($P < 0.01$). Diastolic pressure was also lower than before administration ($P < 0.01$). Obviously, Infuvite Adult caused a decline of blood pressure.

There was no obvious alteration in blood pressure within 30 minutes after the intravenous administration of 13V-LE. In the 13V-LE group, systolic and diastolic pressure were 119 ± 2 mmHg and 68 ± 3 mmHg, respectively, before administration. Systolic pressure was 117 ± 2 mmHg, 112 ± 4 mmHg, and 112 ± 3 mmHg at 10, 20, and 30 minutes, respectively, after intravenous administration of 13V-LE. Diastolic pressure was 71 ± 2 mmHg, 68 ± 4 mmHg, and 64 ± 4 mmHg, respectively. Systolic and diastolic pressure did not show significant differences when compared with before administration (Figure 3). These results further demonstrated that 13V-LE had no influence on blood pressure, but Infuvite Adult did.

Alterations in Plasma-Histamine Concentration

In the Infuvite Adult group, plasma-histamine concentration increased from 3.1 ± 0.5 μ g/mL to 5.7 ± 0.9 μ g/mL after administration, an increase of 45.6% ($P < 0.05$). In the 13V-LE group, there was no significant difference in plasma histamine concentration between before and after

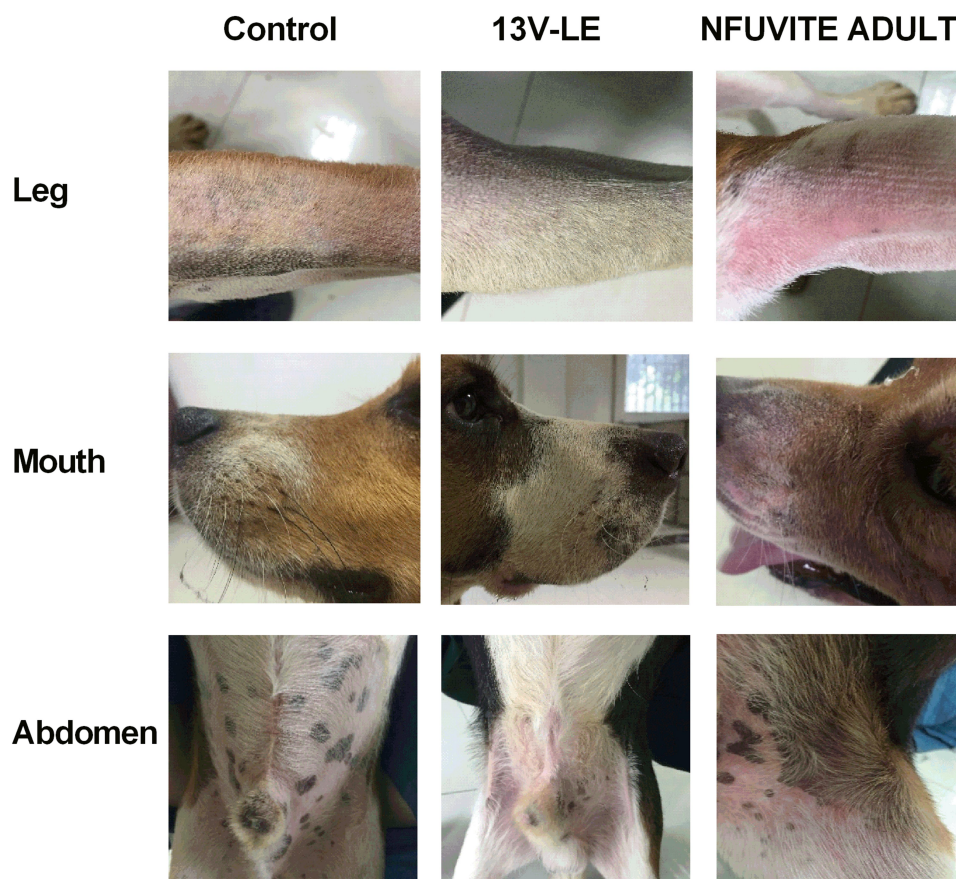


Figure 2 Skin manifestations on beagle dogs after intravenous administration of multiple vitamins. The photos were taken within 30 minutes after the administration, and are representative of six dogs.

administration ($P>0.05$, Table 8). This demonstrated that 13V-LE did not increase plasma histamine, but Infuvite Adult induced an increase in plasma histamine, which is involved in the inflammatory response and itching.

Anaphylactoid reaction, also referred to pseudoallergy or nonallergic drug hypersensitivity, has clinical manifestations that are indistinguishable from anaphylaxis.³⁴ Behavior is considered an important visual and intuitive method to determine anaphylactoid reactions. Beagle dogs are more sensitive and anaphylactoid reactions in them more easily observable than in mice and guinea pigs.³⁵ This is the reason that Beagle dogs were used to investigate anaphylactoid reactions to 13V-LE. From the perspective of animal ethics, we try to minimize the number of animals. Therefore, we reused the dogs on the premise that the accuracy of the results would not be affected. From a pharmacokinetic perspective, 97% of a medicine can be eliminated after five elimination half-lives. In fact, multivitamins cannot be sufficiently cleared from the body after 48 hours, because some vitamins have

a long half-life. For example, vitamins C, D₃, E have half-lives of approximately 10, 15, and 70–90 hours, respectively.^{36,37} However, this had no influence on the anaphylactoid reaction results in this study. Firstly, anaphylactoid reaction is an immediate and transient reaction, and symptoms last for only a few hours. Anaphylactoid reaction is characterized by immediate and transient degranulation of mast cells. Clinical manifestations usually occur within 15 minutes of parenteral injection of the causative agent, reach the maximum usually within 60 minutes, and usually disappear after a few hours.^{38,39} Anaphylactoid reactions to Infuvite Adult disappeared after about 30 minutes. Secondly, the anaphylactoid reactions to Infuvite Adult were induced by Tween 80, but not by the various vitamins. The plasma concentration of Tween 80 was $<0.05\%$ (v:v) within 15 minutes after drug administration.⁴⁰ From a pharmacokinetic perspective, Tween 80 can be sufficiently cleared in the body after 48 hours. Thirdly, in the same group, the results of behavioral observation and plasma histamine concentration on

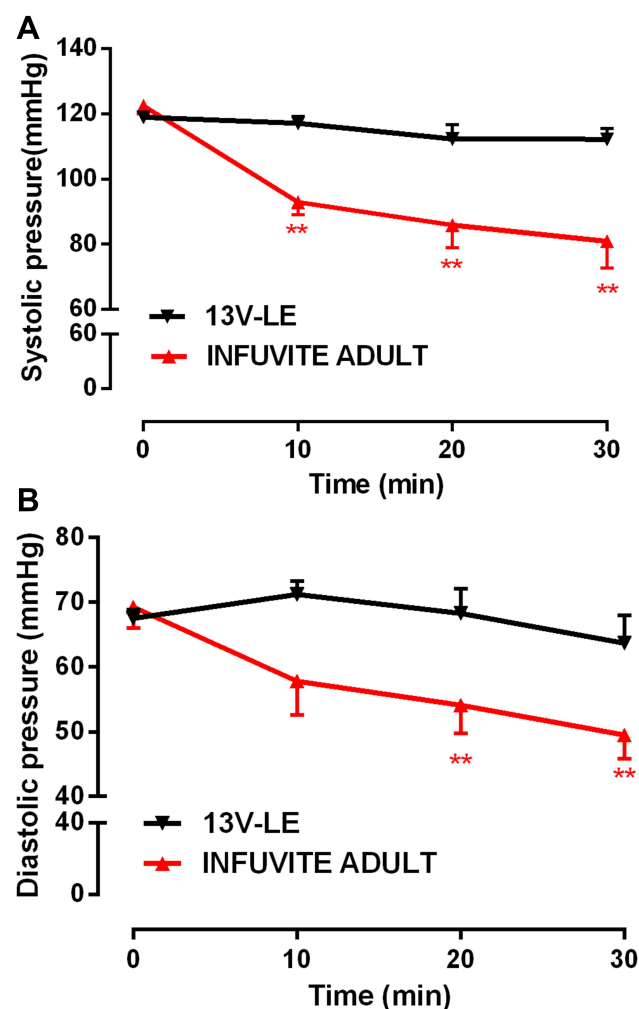


Figure 3 Effect of compound vitamins on blood pressure in beagle dogs. (A) Systolic pressure; (B) diastolic pressure. Means \pm SE, $n=6$. ** $P<0.01$ vs before administration.

the second day were consistent with the results on the fourth day. Therefore, reuse of dogs after 48 hours had no influence on the anaphylactoid reaction test.

Manifestations of anaphylaxis and anaphylactoid reactions are continuous. Weak cutaneous signs, such as rhinorrhea, flushing, and pruritus, often occur at the start.⁴¹ These symptoms in dogs are scratching the nose or head, flinging the ears, sneezing, coughing, skin or mucous

rubeosis, and sialism. The manifestations further aggravate with gastrointestinal symptoms, such as nausea, vomiting, and diarrhea.⁴¹ In more serious cases, hypotension, laryngeal obstruction, severe bronchospasm, shock, and asphyxiation occur.⁴¹ Anaphylactoid reaction is related to the activation of mast cells via non-IgE-mediated degranulation.³⁴ Mast cells can immediately respond to stimulants by degranulation releasing granule-associated mediators, such as histamine, lysosomes, and TNF, into the extracellular milieu, inducing immediate allergic reactions.⁴² Increases in plasma- or tissue-histamine levels have been noted during anaphylaxis and anaphylactoid reactions.⁴¹ Infuvite Adult containing Tween 80 induced anaphylactoid reactions, with serious symptoms (such as pruritus, skin rash, tachypnea, instability of gait, diarrhea, somnolence, hypodynamia, and wheezing), hypotension and high plasma-histamine levels. These reactions were similar to vitamin K₁ injections, with both containing Tween 80.¹⁷ However, behavior, blood pressure, and plasma histamine did not show obvious change in the dogs following administration of 13V-LE, suggesting that it did not induce anaphylactoid reactions.

Conclusion

13V-LE is a novel multivitamin. In this formulation, fat-soluble vitamins and soybean oil were considered as oil phase, egg lecithin was chosen as an emulsifying agent, and water-soluble vitamins, glycerin, and water were considered as water phase. The formulation of 13V-LE displayed high EE with stable ζ -potential, uniform distribution, and small particles, suitable for parenteral administration. Moreover, 13V-LE did not induce anaphylactoid reactions. This formulation of 13V-LE has promise as a new approach to avert hypersensitivity to multivitamins, and to simultaneously improve the stability of vitamins and dissolve water-soluble vitamins and fat-soluble vitamins.

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Table 8 Effect of multivitamins on plasma histamine

	Histamine concentration ($\mu\text{g/mL}$)	
	Before administration	After administration
13V-LE	2.5 \pm 0.2	3.7 \pm 0.4
Infuvite Adult	3.1 \pm 0.5	5.7 \pm 0.9*

Note: * $P<0.05$ vs before administration.

Disclosure

The authors report no conflicts of interest.

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