



Effect of Tween 80 and Span 80 Surfactants Systems on the *Malus domestica* Emulsions for Anti-Cutibacterium acnes

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Abstract: Red apple (*Malus domestica*) extract, rich in hydrophobic quercetin, was formulated into oil-in-water (O/W) emulsion using a 22 factorial design to evaluate Tween 80 (8-10 g) and Span 80 (2-4 g) concentrations, targeting pH (4.5-6.0), transmittance (90-100%), and viscosity (10-2000 cPs). Design-Expert® 13 analysis identified formulations F1, FA, and FAB within acceptable physical property ranges, with all red apple emulsions exhibiting O/W type, skin-compatible pH (5.20-5.48), high transmittance, and suitable viscosity. Freeze-thaw cycling (3 cycles, -15/25°C) and centrifugation showed physical stability with non-significant changes for F1 ($p > 0.05$). The agar well diffusion assay was performed on F1 ($n = 3$), which exhibited optimal physical parameters and met stability criteria, revealing Cutibacterium acnes inhibition zones up to 22.7 ± 0.577 mm. Thus, F1 emerges as a promising nanoemulsion candidate demonstrating antibacterial activity against acne-causing bacteria.

Introduction

Acne vulgaris (AV) is one of the most common dermatological disorders affecting the skin. It affects approximately 75% of adolescents worldwide, typically between the ages of 12 and 15, with peak severity occurring between 17 and 21 years of age. According to the Division of Medical Cosmetics, Department of Dermatology and Venereology, Outpatient Clinic of Dr. Soetomo General Hospital, Surabaya, the highest prevalence of AV cases was found among individuals aged 15-19 years (52.4%) (1). Acne develops due to inflammation of the pilosebaceous follicles, characterized by the appearance of blackheads, pustules, and nodules on the face, shoulders, chest, upper back, and upper arms (2). The occurrence of acne is influenced by various factors, including genetic predisposition, ethnicity, psychological conditions, hormonal imbalance, and more commonly, bacterial infection such as that caused by *Cutibacterium acnes* (3).

Common topical therapies used for treatment of acne include topical retinoids, benzoyl peroxide (BP), antibiotics, clascoterone, salicylic acid, and azelaic acid (4). Systemic therapies such as oral antibiotics, hormonal treatment, and isotretinoin are also widely prescribed for more severe cases. Although these agents are clinically effective, their use is often limited by adverse effects, particularly skin irritation, which may lead to poor patient adherences (5).

Apples are known to contain various compound such as flavonoids, polyphenols, and tannins (6). Quercetin is a flavonoid compound found in apples. It has been

demonstrated to inhibit the growth of acne-causing bacteria such as *Cutibacterium acnes*. The growth of these bacteria is suppressed by quercetin through inhibition of protein synthesis and toxin metabolite production (7). Chandra *et al.* (2022) reported that quercetin gel, yielded a 22.20 mm inhibition zone against *Cutibacterium acnes*. This result affirms the capability of red apple extract emulsion in inhibiting acne pathogenesis, consistent with its antimicrobial properties observed *in vivo* (8).

In line with growing interest in safer and more tolerable alternatives, natural products such as apple extract, a source of the flavonoid quercetin are being explored as active ingredients in cosmetic and dermatological formulations. However, the practical application of quercetin is challenged by its poor aqueous solubility, which can restrict its bioavailability, and consequently, its therapeutic performance in topical products (9).

Nanotechnology offers several advantages in pharmaceutical product formulation, including enhanced solubility of poorly water-soluble compounds and more controlled drug delivery (10, 11). Nanoemulsions (NEs) consist of two immiscible liquids, typically oil and water, stabilized by surfactants (10). NEs are widely applied in dermal delivery systems owing to their small droplet size, fluid properties, and capacity to adsorb onto the stratum corneum lipid layer. NEs are categorized into two primary types: oil-in-water (O/W) and water-in-oil (W/O). O/W NEs effectively deliver hydrophobic agents by enhancing the aqueous solubility of lipophilic substances (12). Apple extract

contain quercetin is a hydrophobic material (13, 14). Thus, O/W NEs represent a suitable approach for formulating apple extract-containing pharmaceutical products to improve active compound solubility in water and optimize formulation absorption.

Tween 80 and Span 80 are widely used in formulations due to their non-ionic properties, which provide superior emulsion stability, high formulation flexibility via HLB matching, and safety for cosmetic products (11). However, emulsions formulated with Tween 80 and Span 80 surfactant systems exhibit inherent limitations that necessitate investigation of their concentration effects on pH, %transmittance, and viscosity for *Malus domestica* formulations. High concentration of surfactants ratios often shift pH toward alkalinity and reduce %transmittance through flocculation or Oswald ripening, while suboptimal ratios yield excessive viscosity that impairs topical spreadability. Examination of these parameters is thus essential to identify stable configurations that enhance the formulation delivery (15). Therefore, the development of a drug delivery system capable of improving the stability and absorption of quercetin is crucial to maximizing its clinical benefit (9). One emerging approach to address these limitations is its formulation in the form of a nanoemulsion serum.

This study aims to evaluate the effect of Tween 80 and Span 80 on pH, %transmittance, and viscosity of *Malus domestica* extract emulsions. The analysis identifies variations in physical properties resulting from different surfactant ratios. Accordingly, an emulsion derived from apple extract holds capability as a nanoemulsion candidate for acne prophylaxis, substantiated by antibacterial assays demonstrating inhibitory effects against acne pathogens.

Methodology or Experimental Section

Materials

The instrument used in this study included a UV-Vis spectrophotometer (Shimadzu UV mini-1240), a laptop (Acer Aspire 3) equipped with Design Expert® 13 software, magnetic stirrer and hotplate (Thermo Scientific), Oven (Mettler), Incubator (Mettler), Laminar Air Flow (Innotech), Rotary Evaporator (Rotavapor R-100 Buchi), and Glassware (Pyrex). The materials used were red apples (*Malus domestica*), Clindamycin gel (Cindala®), Methanol (Smartlab), Tween 80 (Merck), Span 80 (Merck), Virgin Coconut Oil (Smart Organik), Mueller-Hinton Agar (Himedia), Suspension of *Cutibacterium acnes* (agaviLab), and aquadest.

Plant Collection and Determination

Malus domestica fructus was taken from Kota Batu, East Java, in September 2025. The selected fruits were red mature fruit in good condition, without holes and traces of animal bites. Only fruits meeting visual quality criteria, including uniform ripeness and absence of physical damage, were included to ensure sample consistency. Determination was conducted to verify the identity of the plant species used in this study. Determination was performed in UPT Laboratorium Herbal Materia Medica Batu, Kota Batu, East Java, with certificate number 000.9.3/3409/102.20/2025. This taxonomic verification ensured the authenticity of the botanical material prior to further experimental procedures.

Preparation and Extraction of red apple extract

In this study, the extraction of red apple (*M. domestica*) was conducted from its fruit. This extraction was performed utilising maceration technique was adapted from the procedure previously described by Ranjha et al. (2020), with certain modifications. A total of 1 kg of apple was meticulously washed, then subjected to wet-sorting, after which the fruit was thinly sliced and dried in an oven at 45°C. the apple was then milled using a blender and sieved to obtain a powder of uniform size. Subsequently, 15 g of the powder was placed into an Erlenmeyer flask and macerated with methanol at a solid-to-solvent ratio of 1:10. Methanol is highly effective, commonly used maceration solvent for extracting flavonoids from apple materials. This involved the addition of 150 mL of methanol, followed by stirring, covering the flask with aluminium foil, and allowing it to stand for 24 h with intermittent agitation. The mixture was then filtered through the filter paper, after which centrifugation was performed for approximately 10 min at 5000 rpm. Subsequently, the filtrate was subjected to concentration using a rotary evaporator, thereby yielding the extract (16,18).

Preparation of Red Apple Extract Emulsion

The preparation of red apple (*M. domestica*) emulsions was carried out by adapting the method described by Guanse et al. (2025) with several modifications. The formulation process involved preparing two separate mixtures. Firstly, the red apple extract was mixed with virgin coconut oil (VCO). Second, mixture consisting of Tween 80 and Span 80 was blended on a separate hot plate. Both mixtures were stirred using a magnetic stirrer at 1000 rpm and maintained at 45°C until complete homogeneity was achieved. Subsequently, mixture 1 was added dropwise into mixture 2 under continuous stirring at 1000 rpm for 10 min at the same temperature. Following the emulsification step, 100 mL of distilled water was gradually introduced into the homogeneous system while increasing the stirring speed to 1250 rpm for an additional 10 min. The resulting pre-emulsion was then homogenized using a high-speed homogenizer for 2 min and subjected to ultrasonic treatment in a sonicator bath for 45 min with amplitude 65% and pulse on-off every 3 min. All formulations, as presented in **Table 1**, were prepared in triplicate (19). The obtained emulsions were characterized for physicochemical properties (pH, transmittance percentage (%T) by UV-Vis spectrophotometry, and viscosity) and stability test. These evaluations were conducted to assess the physical quality and short-term stability of the prepared emulsion systems.

Table 1. Factorial design scheme.

Material	Function	Formula			
		F1	FA	FB	FAB
Red apple extract	Active substance	10 g	10 g	10 g	10 g
Tween 80	Surfactant	8 g	10 g	8 g	10 g
Span 80	Co-surfactant	2 g	2 g	4 g	4 g
VCO	Oil phase	3 g	3 g	3 g	3 g
Aquadest	Water phase	100 mL	100 mL	100 mL	100 mL

Physical Characterization

Organoleptic Test

All emulsion formulations were subjected to macroscopic physical examination. The observed parameters included colour, odor, and the presence of visually detectable phase separation (20). Data interpret descriptively.

Emulsion Type Test

The emulsion type was determined using the dilution method. Samples were diluted 1:100 (v/v) in both aqueous phase (distilled water) and oil phase (VCO). Formulations exhibiting complete miscibility in the aqueous phase were classified as oil-in-water (O/W) emulsions, whereas those showing complete solubility in the oil phase were identified as water-in-oil (W/O) emulsions (19).

pH Test

The pH value of emulsion was measured using a digital pH meter, with each formulation analyzed in triplicate to determine the mean and standard deviation. The acceptable pH range for the formulation was set between 4.5 and 6.0 (21).

Transmittance Percentage Test

The percent transmittance of the red apple extract emulsions was evaluated using a UV-Vis spectrophotometer at 650 nm wavelength. Each sample was measured in triplicate, with results are expressed as mean transmittance (%) \pm standard deviation to assess optical clarity and emulsion homogeneity. The emulsions were considered to have acceptable optical clarity when the percent transmittance values were within the range of 90-100% (22).

Viscosity Test

Viscosity measurements were performed using an LV-type Brookfield viscometer at room temperature (25°C). Each sample (100 mL) was placed in a glass container, with spindle 2 operated at 3 rpm. Viscosity values in centipoises (cPs) were calculated by multiplying the dial readings by the instrument's correction factor (22).

Data Analysis

All formulas were subjected to a 22-factorial experimental design, utilising Design Expert 13 Software to determine the response of Tween 80 and Span 80 to each physical test parameters, as illustrated in **Figure 1**. Subsequently, the stability results were analysed using a T-test with SPSS software to analysed the optimal formulation, with a confidence level of 95% ($\alpha = 0.05$). The best formula was then subjected to antibacterial testing (23,25).

Stability Test

Emulsion stability was assessed using two complementary methods. In the centrifugation test, samples were placed in Eppendorf tubes and subjected to 5,000 rpm for 5 min. Post centrifugation stability was evaluated by visual inspection for phase separation, precipitation, creaming, or cracking. Additionally, a freeze-thaw cycle test was conducted by storing formulations at -15°C for 48 h, followed by incubation at 25°C for 48 h, with this cycle repeated three times. Stability parameters monitored included pH, %T, and the viscosity (25). These physicochemical measurements were performed before and after the stability tests to determine significant changes in emulsion characteristics, indicating the formulation's physical robustness under stress.

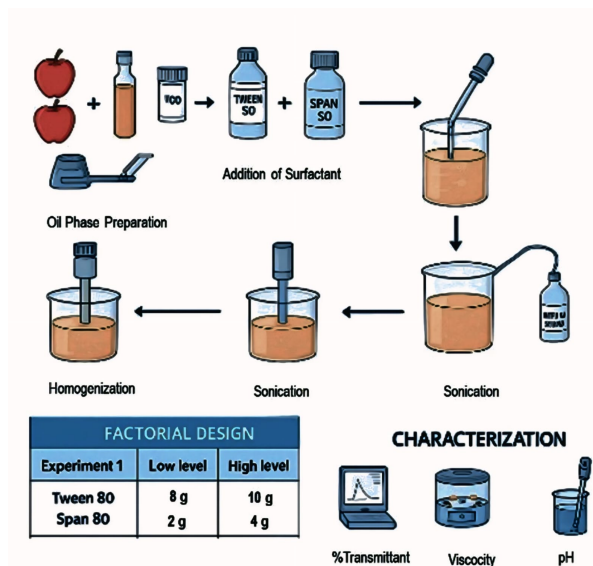


Figure 1. Formulation and characterization scheme

Antibacterial Activity Test

Cutibacterium acnes colonies, cultured for 24 h, were inoculated into 10 mL of 0.9% NaCl solution, vortexed for homogenization, and incubated at 30-37°C for 24 h until turbid. Turbidity was adjusted to match a 0.5 McFarland standard (1.5×10^8 CFU/mL), and this suspension served as the test inoculum. Antibacterial activity of the emulsion formulation was assessed using the agar diffusion method: wells in inoculated agar plates received negative control (aquadest), positive control (clindamycin gel 10 mg/g), and optimal emulsion sample, followed by incubation at 30-37 °C for 24 h. Inhibition zones were measured using calipers (26, 27).

Results and Discussion

Physical Properties and Stability Tests of Red Apple Extract Emulsions

Organoleptic

The formed red apple extract emulsions exhibited transparent and yellowish appearance. The surfactant and co-surfactant combination effectively stabilized the oil-water interface by forming a protective film around the dispersed particles. The HLB values for the four formulations ranged from 11.943 to 13.217.

Emulsion Type

Emulsion types were determined by diluting 1% (v/v) samples in the aqueous phase (20). All four formulations demonstrated complete miscibility in water, they show their classification as oil-in-water (O/W) emulsions. This observation aligns with the calculated HLB values (11,943 to 13,217), which fall within the typical range for O/W emulsion formation.

The HLB value is a calculated parameter derived from the chemical structure of surfactants, allowing pure compounds to be ranked by their relative hydrophilicity-lipophilicity balance (28). HLB values predict both emulsion formation type and stability. Values greater than 10 show oil-in-water (O/W) emulsion, while values below 10 promote water-in-oil (W/O) systems (29). This theory aligns with the experimental findings of this study, which indicated the emulsions as O/W

type.

pH Test

pH measurements were conducted to evaluate the acidity/alkalinity profile of the formulations. Values within the acceptable range for facial skin (4.5-6.0) prevent irritation, dryness, or itching associated pH values ranged from 5.20 to 5.48, meeting skin compatibility criteria.

The relationship between factors, levels, and response values can be specifically interpreted through regression equations and contour plots. The pH response as a function of Tween 80 (X_1) and Span 80 (X_2) levels, derived from factorial design is given by: $Y = 5.34833 - 0.086667X_1 + 0.5X_2 + 0.0116667X_1X_2$. Where Y represents pH, X_1X_2 denotes the Tween 80-Span 80 interaction, and the equation reveals linear relationships with each factor and their interaction, enabling response prediction via substitution of desired factor levels.

Two-way ANOVA analysis revealed significant differences in pH responses across formulations ($p < 0.0001$). Tween 80 exerted the dominant influence (F-value = 515.05; %

contribution = 73.18%), primarily reducing pH values. The interaction effects of Tween 80 and Span 80 on pH responses are illustrated in **Figure 2**.

Transmittance value

Percent transmittance serves as another key indicator for assessing the particle size of nanoemulsion formulations. Particle size trends are indirectly assessed through %T measurements via clarity evaluation principles, though %T alone cannot confirm nano-scale droplet formation. The higher transmittance values correspond to increased clarity and suggest smaller droplet sizes. Values approaching 100% indicate small droplet sizes and good optical clarity. While elevated %T indicates formulation clarity consistent with nanoemulsions, this metric alone cannot distinguish nano-scale droplets from other clear system (e.g., micellar solutions or low-oil microemulsions). Complementary techniques such as droplet size distribution and polydispersity index (PDI) measurements are therefore strongly still required to validate nanoemulsion characteristics beyond transmittance assessment (30, 31).

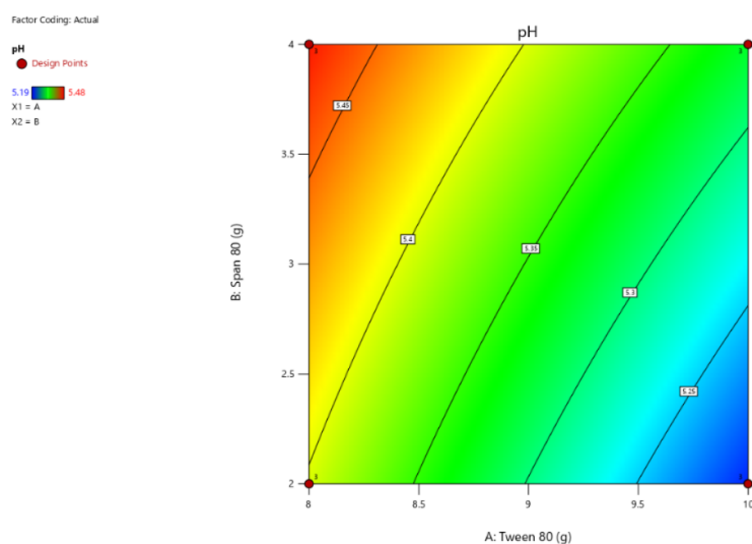


Figure 2. Contour plot illustrating the effect of Tween 80 and Span 80 concentrations on emulsion pH.

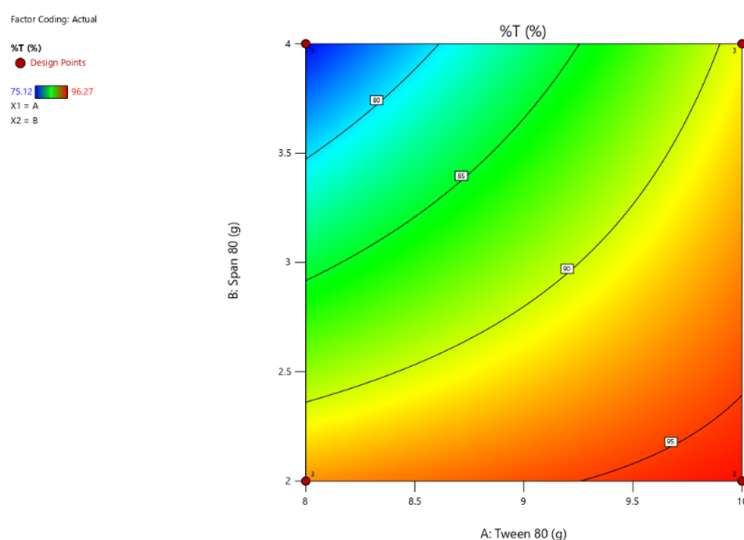


Figure 3. Contour plot showing the influence of Tween 80 and Span 80 concentrations on percent transmittance of red apple emulsion.

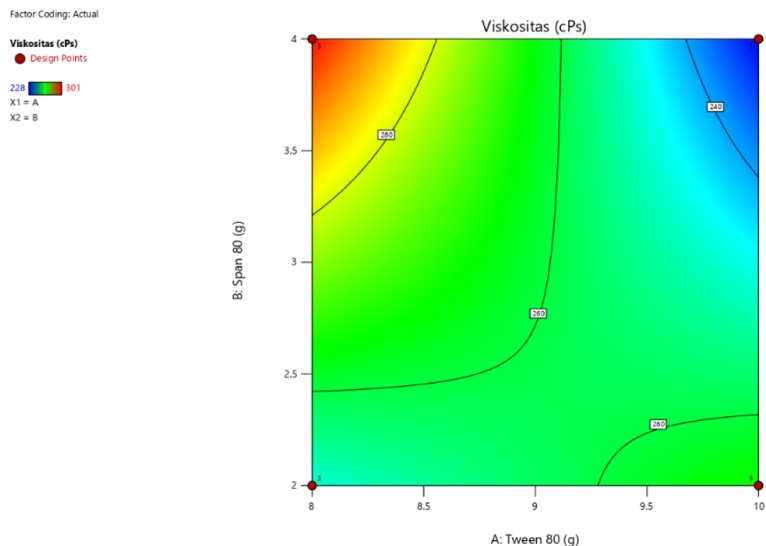


Figure 4. Contour plot depicting the combined effect of Tween 80 and Span 80 concentrations on emulsion viscosity.

Regression equations and contour plots precisely interpret the relationship between factors, levels, and response values by quantitatively describing trends within the experimental domain. The percent transmittance response from Tween 80 (X_1) and Span 80 (X_2) levels in factorial design follows: $Y = 88.8308 + 4.5775X_1 - 5.80917X_2 + 3.1875X_1X_2$. Y represents percent transmittance, X_1X_2 signifies the Tween 80-Span 80 interaction term, and the equation demonstrates linear dependencies on individual factor and their interaction, supporting predictive applications through targeted factor substitution in formulation development.

Measurement results indicated that formula 1, A, and AB fell within the acceptable transmittance range, indicate good optical clarity and homogeneity of the emulsion system. Statistical analysis of formulation effects on transmittance responses revealed significant ANOVA model differences ($p < 0.0001$). Span 80 was the dominant factor influencing transmittance (F-value = 8581.11; %contribution = 52,00%), primarily increasing clarity values. The interaction effects of both surfactants on transmittance responses are depicted in **Figure 3**.

Viscosity

Viscosity measures the internal resistance of a fluid to flow. This resistance arises from frictional forces between the fluid’s constituent molecular volumes, which depend on molecular interactions within the system. In emulsions

system, viscosity is primarily influenced by temperature, droplet size, phase concentrations, and the preparation method employed (21, 32). The optimal viscosity range for emulsion formulations is 10-2000 cPs (33).

Regression equations and contour plots provide precise interpretation of factor, level, and response relationships. The factorial design model for viscosity (Y) as influenced by Tween 80 (X_1) and Span 80 (X_2) is $Y = 260.917 - 13.75X_1 + 3.25X_2 + 22.0833X_1X_2$. Whereas, X_1X_2 signifies the Tween 80-Span 80 interaction term, and the equation demonstrates linear dependencies on individual factors and their interaction, supporting predictive applications through targeted factor substitution.

Statistical analysis of formulation effects on viscosity responses revealed significant ANOVA model differences ($p < 0.0001$). The interaction between Tween 80 and Span 80 exerted the dominant influence on viscosity (8788.12), contributing 70.91% to the response variation. The combined effects of both surfactants on viscosity responses are illustrated in **Figure 4**.

Formulation Evaluation

A 2^2 factorial design evaluated red apple emulsions using Tween 80 (8 g and 10 g) and Span 80 (2 g and 4 g) concentrations. Three key responses were assessed: pH, %T, and viscosity as presented in **Table 2**. Data analysis were performed using Design Expert® software, to generate regression models and response surface plots.

Table 2. Statistical analysis of physical properties.

Formula	pH		Transmittance (%)		Viscosity (cPs)	
	^a F value	^b p value	^a F value	^b p value	^a F value	^b p value
Model	231.94	< 0.0001	5497.59	< 0.0001	4123.79	< 0.0001
Tween 80	515.05	< 0.0001	5328.10	< 0.0001	3403.12	< 0.0001
Span 80	171.43	< 0.0001	8581.11	< 0.0001	190.12	< 0.0001
Tween 80*Span 80	9.33	0.0157	2583.55	< 0.0001	8778.12	< 0.0001

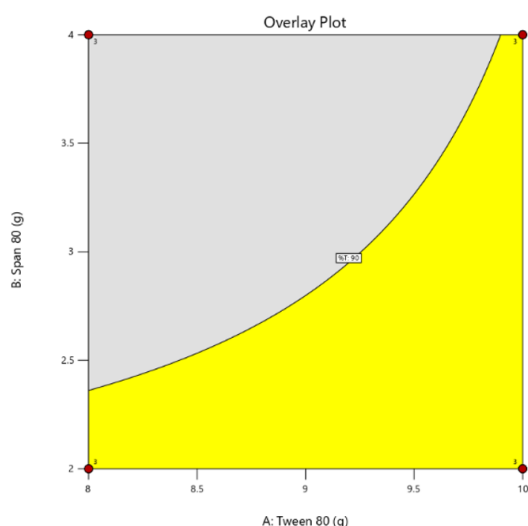
Description: ^aF value determined the variation has the significant impact, two-way ANOVA; ^bp value determined the significant values, two-way ANOVA

Table 3. Red apple extract emulsions physical properties before and after the cycling test.

Formula	Before the Cycling Test			After the Cycling Test		
	*pH	*%T	*Viscosity (cPs)	*pH	*%T	*Viscosity (cPs)
F1	5.40 ± 0.006	93.25 ± 0.262	249.33 ± 0.577	5.39 ± 0.015	92.85 ± 0.084	250.67 ± 0.577
FA	5.20 ± 0.017	96.03 ± 0.216	266.00 ± 1.000	5.00 ± 0.010	90.09 ± 0.085	304.67 ± 4.041
FB	5.47 ± 0.012	75.26 ± 0.135	300.00 ± 1.000	4.88 ± 0.025	14.11 ± 0.472	396.00 ± 6.083
FAB	5.32 ± 0.015	90.79 ± 0.235	228.33 ± 0.577	5.11 ± 0.020	80.38 ± 0.418	285.33 ± 5.686

Table 4. Diameter of red apple extract emulsion inhibition zone against *Cutibacterium acnes*.

Formula	Inhibition Zone (mm)			Zone of Inhibition (mm)	Antibacterial strength criteria
	R1	R2	R3		
Optimal formula (F1)	18.00	16.00	17.00	17.00 ± 1.000	Strong
Positive control (Clindamycin)	23.00	22.00	23.00	22.70 ± 0.577	Strong
Negative control (aquadest)	0	0	0	0 ± 0.000	No inhibition zone

**Figure 5.** Overlay plot of Tween 80 and Span 80 concentrations showing optimal regions for pH, percent transmittance, and viscosity responses.

The overlay plot presented in **Figure 5** illustrates the acceptable response region by the design model. The plot suggests that only formulations F1, FA, and FAB fall within the acceptable range, whereas formulation FB exhibits a transmittance value outside the acceptable limits. Consequently, selection of the best formulation should be based on the lowest surfactant and co-surfactant composition lying within the predicted acceptable region to maximize material efficiency.

Accelerated stability testing

Stability testing was performed using the cycling test method over 3 cycles. Each cycle comprising storage at alternating temperature of -15°C and 25°C for 48 h per condition (25). This procedure was intended to simulate extreme storage conditions that may affect the physical integrity of the emulsion system. Following completion of the 6 cycles, evaluations of organoleptic, pH, %T, and viscosity were conducted on the emulsions formulation. These parameters were selected to monitor potential physical and

chemical instability after temperature stress. The results before and after the cycling test stability are presented in **Table 3**.

It demonstrates that pH and transmittance values decreased significantly ($p < 0.05$), whereas viscosity increased significantly ($p < 0.05$) across all formulations except F1 ($p > 0.05$). The accumulation of unfrozen solutes with increasing cycles accounts for the observed shifts in pH, and viscosity of the formulations (34). Besides, accelerated stability testing using centrifugation method was conducted, revealing no phase separation or flocculation, thereby indicating its stability. Only formulation F1 successfully passed both the cycling test and centrifugation test, demonstrating physical stability. Consequently, F1 was selected as the best formula for subsequent evaluation on its *Cutibacterium acnes* inhibitory activity.

Antibacterial Activity

Antibacterial testing of optimal formula was performed to assess its efficacy against *Cutibacterium acnes* and to determine the appropriate dosage for emulsion preparation. This evaluation provided preliminary insight into the biological performance of the selected formulation. The assay employed the agar well diffusion method under aseptic condition, with sterility controls indicating absence of bacterial growth and clarity of the medium. All procedures were carried out in triplicate to ensure the reliability and reproducibility of the obtained results. Anti-*Cutibacterium acnes* activity testing outcomes are summarized in **Table 4**.

Conclusion

The best red apple extract emulsion (F1) exhibited remarkable physicochemical stability and substantial antibacterial activity. The O/W emulsion obtained possess the potential for effective delivery of the hydrophobic apple extract. However, further investigation is necessary to fully ascertain this. This natural formulation demonstrates considerable promise and potential for advancement in the field of nanotechnology.

Declarations

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Conflict of Interest

The authors declare no conflicting interest.

Data Availability

All data generated or analyzed during this study are included in this published article.

Ethics Statement

Ethical approval was not required for this study.

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Supplemental Material

The supplemental material can be found at the following links:

(<https://view.officeapps.live.com/op/embed.aspx?src=https://etflin.com/file/document/202601220700232042342377.docx>)

,
(<https://view.officeapps.live.com/op/embed.aspx?src=https://etflin.com/file/document/20260122070023536049166.docx>),
(<https://view.officeapps.live.com/op/embed.aspx?src=https://etflin.com/file/document/20260122070023403924151.docx>).
These supplementary files correspond to Paired T Test_pH, Paired T Test_Viscosity, and Paired T Test_%T, respectively.

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Additional Information

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