

Bioactivity Guided Antidiabetic Formulation Development of *Tridax procumbens* Linn Leaves

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ABSTRACT

Purpose: Diabetes mellitus is a global health problem as it is one of the complex disease and main causes of premature illness and death. The present study was focused to scientifically overcome bioavailability drawbacks of *Tridax procumbens* Linn. and improve its medicinal use of for the treatment diabetes by formulating it into microemulsion form. In this study oil in water microemulsion of *Tridax procumbens* Linn leaves extract was prepared using olive oil, Tween 80 and propylene glycol for oral delivery. **Methods:** Phytochemical screening of *Tridax procumbens* Linn. leaves extract was carried out to isolate flavonoids by column chromatography. The o/w microemulsion of whole extract was prepared by water titration method followed by its evaluation for various physical tests and thermodynamic stability. The optimized formulation was tested for drug release *in vitro* and antidiabetic activity by *in vivo* study. **Results:** *Tridax procumbens* Linn. leaves extract isolated for the active moiety showed antidiabetic action in rats. The optimized *Tridax procumbens* Linn. extract microemulsion showed an average particle size 292.4 nm. There was approximately 1.4 fold increase in drug release from microemulsion compared to marketed tablet formulation. At the end of seven days after oral administration of microemulsion a significant reduction in blood glucose level was observed. **Conclusion:** The optimized microemulsion was thermodynamically stable. The enhanced bioavailability of flavonoids in microemulsion proves the traditional claim with regard to *Tridax procumbens* Linn. for its antidiabetic activity.

Key words: *Tridax procumbens*, Microemulsion, Alloxan-induced diabetes, Hypoglycemic activity, Bioavailability.

INTRODUCTION

Diabetes mellitus is the world's third-largest, chronic, non-infectious disease affecting 425 million people globally and this figure is projected to increase up to 629 million by 2045 as per the report published by International Diabetes Federation.¹ Diabetes is one of the complex disease and main causes of premature illness and death. Screening, diagnostic and therapeutic efficacy that favorably affect health outcomes along with cost-effective treatment for diabetes patient is the major focus area for the researchers.² A wild plant named *Tridax procumbens* Linn. commonly known as "Ghamra" and in English "coat butt" belongs to family Asteraceae observed throughout India.³ It has been used since many years

as an indigenous medicine for a variety of ailments and traditionally plant leaves powder has been used since decades in treating diabetic patients in Mandesh Maharashtra and the tribal populations of Udaipur district in Rajasthan.⁴ It has been also reported as anticoagulant, antifungal, hair growth promoter and has insect repellent and wound healing activities in Indian traditional phytotherapy.⁵ It has been reported in literature that orally administered alcoholic extracts of *Tridax procumbens* Linn. leaves at 200 mg/kg dose produces a substantial decrease in blood glucose level in human over the period of 7 days.⁶ TPE contains quercetin as active principles which protects beta cells against

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damage by its anti-inflammatory, anti-apoptotic and antioxidant properties. It regenerates beta cells through stimulation of the ductal stem cell. Thus, quercetin been reported as an adjuvant drug in the management of diabetic mellitus.⁷ The poor solubility of quercetin and crystalline form at body temperatures limit its bioaccessibility and its bioavailability which limit out its use for diabetes treatment.⁸

Natural form of the TPE can be delivered via novel drug delivery system which will be efficient and effective and will have few side effects as compare to synthetic drugs. The term “microemulsion” refers to a thermodynamically stable isotropically clear dispersion of two immiscible liquids, such as oil and water, stabilized by an interfacial film of surfactant molecules.⁹ Microemulsion offer several advantages as drug delivery systems as these are thermodynamically stable along with enhancement of solubility and the bioavailability of less soluble drugs. The oil component penetrate and hence swell the tail group region of the surfactant monolayer leads to solubilization of lipophilic drug and surfactants and co-surfactants lowers interfacial tension to aid dispersion processes and also provide a flexible film that can readily deform around small droplets. Appropriate hydrophile-lipophile character will provide the correct curvature at the interfacial region for the desired microemulsion.¹⁰ The present study was focused to scientifically overcome bioavailability drawbacks of *Tridax procumbens* Linn. and improve its medicinal use for the treatment of diabetes by formulating it into microemulsion from.

MATERIALS

Tridax procumbens Linn. was collected at Kandalgaon Dist. Kolhapur India. Botanical identification was performed by Dr. Yadav, Department of Botany Shivaji University Kolhapur. Quercetin, alloxan monohydrate, polyethylene glycol 400, polysorbate 80 and olive oil were received as gift samples from Loba Chem Pvt. Ltd. (Mumbai, Maharashtra, India). The water was obtained from standard sources.

Animals

Wistar rats weighing 200-250g of both sex were used for the animal experiments and were maintained under standard laboratory conditions at $25 \pm 2^\circ/50 \pm 15\%$ RH and at normal photo period (12 hr dark/12 hr light). Commercial pellet diet [Chakan oil mill, Sangli, India] and water was provided *ad libitum*.¹¹

METHODS

Extraction procedure

The leaves of *Tridax procumbens* Linn. were dried under shade and ground in an electrical grinder to a coarse powder. Powdered leaves (2.5 kg) were extracted for 24 hr with water: ethanol at 70:30 and dried under reduced pressure on rotary evaporator.¹²

Preliminary phytochemical studies

The ethanolic extract of *Tridax procumbens* Linn. leaves was screened by phytochemical test for the presence of flavonoids, alkaloids, phenols, saponins, triterpenoids, glycosides and tannins using modified methods.¹³

Identification of Flavonoids

TLC technique was used to determine the presence of flavonoids in the extract. Toluene, ethyl acetate, formic acid at 5:4:0.2 proportions were used respectively for identification of flavonoids.¹⁴ The extract was run on silica gel 60 F₂₅₄ pre-coated plate with 0.2 mm thickness. Visualization was carried out in iodine chamber by dipping the plate in it, till the color of the spot appears. Retention factor (R_f) was calculated as migration distance of substance to the solvent front.

Isolation of active flavonoid fraction

Ethanolic extract, 10 g, was subjected to column chromatography using 100-200 mesh silica gel (Merck, India) and eluted with 5:4:0.2 solvent system consisting of solvents toluene, ethyl acetate, formic acid to obtain five fractions. Each fraction was isolated twice using TLC and UV spectroscopy, separately, to confirm the presence of active Fraction, Table 1. This process was repeated several times with bulk quantity of the samples until the expected amount of flavonoids have been obtained.¹⁴

Pre-formulation studies

Solubility study

In order to prepare microemulsion, extracted oil and surfactants were selected on the basis of solubility study using equilibration method. A known amount of the

Table 1: R_f value of extract.

Compound	R_f value
Quercetin	0.7
Fraction 1	0.34
Fraction 2	0.5
Fraction 3	0.66
Fraction 4	0.6
Fraction 5	0.5

TPE was dissolved in each of the oils, surfactants and co-surfactants followed by vortexing in stoppered vials and finally shaking at 100 rpm for 72 hr on an orbital shaker at $25 \pm 1^\circ$. These mixtures were further centrifuged for 15 min at 3000 rpm. The supernatant was filtered through 0.45 μm membrane filter. The dissolved drug was diluted within the linearity range and was measured spectrophotometrically using UV-spectrophotometer (Shimadzu UV-1700) at 254 nm.¹⁵

Drug-excipient compatibility studies

FTIR analysis was used to study drug-excipient interaction by scanning samples in the range of 400-4000 cm^{-1} . The surfactant, co-surfactant mixture, oil alone and in combination with extract was analyzed. A spectral comparison was done with FTIR of the pure drug to understand significant functional groups of the drug interacting with the excipients.¹⁶

Optimization of microemulsion components

A series of mixtures were prepared by changing the ratio of optimized oil, surfactant and co-surfactant. These mixtures were prepared and subjected to a simple water titration method. Surfactants and co-surfactants with ratios 1:1, 1:2, 1:3 were mixed together to get different mixtures and fixed volume of oil were added into these mixtures and subjected to gentle shaking and vortexing. The final mixture was titrated against the water with constant stirring using magnetic stirrer. The disappearance of turbidity in the solution was identified as the end point of the titration. Mixtures were kept at room temperature under observation for one day. Milky white and turbid mixtures get separated and specified as unstable emulsions thus termed as biphasic. The clear, transparent or translucent, low viscous mixture indicates the monophasic nature of the system.¹⁴

Monophasic mixtures were prepared with different ratios of oil, surfactant, co-surfactant and water. To construct pseudoternary phase diagram (CHEMIX® School software) these monophasic mixtures were plotted on three diagonals of the triangle. Resulted triangle's shaded area represents monophasic emulsion and considered as the area of a stable microemulsion. All titrations were repeated thrice to get reproducible results. Extract loaded microemulsion prepared by solubilizing extract in the oil and further following the same procedure.

Evaluation of Microemulsion System

Thermodynamic stability studies

TPE subjected for thermodynamic stability with the help of stress tests applied in series. In this test, TPE

microemulsion was checked for thermodynamic stability upon centrifugation at 3000 rpm for 30 min. Stable formulations that did not cream, crack or separate into two layers, were further subjected for the next test. The stability of the formulations at varied temperatures was tested by storing them at 45° and 4° each for a period of 48 hr. The samples were exposed to heating and cooling cycles which was repeated six times for each sample. Formulations which were stable with reference to their physical integrity at these temperatures were exposed to freeze-thaw cycle that involved storing formulations between -21° and $+25^\circ$ each for a minimum period of 48 hr. Thermodynamically stable formulations were selected for further optimization.¹⁷

Physicochemical characterization

Conductivity measurement

Electrical conductivity (σ) of the TPE microemulsion was studied using conductivity meter. Measurements were done to study the effect of the amount of water on the nature of microemulsion and to determine the type of a system; oil-continuous, bi-continuous or water continuous.¹⁸

pH

The pH of TPE loaded microemulsion was measured using a calibrated pH meter at $25 \pm 1^\circ$. The measurements were performed and recorded in triplicate.¹⁵

Viscosity

Viscosity affects the interaction between the different molecules and ultimately effects on the friction in between the molecules present in the fluid. Viscosity measurement of the microemulsion was carried out in triplicates with help of Brookfield viscometer (DV-III+, Programmable Rheometer).⁷

Droplet size, polydispersity index and zeta potential measurement

The droplet size, polydispersity index and zeta potential of the optimized formulation was measured by dynamic light scattering (DLS) using Malvern Zetasizer (Malvern 2000 Hydro SM) and Zetasizer (Micro ZS).

In vitro diffusion study

In vitro release studies gives us information about the amount of drug released from the formulation. It was performed using a dialysis bag method with molecular weight cutoff of 12,000 Da. Dialysis membrane retains microemulsion and molecules released over the time into sodium citrate buffer as with pH 5 used as dissolution medium. Microemulsion was transferred and tightly tied into the dialysis bag which was placed

in a vessel containing dissolution medium in dissolution test USP Type II apparatus rotating at 100 rpm. Aliquots were withdrawn at time intervals of 1 h up to 8 h and analyzed spectrophotometrically using UV visible spectrophotometer (Shimadzu UV-17009) at 254 nm.¹⁹

Accelerated stability study

The accelerated stability study of optimized TPE microemulsion formulation was carried out as per ICH guidelines. Samples of optimized TPE loaded microemulsions were kept in the refrigerator and programmable environmental chamber for three months at $5 \pm 2^\circ$, $25 \pm 2^\circ$ and $60 \pm 5^\circ$ relative humidity (RH) and elevated temperature $40^\circ \pm 2^\circ$ and 75 ± 5 RH. Optimized TPE formulations were withdrawn at 1, 2 and 3 months storage time from the chamber and evaluated for particle size and zeta potential using Malvern instruments (Malvern 2000 Hydro SM) and Zetasizer (Micro ZS), respectively.¹⁴

Determination of drug content

The amount of *Tridax procumbens* Linn present in the optimized microemulsion was determined by UV spectroscopic method. To a 100 ml volumetric flask 1 ml microemulsion was transferred followed by addition of a sufficient amount of methanol and the mixture was shaken. The resulting solution was filtered through a 0.45 μ m Whatman filter paper. Exactly 1 ml of filtrate was transferred to volumetric flask; volume was made to 10 ml by adding methanol and scanned on UV spectrophotometer at 254 nm.

In vivo study

Dried ethanolic extract of *Tridax procumbens* Linn. leaves 200 mg/kg was suspended in 0.5% sodium carboxy methyl cellulose and used to test its hypoglycemic activity in Wistar rats weighing 200-250 g. Animals were divided into 3 different groups, 6 animals were kept in each group and kept at fasting for 18 hr before the experiment. The first control group was given normal saline followed by the second group was administered with the standard formulation and third group received test formulation. Diabetes was induced by the intravenous administration of alloxan monohydrate, 100 mg/kg. A blood sample, 2 ml, was collected after 48 h from the orbital sinus and instantly used for the estimation of glucose level. Treatment was given to the only those animals whose glycemic levels were found to be either or to above 200 mg/dl. These animals were given a daily administration of prepared microemulsion and extract of leaves of *Tridax Procumbens* Linn., 200mg/kg for 7 days. All animal groups were treated at the same time in every morning every day

under the same situation. The blood was collected one 1 h after the last administration and blood glucose level was measured.⁶

RESULTS

Preliminary phytochemical studies

Thin layer chromatography

The ethanolic extract of dried leaves of *Tridax procumbens* Linn. was subjected to thin layer chromatography for confirmation of the presence of quercetin flavonoid. The dark brownish yellow color spot appeared under iodine chamber. Pure quercetin and TPE showed R_f value 0.7 and 0.6 respectively, Table 1, indicating presence of flavonoid in extract.

Isolation of active constituent

Five fractions were collected from column chromatography and were analyzed using TLC and UV spectrophotometry at 254 nm for confirmation of flavonoids. The R_f values are presented in Table 1 indicate the fraction No. 3 showed 0.66 which is very close to that of pure quercetin which shows R_f at 0.7 and thus is an active moiety.

Criteria for the selection of components

One of the factors that involved in the choice of components in the microemulsion system is the solubility of the TPE. Therefore, it is important that the TPE must have high solubility in all components used in the microemulsion system. The results of the solubility study showed that olive oil has the highest solubility 4.2 mg/ml among all the other screened oils namely ethyl oleate, oleic acid, iso-propyl myristate used. Tween 80 had more solubility of 17.38 mg/ml as compare to others to the tween 20, span 80 and span 20. Propylene glycol 400 had more solubility 4.5 mg/ml than ethanol and PG. Thus olive oil, tween 80 and propylene glycol 400 were chosen as oil phase, surfactants and co-surfactant, respectively.

Drug excipient compatibility studies

The FTIR spectrum shows the main functional groups present in the TPE. Drug-excipient compatibility studies were performed to eliminate any compatibility issues of any of the components in the microemulsion system with the TPE. The spectrum in Figure 1 indicates that none of the components interfere with the functional group of the drug.

Pseudo ternary phase diagrams

The pseudo ternary phase diagrams were constructed by varying the concentrations of surfactant Tween 80

with a decrease in oil concentration and increase in water concentration.

Droplet size measurement

The measurement of particle size of microemulsion globules are important to ensure that emulsion was micro droplets. Figure 3 reflects microemulsion formed having droplets size 292.4 nm with 0.252 polydispersity index which is optimized formulation (Q2).

Zeta potential of microemulsion

Zeta potential shows the stability of microemulsion, the optimized Q2 formulation shows -21.6 Mv, Figure 4. Aggregation is not expected to happen, due to the slightly negative charge of the droplets. The values of particle size presented in Table 3 indicate that droplet size is small enough and hence has high zeta potential confirming the stability of microemulsion.¹²⁻¹⁴ It indicates that droplets were stable with no aggregation and

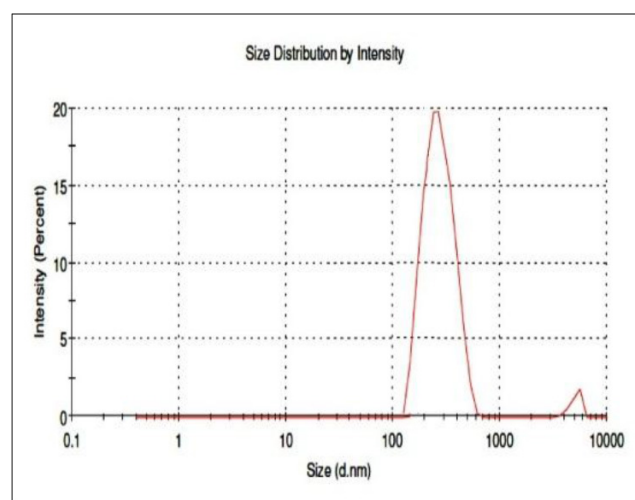


Figure 3: Droplet size measurement of batch Q2.

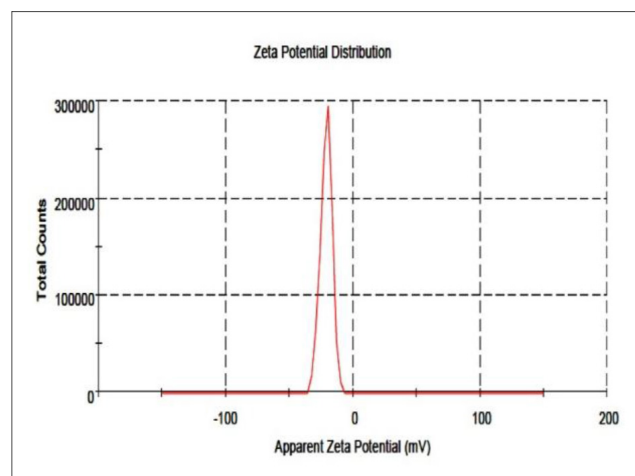


Figure 4: Zeta potential of optimized batch Q2.

Table 3: Droplet size and zeta potential of formulations.

Formulation	Particle Size (d. nm)	PDI	Zeta Potential (mV)	Drug content (%)
Q1	372.7	0.246	-13.6	97.10
Q2	431.1	0.292	-21.6	98.04
Q3	292.4	0.254	-34.5	97.30

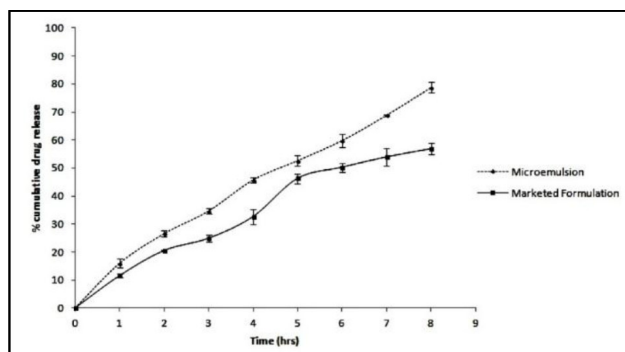


Figure 5: *In vitro* drug release studies of optimized batch Q2 and marketed formulation.

fluctuation in the microemulsion. All batches were having sufficient charge and mobility to inhibit aggregation of particles.

In vitro release studies

Release and diffusion of the drug from the microemulsion were calculated using the data of percentage drug diffused in the given time into the equations of zero order, first order, Higuchi and Korsmeyer pappas model. The *in vitro* drug release studies, Figure 5, shows that the drug release from optimized formulation follows Higuchi matrix model. This was concluded by a higher regression coefficient value in the curve fitting. Drug released from the optimized Q2 formulation and marketed tablet formulation was found to be 80.26% and 59.93%, respectively. Surfactants and co-surfactants solubilize the oil by forming micelles thereby reducing the interfacial tension between oil and water. Thus when TPE is dissolved in oil its solubility improved. Microemulsion showed higher drug release as compared to the marketed formulation which may be attributed to the solubility enhancing component of the surfactant and co-surfactant.

Accelerated stability study

The results of the accelerated stability studies show that the microemulsion did not change significantly in the physical appearance, assay and droplet size. Microemulsion when observed under the microscope

Table 4: *In vivo* experimental study.

Group		I	II	III	IV
Treatment		Control	Test (extract)	Test (formulation)	Standard
Normal Blood Glucose (mg/dl)		75	78	80	84
		82	79	78	79
		85	84	69	82
Diabetic Blood Glucose (mg/dl)		-	287	280	316
			290	285	325
			310	290	298
Blood glucose level (mg/dL)	Day 1	76	272	270	270
		79	265	263	309
		80	300	244	245
	Day 2	75	251	269	215
		72	238	256	295
		82	278	242	222
	Day 3	74	203	201	198
		73	206	200	276
		77	246	182	249
	Day 4	75	138	196	190
		75	200	198	253
		80	219	169	216
	Day 5	76	131	174	180
		74	190	168	234
		75	195	148	186
	Day 6	71	119	132	130
		74	145	156	184
		83	150	125	146
	Day 7	74	102	101	109
		73	103	95	135
		68	123	93	138

there was no droplet aggregation, whereas at the same time the droplets were of uniform size. This indicates the stability of prepared microemulsion.

Determination of drug content

The amount of drug present in the stable microemulsion formulation Q2 was found to be 98.4%.

In vivo experimental study

The results of *in vivo* experimental studies presented in Table 4 reveals that the ethanolic extracts of *Tridax procumbens* Linn. leaves (200 mg/kg) orally administered for 7 days produced a significant decrease in the blood glucose level in the model of alloxan-induced diabetes in rats. The active constituent quercetin has poor bioavailability hence its microemulsion was prepared. The effective surface area of this drug would directly influence its solubility and the dissolution rate. This increased surface area has direct impact on the improved contact with the intestinal mucosa and thus better absorption. Thus microemulsion showed improvement in aqueous solubility and the bioavailability of drug that significantly reduced blood glucose level.

CONCLUSION

The effect of oral administration of ethanolic extracts of *Tridax procumbens* leaves (200 mg/kg) for 7 days produce significant decrease in the blood glucose level in the model of alloxan-induced diabetes in rats. The efficient formation of extract loaded microemulsion with droplet size in the micrometer range has facilitated the incorporation of poorly water soluble extract into the oil in water emulsion system. Effective solubilization was accomplished using the micro-emulsification method. It is evident that this novel formulation improves *in vitro* diffusion and systemic exposure that evidently points towards its better therapeutic efficacy to be used as oral drug therapy.

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CONFLICT OF INTEREST

The authors declare that there is no conflict of interests regarding the publication of this paper.

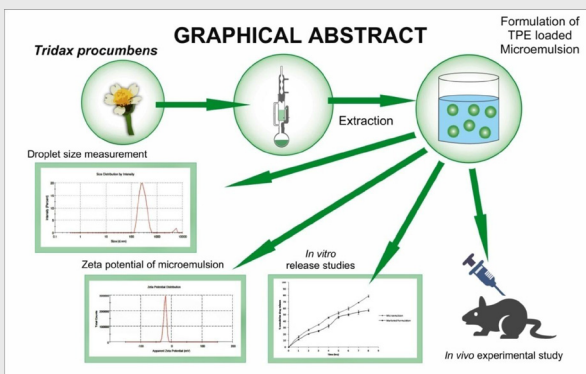
ABBREVIATIONS

UV: Ultra visible; **TPE:** *Tridax procumbens* Linn; **TLC:** Thin Layer Chromatography; **R_f:** Retention Factor; **RH:** Relative humidity; **HLB:** Hydrophilic lipophilic balance; **S_{mix}:** Surfactant; Co-surfactant mixture; **PEG:** Polyethylene glycol; **FTIR:** Fourier transfer infrared.

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PICTORIAL ABSTRACT



SUMMARY

The present study was focused to scientifically overcome bioavailability drawbacks of *Tridax procumbens* Linn. and improve its medicinal use for the treatment of diabetes by formulating it into microemulsion. Water microemulsion of this extract was prepared using olive oil, Tween 80 and propylene glycol for oral delivery to treat diabetes mellitus. The optimized *Tridax procumbens* Linn. extract microemulsion showed an average particle size 292.4 nm. There was approximately 1.4 fold increase in drug release from microemulsion compared to marketed tablet formulation. At the end of seven days after oral administration of microemulsion a significant reduction in blood glucose level was observed. The optimized microemulsion was thermodynamically stable. The enhanced bioavailability of flavonoids in microemulsion proves the traditional claim with regard to *Tridax procumbens* Linn. for its antidiabetic activity.

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