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FORMULATION AND EVALUATION OF INTRANASAL NANOEMULSION CONTAINING RUTIN

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ABSTRACT

Rutin-flavonoid-polyphenolic has gained attention in prevention of brain cancer. The low permeability of Rutin (RU) across the blood-brain-barrier (BBB) leads to its insufficient delivery which in turns result in low therapeutic index. Therefore, developing a novel approaches enhancing the CNS delivery of RU are required for the treatment of Cancer. The aim of this research work was to develop in Nanoemulsion (NE) loaded with RU, for CNS targeting. Rutin (RU) is a poorly water soluble anticancer drug, with oral bioavailability is about 2%. Nanoemulsion (NE) were fabricated by Vortexing technique. Oleic acid was used as oil. Tween 80 was employed as surfactant and Polyethylene glycol 400 was employed as co-surfactant. RU loaded NE for intranasal delivery are considered as promising vehicle for its targeting to CNS to treat the brain cancer.

KEYWORDS

Intranasal delivery, Nanoemulsion, Brain targeting and Vortexing technique.

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INTRODUCTION

The uncontrolled growth of cells and tissue can arises cancer, cancer was a one of the most distressing and life threading disease that serves date worldwide. Cancer like Brain tumors were an abnormal and uncontrolled growth of cells in brain. Modern colloidal nanoparticulate system was novel approach to overcome the problems of chemotherapy. Intranasal drug delivery was the promising strategies for direct deliver drug in nose to brain by passing the BBB viaol factory and trigeminal nerve pathways. Rutin (RU) (polyphenolic compound) has potent antimetastatic and antiproliferative activity against brain tumors. It was important to suppression of nuclear factor-kB is

responsible for tumor proliferation. RU having a promising ability to inhibit angiogenesis, it is process for formation of new blood cell in blood vessel for tumor growth, RU having ability to stop the new blood cell formation in blood vessels responsible for tumor growth and shows antiangiogenic activity. The goal of this study was formulate Nanoemulsion (NE) containing RU for intranasal (nose to brain) delivery to central nervous system (CNS) for the treatment of brain tumor[1-2].

MATERIAL AND METHODS

Material

Rutin (RU) was a gift from Loba Chemie Ltd. (Mumbai, India). Oleic acid, Tween 80 and polyethylene glycol 400 (PEG 400) were purchased from Loba Chemie Ltd. (Mumbai, India).

FORMULATION OF NANOEMULSION (NE)

Selection of Excipients for formulation

The solubility of Rutin (RU) in various oils (castor oil, oleic acid, ethyl oleate, soya oil, coconut oil, oleic acid and clove oil), surfactants (Tween 20 and Tween 80) and co-surfactant (polyethylene glycol 400, polyethylene glycol 200 and propylene glycol) was determined by using Screeningtechnique[3].

Preparation of Nanoemulsion (NE)

Rutin-Nanoemulsion (RU-NE) were prepared by Vortexing technique (Low energy emulsification technique)[4] by slowly pouring the oil, surfactant and co-surfactant mixture using Vortex mixer (Sphinix Ltd, India) into aqueous phase. RU (100mg) was dissolved in mixture of Oleic acid (mL), Tween 80 (mL) and PEG 400 (mL) was slowly added with stirring at 600rpm using magnetic stirrer and formulation composition was reported in Table No.1.

PHYSICOCHEMICAL CHARACTERIZATION OF RUTIN LOADED NANOEMULSION (RU-NE)

Droplet size analysis

The droplet size (DS) were determined by photon correlation spectroscopy (PCS) using a Malvern Zetasizer (Nano ZS 90, Malvern Ltd., Malvern, UK). The measurement using PCS is based on the light-scattering phenomena in which the statistical intensity fluctuations of the scattered light from the particles in the measuring cells are measured. Prior to the measurements, all samples were diluted with double-distilled water to produce a suitable scattering intensity the light scattering was monitored at 25˚C at a 90˚C cangle[5].

Zeta Potential

The ZP, reflecting the electric charge on the droplet surface and indicating the physical stability of colloidal systems, was measured by determining the electrophoretic mobility using the Malvern Zetasizer (Nano ZS 90, Malvern Ltd., and Malvern, UK). The measurements were performed following dilution in double-distilled water. It was measured using the Dip cell by applying a field strength of 20 V/cm and the average of the ZP was given from 30 runs[6].

Drug Content

The drug content of formulation was determined by UV spectrophotometric method. RU from NE formulations (N3) was extracted by dissolving 1 ml of NE in methanol. RU content in the Methanolic extract was analyzed spectrophotometrically (UV 1700, Shimadzu, Japan) at 257 nm, against the standard Methanolic solution of RU.

In vitro Drug permeation studies

In vitro diffusion study of optimized NE (N3) was carried out by Franz diffusion cell having 2.0 cm diameter and 25 ml capacity. Dialysis membrane (Himedia) having molecular weight cut off range 12000-214000 kDa was used as diffusion membrane. Pieces of dialysis membrane were soaked in phosphate buffer saline (PBS) pH 6.4 for 24 h prior to experiment. Diffusion cell was filled with PBS pH 6.4 and dialysis membrane was mounted on cell. The temperature was maintained at 37°c. After a pre-incubation time of 20 minutes, the NE equivalent to 10 µg of RU (N3) was placed in the donor chamber. Samples were periodically with drawn from the receptor compartment for 4 hours and replaced with the same amount of fresh PBS, and assayed by a UV spectrophotometer at 257nm.

RESULTS AND DISCUSSION

Preparation and characterization of Nanoemulsion (NE)

The RU-loaded NE were prepared using Vortexing technique. For the preparation of oleic acid as a liquid lipid. Tween 80 were selected as a surfactant and Polyethylene glycol 400 as a co-surfactant a stabilizer, respectively.

Droplet size analysis and Zeta Potential

The droplet size (nm) and zeta potential (mV) of RU-loaded NE (N3) was found to be 46.85 nm and -37.28 mV respectively.

Drug Content

The concentration of oil and surfactant: co-surfactant was important effect on drug content. The oil content (oleic acid) was increases, to increase drug content and the surfactant and co surfactant concentration (tween 80 + PEG 400) decreases to increase drug content. Because drug having maximum solubility in oil phase and drug content of optimized formulation (N3) was found to be 99.88%.

In vitro Drug permeation studies

The release profile of RU-loaded NE (N3) through the dialysis membrane in PBS (pH 6.4) was found to be 99.87 %. The release pattern of optimized NE (N3) appears to be fast release with negligible burst effect.

Table No.1: Compositions of RU-NE formulations

S.No	Formulation batches	Con. of Oil (Oleic acid) (mL)	Con. of Surfactant (Tween 80) (mL)	Con. of Co-surfactant (PEG 400) (mL)	Droplet size in (nm)
1	N1	3	12	8	119.45
2	N2	3	10	6	147.58
3	N3 (op)	2	6	4	46.85
4	N4	5	8	9	126.25
5	N5	5	10	9	138.26
6	N6	6	10	8	107.48

CONCLUSION

RU have various activities as it may be anticancer, antioxidant, anti-inflammatory drug lipophilic in nature having low oral Bioavaibility, is selected as candidate for the development of RU-NE for its intranasal delivery to target the CNS viaol factory and trigeminal nerve pathway for the treatment of brain tumor.

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CONFLICT OF INTEREST

We declare that we have no conflict of interest.

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