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An Overview of Transferosomes: A Transdermal Drug Delivery System

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Abstract: A lipid bilayer including phospholipids, an edge activator, and an ethanol/aqueous core makes up transferosomes, often referred to as transferosomes, which are ultra-deformable vesicles for transdermal applications. Compared to oral and needle-based methods, this approach has a number of benefits, including self-administration, non-invasiveness, significantly reduced hepatic clearance of the medication, and increased patient compliance. The primary reason for the development of various drug delivery systems was that due to the first pass metabolism, adverse and side effects, nonpatient compliance and invasive procedures, the efficient therapeutic outcomes. The advantage of transferosomes is they are barely invasive without the first pass metabolism. Transferosomes also has several advantages than liposomes due to the active surfactant properties. Due to its penetration enhancer properties, it enables the drug molecule to penetrate into the skin via the stratum corneum than other formulations. So, more research works are encouraged with respect to tranferosomes in order to formulate various new drugs in this type of drug delivery system.

Keywords: Transferosomes, Transdermal drug delivery system, Ethosomes, Invasomes and Barriers.

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INTRODUCTION

A lipid bilayer including phospholipids, an edge activator, and an ethanol/aqueous core makes up transferosomes, often referred to as transferosomes, which are ultra-deformable vesicles for transdermal applications. Compared to oral and needle-based methods, this approach has a number of benefits, including self-administration, non-invasiveness, significantly reduced hepatic clearance of the medication, and increased patient compliance. In addition to being uncomfortable, needle-based procedures produce medical waste that, if improperly disposed of, could endanger public health. For instance, the spread of harmful illnesses like HIV through the

reuse of needles, particularly in developing nations [1]. The word transferome was first set as a brand by the IDEA, a German organization. The 'Transfer some' comes from the Latin word 'Transferre' which means 'to transfer'. In Greek, it means 'body'. Cevc & Blume was the first persons to identify this technology in 1991. They are also involved in the research and published several patents over 3 decades. The primary reason for the development of various drug delivery systems was that due to the first pass metabolism, adverse and side effects, non-patient compliance and invasive procedures, the efficient therapeutic outcomes. The advantage of transferosomes is they are barely invasive without the first pass metabolism [2, 3].

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Advantages of Transferosomes

- 1. Theycan be used for systemic and topical drug delivery system.
- 2. As they are made from natural phospholipids similar to liposomes they are biocompatible and biodegradable.
- 3. They have been widely used as a carrier for various proteins, anti-cancer drugs, anti-fungal drugs, analgesics, anaesthetics, corticosteroids, sex hormone, insulin, albumin etc. for their excellent distribution property [4].

Barriers

- 1. The high cost of raw ingredients and expensive manufacturing equipment has an impact on the cost of the final medication product. As a result, phosphatidylcholine is the most commonly used lipid component due to its low cost in the formulation of transferosomes [5].
- 2. The difficulty in getting pure natural phospholipids is another hurdle to using transferosomes as a drug delivery mechanism. As a result, synthetic phospholipids may be used as a replacement [6].
- 3. Transfersomes are assumed to be chemically unstable due to their proclivity for oxidative degradation. By storing a product at a low temperature and keeping it away from light, transfersome oxidation can be significantly decreased. Post-preparation processing such as freeze-drying and spray-drying can improve transferosome storage performance [7].

Tranferosomes VS Liposomes

- 1. Liposomes are bi-layered lipid vesicles whereas transferosomes are second generation elastic lipid vesicles which act as carriers [8, 9].
- 2. Liposomes contain phospholipids and cholesterol where transferosomes contains phospholipids and edge activator surfactant [10].
- 3. In cases of flexibility liposomes are rigid in nature whereas transferosomes have high deformability due to the surfactant [11].
- 4. In liposomes permeation mechanism involves diffusion, fusion and lipolysis and in transferosomes deformation of vesicles occur [12].
- 5. Liposomes are administered through oral, parenteral and transdermal whereas transferosomes are administered through topical and transdermal [13].
- 6. Liposomes cannot penetrate into the deeper skin whereas transferosomes are stable in gel form only as it causes irritation due to presence of surfactants [14, 15].

Transferosomes under Clinical Trial

Several formulations based on transfersomes are presently undergoing evaluation at various phases of

clinical trials. For instance, a phase III clinical trial was conducted to examine the efficacy and safety of ketoprofen integrated in transfersomes (Diractin®) for the treatment of osteoarthritis in the knees. Over a sixweek treatment period, it has been demonstrated that the medication encapsulated in transfersomal carriers demonstrates higher therapeutic effectiveness in treating knee osteoarthritis pain compared to a placebo and comparably fewer side effects[16]. Similarly, under early-stage clinical trials, the topical administration of insulin-loaded transfersomes (Transfersulin®) for hypoglycemic effects is being studied. In the preclinical investigation, it was discovered that Transfersulin® reduced blood glucose levels in rabbits with alloxaninduced diabetes [17]. A randomized controlled trial was conducted to evaluate the risk-benefit ratio of topical triamcinolone acetonide in transfersomes versus commercially available triamcinolone acetonidecontaining cream and ointment. It has been determined that the risk-benefit ratio of topical triamcinolone acetonide may be considerably enhanced by transfersomes [18]. As a result, transfersomes are recognized as the most exceptional and revolutionary transdermal medication carrier available today [19].

Birth of Ethosomes and Invasomes

By experimenting with vesicle compositions, ethosomes and invasomes were developed in response to the good findings observed with transfersomes. Ethosomes are vesicles that contain phospholipis, water, and a comparatively high percentage of ethanol (20– 50%). Ethamomes have high quantities of ethanol, which changes the skin's lipid bilayer and improves the vesicles' capacity to penetrate the stratum corneum [20, 21]. Phospholipids, terpenes, and ethanol combine to form the elastic phospholipid-based vesicles known as invasomes. By breaking the stratum corneum's tight lipid packing, terpenes and ethanol have demonstrated the capacity to potentially enhance medication penetration [22, 23].

Transferosomes for Susceptible Diseases

Transferosomes is used for the treatment of various susceptible diseases due to its formulation in the form of transdermal delivery system. It is used for the delivery of various drug such as insulin, anesthetics, NSAIDs, interferons, anti-cancer and herbal drugs. It is also used for the delivery of peptides and proteins. It is also used for the drugs with sustained release to improve the therapeutic efficacy of various drugs [24].

CONCLUSION

It is concluded that the discovery of transferosomes have paved a way for the treatment of many conditions and diseases by formulating various drugs in the form of transferosomes. It also makes the patient comfortable with minimal invasive procedures and eliminating adverse effects, drug interactions and some other complications. Transferosomes also has several advanatages than liposomes due to the active surfactant properties. Due to its penetration enhancement properties, it enables the drug molecule to penetrate easily into the skin via the stratum corneum than other formulations. So, more research works are encouraged with respect to transferosomes in order to formulate various new drugs in this type of drug delivery system.

REFERENCES

- 1. Miller, M. A., & Pisani, E. (1999). The cost of unsafe injections. *Bulletin of the world health organization*, 77(10), 808.
- Chaurasiya, P., Ganju, E., Upmanyu, N., Ray, S. K., & Jain, P. (2019). Transfersomes: a novel technique for transdermal drug delivery. *Journal of drug delivery and therapeutics*, 9(1), 279-285.
- Modi, C. D., & Bharadia, P. D. (2012). Transfersomes: new dominants for transdermal drug delivery. *Am J Pharm Tech Res*, 2(3), 71-91.
- Nanda, A., Nanda, S., Dhall, M., & Rao, R. T. (2005). A novel ultra-deformable vesicular carrier for transdermal. *Drug Deliv*, 5, 395.
- Verma, P., Ram, A., Jha, A. K., Mishra, A., & Thakur, A. (2010). Phosphatidylcholine: a revolution in drug delivery technology. *International Journal of Pharmaceutical Sciences and Research (IJPSR)*, 1(2), 1-12.
- 6. Chien, Y. W. (1982). Novel drug delivery systems 1982 New York Marcel Decker Inc, 149–215.
- Rahmi, A. D., & Pangesti, D. M. (2018). Comparison of the characteristics of transfersomes and protransfersomes containing azelaic acid. *Journal of Young Pharmacists*, 10(2s), S11. doi: 10.5530/jyp.2018.2s.3.
- Grit, M., & Crommelin, D. J. (1993). Chemical stability of liposomes: implications for their physical stability. *Chemistry and physics of lipids*, 64(1-3), 3-18.
- Rother, M., Lavins, B. J., Kneer, W., Lehnhardt, K., Seidel, E. J., & Mazgareanu, S. (2007). Efficacy and safety of epicutaneous ketoprofen in Transfersome®(IDEA-033) versus oral celecoxib and placebo in osteoarthritis of the knee: multicentre randomised controlled trial. *Annals of the rheumatic diseases*.
- 10. Kumar, A., Pathak, K., & Bali, V. (2012). Ultraadaptable nanovesicular systems: a carrier for systemic delivery of therapeutic agents. *Drug discovery today*, *17*(21-22), 1233-1241.
- Fesq, H., Lehmann, J., Kontny, A., Erdmann, I., Theiling, K., Rother, M., ... & Abeck, D. (2003). Improved risk-benefit ratio for topical triamcinolone acetonide in Transfersome® in comparison with equipotent cream and ointment: a randomized controlled trial. *British Journal of Dermatology*, 149(3), 611-619.
- Cevc, G. (2002). Transfersomes: Innovative transdermal drug carriers. In: Rathbone M.J., Hadgraft J., Roberts M.S., editors. Modified-Release Drug Delivery Technology. 1st ed. Volume 126. Marcel Dekker, Inc.; New York, NY, USA, 533–546.

- 13. Touitou, E. (1996). U.S. Patent No. 5,540,934. Washington, DC: U.S. Patent and Trademark Office.
- Touitou, E., Dayan, N., Bergelson, L., Godin, B., & Eliaz, M. (2000). Ethosomes—novel vesicular carriers for enhanced delivery: characterization and skin penetration properties. *Journal of controlled release*, 65(3), 403-418.
- Shah, S. M., Ashtikar, M., Jain, A. S., Makhija, D. T., Nikam, Y., Gude, R. P., ... & Fahr, A. (2015). LeciPlex, invasomes, and liposomes: A skin penetration study. *International journal of pharmaceutics*, 490(1-2), 391-403.
- Qadri, G. R., Ahad, A., Aqil, M., Imam, S. S., & Ali, A. (2017). Invasomes of isradipine for enhanced transdermal delivery against hypertension: formulation, characterization, and in vivo pharmacodynamic study. *Artificial cells*, *nanomedicine, and biotechnology*, 45(1), 139-145.
- Sudhakar, C. K., Upadhyay, N., Jain, S., & Charyulu, R. N. (2012). Ethosomes as non-invasive loom for transdermal drug delivery system. *Nanomedicine and drug delivery*, 11(10), 2557.
- Mishra, V., Bansal, K. K., Verma, A., Yadav, N., Thakur, S., Sudhakar, K., & Rosenholm, J. M. (2018). Solid lipid nanoparticles: Emerging colloidal nano drug delivery systems. *Pharmaceutics*, 10(4), 191.
- Paliwal, S., Tilak, A., Sharma, J., Dave, V., Sharma, S., Verma, K., ... & Sadhu, V. (2019). Flurbiprofen-loaded ethanolic liposome particles for biomedical applications. *Journal of microbiological methods*, *161*, 18-27.
- Raj, R., Raj, P. M., & Ram, A. (2018). Nanosized ethanol based malleable liposomes of cytarabine to accentuate transdermal delivery: formulation optimization, in vitro skin permeation and in vivo bioavailability. *Artificial cells, nanomedicine, and biotechnology*, 46(sup2), 951-963.
- El Maghraby, G. M. M., Williams, A. C., & Barry, B. W. (1999). Skin delivery from ultradeformable liposomes: refinement of surfactant concentration. *J. Pharm. Pharmacol*, *51*, 1123-1134.
- 22. Abdulbaqi, I. M., Darwis, Y., Khan, N. A. K., Assi, R. A., & Khan, A. A. (2016). Ethosomal nanocarriers: The impact of constituents and formulation techniques on ethosomal properties, in vivo studies, and clinical trials. *International journal of nanomedicine*, 2279-2304.
- 23. Verma, P., & Pathak, K. (2010). Therapeutic and cosmeceutical potential of ethosomes: An overview. *Journal of advanced pharmaceutical technology & research*, 1(3), 274.
- Fernández-García, R., Lalatsa, A., Statts, L., Bolás-Fernández, F., Ballesteros, M. P., & Serrano, D. R. (2020). Transferosomes as nanocarriers for drugs across the skin: Quality by design from lab to industrial scale. *International journal of pharmaceutics*, 573, 118817.

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