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## (54) Title of the invention : FORMULATION AND DEVELOPMENT OF LIPID BASED DRUG DELIVERY SYSTEM OF GLICLAZIDE

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## (57) Abstract :

The present invention relates to the formulation of lipid based drug delivery system of Gliclazide. Formulations were prepared by mixing the drug and carrier using heat fusion method as it is considered as the mechanism for the enhancement of solubility and dissolution of the drug. The in-vitro releases of the different formulations were studied based on the effect of surfactant and oil including their thermodynamic stability. Formulated drug and adjuvants were characterized by spectrophotometry (UV, FTIR and photon correlation). Dissolution studies showed that F3 had the smallest particle size of 127.6 nm, with values for other formulations ranging from 176.8-248.8 nm. The cumulative percentage release for all formulations ranged from  $21.20\pm1.68\%$  to  $80.92\pm3.82\%$ , with F3 having the highest value of  $80.92\pm3.82\%$ . Soya lecithin, soyabean oil and tween 80 showed no significant influence on formulation's stability. These results confirmed that the prepared formulations increases the oral bioavailability

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