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(54) Title of the invention : NANOCARRIER-BASED CHEMICAL COMPOSITION FOR TARGETED THERAPEUTIC MOLECULE DELIVERY

<p>(51) International classification</p> <p>:B82Y5/00, B82Y30/00, B82Y40/00, A61P35/00, A61K31/704, A61K9/51, A61K47/50, A61K47/56</p> <p>(31) Priority Document No :NA (32) Priority Date :NA (33) Name of priority country :NA (86) International Application No : Filing Date :01/01/1900 (87) International Publication No : NA (61) Patent of Addition to Application Number :NA Filing Date :NA (62) Divisional to Application Number :NA Filing Date :NA</p>	<p>(71)Name of Applicant :</p> <p>1)Dr. Harpreet Singh Address of Applicant :Professor, School of Pharmaceutical Sciences, Faculty of Pharmacy, IFTM University, Moradabad - 244102, Uttar Pradesh, India Uttar Pradesh India</p> <p>2)Dr. Nirmala Sisodia 3)Dr. Preeti Chaudhary 4)Dr. Pratik N Chauhan 5)Debadatta Sahoo 6)Dr. Hemlata Bhatt 7)Dr. Ashish Kumar 8)Prof. Prashant Singh 9)Dr. Abhishek Pathak 10)Dr. Kuna Ramu 11)Prof. Kavi Shanker Varshney</p> <p>(72)Name of Inventor :</p> <p>1)Dr. Harpreet Singh 2)Dr. Nirmala Sisodia 3)Dr. Preeti Chaudhary 4)Dr. Pratik N Chauhan 5)Debadatta Sahoo 6)Dr. Hemlata Bhatt 7)Dr. Ashish Kumar 8)Prof. Prashant Singh 9)Dr. Abhishek Pathak 10)Dr. Kuna Ramu 11)Prof. Kavi Shanker Varshney</p>
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(57) Abstract :

The present invention relates to a nanocarrier-based chemical composition for targeted therapeutic molecule delivery, comprising a biodegradable poly(lactic-co-glycolic acid) (PLGA) polymeric core matrix encapsulating doxorubicin hydrochloride at 5-20% w/w with an encapsulation efficiency of not less than 75%. The nanocarrier surface is functionalized with folate-conjugated polyethylene glycol (PEG-folate, MW 2,000-5,000 Da) and anti-HER2 monoclonal antibody fragments to enable dual-ligand active targeting of folate receptor and HER2-overexpressing cancer cells via receptor-mediated endocytosis. The composition exhibits a hydrodynamic diameter of 80-250 nm, a zeta potential of -20 to -30 mV, and pH-responsive drug release releasing less than 15% at pH 7.4 and more than 80% at pH 5.0 within 72 hours. An optional theranostic variant incorporates Fe₃O₄ magnetic nanocrystals at 10-30% w/w enabling simultaneous MRI-guided drug delivery with r₂ of not less than 150 mM⁻¹s⁻¹. The composition achieves 5.6-fold enhanced cytotoxicity in MCF-7 breast cancer cells compared to non-targeted nanoparticles, demonstrating significant therapeutic advancement for precision oncology applications.

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