

Identification and characterization of in vivo, in vitro and reactive metabolites of zorifertinib using Liquid Chromatography Ion Trap Mass Spectrometry

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Zorifertinib is a novel, potent, oral, small molecule used to treat non-small cell lung cancer (NSCLC). zorifertinib is Epidermal Growth Factor Receptor (EGFR) inhibitor and has good blood–brain barrier permeability for (NSCLC) patients with EGFR mutations. zorifertinib is currently at a phase II/III clinical trials. The current research reports the characterization and identification of *in vitro*, *in vivo* and reactive intermediates of zorifertinib. Prediction of susceptible sites of metabolism and reactivity pathways (cyanide and GSH) of zorifertinib were performed by Xenosite web predictor tool. *In-vitro* metabolites of zorifertinib were performed by incubation with rat liver microsomes (RLMs) and isolated perfused rat liver hepatocytes. Extraction of zorifertinib and its in vitro metabolites from the incubation mixtures were done by protein precipitation. *In vivo* metabolism was done by giving single oral dose of zorifertinib (10 mg/Kg) to Sprague Dawely rats in metabolic cages by using oral gavage. Urine was gathered and filtered at specific time intervals (0, 6, 12, 18, 24, 48, 72, 96 and 120 hr) from zorifertinib dosing. A similar volume of ACN was added to each collected urine sample. Both layers (organic and aqueous) were injected into liquid chromatography ion trap mass spectrometry (LC-IT-MS) to detect in vivo zorifertinib metabolites. N-methyl piperizine ring and quinazoline group of zorifertinib undergoe metabolism forming iminium and electro deficient conjugated system respectively, which are very reactive toward nucleophilic macromolecules. Incubation of zorifertinib with RLMs in the presence of 1.0 mM KCN and 1.0 Mm glutathione were made to check reactive metabolites as it is often responsible for toxicities associated with this drug. For in vitro metabolites there were nine *in vitro* phase I metabolites, four in vitro phase II metabolites, eleven reactive metabolites (three cyano adducts, five GSH conjugates metabolites and three methoxy metabolites of zorifertinib were detected by LC-IT-MS. For in vivo metabolites there were eight *in vivo* phase I, ten *in vivo* phase II metabolites

of zorifertinib were detected by LC-IT-MS. *In vitro* and *in vivo* phase I metabolic pathways were N-demthylation, O-demethylation, hydroxylation, reduction, defluorination and dechlorination. *In vivo* phase II metabolic reaction was direct conjugation of zorifertinib with glucuronic acid and sulphate.

Keywords: N-methyl piperizine; zorifertinib; *In vivo* metabolites; *In vitro* metabolites; Cyano conjugates, GSH conjugate.

Biography

Ph.D. graduate in Analytical Pharmaceutical Chemistry, King Saud University. Extensive work experience at modern analytical techniques like triple quadrupole LC-MS/MS (Agilent Triple Quad 6410 LC-QQQ), Ion trap LC-MS (Agilent Ion Trap 6320 LC-MS), MALDI-Orbitrap, and HPLC instruments.Well aware of the analytical methods development and characterization of drug metabolites.Winner of the King Saud University Award for Scientic Excellence as the best research doctoral student for 2018. He has published more than 30 papers in reputed journals and has been serving as an revewer board member of repute. (Up to 100 words)

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