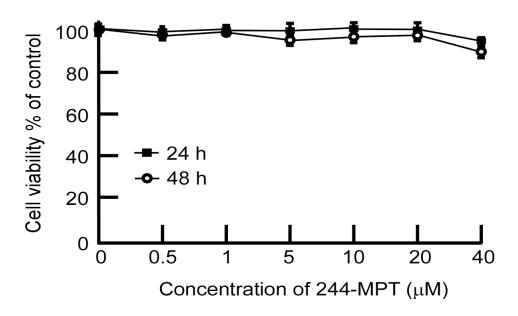
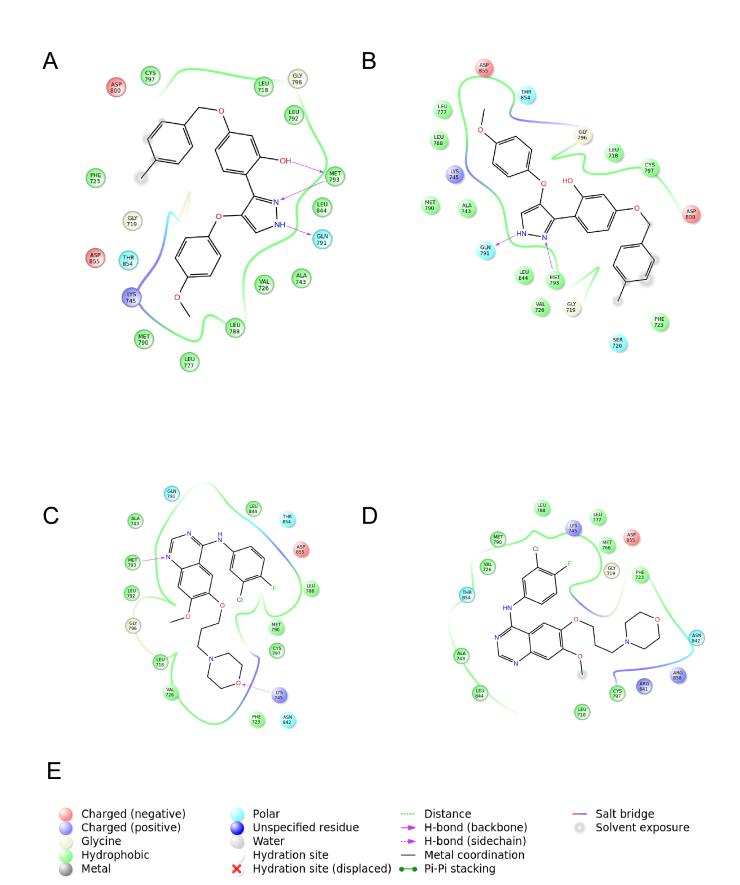
244-MPT overcomes gefitinib resistance in non-small cell lung cancer cells

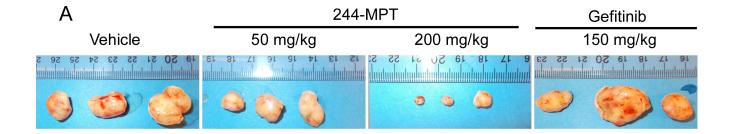
Supplementary Materials

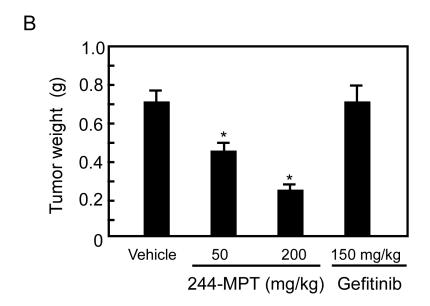


Supplementary Figure S1: 244-MPT is not toxic to cells up to 40 μ M. Cytotoxicity of 244-MPT was measured by MTS assay in MRC-5 cells, a normal lung cell line. The control group was only treated with 0.1% DMSO, the solvent to dissolve 244-MPT. MRC-5 cells were treated with different concentrations of 244-MPT for 24 or 48 h.



Supplementary Figure S2: Ligand interaction diagram of double mutant (L858R/T790M) EGFR ATP-binding pocket with 244-MPT or gefitinib after 5 ns molecular dynamics (MD). Before (A) and after (B) 5 ns MD binding mode of 244-MPT with double mutant (L858R/T790M) EGFR (See also Supplementary Video S1). Before (C) and after (D) 5 ns MD binding mode of gefitinib with double mutant (L858R/T790M). (See also Supplementary Video S2). (E) Figure legend of interaction diagram.





Supplementary Figure S3: 244-MPT inhibits tumor growth in a gefitinib-resistant NSCLC xenograft mouse model. After mice were euthanized, (A) tumor volume and (B) weight were measured. The 244-MPT treated groups exhibited significantly lower tumor volume and weight compared with the vehicle-treated group. Gefitinib treatment had no effect. The asterisk (*) indicates a significant (p < 0.05) decrease in tumor weight of 244-MPT-treated mice compared to vehicle-treated mice.