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Everyone wants to enjoy secure, fast and reliable surfing. Ngagri has worked hard to bring you high-quality service and solutions. Our goal is your success and a beautiful experience! Docking shows binding of azathioprine, but not 6-mercaptopurine, to the active site of mammalian thiopurine methyltransferase. Thiopurine methyltransferase (TPMT) catalyzes the metabolic methylation of the 6-mercaptopurine and azathioprine drugs. Mutations in the gene encoding TPMT are associated with severe toxicity. Therapeutic drugs that are not methylated by this enzyme could be substituted for azathioprine and 6-mercaptopurine during treatment. Pharmacologically active nucleophilic compounds could be used to modulate TPMT activity. To gain information about the possibilities for using this strategy, the binding of azathioprine and 6-mercaptopurine to the active site of TPMT has been investigated. Molecular docking was performed using a homology model of TPMT based on the recently determined X-ray structure of TPMT from *Danio rerio*. Docking studies show that azathioprine binds more favorably than 6-mercaptopurine to the TPMT active site. These results show that the binding of these compounds to mammalian TPMT is feasible, although azathioprine has less favorable binding affinity than 6-mercaptopurine. Although the docking study agrees with experimentally determined activity against cancer cell lines, the calculated binding affinities do not predict TPMT activity correctly. This result could be due to the limitations of both the homology model and the docking protocol, as well as the need for a mammalian TPMT structure for validating the docking results. To further improve the prediction of TPMT activity, the effect of other

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ligands known to be metabolized by this enzyme should be investigated. Development and validation of a monoclonal antibody-based sandwich enzyme-linked immunosorbent assay to detect human placental lact

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