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Formation of a C-Nucleophilic Adduct of Cationic Salicylic Acid Derivatives with Molecular Iodide. [1-(2-Cyano-phenyl)-3-(2,4-dinitrophenyl)urea-acetonitrile]chloride, [ $\alpha$ -2,4-dinitrophenylurea-acetonitrile-3-cyanophenolate], is a product of an in vitro chemistry reaction of the cationic salicylic acid derivative, [2,4-dinitrobenzoyloxyphenyl]-acetyl-L-cysteine hydrochloride, (DNB-chl), with iodide. When DNB-chl is added to a solution of tetrabutylammonium iodide (TBAI), DNB-chl reacts with iodide anion to produce the cationic salicylate, [ $\alpha$ -2,4-dinitrophenylurea-acetonitrile-3-cyanophenolate-N,O]TBA<sup>+</sup>, which is a potent inhibitor of mammalian cystine-glutamate antiporter. In contrast, a reaction of iodide with 2,4-dinitrophenylurea-acetonitrile, a non cationic salicylate analog of DNB-chl, does not lead to the formation of the salicylate ion. A preliminary mechanistic