

# Sulfur Dioxide-Halide Ion Complexes: A Crystallographic Investigation of Bonding

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Sulfur dioxide (SO<sub>2</sub>) is a major air pollutant, and SO<sub>2</sub> emissions result in both acid rain and atmospheric particulates. Little attention has been paid to a relatively simple method for reversibly binding SO<sub>2</sub>, a method which could potentially be used to remove it from waste streams. There is only one example of an “[SO<sub>2</sub>Cl]” anion currently identified in the Cambridge Structural Database (April 2018). The number of “[SO<sub>2</sub>...Cl]” adducts described is also few (less than 10) and the criteria for bonding in such adducts has not been well described. The reaction of SO<sub>2</sub> gas and halide salts was thus used to generate a series of halosulfite (SO<sub>2</sub>X<sup>-</sup>, X = Cl, Br, I) compounds. The structures of the solid compounds obtained, three tetraphenylphosphonium complexes, [PPh<sub>4</sub>][SO<sub>2</sub>X], and [IMesH][SO<sub>2</sub>Cl] (IMesH = 1,3-bis(2,4,6-trimethylphenyl)-imidazolium), were determined using X-ray crystallography. Preliminary investigations suggest that these same products can also be formed *via* solvent-free, solid-gas reactions. It was found that the complexes readily released SO<sub>2</sub> when heated, suggesting that they should be candidates for SO<sub>2</sub> capture.