(c) The representation of diffusion and deposition in the model may introduce unrealistic conditions which affect the chemistry in ways which are not representative of the real atmosphere.

## 4.5 Implications and Conclusions

At this time, there are growing indications that, generally in the northeast U.S., the gas-phase reaction pathway accounts for about 25% and the aqueous-phase reaction pathway about 75% of  $H_2SO_4$  formation in the atmosphere which is dry and wet deposited. The gas-phase oxidation of  $SO_2$  is due to reactions with the photochemically generated radicals HO, HO<sub>2</sub>, and RO<sub>2</sub>. On the regional scale, this  $H_2SO_4$  formation pathway suggests the following implications:

- 1. The reaction rates are non-linear with regard to  $SO_2$  because the free radical concentrations are not constant over time and space. The LRT models, therefore, may not correctly predict the quantity and the deposition patterns of  $H_2SO_4$  formed through the gas-phase reactions;
- 2. Since the gas-phase rates are first order in  $SO_2$ , a reduction in  $SO_2$  concentration will result in a direct reduction in  $H_2SO_4$  formed in the gas phase if the free radicals' temporal-spatial concentrations remain constant;
- 3. The rates are first order in free radical concentrations. If a reduction in free radical concentration (through oxidant precursor control) can be obtained, then this will result in a direct reduction in  $H_2SO_4$  formed in the gas phase;
- 4. Reduction in both  $SO_2$  and free radical concentrations will result in a compounded reduction of  $H_2SO_4$  formation in the gas phase; and