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## Sulphur and iron oxidation state in synthetic fluid-saturated melts

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Sulphur is typically a trace element in silicate melts (rarely more than 5000 ppm) and one of the main con-stituents of volcanic gases. It is an important element in magmatic processes for two main reasons. First, S (as sulphide) controls the behavior of several metals in silicate melts. These metals (which in-clude Cu, Ni, Re, Os, Pd, Pt, Au) are either of economic interest or are used as tracers of large-scale Earth processes. Second, explosive volcanic events are capable of releasing large amounts of S into the atmos-phere, significantly affecting global climate due to injection of sulphate aerosols into the stratosphere. It is well established that the sulfur solubility in silicate melts is a function of redox conditions and melt composition. At oxidizing conditions, sulfate is the dominant species whereas sulfide is the dominant species at reducing conditions. Furthermore, the S solubility at reducing conditions is known to strongly correlate with total Fe content. It is still highly debated to what extent the redox equilibria of the heterovalent elements Fe and S interact during magmatic processes. Here we present data on the S and Fe oxidation state determined by micro-XANES on synthetic fluid and sulfur-saturated hydrous glasses quenched from 200 MPa and 1050°C. Studied compositions range from basalt to rhyodacites. The sulfur oxidation state shows a sharp transition from sulfide to sulfate between oxygen fugacities between QFM and QFM+1 for all studied compositions. The Fe oxidation state varies between 8% and 30% ferric iron in the range QFM-2 to QFM+2 for basaltic and andesitic compositions. In comparison, the Fe oxidation state of the rhyodacitic composition is slightly offset to higher ferric iron contents.