## Abstract Submitted for the TS4CF08 Meeting of The American Physical Society

Raman study of supported molybdenum disulfide single layers WILLIAM DURRER, FELICIA MANCIU, Physics Department, University of Texas at El Paso, PAVEL AFANASIEV, GILLES BERHAULT, Institut de Recherche sur la Catalyse et l'Enironnement de Lyon, IRCELYON, CNRS - University of Lyon, F-69626 Villeurbanne, France, RUSSELL CHIANELLI, Materials Research and Technology Institute, University of Texas at El Paso, TX 79968 — Owing to the increasing demand for clean transportation fuels, highly dispersed single layer transition metal sulfides such as  $MoS_2$ -based catalysts play an important role in catalytic processes for upgrading and removing sulfur from heavy petroleum feed. In its crystalline bulk form, MoS<sub>2</sub> is chemically rather inactive due to a strong tendency to form highly stacked layers, but, when dispersed as single-layer nanoclusters on a support, the  $MoS_2$  becomes catalytically active in the hydrogenolysis of sulphur and nitrogen from organic compounds (hydrotreating catalysis). In the present studies alumina-supported  $MoS_2$  samples were analyzed by confocal Raman spectroscopy. Evidence of peaks at  $152 \text{ cm}^{-1}$ ,  $234 \text{ cm}^{-1}$ , and  $336 \text{ cm}^{-1}$ , normally not seen in the Raman spectrum of the standard bulk crystal, confirms the formation of single layers of  $MoS_2$ . Furthermore, the presence of the 383 cm<sup>-1</sup> Raman line suggests the trigonal prismatic coordination of the formed MoS<sub>2</sub> single layers. Depending on the sample preparation method, a restacking of  $MoS_2$  layers is also observed, mainly for ex-thiomolybdate samples sulfided at 550 °C.

> William Durrer University of Texas at El Paso

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