

Novel Cage Clusters of MoS₂ in the Gas Phase

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Laser evaporation of MoS₂ nanoflakes gives negatively charged magic number clusters of compositions Mo₁₃S₂₅ and Mo₁₃S₂₈, which are shown to have closed-cage structures. The clusters are stable and do not show fragmentation in the post-source decay analysis even at the highest laser powers. Computations suggest that Mo₁₃S₂₅ has a central cavity with a diameter of 4.5 Å. The nanosheets of MoS₂ could curl upon laser irradiation, explaining the cluster formation.

Introduction

Inorganic molecular structures such as polyoxomolybdates are fascinating not only because of their architecture,¹ but also because of their ability to form self-organized structures.² In recent years, large cage structures known as inorganic fullerenes (IFs) have become an active subject of investigation.³ IFs were originally made from layered metal chalcogenides, MX₂, where M = Mo, W and X = S, Se.⁴ Later, layered metal halides such as NiCl₂, CdCl₂, and TiCl₃⁵ and oxides such as TiO₂ and V₂O₅ were also shown to produce IFs.⁶ Boron nitride also forms IFs.⁷ The hollow cage structure of IFs imparts elasticity, and the materials can work as novel lubricants under various conditions.^{4f} The sheetlike structure of these chalcogenides, similar to graphite, suggests that a mechanism similar to that of fullerenes and carbon nanotubes may be operative in the formation of IFs. The unsatisfied valencies present at the periphery of the layered structure makes it unstable at finite dimensions. For the layered structure of graphite to fold and form cages or closed cylinders, high energy is needed, which can be provided by laser vaporization,⁸ arc discharge,⁹ or other means. Ultrasound irradiation can be used for the synthesis of IFs.¹⁰ In all methods described above, the IFs formed were in the solid state. A recent report suggested that the formation of IFs in liquid medium was achieved by laser ablation.¹¹ Formation of onions and nanorods of GaS by laser ablation was reported recently.¹² Zak et al.¹³ reported the synthesis of MoS₂-based IFs in the gas phase, which were subsequently analyzed in the solid state. The encapsulation of WC into an IF of WS₂ was reported, showing that IFs are cage-like.¹⁴ Structures such as octahedron, dodecahedron, and triangular prism have been proposed,¹⁵ but a single molecular cage has not been reported so far.¹⁶ We explored the possibility of making magic cage clusters containing Mo, W, and S in the gas phase. Initial results of this investigation are reported here.

Experimental and Computational Details

Bulk MoS₂ and WS₂ were ground well to form particles of 20-μm diameter, dispersed in acetone, and spotted on the target plate of a Voyager DE PRO Biospectrometry Workstation (Applied Biosystems) matrix-assisted laser desorption ionization time-of-flight (MALDI-TOF) mass spectroscopy (MS) instrument. A pulsed nitrogen laser of 337 nm was used, and the TOF was operated in the delayed extraction mode. For fragmentation by post-source decay (PSD), as well as clustering studies, we used a timed ion selector. Isotope pattern was clearly discernible in all the cluster peaks, and a comparison of the expected and observed patterns was used to assign the peaks. The cluster peaks were significantly higher in intensity when nanosized flakes of MoS₂ were used, which were obtained by sonicating a well-ground MoS₂ sample in acetone for 10 min. A blue-colored dispersion of nanoflakes was formed, which was directly spotted on the target plate, giving a deep blue film upon drying, and laser desorption ionization (LDI) mass spectra were taken from the film. The blue material obtained by evaporation of the solvent in the dispersion was used for various spectroscopic, microscopic, and diffraction studies using standard instrumentation.

To confirm the stability of observed clusters and to understand their structures, we carried out nonempirical calculations¹⁷ of the quantum mechanical ground state of electrons based on density functional theory with a generalized gradient approximation to the many electrons exchange correlation energy given by the Perdew–Burke–Ernzerhof (PBE) functional.¹⁸ We used a standard plane-wave code PWSCF 2.0.1¹⁷ with ultrasoft pseudopotentials¹⁹ to represent the interaction between ions and electrons. An energy cutoff of 25 Ry was used on the plane-wave basis used in the representation of wave functions; the cutoff was 150 Ry for the representation of density. Structures of clusters were determined through total energy minimization using the Broyden–Fletcher–Goldfarb–Shanno (BFGS) algorithm and Hellmann–Feynman forces on atoms. In this procedure, we used different polyhedral structures as initial guesses for smaller clusters and various cage structures of sulfur atoms

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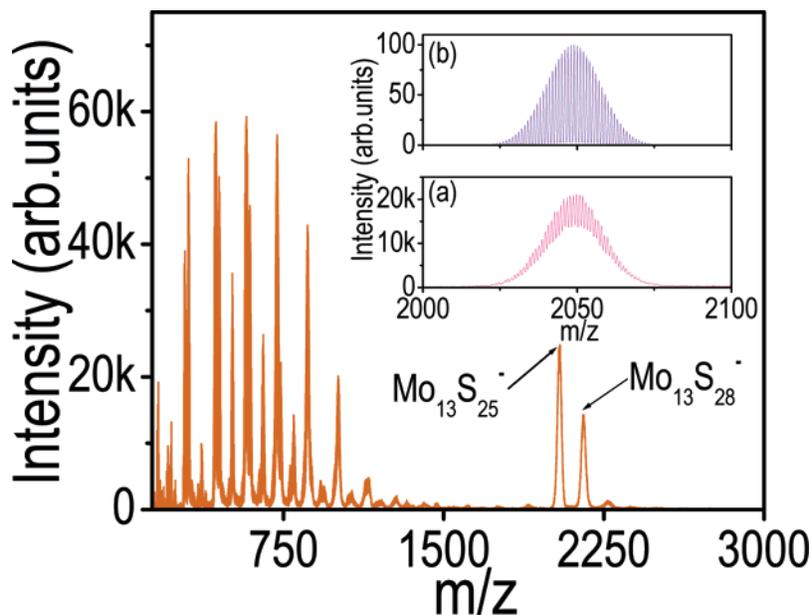


Figure 1. LDI mass spectrum of MoS₂ nanoflakes in the negative mode. Inset: Experimental spectrum taken from nanoflakes (a) shows the expected isotope distribution for Mo₁₃S₂₅⁻ (b).

with Mo atoms located on an inner shell at positions with four- to sixfold sulfur coordination as initial guesses for larger clusters. A large supercell was used with periodic boundary conditions with a vacuum of at least 10 Å separating periodic images and a single *k*-point (0, 0, 0) as Bloch vector.

Results and Discussion

The negative-ion mass spectrum of the MoS₂ clusters obtained from the blue film is shown in Figure 1. The clusters formed are MoS₂⁻, MoS₃⁻, MoS₄⁻, Mo₂S₃⁻, Mo₂S₅⁻, Mo₂S₆⁻, Mo₂S₈⁻, and so on. Isotope patterns matched exactly for all the clusters up to *m/z* ~2000. Even in very pure samples of MoS₂ and WS₂, M_{*n*}O_{*y*}⁻ clusters were observed in poor intensity, possibly due to the presence of oxide impurities in the material. The spectra exhibit a trend: The intensity increases starting from MoS₂⁻, reaches a maximum around Mo₄S₇⁻, and decreases thereafter. Beyond this region, the spectrum is featureless till *m/z* 2000, and suddenly, a magic peak appears with a maximum at *m/z* 2049, followed by another peak at *m/z* 2144. The isotope pattern of the peak at *m/z* 2049 is shown in the inset, which matches with the expected pattern of Mo₁₃S₂₅⁻. On the basis of the expected patterns and the mass numbers, we assigned the magic peaks to Mo₁₃S₂₅⁻ and Mo₁₃S₂₈⁻, respectively. Following these, there are some less intense peaks attributed to clusters of Mo₁₄ and Mo₁₅.

When a finely ground sample dispersed in acetone is spotted on the sample plate, two regions are seen. A typical view of the sample spot as seen by a video camera is shown in Supporting Information. The center region is composed of bulk MoS₂, which is black in color. The periphery is composed of several blue spots due to the nanoflakes. We get a spectrum similar to the one shown in Figure 1 when the flakes are irradiated.

The stability of the ions was checked by PSD. The ion Mo₁₃S₂₅⁻ did not show any fragmentation (Figure 2). Even at the highest laser power, no fragmentation was observable. However, Mo₁₃S₂₈⁻ showed (inset of Figure 2) two fragment peaks at Mo₁₃S₂₆⁻ (*m/z* 2081) and Mo₁₁S₁₉⁻ (*m/z* 1664). All the smaller clusters such as Mo₃S₇⁻, Mo₄S₆⁻, and so on showed well-defined fragmentation patterns (data in Supporting Infor-

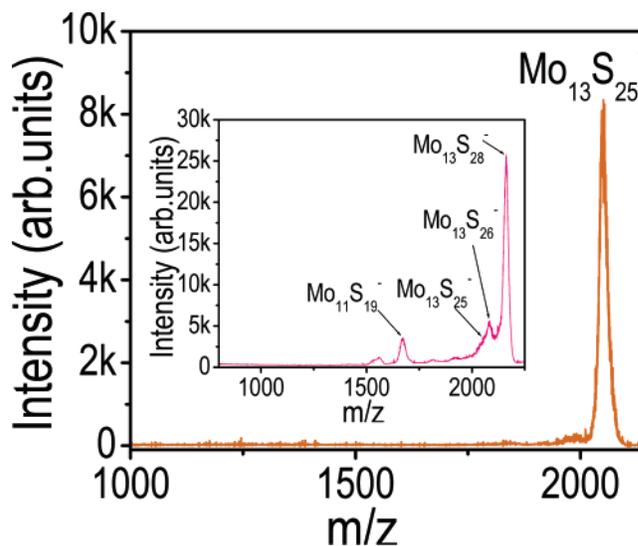


Figure 2. PSD mode LDI mass spectrum of Mo₁₃S₂₅⁻ showing no fragmentation. Inset: PSD mode LDI mass spectrum of Mo₁₃S₂₈⁻ showing fragmentation.

mation). The Mo₁₃S₂₅⁻ and Mo₁₃S₂₈⁻ clusters were unique and were found only in the negative ion mode.

Instead of the flakes, if we irradiate bulk MoS₂, we get a spectrum as shown in Figure 3. Note that the magic peaks are barely detected. Most of the lower clusters are observed, but intensity, in general, is poor.

The blue sample collected, both by scratching the periphery of the sample spot as well as by sonicating bulk MoS₂, was analyzed using UV–visible spectroscopy (UV–vis), powder X-ray diffraction (XRD), transmission electron microscopy (TEM), energy-dispersive X-ray analysis (EDAX), Fourier transform infrared spectroscopy (FT-IR), and Raman spectroscopy under identical conditions. These investigations were done to characterize the nature of the starting material, as other forms of molybdenum blues are known in the literature.¹ The X-ray diffractogram shows the (002) reflection of MoS₂, and Mo and S features in the expected atomic ratio are shown in EDAX (Figure 4a). The material therefore is MoS₂ itself, in the nanodimension with a particle size of ~60–70 nm (Figure 4b).

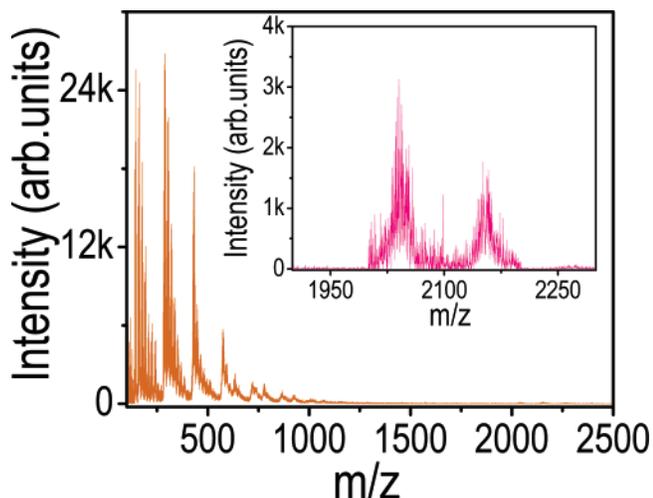


Figure 3. LDI mass spectrum of bulk MoS₂. Inset: Zoomed (25×) portion of the same spectrum showing Mo₁₃S₂₅⁻ and Mo₁₃S₂₈⁻ with very low intensity.

Characteristic E_{2g}¹ and A_{1g} modes of MoS₂ appear in the Raman spectrum at 381 and 406 cm⁻¹, respectively (ref 20). It appears that thinner flakes with fewer layers are easier to get evaporated, which could be curling to form the cagelike structures. All the other data are presented in the Supporting Information.

The possibility of formation of the Mo₁₃ clusters by aggregation of smaller clusters was investigated using the timed ion selector. None of the smaller clusters showed features due to Mo₁₃ (data in Supporting Information). This clearly indicates that the magic clusters around Mo₁₃ are formed by direct laser desorption of MoS₂ layers and not due to aggregation of smaller clusters.

Calculated cohesive energies per atom (Figure 5), while expected to be overestimated, clearly show that Mo₂S₃ and Mo₁₃S₂₅ clusters are relatively more stable than others. Also, we note that the Mo₁₃S₂₅ cluster is most stable among the ones studied here. Energies needed to fragment Mo₁₃S₂₈ and Mo₁₃S₂₅ into the most stable smaller ones such as Mo₂S₃ and Mo₂S₅ are estimated to be 0.45 and 0.57 eV/atom, respectively, according to their stability. In addition, this result demonstrates greater stability for Mo₁₃S₂₅ than Mo₁₃S₂₈, consistent with the experi-

mental observations. As no stable unusual clusters were found in the intermediate region, we did not explore them through computations.

The calculated structure of these clusters exhibits a core of Mo atoms, each with 4–6 neighboring S atoms located in an outer shell, each with 1–3 neighboring Mo atoms. The core of Mo atoms forms a cage with 12 atoms, and the 13th Mo atom is at one end forming a small tetrahedral cap. In the structure, 12 of the Mo atoms have 5 S-neighbors each, while the remaining 1 has 3 S-neighbors. Of the sulfurs, 19 have 3 Mo-neighbors each, 3 have 2 Mo-neighbors each, and the rest (3) have 1 Mo- and 1 S-neighbor each. Average sulfur coordination of Mo in Mo₁₃S₂₅ (Mo₁₃S₂₈) is 4.84 (5.0). The structure of Mo₁₃S₂₅ is like a cage with an almost spherical void space with a diameter of about 4.5 Å (Supporting Information). The Mo₁₃S₂₅ structure has a threefold symmetry (evident in Figure 5), and its core of Mo atoms resembles a curled triangular net of Mo atoms, while that of Mo₁₃S₂₈ has the shape of a short cylinder, revealing significant structural relaxation and deviation from the initial spherical cagelike structure. This indicates that these clusters may have formed by curling of the MoS₂ sheets.

Examination of charge density contours (Supporting Information) reveals mixed ionic and directional bonding between Mo and S atoms. Stability of these clusters arises from mixed ionic-covalent bonding between Mo and S and electrostatic interionic interactions. Comparison of energies of MoS, MoS₂, MoS₃, and MoS₄ clusters shows that the energy of a Mo–S bond is the strongest in MoS₂, while the binding energy per atom is highest for MoS₃. Fragmentation energies of Mo₂S₃ and Mo₂S₅ clusters into MoS and MoS₃ clusters are 2.35 and 1.14 eV/atom, respectively (2Mo₂S₃ = 3MoS + MoS₃ and 2Mo₂S₅ = MoS + 3MoS₃), whereas that for fragmenting Mo₂S₅ and Mo₂S₆ into MoS₂ and MoS₄ (2Mo₂S₅ = 3MoS₂ + MoS₄ and Mo₂S₆ = MoS₂ + MoS₄) are 0.74 and 0.57 eV/atom, respectively. The former are much larger than the latter because MoS itself is not very stable.

In WS₂, magic peaks were found at W₁₂S₂₀O⁻ (*m/z* = 2862) and W₁₃S₂₂⁻ (*m/z* = 3094). There is also an intense ion at W₆S₉O⁻ (*m/z* 1407). These peaks show high stability in PSD (data in Supporting Information). Oxygenation could not be avoided despite repeated baking of the samples in Ar.

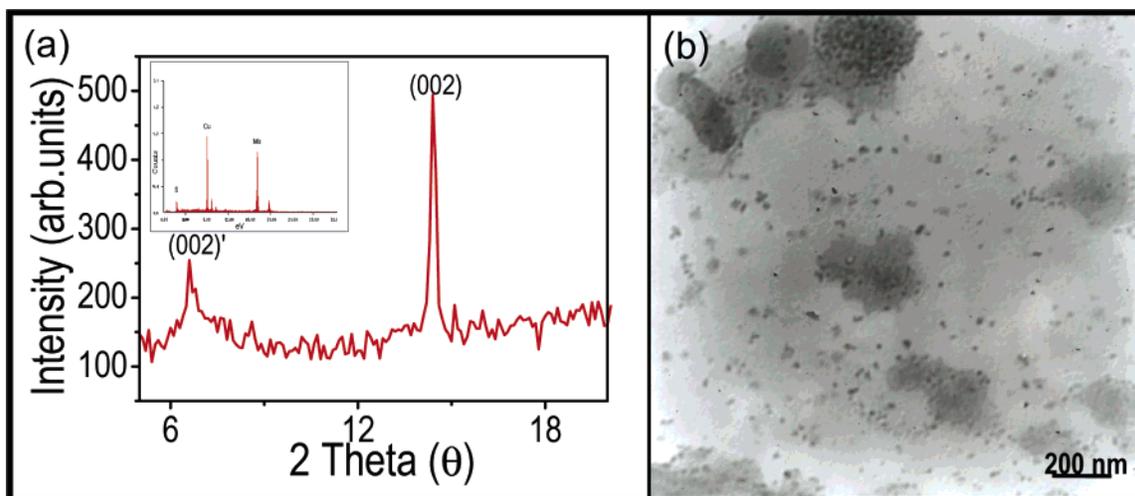


Figure 4. (a) XRD pattern of the acetone extract showing MoS₂ features. The pattern shows the presence of MoS₂ (002) plane (JCPDS no. 87-2416). Inset: EDAX analysis gave Mo and S in the expected ratio. The Cu signal is due to the copper grid used for TEM. (b) TEM of the extract. The material is composed of nanosize flakes of MoS₂. The particle size is estimated to be ~60–70 nm. The analysis was done with a 120 keV Philips CM12 instrument.

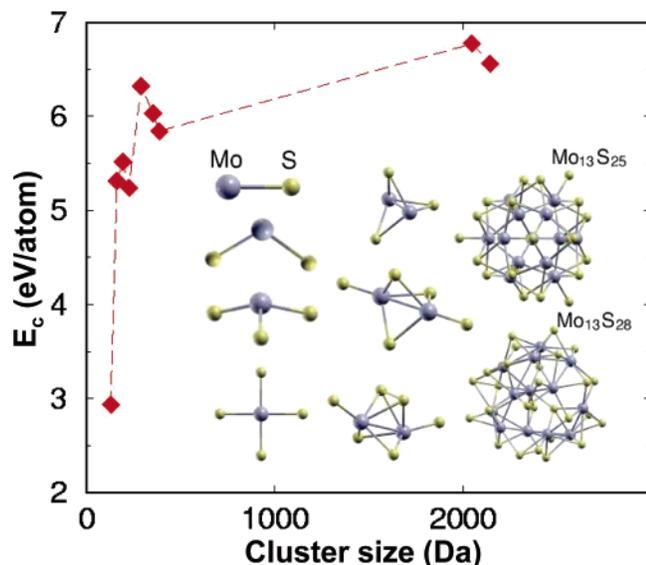


Figure 5. Calculated cohesive energies of Mo_mS_n clusters as a function of their masses. The optimized structures of these clusters are also shown.

In summary, we have shown the existence of stable negatively charged clusters of composition $\text{Mo}_{13}\text{S}_{25}$ and $\text{Mo}_{13}\text{S}_{28}$ in the gas phase. Computations confirm their structural and electronic stability. These clusters are likely to be inorganic fullerenes and may be formed by the curling of nanoflakes of MoS_2 . Considering their stability, they may be formed in the condensed phase under appropriate conditions. Experimental ion mobility experiments may be useful to confirm the proposed structures.

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Supporting Information Available: Schematic view of the sample spot, PSD data of small MoS_2 clusters, their ion selection mass spectra, spectroscopic analysis of MoS_2 nanoflakes,

computational results and WS_2 data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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