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ANNOTATED BIBLIOGRAPHY OF SPECTROPHOTOMETRIC
STUDIES IN THE SYSTEM U-Fe-Se-S-O-H,
1967-78

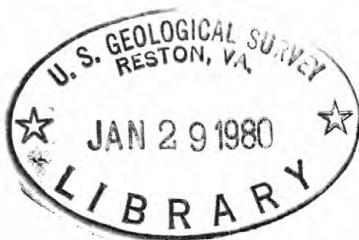
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This report is preliminary and has not been edited or
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INTRODUCTION

Preparation of this bibliography was originally begun to supply background information on previous ultraviolet and visible spectrophotometric studies on various chemical species thought to be important in the formation of sedimentary uranium deposits; we also thought that preparing the bibliography would help direct experimental efforts to determine the association of iron and uranium with several metastable sulfur species. In compiling these references, however, we found that much work had been done on the chemical species of interest by using types of spectroscopy other than ultraviolet and visible spectrophotometry. The dimensions of the bibliography were therefore broadened to include a wide variety of spectral-measuring techniques to make this report useful to a larger audience.

SCOPE OF BIBLIOGRAPHY

This compilation covers the period 1967 through 1978 and is as complete as a reasonable search of the literature would permit. No effort has been made to standardize the units of measurement cited in the annotations, nor has a judgement been made as to the quality of the data.

References listed in this bibliography are limited to spectroscopic studies of chemical species in the U-Fe-Se-S-O-H system. With few exceptions, studies using x-ray or Auger spectroscopy are excluded, and some analytical articles are included. References presenting electronic spectra, which can encompass visible, ultraviolet, or vacuum ultraviolet absorption spectroscopy, are generally included. In the following paragraphs, brief descriptions of the major spectroscopic methods covered by this bibliography are presented.

Ultraviolet and visible spectroscopy is a product of the optical absorbancy of a solution. The intensities of the absorption bands of a solution are used as measures of the concentrations of the various absorbing species present and can be used to obtain information about equilibria in solution. The UV/VIS region of the spectrum ranges from about 200 to 800 nanometers.

Infrared spectroscopy is the absorption of radiation by a molecule that has a changing dipole moment as a result of molecular vibration-rotation. It provides information on bond angles, bond stiffness, and molecular shape.

Moessbauer spectroscopy is the resonance absorption of gamma-ray radiation by a nuclide. Electronic (chemical) differences between the source and sample destroy the resonant condition. Resonance is restored by a relative motion between the source and the absorber (Doppler Shift). The spectrum is the intensity of the gamma-ray signal as a function of this velocity and depends on the molecular structure and oxidation state of the molecule.

Nuclear magnetic resonance spectroscopy (NMR) is the measurement of radio frequencies at which energy is absorbed by a nucleus in an applied magnetic field. The spectra provide information on the location and environment of specific nuclei in a molecule.

Photoelectron spectroscopy is the measurement of the kinetic energy of photoelectrons ejected from a sample by a monochromatic photon source having energies in the x-ray region of the spectrum. From this type of spectra, one can determine ionization potentials and the nature of inner electron orbitals.

Raman spectroscopy results from the interaction of scattered light and a molecule having a changing polarizability because of molecular vibrational-rotational motions. It provides information on bond angles, bond stiffness, and molecular shape.

Microwave spectroscopy results from absorption of microwave radiation inducing changes in rotational energy levels of a gaseous molecule. It provides information on moments of inertia and bond lengths within a molecule.

Electron spin resonance (ESR) or electron paramagnetic resonance (EPR) is the measurement of microwave radiation absorbed by unpaired electrons in an external magnetic field. The spectra provide information on the electronic structure of the molecule.

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Experimental study of the ir transmission spectrum of pure amorphous selenium in the region 0.75-25 μ .
[Translation unavailable to authors. Data taken from Chem. Abs. 73:135530a, 1970.]

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Experimental study on the Raman spectrum of $\text{KH}_3(\text{SeO}_3)_2$ at room temperature as well as the infrared spectra of this crystal both at room temperature and at liquid oxygen temperature. Data are given in tables and on a graph.

Adrian, F.J., Cochran, E.L., and Bowers, V.A., 1967, ESR spectrum of HO_2 in argon at 4.2 K: The Journal of Chemical Physics, v. 47, no. 12, p. 5441-5442.

Experimental study of the electron spin resonance spectrum of HO_2 in argon at 4.2 K with observations on the magnetic Hamiltonian. Data are presented in a figure and a table.

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radical. The spectra of ³²S₀ and ³⁴S₀ are also
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and these are discussed in terms of known crystal
structure. Data are presented in both tables and graphs.

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Experimental study of the infrared and Raman spectra of condensed SO_2 gas at 20 K with comparisons to previous work. Data are presented in a table and typical spectra are shown in a figure.

Anderson, Anthony, and Loh, Y.T., 1969, Low temperature Raman spectrum of rhombic sulfur: Canadian Journal of Chemistry, v. 47, no. 6, p. 879-884.

Experimental study of the Raman spectra of polycrystalline samples of rhombic sulfur at 100 and 300 K, with excitation by the 5145 angstrom line of an argon ion laser. Data are presented graphically and in two tables.

Aring, K., and Sievers, A.J., 1967, Thermal conductivity and far-infrared absorption of UO_2 : Journal of Applied Physics, v. 38, no. 3, p. 1496-1497.

Experimental study of the far-infrared transmission of UO_2 in the frequency range of $5-115 \text{ cm}^{-1}$ at 3 K. Thermal conductivity of the same UO_2 crystal was also measured. Data are presented graphically.

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first-row transition elements: Journal of the Chemical Society, Dalton Transactions, v. 18, p. 1701-1707.

Experimental study of the complex species formed between sulfate ions and several first row transition elements (including Fe II and Fe III) at various ionic strengths. Data are presented graphically and in tables.

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Rotational constants and bond lengths are presented.

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Data are presented on a graph and in the text.

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during the aqueous extraction of the sinter cake obtained in the course of the oxidative sintering of copper electrolytic anode mud with soda. Data are given in tables and on graphs.

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Donovan, R.J., and Little, D.J., 1972, Vacuum ultraviolet spectra of transient molecules and radicals. Part 3.--HSe and HTe: Journal of the Chemical Society, Faraday Transactions. 2 68 pt., v. 10, p. 1812-1818.

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of H_2S : Chemical Physics Letters, v. 13, no. 3, p. 304-306.

Experimental study of the high-resolution photoelectron spectrum of H_2S with calculations of the vibrational structure and bond angle of the first excited state of H_2S^+ . Data are presented graphically.

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Experimental study on the Raman spectra of FeS_2 and other metal sulfides. Data are presented in tables. [Translation unavailable to authors. Data taken from *Chem. Abs.* 71:75975w, 1969.]

Ferraro, J.R., and Fink, Uwe, 1977, Near infrared reflectance spectra and analysis of H_2S frost as a function of temperature: *The Journal of Chemical Physics*, v. 67, no. 2, p. 409-413.

Experimental study on the reflection spectra of H_2S frost in the near infrared region (1.6-4 μm) over the temperature range 62-143 K. Data are given in tables and on a graph.

Fleming, J.W., 1976, Far i.r. rotational absorption spectra of H_2S , SO_2 , NO_2 : *Spectrochimica Acta*, v. 32A, no. 4, p. 787-795.

Experimental study on the far i.r. rotational absorption spectra of H_2S , SO_2 , and NO_2 between 10 cm^{-1} and 40 cm^{-1} at a resolution of 0.05 cm^{-1} with comparisons to earlier studies. Data are presented graphically and in tables.

Fleming, J.W., 1977, Gas phase i.r. absorption spectrometry at extremely long wavelengths (714 μm - 1667 μm): *Spectrochimica Acta*, v. 33A, no. 8, p. 787-791.

Experimental study on the rotational spectra of several gases (including H_2S and SO_2) at extremely long wavelengths using Fourier interferometric spectrophotometry. Data are presented in tables and spectra are shown in several figures.

Fouche, D.G., and Chang, R.K., 1971, Relative Raman cross section for N₂, O₂, CO, CO₂, SO₂, and H₂S: Applied Physics Letters, v. 18, no. 12, p. 579-580.

Experimental study on the Raman cross sections of various gases of interest in atmospheric pollutant studies as measured at 5145 \AA . Data are listed in tabular form.

Fox, M.F., and Hunter, T.F., 1969, Charge-transfer-to-solvent spectra: Nature, v. 223, no. 5202, p. 177-178.

Experimental study on the ultraviolet absorption spectra of simple inorganic anions, including hydroxide, with an explanation for the broad structureless bands characterized as charge-transfer-to-solvent transitions.

Data are presented in a table and a graph.

Furuhashi, Akiko, Noguchi, Kazuo, Kawai, Shigeyuki, and Hayakawa, Yasumasa, 1970, Polarographic and spectroscopic studies on chalcogen dioxides in acid media: Denki Kagaku, v. 38, no. 5, p. 340-343.

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Futekov, L., and Iordanov, N., 1971, [Selenium dioxide-hydrogen chloride-water system studied using Raman spectra with a view to analytical application of the system]: Izvestiia Fizicheskiia Instituts ANEB, Bulgarska Akademii Naukite, Sofia, v. 21, p. 289-292. In Bulgarian.

Experimental study on the Raman spectra of SeO_2 and SeOCl_2 in strongly acidic HCl solutions. [Translation unavailable to authors. Data taken from Chem. Abs. 77:96463s, 1972.]

Garg, C.L., Gupta, S.D., and Narasimham, K.V., 1970, Force constants and bond distances for the U-O bond of uranyl salts: Indian Journal of Pure and Applied Physics, v.8, no. 2, p. 108-110.

Theoretical study on U-O bond distances. Force constants for U-O were calculated using vibrational data from fluorescence and infrared investigations for several uranyl salts. Results are plotted on a graph and listed on a table.

Gerard, A., 1967, [Moessbauer effect and bond nature in some semimetallic iron compounds]: Colloques Internationaux de Centre National de la Recherche Scientifique, No. 157, p. 55-61. In French.

Experimental study on several semimetallic iron compounds (including sulfides and selenides) revealing a Moessbauer spectrum of two lines each arising from quadrupole coupling. Data are presented graphically and in tabular form.

Gerding, H., and Ypenburg, J.W., 1967, Some remarks on the vibrational spectra of liquid and solid sulfur dioxide: Recueil des Travaux Chimiques des Pays-Bas, v. 86, no. 5, p. 458-462.

Tabulation of vibrational spectral data of solid and liquid SO_2 determined from ir and Raman spectral measurements with comparisons to previously published results. Vibrational frequencies of SO_2 in different regions are discussed.

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Experimental study on a newly observed band in the uv absorption spectra of highly alkaline sulfide solutions attributed to the formation of S^{2-} ion. Upper limits for the second dissociation constant of hydrogen sulfide are determined for temperatures as high as 250°C.

Data are presented graphically.

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Experimental study on the equilibrium relations in polysulfide solutions as a function of temperature using absorption spectra. Bisulfide-hydrogen sulfide solutions were used to buffer the pH at various values between 4.7 and 10.6. Data are presented graphically and in the text.

Giguere, P.A., and Herman, Kazimiera, 1970, Studies on hydrogen-oxygen systems in the electrical discharge. IV. Spectroscopic identification of the matrix-stabilized intermediates, H_2O_3 and H_2O_4 : Canadian Journal of Chemistry, v. 48, no. 22, p. 3473-3482.

Experimental study on the infrared absorption of the products from electrically dissociated H_2O and D_2O vapor and other hydrogen-oxygen systems at liquid nitrogen temperature measured between 4000 and 300 cm^{-1} . Data are presented graphically and in tables.

Gillespie, R.J., and Pez, G.P., 1969, Infrared and Raman spectra of the Se_4^{2+} ion: Inorganic Chemistry, v. 8, no. 6,

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Experimental study of the infrared and Raman spectra of a number of compounds containing the polyatomic cation Se_4^{2+} . Data are presented in one table and several figures.

Gorller-Walrand, C., and Vanquickenbourne, L.G., 1971, Identification of the lower transitions in the spectra of uranyl complexes: The Journal of Chemical Physics, v. 54, no. 10, p. 4178-4186.

A qualitative and comparative study of the effects of ligation on the uranyl spectrum. The results are presented in tabular form.

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Compilation of the luminescence spectra of a large variety of uranium minerals plus a tabulation of electron-vibrational frequencies of uranyl in mineral species at 77 K. [Translation unavailable to authors. Data taken from Chem. Abs. 87:104533p, 1977.]

Goulden, J.D.S., and Manning, D.J., 1967, Infra-red spectroscopy of inorganic materials in aqueous solution: Spectrochimica Acta, v. 23A, no. 1, p. 2249-2254.

Experimental study of the ir spectra of several inorganic materials in aqueous solution, including several sulfur species, over the range 1550 to 950 cm^{-1} . Data are presented in both spectra diagrams and tables.

Gribov, L.A., Zubkova, O.B., and Rybal'chenko, I.V., 1977, [One possibility for an approximate calculation of the vibrational spectra of crystals]: Zhurnal Prikladnoi Spektroskopii, v. 27, no. 6, p. 1038-1041. In Russian.

Theoretical study of the vibrational spectra of uraninite (UO_2), and other minerals, using a simplified selection model of the structures studied. [Translation unavailable to authors. Data taken from Chem. Abs. 88:96852v, 1978.]

Gurman, V.S., and Sergeev, G.B., 1970, [Ultraviolet absorption spectrum of a glassy 17.5M solution of hydrogen peroxide in water at 77°K]: Zhurnal Fizicheskoi Khimii, v. 44, no. 3, p. 803-804. In Russian.

Experimental study on the absorption spectrum of a 17.5M solution of H_2O_2 in water from 265-320 nm.

[Translation unavailable to authors. Data from Chem. Abs. 73:9045h, 1970.]

Helminger, Paul, Cook, R.L., and De Lucia, F.C., 1971, Microwave spectrum and centrifugal distortion effects of HDS: Journal of Molecular Spectroscopy, v. 40, no. 1, p. 125-136.

Experimental study on the rotational spectrum of hydrogen deuterium sulfide in the ground vibrational state using microwave techniques. Twenty-seven new transitions were measured in the millimeter and submillimeter wavelength region. Data are presented in tabular format.

Helminger, Paul, De Lucia, F.C., and Kirchhaff, W.H., 1973, Microwave spectra of molecules of astrophysical interest IV. Hydrogen sulfide: Journal of Physical Chemistry Reference Data, v. 2, no. 2, p. 215-223.

Critical review of the available data on the microwave spectrum of hydrogen sulfide. Molecular data such as rotational constants, centrifugal distortion constants, hyperfine coupling parameters, and dipole moments are tabulated.

Helminger, Paul, and De Lucia, F.C., 1975, The ground state rotational spectrum of H₂Se: weighted

microwave-infrared analysis: Journal of Molecular Spectroscopy, v. 58, no. 3, p. 375-383.

Experimental study on transitions of the five major species of hydrogen selenide in the 100-600 GHz region of the microwave spectrum. Data are presented in tabular form.

Helminger, Paul, and De Lucia, F.C., 1977, Pressure broadening of hydrogen sulfide: Journal of Quantitative Spectroscopy and Radiative Transfer, v. 17, no. 6, p. 751-754.

Experimental study on microwave measurements of the self-broadening parameters of four pure rotational transitions of H_2S in the 50-60 GHz and 150-400 GHz regions. Data are given in a table and on graphs.

Henrion, W., 1967, Reflectivity measurements on trigonal selenium single crystals in the spectral region between 1.6 and 6.0 eV: Physica Status Solidi, v. 22, no. 1., p. K33-K37.

Experimental study on Se single crystals grown by sublimation from the vapor phase using absorption reflectivity measurements at 90 and 300 K. Spectra are shown in two figures and data are given in a table.

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spectra of oxides in the region of compositions from
 $\text{UO}_{2.50}$ to $\text{UO}_{2.64}$]: Radiokhimiya, v. 11, no. 4, p.
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Experimental study on the Raman spectrum of K_2SeO_4 .
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Jager, H., Kieschke, H.G., and Lochte-Holtgreven, W., 1968, Absorption spectra of small iron particles dispersed in various gases and observation of absorption bands from highly excited states of the gas molecules: Physics Letters, v. 26A, no. 9, p. 440-441.

Data from previous work on the absorption spectra of small iron particles dispersed in nitrogen, argon, or hydrogen show a continuous absorption which can be fitted to curves calculated according to Mie's theory. In addition, a number of absorption bands due to highly excited molecules of the carrier gas have been described.

Jakobsson, L.R., 1967, The infrared spectrum of the neutral sulphur atom: *Arkiv Foer Fysik*, v. 34, no. 2, p. 19-31.

Experimental study of the infrared spectrum of sulfur over the range 9212 to 34,270 Å by means of a high-frequency discharge through sulfur dioxide. Data are shown in tabular form as well as in three figures.

James, D.W., and Frost, R.L., 1978, Structure of aqueous solutions: *Journal of the Chemical Society, Faraday Transactions 1*, v. 74, n. 3, p. 583-596.

Experimental study of structure making and structure breaking by dissolved electrolytes using infrared librational bands. The concentration dependence of the ir librational band of water was studied for forty-six electrolytes. Data are presented in both tables and graphs.

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Experimental study of the stimulated Raman spectrum of sulfuric acid as a function of concentration. Data are presented in a table and in the text.

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Theoretical discussion on the type of bonding in UO_2^{2+} and the nature of magnetic and absorption spectral data. Two M.O. diagrams are presented.

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Experimental study on the U^{+3} electronic absorption spectrum in anhydrous MeOH, HOAc, and HCO_2H saturated with dry H_2 in the range between 7000 and 26,000 cm^{-1} . Data are presented graphically.

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Experimental study on the far-infrared spectra of liquid hydrogen from 20 to 250 cm^{-1} . Data are shown

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of iron and steels. V. Infrared absorption spectra of
metal carbides, nitrides, and sulfides]: Bunseki Kagaku,
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range $1400-400 \text{ cm}^{-1}$ using the KBr disc technique.

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1050-1055. In Japanese.

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[Paramagnetic relaxation and ESR spectra of aqueous

solutions of trivalent iron]: Doklady Akademii Nauk SSSR, v. 212, no. 5,, p. 1142-1145. In Russian.

Experimental study of the relaxation times of concentrated aqueous solutions of ferric sulfate and ferric nitrate by a nonresonance absorption method. Data are presented graphically.

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Experimental study on the photoelectron spectrum of H₂S. The energics of the observed vibrational and vibronic bands are also reported. Data are presented graphically and in a table.

Karr, Clarence, Jr., and Kovach, J.J., 1969, Far-infrared spectroscopy of minerals and inorganics: Applied Spectroscopy, v. 23, no. 3, p. 219-223.

Experimental determination of the far-infrared spectra, 200 to 50 cm⁻¹, of 18 different minerals and inorganic compounds. Included among these is ferrous sulfate in the surface film on pyrite. Data are presented both graphically and in tabular form.

Kasper, H., and Drickamer, H.G., 1968, High-pressure Moessbauer resonance studies of compounds of iron with group V and group VI elements: Proceedings of the U.S. National Academy of Science, v. 60, no. 3, p. 773-775.

Experimental study on the anomalous behavior of compounds of iron with sulfur, selenium, and other elements using high-pressure Moessbauer resonance techniques. Data are presented in a table and in two figures.

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Experimental study of a number of properties of uranyl selenites. Spectrophotometric studies were conducted in

the 370-500 nm region while ir and Raman spectra were recorded in the 250-4000 cm^{-1} region. Data are given in tables and on graphs.

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Analytical procedure for the photometric determination of thiosulphate using cyanolysis with lanthanum as catalyst at 460 nm. Results are presented in tables and graphs.

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Experimental study on the absorption spectra of uranyl nitrate crystals at 77 K in both ordinary and polarized light. [Translation unavailable to authors. Data taken from Chem. Abs. 71:17311r, 1969.]

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Experimental study on the uv, visible, and ir spectra of Fe^{2+} solutions in H_2SO_4 and $HClO_4$. [Translation unavailable to authors. Data taken from Chem. Abs. 83:16598a, 1975.]

Koster, A.S., 1971, L emission spectra of compounds of iron and manganese: Proceedings Koninklijke Nederlandse Akademie van Wetenschappen, Ser. B, v.74, no. 3, p. 332-339.

Experimental study on the L emission profiles of Fe, FeO, Fe_3O_4 , Fe_2O_3 , FeS, FeS_2 , FeF_3 , Mn, MnO , MnO_2 , and $KMnO_4$. Data are presented in tables and on graphs.

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Experimental study on the binding energies of several compounds containing both iron and sulfur. Data are presented graphically and in the text.

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Experimental study on the electronic emission spectrum of SeO_2 in the 4000-6000 \AA region. Experimental data are given in several tables.

Lakshminarayana, G., and Narasimham, N.A., 1967, A new electronic transition of S_2 in the vacuum ultraviolet region: Current Science, v. 36, no. 20, p. 533.

Experimental study on the emission spectrum of S_2 excited in an electrodeless microwave discharge through sulfur and helium. A new band system in the 1780-1850 Å region is described. Data are given in a table and the spectrum is shown in a figure.

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An indirect atomic-absorption method for the determination of selenium using two reactions that lead to the formation of the $Pd(DanSe)_2Cl_2$ complex and measurement of the palladium absorption. Data are presented in tables and on graphs.

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Theoretical study on the loss of high resolution at low temperatures in the fluorescence spectra of uranyl salts. A model is presented.

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Experimental study on H₂S using quantitative infrared intensity measurements in the 4 μ region at a constant pressure of 20.2 atm. and a temperature range of 295-448 K. The energy of hydrogen bonding in the gas phase was estimated. Data are presented graphically.

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Experimental study on the infra-red and Raman spectra of trigonal, alpha-monoclinic, and amorphous Se and

identification of the frequencies and symmetry characters of the fundamental vibrational modes. Data are presented in tables.

Lutz, H.D., and Willich, P., 1974, [Lattice vibration spectra.

IX. Pyrite structure. Far infrared spectra and normal coordinate analysis of MnS_2 , FeS_2 , and NiS_2]:
Zeitschrift fur Anorganische und Allgemeine Chemie, v. 405, no. 2, p. 176-182. In German.

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Nukleonika, v. 18, no. 7-8, p. 325-336. In Polish.

Experimental study on the absorption spectra (400-500 nm) of uranium (VI) complexes sorbed on anion-exchange resin from sulfate solutions. Data are presented graphically.

Majchrzak, Kazimierz, 1973, [Equilibria in solutions and anionic exchange of uranium (VI) complexes. IV. Stability constants and sorption of uranyl sulfate complexes]:

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Majkowski, R.F., Flint, R.J., and Hill, J.C., 1978, Infrared absorption coefficients of gaseous H_2SO_4 and SO_3 : Applied Optics, v. 17, no. 7, p. 975-977.

Experimental study on the influence of partial pressure variations on the absorption spectrum of gaseous H_2SO_4 and SO_3 in the infrared region. Data and spectra are presented graphically.

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Experimental study on the ir spectra of H_2SeO_4 in various other acids. [Translations unavailable to authors. Data taken from Chem. Abs. 69:54701x, 1968.]

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Experimental spectrophotometric study of thin sections of natural Fe-containing sphalerite in the energy range 10,000-30,000 cm^{-1} . The absorption was resolved into a number of energy bands characteristic of octahedrally bound Fe (III). Data are presented graphically and in two tables.

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Experimental study on the Raman spectra of HSO_4^- and SO_4^{2-} from 1 to 3000 atm in an experimental cell. [Translation unavailable to authors. Data from Chem. Abs. 83:88559n, 1975.]

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Melendres, C.A., and Tani, B., 1978, Moessbauer spectra and lattice parameters of some new sulfide phases of iron: The Journal of Physical Chemistry, v. 82, no. 26, p. 2850-2852.

Experimental study on the Moessbauer spectra of several iron sulfide phases found in the charged and discharged electrodes of molten salt secondary batteries at room temperature and at 77 K. Spectra are displayed graphically and data are listed in a table.

Mendes-Bezerra, A.E., and Uden, P.C., 1968, Determination of sulfate by titrimetric and colorimetric measurement of equivalent displaced zinc ion: Analytical Letters, v. 1, no. 6, p. 355-358.

Two analytical procedures are given for the determination of sulfate in the 0.2 - 100 mg. range. Pertinent information is given in the text.

Meyer, B., Oommen, T.V., and Jensen, D., 1971, The color of liquid sulfur: The Journal of Physical Chemistry, v. 75, no. 7, p. 912-917.

Experimental study on the visible spectrum of liquid

sulfur between the melting point and 700°C at 10 atm., as well as the absorption spectra of various gas phases, solids, and solutions of sulfur. Data are presented graphically.

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Experimental study on the absorption spectrum of liquid sulfur in the temperature range between the melting point and 700°C. Data are given graphically and in the text.

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Experimental study on the Raman spectra of H₂S and D₂S in each of their three solid phases using both laser and conventional excitation. Data are presented graphically and in tabular format.

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Experimental study of the far-infrared spectra of H₂S, D₂S, and H₂S - D₂S mixtures at low pressures with high resolution. Data are presented graphically and in tables.

Miller, R.E., Leroi, G.E., and Eggers, D.F., 1967, Infrared spectrum of deuterium sulfide: The Journal of Chemical Physics, v. 46, no. 6, p. 2292-2297.

Experimental study of the infrared spectrum of deuterium sulfide with investigation of its bending fundamental (v2) at high resolution and calculation of inertial constants. Data are presented in both tables and graphs.

Mohler, E., Stuke, J., and Zimmerer, G., 1967, Some new features of the reflectance spectrum of trigonal selenium single crystals: Physica Status Solidi, v. 22, no. 1, p. K49-K53.

Experimental study on selenium in the energy range 1.85 to 9.2 eV using reflectivity measurements with a qualitative discussion of the observed structure. Spectrum and structural diagrams are given in two figures.

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Experimental study on the ultraviolet absorption spectrum of hydrogen peroxide vapor over the wavelength range 210 to 350 nm at 296 K. Data are presented in a table and in a figure.

Montano, P.A., 1977, Moessbauer spectroscopy of iron compounds found in West Virginia coals: Fuel, v. 56, no. 4, p. 397-400.

Experimental study of iron pyrite and iron (II) sulfate in coal samples from various seams in West Virginia. Evidence for the formation of a new mineral species during the low-temperature ashing process is presented. The data are presented graphically and in a table.

Morillon, C., 1970, [Emission spectra of samarium, neodymium, and uranium between 2.3 and 4.1 micrometers studied with a grating spectrometer]: Spectrochimica Acta, v. 25B, no. 10, p. 513-538. In French.

Experimental study on the emission lines of samarium, neodymium, and uranium. About 300 lines of U in the range 2.3 to 3.4 micrometers were measured. Data are given in tabular form.

Moriyasu, Masataka, Yokoyama, Yu, and Ikeda, Shigero, 1977,
Quenching of uranyl luminescence by water molecule:
Journal of Inorganic and Nuclear Chemistry, v. 39, no.
12, p. 2211-2214.

Experimental study of the quenching of uranyl luminescence by water molecules. The authors conclude that an electron transfer mechanism in terms of oxidation-reduction potentials of uranyl ion and water molecules is involved. Data are presented in tables and one figure.

Moriyasu, Masataka, Yokoyama, Yu, and Ikeda, Shigero, 1977, Anion coordination to uranyl ion and the luminescence lifetime of the uranyl complex: Journal of Inorganic and Nuclear Chemistry, v. 39, no. 12, p. 2199-2203.

Experimental study of the nonexponential uranyl luminescence decay curves of uranyl complexes in the wavelength range 470-620 nanometers at 25°C. Data are presented graphically and in tables.

Moriyasu, Masataka, Yokoyama, Yu, and Ikeda, Shigero, 1977,
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Experimental study of the quenching reaction of uranyl luminescence by metal ions in a lower valence state such as Fe^{2+} . The authors conclude that the reaction is diffusion-controlled and that the dominant process is an electron transfer from the quenchers to the uranyl ion. Data are presented in several tables and two figures.

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The electronic structure of SO_4^{2-} , SO_3^{2-} , SeO_4^{2-} , SeO_3^{2-} , and other ligands was investigated using x-ray photoelectron spectra. [Translation unavailable to authors. Data taken from Chem. Abs. 81:19021b, 1974.]

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Experimental study on the Raman spectra of SeO_2 and $(\text{SeO}_2)_2$ at various temperatures. Data are presented graphically and in tables.

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Experimental study on the pulse radiolytic absorption transients observed in aqueous solutions between 200 and 300 nm. Data are presented in both graphical and tabular format.

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table.

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solutions, and sulfide solutions. The data are presented graphically.

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