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The Nature of the Interior of Uranus Based on Studies of Planetary Ices at High Dynamic Pressure

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Data from the Voyager II spacecraft showed that Uranus has a large magnetic field with geometry similar to an offset tilted dipole. To interpret the origin of the magnetic field, measurements were made of electrical conductivity and equation-of-state data of the planetary "ices" ammonia, methane, and "synthetic Uranus" at shock pressures and temperatures up to 75 gigapascals and 5000 K. These pressures and temperatures correspond to conditions at the depths at which the surface magnetic field is generated. Above 40 gigapascals the conductivities of synthetic Uranus, water, and ammonia plateau at about 20 (ohm-cm)^{-1} , providing an upper limit for the electrical conductivity used in kinematic or dynamo calculations. The nature of materials at the extreme conditions in the interior is discussed.

THE VOYAGER II SPACECRAFT DATA showed that Uranus has a strong magnetic field, which can be described approximately by a dipole of moment 0.23 G R_U^3 , where R_U is the planetary radius (25,600 km). The dipole is centered at $0.3 R_U$ and its axis is tilted 60° from the axis of rotation. The maximum value at the surface is 1.1 G (1), twice the value for Earth, and the minimum surface value is 0.1 G. These unexpected and unusual Voyager data emphasize the need for laboratory measurements of planetary materials at interior conditions in order to derive planetary models consistent with the observational data. We report new shock-wave electrical conductivity and equation-of-state (EOS) data

for planetary "ices" and for a liquid with an atomic composition representative of a mixture of the "ices." Our electrical conductivity data provide density and temperature dependences up to 75 GPa (0.75 Mbar) and 5000 K and provide an upper bound that can be used in kinematic and dynamo calculations to explain the magnetic field. Laboratory data and theory also allow us to hypothesize on the nature of the interior.

The composition of Uranus is estimated from the mass, radius, rotational rate, and gravitational field (2, 3). The combined magnetic-field and radio-emission data yielded an improved measurement of the rotation rate of 17.24 hours (4). This result led to improved values of the density moments, which are derived from the gravitational moments deduced from the observed precessions of the elliptical rings of Uranus. The density moments provide a constraint on the equations of state of candidate materials. Prior to Voyager II, Hubbard and MacFarlane proposed that Uranus consists of three layers: an outer gas layer of $\text{H}_2\text{-He}$;

a middle "ice" layer composed mostly of H_2O , CH_4 , and NH_3 ; and an inner rocky core (2). The "ices" are actually in the fluid phase over a substantial portion of these proposed thermodynamic conditions.

After Voyager II, Kirk and Stevenson proposed that the magnetic field and the density moments derived from improved values of the rotation rate and gravitational moments are consistent with a greater degree of material mixing (5) than in the pre-Voyager models. One model consists of an outer gas-ice-rock region, which is mostly an $\text{H}_2\text{O-H}_2$ mixture, and an inner rocky core. The boundary is at $0.3 R_U$ and 300 GPa and 6000 K. The magnetic field is produced primarily in an H_2O -rich region near $0.7 R_U$ at 50 to 100 GPa, based on a kinematic model that considers equatorial zonal jet flow and uses the electrical conductivity of water extrapolated from Holzapfel's (6) fit to static and shock data up to 10 GPa and 1300 K. The Voyager II data were used by Gudkova *et al.* (7) and they also found more mixing. Their three-layer model has an outer $\text{H}_2\text{-He}$ -ice region, a middle ice-rock layer, and an inner rocky core. The outer-middle boundary at $0.71 R_U$ is at 30 GPa and 2700 K and the middle-core boundary at $0.14 R_U$ is at 600 GPa and 6800 K. Their two-layer model is ice-rock for radii less than $0.72 R_U$ at 25 GPa and 2600 K, with $\text{H}_2\text{-He}$ -rock-ice being the remainder. The vari-

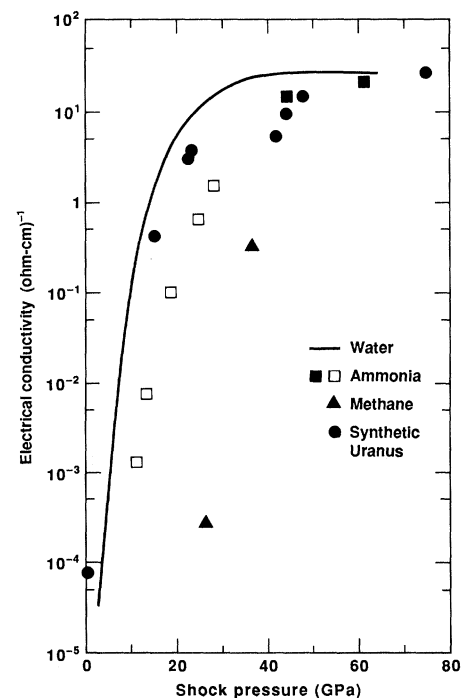


Fig. 1. Electrical conductivity versus shock pressure for planetary fluids. Solid symbols are the present work. Open squares are from (16). Solid curve is smooth curve through the data of (9) and (12). At 40 GPa, temperatures in these fluids are 2600 K in water, 2800 K in synthetic Uranus, 3100 K in ammonia, and 4100 K in methane.

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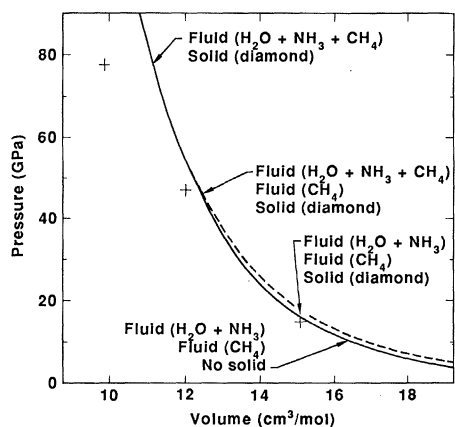


Fig. 2. Shock-compression equation of state of synthetic Uranus of initial density 0.896 g/cm^3 and initial temperature 295 K. Pluses are experimental points with error bars. Solid curve is theory with molecular-fluid phase separation; dashed curve is theory for a homogeneous fluid. Calculated equilibrium phases are indicated at four positions. At 76 GPa temperature is 5100 K.

ous models demonstrate that a unique model for Uranus is not yet possible. These models and the fact that the magnetic field is centered at $0.3 R_U$, which means that the large surface magnetic field of 1.1 G is generated at even larger radii ($\sim 0.7 R_U$), indicate that the most relevant conductivity data are in the range 10 to 100 GPa and several thousand degrees kelvin, the regime in which we have made our measurements.

Shock-wave EOS (8–10), temperature (11), electrical conductivity (9, 12), and spontaneous Raman scattering (13) data were measured for water up to the 100-GPa range. For ammonia (9, 14) and methane (15) EOS data have been obtained up to 70 and 90 GPa, respectively. Conductivities and temperatures of ammonia were measured up to 28 GPa (16) and 61 GPa (17), respectively. Shock data are relevant to planetary interiors because the shock compression of liquid specimens achieves the same high pressures and temperatures as the isentropic compression of low-density gas to form the giant planets (18).

We report here new electrical conductivity and shock temperature data for NH_3 , CH_4 , and the mixture we call “synthetic Uranus.” EOS data were also measured for this mixture, an H-rich liquid with an H:O composition ratio of 3.5:1 and abundance ratios close to the cosmological ones of O:C (7:4) and O:N (7:1). It is a solution of water, ammonia, and isopropanol ($\text{C}_3\text{H}_8\text{O}$) with mole fractions of 0.71, 0.14, and 0.15, respectively. The H:O ratio is similar to that of the ice-gas mixture in the ratio 2.5:1 proposed for Uranus (5), assuming an H_2O - H_2 mix. High densities and temperatures achieved in shock experiments quickly induce chemical and thermal equilibrium, so that our results are expected to be weakly

sensitive to the initial structure of the specimens. Although this mixture is not unique, we believe it is representative of a mixture of the planetary ices. Strong shock waves were generated by the impact of metal plates onto the front wall of liquid-specimen holders. The plates were accelerated up to 7 km/s by a two-stage light-gas gun. The experimental techniques were described previously (9, 17) and details are published elsewhere (19).

The electrical conductivities of shock-compressed liquid ammonia, methane, and synthetic Uranus are plotted versus shock pressure in Fig. 1. Previous data for water are included for comparison. The temperature measurements show that our conductivities are in the range 1500 to 5000 K. The electrical conductivities of synthetic Uranus, water, and ammonia all approach $20 (\text{ohm-cm})^{-1}$ above 40 GPa (20), providing an upper limit on the value of conductivity that can be used in kinematic or dynamo calculations. This limit is only weakly sensitive to chemical species. Below 40 GPa the conductivity of synthetic Uranus is a factor of 2 to 3 less than that of water. Shocked methane has substantially lower conductivity than the other fluids, at least up to 40 GPa and 4000 K, caused probably by the high stability of the CH_4 molecule. Thus, methane can only contribute to the magnetic field if it decomposes significantly into conducting phases at higher pressure and if the depths at which this occurs are sufficiently small so the field generated is significant at the surface.

The maximum conductivity can be used to characterize crudely the flow that produces the magnetic field. The dynamo theorem states that $R_M \equiv uR/\lambda \geq \pi^2$, where R_M is the magnetic Reynolds number, u is the maximum deviation of the velocity field from uniform rotation in a fluid sphere of radius R , $\lambda \equiv c^2/4\pi\sigma$ is the magnetic diffusivity, and σ is the electrical conductivity of the sphere (21). R_M is usually assumed to be in the range 10 to 100 for a steady external field. For $\sigma = 20 (\text{ohm-cm})^{-1}$, $R = 0.7 R_U$, and $R_M = 50$, then $u = 10^{-1} \text{ cm/s}$, which is small compared to the linear rotational velocity of $2 \times 10^5 \text{ cm/s}$ at $0.7 R_U$.

Kirk and Stevenson showed that the magnitude of the magnetic field of Uranus can be calculated kinematically with the conductivity of water suppressed by a factor of 10. They assumed this factor to take account of dissolved insulating H_2 -He (5). The conductivity of synthetic Uranus is still a factor of 3 to 5 larger than they used and allows for the presence of insulating phases. Thus, the shock-wave conductivity data for synthetic Uranus provide more representative experimental data than for pure water and this new data is consistent with the kinematic calculations of the magnetic field.

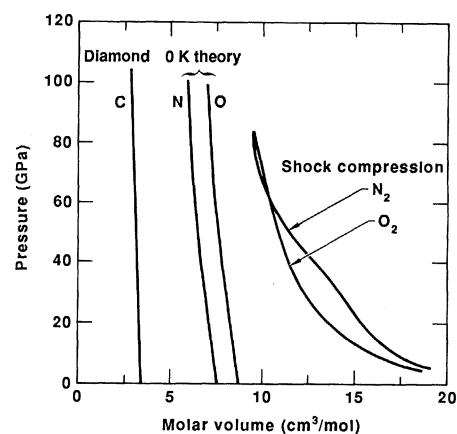


Fig. 3. Shock-compression equations of state for C (diamond) and liquid N_2 and O_2 and calculated 0 K isotherms for monatomic N and O, plotted as pressure versus molar volume. The 0 K curve for diamond is nearly the same as the shock-compression curve shown. The initial molar volumes of liquid N_2 and O_2 are 34.7 and $26.6 \text{ cm}^3/\text{mol}$, respectively.

Three EOS points were measured for synthetic Uranus and are plotted with chemical equilibrium calculations (22) for the shock-compression curve in Fig. 2. The calculation minimizes the free energy of the system with respect to nine molecular-fluid species and diamond. Equilibrium phases formed in the calculation are indicated at four locations on the shock-compression curve. The calculations suggest that at lowest shock pressures, the fluid separates into methane and ammonia-water phases. As pressure and temperature increase, diamond starts to form and methane becomes more soluble in the water-ammonia phase. The discrepancy between theory and experiment at highest pressures is probably caused by neglecting ionic species in the calculations, which cause the high electrical conductivities measured above 40 GPa.

The magnetic field and density moments, together with known material properties, provide evidence that Uranus is indeed composed of ice, gas, and rock. It is interesting to speculate on the nature of materials inside the planet. The high electrical conductivity of shocked water, the major ice constituent, is caused by molecular ionization (12). The same mechanism should apply to ammonia and synthetic Uranus. The range 10 to 20 GPa and 700 to 1200 K is an interesting one in which the ion concentration in H_2O increases from about 0.5% (12) to a saturation value, H bonds between H_2O molecules break (13), and the compressibility increases (10). Above 20 GPa water has been said to be totally ionized into OH^- and H_3O^+ (12). However, Raman scattering experiments at 26 GPa and 1700 K (13) show only a single broad O–H stretch band, which does not distinguish between vibra-

tions in the H₂O molecule and OH⁻ ion. No indication of the H₃O⁺ species is distinguishable in the spectra. Thus, the degree of dissociation of water cannot be determined from the Raman data. A lower bound on ion concentration can be estimated by assuming that mobile protons are the dominant charge carriers. By using a classical conductivity model and a mean free path of a molecular dimension, we estimate the proton concentration to be one proton for every ten original H₂O molecules at pressures above several tens of gigapascals. Thus, we estimate the degree of dissociation of water to be between 10 and 100% above 20 GPa and 1200 K.

At depths where the pressure exceeds 100 GPa and temperatures are several thousand degrees kelvin, molecules and ions are expected to dissociate and form dense, stiff monatomic or H—O—C—N species or both. This statement is based on observations of molecular dissociation at high shock pressures and temperatures. For example, N₂ starts to dissociate at only 30 GPa at 6000 K (23) but remains molecular to 130 GPa at 300 K (24), illustrating that high temperatures inside Uranus are expected to drive dissociation. Shock-compression curves for carbon (diamond) (25), nitrogen (23), and oxygen (26) and theoretical curves for monatomic simple-cubic N (27) and O (28) at 0 K are plotted as pressure versus volume in Fig. 3. The curves for the monatomic phases at 0 K are all steep; the shock-compression curves of liquid nitrogen and oxygen have comparable steep slopes above 60 GPa and 10,000 K. When dissociation is completed, intermolecular repulsion is replaced by stiffer interatomic repulsion. This transition is expected to proceed continuously along the planetary isentrope. In general, elements probably do not phase separate but form mixtures or react with each other to form compounds, which are nearly incompressible as in Fig. 3. Such materials might have high melting temperatures and condense out of the solution of the fluid "ice" mixture. Rock also has a stiff equation of state at high pressures. Thus, at pressures above 100 GPa planetary materials are probably close to their limiting compression, which contributes somewhat to the large volume of Uranus. That is, pressures above 100 GPa probably do not cause significant compression of the phases present because of their small compressibilities. Figure 3 suggests that all materials in the deep interior are dense, stiff materials having a high bulk modulus such as the diamond phase of carbon, which has been suggested to form inside Uranus (29).

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20. The upper limit of detection is ≥ 100 (ohm-cm)⁻¹. Conductivities larger than 20 (ohm-cm)⁻¹ were measured for shocked liquid O₂ [73 ± 10 (ohm-cm)⁻¹ at 43 GPa], liquid N₂ [47 ± 6 (ohm-cm)⁻¹ at 62 GPa], and liquid benzene [46 ± 9 (ohm-cm)⁻¹ at 50 GPa].
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Hand of *Paranthropus robustus* from Member 1, Swartkrans: Fossil Evidence for Tool Behavior

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New hand fossils from Swartkrans (dated at about 1.8 million years ago) indicate that the hand of *Paranthropus robustus* was adapted for precision grasping. Functional morphology suggests that *Paranthropus* could have used tools, possibly for plant procurement and processing. The new fossils further suggest that absence of tool behavior was not responsible for the demise of the "robust" lineage. Conversely, these new fossils indicate that the acquisition of tool behavior does not account for the emergence and success of early *Homo*.

THE GENUS *Paranthropus* GENERALLY has been assumed to consist of small-brained, large-toothed early hominids that subsisted on a vegetarian diet (1). Stone artifacts found with *Paranthropus* have been attributed to their contemporaries of the genus *Homo* (2). With a small brain and a vegetarian diet, it was thought that *Paranthropus* had neither the intellect nor

the impetus to engage in tool behavior (3, 4). Further, the lack of tool behavior was thought to have contributed to the eventual extinction of *Paranthropus* in the early middle Pleistocene. Most inferences about *Paranthropus* have come from studies of craniodental remains (5). Until now the only postcranial fossils referred to *Paranthropus* are seven bones from Swartkrans (6) and four from Kromdraai (7).

There are 22 hand bones among 37 new hominid postcranials from Swartkrans (8).

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