

Bull. Amer. Phys. Soc., 15, 1603 (1970)

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H.E. RORSCHACH, and  
& Rice Univ.,  
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macromolecules within the cell plays an important  
in cellular phenomena. There are two opposing views  
the state of cellular water. The "classical" view  
contains that the bulk of the cell water is in a state  
ivalent to pure water. An opposing view maintains  
a substantial fraction of the cell water differs in  
physical properties from free water. Previous high-  
resolution NMR studies have shown line-broadening that  
has been associated with structural changes in the cell  
water. We have used pulse methods to measure the relaxa-  
tion times and diffusion coefficients for water in rat  
skeletal muscle. Measurements on four different animals  
give the following results.  $T_1 = 30 \pm 70$  ms,  $T_2 = 46 \pm 2$  ms,  
 $D = 1.5 \pm 2 \times 10^{-5}$   $\text{cm}^2/\text{sec}$ . For pure water, we obtain  
 $T_1 = 3.0$  s,  $T_2 = 1.5$  s,  $D = 2.8 \times 10^{-5}$   $\text{cm}^2/\text{sec}$ . The impli-  
cations of these results for the structure of the cell  
water will be discussed.

supported in part by the Robert A. Welch Foundation and  
the U. S. Public Health Service.

### BE 3. Pressure Dependence of Internal Rotation in

$\text{FeSiF}_6 \cdot 6\text{H}_2\text{O}$ \* G. L. Nicolaides and R. W. Vaughan  
California Institute of Technology and D. D. Elleman  
Jet Propulsion Laboratory - Molecular motion in  
ferrous fluorosilicate ( $\text{FeSiF}_6 \cdot 6\text{H}_2\text{O}$ ) has been studied  
as a function of pressure to 80 kilobars using  
conventional wide-line nuclear magnetic resonance  
techniques. This material has a slightly disordered  
 $\text{CsCl}$  structure with  $\text{SiF}_6^{2-}$  and  $\text{Fe}(\text{H}_2\text{O})_6^{2+}$  groups occupy-  
ing the lattice sites. Second moments calculated from  
the  $\text{F}^{19}$  spectra indicate the rapid reorientation of  
 $\text{SiF}_6^{2-}$  octahedra at room temperature and pressure slows  
with the application of pressure. Second moments  
approaching the estimated rigid lattice values are  
obtained near 70 kilobars.

In addition to a detailed discussion of these  
results a brief discussion of the high pressure cell  
developed for these studies will be given.

\*Work supported by the National Science Foundation,  
The Caltech President's Fund, and NAS 7-100.

### BE 4. EPR Studies of $\text{Np}^{4+}$ in $\text{ThO}_2$ \* R.P. RICHARDSON<sup>†</sup>

and J.B. GRUBER, Washington State Univ.--The EPR spectrum  
of  $\text{Np}^{4+}$  in  $\text{ThO}_2$  has been investigated, principally at  
1.7°K in the K-band. Three basic Zeeman resonances are  
energetically isotropic about one [001] crystal axis,  
with g values 2.02, 2.31, and 2.48. These resonances  
appear to come from three sites in which the  $\text{Np}^{4+}$  ions  
are slightly displaced along the [001] axis from the usu-  
al  $\text{O}_h$  point group site. A final basic Zeeman resonance,  
isotropic about the [110] crystal axis with  $g=2.06$ , is  
thought to arise from a fourth site in which the  $\text{Np}^{4+}$   
ion is displaced along the [110] axis. The splittings  
of the cubic field ground  $\text{F}_g$  quartets that arise due to  
these four axial displacements must be between  $5\text{cm}^{-1}$  and  
 $20\text{cm}^{-1}$  in order to fit the EPR and optical data. The  
parameter  $\alpha$ , related to the cubic portion of the crystal  
field, is found to be  $\alpha = -0.667, -0.631, -0.604$  and  
 $-0.661$  for the four nearly cubic sites. The hyperfine  
lines are fit best with  $(390 \pm 30) \times 10^{-4} \text{cm}^{-1}$  for the abso-  
lute value of the hyperfine coupling constant.

\*Work supported by the U.S. Atomic Energy Commission.

<sup>†</sup>Present address: Texaco, Inc., Bellaire, Texas.

BE 5. Nuclear Spin Diffusion Induced by Para-  
magnetic Impurities in Nonconducting Solids. E.  
PHILIP HORVITZ, Rice Univ.\*--It is shown that  
paramagnetic impurities can induce nuclear spin  
diffusion in nonconducting solids inside the so-  
called "barrier radius." The static field cre-  
ated by the impurity spin splits the states  
 $|1/2, -1/2\rangle$  and  $|1/2, 1/2\rangle$ , where the quantum num-  
bers refer to the component in the direction of  
the external magnetic field of two neighboring  
spins. The nuclear dipole-dipole interaction  
mixes these two states so that to first order  
 $\Psi_1 = |1/2, -1/2\rangle + \epsilon|-1/2, 1/2\rangle$ , and  $\Psi_2 = |-1/2, 1/2\rangle$   
 $-\epsilon|1/2, 1/2\rangle$ . The Fourier component of the im-  
purity spin at the frequency corresponding to  
the energy difference of  $\Psi_1$  and  $\Psi_2$  causes  
transitions between these states. This is a spin  
diffusion process because  $\epsilon$  is small. Typically  
the induced nuclear spin diffusion is on the  
order of  $10^{-12} \text{cm}^2/\text{sec}$ . Thus, Bloembergen's  
differential equation should include spin  
diffusion inside the "barrier radius."

\*Present Address: Manned Spacecraft Center,  
Code EE6, Houston, Texas 77058.

### BE 6. Theory of Pressure-Induced Demagnetization of

Porous Polycrystalline Cubic Ferrites.\* DENNIS E. GRADY  
and G. E. DUVALL, Washington State Univ.--A theory is  
presented which predicts the demagnetization of porous  
cubic ferrites induced by hydrostatic pressure for  
values of applied field which would normally saturate  
the magnetic material. The theory considers magneto-  
elastic coupling with the deviation in local strain  
field due to the porosity. Agreement is found with  
existing data on nickel ferrite, manganese ferrite and  
yttrium iron garnet.

\*Research supported by AFOSR Contract #69-1758.

### BE 7.

Clustering and Magnetic Behavior in Au-Fe  
Alloys.\* R. J. BORG and C. E. VIOLET, Lawrence  
Rad. Lab., Univ. of California, Livermore. --

We have been able to induce, by various heat treat-  
ments, significant variations in the magnetic re-  
sponse of Au-16.6 at.% Fe alloys which were exam-  
ined using the Mossbauer effect.

The following are the salient results: (1) there is  
no discrete ordering temperature for either the  
quenched or annealed state, (2) the temperature de-  
pendence of the magnetic hfs cannot be even approxi-  
mated by a Brillouin function, (3) the isomer shift  
does not change detectably with heat treatment, and  
(4) the splitting of the magnetic hfs for the quenched  
state is greater than for the annealed state except at  
the lowest temperature, at which they are equal.

Alloys more dilute in Fe behave otherwise, dem-  
onstrating relatively sharp ordering temperatures,  
Brillouin-like temperature dependence, and no de-  
tectable response to varying temperature.

\*Work performed under the auspices of the U. S.  
Atomic Energy Commission.