

	Observed $D \times 10^4$ cm <sup>2</sup> /sec
0.72	
1.49	
1.60	
1.32	
1.50	
1.43	
1.13	
1.17	
1.47	
0.75	
1.04	
1.24	

verage deviation

t and most  
tem because  
d cover the0.1 N TINO<sub>3</sub><sup>a</sup>

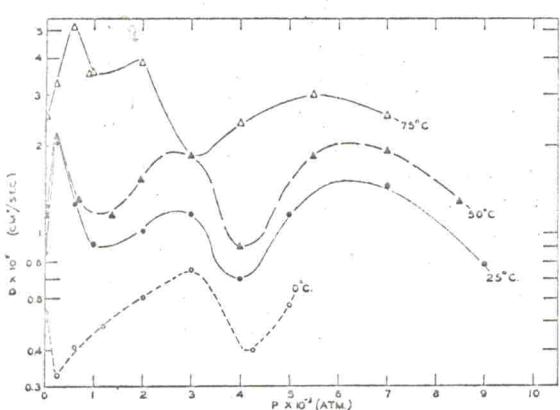
	Observed $D \times 10^4$ cm <sup>2</sup> /sec
2.07	
1.35	
1.80	
1.69	
0.95	
2.00	
1.67	
1.03	
0.88	
1.48	
1.73	

verage deviation

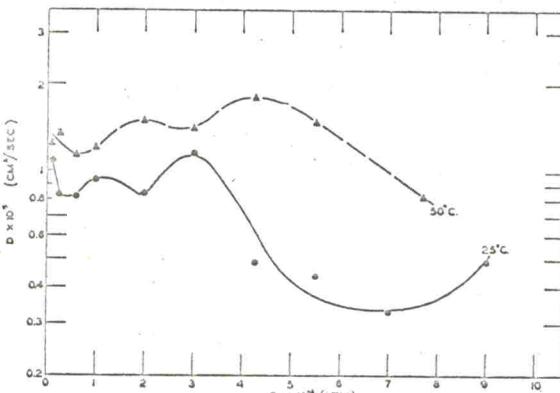
ental work.  
rom Figs. 1  
coefficient  
, and thisOH.<sup>a</sup>

	Observed $D \times 10^4$ cm <sup>2</sup> /sec
1.61	
0.68	
0.40	
0.18	
0.084	
0.041	
3.75	
2.29	
1.26	
0.70	
0.35	
0.088	

verage deviation

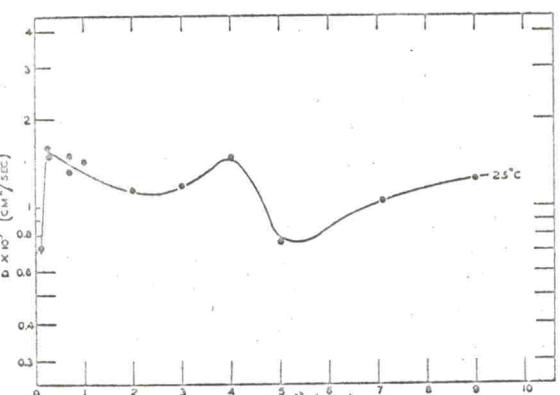
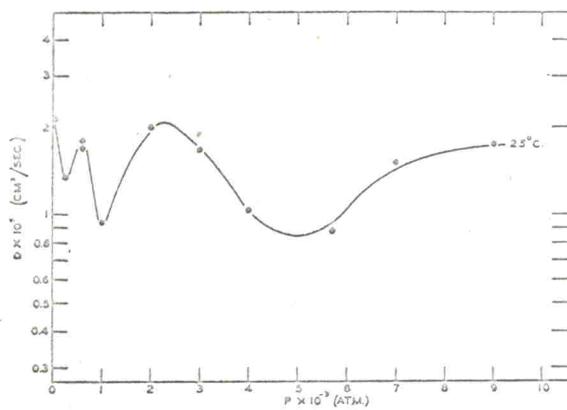
FIG. 1. Diffusion coefficients versus pressure,  
0.1 N Hg(NO<sub>3</sub>)<sub>2</sub>.

minimum shifts toward lower pressure as the temperature increases. The curves also show a maximum in the pressure range 250–600 atmospheres which moves to

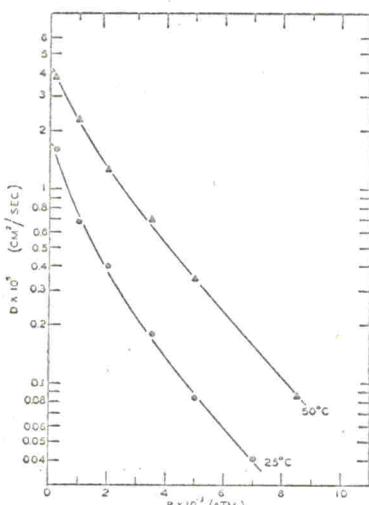
FIG. 2. Diffusion coefficients versus pressure,  
0.1 N CaCl<sub>2</sub>.

lower pressure as temperature decreases and disappears at 0°C.

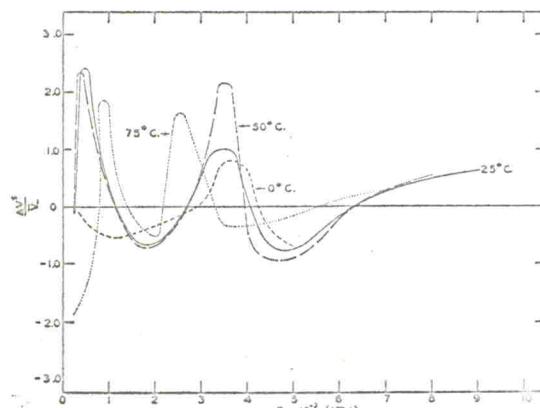
Curves of activation enthalpy and the increase in

FIG. 3. Diffusion coefficients versus pressure,  
0.1 N Ca(NO<sub>3</sub>)<sub>2</sub>.FIG. 4. Diffusion coefficients versus pressure,  
0.1 N TINO<sub>3</sub>.

activation entropy from one atmosphere to  $P$  are shown in Figs. 7 and 8. Any interpretation of these curves

FIG. 5. Diffusion coefficients versus pressure,  
0.01 M HgCl<sub>2</sub> in *n*-butanol.

would be dubious because the activation enthalpy and entropy depend on the displacement of two or more

FIG. 6. Activation volume ratios versus pressure,  
0.1 N Hg(NO<sub>3</sub>)<sub>2</sub>.