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WORKSHOP ON SPACE PHYSICS:
"Materials in Microgravity"
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"Capillarity and Materials Properties" - III

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Please note: These are preliminary notes intended for internal distribution only.

EVALUATION OF FLUXES IN THE MEASUREMENT OF THE SURFACE TENSION OF LIQUID METALS

by

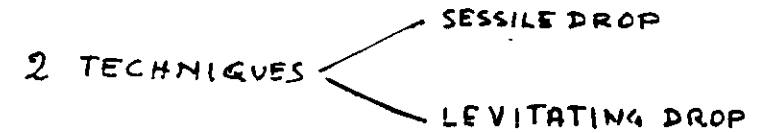
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- MEASUREMENTS OF THE SURFACE TENSIONS OF
LIQUID METALS IN THE PRESENCE OF OXYGEN



- COMMON FEATURES :

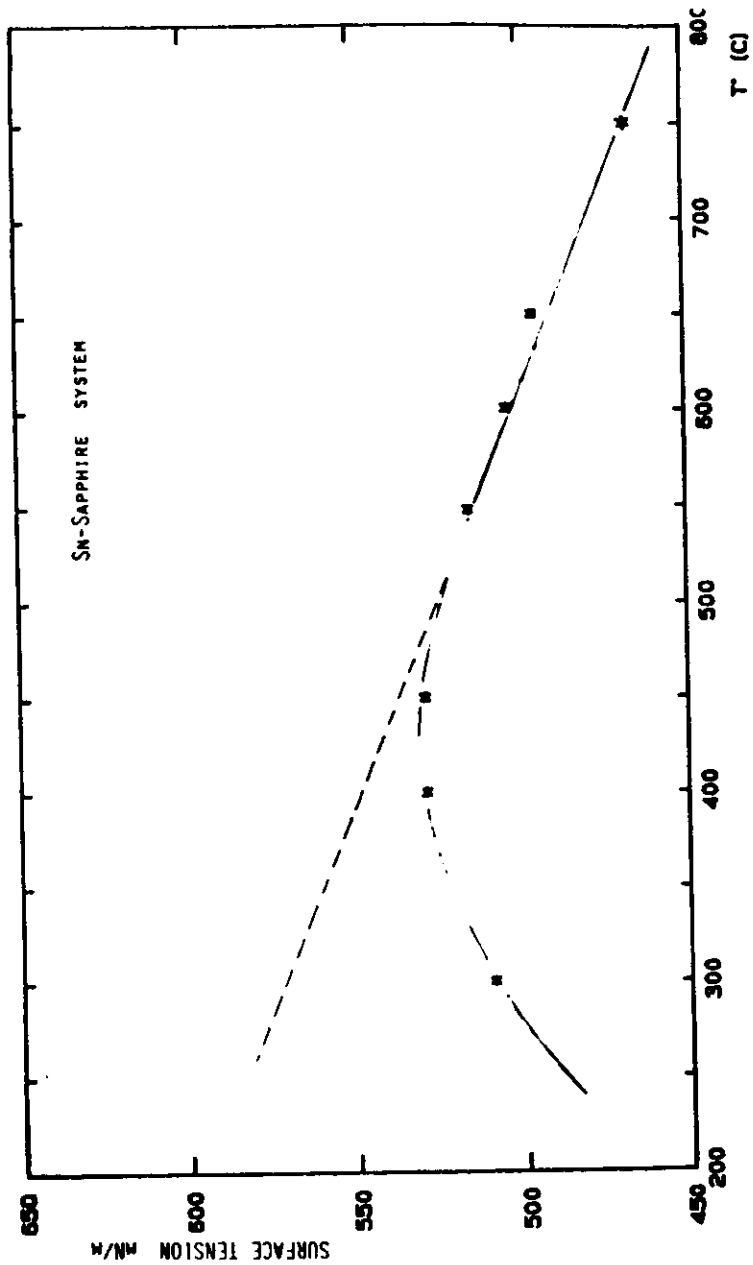
- 1) ISOTHERMAL CONDITIONS
- 2) VERY PURE METALS
- 3) FLUX OF INERT GAS (He, Ar) + OXYGEN
OR VACUUM

- AIM : TO MEASURE $\sigma = f(P_{O_2}, T)$

- TWO ASSUMPTIONS :

- 1) THERMOD. EQUILIBRIUM BETWEEN FLOWING
GAS AND METAL \rightarrow compute $X_{O, BULK}$
- 2) INITIAL "ZERO" OXYGEN CONTENT IN THE

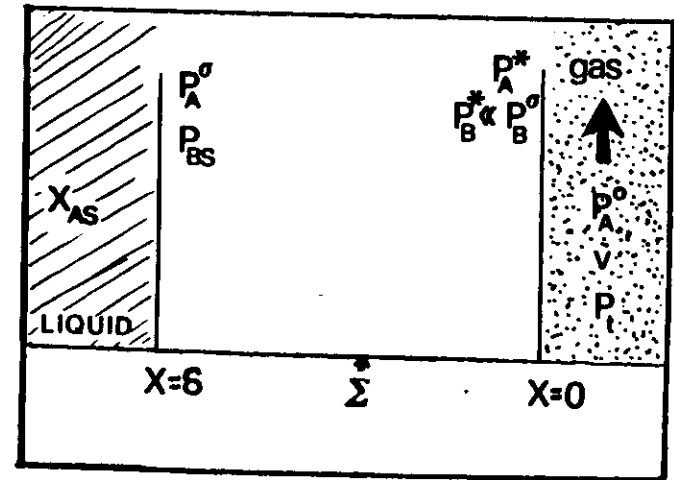
BUT CONDITIONS 1) & 2) are nearly never met
IN FACT : LARGE SCATTER OF EXP. DATA. (See also \rightarrow TIT)



- THE MODEL .

- PURE LIQUID METAL (i.e. $P_A^g = 0$)
- SMALL (Laminar), CONSTANT FLOW OF CARRIER GAS
- ONLY ONE OXIDE FORMS

$$(\alpha A)_g + (B)_g \rightarrow (BA_\alpha)_{sol}$$
- NEGLIGIBLE VAPOUR PRESSURE OF BA_α
- TOTAL PRESSURE P_c CORRESPONDING TO MEAN-FREE PATHS SMALLER THAN δ .



Theory.

- FROM REACT. ①

$$P_A^\alpha \cdot P_B = P_{As}^\alpha \cdot P_{Bs} \quad 2$$

- CONSIDERING A LINEAR VARIATION OF P_A and P_B FOR $0 \leq x \leq \delta$ THE CONDITION :

$$P_A^* \gg P_{As}$$

IS OBTAINED, IN ORDER TO HAVE REACT. ①.

IN THIS CASE, WE DEFINE:

$$\Phi = \delta \sqrt{\frac{\alpha k P_{Bs}}{P_A^* D_A}} \quad \text{THIELE MODULUS}$$

$$\epsilon = \frac{\alpha P_{Bs} D_B}{P_A^* D_A}$$

FOUR REGIONS ARE IDENTIFIED :

a) SLOW REACTIONS $\Phi^2 \ll 1$

- REACT. TIMES \gg DIFF. TIMES

- FLUXES DEPEND ONLY ON D_A, D_B

\approx SAME RESULTS AS FOR NO-REACTIONS

$$\Phi^2 \ll \epsilon$$

b) FAST REACTIONS WITH EXCESS OXYGEN.

- DIFF. TIMES OF B $>$ REACT. TIMES

- LARGE AVAILABILITY OF A

- REACT. ① CONSUMES ALL B

- OXYGEN FLUX UNAFFECTED BY REACT. ①

- FLUX OF B HIGHLY ENHANCED

$$\Phi^2 \gg \epsilon$$

$$\Phi^2 \ll \frac{1}{\epsilon}$$

c) FAST REACTIONS WITH EXCESS METAL VAPOURS

- DIFF. TIMES OF A $>$ REACT. TIMES

- LARGE AVAILABILITY OF B vapour

- REACT. ① CONSUMES ALL A

- OXYGEN FLUX ENHANCED

BY THE FACTOR $\phi \approx \Phi$

$$\Phi^2 \gg 1$$

$$\Phi^2 \ll \epsilon^2$$

d) INSTANTANEOUS REACTIONS

- REACT. TIMES \ll (DIFF. TIMES)_{A, B}

- REACT. ① CONFINED IN A

"REACTION SURFACE" Σ AT

$$X = \frac{\delta}{1 + \epsilon}$$

- OXYGEN FLUX ENHANCED BY THE

FACTOR $\phi \approx 1 + \epsilon$

$$\Phi^2 \gg \frac{1}{\epsilon}$$

$$\Phi^2 \gg \epsilon^2$$

ESTIMATION OF FLUXES.

FOR EACH REGION, THE FLUX OF OXYGEN AT THE LIQUID-VAPOUR INTERFACE CAN BE CALCULATED FOR REGIONS a AND b:

$$N_{AS} = \frac{f \cdot v \cdot P_A^0}{RT(1+fP_e)} \quad P_e = \frac{v \cdot S}{D_A}$$

FOR REGION c:

$$N_{AS} = \frac{f v P_A^0}{RT} \frac{\Phi}{\Phi + f P_e} \frac{1}{1 + \frac{D_B P_{B3} d^2}{D_A P_{AS} \approx (\alpha+1)}}$$

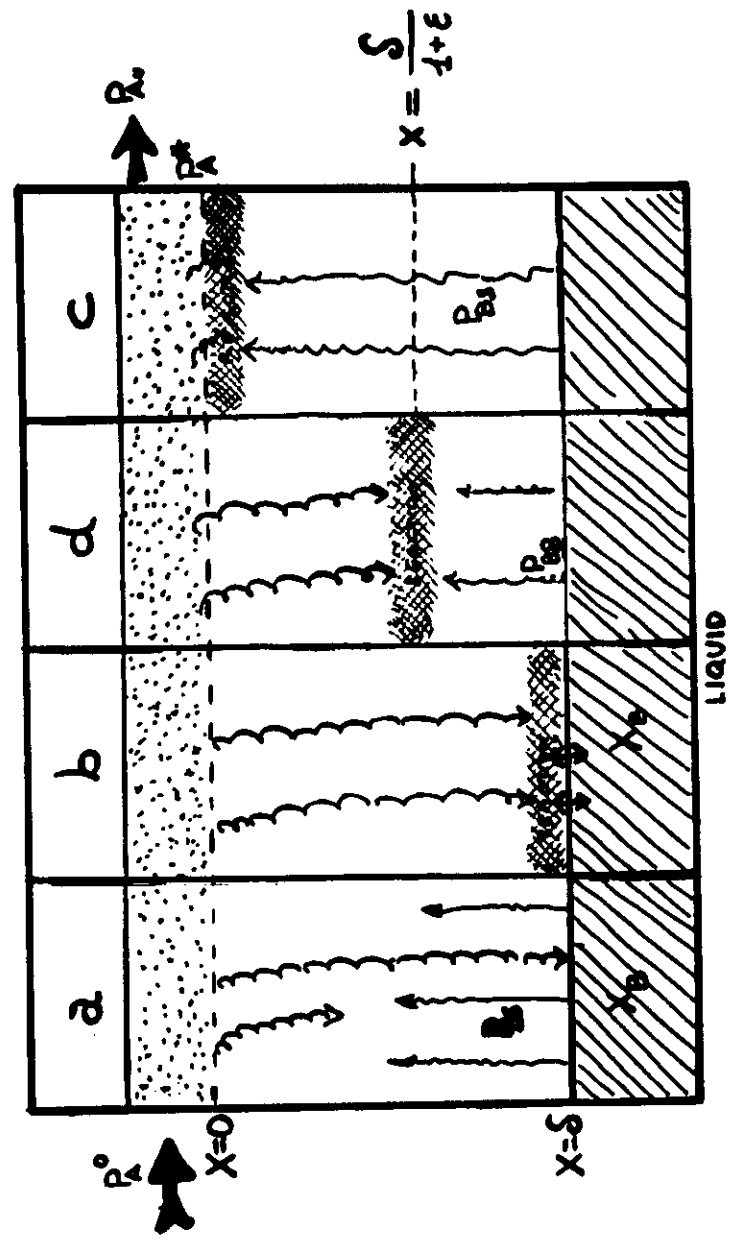
FOR REGION d:

$$N_{AS} = \frac{f v P_A^0}{RT} \frac{1 + \epsilon_0}{1 + f P_e} \frac{1}{1 + \frac{D_B P_{B3} \alpha^2}{D_A P_{AS} \approx (\alpha+1)}}$$

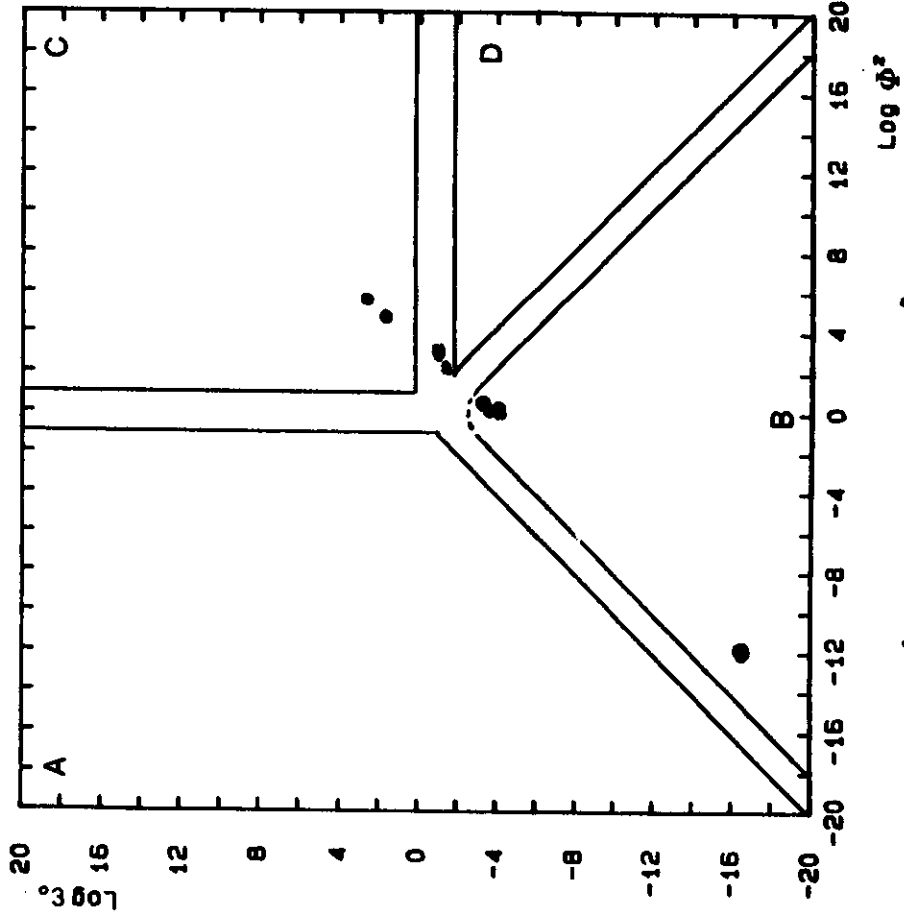
FROM N_{AS} THE TIMES NECESSARY TO SATURATE THE BULK AND THE SURFACE OF THE DROPS CAN BE CALCULATED:

$$t_b = \frac{K_{AS} P_B V_d}{2 M_b S_d N_{AS}}$$

$$t_H = \frac{M}{N_{AS}} \quad \text{WHERE } M \approx 3 \cdot 10^{-9} \text{ mole/cm}^2$$



METAL	OXIDE	T K	REACTION	N_{AS} MOLE CM ² /SEC	θ_b SEC	θ_M SEC
AG	Ag ₂ O	1273	ABSENT	9.6 E-13	1.5 E9	3.2 E3
AL	Al ₂ O ₃	973	B	1.2 E-12	1.1 E1	2.5 E3
			C	2.8 E-51	4.6 E39	1.1 E42
CU	CuO	1473	ABSENT	7.9 E-13	1.8 E9	3.8 E3
CO	CoO	1823	ABSENT	6.4 E-13	1.6 E8	4.7 E3
FE	FeO	1823	C	3.3 E-16	4.0 E11	8.9 E6
NI	NiO	1773	ABSENT	6.6 E-13	5.8 E8	4.6 E3
PB	PbO	650	B	1.8 E-12	1.4 E4	1.7 E3
			C	2.2 E-26	1.1 E18	1.3 E17
SN	SnO ₂	550	B	2.1 E-12	2.8 E1	1.4 E3
"	"	1023	B	1.1 E-12	2.7 E6	2.6 E3
			C	9.9 E-21	4.3 E14	3.0 E11



$$\log \epsilon_0 = 1.07 \log \Phi^2 - 3.9$$

