2/18/2019 last time ... 2H, (g) + 2NO(g) -> N2(g) + 2H2O(g) - determined rate = K [H2] [NO] via XPTs! RATELAW Q: if [H2] = 0.DIOM and [NO] = 0.0025M and experimentally rate = 3.5×10-6 M·s-1 - What is the rate constant? K K = rate = 3.5×10-6 M.s-1 [H2][N0]2 [0.010M][0.0025M]2 = 56 M.s or s or M-2 -1 OR 1 OR 1 OR L2 OR L2.mol-25-1 ex: What would rate be if [H\_]=[NO] = 0.50M? rate = K[H2][N0] = 56 M-3-1 x (0.50M) x (0.50M)  $= 7 M^{-2} s^{-1} M^{3}$ = 7.0 Ms-1

How do we determine a rate law?	
Do XPTS! - Common to use method of initial rates	•
-need to know init cones.	
- mix reactants.	
- measure init. rate of rxn!	
- we will run multiple trials + try to figure out orders and K from our data!	
ex: $NO_2(g) + (O(g) \longrightarrow NO(g) + (O_2(g))$	
assume rate law is: rate = $K[NO_2][CO]$ $time = 0$ XPT# [NO <sub>2</sub> ] (M) [CO] <sub>0</sub> (M) Init (M) rate	. · s-')
1 0.10 0.10 0.0021 2 5 0.20 5 0.10 5 0.0082 3 0.20 same 0.20 doubles 0.0083	Same
rate & [NOz] -> if m=0, rate & [NOz] shoulds	$\hat{x}$
=> [0] -> if m=1, rate ~ [NO2] : double -> d NO2 come ro	
→ if m=2, rate or [NOz]?: double → que  ⇒ 2nd order wrt NOz!  NOzrane rate	adruple's
=> 2nd order wit No.!	" (V)

rate & [NO2] (why? x2[NO2], x4 rate) at x [(0]° (why? ,2(0), x1 at.) => rate = K[NO<sub>2</sub>]<sup>2</sup>[co]<sup>o</sup>

rate = K[NO<sub>2</sub>]<sup>2</sup> K? K = rati #id 0.0021 M·s-1 = 0.21 M's-1

[0.10M]2 = always for 2nd order! We'll do a bunch of these cales in lab!

- we'll also need to use logarithms

(see p633) The integrated rate law: dependence of [] on hime! -useful to know how [] varies wy time