<u>Reaction kinetics notes</u>





Reaction kinetics notes

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Vote = k [a] m [B] n where: K is a constant, and m, n shows the orders of the reaction regarding each reagent. It is an experimentally determined equation as information (n, m, k etc.) can be found only through theoretical considerations. The equation of the rate shows the relationship between the speed of a reaction and the concentrations of individual reagents. Once you find orders, they provide information about the mechanism of the specific reaction. The total order of the reaction is the total order of the reaction is the total order of the reaction. catalysts influence the value of the speed constant. Graphics results If a time reaction concentration chart is drawn a curve is obtained while the reagent are exhausted the rate decreases. If a rate chart is drawn against time, the form obtained will depend on the overall order of the reaction. 0 ° graphic order: - straight flat line 1st order chart: - straight line All other orders: curve If a curve is obtained, further mathematical treatment of results is required. How: Vote = K[A] m[B] n If [B] is kept constant, then: Log Rate = Log K '+ M Log [A] and a log frequency chart against the register [A] will give a line of gradient M (the graph has the Y = MX + C) Maintain constant and treating the speed results when [B] varies allow a similar determination of the order compared to B. Once the two orders are Accerted then the constant K tariff can be found. Half-life Half-life is the time used for concentration of reagents to reach half of its original value. For most reactions, half of life changes as reaction procedures, but this is not the case of reactions of the first orders where half of life is constant rate is possible to find from a graph of concentration/time taking a point, finding its concentration, then finding a point on the chart that corresponds to half this concentration. Half of life is time between these two points. Half of life is also equal to LN2 / K where K is the constant rate (equation provided in the data book). Solve the rate equation for inspection If a number of experimental results are obtained at rate at different reatrimony concentrations, the rate variation can be ascertained for such reactions when the concentration of a low concentration b 1 0.1m 0.1m 6 x 103 2 0.2m 0.1m 1.2 x 104 3 0.4m 0.1m 2.4 x 104 4 0.1m 0.2m 6 x 103 5 0.2m 0.2 M 1.2 x 104 In experiments 1.2 and 3 the concentration of a change while the concentration of B is maintained constant. means that the rate equation can be written as: Rate = k'[A]m for these three experiments. If we check the rate of experiments 1,2 and 3 we see that as the concentration of A is doubled so that the rate doubles. In other words, the reaction order must be 1 so that whatever happens to the concentration must also occur in quantities equal to the rate. Similar inspection of B is changed. The order with respect to B must be 0. Orders can now be substituted in the rate equation: Rate = k[A]1[B]0 To get a value of the rate constant, we simply substitute the values of one of the above experiment 1) Values from experiment 1 6 x 103 = $k \times [0.1]1 \times [0.1]0$ k = 6 x 103 / 0.1 k = 6 x 104 Solve the tariff equation - test yourself 16.2 - Reaction mechanism The mechanism of a reaction is a series of reaction which eventually lead to the final products. A reaction which eventually lead to the final products of a reaction which eventually lead to the final products. Molecularity The number of particles that react in the rate determining the stage of a reaction. Activated Complex...not literally a chemical, but an intermediate in which the bonds are in the process of breaking and forming. The Order of Reaction This provides information about the particle sinvolved in the velocity determination step (which is a step in the mechanism). For example, if two of a particle type collide, the order relative to that particle will be 2 (and zero to any other). 16.3 - Energy of activation Arrhenius equation: k = Ae (-Ea/RT) (data book) where: A is a constant relative to the number, orientation and frequency of collisions occurring between the particles in the results can be plotted on a graph to get a value for the activation energy for a specific reaction. k = Ae (-Ea/RT) then: ln k = lnA - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient - Ea/RT A natural register plot of k versus (1/T) will give a straight line of gradient The reagents and products are of Chemical energy (with the distance between the highest point) high) be equal to the energy of activation of the reaction forward.) an energy chart showing an esoteric reaction reactions occur only when the reactivative particles have greater energy than the activation energy and are able to overcome the energy barrier of activation. catalysis catalysts provide an alternative mechanism with a lower activation energy. This means that a greater number of collisions will be successful and the reaction proceeds at a faster rate homogeneous catalysts -- catalysts in the same state (that is, solid, liquid or gas) as reactionaries. etrogenous catalysts -- catalysts in a different phase (usually a solid) from reagents and finally producing a path of reaction of less activation energy (and also regenerated at the end of this process.) etrogenous catalysts provide a reactive site on which complex forms are activated, weaken ties and increase the collision rate thus increasing the reaction rate. Examples of catalyzed processes type of oate contact catalyst process (sulphur trioxide) vanadium pentoxide heterogen haber (ammonia) heterogenous alcheni iron hydrogenation nichel heterogen polymerization pt (zeigler natta) heterogenous oxygen polymerization homogeneous catalysts rice resources solve the equation - test 1. Why the slowest step of a mechanism said to be the determination of speed? acts as a bottleneck as it prevents other processes from reaching the end of the reaction and thus determines the overall rate. is similar to the idea of a car journey that must go through a zone of road works. the car will be slowed down directly from the roadworks and this is therefore the crucial factor in determining the overall time of the trip. return this is a digital download, no physical product will be slowed down directly from the roadworks and help you know when and how to use all formulas. then we cut your study time in half and help you pass the test! diFREE preview: kinetics notes preview page 12 50 + pages of examples and explanations on: rates of reaction factors affecting reacting reaction factors affecting reaction factors affectin find the constant rate find the law of rate with three responsives the integrated laws of evaluation mechanisms reaction each page is full of: detailed examples explaining each step common talk and questions Note: There are no refunds on digital notes since there is no way to download them. if you're not sureIt's what you need, send me your program here and e-mail my recommendations. 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