### SPECTROPHOTOMETRIC DETERMINATION OF AN EQUILIBRIUM CONSTANT Lab report

Name: ID #: TA name: Lab Section: Lab Date:

Looking at Table 5 (data sheet), how is K<sub>c</sub> related to the initial concentrations ( [Fes\*]<sub>o</sub> and [SCN-]<sub>o</sub>)? Is it what you expected? (Explain briefly) (5 marks)

Looking at Table 5 in the results, it is somewhat evident that Kc is actually independent of the initial concentrations of the reactants since they are relatively consistent among the five test tubes. Furthermore, the Kc values are much more relative as opposed to the significant changes in Kc due to temperature variation which holds true because the equilibrium constant is affected by temperature.

However, looking at the data for any trends, it appears that Kc drops as the concentration of [SCN-] increases, but this should not be true. So, accounting for relative human error and limited machine capabilities, the value only fluctuated minimally from the average which was 204 (M-1). Thus the results are not exactly as I had expected, but ignoring the small deviations, the results are based on Le Chatelier's Principle which states that when a chemical system in equilibrium is disturbed there will be a shift in the system to reduce the effect of that disturbance. So, by increasing [SCN-] from tube 1 to 5, there should be no change of Kc since equilibrium will shift to the right of the reaction to increase the concentration of [FeSCN<sup>2+</sup>] and balance out.

$$Kc = \frac{[FeSCN^{2+}]eq}{[Fe^{3+}] * [SGN^{-}]}$$

To obtain [FeSCN2+]eq in tube 5, you made the assumption that 100% of the ions SCN- had
reacted. Now that you know the value for Ke (average in table 5), build an ICE diagram to
calculate the true percentage of SCN- reacted in tube 5. Comment on the validity of the
assumption. (Make sure you include the ICE diagram in your answer) (10 marks)

	[Fe-3]	+ [SCN-]		
Initial	0.180 M	0.000199 M	0 M	
Change	-x	-x	+x	
Equilibrium	(0.180 - x) M	(0.000199 - x) M	x M	

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#### **Table of Contents Equilibrium Constant Lab Report Discussion**

- 1. Understanding the eBook Equilibrium Constant Lab Report Discussion
  - The Rise of Digital Reading Equilibrium Constant Lab Report Discussion
  - Advantages of eBooks Over Traditional Books
- 2. Identifying Equilibrium Constant Lab Report Discussion
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an Equilibrium Constant Lab Report Discussion
  - User-Friendly Interface
- 4. Exploring eBook Recommendations from Equilibrium Constant Lab Report Discussion
  - Personalized Recommendations
  - Equilibrium Constant Lab Report Discussion User Reviews and Ratings
  - Equilibrium Constant Lab Report Discussion and Bestseller Lists
- 5. Accessing Equilibrium Constant Lab Report Discussion Free and Paid eBooks
  - Equilibrium Constant Lab Report Discussion Public Domain eBooks
  - Equilibrium Constant Lab Report Discussion eBook Subscription Services

- Equilibrium Constant Lab Report Discussion Budget-Friendly Options
- 6. Navigating Equilibrium Constant Lab Report Discussion eBook Formats
  - o ePub, PDF, MOBI, and More
  - Equilibrium Constant Lab Report Discussion Compatibility with Devices
  - Equilibrium Constant Lab Report Discussion Enhanced eBook Features
- 7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of Equilibrium Constant Lab Report Discussion
  - Highlighting and Note-Taking Equilibrium Constant Lab Report Discussion
  - Interactive Elements Equilibrium Constant Lab Report Discussion
- 8. Staying Engaged with Equilibrium Constant Lab Report Discussion
  - o Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - Following Authors and Publishers Equilibrium Constant Lab Report Discussion
- 9. Balancing eBooks and Physical Books Equilibrium Constant Lab Report Discussion
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection Equilibrium Constant Lab Report Discussion
- 10. Overcoming Reading Challenges
  - $\circ\,$  Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
- 11. Cultivating a Reading Routine Equilibrium Constant Lab Report Discussion
  - Setting Reading Goals Equilibrium Constant Lab Report Discussion
  - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Equilibrium Constant Lab Report Discussion
  - Fact-Checking eBook Content of Equilibrium Constant Lab Report Discussion
  - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
  - Utilizing eBooks for Skill Development
  - Exploring Educational eBooks
- 14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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