國立交通大學 104 學年度碩士班考試入學試題

新4544 総積分同試製品用:104年2月7日第3節分學和別:出科系丙組、分醫所第1頁,共≥頁

系所班別:生物科技學系分醫組別:生科系丙組、分醫門

【不可使用計算機】*作答前請先核對試題、答案卷(試卷)與准考證之所組別與考科是否相符!

1. To calculate

(a) (3%) Given
$$p > 1$$
, $\int_{1}^{\infty} \frac{1}{x^{t}} dx = ?$

(b) (3%)
$$\lim_{n \to \infty} \frac{2n^5}{1^2 + 2^2 + \dots + n^2} = ?$$
(c) (4%)
$$\lim_{x \to -\infty} \frac{\sqrt{x^2 + 9}}{3x + 6} = ?$$

(c) (4%)
$$\lim_{x \to -\infty} \frac{\sqrt{x^2 + 9}}{3x + 6} = 7$$

2. To evaluate

(a) (4%)
$$\lim_{x\to 0} \frac{x-\sin x}{x^3}$$

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(b) (4%) $\lim_{x\to 0} \frac{(s^{2X}-1)\ln(1+x^5)}{(1-\cos 3x)^2}$

3. To determine whether the given series converges or diverges by any appropriate test.

(a) (4%)
$$\sum_{m=1}^{\infty} \left| \sin \frac{1}{n^2} \right|$$

(b) (4%) $\sum_{m=1}^{\infty} \frac{1+n^{4/5}}{2+n^{5/5}}$
(c) (4%) $\sum_{m=1}^{\infty} \frac{n!}{n^2 e^n}$

(b) (4%)
$$\sum_{n=1}^{\infty} \frac{1+n^{4/3}}{2+n^{5/3}}$$

(c) (4%)
$$\sum_{n=1}^{\infty} \frac{n!}{n^2 e^n}$$

4. Let

$$f(x) = \sum_{k=0}^{\infty} \frac{2^{2k} k!}{(2k+1)!} x^{2k+1} = x + \frac{2}{3} x^3 + \frac{4}{3 \times 5} x^5 + \frac{8}{3 \times 5 \times 7} x^7 + \cdots$$

(a) (5%) Find the radius of convergence of this power series.

(b) (5%) Show that f'(x) = 1 + 2xf(x).

(c) (5%) What is $\frac{d}{dx} (e^{-x^2} f(x))$?

(d) (5%) Express f(x) in terms of an integral.

5. (14%) At the critical point, gases obey the following equations:

$$\left(\frac{\partial P}{\partial V_m}\right)_T = 0$$
 and $\left(\frac{\partial^2 P}{\partial V_m^2}\right)_T = 0$

where P is pressure, T is temperature and V_m is molar volume. If a certain gas behaviors like van der Waals gas, $P = \frac{RT}{(V_m - b)} - \frac{a}{V_m^2}$, show that the relation between critical point constants $(V_{m_C}; T_C; P_C)$ and van der Waals coefficients (a and b) as following:

$$V_{m_C} = 3b; T_C = \frac{8a}{27Rb}; P_C = \frac{a}{27b^2}$$

國立交通大學 104 學年度碩士班考試入學試題

科目:微積分(4531)

考試日期:104年2月7日 第 3節

系所班別:生物科技學系

系所班別:生物科技學系 組別:生科系丙組 第 / 頁,共 ≥ 頁 【不可使用計算機】*作答前請先核對試題、答案卷(試卷)與准考證之所組別與考科是否相符!!

6. (12%) The change of the enthalpy H could be represented as following:

$$dH = \left(\frac{\partial H}{\partial T}\right)_{p} dT + \left(\frac{\partial H}{\partial P}\right)_{T} dP$$

where $\left(\frac{\partial H}{\partial p}\right)_{\perp}$ indicates how enthalpy changes with pressure under constant temperature and it obeys the equation shown below:

$$\left(\frac{\partial H}{\partial P}\right)_T = V - T \left(\frac{\partial V}{\partial T}\right)_P$$

where P is pressure, T is temperature and V is volume.

- (a) Assume argon is an ideal gas, show that $\left(\frac{\partial H}{\partial P}\right)_{\tau} = 0$. (Ideal gas law: PV=nRT, where R is gas constnat)
- (b) Calculate ΔH for the isothermal reversible expansion of 1 mole argon for an initial volume of V_1 to final volume of V_2 at constant temperature of T.
- 7. (24%) The restriction enzyme EcoRI catalyses the cleavage of DNA and bring about the sequence of reactions:

Supercoiled DNA \rightarrow open circle DNA \rightarrow linear DNA

To illustrate the kinds of consideration involved, we suppose the a reaction takes place in two steps. First an intermediate I (the open circle DNA) is formed from the reactant A (the supercolled DNA) in a first-order reaction with reaction k₁. Then I decays in a first-order reaction with rate k₂ to form the product P (linear DNA):

$$A \stackrel{k_1}{\to} I \stackrel{k_2}{\to} P$$

The initial conditions for this reaction are $[A]_0 = A_0$ and $[I]_0 = [P]_0 = 0$. Also at all time, the relationship between the concentrations of all species is $[A]_t + [I]_t + [P]_t = A_0$.

- (a) All the steps in the reaction scheme are elementary reactions. Express $\frac{d[A]}{dt}$, $\frac{d[I]}{dt}$, and $\frac{d[P]}{dt}$ in terms of reaction constants (k1 or k2) and the concentration of all related species ([A] or [I]).
- (b) Derive that $[A]_t = A_0 e^{-k_1 t}$
- (c) From (a) and (b), then derive that:

$$[I]_t = \frac{k_1}{k_2 - k_1} A_0 (e^{-k_1 t} - e^{-k_2 t})$$

- (d) Derive $[P]_t$ in term of A_0 , k_1 , and k_2 , using a relationship of $[A]_t + [I]_t + [P]_t = A_0$.
- (e) In the case of $k_1 \gg k_2$, all of the A initially present is rapidly converted into I, which is then slowly used up to

form C. Show that $[C]_t = [1 - \exp(-k_2 t)]A_0$.

(f) From (c), when would the intermediate reaches to its maximum concentration.