Determination of Half-life of Dissolution Process of Oxaliplatin in DMSO

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The enthalpies of dissolution for oxaliplatin in dimethyl sulfoxide were measured using a RD496-CK2000 calve microcalorimeter at 309.65 K under atmospheric pressure. Differential enthalpies and molar enthalpies were determined for oxaliplatin in dimethyl sulfoxide, then the relationship of the heat effect of each process with the amount of the substance was setted up. On the basis of the dynamics equation, the process of dissolution is a pseudo first order reaction was indicated. Meanwhile, the half-life period $t_{1/2} = 59.32$ min, the molar enthalpy $\Delta_{\text{sol}}H_m = 14.996$ kJ mol⁻¹, the molar gibs free energy $\Delta_{\text{sol}}G_m = 97.93$ kJ mol⁻¹, the molar entropy is $\Delta_{\text{sol}}S_m = -267.81$ J mol⁻¹ K⁻¹ were calculated. The result not only provides a simple method for the detemination of the half-life for drugs, but also offer a theoretical reference for the clinical application of oxaliplatin

Keywords: Thermodynamics, Kinetics, Dissolution heat, Half-life, Oxaliplatin.

INTRODUCTION

The systematic name of oxaliplatin is oxalato (*trans*-(-)-1,2-cyclohexanediamine) platinum (II), for short L-OHP. It was synthesized by the Japan's Kidani in 1978 at first. Following the cisplatin and carboplatin, it was called the third generation of platinum anticancer drugs, which was developed by Swiss Debiopharm company, produced and soled by Sanofi company in France in 1996. Now it is applied by many countries including the United States¹⁻⁶. Meanwhile, capecitabine platinum developed successfully by Nanjing pharmaceutical in China, which was qualified for new drug registration and quality certification. But its unique mechanism of action and high anticancer activity were focused by the international medical community on tumor. Academic researchers on drugs have obtained the half life based on the principle of pharmacokinetics using HPLC to detect the drug concentration in blood. It is an effective method, but the procedure is complicated^{7,8}. So a simple and feasible method to carry out experiments and to draw reliable data is particularly important.

Aiming at the above problems, we have gained the half-life of the drug by calorimetry. The conclusions showed that the results obtained by the calorimetry method were tallied with what by the pharmacokinetics. It is easy to be operated. Therefore, it will have an important significance to enhance the quality of medicines by studying its thermodynamic functions and determining the kinetic parameters.

EXPERIMENTAL

Oxaliplatin (99 %, Kunming Guiyan Pharmaceutical Co., Ltd.). DMSO: analytically pure.

Equipment and conditions: The experiment was performed using a RD496-2000 Calvet Microcalorimeter (Mianyang CAEP Thermal Analysis Instrument Company, China). The microcalorimeter was calibrated by Joule effect and its sensitivity was 64.22 ± 0.04 $\mu\text{V mW}^{-1}$ at 309.65 K. The enthalpy of dissolution of KCl (spectrum purity) in distilled water (about 20 mg/2.00 g) measured at 298.15 K was 17.535 kJ mol⁻¹, which was in an excellent accordance with the literature value⁹ 17.536 kJ mol⁻¹, showing that the device of measuring the enthalpy used in this work was reliable.

The proper quality of oxaliplatin (12.65 mg, 13.22 mL, 14.43 mL, 15.30 mL, 15.95 mL) were dissolved in DMSO with 1.50 mL, respectively, at 309.65 K under the atmospheric pressure. The enthalpy change of the process was detected by the RD496-2000 Calvet Microcalorimeter.

RESULTS AND DISCUSSION

Thermochemical behaviours of the dissolution of oxaliplatin: The certain quality of oxaliplatin was dissolved in DMSO at 309.65 K. There are five concentration gradients to carry out in this experiment. The curve describing the entire dissolution process of oxaliplatin in DMSO is shown in Fig. 1. The dissolution is of an exothermic process. The entire process

was repeated three times. The heat flow curves obtained under the same conditions overlap with each other, indicating that the reproducibility of the test is satisfactory.

Fig. 1. Heat release rate ($\Delta H/\Delta t$) in the entire dissolution process of oxaliplatin in DMSO

Table-1 shows the experimental data obtained from the typical thermogram curve of the dissolution with different volume mixture solution B in 1.5 mL DMSO.

Sample mass (mg)	$n/(\times 10^{-3} \text{ mol})$	Q (mJ)	$\Delta_{\text{sol}}H_m$ (J mol $^{-1}$)
12.65	0.032	470.87	14713.11
13.22	0.033	495.67	14820.23
14.43	0.037	545.91	14953.70
15.30	0.039	591.53	15281.96
15.95	0.040	613.96	15215.05
Average			14996.81

n : the amount of paclitaxel; Q : the heat effect of the dissolution process; $\Delta_{\text{sol}}H_m$: the molar enthalpy the dissolution process.

Table-1 showed the concentrations of the solution almost have little influence on the values of the molar enthalpy ($\Delta_{\text{sol}}H_m$) at 309.65 K. So the average value of $\Delta_{\text{sol}}H_m$ which is 14.996 kJ mol $^{-1}$ can represent the molar enthalpy of the infinite diluted DMSO at 309.65 K.

The heat effect vs. the amount of the substance relationships of oxaliplatin in DMSO is shown in Fig. 2.

The linear equation for the DMSO is as follows:

Fig. 2. Relationship between the amount of oxaliplatin (n) and the heat Q dissolved in DMSO

$$Q = 17416000 n - 86.935, r = 0.9989 \quad (1)$$

where, r is correlation coefficient.

The differential enthalpy ($\Delta_{\text{dif}}H_m$) is obtained from the equation (1).

Kinetics of dissolution process of oxaliplatin in DMSO:

The kinetic equation describing the dissolution of oxaliplatin in DMSO⁸ is eqns. 2 and 3 is chosen as the model function describing the process:

$$\frac{d\alpha}{dt} = kf(\alpha) \quad (2)$$

$$f(\alpha) = (1 - \alpha)^n \quad (3)$$

Combining eqns. 2 and 3, substituting $\alpha = \frac{H_1}{H_0}$ into the equation and then get a logarithmic converter:

$$\ln \left[\frac{1}{H_0} \left(\frac{dH}{dt} \right)_i \right] = \ln k + n \ln \left[1 - \left(\frac{H}{H_0} \right)_i \right] \quad i = 1, 2, \dots, L \quad (4)$$

In these kinetic function; H represents the heat at time of t ; H_0 is the heat of the whole process; k is the dissolution rate constant of oxaliplatin in DMSO; n is the reaction order and L is the counting number.

By substituting the data taken from Table-2, $(dH/dt)_i$, $(H/H_\infty)_i$, H_∞ , $i = 1, 2, \dots, L$, into the kinetic eqn. 4, the obtained values of n and $\ln k$ are listed in Table-2.

m (mg)	t (s)	dH/dt (mJ s $^{-1}$)	H_t (mJ)	H_t/H_0	H_∞ (kJ mol $^{-1}$)
	0	0.079	172.2	0.366	
	20	0.075	196.1	0.416	
	40	0.071	218.8	0.465	
	60	0.067	240.2	0.510	
	80	0.063	260.4	0.553	
	100	0.059	279.3	0.593	
	120	0.055	297.0	0.631	
	140	0.051	313.4	0.666	
	160	0.047	328.6	0.698	
	180	0.044	342.7	0.728	
12.65	200	0.040	355.7	0.755	14.71

Note: The data of Oxaliplatin (13.22 mg, 14.43 mg, 15.30 mg, 15.95 mg) dissolved in DMSO with 1.50 mL were omitted for verbosity.

Substituting the values of n and k in Table-3 into eqns. 2 and 3 and then uniting them, the kinetic equation of the dissolution process can be deduced to be

$$\frac{d\alpha}{dt} = 10^{-3.71} (1 - \alpha)^{0.93} \quad (5)$$

Sample mass (mg)	n	$\ln k$	r
12.65	0.72	-8.36	0.9996
13.22	1.12	-8.85	0.9996
14.43	1.02	-8.56	0.9996
15.30	0.91	-8.48	0.9995
15.95	0.89	-8.46	0.9995
Average	0.93	-8.54	0.9996

The kinetic equation is similar to quasi-first order reaction of the dissolution process. So the half-life period can be calculated with eqn. 5 and its value is 59.32 min.

$$t_{1/2} = \frac{\ln 2}{k} \quad (6)$$

Thermodynamics of oxaliplatin in DMSO: On the basis of these experimental data and calculated results, the kinetic parameters of the dissolution process were inferred through eqn. 7.

$$\ln \frac{k \cdot h}{k_B \cdot T} = \frac{\Delta_{\text{sol}} S_m}{R} - \frac{\Delta_{\text{sol}} H_m}{RT} \quad (7)$$

Substituting $k_B = 1.380 \times 10^{-23} \text{ J K}^{-1}$, $h = 6.626 \times 10^{-34} \text{ J s}$, $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$, $\Delta_{\text{sol}} H_m = 14.996 \text{ kJ mol}^{-1}$, $T = 309.65 \text{ K}$, so $\Delta_{\text{sol}} S_m = -267.81 \text{ J}$.

And then putting $\Delta_{\text{sol}} H_m$ and $\Delta_{\text{sol}} S_m$ into the following formula

$$\Delta_{\text{sol}} G_m = \Delta_{\text{sol}} H_m - T \Delta_{\text{sol}} S_m \quad (8)$$

We can ascertain $\Delta_{\text{sol}} G_m = 97.93 \text{ kJ mol}^{-1}$.

Conclusions

- The experiment results show that microcalorimeter method can obtain the half-life of drug easily.

- The kinetic equation of the dissolution process of oxaliplatin in DMSO at 309.65 K is $\frac{d\alpha}{dt} = 10^{-371} (1 - \alpha)^{0.93}$. It is a quasi-first order reaction and its half-life is $t_{1/2} = 59.32 \text{ min}$, the rate constant $k = 1.96 \times 10^{-4} \text{ s}^{-1}$.

- The dissolution of oxaliplatin in normal saline is an exothermic process. The molar enthalpy ($\Delta_{\text{sol}} H_m$) is $14.996 \text{ kJ mol}^{-1}$ and $\Delta_{\text{sol}} S_m$ is $-267.81 \text{ J mol}^{-1} \text{ K}^{-1}$. The negative value of entropy of activation indicates that the dissolution of oxaliplatin in DMSO get a more ordered system.

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REFERENCES

1. Y. Kidani and K. Inagaki, *cis*-Platinum(II) complex of *trans*-1,1,2-diaminocyclohexane, US Patent 4169846 (1979).
2. J. Capdevila, E. Elez, S. Peralta, T. Macarulla, F.J. Ramos and J. Tabernero, *Expert Rev. Anticancer Ther.*, **8**, 1223 (2008).
3. N. Sawabu, *Nippon Shokakibyo Gakkai Zasshi*, **100**, 645 (2003).
4. J. Trédaniel, C. Becht, M. Bekradda, H. De Cremoux, J. Alexandre, F. Chomy, S. Szyldergemajn, Y. Yataghene and S. Culine, *Lung Cancer*, **63**, 259 (2009).
5. A. Gunnlaugsson, H. Anderson, E. Fernebro, E. Kjellén, P. Byström, Å. Berglund, M. Ekelund, L. Pählman, T. Holm, B. Glimelius and A. Johnsson, *Eur. J. Cancer*, **45**, 807 (2009).
6. Z. Wang, J. Zhou, J. Fan, S.-J. Qiu, Y. Yu, X.-W. Huang, J. Sun, C.-J. Tan and Z. Dai, *Expert Opin. Investig. Drugs*, **18**, 1595 (2009).
7. W.L. Lin, D.G. Li, Q. Chen, H.M. Lu, X.M. Ma and P.L. Sun, *World Chin. J. Digestol.*, **11**, 1535 (2003).
8. M. Galanski, A. Yasemi, M.A. Jakupiec, N. Grafv. Keyserlingk and B.K. Keppler, *Monatsh. Chem.*, **136**, 693 (2005).
9. R.L. Montgomery, R.A. Melaugh, C.-C. Lau, G.H. Meier, H.H. Chan and F.D. Rossini, *J. Chem. Thermodynamics*, **9**, 915 (1977).