

Thermodynamics and Mutual Distribution of Macromolecules in the Systems: Chlorine-Chlorocontaining Polymers



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Abstract: In given work a scientific base compatibility of polymers, produced by industry have been studied by revealing of thermodynamically regularities of polymer mixtures.

The most significant thermodynamic regularities of the mixtures of polyolefin, produced by industry and heterochained polymers have been revealed and on this base the scientific bases of their compatibility have been developed for creation of multicomponent compositional materials with given complex of operating properties.

Key words: polymer-polymer mixtures, chlorinated polyethylene (CPE), polymethylmetacrylate (PMMA), polyethylene of low density (PELD), chlorocarboxylated polyethylene of low density (CCPE) chlorinated PELD (CPE), polyvinyl chloride (PVC), criteria of compatibility, chlorination, upper and lower critical temperature, specific holding of volume, Gibbs mixing energy, parameter of solubility, distribution of macromolecules, chlorine-chlorocontaining polymers, polar groups, thermodynamically compatibility, sorbate, gas-carrier.

I. INTRODUCTION

Need of industry in new polymer materials, possessing by necessary properties for work in extremal conditions continuously increase. It is may be achieved by synthesis of the new types of polymers and creation of polymer compositions where properties of the separate components additively are the complement of one another [1-5]. Therefore development of different polymer compositions by simplification of technology its production and processing, shortening expenditures on production of materials with necessary complexes of properties is extremely actual. For production of the new types of polymer materials by use of polymer-polymer mixtures determination of its thermodynamically stability as one of the factors, deterring a structure and properties of the final product acquire essential interest [6-8]. Use of chlorcontaining polymers in industry, stipulated by complex of enough good properties determine the desire to improvement of series them indices. From this point of view research of thermodynamical conduct of chloropolymers mixtures with another chloropolymers, having in their composition the polar groups of chloropolymers present a special interest.

In the work [9] the thermodynamically compatibility of chlorinated polyethylene (CPE) with different chlorination degree, shown that by dominant of dispersion forces compatibility is possible by proximity parameters of solubility and interaction. Analysis of compatible and incompatible pairs shown that:

- by difference in chlorine content, reaching
- 18 mas % ΔH_m is relatively high (+0.519 Jou/g)

Owing to big difference in thermal coefficients expansion - $T\Delta S_m$ is positive (+0.276 Jou/g). The total energy ΔG_m is positive (+0.795 Jou/g). From this it is concluded that a mixture must be inmixing (incompatible);

- by approximately near content of chlorine (27 and 35 mas%) the dispersion forces between molecule, owing to similarity lead to little interaction degree. By calculation, a big difference in thermal coefficients expansion gives a little negative value ΔH_m (-0.108 Jou/g). But this difference causes a big positive - $T\Delta S_m$ (+0.761 Jou/g) and in result ΔG_m (+0.653 Jou/g) doesn't favorable to mixing.

II. METHOD

It have been determined, that uniting criteria of compatibility $\Delta G_m < 0$ [7, 8] in total case is carried out, although a structural compatibility is possible also by little positive values of ΔG_m . In all cases the surplus energy ΔS_m doesn't favorable to mixing, but by proximity of percent content of chlorine and coefficient of thermal expansion the compatibility is possible.

In the work [9] the system (CPE - PMMA) have been studied. It was shown that polymethylmetacrylate (PMMA) is compatible with chlorinated polyethylene by appointed chlorination degree of the last and only by appointed temperatures. It have been determined that a mixtures have an upper and lower critical temperatures of solubility.

The present work devotes to study of thermodynamic interaction between components mixture of chlorocontaining polymers, one of which in its composition have a polar carboxylic groupings. By that the method of turned gas chromatography have been used, which at the last time is widely applied for study of polymers compatibility.

As initial raw materials the industrial samples of polymers: chlorocarboxylated polyethylene of low density PELD (CCPE) with contact of chlorine equal 27 mas % and carboxylic groups 1.2 mas. % correspondingly, chlorinated PELD (XPE) with chlorine content 35 mas. % and polyvinylchloride (PVC) by brand C-70 have been used, the characteristics of which are presented in the table 1

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Table1 Characteristic of investigated polymers

polymer	content of chlorine mas. %	content of carboxylic groups, mas. %	alcohol ethyl propyl alcohol	D:B:O	$M \cdot 10^{-4}$	$M \cdot 10^{-3}$	M/M	by 298 K, g/sm ²	$k \cdot 10^{-4}$	T _c K
CCPE	2.7	1.2	-	-	3.5	5.7	6.11	1.0713	8.64	239
CPE	35	-	-	-	3.5	5.8	6.08	1.0713	5.80	263
PVC	57	-	-	-	15.0	47.0	3.19	1.3446	3.09	349
packing products	-	-	-	4:3:1	3.0	10.2	2.95	1.2284	4.63	238
SKEP	-	-	1.0	-	9.0	30.2	2.98	0.8723	8.59	2.11

D:B:O* – 4.4 – diphenylmethanediisocyanate;
1.4

butadiol:oligoethylenebutyleneglycoladipinate

For research the chromatographical columns with individual polymers and its mixtures by correlation of components 30:40, 50:50, 70:20 have been prepared/ In the work the chromatograph “Tsvet-100” with flame ionization detector have been used. As a gas-carrier the nitrogen and as sorbate ten solvents, relating to different classes (alcohols, ketons and chloroderivatives of aliphatic hydrocarbons) have been used.

The temperatures of chromatographical measuring were higher of temperatures T_c of investigated polymers and made up for all systems 353, 373 and 393 K. Method measuring is in detail stated in the work [10].

III. RESULTS

In all investigated systems dependence of specific holding volume of sorbate V_g^q from value of sample within 0.01-0.1 mol doesn't observed. In case of dependence V_g^q from rate of gas-carrier the extrapolation to zero rate of gas-carrier have been conducted. Contribution of adsorption effects to value of holding volumes wasn't taken into consideration because content of polymer phase on Teflon carrier was in all cases more than 6.5%.

On base of experimental data by absolute specific holding volumes of sorbates V_g^0 and specific volumes of polymers the parameters interaction of sorbate with every polymer X_{12}^∞ - X_{13}^∞ and their mixtures $X_{1(23)}^\infty$ have been calculated. The parameters interaction X_{23}^∞ of polymer with one another have been determined. Calculation have been conducted by equations, offered by Patterson with collaborators [11, 12].

$$X_{12}^\infty = \ln \left[\frac{273.2 \cdot (\omega_2 v_2 + \omega_3 v_3)}{P_1^0 \cdot V_g^0 \cdot V_1} \right] - \left(1 - \frac{V_1}{M_2 v_2} \right) - \frac{P_1^0}{RT} \cdot (B_{II} \cdot V_1)$$

$$X_{12}^\infty = \ln \left[\frac{273.2 \cdot g_R \cdot v_2}{P_1^0 \cdot V_g^0 \cdot V_1} \right] - \left(1 - \frac{V_1}{M_2 v_2} \right) - \frac{P_1^0}{RT} \cdot (B_{II} \cdot V_1) \quad (1)$$

Where R – is universal gas constant; $v_2, v_3, V_2 \cdot V_3$ – specific molar volumes of polymers and molecular mass of thermodynamically segments; $\omega_2, \omega_3, Y_2, Y_3$ – mass. volume parts of polymers; P_1^0, V_1, B_{II} – pressure of saturated vapour, molar volume and the second variable coefficient of sorbet by temperature T.

As sorbates it is recommended to use solvents, which minimally combine conduct of both polymers, that is close to ideal solvents of both polymers [10].

The received in that case values $X_{1(23)}^\infty - X_{23}^\infty$ are correct and may be used for valuation of thermodynamical compatibility of polymers [11]. Thus from all spectrum of sorbate by compatibility two TQF and chloroform (CF) have been selected, which for investigated samples of polymers which by temperature of experiment turned out to be the most close to ideal solvents.

On Figure 1 change of the value $X_{1(23)}^\infty$ from composition of both systems is presented. As it is evident, in both cases values differ from additive, moreover for system CCPE-CPE value in the all sphere of investigated composition is lower of additive by increase of difference by rise of CCPE content. For system CCPE-PVC by correlation of components 50:50 value of X_{23}^∞ coincide with additive, that is in presence of total solvent the thermodynamically interaction between components is getting worse.

The received values have been used for calculation of parameters interaction with one another by equation

$$X_{23}^\infty = \frac{X_{12}^\infty Y_2 + X_{13}^\infty Y_3 - x_{1(23)}}{Y_2 Y_3} \quad (2)$$

For both systems in investigate fields concentrations values $X_{23}^\infty > 0$ and strongly are changed by value with change of composition that indicate on change of conduct of composition mixture.



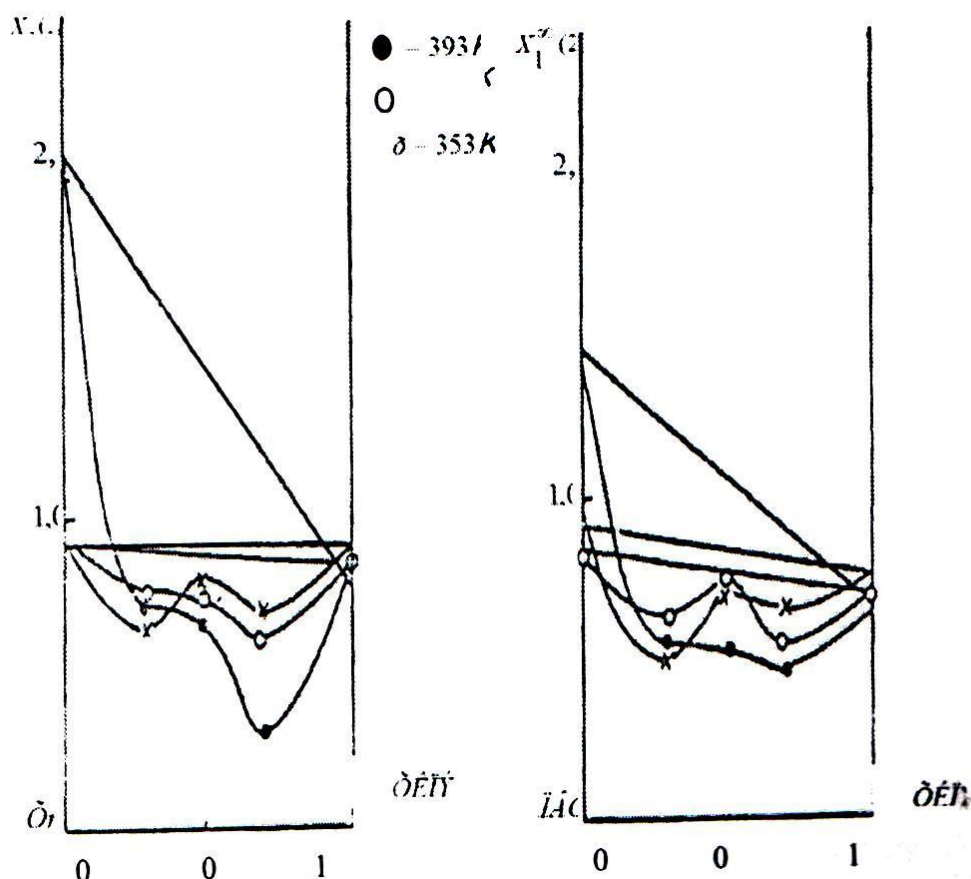


Figure 1. Change of value 1(23) with composition for systems CCPE-CPE-PVC

By little content one of component in mixture the investigation wasn't conducted, because as was indicated in [14] by division of numerator fraction of equation (2) on product $Y_2 \cdot Y_3$ by little values Y_2 or Y_3 the quotient is sharply increased, leading to inequivalent values.

Taking into consideration that value of X_{23}^T equal 0.3-0.4 determine compatibility of the systems from the received experimental data it follows that systems CCPE-CPE and CCPE-PVC are partly compatible.

By experimental received values V_g^0 for sorbates on individual and mixed motionless phases the excess mol mixing energy G^E of polymers, calculated by equation (3) have been determined.

$$G^E = RT(\ln V_{M1} - x_2 \ln V_{M2} - x_3 \ln V_{M3}) \quad (3)$$

Where V_{M1} , V_{M2} , V_{M3} - a model holding volumes of sorbate by mixture and individual components; R , T - universal gas constant and absolute temperature. It is expected that for components of mixture with big molecular mass ΔG^E ideal - 0, the average Gibbs energy, related to 1 g of mixture may be determined by equation (4):

$$\Delta g_x = \frac{G^E}{M} \quad (4)$$

Where M - the average molecular mass of mixed phase.

Having the temperature dependence of quantity Δg_x , a values of enthalpy and entropy have been calculated by equations (5)

$$\Delta h_x = \frac{\partial(g^x/T)}{\partial(1/T)} \quad \Delta S_x = \frac{\Delta h_x - \Delta g_x}{T} \quad (5)$$

On Figure 2 the concentration dependences of received thermodynamical parameters are shown. As it is evident both investigated systems behave itself in analogous way, but for system CCPE-PVC correlation Δh_x and $T\Delta S_x$ is that Δg_x is essentially less by value, than in case of system CCPE-CPE achieving in maximum +0.4 Joul/g, at the same time as for second system +1.6 Joul/g, ie. a system CCPE-PVC is more thermodynamically stable than system CCPE-CPE.

By increase of temperature correlation Δh_x and $T\Delta S_x$ for different compositions undergo essential changes, moreover observed changings lead to changing of Δg_x mark. These data testify about that criterion of availability of lower and upper critical temperature (thermodynamical criterion LCTM - $\Delta h_x < 0$, $\Delta S < 0$ and UCTM $\Delta h_x > 0$, $\Delta S > 0$ are by appointed composition and temperature for both systems).

Phase diagrams by investigated systems have been calculated by equation spinodally:

$$\left(\frac{\partial g^2}{\partial^2 \omega}\right)_{p=0} \quad (6)$$

By help of the plant (Figure 3) analogous with [18] for determination of thermodynamical parameters of interaction sorbate-polymer or polymer-polymer by us the

first derivatives Δg_x by composition, then – the second derivatives by composition have been calculated and graphical dependence $\left(\frac{\partial g^2}{\partial^2 \omega}\right)_{p,p}$ from temperature have been created.

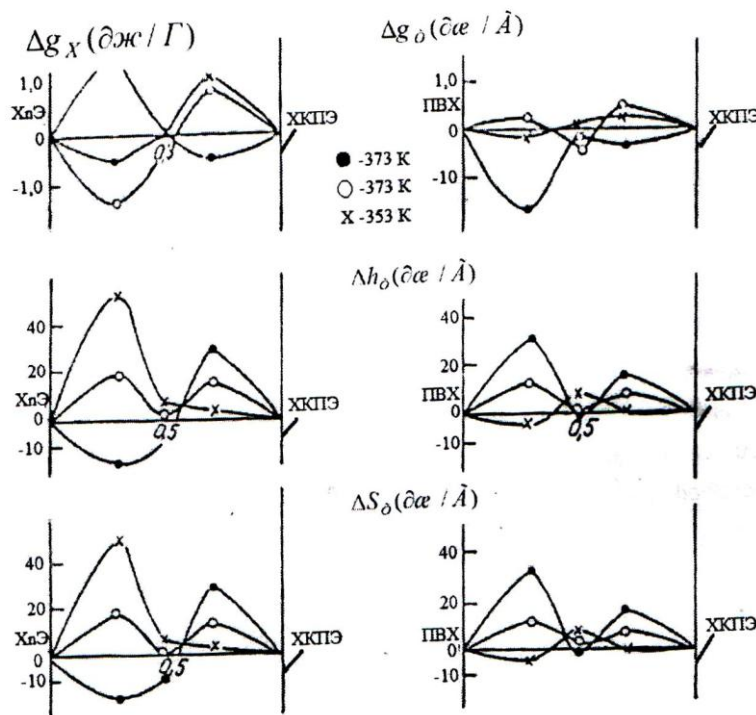


Figure 2. Concentration dependences of thermodynamical parameters in systems CCPE-CPE and CCPE-PVC.

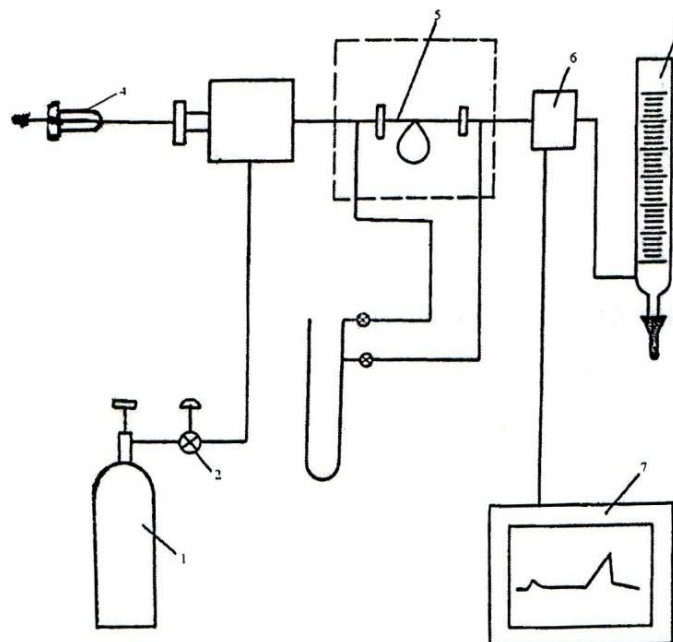


Figure 3. Scheme of plant for determination of thermodynamical parameters of interaction of polymer-sorbent

1-container with gas-carrier; 2-regulator of gas-carrier pressure; 3-microsyringe for introduction in gas line of sorbate microsamples; 4-evaporator of the block sorbate injection; 5- columns with polymer, piled on carrier, put in thermostate; 6-detector; 7-memory apparatus; 8-soap flowmeter

By extrapolation of these dependences to $(\partial g^2 / \partial^2 \omega)_{p=0}$ the temperatures of phase division have

been determined. The received diagrams are shown on Figure4. As it was expected, according to supposition both systems have two critical temperature dissolution – upper and lower. Moreover for system CCPE-PVC the LCTM more than (correspondingly 374 and 366 K). This system within critical points is compatible by all composition field.

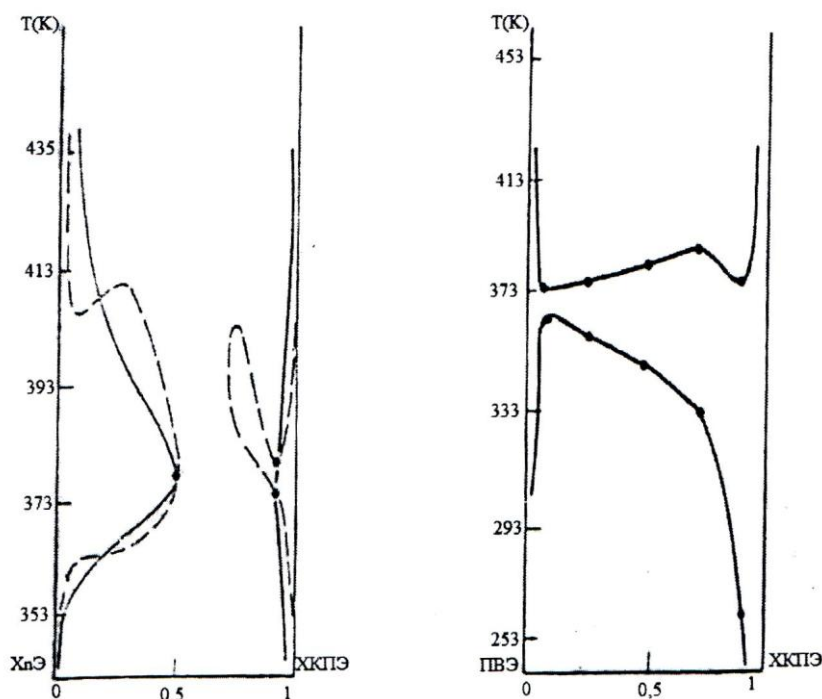


Figure 4. Diagram of the phase division of temperatures

Asymmetry of the phase diagram as was indicated for example in [13], is characteristic for mixture of polymers with considerable difference in values of molecular mass. Observed bipod stance is characteristic by polymers with strong polydispersity [14].

IV. CONCLUSIONS

Analysis of the phase diagram of CCPE-CPE system testify about more the worst compatibility, than in case of CCPE-PVC system. It is interesting to note, that for the first system the UCTM more than LCTM that lead to phase diagram “sand glass” (spinodales become one, giving isthmus) ie. by any temperature it is impossible to get a system being compatible by all field of composition. Compatibility is considerably better by little content of CCPE whereas for system CCPE-PVC compatibility improves by increase of content CPE.

Thus, an experimental comparison of values parameters solubility of polymers, received by two methods: spectra-gas chromatography (SQC) and turbidimetric titration (TDT), characterizing by limit opposite state of polymer by relation to solvent shown the significance difference in parameters solubility for polymers with strong specific interaction between macromolecules, that require attention by use of theory of parameters solubility for supposition of compatibility of polymer pair to method, by which a used value was received. Apparently, that for

crystallizing polymers also the same difference in parameters of solubility must be observed.

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