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137-58-2-2809

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 2, p 87 (USSR)

AUTHORS: Kurilekh, D. G., Psarev, V. I.

The Nature of the Drop in Yield Point in the Finishing of Sheet TITLE:

Steel (O prirode ponizheniya predela tekuchesti pri otdelke

stal'nykh listov)

PERIODICAL: Nauchn. zap. Dnepropetr. un-ta, 1956, Vol 45, pp 69-71

A study was made of the mechanical properties of annealed ABSTRACT:

and leveled sheet steel, and an attempt was made to determine the cause of the drop in the T value after leveling. Used in the tests were sheets of steel 08KP, annealed at 680-710° for 8 hours. The sheets, 2000-2500 mm in length, were cut in half. From the one half samples were cut to be tensile-tested to destruction at angles to the direction of rolling of 0, 30, 45, 60, and 90°; the other half was sent on for leveling, then for mechanical testing. Reduction amounted to 0.85-1.0 percent. Tested were 6 sheets, 120 samples; the thickness of the samples was 0.82 mm. Chemical composition and test-result

tables are included. The penetration depth of the leveling de-

formation was studied by X-ray. It was found that the solid Card 1/2

137-58-2-2809

The Nature of the Drop in Yield Point (cont.)

background of the line (110) receded from the surface to a depth of up to 0.05-0.07 mm in sheets reduced by 1.0 percent, and to a depth of up to 0.08-0.12 mm in sheets reduced by 1.8 percent. In the leveled sheets the O_S value dropped from 24.05-26.7 kg/mm² to 19.35-2075 kg/mm², with a different value for each direction tested. Changes in O_B and O_B were insignificant. It is assumed that the increase in the surface area of a sheet resulting from a plastic flow of the ferrite grains in the surface layers of the sheet should lead to an elastic deformation of the individual grains not only on the surface but throughout the thickness of the sheet. The effect of leveling on the sheet steel was such that the elastic tensile deformation of the inner ferrite grains was balanced out by the elastic compression deformation of the surface layers. When the samples were tensile-tested, the pre-existing elastic deformation of the grains was made evident by its absorption of some of the stress---which was apparent from the drop in the O_S value.

D.M.

1. Sheets-Properties 2. Sheets-Steel-Analysis

Card 2/2

5/126/63/015/002/012/033 E111/E151

AUTHOR:

Psarev, V.I.

TITLÉ:

Coagulation of the carbide phase during tempering of

a molybdenum steel

FERIODICAL: Fizika metallov i metallovedeniye, v.15, no.2, 1963,

234-238

The coagulation of carbides has been studied in two TEXT: molybdenum steels: 1) 0.04% C, 0.34 Mn, 0.17 Si, 1.39 Mo; 2) 1.27% C, 0.46 Mn, 0.21 Si and 1.04 Mo. Specimens with a suitable carbide structure were selected from batches tempered at 600 °C. Coagulation was studied at 600, 650 and 700 °C by the microscopical examination of specimens after quenching and surfacestripping. Coagulation during continuous heating at 10, 50 and C/hour was also studied on steel (1). Crystal-structure changes were studied by X-ray diffraction on carbides isolated electrolytically, the structure obtained under the test conditions being that of cementite. Redistribution of molybdenum between the crystal lattices of alpha-iron and cementite leads to breaks on the curves correlating average carbide-particle radius and Card 1/3

Coagulation of the carbide phase... S/126/63/015/002/012/033

isothermal holding time at 600 and 650 °C; at 700 °C coagulation dies away, virtually ceasing after about 80 hours. The hardness to the changes in the dispersion of the changes correspond carbide phase and in the state of the alpha-solid solution. Comparison of these results and those with carbon steel indicate that molybdenum increases the dispersion of the carbide phase and at the same time greatly retards the coagulation. The results show that carbide coagulation is governed by diffusion of carbon, but also that the state of the matrix crystal lattice greatly affects both the dispersion of the carbide particles and their coagulation. The known fact that alloying steel with a very small amount of Mo does not lead to a large change in the carbon diffusion coefficient although the carbide phase is considerably more dispersed and the carbide coagulation parameters greatly changed, is explicable in terms of the equilibrium conditions in a simplified model. Similar results should be obtained with steels alloyed with other carbideforming elements in quantities insufficient to form special carbides.

There are 4 figures and 1 table.

Card 2/3

S/126/63/015/C02/012/033 E111/E151 Coagulation of the carbide phase ...

ASSOCIATION: Chernovitskiy gosudarstvennyy universitet (Chernovitsy State University)

June 21, 1962 SUBMITTED:

Card 3/3

PSAREV, V.I.; MAKOVIYCHUK, Yu.I.; STEFANOVSKAYA, N.B.

Coagulation of the carbide phase in molybdenum and vanadium steels at temperatures lower than the A₁ point. Izv. vys. ucheb. zav.; chern. met. 6 no.8:120-127 '63. (MIRA 16:11)

1. Chernovitskiy gosudarstvennyy universitet.

S/0137/63/000/012/1034/1034

ACCESSION NR: AR4014151

SOURCE: RZh. Metallurgiya, Abs. 121211

AUTHOR: Psarev, V. I.; Galichuk, Ya. D.; Dobry*den', K. A.

TITIE: On the intergrowth of crystals of the compounds CdSb and Zn3Sb2 in alloys

CITED SOURCE: Nauchn. yezhegodnik za 1959 g. Chernovitsk. un-t. Fiz.-matem. fak. Chernovitsy*, 1960, 617-619

TOPIC TAGS: Cadmium antimonide crystal, Zinc antimonide crystal, crystal aggregation, crystal intergrowth, centrifugal crystallization

TRANSLATION: An investigation was made into the feasibility of consolidating crystals of CdSb and Zn₃Sb₂ in binary alloys of Cd with 25% Sb and Zn with 35% Sb by the mechanical effect of centrifugal forces on a crystallizing casting. The alloys were prepared from c.p. Zn, Sb of grade Su-O, and Cd of grade Kd-O, turn the purified by triple vacuum distillation. The crystallization was carried

Card 1/2

ACCESSION NR: AR4014151

out in evacuated pyrex ampoules, which were cooled together with the furnace after the metal had melted, at rest and with rotation. During the rotation, and in the course of the solidification, the CdSb and Zn₃Sb₂ crystals, which differ considerably in density from the adjoining liquid, grow together into large aggregates. This is illustrated by a comparison of the microstructures of the specimens cooled with and without rotation. V. Zolotorevskiy.

DATE ACQ: 09 Jan64

SUB CODE: ML, PH, CH

ENCL: 00

Card 2/2

KOSTUR, N.L.; PSAREV, V.I.

Methods of doping the crystals of CdSb compounds through alloys of the system Cd - Sb. Fiz. met. i metalloved. 16 no.1:71-79 J1 '63. (MTRA 16:9)

Chernovitskiy gosudarstvennyy universitet.
 (Metal crystals) (Cadmium-antimony alloys)

Methodology involved in heating and cooling according to a previously given law. Inzh.-fiz.zhur. no.1:98-102 Ja '60. (MIRA 13:4)

1. Gosudarstvennyy universitet, g.Chernovtsy. (Furnaces)

BIYUMENFEL'D, V.N.; LYUFUR, S.L.; LIVSHITS, B.S.; PARILOV, V.P.;

PSAREV, S.A.; RODZYANKO, V.Ye.; GOLUBTSOV, I.Ye., otv. red.;

KIRILLOV, L.M., red.; SLUTSKIN, A.A., tekhn. red.

[Methodology for designing the equipment of crossbar automatic telephone exchanges] Metodika rascheta oborudovaniia ATS koordinatnykh sistem; informatsionnyi sbornik. Moskva, Gos. izd-vo lit-ry po voprosam sviazi i radio, 1961. 130 p. (MIRA 15:4) (Telephone, Automatic-Equipment and supplies)

AUTHOR:

Psarev, V. I.

sov/78-3-9-36/38

TITLE:

Computation of the Line of Cementite-Austenite Equilibrium in Fe-C Alloys (Raschet linii tsementitno-austenitnogo ravnovesiya Fe-C splavov)

PERIODICAL:

ABSTRACT:

Zhurnal neorganicheskoy khimii, 1958, Vol 3, Nr 9, pp 2217-2218 (USSR)

The paper under review is an attempt at computing the line of cementite-austenite equilibrium in Fe-C alloys with consideration to the mathematical dependence of the n-grain size of the Fe₃C particle on temperature and concentration. The dependence of the n-grain size of the Fe₃C particle on temperature and concentration

is given in the equations 1 and 2:

$$r_{2} = \frac{r_{1}}{1 - \frac{r_{1}}{a} \left[\ln \frac{c_{0}}{c_{2}} (T_{2} - T_{1}) + T_{1} \ln \frac{c_{1}}{c_{2}} \right]}$$
(1)

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SOV/78-3-9-36/38 Computation of the Line of Cementite-Austenite Equilibrium in Fe-C Alloys

$$r = \frac{r_0}{1 - \left(\frac{r_0}{a} \ln \frac{c_0}{c}\right)T}$$
 (2)

In these equations r_1 represents the n-grain size of the Fe_3C particle at the temperature T1, r2 the medium grain size of $\mathrm{Fe_3^{\,C}}$ at $\mathrm{T_2^{\,\prime}}$, and $\mathrm{c_2^{\,\prime}}$ the respective concentrations at the limits of the particle with the radii r_1 and r_2 .

The above formulae were tested in numerous experiments with highly disperse Fe-C alloys. The dependence obtained between the solubility of the compound $\operatorname{Fe_3} C$ at different temperatures is used for the computation of carbon concentrations in Fe-C systems. There are 2 tables and 3 references, 3 of which are Soviet.

ASSOCIATION:

Chernovitskiy gosudarstvennyy universitet (Chernovtsy State University)

Card 2/3

AUTHOR	Psarev, V.I.	SOV-21-53-4-7/29
TITLE:	Computation of the Activation ation Process in Steel (Raschët koagulyatsii karbidov v stali)	t energii aktivatsii protsessa
PERIODICAL	Dopovidi Akademii nauk Ukrains pp 386-389 (USSR)	s'koi RSR, 1958, Nr 4,
ABSTRACT:	The diffusion character of compersed carbide particles in sociate the activation energy derives a formula for this activation $Q = \frac{R \ln \frac{L}{L}}{L}$ where R is a universal gas convalues of the linear growth respectively.	teels makes it possible to cal- of this process. The author tivation energy:
Card 1/2	T_2 at the same size of the parexperimental investigations f from 1.2 to 1.7% C, at a wide 750 to 950 C, and treated the method. The final result was	rticles. The author carried out or a series of carbon steels, range of temperatures, from results by the least-square

SOV-21-58-4-7/29

Computation of the Activation Energy of the Carbide Coagulation Process in Steel

是这种的大型,我们就是这种的人,我们就是这种的人,我们就是这种的人,但是这种的人,这种是是这种的人,我们也不是这种的人,我们就是这种的人,我们就是这种的人,也可

activation energy (above the critical point A_1) closely agreed with that of carbon diffusion in austenite determined by M.Ye. Blanter \int Ref. 2 \int and others. Similar computations were also performed by the author for various steel alloys. There is 1 graph and 5 references, 4 of

which are Soviet and 1 American.

ASSOCIATION:

Dnepropetrovskiy gosudarstvennyy universitet (Dneprope-

trovsk State University)

PRESENTED:

By Member of the AS UkrSSR, V.N. Svechnikov

SUBMITTED:

July 5, 1957

NOTE:

Russian title and Russian names of individuals and institutions appearing in this article have been used in the

transliteration.

1. Carbides--Diffusion 2. Diffusion--Mathematical analysis

3. Steel

Card 2/2

5/148/62/000/012/003/008 E071/E151

AUTHORS:

Psarev, V.I., and Korobiy, O.I.

TITLE:

Influence of the state of the solid solution on the kinetics of coagulation and dispersion of the carbide

phase in steels

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy. Chernaya metallurgiya, no.12, 1962, 103-109

The influence of the state of the matrix phase, the grain boundaries and of grain size on the coagulation of carbides was investigated for the steels WX15 (ShKh15) (1.13% C, 1.53% Cr, 0.44% En), 17 (U17) (1.67% C), and for a nickel steel (1.17% C. 1.5% Ni) made from Armco iron in a high frequency furnace in argon atmosphere. The specimens were water-quenched from 1100 °C. annealed at 620-630 °C and given isothermal treatments at various temperatures; in addition they were reheated and quenched from various temperatures between 950 and 1150 °C. From the number and mean radii of the carbide particles, calculations were made of the linear velocity of their growth and of the two parameters of the coagulation process: a - parameter characterising the slowing card 1/3

5/1/8/62/000/012/003/008 E071/E151

Influence of the state of the solid ... down of the coagulation in the later stages of the process; - parameter characterising the acceleration of the coagulation in the early stages. It was shown that the velocity of coagulation, the degree of dispersion of the carbide phase and the degree of slowing down of the process largely depend on the state of the a and y solid solutions. The higher the hardening temperature, the larger will be the grain size of the solid solution, the more dispersed will be the carbide phase and the higher will be the hardness of the steel throughout the course of the coagulation process and vice versa. The values of parameter p are higher for specimens hardened from lower temperatures, and those of parameter a are higher for specimens hardened from higher temperatures. Conclusions; the process of coagulation of carbides along grain boundaries and other weak places is determined not only by the diffusion coefficient, but also by changes in the solid solution. It is necessary to consider the influence of an alloying element on the state of the matrix of the solid solution (bond forces, presence of defects, grain size, etc) as well as on changes of the latter (intensity of healing Card 2/3

5/148/62/000/012/003/008 Influence of the state of the solid ... E071/E151

of lattice defects, mobility of matrix grain boundaries, etc) during the process of heating. There are 5 figures and 1 table.

ASSOCIATION: Chernovitskiy gosudarstvennyy universitet (Chernovitsy State University)

SUBMITTED: January 23, 1962

Card 3/3

PSAREV, V.I.; SALLI, I.V.

Coalescence of finely dispersed particles and determining surface tension on solid phase boundaries. Fiz. met. i metalloved 5 no.2: 268-278 '57. (MIRA 11:3)

1. Dnepropetrovskiy gosudarstvennyy universitet.
(Surface tension) (Alloys--Metallography)

AUTHORS: Psarev, V. I., and Salli, I. V.

126-2-12/35

TITLE:

Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases. (Koalestsentsiya melkodispersnykh chastits 1 opredeleniye poverkhnostnogo natyazheniya na granitse tverdykh faz).

PERIODICAL: Fizika Metallov i Metallovedeniye, 1957, Vol. 5, No. 2, pp. 268-278 (USSR)

ABSTRACT: Coalescence is a process of coarsening of the particles (grains or crystals) of the disperse phase brought about by displacement of matter from the small particles to the larger ones. This process is of great importance during tempering of hardened steels, during heat treatment of etching alloys, etc. An attempt to describe quantitatively this process has been made in earlier work of the authors of this paper and also of other authors (Refs. 4-7,9). It is known from thermodynamics that, in the case of coalescence, the change in the free energy of the system ΔF is proportional to the change of the total surface of division between the phases and $\Delta F = \delta \Delta s$, whereby the coefficient σ represents the specific free energy on the surface of division of two phases. The fundamental relations of the molecular-kinetic theory of phase

Card 1/7

126-2-12/35

Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

transformations include this important thermodynamic magnitude but so far no reliable data are available even on the order of magnitude of σ for solid phase boundaries. In this paper a calculation is presented with the aim of determining the order of magnitude of the surface tension at the boundary of the solid phases using experimental data on the kinetics of coalescence. During coalescence all the particles of the disperse phase can be sub-divided into the following two classes: large, growing particles, i.e., particles which continue to grow all the time until the process is stopped; dissolving particles, i.e., smaller particles, the smallest of which dissolve during the first stages of coalescence of which dissolve during the first stages of coalescence whilst the larger ones may even grow in the initial stage and still dissolve later. Bokshteyn, S. Z. (Ref.3) has found that during coalescence the average dimensions of the particles of both classes increase all the time. Konobeyevskiy, S. T. (Ref.1) proposed a formula for the growth of the average particle size of the disperse phase as a function of time during isothermal annealing in which Card 2/7 the rate of growth is expressed by a linear relation; for

126-2-12/35 Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

a better agreement between this formula and the experimental results, the authors of this paper introduce a decelerating term of the rate of growth thus obtaining Eq.(1), p.269. The deviation of the coagulating system from the equilibrium position can be characterized by a linear speed of growth of the average dimension of coarsening particles at a given instant of time. However, with the progress of time the particles grow bigger and the speed of growth will decrease and tend to become zero. In earlier work of one of the authors (Ref.7), a rormula was proposed for calculating the average dimension of the particles which contains a term with an exponential dependence on time. An equation is also derived for calculating the surface tension o at the boundary of solid phases, Eq.(7), p.271. If experimental data are available enabling plotting average radius of the disperse phase particles on the duration of isothermal annealing (Fig.1, p.70), it is possible on the basis of the here derived relation to evaluate the surface tension at the boundary of solid For verifying the proposed formula the authors Card 3/7 phases.

126-2-12/35 Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

used the experimental data of Bokshteyn, S.Z. (Ref.3), Schimura, S. et alii (Ref.8); the basic data extracted from the experimental results of these authors, which are necessary for calculating the surface tension, are entered in Table 1, p.272. In addition, the method of calculation is verified on extensive experimental data which permits determining the surface tension at the boundaries graphite-austenite, carbide-austenite and carbide-ferrite. The process of coalescence of graphite particles in iron was investigated on two groups of specimens, the first consisting of iron with a nearly sutectoidal content of admixtures, i.e., 1% Si, 0.3% Mn, 0.15% P and 0.1% S, the second was prepared from pure synthetic iron with the same C and Si contents as for the first group but having a minimum content of the The experimental technique was other admixtures. described in an earlier paper (Ref.6). Tables 2 and 3 give the data on the dependence of the numbers of grains per unit of volume on the time and temperature of annealing of various specimens of malleable iron. Card 4/7results are entered into tables and they are discussed

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126-2-12/35 Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

The data contained in the paper justify in some detail. the assumption that the presented method for determining the order of magnitude of surface tension at the boundary of the solid phases is fully satisfactory and this view is supported by a number of facts. The same value was obtained for the surface tension of cementite at the boundary with ferrite by determining it from three different sources. For the steel Y-8 at 650°C σ = 15 dyn/cm, for below-eutectoidal steel at 630°C Bokshteyn (Ref.3) obtained σ = 13.2 dyn/cm, whilst for steels with various carbon contents at 720°C Schimura et alii (Ref.8) quoted the value $\sigma = 7.5$ dyn/cm. If the temperature is taken into consideration these values can be considered as being fully in agreement. According to Schimura et alii (Ref.8) the surface tension of cementite at the boundary with ferrite remains constant when changing the carbon concentration. For all the investigated alloys the magnitude of the surface tension decreases with increasing temperature, whereby the decrease is almost inversely proportional with the solubility of the

Card 5/7 disperse phase in the mother phase (in the temperature

CIA-RDP86-00513R001343510003-1"

APPROVED FOR RELEASE: 06/15/2000

126-2-12/35

Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

range where no phase transformations occur). The surface tension of cementite at the boundary with austenite is almost five times lower than the surface tension of graphite at the boundary with austenite. This fact is confirmed by phenomena which accompany solidification of Investigation of alloy steels showed that for a relatively small content of the alloying element, of the order of 2%, the order of magnitude of the surface tension does not change. However, a definite tendency towards increasing can be observed in the surface tension of carbide bounding on ferrite on introducing carbide forming elements (Cr,Ti) and a tendency can be observed towards lowering of the surface tension on introducing non-carbide forming elements, cobalt for instance. The obtained values of surface tension appear reasonable also size of the critical as regards the determined germination nucleus during phase transformation, for instance for steel containing 1.2% C and super-cooled to 50° C below the A₁ point, the critical size of the cementite germ is about 10^{-7} cm, which is a fully

Card 6/7 realistic value for such undercooling.

126-2-12/35

Coalescence of finely dispersed particles and determination of the surface tension at the boundary of solid phases.

There are 4 figures, 10 tables and 10 references, 8 of which are Slavic.

SUBMITTED: October 11, 1955 (Initially), April 10, 1957 (after revision.)

ASSOCIATION: Dnepropetrovsk State University.
(Dnepropetrovskiy Gosudarstvennyy Universitet).

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Card 7/7

_PSAREV, V.I.

Strengthening of crystals of chemical compounds in alloys of the systems $Z_n - S_b$, $Cd - S_b$, $In - S_b$. Kristallografiia 5 no.3: 479-481 My-Je '60. (MIRA 13:8)

1. Chernovitskiy gosudarstvennyy universitet.
(Zinc antimonide)
(Cadmium antimonide)
(Indium antimonide)

PSAREV, V. I.

"On Calculation of Kinetics Curves of Heating and Cooling.:

Report submitted for the Conference on Heat and Mass Transfer, Minsk, BSSR, June 1961.

ISAREV, V.L

s/070/60/005/03/008/008 E132/E360

AUTHOR: ___ Psarev, V.I.

On Increasing the Sizes of Crystals of the Chemical Compounds in the Alloy Systems Zn-Sb, Cd-Sb and In-Sb

Kristallografiya, 1960, Vol. 5, No. PERIODICAL:

TEXT: Investigations have been made of the possibilities of increasing the sizes of the particles of Zn,Sb, 2 CdSb, Vand InSb)

occurring in alloys of the corresponding elements. This increase occurred in the process of coalescence under the action on the alloy of mechanical forces (centrifugal). Specimens were made from the elements by heating under vacuum in pyrex tubes, first at 40 above the liquidus temperature then at it for 6-7 hours. The other alloys were made similarly. The rate of enlargement of the particles was greater the higher the temperature. A graph is given of the rate of growth of Zn3Sb2 crystals from a melt of 93%

Zn with 7% Sb at 440 °C. After some 60 hours the growth had almost finished. Where there is a difference in density between the solid and the liquid phases centrifugation may assist coalescence.

Card 1/2

TITLE:

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L 11,611,-66 EWT(m)/T/EWP(t)/EWP(b) IJP(c) JD ACC NR: AT6002264 SOURCE CODE: UR/2564/65/006/000/0288/0295		,
AUTHOR: Kostur, N. L.; Psarev, V. I.		
ORG: none		
TITLE: Alloying crystals of some intermetallic compounds by means of alloys [Paper presented at the Third Conference on Crystal Growing held in Moscow from 18 to 25 November, 1963]		
SOURCE: AN SSSR. Institut kristallografii. Rost kristallov, v. 6, 1965, 288-295		
TOPIC TAGS: cadmium compound, antimonide, indium compound, electric conductivity, Hall constant, thermoelectromotive force		
ABSTRACT: CdSb and InSb compounds were alloyed with alloys of the corresponding systems. The alloying elements chosen for CdSb were Ag? Au, 2b, and Bi, and for InSb,		
Bi and Hg. Crystals of the alloyed compounds were then used for growing large single crystals by zone recrystallization. The degree of alloying was determined in these crystals measuring the temperature dependence of the electrical conductivity σ, Hall		
constant R, and thermo-emf α . The results shows that the method of alloying intermetallic		
Card 1/2		
	1. 255	er Same

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ompounds with alloys of systemy valuable because data on the sign and are lacking for intermosas: 5 figures.	solubility o	f elements.	which are	e necess	ary in di	rect amoy-	
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SUB CODE: 20 / SUBM DATE	: none / OR	IG REF: U	,,				
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PSAMEV, V.I.

Mechanism of the coalescence of finely divided phases in metal alloys. Fiz. met. i metalloved. 18 no.3:471-473 S 164. (MIRA 17:11)

1. Chernovitskiy gosudarstvennyy universitet.

45222-65 CCESSION NR:	ENT(m)/ENP(z)/ENA(c) AP5010985)/t/ewp(b)/ema(d)/i U	MP(t) IJP(c) R/0148/65/000/004/0	142/0149
UTHOR: Psare	34, V. I.			, 8 , 17
ITLE: Coagul	lation of carbide parti	cles in <u>high-mangan</u>	se steel 4	
OURCE: IVUZ.	. Chernaya metallurgiy	/a, no. 4, 1965, 142	149	B
OPIC TAGS:	carbides, steel alloys,	coagulation		
n 12-13%-Mn s f steel (see	re results of the investeel are presented. Stable 1 of the Enclosu	amples were taken f ire) and subjected t	rom forged rods of 5 20-25% precipitat	two grades ion
750-850°C) ar ispersed part	lowed by tempering at 5 nd quenching in water, ticles and other coagul amples. It is shown the	the number of carbi Lation parameters we	le particle, radii re measured in defo	of the rmed and
n the kinetic arbide partic f manganese a	imples. It is shown the confidence of carbide coagulation is atoms the concentration olution. The character	on in high-manganes not accompanied by n of which remains h	a steel. The growt my considerable mi ign in the surround	h of gration ing y-

45222-65 CESSION NR: AP5010985			
tributed to a lower number of a steels than on nonde formulas.	ring the growth of the carbone rapid growth of the par	ide particles while the ticles after deformation is undaries on the surface of has: 2 figures, 1 table,	
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45222-65 CESSION NR:	AP5010985					EKCLO	SUNE: 01	o l	
			Table						
	Compositio	m of ty	pes of	steel i	nvestig	ated			
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1	1.30 1.34	0.69 0.67	12.62	0.090	0.01				
医三哥									

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EWP(q)/EWT(m)/BDS AFFTC JD 17083-

ACCESSION NR: AP3004594

S/0126/63/016/001/0071/0079

AUTHORS: Kostur, N. L.; Psarev, V. I.

TITLE: A procedure for growing CdSb crystals from alloys of Cd-Sb system

SOURCE: Fizika metallov i metallovedeniye, v. 16, no. 1, 1963, 71-79

TOPIC TAGS: CdSb, crystal growing, Cd-Sb alloy

ABSTRACT: A new method for growing large CdSb monocrystals is offered. The procedure consisted of three steps. 1) Binary alloys Cd-Ag (Ag to 2%) and Cd-Hg (Hg to 20 by weight) were prepared. They had a uniphasal structure of solid solution on the Cd base. 2) Various amounts of antimony (from 10 to 40% weight) were added to these binary solutions. The solutions were melted and held at 6200 for 5 hours while being stirred to insure a good distribution of the alloy components. After that, the melt was cooled to 5000, aged at that temperature for 20 hours, and cooled again in the oven. The resultant ternary alloys Cd-Ag-Sb and Cd-Hg-Sb had a two-phase structure consisting of the CdSb crystals and a cadmium eutectic. 3) The CdSb crystals were separated centrifugally from the melt. They contained from 20 to 40% weight of Sb. These crystals were used

Card 1/2

ACCESSION NR: AP3004594		watala by the method	
as the initial material for t of zonal recrystallization.	he growing of large CdSb monorry Orig. art. has: 2 tables and 5	figures.	
ASSOCIATION: Chernovitskiy g University)	osudarstvenny*y universitet (Ch	ernovtsy State .	
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PSAREV, V.I.; Prinimali uchastiye: MAKOVIICHUK, Iu.I., student; REVA, B.F., stadent

Coagulation of the carbide phase during the tempering of malybdenum steel. Fiz. met. i metalloved. 15 no.2:234-238 F 163. (MIRA 16:4)

1. Chernovitskiy gosudarstvennyy universitet.
(Molybdenum steel---Heat treatment)
(Phase rule and equilibrium)

PSAREY, W.I. [Foar for, V.I.]; KOSTUR, N.L. [Kostur, M.L.]; (BSTRA, A.V.

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Phase separation in alloys of the systems Cd - Sb and In - 3b by centrifuging the melt. Ukr. fiz. zhur. 8 no.6:694-699 Je '63. (MIRA 16:7)

 Chernovitskiy (osudarstvennyy universitet. (Cadminm-indium-antimony alloys) (Crystils--Growth)

PSAREV, V. I.; KOROBIY, O. I.

Effect of the state of a solid solution on the kinetics of the coagulation process and dispersion of the carbide phase in steels. Izv. vys. ucheb. zav.; chern. met. 5 no.12:103-109 62. (MIRA 16:1)

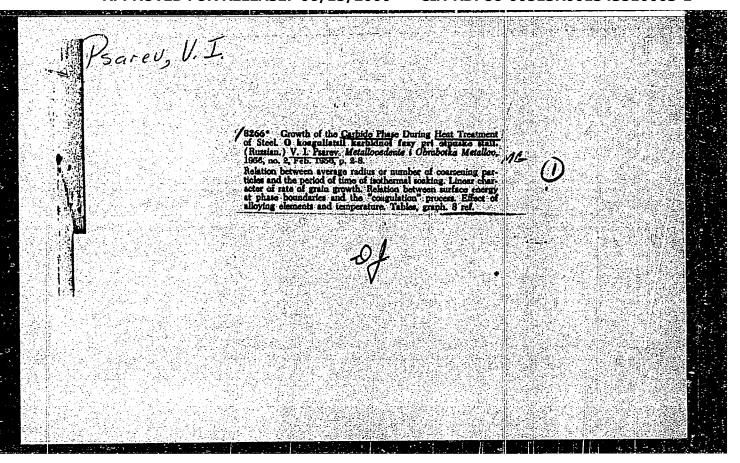
1. Chernovitskiy gosudarstvennyy universitet.

(Steel-Metallography)
(Phase rule and equilibrium)

PSAREV, V.I.; DOBRYDEN', K.A.; KAZYUK, F.I.

Investigating the coagulation process of finely divided phases in Zn-Sb alloys. Fiz. met. i metalloved. 14 no.1:130-133 J1 '62. (MIRA 15:7)

1. Chernovitskiy gosudarstvennyy universitet.
(Zinc-antimony alloys-Metallography)



18(3)

SOV/148-59-2-11/24

AUTHOR:

Psarev, V.I., Candidate of Technical Sciences

TITLE:

On Regularities of the Coagulation Process of Carbide Particles in Steel Above the Al Foint (O zakonomernostyakh protsessa koagulyatsii karbidnykh chastits v stalį vyshe

tochki Al)

PERIODICAL:

Izvestiya vysshikh uchebnykh zavedeniy, Chernaya metallurgiya,

1959, Nr 2, pp 93-99 (USSR)

ABSTRACT:

An attempt is made to complete the existing data on coagulation processes in steel and to determine the interdependence between the coagulation parameters and the carbon content or the coagulation temperature. Information is given on general characteristics of coagulation kinetics of simple carbide systems in carbon and alloy steels. The correlation between the coagulation parameters and the equilibrium curve of iron carbon alloys is determined. The connection between the carbide phase dispersion and the shifting of the carbideaustenite equilibrium line due to the effect of an alloying

Card 3

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On Regularities of the Coagulation Process of Carbide Particles in Steel Above the Al Point

> element is established. It was stated that the linear coagulation rate in carbon steels depended on the time and temperature of isothermic heating. A higher carbon content in the steel or higher temperatures caused the reduced dispersion of the carbide phase. A formula is given to compute the dependence of the end radius of carbide particles on the coagulation temperature. In alloy steels an increased content of the alloy component created special carbide phases. Their coagulation depended on various factors, such as surface tension in the carbide-austenite border, diffusion of carbon and of alloying components etc. The experimental part of the work was carried out in cooperation with Ye. V. Akhtemiychuk and S. I. Ivanenko. There are 2 tables, 1 graph, 3 sets of microphotos and 6 Soviet

> references.

Card 2/3

APPROVED FOR RELEASE: 06/15/2000 CIA-RDP86-00513R001343510003-1"

SOV/148-59-2-11/24

On Regularities of the Coagulation Process of Carbide Particles in Steel Above the A₁ Point

ASSOCIATION:

Chernovitskiy gorudarstvennyy universitet (Chernovtsy State

University)

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50V/126-7-4-9/26

AUTHOR: Psarev, V.I.

TITLE: Investigation of the Process of Coalescence of Carbides

in Carbon Steels

PERIODICAL: Fizika metallov i metallovedeniye, 1959, Vol 7, Nr 4,

pp 551-558 (USSR)

ABSTRACT: Although the mechanical properties of steels depend

to a large extent on the size of the carbide particles, little has been done in the way of establishing the laws governing the rate of coalescence and the final size attained by these particles during changes taking

place in steels either in the course of thermal treatment, or in service at elevated temperatures. The problem of the formulation of general laws of this nature is the subject of the present paper. A two-phase system in which one of the phases is present in the form of small, dispersed particles will tend to attain the state of equilibrium by the process of growth and coalescence of the finely dispersed phase. In the previously published works (Ref 1 and 2), the author of the present paper derived an expression describing the

Card 1/11 relationship between the mean radius of the coalescing

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

particles and the time of isothermal heating

$$r = r_0 + \frac{v_0}{\alpha} \left[1 - \exp(-\alpha t) \right]$$
 (1)

where: r - mean particle size at time t; r_o - initial mean particle size at t - 0; v_o - initial rate of growth of the particles; α - coalescence constant for isothermal heating. It was shown experimentally (Ref 1) that this equation describes the relationship under consideration only starting from a certain time t^1 , when the condition $\ln v_o/v = \alpha t$ (where α is the angle coefficient) is fulfilled (compare Fig 1). It was found also that for various steels tempered at 500 to 600°C, t^1 is equal 3 to 4 hours; at 700°C, t^1 is equal 2 to 1.5 hours and it approaches zero at temperatures above A₁. A more accurate expression for the relationship described by Eq (1) can be obtained by solving this problem in the second approximation; if the function

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

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of the variation of the rate of growth of the particle is expanded into series and if the second term of this scries is taken into account, the following expression is obtained:

$$-\frac{dv}{dt} = \alpha v + \beta v^2 = f(v)$$
 (2)

The coefficients α and β in the expanded function $\ell(\mathbf{v})$ are constant at T= const but are affected by the variation of temperature and other factors; these coefficients are termed by the present author coalescence parameters. From Eq (2), and bearing in mind that $\mathbf{v}=$ dr/dt, an expression can be derived describing the relationship between the mean particle size (r) and the duration (t) of isothermal heating

$$r = r_0 + \frac{1}{\beta} \ln \left\{ 1 + \frac{\beta v_0}{\alpha} \left[1 - \exp(-\alpha t) \right] \right\}$$
 (3)

Card 3/11 This equation satisfies the limiting conditions

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

$$t = \infty r_m = r_0 + \frac{1}{\beta} \ln \left(1 + \frac{\beta v_n}{\alpha}\right)$$
 (4)

where: r_m - the final particle size. From the latter expression the function v = f(r) can be found in the form of

$$v = \frac{\alpha}{\beta} \left[\exp \beta (r_m - r) - 1 \right]$$
 (5)

From the analysis of the possible values of the roots of Eq (5) it follows that it can describe the coalescence process when β has positive values; if β approaches zero, Eq (1) is obtained. Regardless of the order of magnitude of β , the second term in Eq (2) can be left out starting from time t¹ (see Fig 1), when the condition $\alpha v \gg \beta v^2$ is fulfilled. Starting from this moment, the time dependence of the particle size can be adequately described by Eq (1). The values of β for the process of growth of the carbide particles in steels was calculated for temperatures both above (Ref 1, 3, 4, 5)

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

and below (Ref 2,4,5,6,7) the critical point A1; In the former case, the values of $\boldsymbol{\beta}$ are so small that the value of αv in Eq (2) is much larger (by one or two orders of magnitude) than that of βv^2 already in the initial stages of the process of coalescence; consequently, the time dependence of the carbide particle size at temperatures above Al is adequately described by Eq (1). For changes occurring at temperatures below A_1 , when the values of β are relatively large, Eq (5) has to be used. Fig 2 shows the relationship d_{cp} (mean particle_diameter, 105cm) versus vcp (mean rate of growth, 105cm/h) for 0.4% C (curve 1) and 0.7% C (curve 2) steels heated isothermally at 630°C. Both curves were plotted from Eq (5), taking $\beta = 1.75 \times 10^5 \text{cm}^{-1} \text{ (curve 1) and } = 2 \times 10^5 \text{cm}^{-1} \text{(curve 2)}$ and experimental values for other magnitudes appearing in this equation. Good agreement between the theoretical curves and experimental points (shown in Fig 2 by circles and crosses) indicates that by taking into account the second term of the expanded function f(v), it is possible to describe with sufficient accuracy the process of

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

coalescence of carbides in the early stages of their growth i.e. before time t^1 . The problem of determination of the coalescence parameters is discussed in the second part of the present paper. It has been shown experimentally (Ref 1) that the final particle size of carbides in steels increases with the heat treatment temperature and that at the same temperatures, it can attain different values in different steels from which it was concluded that r_m depends on the concentration of the constituent atoms at the phase boundary. The relationship $r_m = f(c, T)$, where c - concentration; T - temperature, can be obtained from the known Thompson (Kelvin) formula by applying it to the limiting (initial and final) values of r_m , and then excluding the equilibrium concentration c_{∞} . The following expression is obtained:

$$c_{\min} = c_0 \exp \left[-\frac{a}{T} \left(\frac{r}{r_0'} - \frac{1}{r_m} \right) \right]$$
 (6)

Card 6/11 where a = $2\sigma Mv/R$ (σ - specific surface energy at the

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

phase boundary; M - molecular weight of the dispersed phase; R - gas constant). From Eq (6) the expression for the final particle size can be obtained

$$\mathbf{r}_{m} = \frac{\mathbf{r}_{0}^{\prime}}{1 - \mathbf{C}_{1}^{\prime} \mathbf{T}} \tag{7}$$

where $C^{\dagger} = r_0/a \ln c_0/c_{\min}$. At present, it is not possible to determine C' which, however, obviously is a function of temperature. Fig 3 shows the temperature dependence of r_m plotted for cases when C^* increases with the temperature (curve 1) when it remains constant (curve 2) and when its value decreases with rising temperature (curve 3). Calculations based on experimental data (Ref 1 to 4) show that for practical purposes C' changes very little with temperature and that the temperature at which 1/C' = T' corresponds almost exactly to the temperature of the phase transformation. (These calculations were based on experimental data on coalescence of carbides in carbon and alloy steels (Ref 1 to 3) and graphite particles in malleable cast Card 7/11 iron (Ref 4).) The values of r_m calculated from the

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Investigation of the Frocess of Coalescence of Carbides in Carbon Steels

formula $r_m = r_0 + v_0/\alpha$ (Ref 1) and values of the same magnitude calculated from Eq (7) are given in Table 1, for steel with 1.7% C, steel Ul2A and malleable cast iron. (The values of C' and r_o for each alloy are given at the bottom of the table). The temperature dependence of r_m in alloy steels is illustrated in Fig 4. (Curve 1 chromium steel, C' = 6.68×10^{-4} °C⁻¹, r₀ = 3.75×10^{-5} cm; Curve 2 - manganese steel, C' = 7.5×10^{-4} °C⁻¹, $r_0 = 5.1 \times 10^{-5}$ cm). Taking into account the fact that with a sufficient degree of accuracy $C^2 = 1/T^2$ (where T^2 is the temperature of the phase transformation in ${}^{\circ}K$) the expression describing the temperature dependence of the coalescence parameter α can be derived. (It is pointed out at this stage that the above consideration applies to the case when, as a result of complete dissolution of the finely dispersed phase, a two-phase system is changed into a single phase system; consequently, it does not apply to coalescence of carbides in steels at temperatures below A1). The relationship between α and r_m is given by $\bar{\alpha} = -D/\beta$ de/dx,

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where D - diffusion coefficient, ρ - density of the coalescing phase and dc/dx - concentration gradient. After substituting the values for r_m , v and D, the temperature dependence of α is obtained in the form of

$$\alpha = \frac{k}{T^{\frac{1}{2}}e^{-\frac{Q}{RT}}(\frac{T^{\frac{1}{2}}}{T} - 1)$$
 (8)

where $k=-D_0/p$ de/dx; T^{\pm} - the temperature at which carbides are completely dissolved. Analysis of this expression shows that α increases with rising temperature (see Fig 5), reaches a maximum at $T_1=T^{\pm}/(1-RT^{\pm}/Q)$, and then falls to zero at $T=T^{\pm}$. The validity of Eq (8) can be checked with the aid of experimental data if the values of Q and k are known. The results of such calculations for a carbon steel, a chromium steel and a manganese steel are plotted in Fig 6, curves 1,2 and 3 respectively. (The values of Q cal/g-atom, k cm/sec and T^{\pm} h, are given in the caption.) Similar calculations were carried out for a wide range of carbon

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steels and in every case there was good agreement between the experimental and calculated values of a corresponding to the ascending portion of the α = f(T) curve; after passing the maximum, the calculated values of α are higher than those obtained experimentally. Table 2 gives experimental (column 2) and calculated (column 3) values of α at various and calculated (column 3) values of α at various temperatures. The problem of deriving an expression for the coalescence parameter β is more complex, since in this case it is necessary to know the form of the function $\nu = f(r)$. Then, by differentiating Eq (5) in respect to r, the following formula is obtained:

 $\beta = \frac{\ln\left(-\frac{1}{\alpha}\frac{dv}{dr}\right)}{r_{m}-r} \tag{9}$

The value of the magnitude dv/dr is best obtained graphically from the relationship between v and r; a curve of this type, plotted from experimental data is shown in Fig 2. Incidentally, it follows from that

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Investigation of the Process of Coalescence of Carbides in Carbon Steels

> curve that at the point where $r = r_m$, parameter $\alpha = -(dv/dr)_r = r_{m^\circ}$ It is stated in the conclusion that although the analytically derived expressions describing the temperature dependence of the coalescence parameters are in fairly good agreement with the experimental data on the coalescence of the carbide particles in steels at temperatures above A1, certain assumptions made in deriving these expressions should be verified by further experimental work. There are 6 figures, 2 tables and 8 references, 7 of which are Soviet and 1 German.

ASSOCIATION: Dnepropetrovskiy gosudarstvennyy universitet

(Dnepropetrovsk State University)

SUBMITTED:

April 14, 1958

Card 11/11

PSAREV, V.I.

Solubility of carbon in ferrites. Fiz. met. metalloved.ll no.6:944-945 Je '61. (MIRA 14:6)

1. Chernovitskiy gosudarstvennyy universitet. (Iron—Metallography)

CIA-RDP86-00513R001343510003-1 'APPROVED FOR RELEASE: 06/15/2000

PSAREV, V.J.

137-58-3-5842

Translation from: Referativnyy zhurnal, Metallurgiya, 1958, Nr 3, p 196 (USSR)

AUTHORS: Psarev, V. I., Salli, I. V.

Kinetics of Coalescence and the Determination of Surface Tensions TITLE:

on the Boundaries of Solid Phases (Kinetika koalestsentsii i

opredeleniye poverkhnostnogo natyazheniya na granitse tverdykh

faz)

Nauchn. zap. Dnepropetr. un-t, 1956, Vol 45, pp 61-67

The authors present data on the kinetics of coalescence of PERIODICAL: globular cementite over a wide range of temperatures and for ABSTRACT:

various grades of pure and alloyed steels. Investigations are carried out to obtain experimental data permitting the calculation

of the surface tension of on the boundaries of solid phases according to the method proposed by Salli (see RzhMet, 1958, Nr

3, abstract 5841). Values of on the cementite-ferrite

boundary, calculated from various sources (including allowances for the temperature t), are fairly close: steel U-8 at t=650°C,

O=15 dynes/cm; subeutectic steel at t=630°, O=13.2 dynes/cm.

The value of of is not affected by the C content, but is decreased

at higher temperatures. Although the order of magnitude of

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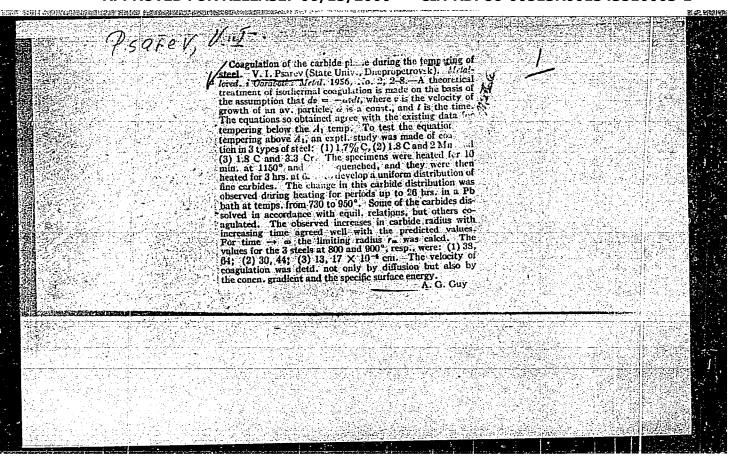
137-58-3-5842

Kinetics of Coalescence and the Determination of Surface Tensions (cont.)

is not affected by the introduction of alloying materials into the steel, a trend toward higher values of $\mathcal O$ on the carbide-ferrite boundaries is observed when carbide-forming additives (Cr. Ti) are introduced. The introduction of additives which do not form carbides (Co) produces a reduction in the value of $\mathcal O$.

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Card 2/2



PSAREV, V. I.

FSAREV, V. I. "The Kinetics of Coalescence of the Carbides of Iron-carbide Alloys." Acad Sci Ukrainian SSR. Inst of Ferrous Metallurgy. Dnepropetrovsk, 1956. (Dissertation for the Degree of Candidate in Technical Science)

So: Knizhnaya Letopis', No. 18, 1956,

PSAREV, V.I.; YURCHUK, I.A.

了对外的现在分词 经企业分别,但他正规决划的现在分词的成功的的现在分词的对象的的是是不是是的一个人的对象的 医结合性 化双氯化乙酰胺 医克克克氏征 医拉克克氏征 医拉克克氏征 医多种

Coalescence of carbide particles in the process of continuous heating. Izv. vys. ucheb. zav.; chern. met. 4 no.10:82-87 '61. (MIRA 14:11)

1. Chernovitskiy gosudarstvennyy universitet.
(Steel--Heat treatment)

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AUTHOR:

Psarev, V. I.

TITLE:

On the coagulation of carbide particles in carbon steels below

the A₁ point

PERIODICAL:

Izvestiya vysshikh uchebnykh zavedeniy. Chernaya metallurgiya,

no. 12, 1960, 91 - 96

TEXT: The purpose of the described experiments was to obtain more data on the coagulation of carbide particles below the critical A₁ point in carbon steel, for the data published hitherto vary and no systematical studies of the process had been undertaken. The test method had been described in two previous articles by the author (Ref. 6: V. I. Psarev, "Metallovedeniye i obrabotka metallov", 1956, no. 2; and Ref. 7: V. I. Psarev, "Fizika metallov i metallovedeniye", v. 7, no. 4, 1959). The investigated commercial steel grades y10 (U10), y12 (U12) and y17 (U17) were smelted from armod iron in an electric high-frequency furnace in an argon medium. Cylindrical specimens 12 - 15 mm in diameter and 15 - 20 mm in length were heated to 1,000°C ("U17" to 1,100°C), soaked in this heat for 5 - 8 min,

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On the coagulation of cartide particles in...

quenched in water, and subsequently tempered at 620 - 630°C for 3 hrs, then additionally heated for 1 hr to different temperatures in which the coagulation had to be determined. The decarbonized layer was removed from the specimen after each isothermic heating (in a lead bath), the number of carbible particles was seen total at x1440 magnification with the aid of a grid placed in a microscope coular and the Brinell hardness was measured. The final radius of the particles was determined using the formula (from Ref. 6, 7)

$$\mathbf{r}_{tt} = \mathbf{r}_{0} + \frac{\mathbf{v}_{0}}{\mathbf{c}^{2}} \tag{1}$$

where r_0 is the initial particle radius; ν_0 - the initial linear particle growth rate; α - the coagulation factor characterizing the damping of the coagulation process and equal

$$\alpha = \frac{1}{t} \ln \frac{v_0}{v} . \tag{2}$$

The mean radius r of the carbide particles was determined by the formula (also from Ref. 6, 7);

$$r_{r} = r \frac{v_{0}}{\alpha} \left[1 - \exp(-\alpha t) \right], \qquad (3)$$

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On the coagulation of carbide particles in ...

where t is the isothermic soaking time, and $v = \frac{dr}{dt}$ the experimentally determined rate of coagulation. The dependence of α on the tempering temperature below the A1 point was similar to the dependence above A1, and the mean radius of cementite particles increased with an increase in the carbon content in steel (Fig. 1). The quantity of cementite particles (during the same time of isothermic soaking) increased at the beginning and reached a certain maximum at a certain carbon content in steel, then decreased (Table 2). The growth rate of the constant particle size increased with the carbon content with linear regularity (Fig. 1). The final particle radius increased with the coagulation temperature. The relation curve is hyperbolic, but a quantitative regularity could only be stated for relatively pure binary stael (not for commercial steel). A linear relation between the steel hardness and the total carbide phase surface area was found in the interval 100 - 150 hrs of isothermic heating; during longer heating (150 - 250 hrs) it changed (Fig. 3). The results led to the conclusion that the coagulation of carbides must cease faster in high-carbon steel than in low-carbon, and same result can be attained by a higher coagulation temperature, but the linear coagulation rate grows rapidly with a rise in C-content and tempera-

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On the coagulation of carbide particles in. ...

bura, and the result is a larger final size of particle. The growth of particles cannot be explained by easier coagulation at higher C-content, for the curve maximum shifted toward a low C-content after 1-hour isothermic heating. A further analysis is necessary. There is no linear relation between the steel hardness and the aggregate carbide phase surface, for the oversaturation of the alpha phase with carbon varies drastically with time. The relation may be approximately linear only in later coagulation stages. Long soaking (250 - 300 hrs), even in a protective medium, leads to partial decarbonization. The approximate relation between the steel hardness H and total earlies surface S may be expressed as

 $H = H_m + k(S - S_m)$

where S_m is the minimum total carbide surface $S_m=4\pi r_m^2~N_m$; N_m - the minimum number of carbide particles in a cm³ after the end coagulation process; k - the proportionality factor depending on the variations of the alpha state, the rate of variation of the total carbide phase surface, C content in steel, and more factors. Ya. N. Balagutrak and M. D. Karliychuk participated in the work. There are 3 figures and 7 references: 6 Soviet-bloc and

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CIA-RDP86-00513R001343510003-1

On the coagulation of carbide particles in ...

1 non-Soviet-bloc.

ASSOCIATION: Chernovitskiy gosudarstvennyy universitet (Chernovitsy State

University)

SUBMITTED: April 4, 1960

Table 2.

Scaking Bpems, Budepm-	Carbon content, Содержание углерода. % in %						
Tine ku, aacu nrs.	0.4 [1]	0,7 [1]	0.95	1,15	1,67		
1 25	142.7 69.2	192.3 67.6	100,4 52,4	86,35 57,0	66.0 42,8		

Particles numbers in $(N \cdot 10)^{-10}$ cm⁻³].

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PSAREV.V.I. Cealescence of the carbide phase in steel tempering. Metalloved. i obr. met. no.2:2-8 F 156. (MIRA 9:7) 1.Dnepropetrovskiy gesudarstvennyy universitet. (Steel--Metallography) (Tempering)

L 47341-66 EWI(1)/EWT(m)/T/EWP(t)/ETI IJP(c) GG/JG/JD/WW SOURCE CODE: UR/0058/66/000/004/A075/A075	
L 47341-66 EWT(1)/EWT(m)/T/EWP(1)/ETT TSP(1) GO, CO. UR/0058/66/000/004/A075/A075 ACC NR: AR6025 55 SOURCE CODE: UR/0058/66/000/004/A075/A075	
	a e
AUTHOR: Kostur, N. L.; Psarev, V. I. 17 17	
TITLE: Features of crystallization of the Cust compound	
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REF SOURCE: Sb. Simpozium. Proteessy sincessy si	
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chiometry, single crystal growing, of a melt. which	
ABSTRACT: An immestigation was made of the influence of supercooling of and the content depends on the rate of its cooling, the degree of prior superheating, and the content depends on the rate of its cooling, the degree of prior superheating, and the content depends on the rate of its cooling. The is	
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was superheated by 10 - 50 . you will all of stoichiometric composition	
have a single-rase solution. On the basis of the obtained results, the most openation was superheated by 10 - 30°. On the basis of the obtained results, the most openation regimes are proposed for growing CdSb single crystals of stoichiometric composition and of crystalr doped with various impurities. [Translation of abstract]	1
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CIA-RDP86-00513R001343510003-1 "APPROVED FOR RELEASE: 06/15/2000

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EWT(m)/T/EWP(t)/ETI IJP(c) L 04302-67 UR/0363/66/002/008/1383/1389 SOURCE CODE: ACC NR: AP6029813 (A)38 31 AUTHOR: Kostur, N. L.; Psarev, V. I. ORG: Chernovtsy State University (Chernovitskiy gosudarstvennyy universitet) TITLE: Conditions for crystallization of the intermediate phases in the Cd-Sb system SOURCE: AN SSSR. Izvestiya. Neorganicheskiye materialy, v. 2, no. 8, 1966, 1383-1389 TOPIC TAGS: crystallization, phase diagram, solution property, phase composition, phase analysis, cadmium, antimony ABSTRACT: The Cd-Sb system was studied in the 41.93-52.0 wt % Sb range by a combination of metallographic-, thermographic-, and x-ray techniques in order to define the conditions of crystallization of the stable and metastable phases of the system. Samples were prepared by fusing mixtures of pure components in sealed ampoules for 5 hr at 630-640°C and 10⁻³-10⁻⁴ mm Hg. For a given rate of cooling (100-300 deg/hr) the formation of the intermediate phases was found to depend upon the maximum temperature of the melt. The variation in the formation of these phases depends apparently upon the presence of undissociated CdSb molecules which act as seeds during the crystallization process. A compound with a composition close to that of CdSb was found to crystallize out at 420-450°C. The CdSb, Cd4Sb3, and Cd3Sb2 crystallize only from melts heated up to 630°C. The metastable phases can be stabilized by means of rapid cool-UDC: 546.48+546.86 Card 1/2

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APPROVED FOR RELEASE: 06/15/2000

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Council : loss Carscony : Form Animals. General Problems. Q-1

A38. JOUR. : RZBiol., Rc. 4, 1959, No. 16595

AUTHOR

: Pshenichnyy, P. D.

INST. TITLE

: Breeds and Productive Types of Ferm Animals.

ORIG. PUB.: Zhivotnovedstvo, 1958, No 7, 55-56

ABSTRACT : No abstract.

CARD:

1/1

PSARFY, V.I.

Category: USSR/Solid State Physics - Phase Transformation in Solid Bedies E-5

Abs Jour : Ref Zhur - Fizika, No 2, 1957 No 3855

Author Fsarev, V I

Inst Deepropetrovsk Uciversity, USSR

Title Goagulation of Carbide Phase in the Tempering of Steel.

Orig Fub : Metallivedeniye i obrabotka metallov, 1956, No 2, 2-8

Abstract : Assuming that the quartity that characterizes the deviation of the

system from the equilibrium state is the linear velocity of the growth of the thickening particles, a relationship was derived between the average radius of the particle and the time of the isothermal heating of the steel. Experimental data are given (over a wide temperature range) on the investigation of the coagulation of carbide particles above the critical point A_1 in three types of steel: carbon, mangamese, and chrome. Analysis of the data made it possible to obtain the temperature dependence of the final dimension of the thickening

carbide particles.

Card : 1/1

L 15946-66 EWT(m)/T/EWP(t)/EWP(b) IJP(c) JD

ACC NR: AT6002257 (N) SOURCE CODE UR/2564/65/006/000/0247/0254 33

AUTHOR: Psarev, V.I.; Dobryden', K.A.

网络埃勒克莱亚亚 医科内里 医外性外视器 医胆囊 医生物 医生殖性 医红斑 医皮肤 医皮肤 医皮肤 医皮肤 计算术 经分别 医阿里克氏 医甲氏试验 经保险

3+1

ORG: none

TITLE: Principles of crystallization and stabilization of the structure of compounds CdSb and Cd₃Sb₂ in alloys of the Cd-Sb system. [Paper presented at the Third Conference on Cystal Growing held in Moscow from 18 to 25 November, 1963]

SOURCE: AN SSSR. Institut krists ografii. Rost kristallov, v. 6, 1965, 247-254

TOPIC TAGS: cadmium compound, antimony compound, cadmium alloy, antimony alloy, crystallization

ABSTRACT: The study was devoted to the determination of the conditions under which the compounds CdSb, Cd₃Sb₂, and Cd₄Sb₃ are formed during crystallization of Cd-Sb alloys. The structural state and composition of the samples was studied by thermographic, electrolytic, metallographic, and x-ray analyses. It was found that during crystallization of melts containing from 41.93 to 52.00%, CdSb crystals supersaturated with cadmium precipitate first, as a result of which the melt becomes rich in antimony, and suitable supercooling creates conditions for the nucleation of Cd₃Sb₂ crystals. When the melt is Card 1/2

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rapidly cooled, Cd and the structure of phase Cd ₃ Sb ₂ . Th on Cd ₃ Sb. Orig. 2	of the solidifie e compound C	d alloy consis daSba is appa	sts primarily or rently a subtr	of particles of active solid	i the meta	stable	
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PSAREVA, T.V.; RIKHTER, G.D.; CHIZHOV, C.P.

Second scientific conference of Kazakhstan geographers. Izv.
AN SSSR. Ser. geog. no. 4:166-167 Jl-Ag '61. (MIRA 14:7)
(Kazakhstan--Geography--Congresses)

GROSVAL'D, M.G.; PSANEVA, T.V.; AVSYUK, G.A., otv. red.; OGANOVSKIY, P.N., red.

[Franz Josef Land]Zemlia Frantsa-Iosifa. Moskva. (Its Materialy gliatsiologicheskikh issledovanii). [Ice structure]Struktura 14. (MIRA 16:2) 1962. 99 p.

1. Akademiya nauk SSSR. Institut geografii. (Franz Josef Land--Ice)

KRENKE, A.N.; PSARKVA, T.V.; AVSYUK, G.A., otv.red.; OGANOVSKIY, P.N.,

[Franz Josef Land; snow cover] Zemlia Frantsa-Ioaifa; snezhnyi pokrov. Sost.A.N.Krenke i T.V.Psareva. Moskva, 1960. 151 p. (Materialy gliatsiologicheskikh issledovanii).

1. Akademiya nauk SSSR. Institut geografii. (Franz Josef Land--Snow)

PSAREVA, Ye.N., kand.sel'skokhoz. nauk; MATVEYENKO, T.M.

Role of intravarietal selection in the resistance of tobacco to the cucumber mosaic virus CMV-1. Agrobiologiia no.3:413-418 My-Je '63. (MIRA 16:7)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut tabaka i makhorki, g. Krasnodar. (Tobacco--Disease and pest resistance) (Gucumber mosaic virus)

PSAREVA, Ye.N., kand.sel'skokhozyaystvennykh nauk; MATVEYENKO, T.M.

Pathogenicity of the root rot of tobacco in various tobacco growing regions of the U.S.S.R. Agrobiologiia no.3:388-396 My-Je '62. (MIRA 15:10)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut tabaka i makhorki, Krasnodar.

(TOBACCO--ROOT ROT)

	ACCESSION NR: AP3002125 EWP(q)/FWT(m)/RDS AFFTC/ASD JD	
	AUTHOR: Psar'ov V. I., Kostur M. L., Obstra A. V.	k og i
e"	the melt.	•
	SOURCE: Ukrains'kyy fizychnyy zhurnal, v. 8, no. 6, 1963, 694-699	•
	alloy, melt, crystal growth, silver alloy, electrical conductivity, thermal electromotive force, thermal E.M.F., transport phenomena, mercury alloy, Insb,	The second second
	ABSTRACT: The authors suggested a method for separation of crystals of CdSb and InSb compounds from the liquid excess phase (Cd, In) by means of centrifuging the liquid melt. The composition of compounds and cooling conditions are given chases take place simultaneously as the melt is moving. The method was used for alloying CdSb crystals through alloys of Cd Sb and their subsequent separation alloyed with up to 1 to 1.5% of silver. This resulted in an increase in	
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ACCESSION NR: AP3002125

clectrical conductivity and decrease in thermal electromotive force. The CdSb compound was also alkoyed with mercury, with no appreciable effect on either compound was also alkoyed with mercury, with no appreciable effect on either compound was also alkoyed with mercury. The results are shown on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The results are shown on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electrical conductivity or thermal electromotive force. The cdSb compound on electromotive force for

PSAR'OV, V.I.

Calculating the activation energy value of the carbide-coagulation process in steels [with summary in English]. Dop. AN UESR no. 4:386-389 '58.

1. Dnioropetrovs'kiy dershavniy universitet. Predstaviv AN UESR V.M. Sviechnikov [V.N.Svachnikov].

(Steel)

(Gementation(Metallurgy))

L 1637C-65 EWT(m)/EWP(t)/EWP(b) IJP(c)/ESD(t)/AFWL/ASD(a)-5 JD ACCESSION NR: AP4044172 S/0185/64/009/008/0900/0907

AUTHOR: Kostur, M. L. (Kostur, N. L.), Tsanta, V. I. (Psarev. V. I.)

TITLE: Solubility and effect on certain elements of the physical properties of InSb and CdSb

SOURCE: Ukrayins'ky*y fizy*chny*y zhurnal, v. 9, no. 8, 1964, 900-907

TOPIC TAGS: elements solubility, InSb alloy, CdSb alloy, semiconductor, emflattice parameter $\sqrt[n]{1}$

ABSTRACT: The solubilities of Ga, Hg, and Bi in InSb, and of Bi in CdSb have been investigated. The concentration of the admixture was determined by x-ray diffraction analysis from the changes of the lattice parameters. The solubilities of Ga up to 10.1 at.%, of Hg up to 6.2 at.%, and of Bi up to 0.26 at.% in the InSb crystals were found to depend on their dispersion. An expression was found for the change of the InSb lattice as a function of Ga content. Alloying of InSb with Ga and Hg produces a drop of thermal emf and an increase of the concentra-

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ACCESSION NR: AP404417	는 이 교육을 위한 수 있는 이 경우 사람이 모르는 것 같다. 그는 기술 가장이 가는 그는 이 경우를 가져왔다.	4 10 2 10 10 10 12 12 12 12 12 12 12 12 12 12 12 12 12	
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CIA-RDP86-00513R001343510003-1

L 15750-65 EWT(m)/EWP(t)/EWP(b) IJP(c)/ESD(t)/ESD(gs)/AFWL/ASD(a)-5/ASD(f)-2/ACCESSION NR: AP4042807 ASD(m)-3 JD S/0126/64/018/001/0047/0054

AUTHOR: Psarev, V. I.; Dobry*den', K. A.

TITIE: Phase transformations in Cd - Sb alloys

SOURCE: Fizika metallov i metallovedeniye, v. 18, no. 1, 1964, 47-54

TOPIC TAGS: phase transformations, Cd, Sb, crystallization, eutectic, stability, spheroidization, coalescence, equilibrium, metastability, isothermal holding, lattice distortion

ABSTRACT: Data available on the formation and transformation of metastable Cd₃Sb₂ compounds into the stable state are highly controversial. The authors investigated the conditions under which Cd-So alloys form, their structural state and the character of transformation. The formation of metastable phases is explained by the fact that insofar as supersaturation of CdSb crystals with Cd depends on their dispersity, the crystallization of the stable sutectic (CdSb plus Cd) or the dispersity of primary CdSb crystals under conditions of non-equilibrium leads to the dislocation of the composition towards a higher Sb contents. The Cd₃Sb₂ compound crystallizes from Cd - Sb melts in equilibrium. Specimens were

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L 15750-65 ACCESSION NR: AP4042807

methods. The crystallographic examinations combined with thermographic and x-ray methods. The crystallographic structure of the particles was studied by electrolytic separation. After an 11-hour isothermal holding period at 290 C, the crystal lattice of impure CdSb was almost identical to that of the pure compound. In rapidly cooled specimens, the degree of lattice distortion was affected by the dispersity of particles. Subsequent heating led to the stabilization of the non-equilibrium lattice of CdSb particles. However, a complete restoration of the lattice equilibrium is achieved in the process of coalescence and spheroidization of the non-equilibrium particles. Rapid cooling from 400 to 410 C produces a three-phase structure, CdSb crystals having the highest microhardness followed by Cd₃Sb₂ crystals and the metastable eutectic. The result of tests designed to separate crystallization of the stable and the metastable eutectic confirm the assumption that the Cd₃Sb₂ phase forms in the melt of a compound that is changed and in a state of equilibrium. Orig. art. has: 6 figures.

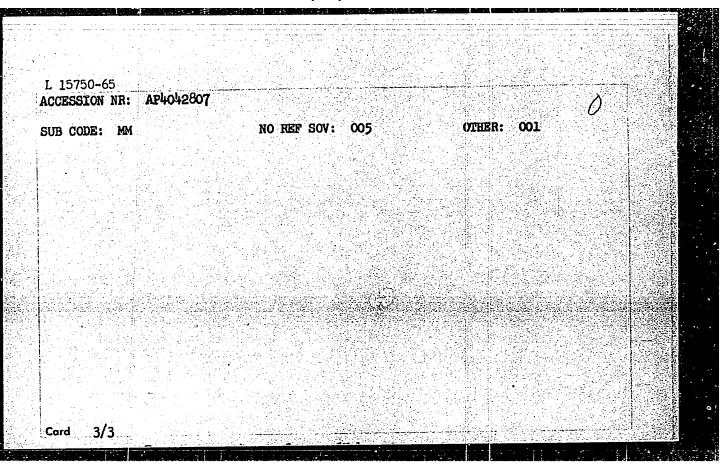
ASSOCIATION: Chernovitskiy gosuniversitet (Chernovitsy State University)

SUBMITTED: 28Apr63

FNCL: 00

Card 2/3

"APPROVED FOR RELEASE: 06/15/2000 CIA-RDP86-00513R001343510003-1



USSR/Cultivated Plants - Commercial. Cil-Bearing. Sugar-Bearing.

: Ref Zhur Biol:, No 18, 1953, 82463 Abs Jour

: Psareva, Ye. Author

: All-Union Scientific Research Institute of Tobacco Inst

: Differences in the Quality of Tobacco Seeds and Morpho-Title

logical Characteristics of the Plants.

: Byul. nauchno-tekhn. inform. Vses. n.-i. in=t tabaka i Orig Pub

makhorki, 1957, 3, 32-38

: In 1946-1948, heterogeneity of the biological properties Abstract

of tobacco seeds was studied in relation to their position on the raceme. Tests were conducted inder field conditions with the selected varieties Dynbek 44 and Tyk-Kulak 235. Results showed that the difference in the quality of the seeds explained by the position of the seed vessel on the raceme (in the center or on the

card 1/3

USSR/Centivated Plants - Commercial. Oil-Bearing. Sugar-Bearing. M

Abs Jour : Ref Zhur Biol., No 18, 1958, 82463

periphery), shows itself both in the absolute weight of the seeds and in a number of variety characteristics: length of the vegetation period, yield and the form of the leaves. It does not depend on the difference in the period of blossoming and seed formation and is a biological characteristic of the tobacco. The farther the seed vessel is located from the center of the raceme the stronger the deviation in the variety characteristics (in the undesirable direction) of the plants grown from them. In contrast to the principal phenotype, plants obtained from the periphery seeds are characterized by retarded maturing, lowered yield and a narrower and more elongated form of the leaves. These data confirm the need for using the "shaping" of the racemes, that is the removal of the side seed vessels. Leaving them on the plants lowers the miformity of the variety plantings by creating forms deviating from the main mass of the plants

Card 2/3

- 101 -

Paramag. stism of Mm-Cu alloys at temperatures higher than that of melting. Fiz. met. i metallowed. 14. no.3:378-382 S '62. (MIRA 15:9) 1. Moskovskiy gosudarstvennyy universitet imeni M.V.Lomonosova. (Manganese-copper alloys---Magnetic properties) (Metals at high temperatures)

PSCHENICHNYY, I.P.; SHTKYGARDT, Yu.N.; MESHCHERYAKOV, A.V.; VASIL'YEV, V.N.; SOKOLOVA, E.F.; EROVKOVICH, E.D.; RUBANOVSKIY, B.R.; LUR'YE, R.G.; PARAKHONYUK, Z.M.; GOROKHOVSKIY, B.I.; ZHDANOV, V.S.; GORBUNOVA, Z.V. GLIKIN, M.I.; TAVAR'YAN, E.A.; SUKHODOLYA, Ye.I.

Abstracts. Kardiologiia 4 no.4:87-90 Jl-Ag * 64. (MIRA 19:1)

3):[03 Z/009/60/010/05/005/040 **B**112/**B**153

5,3600

AUTHORS: M. Repáš and J. Pschera Preparation of Anhydrous Ethylene Chlorhydrine

PERIODICAL: Chemický Průmysl, Vol 10, 1960, Nr 5, pp 238-240

ABSTRACT: The authors present a study of the laboratory preparation of anhydrous ethylene chlorhydrine from ethylene oxide and hydrochloric acid. They have adopted the preparation in the liquid phase and used ethylene chlorhydrine, the final reaction product as solvent, utilising the good solubility of HCl in that medium. They have studied the effect of temperature and concentration of reactants on the conversion of ethylene oxide and the formation of byproducts. For the study of temperature effects on yields, equimolar ratios of ethylene oxide and HCl were chosen. It was seen that a 92.5% conversion was obtained at 30 oc. At higher temperatures yields of ethylene chlorhydrine decreases rapidly while the formation of byproducts increases. Results of this series of experiments are summarized in graphs. Another series of experiments included the effect of varying proportions of reactants on yields. An excess of HCl suppresses the formation of byproducts, equilibrium being reached at

Card 1/2

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Z/009/60/010/05/005/040 E112/E153

Preparation of Anhydrous Ethylene Chlorhydrine

50% excess. An excess of ethylene oxide, on the other hand, enhances byproduct formation, according to equation

 ${\rm C1}-{\rm GH}_2-{\rm CH}_2-{\rm 0}-{\rm CH}_2-{\rm CH}_2-{\rm OH}$

The authors have also studied the effect of batch sizes on yields and the effect of refrigeration arrangements on the efficiency of conversion.

There are 4 figures and 5 references, of which 2 are English, 1 French, 1 German and 1 Gzech. Card 2/2

ASSOCIATION: Výzkumný ústav pre petrochémiu, Nováky (Research Institute for Petrocarbon Chemistry, Nováky)

SUBMITTED: October 12, 1959

PSCHERA, Jiri

"Processes and equipment in the chemical industry" by Jaroslav Schneider and others. Reviewed by Jiri Pschera. Chem prum 13 no.9:487-488 S '63.

1. Vyzkumny ustav organickych syntez.

CIA-RDP86-00513R001343510003-1 "APPROVED FOR RELEASE: 06/15/2000

5/081/63/000/002/050/088 B171/B102

AUTHORS:

Repás, Milan, Mistrik, Edmund Juraj, Pschera, Jiří

TITLE:

Liquid- phase-method and plant for continuous preparation of ethylene chlorohydrin from ethylene oxide and hydrogen

chloride :

PERIODICAL:

Referativnyy zhurnal. Khimiya, no. 2, 1963, 401, abstract 2N23 (Czechosl. patent 100737, Aug. 15, 1961)

TEXT: A patent has been granted for the preparation of concentrated (95-100%) ethylene chlorohydrin (I). Ethylene oxide (II) and Hol are fed through perforated pipes or ceramic bubblers into a cylindrical reactor (RT), filled with a liquid which, although miscible with I and solvent of II and HCl, does not react with any of these substances. Raw materials are fed at the rate of 100-1000 1/hr per 1 1 of RT capacity, at a temperature from -30 to 100°C (preferably ~70°C), using 1.005-1.01 mole of HCl to 1 mole of II. The incoming minute bubbles of II and HCl vigorously react to form I. The mixture, warmed by the heat of the reaction is water-cooled in a thermo-siphon and returned into the RT. The

Liquid - phase-method an		B171/B102		
generated I is continuo 175-1750 l/hr per liter with respect to II. Fo are fed at 0-25°C into of 350 mm and a capacit 99%). A flow sheet is	r example, if 100 the lower part of	showing thus a yi kg/hr of II and the RT (having a	.eld of $90-100$ 51.2 m ³ /hr of in inside diam	% HCl

PSCZOLOWSKI, T.

PSCZOLOWSKI, T. Radiophony in the Palace of Culture and Science. p. 18, Vol. 6, no. 1, Jan. 1956 RADIOAMATOR. Warszawa, Poland

SOURCE: East European Accessions List (EEAL) Vol. 6 No. 4 April 1957

F. M. The section

(MIRA 17:6)

PSEHKOVA, V.M.; KHROMOV, S.I.; SHAKHPARONOV, M.I.

December Flenum of the Central Committee of the CPSU and the chemical science at Moscow university. Vest. Mosk. un. Jer. 2:

Khim. 19 no.1:3-19 Ja-F '64.

AMASTASIU, St.; STOICA, Rodica; JELESCU, Eugenia; PSEMESCHI, Valeria; BERCCVICI, Rodica

New aspects in the technology and control of alkyl phenol production. Rev chimie Min petr 15 no.2:71-75 F 164.