

Pb-Zn, Cd, Hg

$M_2 X$, $M X_3$, $M_2 X_3$,
 $M = \text{Pr}, \text{Ce}, \text{La}$. $X = \text{Sn}$, (Tur)

$M'Y$, $M'Y_2$, $M'Y_3$
 $M' = \text{Pr}, \text{La}, \text{Ce}$ $Y = \text{Ag}, \text{Au}$ (Tur)

Pr Pb , Pr Pb_3 , $\text{Pr}_2 \text{Pb}$, Pr Cu_2 , Pr Cu_6 , Pr Al_2 , Ce, La ,
 Pr Tl , Pr Tl_3 , $\text{Ce}_2 \text{Pb}$, Ce Pb_3 , Ce Tl , Ce Tl_3 , Ce Cu_2 ,
 Ce Cu_6 , Ce Al_2 , La Pb , La Pb_3 , $\text{La}_2 \text{Pb}$, La Tl , La Tl_3 ,
 La Cu_2 , La Cu_6 , La Al_2 (Tur)

Rolla L., Tandelli A., Cannari G., Vozel R.,
 Z. Metallkunde, 1943, 35, 29-42 5

Соединение в системах — 1963
РЗ-2-Т.

IT = Mg, Zn, Al, Fe, Cu)
1 кр. спр-ре)

Тиагоминский Е. И., Кричевский
Н. И., Черкашин С. С., Заречник
О. С., Заслуцкий Н. Н., Евдокименковы
В. С., Редкоземельн. минералы;
М., АН СССР, 1963, 67-70

РХ. 1966 25263 № 1000 ф-к

ThCr₂ / JA - piezokristallografiia 1963
releevančeskiy

(kristal. cečip-pa) VIII 4365

Storm A. R., Benson K. E.

Acta crystallogr.

1963, 16, N^o 7, 401-403.



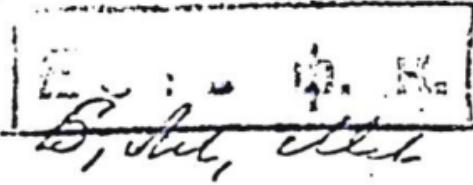
PX, 1964, 16 B 234

clced. / all = y, Ce, Pr, Nd, Sm, Eu
Lu, Gd, Tb, Dy, Ho, Er, Tm, Yb,
Lu).

VIII 4262

Johnson L, Schablaeke R,
Takai B, Anderson K,
Trans. Metallurg. Soc. AIME,
1964, 230, n6, 1485-1486

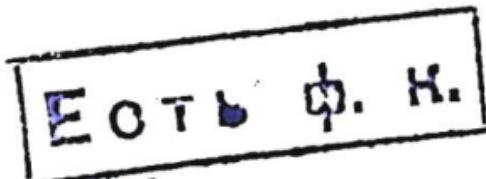
P.M., 1965, 321.41



ELIX (el = P-32), spouse Ce 2, Pen; 1965
X = Cei, Ing., En, Cal, Hz u. Sl)
(spouse! esp-pa) VIII 4235

Tandelli A; Palenzona A;

Y. Less - Common vultals,
1965, ♀, mt, 1-6



Чка

P.K., 1966, 33348

$\text{La}_2\text{Cd}_{17}$; CeCd_6 ; PrCd_6 ; NdCd_6 ; VIII 119
 SmCd_6 ; LaCd_{11} ; CeCd_{11} ; PrCd_{11} ; 1966
 NdCd_{11} . Kremn. cíp-pa Tu

Johnson J., Anderson K. G.,
Blomquist R.A., Trans. Amer.
Metals, 1966, 159, 352-5

B, Am. U.S.A. 1948 B, H.

Y_{Hg_4} , La_{Hg_4} , Ce_{Hg_4} , Pr_{Hg_4} , Nd_{Hg_4} , Sm_{Hg_4} ,
 Gd_{Hg_4} , Y_{Hg_3} , La_{Hg_3} , Ce_{Hg_3} , Pr_{Hg_3} , Nd_{Hg_3} , Sm_{Hg_3} , Gd_{Hg_3} , Dy_{Hg_3} , Ho_{Hg_3} , Er_{Hg_3} , Y_{Hg_2} ,
 La_{Hg_2} , Ce_{Hg_2} , Pr_{Hg_2} , Nd_{Hg_2} , Sm_{Hg_2} , Gd_{Hg_2} ,
 Dy_{Hg_2} , Ho_{Hg_2} , Er_{Hg_2} , Y_{Hg} , La_{Hg} , Ce_{Hg} ,
 Pr_{Hg} , Nd_{Hg} , Sm_{Hg} , Gd_{Hg} , Dy_{Hg} , Er_{Hg} ,
 Ho_{Hg} . (Th)

VIII 4278

Kirchmayer H. R., Lutzscheider W.;
Z. Metallkunde

1966, 57, nr 10, 425-731

E. 1. 4. N.

P.M., 1967, 3210 6. Aufl.

$Zn_6 Zn_{13}$ ($Ln = Gd, Dy, Tm, Ln$) 1966.

$Zn_2 Zn_{17}$ ($Ln = Pr, Nd, Sm, Er, Yb, Lu$)

$Ln Zn_{12}$ ($Ln = Sm, Gd, Ho, Tm, Er$)

$Ln Zn_{13}$ ($Ln = La, Eu, Yb$)

открытие вест. Н.У. Урал. А.С.,

изд. АН СССР. Челябинск. из
издательства, 1966, 2, № 4, 630.

РХ, 1966, 205377

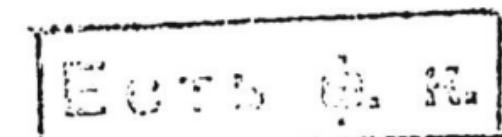
111

R₂N₂ / R - pegmatites (seeir.) (P_u)¹⁹⁶⁶

Michel D.Y., Ryba E, VIII 375

Kjivival P.K.

J. Less-Common Metals,
1966, 11, pt. 67-69



5 Oct. 1966

PX, 1967, 2B 600

AB_2 /A-Sc, Y, P₃M; B-Cu, Ag, Al¹⁹⁶⁷
(spurcii. cusp-pa) VIII 4440

Dwight A.E., Downey Y. D.,

Conner R.A., Jr.,

Acta crystallogr.,

1967, 22, n⁵, 445.

P.M., 1967, 102131 clse

VIII

M_{Zn}₂

M = La, Pr, Nd, Sm, Gd, Tb, Dy,
Ho, Er, Tm, Lu.

1968.
(1967)

abc

VII 718

Foznasini Maria L., Mezlo Franco.

Atti Accad. naz. Lincei. Rend. Cl. sci. fis., mat. e
natur., 1967 (1968), 43, n° 5, 357-363.

Sui composti di formula MX₂ formati dalle
terze rare con lo zinco.

E... • Φ H.

61

19

PNR, 1969, 45860.

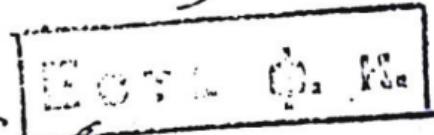
M_{Zn}₁₃ (M-Sr, Ca, La, Eu, Y) 1967

M_{Zn}₁₂ (M-Sm, Gd, Tb, Dy, Ho, Er,
M_{Zn}₁₁ (Tb, Eu)) VIII 338

M_{Zn}₁₇ (M-Th, 'u, see P3 et spacie' Pm, Eu)

(spicciu. ciup-pa)

Tandelli A, Palenzona A.,
J. Less-Common Metals,
1967, 12, n^o 5, 333-343



P.M. 1967, 112135 5, M.I., U.S.

$\text{La}_{\text{Zn}}_{13}$; $\text{Zn}_{1,2}\text{Y}$, $\text{La}_{\text{Zn}}_{11}$; $\text{Ce}_{\text{Zn}}_{11}$, $\text{Ce}_{\text{Zn}}_{11}$, VIII 4896
 $\text{Pr}_{\text{Zn}}_{11}$, $\text{Nd}_{\text{Zn}}_{11}$, $\text{La}_2\text{Zn}_{17}$, $\text{Ce}_2\text{Zn}_{17}$, $\text{Pr}_2\text{Zn}_{17}$, $\text{Nd}_2\text{Zn}_{17}$, 1967
 $\text{La}_{\text{Zn}}_{7,3}$; Ce_{Zn}_7 , Pr_{Zn}_7 ; $\text{Nd}_{\text{Zn}}_{6,5}$, $\text{La}_{\text{Zn}}_{5,2}$, $\text{Ce}_{\text{Zn}}_{5,2}$,
 $\text{Pr}_{\text{Zn}}_{5,3}$; Y_{Zn}_5 ; $\text{Ce}_{\text{Zn}}_{4,5}$; $\text{Pr}_{\text{Zn}}_{4,5}$; $\text{Nd}_{\text{Zn}}_{4,5}$, La_{Zn}_4 ,
 $\text{Y}_{\text{Zn}}_{4,5}$; Ce_{Zn}_3 ; Pr_{Zn}_3 ; Nd_{Zn}_3 ; Y_{Zn}_3 ; Y_3Zn_{11} ;
 $\text{Ce}_3\text{Zn}_{11}$, $\text{Pr}_3\text{Zn}_{11}$, $\text{Nd}_3\text{Zn}_{11}$, La_{Zn}_2 , Ce_{Zn}_2 ; Pr_{Zn}_2 ;
 Nd_{Zn}_2 ; Y_{Zn}_2 ; La_{Zn} , Ce_{Zn} , Pr_{Zn} , Nd_{Zn} , Y_{Zn} ,
Kjær. CIP-pa

Velleckis E., Schablaške R.V., Johnson T.,
Feder H.M., Trans. Metallurgy. Soc. AIMM,
1967, 239, N1, 58-63

PM70

M1

LiFe_4 , LiFe_3 , LiFe_2 , LiFe 1968

VIII 4280

Prieschnagk H.R., Zugsehider 25,

Z. Metallkunde, 1968, 59, 4,

296-297.

EATB B. H.

5, III, (G) 102

PKC Xue, 205854 (1968)

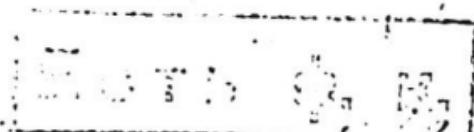
RX₂, P = P₃ H; X = Lg, lg, M, K, ec, ¹⁹⁶⁸
Lg, M, K, ec, ¹⁹⁶⁸
(spars. comp-pat)

VIII 4231

Pandelli R, Palenzona Q.

J - Lg - Common 1968

1968, 15, d3, 273 - 284



Pub, 1968, 122164

$\text{Co}_3\text{Zn}_{11}$, $\text{Pd}_3\text{Zn}_{11}$, $\text{Nd}_3\text{Zn}_{11}$, Y_3Zn_{11} , $\text{Tb}_3\text{Zn}_{11}$, $\text{Sm}_3\text{Zn}_{11}$, $\text{Dy}_3\text{Zn}_{11}$

VII 503b/1970
Tifves. Cf-p;

Brunzzone G., Fornasini M.L., Merlo F.

Colloq. int. CNRS, 1970/180/1, 125-133 (pp.).

OS unrefined amorph. cagedularz. P3 11 C
 Zn .

① Mi 14

PM, 1971, 12/45

LaCd_{II}, CeCd_{II}, PrCd_{II}, LaZn_{II}. 1970

CeZn_{II}, PrZn_{II} (OGr) 8 VIII 3735

Johnson Irving, Yonco Robert & St.

Steel. Trans., 1970, 1, NY, 905-910 (ann.)

Thermodynamics of cadmium- and zinc-rich alloys in the Cd-60, Cd-Ce, Cd-Pr, Zn-60
Zn-Ce and Zn-Pr systems

PMK Nov, 1970

225847

O

10

DEC 10 1970
MOP

MOP

111 - 3495

1970

PrZn
PrZn₂
Pr₂Zn₁₇

T_m

(70804w) Phase relations and crystallographic data for the praseodymium-zinc system. Mason, John T.; Chiotti, Premo (Ames Lab., U.S. At. Energy Comm., Ames, Iowa). *Met. Trans.* 1970, 1, 2119-23 (Eng). Metallographic, thermal, x-ray, and resistivity data were employed in establishing the phase diagram of the Pr-Zn system. Eight compds., 3 eutectics, and 1 eutectoid were obsd. The compds. PrZn, PrZn₂, and Pr₂Zn₁₇ melt congruently at 882, 896, and 978°, resp. The compds. PrZn₃, Pr₃Zn₁₁, PrZn_{4.46}, PrZn₂₂, and PrZn₁₁ undergo peritectic decomprn. at 833, 855, 891, 956, and 743°, resp. The eutectic temps. and compns. in wt. % Zn are 576° at 11.9, 833° at 39.0, and 830° at 56.8%. The eutectoid reaction occurs at 558°, and 5.2% Zn. The lattice parameters of the compds. in the system were detd. by x-ray powder-diffraction methods. Single-crystal x-ray techniques were used to show that PrZn₃ has the YZn₃ (space group *Pnma*)-type structure. Nat L. Shepard

C.S. 1970 73.14

Pr Zn X

VIII-4438

1941

115001w Thermodynamic properties of praseodymium-zinc alloys. Chiotti, Premo; Mason, John T. (Ames Lab., Iowa State Univ., Ames, Iowa). *Met. Trans.* 1971, 2(4), 967-73 (Eng). Zn vapor pressure data obtained by the dewpoint method were employed in calcg. the thermodynamic properties of both liq. alloys and solid compds. The liq. alloy data were fitted to a relation which was then transformed into a 2nd expression by integration of the Gibbs-Duhem equation. The std. free energy changes for the reactions $\text{Pr}(\alpha) + x\text{Zn}(l) \rightarrow \text{PrZn}_x(s)$ are: $\Delta F^\circ(\text{PrZn}) = -17,210 + 3.80T$; $\Delta F^\circ(\text{PrZn}_2) = -29,160 + 7.56T$; $\Delta F^\circ(\text{PrZn}_3) = -38,580 + 12.34T$; $\Delta F^\circ(\text{PrZn}_{3.67}) = -44,600 + 15.30T$; $\Delta F^\circ(\text{PrZn}_{4.46}) = -51,200 + 18.75T$; $\Delta F^\circ(\text{PrZn}_{7.33}) = -72,830 + 31.72T$; $\Delta F^\circ(\text{PrZn}_{8.5}) = -80,340 + 36.65T$; and, $\Delta F^\circ(\text{PrZn}_{11}) = -89,350 + 45.48T$.

Nat L. Shepard

C.A. 1941. 44. 2d

(P3H) Zn_{~5} - крист. сим. ~~№ 2394~~ 1973

"J. Less-Common Metals", 1973, 32, N3,
391-394 (ascr.), Green M. L.

XVIII-1004

Неподвижные соединения тунна
P3H (Zn)_{~5}, некоторые спутники [тунна]
CaCu₅ (Дад).

Прим., 1960, 1974 (Mrs.)

$\text{Li}_3\text{Mg}_{2.5}\text{Ge}_{2.5}\text{O}_{12}$ u gp (Tm) VII 5380
1974
 $\text{3d}_2\text{Co}_3$; Ge_2O_{12} min a $\text{M}_3\text{M}_{2.5}\text{Ge}_{2.5}\text{O}_{12}$
zge M = Feegkozeum (Pr) eit. M' = Mg, Co, Ni, Zn

Bayer 6.,
Matierwissenschaften, 1974, 58,
N12, 622

5

(P)

①

20

CA72

PrCu₂ (C_P) XVIII 280 1974

Wun M., Phillips N. E.,

Phys. Lett A 1974, 50A (3) 195-6.

Low-temperature heat
capacity of praseodymium-
copper.

C.A. 1975, 82 n 20. 130045x.

5



Pr^N₂Zn,
Nd^N₂Zn

XVIII-536

1975

+1

Td

D 1410836 Jagodic properties, magnetic structure, and crystal field, in praseodymium-zinc and neodymium-zinc. Marin, Pa. Pierre, J. (Lab. Magie., CNRS, Grenoble, Fr.). *Phys. Status Solidi A*, 1975, 39(2), 549-59 (Eng). The magnetic properties of PrZn and NdZn compds. were studied by means of magnetoo measurements and neutron diffraction. Both compds. are antiferromagnetic below 52 and 70°K, resp. The ordered moments are reduced to 2.35 and 2.55 μ B at low temps. by crystal field effects. Dilatometric measurements reveal a large tetragonal distortion, existing at the Neel temp. and reaching 1.4% & 4.2% in PrZn. The crystal field parameters are coherent in sign with those of the heavy rare earth compds. In PrZn, the magnetostriction is sufficient to explain the 4-fold direction of the moment. The ambiguities in the magnetic structure are discussed.

C.A. 1975 83 N/6



$ZnLn_4[SiO_4]_3$ [Ln = La, Ce, P₂, | 1976
Nd, Sm, Eu Gd, Tb, Dy] (T_m)

Федоров Н.В., Тунчик Г.Я. VIII-1067
Судоров П.Н.

Ж. неорганической химии, 1976, 21, N3, 666-669

Силикат-апатиты редкоземельные
Элементы и уранка состава $ZnLn_4[SiO_4]_3$.

РДИХИ, 1976

1976

5(φ) Ии

PrCd

1983

(T_C)

198: 208386m Electrical and thermal transport properties of praseodymium cadmium single crystals. Sousa, J. B.; Pinto, R. P.; Braga, M. E.; Aleonard, R.; Morin, P.; Schmitt, S. (Cent. Fis., Univ. Porto, Porto, Port. 4000). *J. Phys. F* 1983, 13(4), 827-38 (Eng). The temp. dependences of the elec. resistivity (ρ), thermopower (S), thermal cond. (K), and temp. derivs. $d\rho/dT$ and dS/dT were measured in single crystals of PrCd, with the elec. field and temp. gradient applied along the $\langle 110 \rangle$ direction. Three phase transitions were identified: a structural transition at $T_s = 125$ K, where the high-temp. CsCl structure ceases to exist, an antiferromagnetic-paramagnetic transition at $T_N = 40$ K, and an unknown phase transition at $T^* = 20$ K. Particular attention was focused on the details of the structural transition at T_s , namely on the peculiar irreversible behavior obserd. there. Different contributions to the transport coeffs. of PrCd were estd. The lattice stability in CsCl structure intermetallic compds. is discussed.

C.A. 1983, 98, n24

1984

CdPr₂Se₄
CdPr₄Se₄

16 Б3089. Исследование системы CdSe—Pr₂Se₃.
Агаев А. Б., Кулниева С. А., Кулниева У. А.
«Азерб. хим. ж.», 1984, № 6, 95—97 (рез. азерб.)

Методами ДТА, РФА, микроструктурного анализа, а также измерением микротвёрдости и определением плотности исследовано взаимодействие в системе CdSe—Pr₂Se₃. Построена диаграмма состояния системы. Установлено, что система CdSe—Pr₂Se₃ является квазибиарным сечением тройной системы Cd—Pr—Se. Отмечено образование конгруэнтно плавящегося соединения CdPr₂Se₄ и инконгруэнтно плавящегося CdPr₄Se₄. Тройное соединение CdPr₂Se₄ образуется по перитектической реакции.

По резюме

X. 1985, 19, n 16

CdPr₂Se₄

1984

103: 28073v Study of the cadmium selenide-praseodymium selenide ($\text{CdSe-Pr}_2\text{Se}_3$) system. Agaev, A. B.; Kulieva, S. A.; Kulieva, U. A. (Azerb. Gos. Univ., Baku, USSR). *Azerb. Khim. Zh.* 1984, (6), 95-7 (Russ). The phase diagram was constructed from DTA, microhardness, x-ray phase anal., and microstructural data. The $\text{CdSe-Pr}_2\text{Se}_3$ system is a quasi-binary section of the Cd-Pr-Se section. The compd. CdPr_2Se_4 congruently m. $\sim 1150^\circ$ and CdPr_4Se_7 incongruently m. $\sim 1130^\circ$ and 66.7 mol % Pr_2Se_3 . Solv. of Pr_2Se_3 in CdSe is 3 mol %.
(Pm)

C. A. 1985, 103, N.Y.

PrCd 1987
Kadomatsu H.,
Karisu M., et al.

η_{tr} ; J. Phys. F: Met. Phys.
1987, 17(12), L305-L309.

(c.c. LaAg; $\overline{1}$)

$\text{Pr}_2\tilde{\text{Zn}}_{17}$

(On 37236)

1993

118: 220757x Specific heats of rare earth zinc (R_2Zn_{17}) intermetallic compounds. Marquina, C.; Kim-Ngan, N. H.; Bakker, K.; Radwanski, R. J.; Jacobs, T. H.; Buschow, K. H. J.; Franse, J. J. M.; Ibarra, M. R. (Fac. Cienc., Univ. Zaragoza, Spain). *J. Phys.: Condens. Matter* 1993, 5(13), 2009-16 (Eng). Sp. heat measurements were performed on polycryst. R_2Zn_{17} compds. ($\text{R} : \text{Pr, Nd, Tb, Dy, Ho, Er, and Y}$) at 0.3-150 K. For all the compds., except for $\text{R} : \text{Y}$, magnetic anomalies in the sp. heat were obsd., which indicates the existence of magnetic order at low temps.

(C_p)

□

(46)

$\text{Nd}_2\tilde{\text{Zn}}_{17}$, $\text{Pr}_2\tilde{\text{Zn}}_{17}$, $\text{Dy}_2\tilde{\text{Zn}}_{17}$,

$\text{Ho}_2\tilde{\text{Zn}}_{17}$, $\text{Er}_2\tilde{\text{Zn}}_{17}$,

$\text{Y}_2\tilde{\text{Zn}}_{17}$

C.A. 1993, 118, N22

Pr₃Zn₂As₆

1999

F: $\frac{\text{Pr}_3\text{Zn}_2\text{As}_6}{1}$ (структура)
P:

02.02-19Б2.32. Арсенид празеодима цинка $\text{Pr}[3]\text{Zn}[2]\text{As}[6]$: кристаллизующийся с структурном типа $\text{HfCuSi}[2]$, вариант с вакансиями. The praseodymium zinc arsenide $\text{Pr}[3]\text{Zn}[2]\text{As}[6]$: crystallizing with a vacancy variant of the $\text{HfCuSi}[2]$ type structure / Nientiedt Andre T., Jeitschko Wolfgang // J. S State Chem. - 1999. - 142, N 2. - С. 266-272. - Англ.

Структура

Получено соединение $\text{Pr}[3]\text{Zn}[2]\text{As}[6]$ (I) взаимодействием элементарных компонентов в расплаве $\text{NaCl}-\text{KCl}$. I кристаллизуется с ярко выраженной субъячейкой структурного типа $\text{HfCuSi}[2]$ с двумя формульными единицами $\text{PrZn}[2/3]\text{As}[2]$ в тетрагональной субъячейке с а 400,6, с 1019,7 пм. Упорядоченное распределение занятых и незанятых позиций цинка приводит к решетке в три раза большей. Определены параметры решетки: ф. гр. Pmmn , а 399,89, b 1204,2, с 1019,7 пм, z 2, R 0,039 для 2663 отражений. Обсуждены химические связи в I.