New Penta Bismuth Based Alloy for Shielding Blocks in Mega-Volt Radiotherapy

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Abstract: Microstructure, thermal, electrical and mechanical properties of penta Bi- Sn- Pb based alloys have been investigated. Matrix structure (Formed crystalline phases) and measured physical properties of Bi- Sn- Pb- In- X (X= Cd/or Zn) penta alloys changed with varying composition. The new penta fusible, $Bi_{50}Pb_{15}Sn_{22}Cd_{3}In_{10}$ alloy has best properties such as high density, low melting point and friendly environmental, (reduced toxicity elements Pb and Cd by 40% and 75%, compared used alloys), for shielding blocks in mega-volt radiotherapy. The melting temperature of $Bi_{50}Pb_{15}Sn_{22}Cd_{3}In_{10}$ alloy is ~58 °C and it is density is 10.117 gm/cm³. The elastic modulus of $Bi_{50}Pb_{15}Sn_{22}Cd_{3}In_{10}$ is 29.03 Gpa. Vickers hardness and internal friction values of $Bi_{50}Pb_{15}Sn_{22}Cd_{3}In_{10}$ alloy are 9.72 Kg/mm² and 0.085. The $Bi_{50}Pb_{15}Sn_{22}Cd_{3}In_{10}$ alloy consists of rhombohedral Bi phase, tetragonal Sn phase, face centered cubic Pb phase, hexagonal Cd phase, face centered cubic In phase, Pb7Bi₃ and SnBi intermetallic compounds.

Key words: shielding blocks, fusible alloys, thermal and mechanical properties, electrical resistivity

1. INTRODUCTION

Over the past few years fusible alloys have become a hot subject because they important for industrial and medical applications. Now is the time to understand the correlation between microstructure and physical properties of fusible alloys for solder, bearing and shielding blocks applications. There are an enormous number of alloys made by metals as bismuth, lead; tin and cadmium have in common the feature of a low melting point. In the past they were commonly known as Lipowitz's, Wood's, D'Arcet and Rose alloys and now they are known as Arconim's alloys. In our days other alloys having slightly different compositions in order to fatherly improve their peculiarities had replaced them. Structure, growth properties and physical metallurgy of a series of Pb-Sn-Cd alloys containing up to 60- wt.% Bi and Pb-Sn-Bi were investigated [1]. The results showed that, formation of metastable crystalline phase in the range of composition investigated causes a pronounced increased the electrical resistivity. Adding bismuth to PbCdSn had to the appearance of the crystalline metastable phase which produced hardening effect [2]. Also bismuth atoms act as scattering centers and increasing their concentrations caused an enhancement of resistivity. The physical characteristic of lipowitz's metal and bismuth-lead eutectic allovs as a shielding block for mega voltage therapy machine were studied and analyzed. Internal friction of irradiated and non-irradiated alloys is sensitive to the chemical composition used alloys [3]. Structure, mechanical and electrical transport properties of $Pb_{60}Sn_{38}X_2(X = Sb, Bi, or Ag in weight percent$ as ternary additions) were investigated. Ledbetter's theoretical values of the ratio of shear modulus to elastic modulus, μ/E , are in a good agreement with the experimental results [4]. The crystalline metastable γ (Pb-Bi) phase was appeared in Pb₅₀Sn_{50-x}Bi_x (x= 30 and 50 wt. %) alloys and the lowest value of Vickers hardness was attributed to the formation of intermediate metastable phases [5]. Mechanical and electrical properties of PbBiSnCd were dependence on tin content obtained [6]. The ductility of the binary Bi-Sn eutectic alloy has significantly improved by adding small amount Ag [7]. Metastable shift of the solubility limit in Sn-Bi alloys containing 15, 20 and 25 at. % bismuth was produced by splat quenching [8]. The effect of splat cooling on crystal structures and heats of formation of

non-equilibrium intermediate phases of Pb-Bi alloy was studied [9]. Solid solubility extension of Pb in Bi and formation and lattice parameter of several new metastable crystalline solid phases in Pb-Bi including complex Pb-Bi phases was reported [10]. The crystallographic relation-ship between the phases in the Cd-Zn eutectic alloys using standard x-ray techniques on selected areas of bulk eutectic specimens was examined [11]. Also the orientation characteristics of eutectic alloys of Bi-Cd, Cd-Sn, Sn-Zn and Al-Si were studied [12]. Microstructure, electrical, mechanical and thermal properties of rapidly solidified Bi58Sn42 eutectic alloy have been investigated [13]. Thermal properties and microstructure of 58% Bi-42% Sn, 53% Bi-26% Sn-21% Cd,70% In-30% Sn, 50% Sn-50% In and 3% Sn-37% Bi-10% In solder alloys have been studied and analyzed [14]. Attenuation coefficients, structure and physical properties of Bi-Pb-Sn fusible alloys were studied [15]. Microstructure, electrical, mechanical and thermal properties of melt spun bismuth- tin and bismuth- lead- tin- eutectic alloys also investigated [16, 17]. Optical microscopy, X-ray diffractometry, double bridge method, Vickers microhardness testing and dynamic resonance techniques have been used to investigate structure, electrical resistivity, hardness, internal friction and elastic modulus of quenched Bi-Pb-Sn-Cd-Sb penta-alloys, Bi-Pb, Bi-Pb-Sn, Bi-Pb-Cd and Bi-Pb–Sn–Cd fusible alloys [18, 19]. The effect of the quenching rate on structure and some physical properties of the Pb-Sn-Cd melt spun fusible alloys have been investigated by El-Bediwi [20].

The aim of our research was to produce new bismuth based alloy with superior properties as shielding blocks in mega-volt radiotherapy

2. EXPERIMENTAL WORK

Using elements bismuth, tin, lead, indium, cadmium and zinc have a high purity, more than 99.95%. The used alloys, $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_{4}X_{3}(X=Cd \text{ or } Zn)$, were molten in the muffle furnace. The resulting ingots were turned and re-melted several times to increase the homogeneity of the ingots. From these ingots, long ribbons of about 3-5 mm width and ~ 70 µm thickness were prepared as the test samples by directing a stream of molten alloy

onto the outer surface of rapidly revolving copper roller with surface velocity 31 m/s giving a cooling rate of 3.7×10^5 k/s. The samples then cut into convenient shape for the measurements using double knife cuter. Structure of used alloys was performed using an Shimadzu x–ray diffractometer (Dx–30, Japan) of Cu–K α radiation with λ =1.54056 Å at 45 kV and 35 mA and Ni–filter in the angular range 2 θ ranging from 20 to 100° in continuous mode with a scan speed 5 deg/min. Scanning electron microscope JEOL JSM-6510LV, Japan was used to study microstructure of used samples. The melting endotherms of used alloys were obtained using a SDT Q600 V20.9 Build 20 instrument. A digital Vickers micro-hardness tester, (Model-FM-7- Japan), was used to measure Vickers hardness values of used alloys. Internal friction Q⁻¹ and the elastic constants of used alloys were determined using the dynamic resonance method [21-23].

3. RESULTS AND DISCUSSIONS

Microstructure

X-ray diffraction patterns of Bi50Pb15Sn22In10X3 and Bi50Pb15Sn28In4X3(X=Cd or Zn) alloys have lines corresponding to rhombohedral Bi phase, tetragonal Sn phase, face centered cubic Pb phase, hexagonal Cd phase, face centered cubic In phase, Pb7Bi3 and SnBi intermetallic compounds as shown in Figure 1. X-ray analysis of Bi50Pb15Sn22In10X3 and Bi50Pb15Sn28In4X3(X=Cd or Zn) alloys show that, the change in feature of formed phases (such as intensity, broadness of peak, miller indices, position (2θ) , and area under peaks) correlates to the alloy composition. Lattice parameters, (a and c), and unit volume cell (V) of rhombohedral Bi phase in Bi50Pb15Sn22In10X3 and Bi50Pb15Sn28In4X3(X=Cd or Zn) alloys were determined and then listed in Table 1a. Adding Cd/ or Zn to Bi- Pb- Sn- In alloys caused a little variation in Bi lattice parameters and unit cell volume. Crystal particle size of rhombohedral Bi phase in Bi50Pb15Sn22In10X3 and Bi50Pb15Sn28In4X3(X=Cd or Zn) alloys are seen in Table 1b. Adding Cd to Bi- Pb- Sn- In alloys produced higher crystal size of Bi phase than Zn.





Figure 1:- x-ray diffraction patterns of penta Bi- Pb- Sn based alloys

Table 1a:-lattice parameters and unit cell volume of Bi in penta Bi- Pb- Sn based alloys

Samples	a _{rho} Å	сÅ	V Å ³
$Bi_{50}Pb_{15}Sn_{22}Cd_3In_{10}$	4.748	11.87	70.766
$Bi_{50}Pb_{15}Sn_{22}Zn_3In_{10}$	4.816	12.106	71.962
Bi50Pb15Sn28Cd3In4	4.754	11.886	70.994
Bi50Pb15Sn28Zn3In4	4.753	11.879	71.014

Table 1b:- crystal particle size of Birho in penta Bi- Pb- Sn based alloys

Samples	Particle size Å
Bi50Pb15Sn22Cd3In10	357.64
$Bi_{50}Pb_{15}Sn_{22}Zn_3In_{10}$	264.646
Bi50Pb15Sn28Cd3In4	372.43
Bi50Pb15Sn28Zn3In4	339.46

Scanning electron micrographs, SEM, of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys show heterogeneous structure as shown in Figure 2 and that agreed with x-ray analysis. Adding Cd/ or Zn to Bi- Pb- Sn- In alloys caused a change in matrix microstructure of Bi- Pb- Sn alloy.



Figure 2:- SEM of penta Bi- Pb- Sn based alloys

Thermal properties

Thermal analysis is often used to study solid state transformations as well as solid-liquid reactions. Figure 3 shows DSC thermographs of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys. Little variation occurred in exothermal peaks of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys after adding Cd/ or Zn. The melting temperature and other thermal properties of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys are listed in Table 2. Variation on melting temperature, specific heat, enthalpy and thermal conductivity of used alloys depend on its compositions.





Figure 3:- DSC of penta Bi- Pb- Sn based alloys

Table 2:- melting point and other thermal properties of
penta Bi- Pb- Sn based alloys

Samples	Melting	CP	ΔS	K
	point °C	J/g. °C	J/g. °C	W.m ⁻¹ .K ⁻¹
Bi50Pb15Sn22Cd3In10	58.22	0.596	0.224	0.398
Bi50Pb15Sn22Zn3In10	69.09	0.314	0.059	0.701
Bi50Pb15Sn28Cd3In4	69.87	0.743	0.157	0.700
Bi50Pb15Sn28Zn3In4	67.94	1.387	0.274	0.429

Electrical resistivity

Plastic deformation raises the electrical resistivity as a result of the increased number of electron scattering centers. Also crystalline defects serve as scattering center for conduction electrons in metals, so the increase in their number raises the imperfection. The measured electrical resistivity of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys are shown in Table 3. Electrical resistivity of Bi-Pb- Sn- In alloys varied after adding Cd/ or Zn, which depend on alloys compositions.

Samples	ρx10 ⁻⁸ Ω.m	$\sigma x 10^5 \Omega.m$
$Bi_{50}Pb_{15}Sn_{22}Cd_3In_{10}$	254.8	2.421
Bi50Pb15Sn22Zn3In10	226.55	4.414
Bi50Pb15Sn28Cd3In4	226.87	4.408
Bi ₅₀ Pb ₁₅ Sn ₂₈ Zn ₃ In ₄	380.92	2.625

Table 3:- electrical resistivity and electrical conductivity of penta Bi- Pb- Sn based alloys

Mechanical properties

The elastic constants are directly related to atomic bonding and structure. Elastic modului of $B_{150}Pb_{15}Sn_{22}In_{10}X_3$ and $B_{150}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys are listed in Table 4. Elastic modulus values of Bi- Pb- Sn- In alloys changed after adding Cd/ or Zn due to matrix structure change.

The resonance curves of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or } Zn)$ alloys are shown in Figure 4. Calculated internal friction and thermal diffusivity $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3(X=Cd \text{ or } Zn)$ alloys are listed in Table 4. Internal friction of Bi- Pb- Sn- In alloys varied after adding Cd/ or Zn.

Table 4:- elastic modului, internal friction and thermal diffusivity of penta Bi- Pb- Sn based alloys

Samples	Е	μ	В	Q-1	Dth x10-8
	GPa	GPa	GPa	_	m ² \sec
$Bi_{50}Pb_{15}Sn_{22}Cd_3In_{10}$	29.3	10.84	33.05	0.085	35.2
$Bi_{50}Pb_{15}Sn_{22}Zn_3In_{10}$	31.63	11.71	35.31	0.025	34.17
Bi50Pb15Sn28Cd3In4	24.40	9.01	27.86	0.14	43.98
Bi50Pb15Sn28Zn3In4	25.38	9.38	28.67	0.059	41.09



based alloys

Vickers microhardness and minimum shear stress

The hardness is the property of material, which gives it the ability to resist being permanently deformed when a load is applied. Vickers hardness of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or }Zn)$ alloys at 10 gram force and indentation time 5 sec are exposed in Table 5. The minimum shear stress (τ_m) of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or }Zn)$ alloys was calculated and then listed in Table 5. Vickers hardness of Bi- Pb- Sn- In alloys depend on its compositions.

Table 5:- Vickers hardness and minimum shear stress of penta Bi- Pb- Sn based alloys

Alloys	H _v kg/mm ²	$\mu_n kg/mm^2$
Bi50Pb15Sn22Cd3In10	9.72±1.1	3.21
Bi50Pb15Sn22Zn3In10	5.61±0.8	1.85
Bi50Pb15Sn28Cd3In4	21.82±1.13	7.2
Bi50Pb15Sn28Zn3In4	20.92±1.02	6.9

4. CONCLUSIONS

Microstructure (formed phases) of $Bi_{50}Pb_{15}Sn_{22}In_{10}X_3$ and $Bi_{50}Pb_{15}Sn_{28}In_4X_3(X=Cd \text{ or }Zn)$ alloys depend on alloys compositions. Physical properties (thermal, electrical and mechanical) of Bi- Pb- Sn-In- X (X=Cd/ or Zn) alloys effected by alloys compositions. The new alloy, $Bi_{50}Pb_{15}Sn_{22}Cd_3In_{10}$, has better properties for shielding blocks in mega-volt radiotherapy.

5. REFERENCES

- [1] Kamal M, El-Bediwi A.B and Karman M.B, J. Mater. Sci.: Mater. Electro. 9 (1998) 425
- [2] Kamal M, Ewaida M. A, Elleithy M. A and Dawod T. A, Mans. Sci. Bull C. Nat. Sci. and Phys. Sci 27: 1(2000)
- [3] Dawod T. A, M. SC. Thesis, Faculty of Science, Mansoura University, (2000)
- [4] Kamal M, Mazen S, El- Bediwi A. B and El- Naggar M, Radia. Eff. & Def. in Sol. 157 (2002) 467-474
- [5] Kamal M, El-Bediwi A. B, J. Mater. Sci.: Mater. Electro. 11 (2000) 519-523
- [6] Kamal M, Karman M. B and El-Bediwi A. B, U. Scientist Phyl. Sciences, 9: 2 (1997) 164
- [7] Mc Cormack M, Chen H. S, Kammalott G.W, Jin S. J. Electron. Mater. 26: 8 (1997) 954
- [8] Laine E, Lähteenmäkl I, Lehtoranta I, J. Mater. Sci. 13 (1978) 108-112
- [9] Suryanara Yana C, Scripta Metall. 5 (1971) 337-40
- [10] Borromêe-Gautier C, Giessen B. C and Grrant N. J, J. Chem. Phys. 48 (1968) 1905-11
- [11] Straumanis W and Brakss N; Z. Phys. Chem., 30B (1935) 17
- [12] Straumanis W and Brakss N;; Z. Phys. Chem. 38B (1937) 140
- [13] Kamal M, Mazen S, El-Bediwi A, Kashita E, Radia. Eff. & Def. in Sol. 161 (2006) 143-48
- [14] Chriastelova J, Ozvold M, J. of alloys and compounds 457 (2008) 323-328
- [15] Kamal M, Moharram B. M, Farag H, El-Bediwi A and Abosheiasha H. F, Radia. Eff. & Def. in Sol. 161 (2006) 137-142
- [16] Kamal M, Mazen S, El- Bediwi A. B, Kashita E, Radia. Eff. & Def. in Sol. 161 (2006) 143- 148
- [17] Kamal M, Moharram B. M, Farag H, El-Bediwi A and Abosheiasha H. F, Radia. Eff. & Def. in Sol. 161 (2006) 421- 425
- [18] Kamal M, El-Bediwi A. B, Radia. Eff. & Def. Sol. 159: 11- 12 (2004) 651- 657
- [19] El-Bediwi A. B, A.M.S.E., 77: 4 (2004) Modelling A
- [20] El-Bediwi A.B, A.M.S.E., 75: 3 (2002) 1
- [21] Cullity B. D, "Element of x-ray diffraction" Ch.10 (1959) 297
- [22] Sppinert S and Teffit W. E, ASTM, Proc. 61 (1961) 1221
- [23] Schreiber E, Anderson O. L and Soga N, Elastic Constants and their Measurement, McGraw-Hill Book Company, Ch. 4 (1973)