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Fabrication and characterization of Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (x = 0, 2.5, 4, 6, and 8) metallic glasses with high thermal stability and good soft magnetic properties

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Atomic structure and thermal behavior of Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (x = 0, 2.5, 4, 6, and 8) metallic glasses with good soft magnetic properties have been investigated by high-energy synchrotron X-ray diffraction and differential scanning calorimeter, respectively. It has been shown that the extension of the supercooled liquid region first increases and reaches a large value of 95 K and subsequently decreases as a function of Y content. Analysis of the structure factors and pair correlation functions in the reciprocal-space and real-space have indicated that the addition of Y noticeably changes the atomic structure and reduces the degree of the medium-range order. Magnetic measurements have implied that the introduction of Y enhances both saturation magnetization and Curie temperatures of the ribbons, while keeping their coercivity very small. The underlying mechanisms for changes in the atomic structure, improving the thermal stability and magnetic properties upon Y addition have been discussed. © 2014 AIP Publishing LLC.

I. INTRODUCTION

Metallic glasses (MGs) exhibit promising physical and mechanical properties compared to their crystalline counterparts.1, 2 They can show a super high strength, large elastic limit, good wear and corrosion resistance, and excellent soft magnetic properties.3–5 It is known that many properties of such materials strongly depend on their electronic and atomic structural features such as the bonding nature among the constituent elements and the degree of the ordering over the short-range or medium-range scales.3 One of the important methods to improve the glass forming ability (GFA), thermal stability and properties of MGs is microalloying or changing the chemical composition through a minor addition of some elements.4 The microalloying can result in formation of the glassy structure with a higher packing density and new local atomic configurations with strong interactions, which can improve the GFA and thermal stability.4 On the other hand, the microalloying can influence the bonding nature, which affects the local ordering, elastic constants, and in some cases makes the glassy structure heterogeneous which altogether improve the mechanical properties of the MGs.5–7

Co-based MGs are well-known alloys owing to their excellent soft magnetic properties, e.g., very small magnetic coercivity, high magnetic permeability, and moderate saturation polarization.8–10 Such properties make them as an ideal candidate for some applications like magnetic sensors, memories, and power devices.10, 11 It has been demonstrated that the increase of metalloid content especially B up to about 30 at. % can enhance their thermal stability, electrical resistivity, and mechanical properties.12, 13 For instance, Inoue et al.14 discovered 2 mm Co$_{33}$Fe$_{26}$Ta$_{5.5}$B$_{31.5}$ glassy rods with very high fracture strength of about 5 GPa and maximum permeability of 550 000. Recently, we have found that the increase of Ta up to 8 at. % results in formation of the Co$_{40}$Fe$_{22}$Ta$_{8}$B$_{30}$ MGs with a significantly better thermal stability but almost the same magnetic properties as the well-known Co$_{43}$Fe$_{20}$Ta$_{5.5}$B$_{31.5}$ MG.15 Moreover, the Co$_{40}$Fe$_{22}$Ta$_{8}$B$_{30}$ bulk glassy alloy with 10 mm diameter indicating ultra-high hardness, excellent thermal stability, and soft magnetic properties has been fabricated by powder metallurgy method.16 It would be possible that thermal stability and magnetic properties of the Co$_{40}$Fe$_{22}$Ta$_{8}$B$_{30}$ MG can be further improved by microalloying through (partial) substitution of Ta with another element. In the current research, Y was selected as substituting atom owing to its larger atomic radii (180 pm (Ref. 17)) compared to other constituent elements and its large negative heat of mixing with B and Co.18 We expected that the addition of Y can produce new atomic pairs and enhances the sequential variations in the atomic size, which results in a denser topological configuration of atoms in the glassy phase and consequently a better thermal stability. As a result, Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (x = 0, 2.5, 4, 6, and 8) glassy ribbons were produced and their atomic structure, thermal stability, and magnetic properties were investigated.

II. EXPERIMENTAL PROCEDURE

The master alloys with nominal compositions Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (at. %) (x = 0, 2.5, 4, 6, and 8) were prepared by arc-melting of the pure Co (99.9%), Fe (99.9%), Ta (99.9%),...
Y (99\%), and crystalline B (99\%) under a Ti-gettered argon atmosphere. The obtained ingots were remelted at least three times to improve their homogeneity. Glassy ribbons were obtained under an argon flow by a single-roller Bühler meltspinning on a copper wheel at 41 m/s tangential wheel velocity.

The phase analysis of the ribbons was carried out by X-ray diffraction (XRD) in reflection geometry, using Co Kα radiation (λ = 0.17889 nm) and in transmission geometry, using a monochromatic high-energy synchrotron radiation (λ = 0.0123984 nm) of 100 keV, at the P07 beam line of PETRA III electron storage ring (DESY, Hamburg, Germany). The illumination of the ribbons at room temperature was carried out for 20 s using an incident beam collimated properly with a cross-section of 1 × 1 mm². A 2-dimensional (2D) detector (Perkin Elmer 1621) placed orthogonal to the X-ray beam was used to record the XRD patterns. The distance of 60 cm was selected between sample and detector in order to achieve the large wave vectors q (q = πx sin θ/2) up to 180 nm⁻¹. The 2D XRD patterns were integrated to the q-space, using FIT2D software package. The elastically scattered intensities were obtained by PDFgetX2 software from the integrated data after performing the different corrections described elsewhere. The total structure factors, S(q), of the as-cast ribbons were calculated from their elastically scattered intensities according to the Faber-Ziman method. The reduced pair correlation functions, G(r), of the as-quenched ribbons were determined from the Fourier transform of S(q) as

\[
G(r) = \frac{2}{\pi} \int_{q_{\text{min}}}^{q_{\text{max}}} q(S(q) - 1) \sin(qr) dq. \tag{1}
\]

The thermal behavior of the ribbons was evaluated by a differential scanning calorimeter (DSC, NETZSCH 404) at a constant heating rate of 20 K/min under a purified Ar. The Hysteresis curves and saturation magnetization of the ribbons were measured by a vibrating sample magnetometer (VSM). The coercivity was determined by a Foerster Coercimat under a magnetic field sufficient for saturating the ribbons. The thermomagnetic behavior and Curie temperature of the ribbons were recorded by a Faraday magnetometer at a constant heating rate of 10 K/min. The measurement errors are within ±1.5 K for the DSC and Curie temperature, ±0.1 A/m for coercivity, and ±0.5 Am²/kg for saturation magnetization.

III. RESULT AND DISCUSSIONS

A. Thermal behavior

The XRD patterns of the as-quenched CoₓFe₂₂Taₓ₈₋ₓ₂B₃₀ ribbons are shown in Fig. 1. Formation of only broad halo pattern without any appreciable Bragg peak for each composition reveals that the prepared ribbons have a fully glassy structure and addition of Y element does not induce any crystallization. Figure 2 shows the DSC plots of the as-cast CoₓFe₂₂Taₓ₈₋ₓ₂YₓB₃₀ ribbons measured at a constant heating rate of 20 K/min. All the compositions except the Ta free alloy (x = 8) indicate a distinct endothermic peak corresponding to the glass transition temperature, T_g, followed by a supercooled liquid region (SLR), ΔT_s (ΔT_s = T_g–T_c, that T_s is the crystallization temperature, measured as the onset of the first exothermic event). According to Fig. 2, the crystallization in Y free alloy (x = 0) proceeds through three stages of exothermic reactions. However, by addition of Y, the second and third exothermic events become weaker and the crystallization takes place through almost a single sharp exothermic peak. The thermal parameters of the as-quenched ribbons are listed in Table I. From this table, it is observed that by addition of Y, the T_g progressively decreases, while the T_c is first enhanced and reaches a maximum value of 984 K for x = 2.5 and subsequently decreases. It is seen that the glassy alloy with x = 2.5 exhibits a largest ΔT_s of 95 K and consequently the highest thermal stability among other compositions investigated in the present work.

In order to identify the phases formed during crystallization, the ribbons were heated isochronally up to the end of primary crystallization peak and subsequently cooled down to room temperature. Figure 3 shows the XRD patterns of the partially devitrified CoₓFe₂₂Taₓ₈₋ₓ₂YₓB₃₀ ribbons. For the Y free alloy, dominant phase is the (Co,Fe)₂Ta₂B₆ with a face-centered cubic (FCC) structure. In addition, small

![FIG. 1. The XRD patterns of the CoₓFe₂₂Taₓ₈₋ₓ₂YₓB₃₀ (x = 0, 2.5, 4, 6, and 8) glassy ribbons measured in a reflection mode using Co Kα radiation.](image)

![FIG. 2. The DSC plots of the CoₓFe₂₂Taₓ₈₋ₓ₂YₓB₃₀ (x = 0, 2.5, 4, 6, and 8) glassy ribbons measured at a constant heating rate of 20 K/min.](image)
amount of the (Co,Fe)B<sub>2</sub> phase forms near the end of the first crystallization event. The (Co,Fe)<sub>21</sub>Ta<sub>2</sub>B<sub>6</sub> phase has the Cr<sub>23</sub>C<sub>6</sub> prototype structure with the Fm3m space group and a large lattice parameter of 1.055 nm, which contains 116 atoms in each unit cell. By introduction of 2.5 at. % Y, the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase appears after heating above the first crystallization event. The participation of Y in the complex boride phase can be found from a small shifting in the position of the (Co,Fe)<sub>21</sub>Ta<sub>2</sub>B<sub>6</sub> peaks to the lower angles (see Fig. 3) owing to the increased lattice parameter of this phase. In the Cr<sub>23</sub>C<sub>6</sub> prototype structure, it has been shown that the transition or rare-earth elements with large atomic radii such as Ta and Y prefer to occupy the 8c sites, while Fe/Co atoms occupy 4a, 32f, and 48h sites. Adding more Y (x > 2.5) is responsible for formation of the Co<sub>1</sub>Y<sub>2</sub>B<sub>2</sub> compound besides the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase and the fraction of the former phase enhances with increasing Y (see Fig. 3). For the alloy with x = 6, a new phase corresponding to the z-(Fe,Co) solid solution precipitates. For x = 8, the amorphous phase transforms in an eutectic mode to a mixture of Co<sub>1</sub>Y<sub>2</sub>, (Fe,Co) and tetragonal (Co,Fe)<sub>2</sub>B phases after crystallization. According to Fig. 3, it is obvious that the fraction of the complex (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase reduces with increasing Y and the Ta free composition does not show any boride compound with the Cr<sub>23</sub>C<sub>6</sub> structure. It has been suggested that the presence of some elements such as Nb or Ta in Fe-based or Co-based glasses plays a crucial role in formation of borides with the Cr<sub>23</sub>C<sub>6</sub> structure.

Formation of the complex FCC (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase upon primary crystallization is correlated with high thermal stability in the present glassy alloys. Due to a complex crystal structure and a large number of atoms in the unit cell of the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase, its precipitation requires long-range rearrangements of the constituent elements especially when there is a significant difference between the atomic structure of the glassy phase and the precipitating crystals. It is known that the metal-metalloid type glassy alloys usually have a network-like short-range order (SRO), which is fundamentally different with respect to the crystals with Cr<sub>23</sub>C<sub>6</sub> structure.

Since Y has a largest atomic radii (180 pm) among the constituents in this alloying system, small addition of Y can give rise to a more consecutive change in the atomic size in the order of Y > Ta > Co > Fe > B. Furthermore, Y has a large negative heat of mixing with B (−50 kJ/mol) and Co (−22 kJ/mol) and small mixing enthalpy with Fe (−1 kJ/mol). Hence, a minor addition of Y results in formation of an amorphous phase with a larger thermal stability due to a high packing density and new local atomic configurations with strong interactions, which can dramatically suppress the diffusivity of the elements. Poon et al. have suggested that the large (L) and small (S) atoms can form strong L-S percolating network or reinforced "backbone" in the amorphous structure.

In the current alloying system, due to a large negative heat of mixing between the large Y or Ta and small B atoms, strong Y-B and Ta-B percolating networks can be created. It has been shown that in the (Co,Fe)<sub>21</sub>Ta<sub>2</sub>B<sub>6</sub> structure, the Ta atoms located in 8c sites have the largest interatomic spacing with B atoms (0.3737 nm), which occupy the 24 e site and consequently the Ta-B bond is absent. As Y atoms are also located in the 8c site, the absence of the Y-B bond is expectable in the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase. Hence, formation of the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase upon devitrification needs breaking strong Y-B and Ta-B pairs in the glassy structure before long-range diffusion of Y and Ta atoms with small diffusivity, which contributes to a large thermal stability of the present glassy alloys.

According to Table I, excessive addition of Y (x > 2.5) decreases the thermal stability of the ribbons probably due to decreased fraction of the (Co,Fe)<sub>21</sub>(Ta,Y)<sub>2</sub>B<sub>6</sub> phase and formation of less complex phases such as Co<sub>1</sub>Y<sub>2</sub>B<sub>2</sub>, (Co,Fe) and (Co,Fe)<sub>2</sub>B, which can be precipitated easier. The decrease in \( T_g \), upon increasing the Y fraction (see Table I) can be correlated with the changes occurred in the bonding nature and atomic interactions. The mixing enthalpies between Ta and Co, Fe, and B are −24, −15, and −54 kJ/mol, respectively. These values are larger than the mentioned mixing enthalpies between Y and other constituents in the present alloying system. This means that a large Y addition results in formation of an amorphous structure with weaker atomic interactions and consequently there is a lower activation barrier against configuration changes of the glass to occur, causing a decrease in \( T_g \).
B. Atomic structure

Figure 4 shows the total structure factors $S(q)$ of the as-quenched Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ glassy ribbons. It can be seen that all patterns indicate prominent oscillations up to $q \approx 160$ nm$^{-1}$. Furthermore, addition of Y affects the shape and position of the diffuse peaks in the entire $q$ range, indicating the changes in atomic structure of the glassy phase. According to Eq. (1), the second diffuse peak of $S(q)$ for the Y free alloy has a strong shoulder around $q = 60$ nm$^{-1}$ and the intensity of this shoulder decreases progressively by addition of Y. Moreover, the Ta free alloy exhibits a shoulder on the third peak of the $S(q)$ around $q = 70$ nm$^{-1}$, which is not appearing for the Ta bearing alloys. The vanishing of the shoulder of the second $S(q)$ peak was observed by partial replacement of Fe with lanthanide elements such as Sm, Tb, and Dy in the Fe$_70$Co$_{10}$B$_{20}$ amorphous alloy. It is known that the presence of shoulder on the second diffuse peak of $S(q)$ is an evidence for a certain structural ordering in the amorphous phase. The characteristic shoulder can be absent even for non-magnetic amorphous alloys like Pt-Ni-P, Pt-P, and La-Al-Ni. Hence, the partial replacement of Ta with Y may increase the extent of the disorder in the Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ glassy ribbons.

Besides the shape, addition of Y can affect the position of the diffraction maxima. Figure 5 shows the variation in positions of the first diffuse maxima ($q_1$) and the second one ($q_2$) of the structure factors $S(q)$ as a function of Y content. Those positions were determined by fitting the top of the $S(q)$ peaks to the pseudo-Voigt function. It is observed that $q_1$ decreases continuously by introduction of Y, however, $q_2$ indicates a reverse behavior and it increases upon Y addition. As mentioned above, addition of Y affects the local ordering of the amorphous phase. It is known that the diffuse maxima of $S(q)$ with large $q$ values determine the SRO constructed by solute-centered quasi-equivalent clusters. On the other hands, the low-$q$ part of $S(q)$ characterizes the packing and connection of the quasi-equivalent clusters or the medium-range order (MRO). As inferred from Fig. 4, the observed changes in position and shape of the diffuse maxima particularly for the first and second peaks, can manifest that changes occurred in the MRO.

The structural evolution of the Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ glassy ribbons can be further investigated in the real-space by calculating the reduced pair correlation functions $G(r)$ according to Eq. (1). Figure 6 plots the $G(r)$ of the Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ glassy ribbons. It is obvious that the Y addition induces significant changes in shape and intensity of the $G(r)$ peaks in the short-range and medium-range atomic correlations. As the figure shows, the Y free alloy indicates pronounced oscillations up to $r \approx 1.6$ nm, demonstrating a high degree of MRO. However, addition of Y decreases noticeably the intensity of $G(r)$ peaks in the medium-range scale, which results in a faster decay of oscillations and decreasing the corresponding correlation length (Fig. 7). In other words, the degree of the MRO decreases in the Y bearing glassy alloys, in a good agreement with the reciprocal-space results (Fig. 4). Furthermore, the splitting of the second $G(r)$ peak can be observed in each composition. It is obvious that the intensity of the right shoulder of the second $G(r)$ increases by introduction of Y. This effect implies that the atoms prefer to place at larger distances in the second near-neighbor shell, probably because of large atomic radii of Y compared to other elements.

![Image 4](https://example.com/image4.png)

**FIG. 4.** The XRD total structure factors $S(q)$ of the Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (x = 0, 2.5, 4, 6, and 8) glassy ribbons.

![Image 6](https://example.com/image6.png)

**FIG. 6.** The reduced pair correlation functions $G(r)$ of the Co$_{40}$Fe$_{22}$Ta$_{8-x}$Y$_x$B$_{30}$ (x = 0, 2.5, 4, 6, and 8) glassy ribbons.
A minor introduction of the In the current alloying system, With decreasing depicts the evolution of The type of such polyhedrons. In addition, the inter-atomic bond lengths, calculated from sum of the Goldschmidt’s radii, are presented. All the plots indicate a pre-peak at \( r \approx 0.21 \text{ nm} \), due to formation of \((\text{Co,Fe})-\text{B}\) pairs in the glassy structure. It should be emphasized that the contributions of Fe and Co atoms are indistinguishable owing to their similar atomic radii. According to the type and concentration of the constituent elements, it can be inferred that the maximum of the first \(G(r)\) peak in all compositions dominantly consists of \((\text{Co,Fe})-(\text{Co,Fe})\) pairs. In the 5 component alloys containing both Y and Ta, there are 15 overlapping atomic pairs in the first coordination shell. It is worth to note that with only one total atomic pair density distribution function, it is not possible to distinguish all of partial atomic pair correlations. The overlapping of the atomic pairs results in a non-asymmetric first \(G(r)\) peak and appearance of a strong right-hand shoulder, particularly for Ta free alloy, as can be seen in Fig. 8. Besides the significant changes in the shape of the main \(G(r)\) peak, addition of Y changes the position of its maximum. According to Fig. 9, the position of the maximum of the first \(G(r)\) peak decreases almost linearly with increasing the Y fraction. This may imply that the \((\text{Co,Fe})-(\text{Co,Fe})\) pair separation decreases by Y addition and it approaches the theoretical \((\text{Co,Fe})-(\text{Co,Fe})\) bond length (0.25 \text{ nm}), obtained as a sum of atomic radii. It is worth to note that a similar contraction of the dominant inter-atomic distance happens for Fe-Fe atomic pairs in the Fe-\(30\)Co-\(10\)B\(20\) glasses upon partial replacement of the Fe atoms with some lanthanide elements.

It is known that the building blocks or SRO in MGs can be identified as the Kasper-type (quasi-equivalent) solute-centered coordination polyhedra. The type of such polyhedral is controlled by the effective atomic size ratio between the solute and solvent atoms, \(R^*\). With decreasing \(R^*\), the preferred polyhedral type changes from the Frank–Kasper type (for \(R^* > 1.2\)) to the icosahedral type (for \(R^* < 0.902\)), then to the bi-capped square Archimedean antiprism (BSAP) type (for \(R^* < 0.835\)) and finally to the tri-capped trigonal prism (TPP) type (for \(R^* < 0.732\)). In the current alloying system, the \(R^*\) between B as the main solute atom and Co as the solvent atom is 0.69. Hence, it is anticipated that the chemical SRO in the Co-\(40\)Fe-\(22\)Ta-\(8-x\)Y-\(x\)B\(30\) glassy ribbons is probably made of B-centered TPP polyhedra. It has been shown that B-centered-TPP polyhedra with a Voronoi index of \(\{0,3,6,0\}\) is the dominant polyhedra in the SRO of the Co-\(31\)Fe-\(26\)Ta-\(35.5\)B\(31.5\) MG, which has a similar composition to the glassy alloys in this work. The TPP polyhedra in metal-metalloid type MGs are the ideal blocks for constructing the network-like MRO. Formation of the network-like structure can be correlated not only to the necessity of dense packing but also to the chemical interactions and the directionality of the covalent bonds in metal-metalloid pairs.

For the Co-\(43\)Fe-\(26\)Ta-\(35.5\)B\(31.5\) MG, it has been shown that the B-centered network-like MRO can be built through the connection of the TPP polyhedra via vertex-, edge-, face-, and intercross-shared atoms and all four kinds of component elements can serve as glue atoms. A minor introduction of the glue elements with very large atomic radii like Y can distort

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**FIG. 7.** Evolution of the correlation length for the Co-\(40\)Fe-\(22\)Ta-\(8-x\)Y-\(x\)B\(30\) (\(x = 0, 2.5, 4, 6, \text{ and } 8\)) glassy ribbons with respect to Y fraction.

**FIG. 8.** The reduced pair correlation functions \(G(r)\) of the Co-\(40\)Fe-\(22\)Ta-\(8-x\)Y-\(x\)B\(30\) (\(x = 0, 2.5, 4, 6, \text{ and } 8\)) glassy ribbons depicted in the range of the first coordination shell.

**FIG. 9.** Variations in position for the first peak of the pair correlation functions \(G(r)\) of the Co-\(40\)Fe-\(22\)Ta-\(8-x\)Y-\(x\)B\(30\) (\(x = 0, 2.5, 4, 6, \text{ and } 8\)) glassy ribbons as a function of Y fraction.
the TTP polyhedra and slightly change their arrangement. On the other hand, the excessive addition of Y could introduce large internal strains and hence break the network-like structure and consequently decrease the degree of such MRO (see Figs. 6 and 7). A continuous decrease of thermal stability upon addition of Y beyond 2.5 at. % (Table I) could be attributed to decrease in the extent of the network-like MRO, which has a different atomic configuration compared to the precipitating (Co,Fe)$_{21}$(Ta,Y)$_2$B$_6$ crystal.

C. Magnetic properties

Figure 10 shows the evolution of the saturation magnetization (normalized) as a function of temperature, measured at a constant heating rate of 10 K/min. As the figure shows, the magnetization drops for each sample and passes through an inflection point attributed to the Curie temperature $T_c$ of the glassy phase and then it becomes spontaneously zero. According to this model, the VBS for the transition metals with strong perturbing potential (with a valence difference $\Delta Z > 2$) could move above the Fermi energy. Hence, it is anticipated that the charge transferred to the d bands of Fe or Co atoms may decrease by introduction of Y atoms and consequently the ferromagnetic atoms have larger magnetic moments in compositions with larger Y content. The suppression of the magnetic moment of the Co atoms in the Co-based alloys after alloying with non-ferromagnetic transition metals has been described by Friedel’s virtual-bound-state (VBS) model. According to this model, the VBS for the transition metals with strong perturbing potential (with a valence difference $\Delta Z > 2$) could move above the Fermi energy. Consequently, the magnetic moment of Co atoms decreases due to the charge transfer from the VBS of the non-magnetic transition metals (like Ta and Y) into the d-band of Co based on the following equation:

$$\frac{d\mu_{Co}}{dC} = -(10 + \Delta Z)\mu_B,$$  \hspace{1cm} (2)
coefficient constant, which is in the order of $\lambda \approx 10^{-7}$ ppm for the Co-based glassy alloys.\footnote{1}

According to above results, alloy with $x = 2.5$ due to a large SLR and very good soft magnetic properties can be promising for the spark plasma sintering (SPS) or hot pressing processes, in order to produce the bulk glassy alloys with large size and complex shape by consolidation of the glassy powders in the temperature range of SLR, where the viscosity drops significantly.\footnote{1}

IV. CONCLUSIONS

The influence of Y addition on atomic structure, thermal stability, and magnetic properties of the Co$_{90}$Fe$_{22}$Ta$_8$Y$_3$B$_{30}$ MGs was investigated. The results indicated that the addition of 2.5 at.\% Y noticeably improves the thermal stability through increasing the extension of the SLR, while the excessive alloying by Y deteriorates the thermal stability. Moreover, addition of Y decreases the degree of the network-like MRO probably due to the introduction of large inter-atomic strains. The decrease in thermal stability after extra addition of Y was attributed to the decrease in the extent of network-like MRO and fraction of the (Co,Fe)$_2$Ta$_3$Y$_2$B$_8$ phase appeared upon primary crystallization. It was shown that the (Co,Fe)-(Co,Fe) pairs have shorter inter-atomic distance in the Y bearing alloys. This gives rise to stronger interactions between the ferromagnetic elements, which results in improving the saturation magnetization and Curie temperature.

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