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# Structure, magnetocaloric effect and critical behaviour in $\text{Ni}_{50}\text{Mn}_{30}(\text{Sn},\text{In})_{20}$ Heusler alloys

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## Abstract

A detailed investigation of structure, critical behaviour and magnetocaloric properties of  $\text{Ni}_{50}\text{Mn}_{30}\text{Sn}_{20}$  (Sn20) and  $\text{Ni}_{50}\text{Mn}_{30}\text{In}_{20}$  (In20) alloys has been investigated by means of X-rays diffraction and magnetic measurements.  $\text{Ni}_{50}\text{Mn}_{30}\text{Sn}_{20}$  alloy shows a cubic austenite  $L2_1$  structure, and undergoes a second order magnetic transition at a Curie temperature of  $T_{c,1}^A(\text{Sn}20) = 333 \text{ K}$ . However, the  $\text{Ni}_{50}\text{Mn}_{30}\text{In}_{20}$  alloy exhibits a mixture of cubic  $L2_1$  and  $B2$  austenite structures having Curie temperatures of  $T_{c,2}^A(\text{In}20) = 285 \text{ K}$  and  $T_c^*(\text{In}20) = 330 \text{ K}$ , respectively. The modified Arrott plots, Kouvel-Fisher curves and critical isotherm analysis have been used to estimate the critical exponents ( $\beta$ ,  $\gamma$ , and  $\delta$ ) around the Curie temperature. For Sn20 alloy, the reliable exponents are consistent with the mean field model, revealing long-range ferromagnetic interactions. Nevertheless, the critical exponents of In20 alloy around 330 K cannot be arranged into any of the universality classes of well-known classical standard models. The maximum entropy change under 5 T of Sn20 ( $\Delta S_M^{max} = 2.43 \frac{\text{J}}{\text{kg}} \cdot \text{K}$ ) is slightly higher than that of In20 ( $\Delta S_M^{max} = 2.05 \frac{\text{J}}{\text{kg}} \cdot \text{K}$ ). The experimental results of entropy changes are in good agreement with those calculated using Landau theory.

**Keywords:** NiMn-based alloys; Magnetic transition; Critical behavior; Magnetocaloric effect; Landau theory.

# 1. Introduction

Heusler alloys have attracted considerable attention due to their several technological applications such as sensors, spintronic devices and magnetic refrigeration [1-3]. Several experimental [4, 5] and theoretical [6, 7] studies have been carried out to investigate the properties of Heusler alloys. Off-stoichiometric NiMn-based ferromagnetic alloys exhibit an exchange bias phenomena (*EB*) [8], magnetoresistance [9, 10], magnetocaloric effect (*MCE*) [11, 12], magnetic super-elasticity [13] and piezoresistance [14]. It has been reported that  $\text{Ni}_{50}\text{Mn}_{50-x}\text{Sn}_x$  series crystallize in a cubic austenite  $L2_1$  structure, space group  $Fm\bar{3}m$ , for  $x \geq 15$ , and a martensite structure for  $x < 15$  [15, 16]. An extended annealing of  $\text{Ni}_{50}\text{Mn}_{39}\text{Sn}_{11}$  and  $\text{Ni}_{50}\text{Mn}_{37}\text{Sn}_{13}$  alloys at 773 K, involves a phase decomposition of the eutectoid-type structure into  $\text{Ni}_{54}\text{Mn}_{45}\text{Sn}_1$  and  $\text{Ni}_{50}\text{Mn}_{30}\text{Sn}_{20}$ . Wherein, the Curie temperature was about 350 K [17]. The standard enthalpy of formation,  $\Delta_f H^\circ$ , of  $\text{Ni}_{50.6}\text{Mn}_{29.6}\text{Sn}_{19.8}$  alloy was  $-21.0 \text{ kJ/mol of atom}$  [18]. A linear shift of the reverse and forward martensitic transition temperatures in  $\text{Ni}_{46.4}\text{Mn}_{32.8}\text{Sn}_{20.8}$  alloy occurs by high magnetic field application up to 14 T [19]. Moreover, a spin glass behaviour was found in  $\text{Ni}_{50}\text{Mn}_{30}\text{Sn}_{20}$  alloy [20]. For Ni-Mn-In alloys, the structural transition was observed in Mn-rich composition where both the austenite and martensite phases exhibit a ferromagnetic order [21, 22]. Krenke *et al.* reported that  $\text{Ni}_{50}\text{Mn}_{50-x}\text{In}_x$  alloys with  $0.16 \geq x \geq 0.05$  show a first order structural transformation, and the Curie temperature decreases from 310 K for  $x = 20$  to 290 K for  $x = 25$  [23]. Furthermore, the martensitic transition temperature *MT* is influenced by change in elemental composition through the change of valence electron concentration per atom (*e/a*).

Depending on the nature of the magnetic transition, the magnetocaloric effect can be categorised. For the second order magnetic transition, the resulted negative entropy change around the Curie temperature of the austenite structure,  $T_c^A$ , produces a direct *MCE*. However, an inverse magnetocaloric effect, *IMCE*, is engendered with a positive value of the entropy change around the martensitic transformation, *MT*. The existence of a universal behavior in materials can be investigated through the critical exponents ( $\beta$ ,  $\gamma$  and  $\delta$ ) associated with the phase transitions. By using several models [24] such as the mean field model associated with long-range mean field theory where  $\beta = 0.5$  and  $\gamma = 1$ , short-range interactions as specified by the Heisenberg model with  $\beta = 0.365$  and  $\gamma = 1.336$ , 3D-Ising model with  $\beta = 0.325$  and  $\gamma = 1.241$ , and tricritical mean field model with  $\beta = 0.25$  and  $\gamma = 1$ , the critical exponents  $\beta$ ,  $\gamma$  and  $\delta$  can be estimated from the isothermal magnetization curves,  $M(H)$ . The exponent  $\beta$ , which is linked to the evolution of the spontaneous magnetization with respect to

1 the temperature ( $M_s \propto (T - T_c)^\beta$ ), describes the growth of the ordered moment below  $T_c$ ;  $\gamma$  is  
2 related to the temperature dependence of the initial magnetic susceptibility ( $\chi_0^{-1} \propto (T -$   
3  $T_c)^\gamma$ ). It defines the divergence of  $\chi_0$  at  $T_c$ , and  $\delta$  is associated to the critical isothermal  
4 magnetization. It describes the curvature of the isothermal magnetization curves  $M(H)$  at  $T_c$ .  
5 The mechanisms of magnetic phase transitions are to some extent vague due to the influence  
6 of many parameters such as the preparation method, the alloy composition, the thermal  
7 history, etc. Consequently, the aim of the present work is to investigate the magnetocaloric  
8 effect and the critical behaviour near the Curie temperature in the melt spun  $\text{Ni}_{50}\text{Mn}_{30}(\text{Sn},$   
9  $\text{In})_{20}$  ribbons.  
10

## 11 2. Experimental details

12 Polycrystalline alloys with nominal compositions of  $\text{Ni}_{50}\text{Mn}_{30}\text{Sn}_{20}$  and  $\text{Ni}_{50}\text{Mn}_{30}\text{In}_{20}$ ,  
13 henceforth called Sn20 and In2, respectively, have been prepared by arc melting high purity  
14 elements Ni, Mn, Sn and In (99.9 %) under an argon atmosphere. The ingots were re-melted  
15 five times to ensure homogeneity. Sn20 and In20 alloys ribbons have been elaborated by melt  
16 spinning onto a surface of copper wheel rotating at a linear speed of 40 m/s under argon  
17 atmosphere. The crystal structure was checked by X-ray diffraction (XRD) on a Brüker D8  
18 Advance diffractometer using Cu-K $\alpha$  radiation ( $\lambda_{\text{Cu}} = 0.154056$  nm) in a  $\theta - 2\theta$  Bragg  
19 Brentano geometry. The XRD patterns were computer refined using the MAUD program  
20 which is based on the Rietveld method [25]. Magnetization measurements were carried out in  
21 a superconducting quantum interference device (SQUID) magnetometer and a BS2  
22 magnetometer developed at the Néel Institute under an applied magnetic field of 0.05 T. The  
23 critical exponents and the magnetic entropy changes around the Curie temperature have been  
24 determined from the isothermal magnetization measurements as a function of the applied  
25 magnetic field.  
26

## 27 3. Results and discussion

28 **Figure 1** and **Figure 2** show the Rietveld refinement of the XRD patterns of Sn20 and  
29 In20 alloys, respectively. The XRD patterns exhibit thin and intense peaks characteristic of a  
30 fully crystalline microstructure due to the fast crystallization and growth kinetics of the  
31 samples during the melt spinning process. One observes also that the peak intensities of Sn20  
32 are slightly high than those of In20. The high order of  $L2_1$  austenite in both alloys is  
33 confirmed by the superlattice (111), (311) and (311) reflections. In addition, Sn20 is a  
34 textured alloy as shown by (311) reflection intensity. Such preferential orientation might be  
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1 due to fact that during the rapid solidification, the heat removal induces directional growth of  
 2 the crystalline phase. The importance of (111) and (311) peaks intensity compared to those  
 3 obtained in the Ni<sub>50</sub>Mn<sub>30</sub>Sn<sub>20</sub> Heusler alloy prepared by arc melting followed by annealing at  
 4 1000°C for 24 h [20], confirms the effect of the preparation method on the structural  
 5 properties. Usually the non-stoichiometry of the alloy with a low content of Z element gives  
 6 rise to weak minor peaks of the superstructure in Heusler alloys. While the Sn20 alloy show a  
 7 single austenite  $L2_1$  structure (lattice parameter  $a=0.5994(3)$  nm, space group  $Fm\bar{3}m$ ), the  
 8 In20 alloy exhibit a mixture of an ordered  $L2_1$  phase, (volume fraction ~85 %, and lattice  
 9 parameter  $a = 0.6088(3)$  nm), and a partially ordered  $B2$  phase (space group  $Pm\bar{3}m$ ,  
 10 volume fraction ~15 % and lattice parameter  $a = 0.3011(2)$  nm). The formation of  $B2$   
 11 structure can be linked to the degree of atomic order.

12 The temperature dependence of the magnetization,  $M(T)$ , recorded at 0.05 T for both  
 13 Sn20 and In20 alloys, is displayed in Fig. 3. The Curie temperature was inferred from the  
 14 minimum in  $dM/dT$  versus temperature curve as shown by the inset in Fig. 3. For the Sn20  
 15 alloy, the ferromagnetic to paramagnetic transition  $T_c^A(Sn20)$  occurs at 333 K, whereas the  
 16 In20 alloy displays two magnetic transition temperatures at 285 K and 330 K that can be  
 17 attributed to  $L2_1$  and  $B2$  structures, respectively. This result agrees well with the XRD one  
 18 and confirms thus the biphasic character of In20 alloy. The magnetic transitions of both  
 19 phases are different from those reported in the literature [26-28]. Those discrepancies might  
 20 be ascribed to the composition change and/or the preparation conditions. For example, the  
 21 obtained lower Curie temperature ( $T_c \sim 222$  K) for the austenite phase by R.C. Bhatt et al.  
 22 [20], has been attributed to deficiency of Sn atoms in the system.

23 The isothermal magnetization curves of In20 and Sn20 alloys,  $M(H)$ , around the Curie  
 24 temperature are presented in Fig. 4 as a function of the applied magnetic field up to 6 T. For  
 25 both alloys, the magnetization does not saturate even at 5 T. In the magnetic system, the  $MCE$   
 26 is related to the magnetic entropy change,  $\Delta S_M$  which might be induced by an applied  
 27 magnetic field that leads to the change in the magnetic order. The magnetic entropy change  
 28  $\Delta S_M$  can be deduced from the isothermal magnetization,  $M(H)$ , curves via the following  
 29 Maxwell relation:

$$30 \Delta S_M(T, \Delta H) = S_M(T, H_2) - S_M(T, H_1) = \int_{H_1}^{H_2} \left( \frac{\partial M}{\partial T} \right)_T dH \quad (1)$$

31 Where  $H_1$  and  $H_2$  are the applied magnetic fields with  $H_1 < H_2$ , and  $\Delta H = H_2 - H_1$ .  $\Delta S_M$  can  
 32 be calculated by integrating the  $M(H)$  curves at small discrete magnetic field and temperature  
 33 intervals, and can be approximated by the succeeding equation:

$$\Delta S_M(T, M) = \sum_i \frac{M_{i+1}(T_{i+1}, H_{i+1}) - M_i(T_i, H_i)}{T_{i+1} - T_i} \Delta H \quad (2)$$

Where  $M_i$  and  $M_{i+1}$  are the experimental data of the magnetization at  $T_i$  and  $T_{i+1}$ , respectively, under the magnetic field  $H_i$ . The variation of  $-\Delta S_M$  as a function of the temperature is shown in Fig. 5. The maximum magnetic entropy change,  $-\Delta S_M^{max}$ , increases with increasing the applied magnetic field and reaches 2.43 J/kg.K for Sn20, and 2.05 J/kg.K for In20 (see the inset in Fig. 5). One notes that  $-\Delta S_M^{max}$  value of Sn20 alloy is  $\sim 16\%$  higher than that obtained in Ni<sub>50</sub>Mn<sub>35</sub>Sn<sub>15</sub> alloy prepared in the same conditions [29].

The relative cooling power,  $RCP$ , associated with the magnetic entropy change is an important feature to evaluate the magnetocaloric efficacy of materials. The relative cooling power is given by the following equation:

$$RCP = - \int_{T_1}^{T_2} \Delta S_M(T) dT \quad (3)$$

Where  $T_1$  and  $T_2$  temperatures are defined by the full width at half maximum of  $\Delta S_M(T)$  peak. For Sn20 and In20 ribbons, the calculated  $RCP$  values are of about 172.88 J/kg and 240 J/kg, respectively, for an applied magnetic field of 5 T. The  $RCP$  value of Sn20 is much higher than that obtained for the same composition by Bhatt et al. ( $RCP = 54.6 \text{ J/kg}$ ) under a magnetic field of 3 T [20].

Amaral *et al.* proposed a theoretical model based on the contribution of the magneto-elastic interactions of electrons [30] and on Landau theory in the critical region [31]. The free energy ( $G$ ) is developed as a function of the magnetization ( $M$ ):

$$G(M, T) = G_0 + \frac{a(T)}{2} M^2 + \frac{b(T)}{4} M^4 + \frac{c(T)}{6} M^6 + \dots - HM \quad (4)$$

For the equilibrium condition,  $\partial G / \partial M = 0$ , the total magnetization in the critical region is given by the following relationship:

$$H = a(T)M + b(T)M^3 + c(T)M^5 \quad (5)$$

The Landau's coefficient  $a(T)$ ,  $b(T)$  and  $c(T)$  are determined from the polynomial fit of the experimental  $M(H)$  data according to the Eq.5. The variation of  $a(T)$  shows a minimum around  $T_c^A$ , and  $b(T_c^A) = 0$  proves the second order character of the magnetic transition. The derivative of the free energy with respect to the temperature gives a theoretical model of the magnetic entropy change:

$$-\Delta S_M(T, H) = \left( \frac{dG}{dT} \right)_H = \frac{a'(T)}{2} M^2 + \frac{b'(T)}{4} M^4 + \frac{c'(T)}{6} M^6 \quad (6)$$

where  $a'(T)$ ,  $b'(T)$  and  $c'(T)$  are the derivative of Landau's coefficients with respect to the temperature. The calculated  $-\Delta S_M$  curves are in a good agreement with the experimental results (Fig. 7). Thus, the calculated Landau's coefficients are accurate.



In order to understand the interaction type of the magnetic moments and the universality class of Sn20 and In20 ribbons, the critical exponents have been analysed through different techniques such as the modified Arrott plots (*MAP*) [32], Kouvel-Fisher plots (*K – F*) [33, 34] and critical isotherm (*CI*). Hence, the critical exponents  $\beta$ ,  $\gamma$  and  $\delta$  can be determined using Equations 7-9:

$$M_S(T) = M_0(-\varepsilon)^\beta; \quad \varepsilon < 0, T < T_c \quad (7)$$

$$\chi_0^{-1}(T) = \left(\frac{h_0}{M_0}\right) \varepsilon^\gamma; \quad \varepsilon > 0, T > T_c \quad (8)$$

$$M = DH^{1/\delta}; \quad \varepsilon = 0, T = T_c \quad (9)$$

where  $M_S(T)$  and  $\chi_0^{-1}(T)$  are the spontaneous magnetization and the inverse initial magnetic susceptibility, respectively;  $\varepsilon = (T - T_c)/T_c$  is the reduced temperature;  $M_0$ ,  $h_0$ , and  $D$  are the critical amplitudes.

The magnetic interactions that characterize a magnetic system can be described by four distinct conventional models: the mean field model, 3D-Heisenberg model, 3D-Ising model, and tricritical mean field model [24]. The modified Arrott plots around the Curie temperature of  $L2_1$  structure are shown in Figures 8 and 9. The calculated relative slopes, *RS*, identified as the slope at each temperature  $T$  to the slope at the critical temperature ratio, are plotted using the linear fit from the high field region of each curve (Fig. 10). In the ordered  $L2_1$  structures, the mean field model is the best model that describes both In20 and Sn20 ribbons since *RS* are much closer to 1. By fitting the spontaneous magnetization  $M_S(T, 0)$  and the inverse of the initial susceptibility  $\chi_0^{-1}(T, 0)$  curves using equations 7 and 8, respectively, the new values of  $\beta$ ,  $\gamma$  and  $T_c^A$  (table 1) can be determined (Fig. 11). One notes that the  $\beta$  and  $\gamma$  values are close to the mean field model with a long range order of magnetic interactions, and the Curie temperatures are close to those obtained from the  $M(T)$  curves.

The Kouvel-Fisher plots of  $M_S(T)(dM_S(T)/dT)^{-1}$  and  $\chi_0^{-1}(T)(d\chi_0^{-1}(T)/dT)^{-1}$  against the temperature (Fig. 12) show straight lines with slopes  $1/\beta$  and  $1/\gamma$ , respectively. Moreover, the  $\beta$  and  $\gamma$  exponent values are interconnected to the third exponent  $\delta$  via the Widom scaling relation ( $\delta = 1 + (\gamma/\beta)$ ) [35]. The calculated  $\delta$  values (table 1) are comparable to those estimated from the *CI* curves (Fig. 13). The validity of the critical exponents and the Curie temperature values are confirmed by the scaling hypothesis [36] through the following equation:

$$M(H, \varepsilon) = \varepsilon^\beta f_\pm(H/\varepsilon^{\beta+\gamma}) \quad (10)$$



The regular analytic functions  $f_+$  and  $f_-$  are undertaken for  $T > T_c^A$  and  $T < T_c^A$ , respectively. The  $M|\varepsilon|^{-\beta}$  as a function of  $H|\varepsilon|^{-(\beta+\gamma)}$  are plotted in the vicinity of the Curie temperature (Fig. 14). The presence of two distinct branches below and above  $T_c^A$  confirms the fact that the predicted critical exponents and Curie temperatures are reasonably accurate.

The modified Arrott plots of the disordered cubic  $B2$  structure around  $T_c^*(In20) = 330\text{ K}$  show curved lines rather than straight-lines (Fig. 15), and the linear fit at high field region has negative intercept. Accordingly, those models are unable to describe the system. The exponents values were obtained from the optimum fitting of  $M_s(T, 0)$  and  $\chi_0^{-1}(T, 0)$  curves using Eq.(7) and Eq.(8), respectively, which were determined from the high field region of  $M^{1/\beta}$  against  $(\mu_0 H/M)^{1/\gamma}$  plot. This process was repeated several times until the convergence. The nonlinear fit to the  $M_s(T, 0)$  and  $\chi_0^{-1}(T, 0)$  plots gives  $\beta = 0.434 \pm 0.037$ ,  $\gamma = 0.737 \pm 0.490$  and  $T_c^* = (331.328 \pm 0.919)\text{ K}$  values (Fig. 16). As shown in Fig. 17, the modified Arrott plot presents almost parallel isotherms near  $T_c^*$  by using the unconventional exponents. The Kouvel-Fisher plots are presented in Fig. 18, and the convergence was reached with nearby values to those obtained from the modified Arrott plots with  $\beta = 0.466 \pm 0.050$ ,  $\gamma = 0.702 \pm 0.043$  and  $T_c^* = (331.420 \pm 1.142)\text{ K}$ . The validity of those unconventional critical exponents and  $T_c^*$  values were confirmed by the scaling hypothesis as shown in Fig. 19. The third exponent  $\delta$  was determined experimentally from the critical curve  $M = f(H, T_c^*)$  (Fig. 20), and analytically by the Widom scaling relation. The experimental and calculated  $\delta$  values were  $\delta = 2.645 \pm 0.003$  and  $\delta_{cal} = 2.698 \pm 0.453$ , respectively. The critical exponents of Sn20 and In20 alloys as well as those of the different models are reported in Table 1. One observes that  $\beta$  have smaller values indicating a faster growth of the ordered moment. The initial susceptibility has a softer divergence at  $T_c^A(Sn20)$  and  $T_c^A(In20)$ , and sharper divergence at  $T_c^*(In20)$ . The  $\delta$  values reflect the faster saturation of  $M(H)$  curves at  $T_c^A$ , the less curvature and slower saturation at  $T_c^*$ .

#### 4. Conclusion

Structure, magnetocaloric effect and critical behaviour of the  $Ni_{50}Mn_{30}(Sn,In)_{20}$  alloys were studied. Sn20 ribbons exhibit a single cubic  $L2_1$  structure and undergo a second order magnetic transition at 333 K. The presence of  $L2_1$  and  $B2$  structures in the In20 ribbons is confirmed by the existence of two Curie temperatures ( $T_c^A = 285\text{ K}$  and  $T_c^* = 330\text{ K}$ ). Magnetic entropy change,  $RCP$  and critical exponents' values of Sn20 are higher than those of In20. The mean field model, with long-range ferromagnetic interactions, was the best model

1  
2 that describes the magnetic interactions around  $T_c^A$ . However, the critical exponents values  
3 around  $T_c^*$  are different from the well-known universality class.

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## Figure captions

**Figure 1:** Rietveld refinement of the XRD pattern of Sn20 alloy.

**Figure 2:** Rietveld refinement of the XRD pattern of In20 alloy.

**Figure 3:** Temperature dependence of the magnetization measured at 0.05 T. The *insets* display the characteristic magnetic transition.

**Figure 4:** Isothermal magnetization around the Curie temperatures at different temperatures.

**Figure 5:**  $-\Delta S_M(T)$  curves under an applied fields up to 5 T (left), and the corresponding magnetic field dependence of  $-\Delta S_M^{max}$  (right).

**Figure 6:** Superposition of experimental  $-\Delta S_M(T)$  curves and calculated ones under an applied fields up to 5 T, and the corresponding Landau's coefficient  $a(T)$ ,  $b(T)$  and  $c(T)$ , respectively.

**Figure 7:** Modified Arrott plots for Sn20 alloy: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_c^A$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.

**Figure 8:** Modified Arrott plots for In20 alloy: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_c^A$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.

**Figure 9:** Relative slope ( $RS$ ) as a function of temperature ( $RS = S(T)/S(T_c)$ ) deduced from different models.

**Figure 10:** Temperature dependence of spontaneous magnetization  $M_S(T, 0)$  and the inverse of initial susceptibility  $\chi_0^{-1}(T, 0)$  around  $T_c^A$ .

**Figure 11:** Kouvel-Fisher plots for the spontaneous magnetization and the inverse of initial susceptibility around  $T_c^A$ .

**Figure 12:** Critical isotherms with the log-log plot shows in *insets*.

**Figure 13:** Scaling plots indicating universal curves above and below Curie temperatures for Sn20 and In20 alloys.

**Figure 14:** Modified Arrott plots for In20: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_c^*$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.

**Figure 15:** Temperature dependence of spontaneous magnetization  $M_S(T, 0)$  and the inverse of initial susceptibility  $\chi_0^{-1}(T, 0)$  around  $T_c^*$ .

**Figure 16:** Modified Arrott plots: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  with  $\beta = 0.43$  and  $\gamma = 0.73$ .

**Figure 17:** Kouvel-Fisher plots for the spontaneous magnetization and the inverse of initial susceptibility around  $T_c^*$  (*In20*).

**Figure 18:** Scaling plots indicating universal curves above and below  $T_c^*$  (*In20*).

**Figure 19:** Critical isotherm of  $M(\mu_0 H, T_c^*)$  with the log-log plot shown in the *inset*.

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**Table caption**

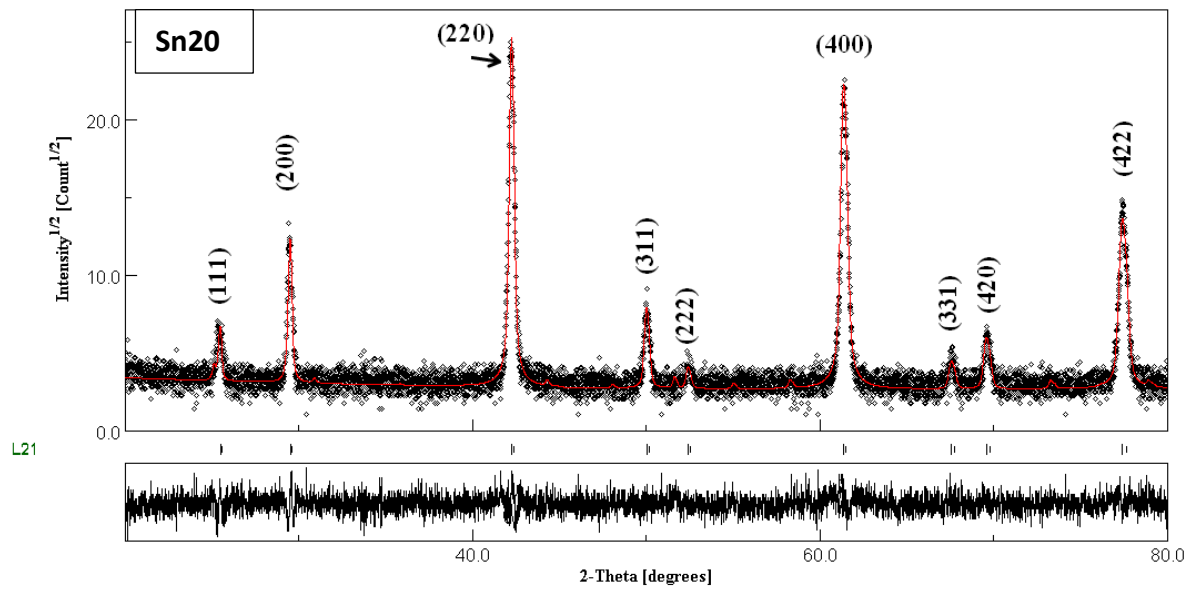
**Table 1:** Critical exponents values of Sn20 and In20 alloys with those of the theoretical models.

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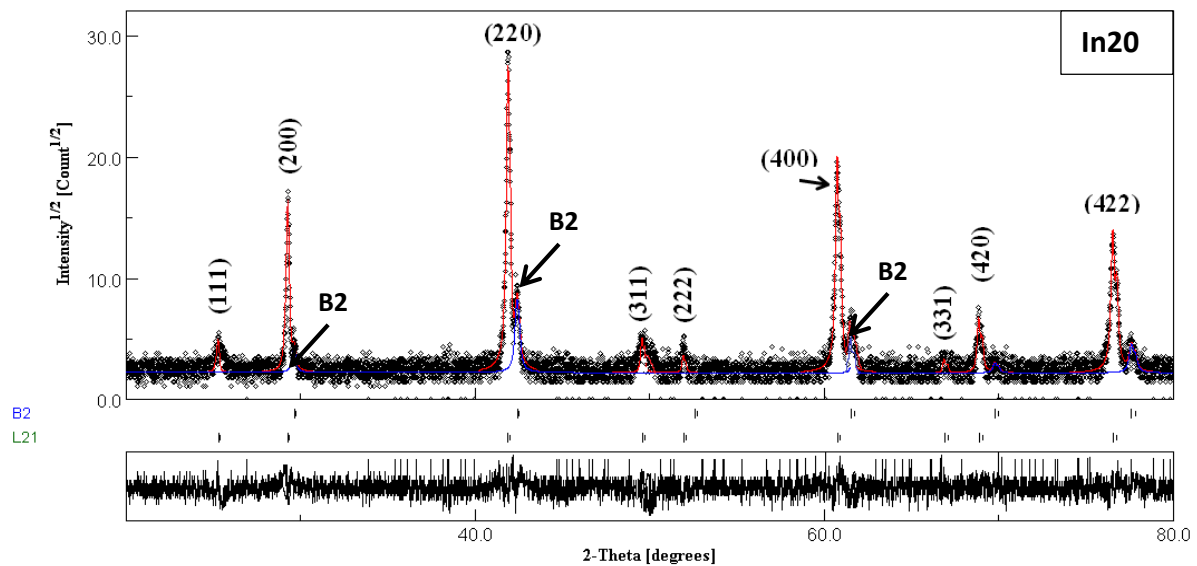


**Table 1**

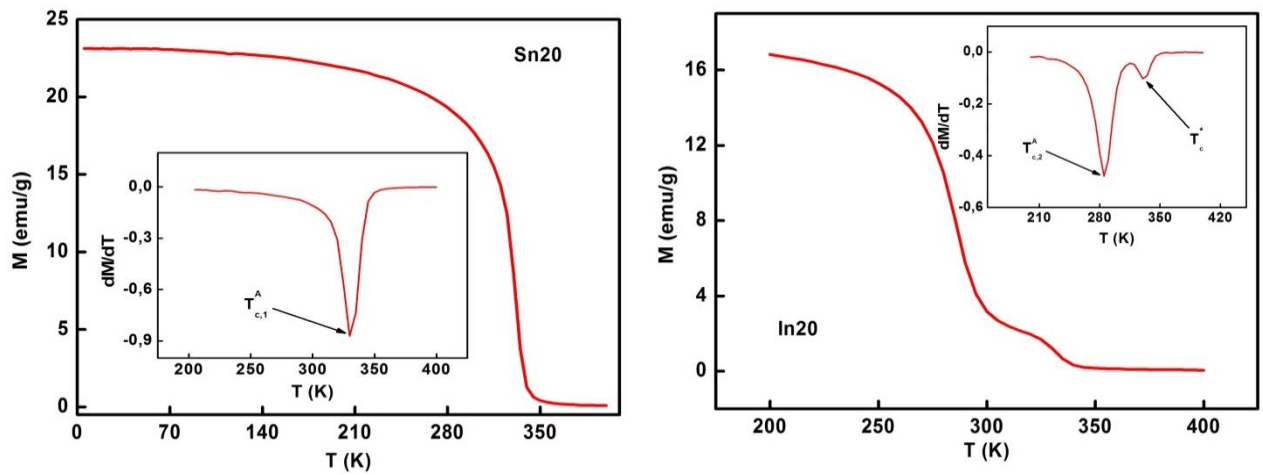
Sample	Technique	$\beta$	$\gamma$	$\delta$	Ref.
Sn20	MAPs	0.421±0.076	1.078±0.142	3.560±0.0661.	<b>This work</b>
	K-F	0.349±0.040	1.171±0.118	-	
	CI			3.710±0.007	
In20	MAP	0.461±0.032		3.537±0.174	<b>This Work</b>
	K-F	0.494±0.045	1.170±0.206	-	
	CI		1.159±0.146	3.439±0.007	
	MAP	0.434±0.037	0.737±0.490	2.698±0.453	
	K-F	0.466±0.050	0.702±0.043	-	
	CI			2.645±0.003	
Mean field model		0.5	1.0	3.0	
3D-Heisenberg model		0.365	1.336	4.80	<b>[24]</b>
3D-Ising model		0.325	1.241	4.82	
Tricritical mean field		0.25	1.0	5.0	



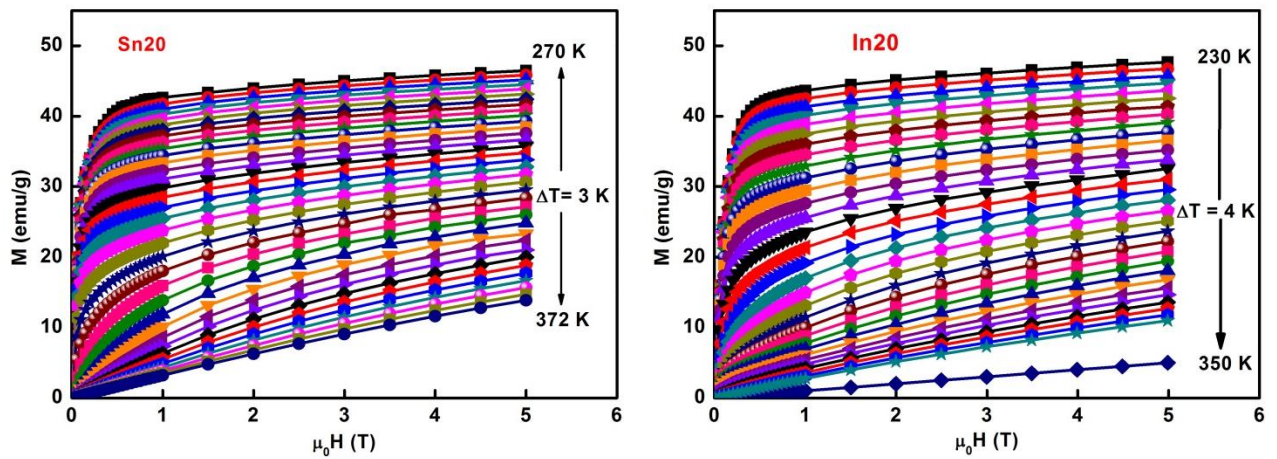
**Fig.1:** Rietveld refinement of the XRD pattern of Sn20 alloy.



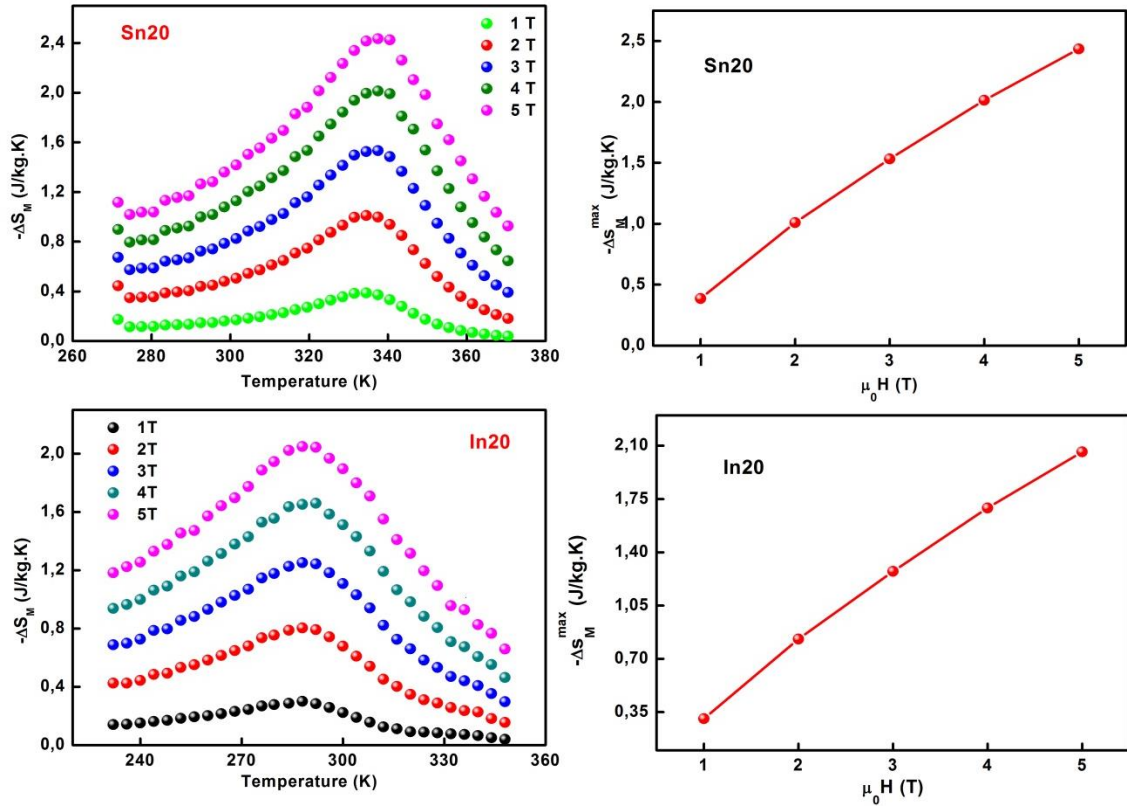
**Fig.2:** Rietveld refinement of the XRD pattern of In20 alloy.



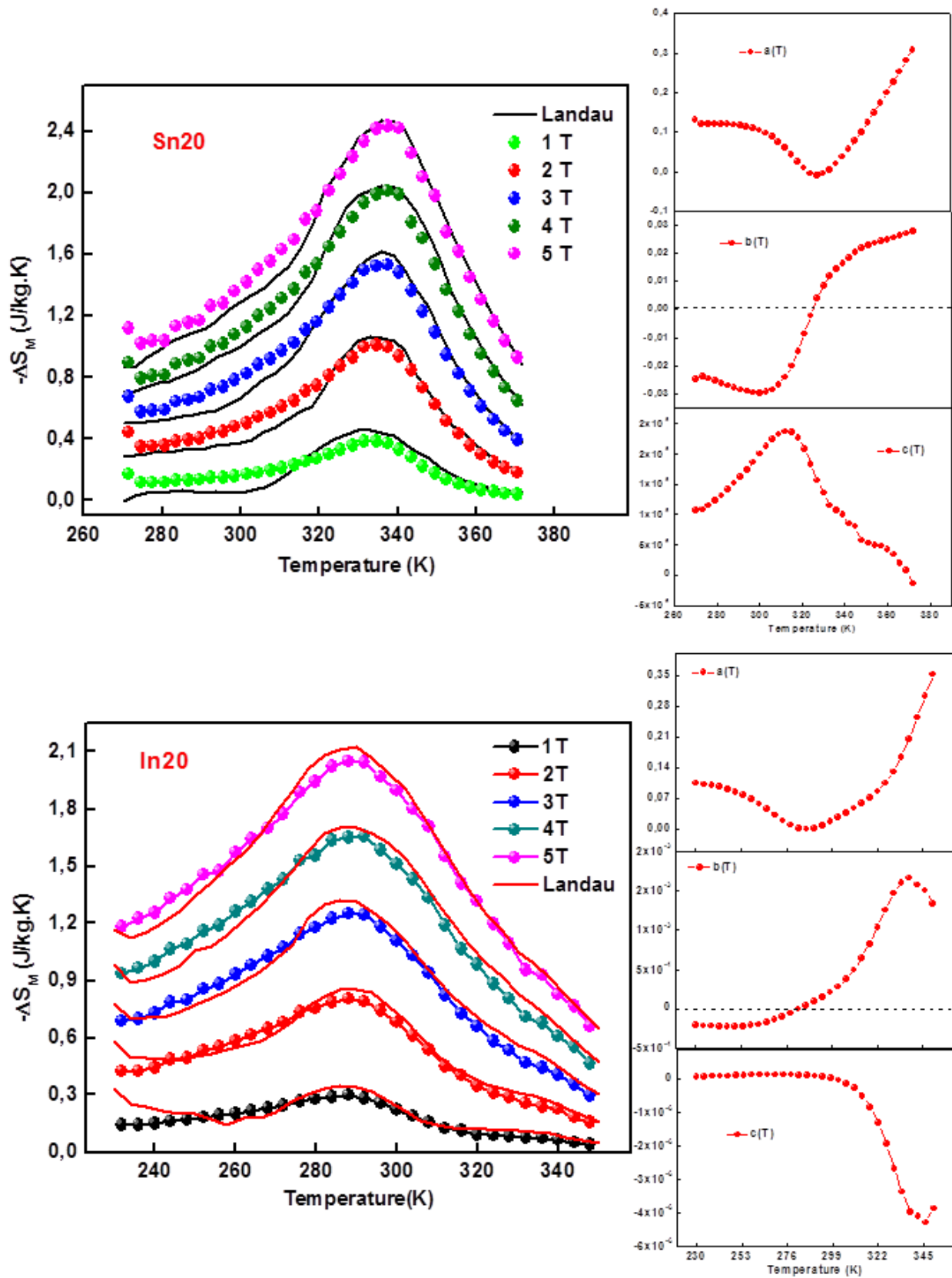
**Fig. 3:** Temperature dependence of the magnetization measured at 0.05 T. The characteristic magnetic transition is shown in the inset.



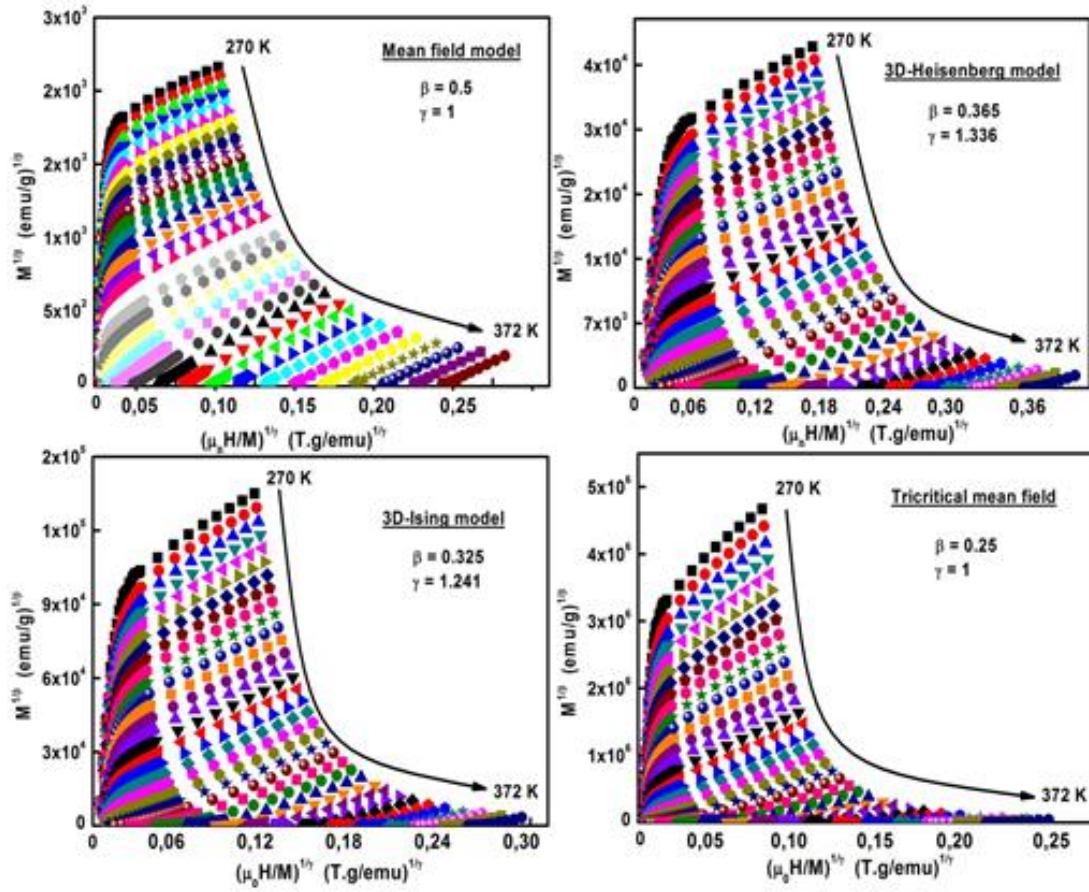
**Fig. 4:** Isothermal magnetization of In20 and Sn 20 alloys, around the Curie temperatures, at different temperatures.



**Fig. 5:**  $-\Delta S_M(T)$  curves under applied magnetic fields up to 5 T (left), and the corresponding magnetic field dependence of  $-\Delta S_M^{\max}$  (right).

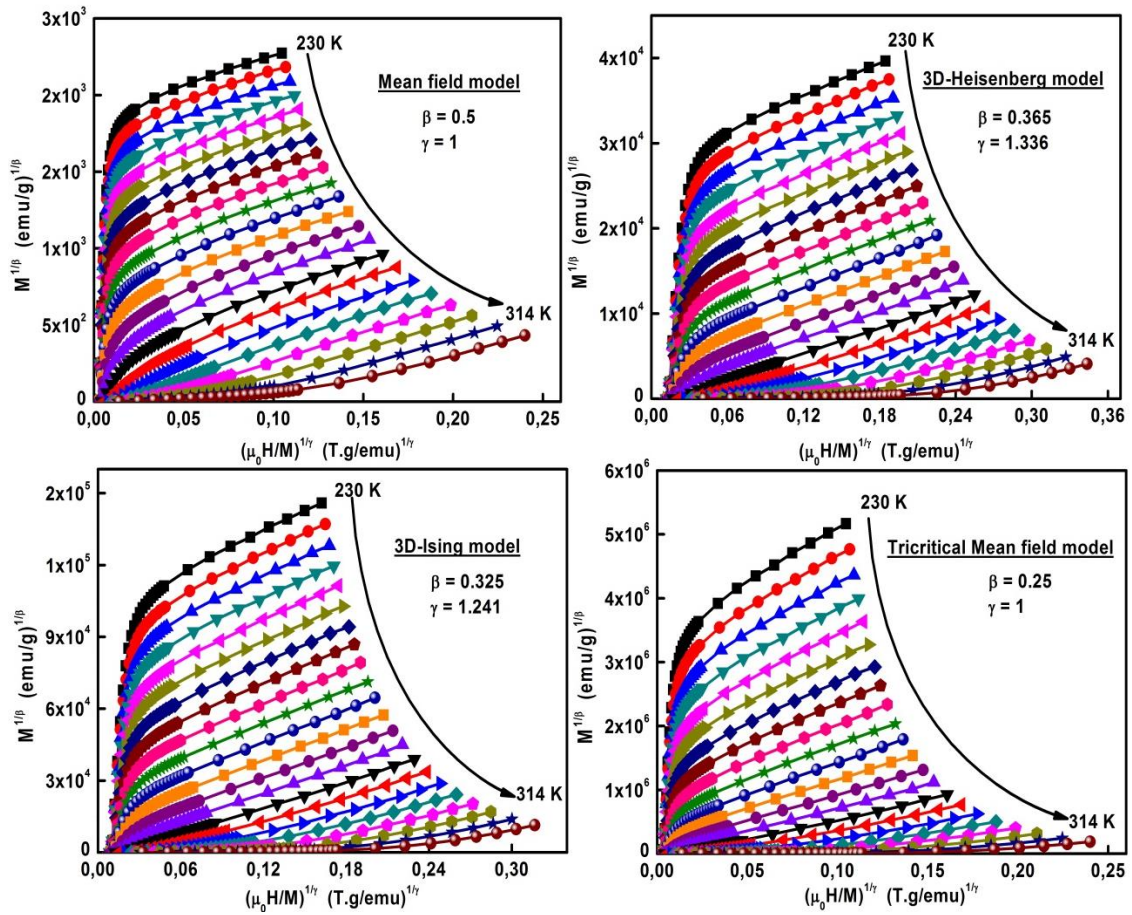


**Fig. 6:** Experimental and calculated  $-\Delta S_M(T)$  curves under an applied field up to 5 T, and the corresponding Landau's coefficient  $a(T)$ ,  $b(T)$  and  $c(T)$ .

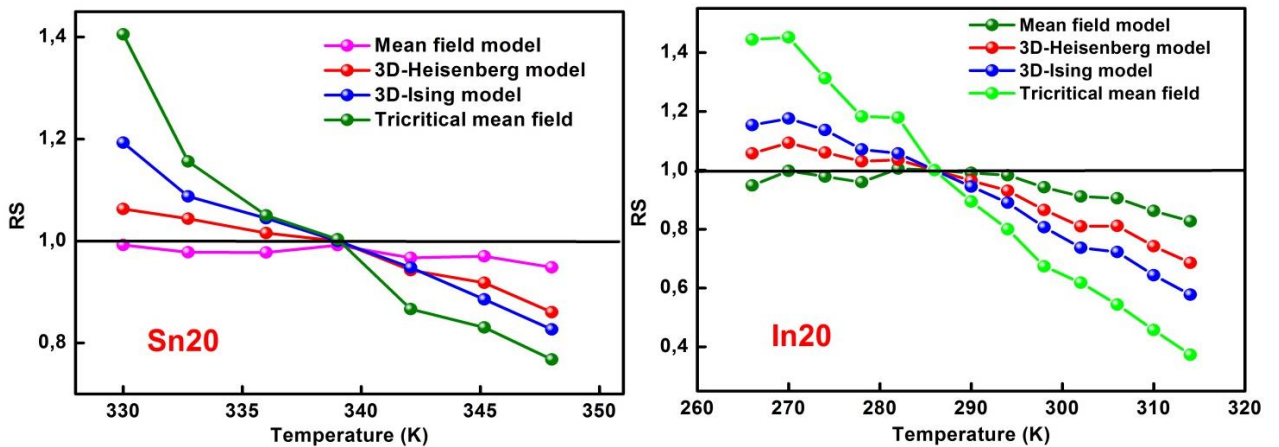


**Fig. 7:** Modified Arrott plots for Sn20 alloy: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_{c,1}^A$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.



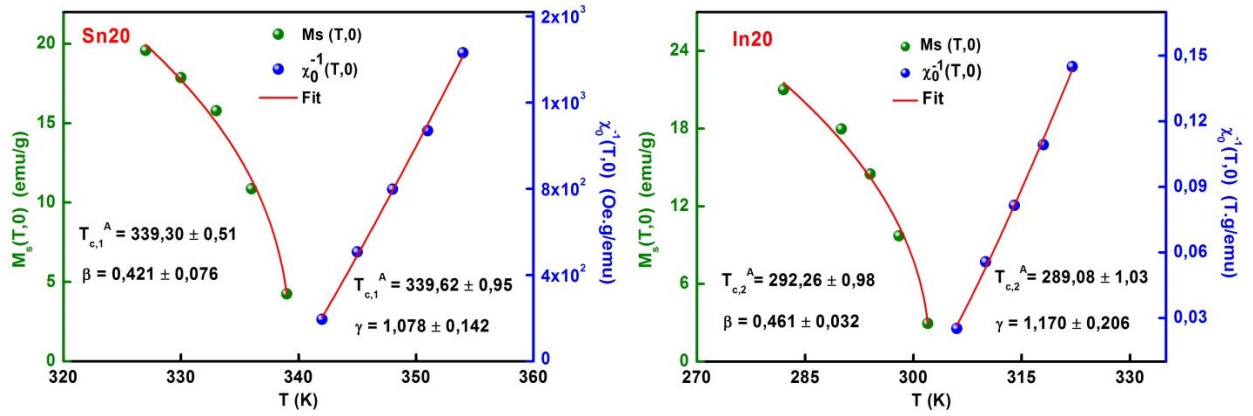


**Fig. 8:** Modified Arrott plots for In20 alloy: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_{C,2}^A$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.

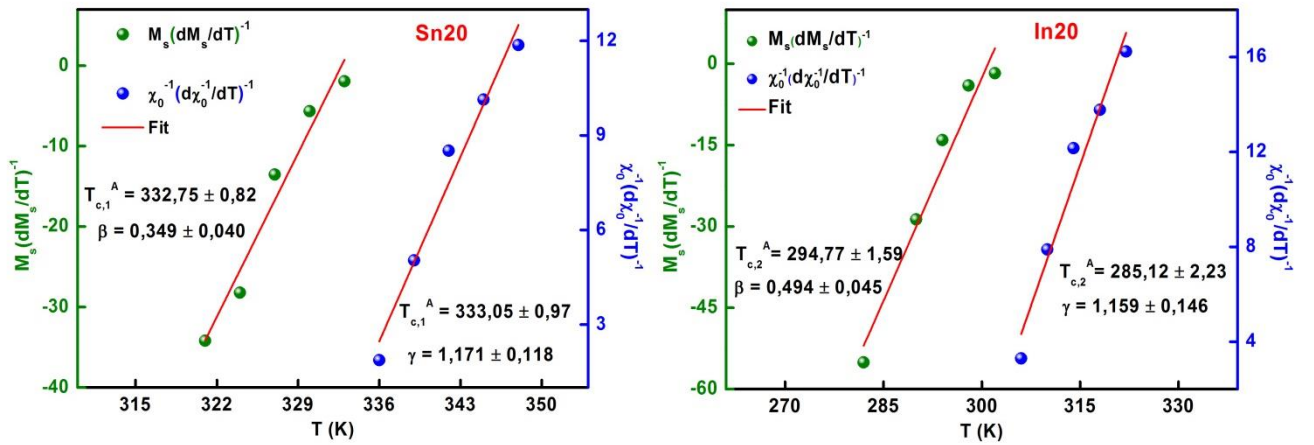


**Fig. 9:** Relative slope ( $RS = S(T)/S(T_C)$ ) as a function of temperature using different models.

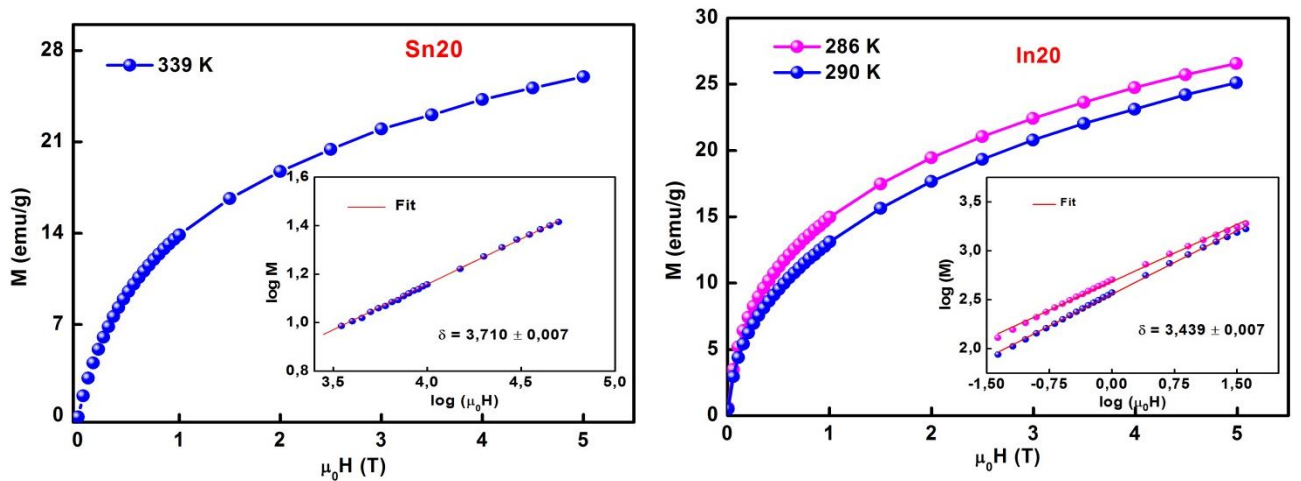




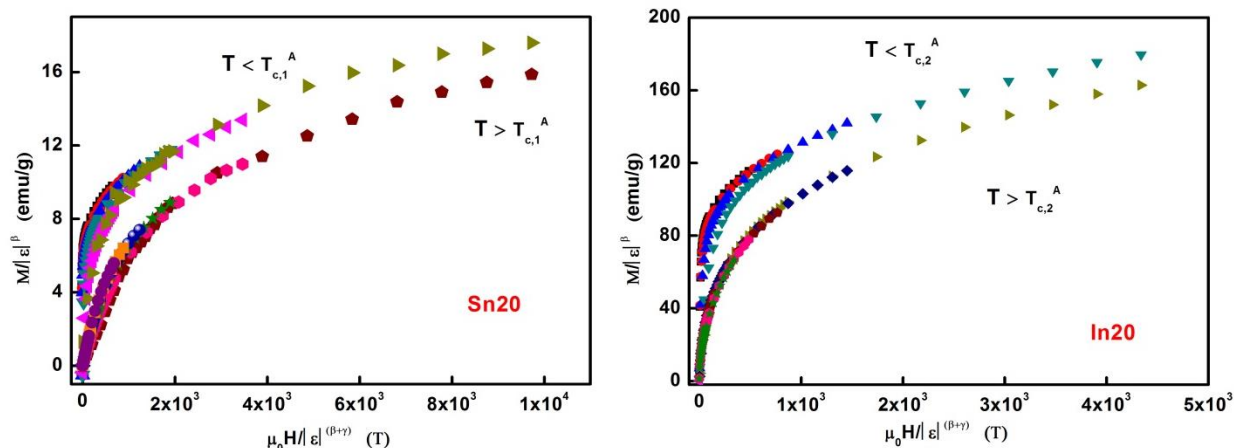
**Fig. 10:** Temperature dependence of spontaneous magnetization  $M_S(T, 0)$  and the inverse of initial susceptibility  $\chi_0^{-1}(T, 0)$  around  $T_C^A$ .



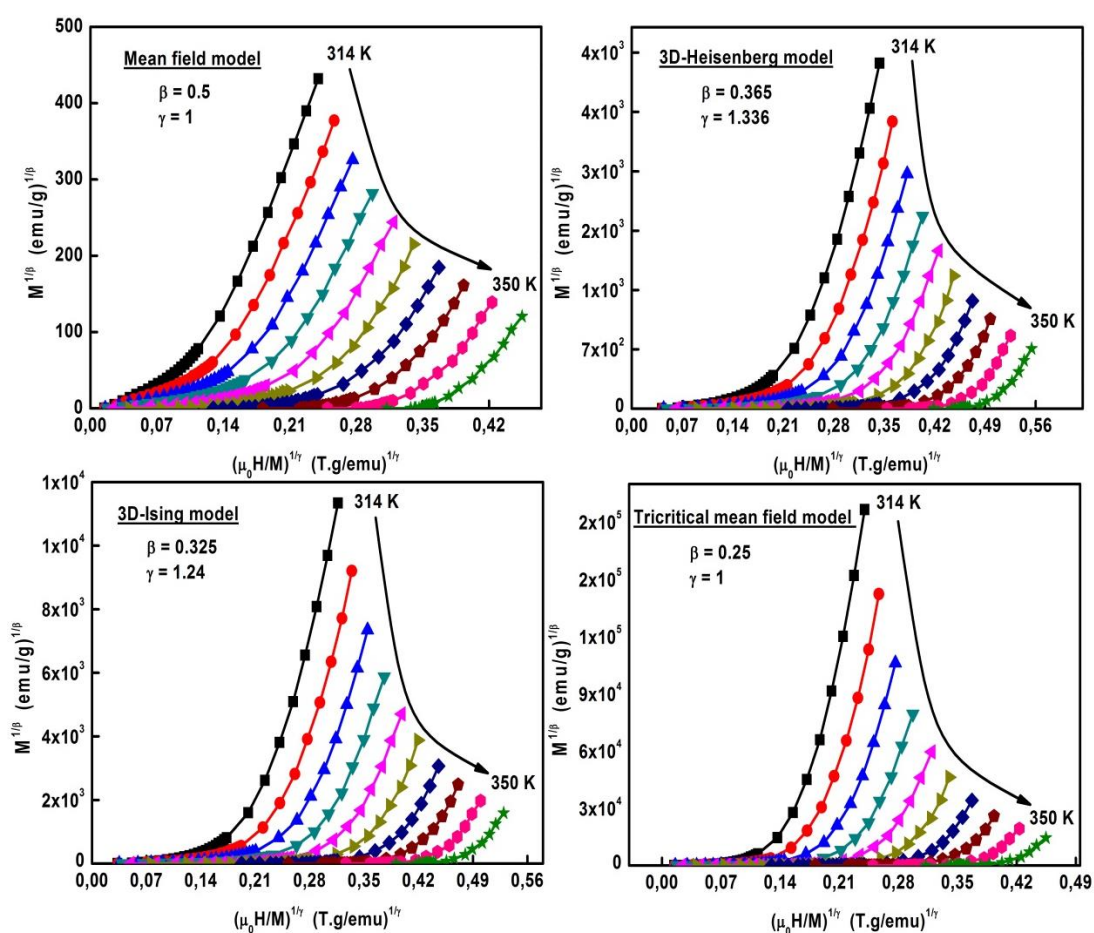
**Fig. 11:** Kouvel-Fisher plots for the spontaneous magnetization and the inverse of initial susceptibility around  $T_C^A$ .



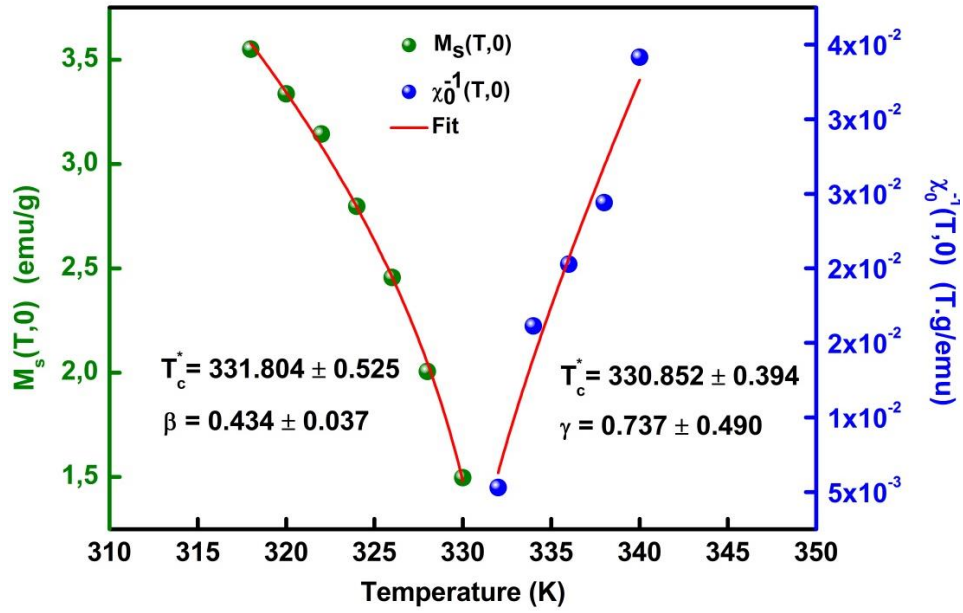
**Fig. 12:** Critical isotherms. Insets show the log-log plot.



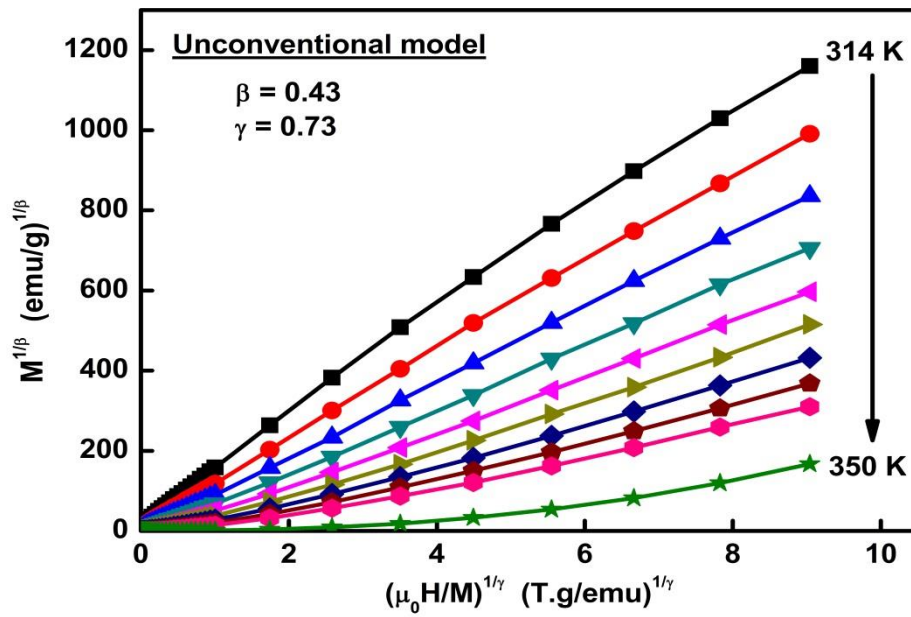
**Fig. 13:** Scaling plots indicating universal curves above and below the Curie temperature for Sn20 and In20 alloys.



**Fig. 14:** Modified Arrott plots for In20 alloy: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  around  $T_c^*$  with the mean field model, 3D-Heisenberg model, 3D-Ising model and tricritical mean field model.



**Fig. 15:** Temperature dependence of spontaneous magnetization  $M_S(T, 0)$  and the inverse of the initial susceptibility  $\chi_0^{-1}(T, 0)$  around  $T_c^*$ .



**Fig. 16:** Modified Arrott plots: isotherms of  $M^{1/\beta}$  against  $(H/M)^{1/\gamma}$  with  $\beta = 0.43$  and  $\gamma = 0.73$ .

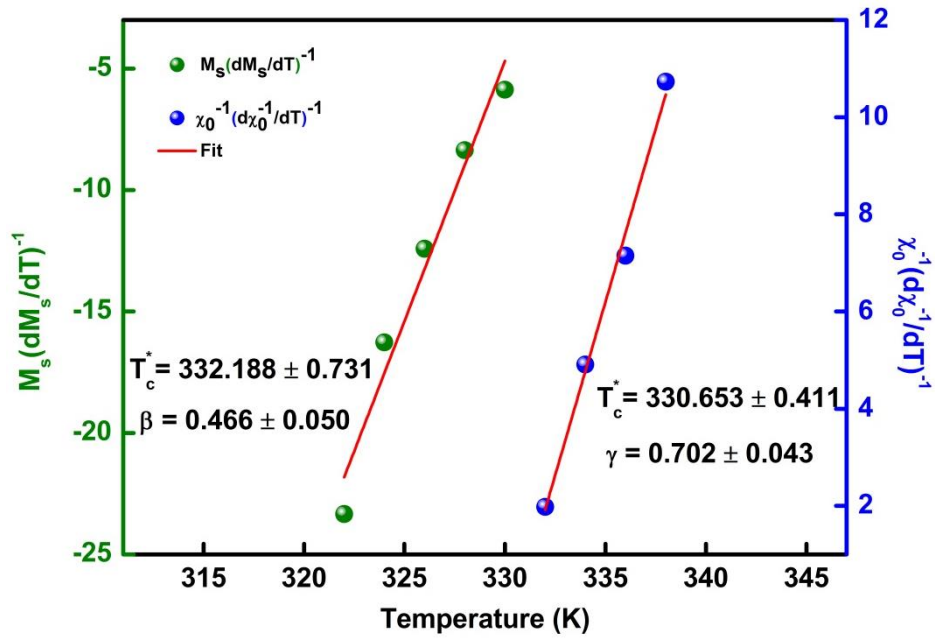


Fig. 17: Kouvel-Fisher plots for the spontaneous magnetization and the inverse of the initial susceptibility around  $T_c^*$ .

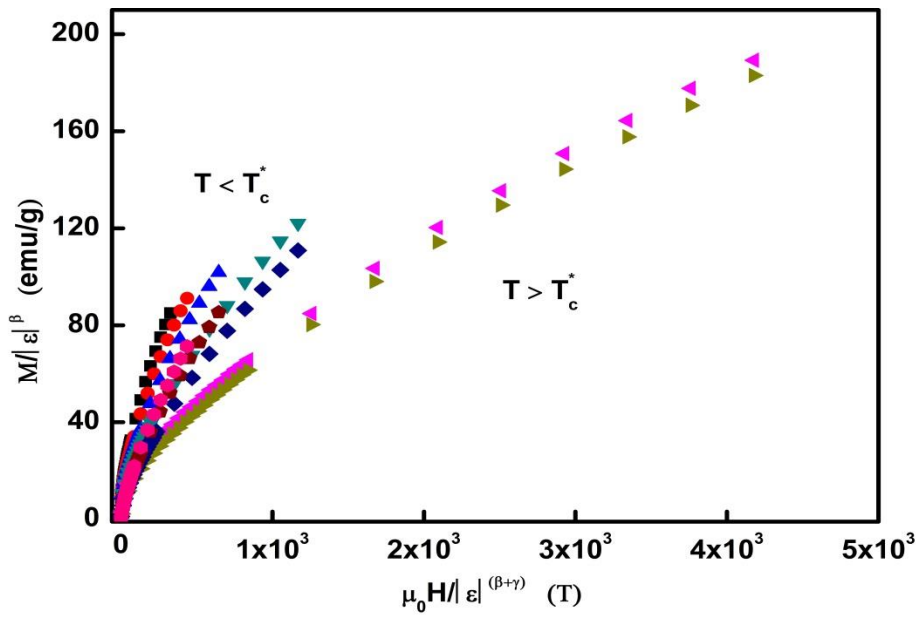
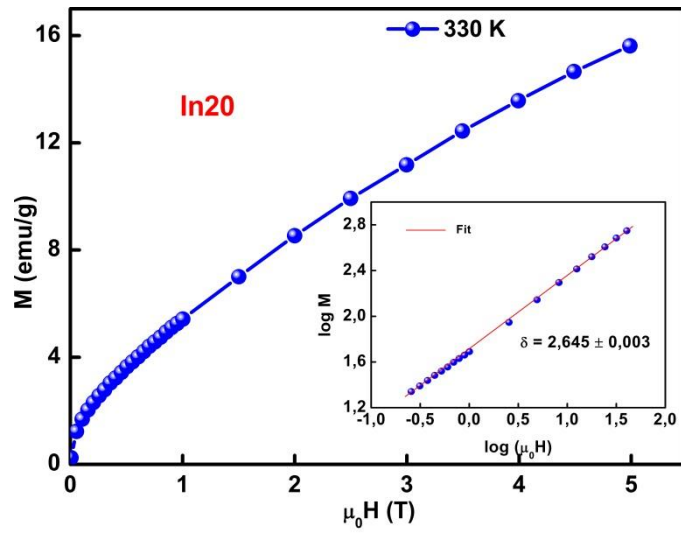


Fig. 18: Scaling plots indicating universal curves above and below  $T_c^*$ .



**Fig. 19:** Critical isotherm of  $M(\mu_0 H, T_c^*)$ . The log-log plot is shown in the *inset*.