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Influence of Concentration Fluctuations on Relaxation Processes in Spin Glasses

Julia N. Wagner ^{1,2,3,4}, Wolfgang Häußler ^{1,2}, Olaf Holderer ⁵ , Andreas Bauer ¹,
Stephen M. Shapiro ⁶ and Peter Böni ^{1,*}

¹ Physik-Department E21, Technische Universität München, 85747 Garching, Germany; juna.wagner@gmail.com (J.N.W.); whaeussler@arcor.de (W.H.); Andreas.Bauer@frm2.tum.de (A.B.)

² Heinz-Maier-Leibnitz-Zentrum (MLZ), Technische Universität München, Lichtenbergstr. 1, 85748 Garching, Germany

³ Institute for Materials Science, University of Stuttgart, Heisenbergstr. 3, 70569 Stuttgart, Germany

⁴ Karlsruhe Institute for Technology, KNMF and IAM-WK, Hermann-von-Helmholtz-Platz 1, 76344 Eggenstein-Leopoldshafen, Germany

⁵ Forschungszentrum Jülich GmbH, JCNS Outstation at MLZ, Lichtenbergstr. 1, 85747 Garching, Germany; o.holderer@fz-juelich.de

⁶ Brookhaven National Laboratory, Department of Physics, Upton, NY 11973, USA; sshapiro@bnl.gov

* Correspondence: peter.boeni@frm2.tum.de; Tel.: +49-89-289-14711

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Abstract: Using the unique combination of atomically resolved atom probe tomography (APT) and volume averaged neutron (resonance) spin echo (NRSE and NSE) experiments, the influence of nano-scaled clusters on the spin relaxation in spin glasses was studied. For this purpose, the phase transition from the paramagnetic phase to the spin glass phase in an Fe-Cr spin glass with a composition of Fe_{17.8}Cr_{82.2} was studied in detail by means of NRSE. The microstructure was characterised by APT measurements, which show local concentration fluctuations of Fe and Cr on a length scale of 2 to 5 nm, which lead (i) to the coexistence of ferro- and anti-ferromagnetic clusters and (ii) a change of the magnetic properties of the whole sample, even in the spin glass phase, where spins are supposed to be randomly frozen. We show that a generalized spin glass relaxation function, which was successfully used to describe the phase transition in diluted spin glasses, can also be used for fitting the spin dynamics in spin glasses with significant concentration fluctuations.

Keywords: spin glasses; disordered systems; magnetism; neutron spin echo

1. Introduction

Spin glasses possess no long-range magnetic order due to frustrated magnetic interactions and disorder, which leads to stochastically oriented magnetic moments. Typically, phase transitions in spin glasses occur at temperatures below 60 K. In spin glasses, the disorder in the system at high temperatures, which is typically paramagnetic, reappears at low temperatures in a frozen state. Due to the frustration of the magnetic states in the spin glass phase, slow decay processes on large time scales may be observable [1]. Inelastic and quasi-elastic neutron scattering capture an important part of these slow relaxation processes from the picosecond to the nanosecond regime [2,3]. Muon spin relaxation (μ SR) allows for extending the time scale to 1 μ s [4]. However, being a local probe μ SR does not allow for obtaining information on the spatial correlations of the relaxation processes.

Based on the neutron scattering data, Pickup et al. suggested a generalized spin glass relaxation process [5] by connecting the probabilistic Weron model [6] and a model that describes highly disordered systems on the basis of Lévy-stable distributions, firstly introduced by Tsallis [7]. Pickup used this model to describe successfully the relaxation processes in the diluted spin glass

systems $\text{Au}_{1-x}\text{Fe}_x$ and $\text{Cu}_{1-x}\text{Mn}_x$ [5]. In this way, it was shown that processes in spin glasses are similar to those in other complex disordered systems. In a few spin glass systems, the so-called cluster spin glasses, clusters of magnetic order coexist with the magnetic amorphous spin glass phase [8]. However, only little is known about the relaxation processes in these cluster spin glasses.

In this contribution, we study the relation between microstructure leading to magnetically ordered clusters and the averaged spin relaxation in an Fe-Cr sample, which shows a temperature driven phase transition to a cluster spin glass (SG) phase. For this purpose, we combine two complementary high resolution methods, namely the atomically resolved atom probe tomography (APT) [9–11] and the high resolution neutron (resonance) spin echo spectroscopy (N(R)SE). Additionally, magnetization measurements complete the characterization of the specimen.

Atom probe tomography (APT) measurements show that nanometer sized Cr rich regions coexist next to Fe rich regions. These may result in magnetically ordered clusters even in the spin glass phase. The shown results support the assumption that the Weron model introduced by Pickup et al. [5] for diluted spin glasses like $\text{Au}_{1-x}\text{Fe}_x$ and $\text{Cu}_{1-x}\text{Mn}_x$ still works for spin glasses with inhomogeneities in the microstructure, if some of the parameters are restricted; however, the physical interpretation of the results may differ. The model will be explained in detail below. It is shown that even small scale inhomogeneities in the microstructure may change the overall relaxation processes of spin glasses significantly. The relaxation process is then governed by different contributions of hierarchically ordered cluster interactions and parallel spin interactions compared to diluted spin glasses.

2. Experimental Details

For this study, an Fe-Cr alloy with a nominal composition of $c_{\text{Fe}} = 14.5$ and $c_{\text{Cr}} = 85.5$ at.% was used. The sample was prepared from starting materials with a purity of 99.99%, arc melted, annealed for four days at 1100 °C and quenched in water. To relax the occurring strains, the sample was annealed at 1000 °C for one day. The cylindrical sample has a height of approximately 20 mm and a diameter of about 10 mm. Atom probe tomography revealed an averaged chemical composition of $c_{\text{Fe}} = 17.8$ and $c_{\text{Cr}} = 82.2$ at.%. The concentration of impurities from the production process is approximately 0.01 at.% and thus neglectable. The phase diagram of the Fe-Cr-system is given by Burke et al. [12]. Based on the found chemical composition c_{Fe} and c_{Cr} , the material is supposed to show no magnetic order and to transfer directly from the paramagnetic regime to the spin glass phase.

The alternating current (AC) susceptibility measurements were performed using a Quantum Design physical property measurement system (PPMS). An excitation amplitude of 1 mT with an excitation frequency between 10 Hz and 10 kHz was used.

The relaxation measurements were carried out on the neutron (resonance) spin echo spectrometers RESEDA (REsonance Spin Echo spectrometer for Diverse Applications) [13] and J-NSE (Jülich Neutron Spin Echo spectrometer) [14] at the neutron source FRM II (Forschungs-Reaktor München II). The wavelength chosen was 5.5 Å for RESEDA and 5 Å for J-NSE. At RESEDA, a range of spin echo times τ from approx. 0.05 ns to 1 ns was accessible. With J-NSE, the range could be extended to 10 ns. The τ -dependence of the polarization P of the neutrons was independent of the momentum transfer q . Thus, to improve measurement statistics, the experimental results are averaged for q values between $0.04 \text{ \AA}^{-1} \leq q \leq 0.08 \text{ \AA}^{-1}$. The intermediate scattering function $S(q, \tau)$ was determined for temperatures $T = 12.1, 14.5, 19.4, 25.1, 30.1$ and 34.2 K. After correcting the data for depolarization by the sample and the spectrometer at the different temperatures, the data sets are normalized to the intermediate scattering function as measured at $T = 3.6$ K, where the dynamics appear to be frozen on the time scales achieved by the spectrometers.

The atom probe tomography measurements were carried out with a Cameca LEAP 4000X HR system from Cameca (CAMECA, Madison, WI, USA) located at the Karlsruhe Nano Micro Facility (KNMF). The atom probe tips were prepared by means of a focused ion beam with a Strata 400 system (ThermoFisher Scientific, Hillsboro, OR, USA). The atom probe tips were measured in the voltage mode

with a pulse rate of 200 kHz and a pulse fraction of 20% at a temperature of about 55.8 K. Analysis and reconstruction of the data was done with the IVAS 6.8.10 software package from Cameca [15].

3. Generalized Spin Glass Relaxation Model Based on Weron

In the Weron model [6] the fractal character of the spin glass is described by the exponent β , whereas k is a measure for the contribution of hierarchical parallel and thus independent relaxation processes. Combined with the relaxation time τ , the generalized relaxation function based on the Weron model is given by

$$\varphi(t) = [1 + k(t/\tau)^\beta]^{-1/k}. \quad (1)$$

In the Tsallis model [7], the sum of the entropy of two independent processes is larger or smaller than the direct sum of their entropies. The difference is scaled with the non-extensivity parameter q_τ . Brouers and Sotolongo-Costa [16] showed that there is a direct relation between the Tsallis and the Weron model when k is replaced by $k = (q_\tau - 1)/(2 - q_\tau)$. The application to spin glasses will be shortly described in the following and is explained in greater detail by Pickup et al. [5]. Generally, the dynamics in a system slows down with decreasing temperature. This statement also applies for spin glasses. Thus, the lifetime of relaxation processes, τ , must increase with decreasing temperature. For the temperature limit $T \rightarrow 0$, $\tau \rightarrow \infty$ must apply.

Above the spin glass transition temperature, T_{SG} , spin–spin interactions are dominant. In the Weron model, this fact is expressed by a relatively large value of β ($\beta_{\max} = 1$). Because the inter-cluster interactions are suppressed in the regime above T_{SG} , the value of k will be small. For high temperatures, where relaxation processes are dominated by thermal activation, $\beta = 1$ and $k \rightarrow 0$, i.e., the Weron model describes a pure exponential decay. For decreasing temperatures, the spins are slowed down (until they are frozen) and the contribution of parallel, independent relaxation processes increases: the value of k increases, while β decreases. If all relaxation processes occur in single spin clusters, which do not interact with each other, k converges to infinity. In this limit, $q_\tau = 2$ applies.

4. Results

The AC susceptibility measurements shown in Figure 1 demonstrate that the sample exhibits a temperature driven spin glass (SG) phase transition near $T_{SG} \approx 25$ K. This value is consistent with results shown by Burke et al. for an iron concentration of $c_{Fe} \sim 17.8$ at.% [12]. As expected from the chemical composition, there is no clear phase transition from the paramagnetic (PM) to a magnetically ordered phase for temperatures between 50 and 200 K in the AC susceptibility measurements.

Figure 2a shows the reconstructed APT with the elements Fe and Cr. The black box represents the area (120×30 nm², length $\times$ height) in which the 2D concentration maps of Fe and Cr are shown after integration of the concentration along the depth (2.5 mm) of the sample. The resulting 2D concentration maps of Cr, Fe and Cr + Fe, as analyzed by the software IVAS [15] are shown in Figure 2b–d). Impurities in the material, such as N and C from the production process and Ga from the FIB cut of the atom probe tip are locally far less than 0.05 at.%.

Figure 3 shows the intermediate scattering function versus the relaxation time as measured at RESEDA and J-NSE together with fits to the Weron model given by Equation (1). It is clearly seen that the spin fluctuations freeze with decreasing temperature T . For the fitting procedure, the following two obvious assumptions were made: (i) $0 \leq \beta \leq 1$. Of course, $\beta = 1$ above T_{SG} in the paramagnetic (PM) phase. Therefore, β can only decrease with decreasing T . (ii) τ has to monotonically increase with decreasing T due to freezing. If the parameter k was not restricted, it increased with decreasing T , as one would expect because more and more independent relaxation processes can contribute to the dynamics. However, τ decreased, which contradicts the physical intuition (dashed lines in Figure 4). Meaningful fits, i.e., a decreasing τ , was only obtained if k was constrained to a constant value for all $T \leq T_{SG}$. The exact value was hard to determine and was found to be $k \leq 4$. All values $k \leq 4$ led to a physically meaningful decrease of τ with decreasing temperature.

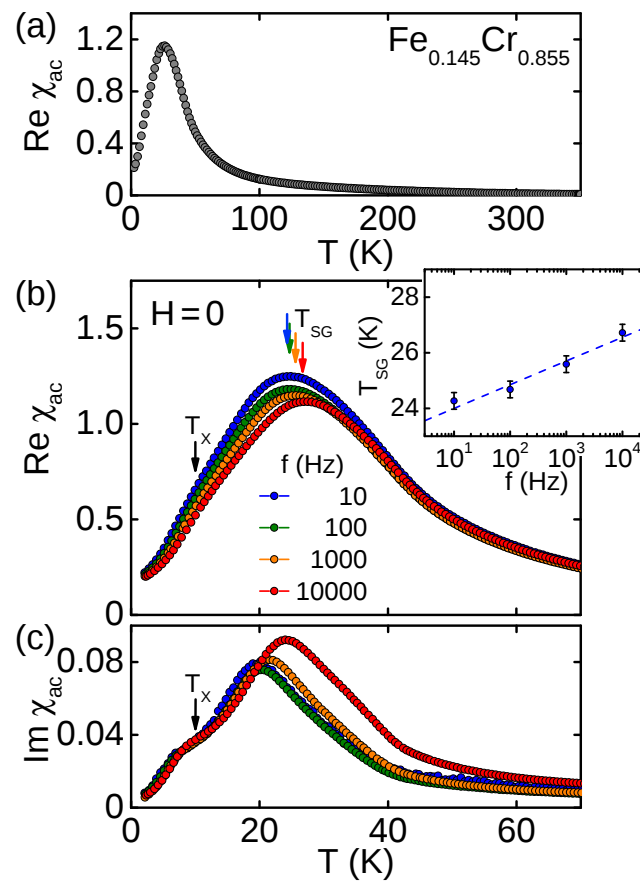


Figure 1. Temperature dependence of the alternating current (AC) susceptibility of the studied sample. (a) real part of the AC susceptibility between 2 K and 350 K measured at 911 Hz. The absolute value is large, when compared with a typical antiferromagnetic compound; (b) real part of the AC susceptibility, $\text{Re } \chi_{ac}$, as measured at low temperatures for excitation frequencies between 10 Hz and 10 kHz. As depicted in the inset and characteristic for a spin glass, the position of the maximum at T_{SG} shifts clearly as a function of the excitation frequency; (c) imaginary part of the AC susceptibility, $\text{Im } \chi_{ac}$, at low temperatures. A change of slope at $T_x \sim 10$ K in $\text{Re } \chi_{ac}$ and, in particular, in $\text{Im } \chi_{ac}$ suggests the presence of an additional transition, which is not captured in the neutron data and will not be considered further.

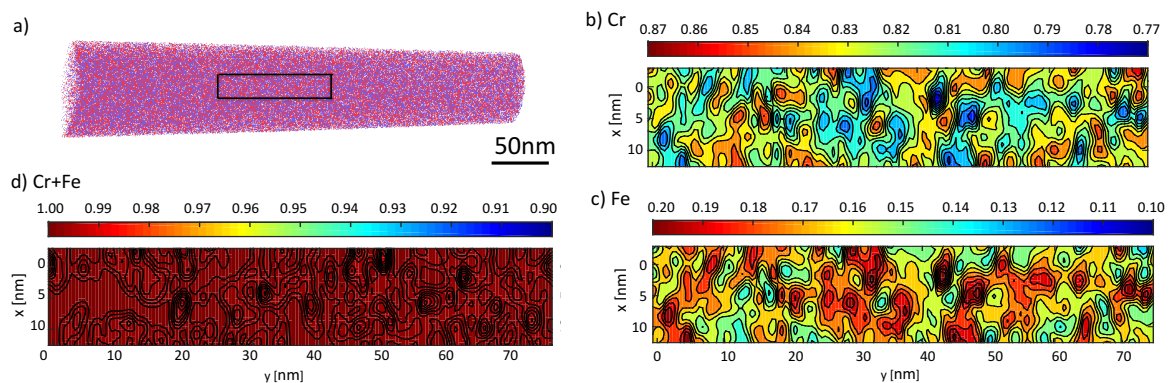


Figure 2. (a) Atom probe reconstruction of the Fe-Cr alloy, measured at 50 K. The black box marks the area where the data for the 2D concentration maps (b,c) are taken from. Clearly visible are small clusters of Cr and Fe, respectively. The concentration distribution is integrated over a sample slice with a thickness of 2.5 nm; (d) shows the summation of (b,c), which should sum up to 100%. The small differences of less than 1% are caused by impurities such as P, N, and Ga.

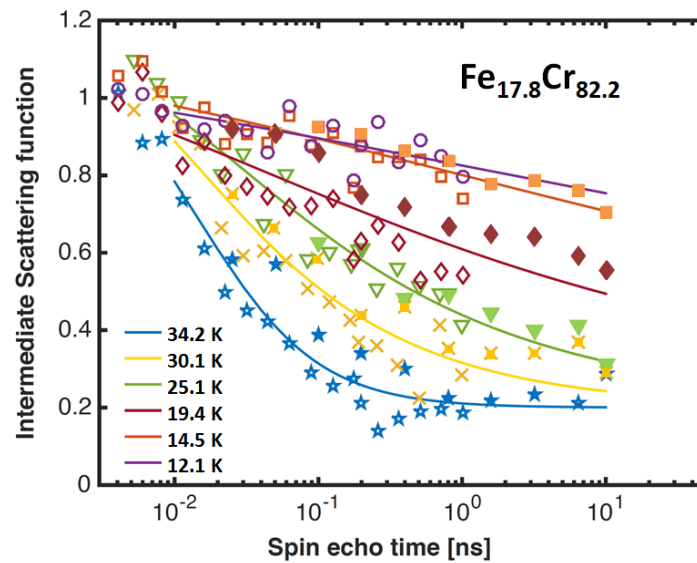


Figure 3. Intermediate scattering function as measured by neutron (resonance) spin echo at RESEDA (open symbols) and at J-NSE (filled symbols) for $q \simeq 0.06 \text{ \AA}^{-1}$. The solid lines show the restricted Weron model fitted to the experimental data (symbols) assuming $k = 2.5$ for $T \leq T_{\text{SG}}$. The error bars of the individual data points are given in Table 1. The parameters of the fit are shown in Figure 4.

Of course, the exact values of the fit parameters depend strongly on the chosen value of k , thus exact values of the fit parameters cannot be extracted. The following discussion is based on fit results obtained with a constant $k = 2.5$ for $T \leq 25.1 \text{ K}$. Figure 4 shows the resulting temperature dependence for k , β and τ with the restriction $k = 2.5$ (solid lines and closed symbols) compared to fit results with no boundary conditions except $0 \leq \beta \leq 1$ (dashed lines with open symbols). Table 1 shows all values of the intermediate scattering function dependent on the spin echo times for all temperatures, so that the fit values might be reproduced by interested readers.

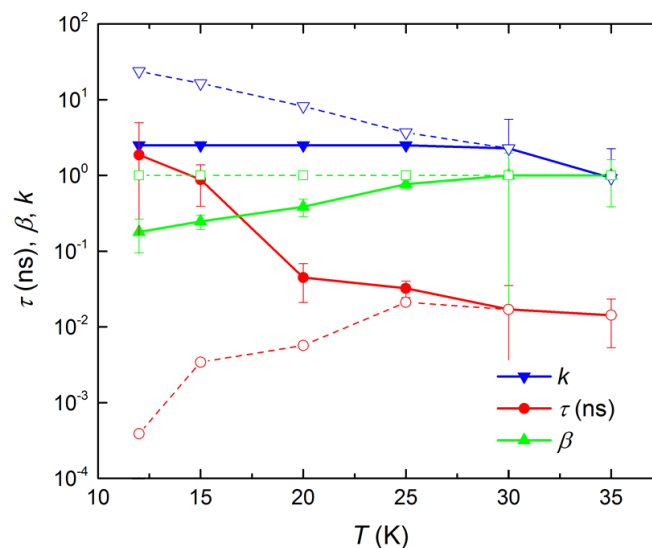


Figure 4. Parameters τ , β and k analysed by a fit of the Weron function to the intermediate scattering functions observed for different temperatures by neutron (resonance) spin echo spectroscopy (N(R)SE) measurements with no restrictions except $0 \leq \beta \leq 1$ (dashed lines and open symbols) and with restrictions (i) $0 \leq \beta \leq 1$, (ii) τ constantly increasing with decreasing T and (iii) $k = 2.5$ for $T \leq 25 \text{ K}$ (solid lines and closed symbols). The error bars for the non-restricted fits (not shown) are very large because the parameters are strongly correlated.

Table 1. Spin echo time τ_{NSE} and values of the intermediate scattering function ($I(\tau_{NSE})/\text{error}$) for all measured temperatures ($12.1 \leq T \leq 34.2$ K). The data from RESEDA covers spin-echo times in the range $0 < \tau_{NSE} < 1$ ns (roman font). The data from J-NSE extends the spin-echo times to approximately 10 ns (slanted font). The data overlap between approximately 0.1 ns and 1 ns.

τ_{NSE} (ns)	$I(\tau_{NSE})$ 12.1 K	Error	τ_{NSE} (ns)	$I(\tau_{NSE})$ 14.5 K	Error	τ_{NSE} (ns)	$I(\tau_{NSE})$ 19.4 K	Error
0.00409	1.02276	0.06006	0.00409	1.05768	0.05938	0.00409	0.98686	0.05903
0.00602	1.00878	0.05935	0.00602	1.09378	0.06067	0.00602	1.06622	0.06179
0.00813	0.96591	0.05791	0.00813	1.01485	0.05781	0.00813	0.95992	0.05795
0.01145	0.92834	0.0538	0.01145	0.92272	0.05197	0.01145	0.82503	0.05063
0.01617	0.91736	0.05504	0.01617	0.97386	0.05513	0.01617	0.88892	0.05423
0.02253	0.93979	0.05484	0.02253	0.88035	0.05109	0.02253	0.79933	0.05037
0.03181	0.91483	0.05615	0.03181	0.90611	0.05406	0.03181	0.77239	0.05158
0.04465	0.85836	0.05449	0.04465	0.88378	0.0536	0.04465	0.74521	0.05102
0.06279	0.97755	0.06212	0.06279	0.95468	0.05939	0.06279	0.71707	0.05341
0.08888	0.87447	0.05982	0.08888	0.87917	0.0582	0.08888	0.72044	0.05503
0.12548	0.92691	0.06301	0.12548	0.90804	0.06025	0.12548	0.7405	0.05691
0.17724	0.78653	0.05843	0.17724	0.76687	0.05573	0.17724	0.5812	0.05268
0.19732	0.87647	0.06422	0.19732	0.90613	0.0629	0.19732	0.63119	0.05613
0.26199	0.93792	0.06647	0.26199	0.84559	0.06081	0.26199	0.66925	0.05729
0.36492	0.8335	0.0552	0.36492	0.84718	0.05369	0.36492	0.62548	0.04892
0.51595	0.89125	0.05869	0.51595	0.8401	0.05497	0.51595	0.52864	0.048
0.71752	0.85023	0.07396	0.71752	0.79749	0.06927	0.71752	0.55169	0.06368
1.01391	0.79711	0.07074	1.01391	0.73849	0.06605	1.01391	0.54046	0.06272
			<i>0.02529</i>	<i>0.92578</i>	<i>0.0217</i>	<i>0.02543</i>	<i>0.91805</i>	<i>0.02341</i>
			<i>0.04978</i>	<i>0.90893</i>	<i>0.02173</i>	<i>0.04989</i>	<i>0.90867</i>	<i>0.02357</i>
			<i>0.0998</i>	<i>0.92581</i>	<i>0.02124</i>	<i>0.1003</i>	<i>0.8605</i>	<i>0.02246</i>
			<i>0.19972</i>	<i>0.8752</i>	<i>0.01941</i>	<i>0.19945</i>	<i>0.74776</i>	<i>0.0202</i>
			<i>0.39854</i>	<i>0.86228</i>	<i>0.01845</i>	<i>0.39976</i>	<i>0.71885</i>	<i>0.0191</i>
			<i>0.79839</i>	<i>0.83665</i>	<i>0.01753</i>	<i>0.79888</i>	<i>0.66614</i>	<i>0.01803</i>
			<i>1.59816</i>	<i>0.7778</i>	<i>0.01495</i>	<i>1.59894</i>	<i>0.64925</i>	<i>0.01571</i>
			<i>3.19855</i>	<i>0.7861</i>	<i>0.01559</i>	<i>3.19747</i>	<i>0.64314</i>	<i>0.0161</i>
			<i>6.39774</i>	<i>0.76103</i>	<i>0.01594</i>	<i>6.39856</i>	<i>0.59164</i>	<i>0.01658</i>
			<i>9.9985</i>	<i>0.70518</i>	<i>0.02048</i>	<i>9.99848</i>	<i>0.55466</i>	<i>0.0217</i>

Table 1. Cont.

τ_{NSE} (ns)	$I(\tau_{NSE})$ 25.1 K	Error	τ_{NSE} (ns)	$I(\tau_{NSE})$ 30.1 K	Error	τ_{NSE} (ns)	$I(\tau_{NSE})$ 34.2 K	Error
0.00526	1.09754	0.06667	0.00525	0.96771	0.0617	0.00409	1.01164	0.06198
0.00754	1.03817	0.06788	0.00765	1.00928	0.06715	0.00602	0.88442	0.05747
0.01079	0.99185	0.06757	0.01082	0.92648	0.06523	0.00813	0.89517	0.05781
0.01512	0.88966	0.06278	0.01508	0.88101	0.06266	0.01145	0.73806	0.04983
0.02125	0.80296	0.05852	0.02125	0.66394	0.05382	0.01617	0.61188	0.04774
0.03002	0.85592	0.06467	0.02987	0.59078	0.05516	0.02253	0.49692	0.04387
0.04216	0.67408	0.05884	0.04217	0.60437	0.0567	0.03181	0.44948	0.04482
0.05965	0.8016	0.06767	0.05955	0.57911	0.05997	0.04465	0.42374	0.04469
0.08404	0.58313	0.06082	0.08394	0.50844	0.05899	0.06279	0.36484	0.04685
0.11873	0.60005	0.06782	0.11872	0.4726	0.06416	0.08888	0.28966	0.04716
0.16803	0.57163	0.08267	0.16771	0.42479	0.0778	0.12548	0.25518	0.04778
0.19188	0.60791	0.09113	0.19179	0.37019	0.0829	0.17724	0.27402	0.04849
0.25498	0.50625	0.07413	0.25486	0.36084	0.07005	0.19732	0.21112	0.05007
0.355	0.56178	0.06561	0.35483	0.30882	0.059	0.26199	0.14018	0.04911
0.50215	0.51515	0.07345	0.50191	0.22345	0.06644	0.36492	0.1709	0.04189
0.70143	0.49382	0.10095	0.70109	0.41314	0.09901	0.51595	0.19155	0.04417
0.9911	0.41238	0.09801	0.99062	0.28469	0.09488	0.71752	0.19639	0.05956
0.09993	0.62805	0.02397	0.02534	0.75225	0.02896	1.01391	0.18747	0.05861
0.20028	0.61087	0.02242	0.04913	0.66412	0.02883	0.02533	0.58292	0.03211
0.39978	0.48247	0.02087	0.09842	0.58733	0.02762	0.05043	0.56945	0.03201
0.79925	0.49432	0.02003	0.19907	0.43724	0.02542	0.09976	0.38865	0.03089
1.59864	0.44597	0.01732	0.40046	0.45993	0.02417	0.19893	0.34034	0.02876
3.19816	0.40069	0.01767	0.79944	0.3529	0.02284	0.39957	0.29977	0.02729
6.3974	0.41392	0.01823	1.59859	0.34077	0.01993	0.79908	0.2258	0.02593
9.99879	0.31263	0.02397	3.19825	0.34015	0.02046	1.59863	0.21929	0.02263
			6.39851	0.36829	0.02118	3.19827	0.23321	0.0233
			9.99884	0.29178	0.02805	6.3991	0.21245	0.02436
						9.99846	0.28799	0.03217

5. Discussion

The spin glass transition temperature $T_{SG} \sim 25$ K as measured by means of AC susceptibility and neutron scattering are consistent with each other. For N(R)SE, the transition might be identified due to the observation that the time dependence becomes more exponential with increasing temperature. $T_{SG} \sim 25$ K is also consistent with the value given by Burke et al. [12] for an iron concentration of $c_{Fe} = 17.8$ at.%. The susceptibility measurements show that there is no other phase transition for $T > T_{SG}$, indicating that there is no magnetically ordered phase in the material (Figure 1). A change of slope at $T_x \sim 10$ K in $\text{Re } \chi_{ac}$ and, in particular in $\text{Im } \chi_{ac}$, suggests the presence of an additional transition, which shows no signature in the neutron scattering data and will not be considered further.

The atom probe tomography studies clearly show that Fe rich regions coexist with Cr rich regions in the material (Figure 2). In those zones, the concentration of Fe may be high or low enough that ferromagnetic (FM) or antiferromagnetic (AFM) ordering will be observed. According to Burke et al. [12], an Fe concentration of about 19 at% for FM ordering and of 16 at% for AFM ordering is necessary for temperatures lower than ~ 50 K. Figure 2 shows that those values are reached locally over a length scale of approximately 2–5 nm.

According to Burke et al. [12] and Fincher and Shapiro et al. [17,18], the Néel temperature for concentrations of c_{Fe} of 14 at.% is supposed to be $T_N \sim 60$ K. The Curie temperature for $c_{Fe} = 20$ at.% is $T_C \sim 80$ K. Thus, with decreasing T , a change in polarization and intensity of the neutron signal around $q = 0.06 \text{ \AA}^{-1}$ is expected as follows: approaching T_C from high temperatures will initiate the formation of ferromagnetic clusters, which reduces the polarization of the scattered neutrons. Decreasing T further towards T_N leads to the formation of antiferromagnetic ordering in regions surrounding the ferromagnetically ordered regions. Thus, the size of the ferromagnetic clusters decreases, resulting in an increase in polarization. Lowering T further favors the spin glass phase, thus shrinking the regions of both ferromagnetic and antiferromagnetic phase further, thus leading to a continuing increase of intensity and polarization with decreasing T .

Indeed, these trends can be qualitatively observed in the neutron scattering data. Figure 5 shows the neutron polarization and the intensity of the neutrons at $q = 0.06 \text{ \AA}^{-1}$ with decreasing T . The polarization decreases with decreasing T , reaches a minimum around 120 K and increases again. The deviation from the expected minimum around 80 K may be due to the volume averaging and q -weighting by the neutrons. Overall, an increase in neutron intensity is observed as expected.

To put our results in a more quantitative basis, we estimate the size of the ferromagnetic clusters using the expression for the polarization of the neutrons $P = P_0 \exp(-\frac{1}{3}\gamma^2 \langle B^2 \rangle \frac{\delta}{v^2} d_{eff})$ derived by Halpern and Holstein [19] in the limit of small fields and domains. P is a function of the gyromagnetic ratio $\gamma = 2.916 \text{ kHz/Gauss}$, the magnetic field B , the domain size of the ferromagnetic clusters δ , the effective sample thickness for iron d_{eff} , and the velocity of the neutrons v . The velocity of neutrons with a wavelength of $\lambda = 5.5 \text{ \AA}$ is $v \approx 720 \text{ m/s}$. For the magnetization of the Fe clusters, we assumed the bulk magnetization of Fe $\mu_M = 1.74 \text{ T}$ [20]. With $d_{eff} = d_{bulk} c_{Fe} = 10 \text{ mm} \cdot 0.145 \sim 1.38 \text{ mm}$ an approximate cluster size of $\sim 200 \text{ \AA}$ for temperatures between 250 K and 120 K is obtained. For the final polarization measured in our experiment (about 28%) at 8 K, this assumption corresponds to a size of the ferromagnetic clusters of $\delta \sim 140 \text{ \AA}$. This is still considerably larger than the regions of high Fe concentrations in the APT analysis, which scale to 2–5 nm. However, the conclusion, which can be drawn from APT and N(R)SE is similar: in regions of high Fe concentration, the ferromagnetic order might still be present, even if those regions are surrounded by pure spin glass phase. The same holds true for the regions of high Cr concentration, where the antiferromagnetic ordering might still be present. Both phenomena lead to islands of magnetic ordering in the spin glass phase.

Based on the above discussion, the results of the N(R)SE experiments can be interpreted as follows: for $T > T_{SG}$, the sample is magnetically ordered in locally restricted areas as indicated by the ATP-data, leading to a maximum value $\beta = 1$ and a minimal value of k , as seen in Figure 4. Regions of magnetic fluctuations are present, but their parallel hierarchical relaxation processes have only a small weight when compared to the spin-fluctuations within the ordered clusters. With decreasing T , the volume

fraction of the sample, where the spin fluctuations slow down increases and thus the contribution of the parallel relaxation processes increases due to the decreasing size of magnetically ordered clusters. The value of k increases, while β decreases. In diluted spin glasses, k is supposed to diverge to infinity ($q_T \rightarrow 2$) because relaxation processes on all time scales occur. In our sample, however, the value of k converges to a constant value (Figure 4). The value of k is constrained because of the limited number of parallel processes due to the remaining clusters, which are directly observed by ATP and indicated by the finite polarization of the neutrons at low T .

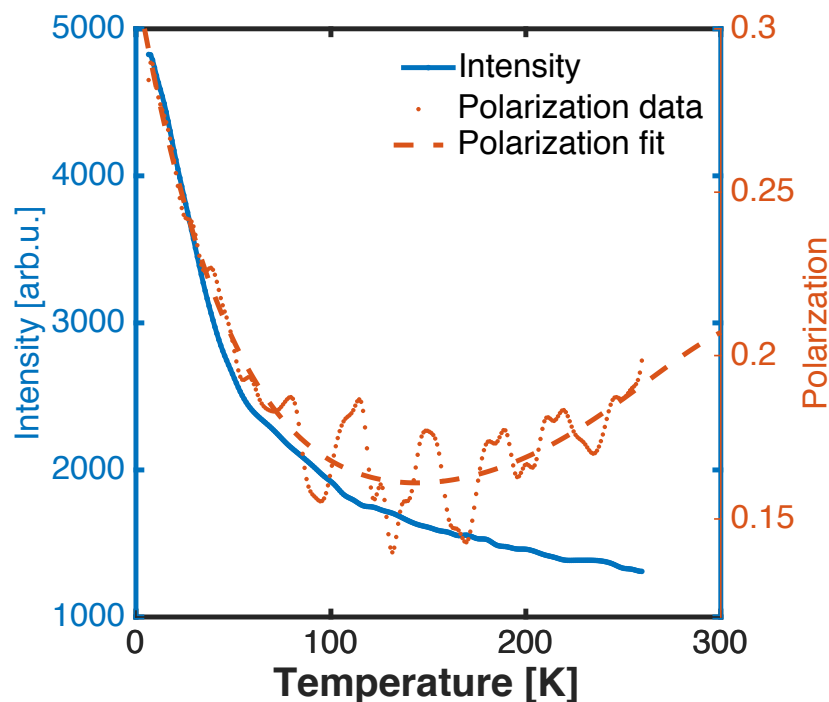


Figure 5. Temperature dependence of the intensity (blue) and the polarization (orange) as measured at $q = 0.06 \text{ \AA}^{-1}$ at RESEDA. The data are recorded during the course of cooling down the sample. Data recorded during heating up the sample give similar results (not shown), thus hysteresis effects do not play a role.

Figure 6 shows a comparison of the results obtained from the diluted spin glass systems $\text{Au}_{1-x}\text{Fe}_x$ and $\text{Cu}_{1-x}\text{Mn}_x$ as studied by Pickup [5] and the results of our work. The major difference between the two data sets below T_{SG} reflects the different mechanisms taking place during the spin freezing. Whereas in diluted spin glasses all spins are frozen in random orientations, in the Fe-Cr system ferromagnetic and antiferromagnetic clusters coexist within a sea of disordered spin glass phase. Therefore, the hierarchical order of relaxation processes changes from only parallel relaxation processes to spin-spin and parallel relaxation processes. Since our study already showed the complementation of scattering methods and atom probe tomography, the further investigations must concentrate on the study of the short range order of Fe-Cr alloys dependent on the found clusters. Here, special emphasis will be laid on the influence of the heat treatment of the material on both the phase transition and the short range order in the spin glass phase, because Fe-Cr alloys are known for their clustering and spinodal decomposition [21]. In addition, the phase transition at $T \sim 10 \text{ K}$ needs further attention to be fully identified.

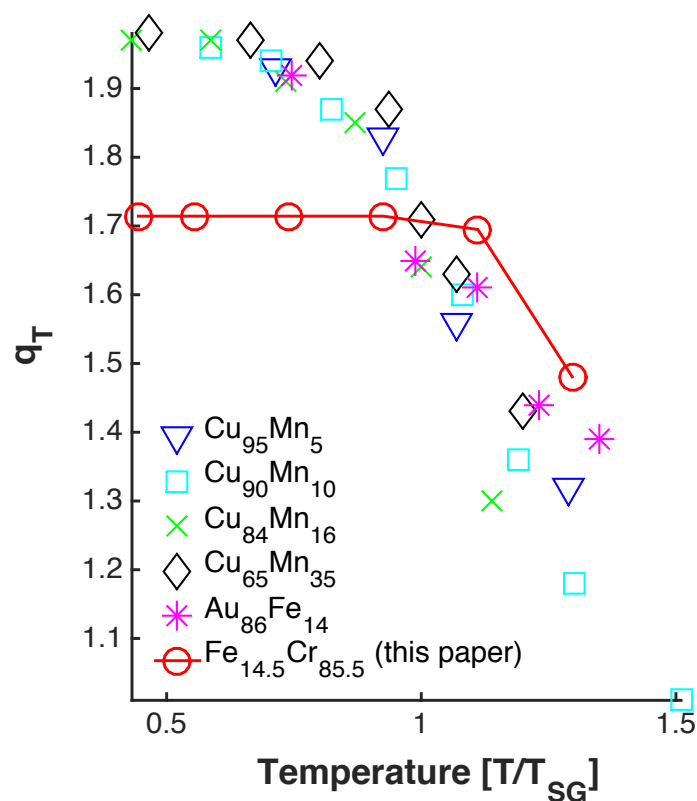


Figure 6. Comparison between data published by Pickup [5] on diluted spin glasses $\text{Au}_{1-x}\text{Fe}_x$ and $\text{Cu}_{1-x}\text{Mn}_x$ and the results observed in the present study on the cluster spin glass $\text{Fe}_{17.8}\text{Cr}_{82.2}$.

6. Conclusions

By combining atomically resolved (APT) with quasielastic neutron scattering using N(R)SE, we have characterized the evolution of the spin glass phase in the presence of ferro- and antiferromagnetic clusters in the cluster spin glass $\text{Fe}_{17.8}\text{Cr}_{82.2}$. The measured data can be interpreted in terms of the Weron model [6] that assumes the decay of spin fluctuations in parallel processes in the presence of magnetically ordered clusters. The finite value of the k -parameter indicates that the number of parallel processes in the cluster spin glass FeCr is limited. We speculate that cluster-type spin glasses may generally be classified based on the value of the parameter k that is a measure for the number of parallel processes at low T .

Author Contributions: J.N.W. was the leading scientist for experiments (neutrons and APT), analysis and interpretation of the data; W.H. and O.H. were responsible for RESEDA and N(R)SE, respectively, supported before, during and after the neutron experiments, discussed analysis and interpretation of the data; A.B. conducted the AC susceptibility measurements and interpreted them; S.M.S. provided the sample material; P.B. supported all steps of the experiment, the interpretation of the data and improved the paper. All authors reviewed the manuscript.

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Abbreviations

The following abbreviations are used in this manuscript:

APT	atom probe tomography
FRM II	Forschungs Reaktor München II (research neutron source Heinz Maier-Leibnitz)
FWHM	full width at half maximum
ISF	intermediate scattering function
J-NSE	Jülich Neutron Spin Echo spectrometer
KNMF	Karlsruhe Nano Micro Facility
N(R)SE	neutron (resonance) spin echo
NSE	neutron spin echo
PM	paramagnetic
PPMS	physical property measurement system
RESEDA	REsonance Spin Echo spectrometer for Diverse Applications
RF	radio frequency
SET	spin echo time
SG	spin glass

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