Element-specific soft x-ray spectroscopy, scattering, and imaging studies of the skyrmion-hosting compound $Co_8Zn_8Mn_4$

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A room-temperature skyrmion-hosting compound $Co_8Zn_8Mn_4$ has been examined by means of soft x-ray absorption spectroscopy, resonant small-angle scattering, and extended reference holography. An elementselective study was performed by exciting the 2*p*-to-3*d* transitions near Co and Mn *L*²,³ absorption edges. By utilizing the coherence of soft x-ray beams the element-specific real-space distribution of local magnetization at different temperatures has been reconstructed using iterative phase retrieval and holography with extended reference. It was shown that the magnetic moments of Co and Mn are ferromagnetically coupled and exhibit similar magnetic patterns. Both imaging methods provide a real-space resolution of 30 nm and allowed us to record a magnetic texture in the temperature range between $T = 20$ K and $T = 120$ K, demonstrating the elongation of the skyrmions along the principal crystallographic axes at low temperatures. Micromagnetic simulations have shown that such deformation is driven by a decreasing ratio of symmetric exchange interaction to antisymmetric Dzyaloshinskii-Moriya interaction in the system and effect of the cubic anisotropy.

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I. INTRODUCTION

Magnetic properties of transition metals (TMs) are generally determined by the 3*d* valence electrons. Resonant soft x-ray scattering at *L*2,³ absorption edges of TMs involves 2*p*-to-3*d* transitions, thus being an element-selective probe with the possibility to distinguish magnetic signal from different elements in multicomponent magnets [\[1\]](#page-10-0). Moreover, the spatial coherence of the polarized soft x-ray beams provided by modern synchrotron radiation sources and free-electron lasers give vast opportunities for the lensless imaging using coherent diffraction imaging $[2-4]$, ptychography $[5,6]$, and holographic techniques [\[7,8\]](#page-10-0). Both coherent resonant soft x-ray scattering (RSXS) imaging and holography allow us to perform real-space imaging of the magnetization distribution in thin samples with various environments, such as in high magnetic fields or at low temperatures. Flexibility of the environment and enhanced robustness of these methods against the specimen displacements are a significant advantage of the lensless techniques compared to scanning transmission magnetic x-ray microscopy (STXM) [\[9\]](#page-10-0), although the

possibility of cryogenic STXM imaging also has been recently demonstrated [\[10\]](#page-10-0). Coherent diffraction allows the solution of classical crystallographic inverse problem of phase retrieval by using the iterative reconstruction algorithms applied to the resonant diffraction intensities [\[11\]](#page-10-0). X-ray magnetic holography is based on the utilization of the interference between magnetic scattering from the object under investigation and the reference wave generated by charge scattering from the prepared source. Imaging experiments using holographic approaches can be realized in a few different ways: Fourier transform holography (FTH) is based on a reference wave scattered from one or multiple small (30–150 nm) pinholes placed near the sample aperture [\[7\]](#page-10-0). Alternatively, holography with extended reference by autocorrelation linear differential operation (HERALDO) [\[12\]](#page-10-0) can be performed. In contrast to FTH, the HERALDO technique implies the scattering from an extended reference object, such as a narrow slit or a sharp corner, which allows us to improve the contrast of the real-space image without compromising the resolution [\[13\]](#page-10-0). Moreover, fabrication of the extended reference is less challenging than the array of the reference pinholes [\[13,14\]](#page-10-0). Previously, FTH and HERALDO techniques were successfully applied for imaging of the element-specific magnetic domain patterns in thin films and multilayers with perpendicular magnetic anisotropy [\[15–17\]](#page-10-0). Both FTH and HERALDO with references milled at an oblique angle into

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the masks also allow the imaging at a tilted angle, which is relevant for the spintronic devices with an in-plane magnetic anisotropy, such as spin valves [\[18\]](#page-10-0) and magnetic nanoelements [\[19–21\]](#page-10-0). Thus the coherent soft x-ray scattering and imaging are powerful tools to study the spin ordering in multicomponent magnetic compounds with element selectivity.

Recently, several bulk materials that exhibit nontrivial topological spin textures and contain two or more magnetic elements have been discovered: doped B20-type alloys [\[22,23\]](#page-10-0), Co-Zn-Mn compounds with β -Mn structure [\[24\]](#page-10-0), molybdenum nitrides [\[25\]](#page-10-0), and Heusler alloys [\[26–28\]](#page-10-0). Competition between the magnetic interactions in noncentrosymmetric compounds results in the complex phase diagram. The interplay between exchange interaction, antisymmetric Dzyaloshinskii-Moriya interaction (DMI), and magnetocrystalline anisotropy may cause incommensurate spin phases such as helical, conical, and Bloch-type skyrmion lattice states [\[29,30\]](#page-10-0). The typical size of a magnetic skyrmion varies in a range from a few to a few hundred nanometers which makes them promising candidates for future spintronic applications such as skyrmion racetrack memory and logic devices [\[31\]](#page-10-0). The skyrmions can be manipulated by current pulses with ultralow current densities [\[32\]](#page-10-0), electric [\[33,34\]](#page-10-0) and microwave fields [\[35–37\]](#page-10-0), and temperature gradients [\[38–40\]](#page-11-0). In the past decade skyrmion textures have been extensively studied by means of small-angle neutron scattering (SANS) [\[41–43\]](#page-11-0) and Lorentz transmission electron microscopy (LTEM) [\[44,45\]](#page-11-0). Also, several groups reported on the resonant x-ray diffraction [\[46,47\]](#page-11-0), small-angle scattering [\[48–51\]](#page-11-0) studies of Bloch-type skyrmions in the chiral magnets and imaging of Néel-type skyrmions stabilized by interfacial DMI [\[52–55\]](#page-11-0). Since xray magnetic circular dichroism (XMCD) is sensitive to the component of the magnetization parallel to the incident x-ray beam, transmission soft x-ray imaging is a complementary method to LTEM, which is sensitive to the in-plane magnetic flux inside the sample [\[56\]](#page-11-0). Hence, for the Néel-type skyrmions, where the curl of magnetization lies in the sample plane and produce no contrast for LTEM without tilting the sample [\[57,58\]](#page-11-0), soft x-ray imaging has been successfully employed for the room-temperature Néel-type skyrmion-hosting thin films [\[52–55\]](#page-11-0), but not yet for the thin plates of polar magnets that order magnetically at cryogenic temperatures [\[59,60\]](#page-11-0).

Room-temperature magnetic ordering of the chiral β -Mntype Co-Zn-Mn alloy [\[24\]](#page-10-0) makes these materials promising for applications. The β -Mn-type compound Co₈Zn₈Mn₄ exhibits a transition from the paramagnetic state to a helical or Bloch-type skyrmion lattice state at $T_c \approx 300$ K with a magnetic modulation period of 125 nm [\[61\]](#page-11-0), and undergoes a spin glass transition at $T_e \approx 8$ K, probably due to freezing of Mn spins [\[62\]](#page-11-0). Frustration at the Mn site ultimately results in increment of a spin-glass transition temperature ($T_g \approx 30$ K) in the compound with higher Mn concentration $Co₇Zn₇Mn₆$. Moreover, a low-temperature frustration-induced equilibrium skyrmion phase has been recently found in the latter [\[62\]](#page-11-0). In the present work we utilized the polarization-dependent soft x-ray magnetic spectroscopy, coherent RSXS, and HER-ALDO techniques to perform an element-selective study of the magnetic interactions and long-range ordering in $Co_8Zn_8Mn_4$ compound. The coherence of the synchrotron

radiation allowed us to successfully combine small-angle scattering in transmission geometry with coherent diffraction imaging and employ the small-angle scattering patterns for the real-space reconstruction of the local magnetization distribution via an iterative phase retrieval algorithm. The coherent diffraction imaging results were compared to the real-space reconstruction results provided by HERALDO.

II. EXPERIMENT

X-ray spectroscopy, scattering, and imaging experiments were performed at the variable-polarization soft x-ray beamline BL-16A of the Photon Factory (KEK, Japan) [\[63\]](#page-11-0) and BL29 BOREAS of the ALBA synchrotron radiation laboratory (Cerdanyola del Vallés, Spain) [\[64\]](#page-11-0).

Experimental geometry of soft x-ray absorption (XAS) and XMCD experiments are shown in Fig. $1(a)$. A bulk polycrystalline $Co_8Zn_8Mn_4$ specimen was obtained from an ingot grown by the Bridgman method as described in Ref. [\[61\]](#page-11-0). The sample was polished to remove the potentially oxidized surface layer prior to the experiment. The sample was placed in the vacuum chamber with a pressure of 10^{-9} Torr equipped with a 5 T superconducting magnet. XAS and XMCD signals were measured with an energy resolution of 0.1 eV using the surface-sensitive total electron yield (TEY) method near Co and Mn $L_{2,3}$ absorption edges with right and left circularly polarized (RCP and LCP) x rays.

RSXS and HERALDO experiments were performed in the transmission geometry [Figs. $1(b)$ and $1(c)$]. Commercial silicon nitride membranes from Silson Ltd. (Southam, UK) were processed for soft x-ray experiments. The front side of each membrane was coated with an \approx 4- μ m-thick layer of gold to absorb the incoming x-ray beam. Further treatment of the membranes and thin plate fabrication were carried out by using a focused ion beam (FIB) setup Hitachi NB5000 equipped with a scanning electron microscope (SEM). Since the attenuation length for soft x rays in $Co_8Zn_8Mn_4$ alloy is of about 100 nm, the scattering and imaging experiments were carried out on thin plates of $Co_8Zn_8Mn_4$ with thickness of 200 and 150 nm, respectively. Two FIB-thinned plates of $Co_8Zn_8Mn_4$ containing a (001) plane were cut from the bulk single crystal. For the RSXS experiment a thin plate was attached directly to the $Si₃N₄$ membrane [Figs. [1\(d\)](#page-2-0) and [1\(e\)\]](#page-2-0). In the case of the RSXS sample the aperture with a diameter of 4.5 μ m and asymmetric shape, which provides a better convergence of the phase retrieval algorithm [\[65\]](#page-11-0), was drilled in the gold coating [Fig. $1(d)$].

In the case of the HERALDO sample we fabricated the sample aperture by a different fabrication approach: a large hole with a diameter of 6 μ *m* was drilled in the gold-coated membrane, and covered by a $1-\mu$ m-thick gold plate fabricated by FIB from a bulk specimen. Then, a circular sample aperture with a diameter of 700 nm and a reference slit with a length of 1 μ m and width 40 nm was milled in the Au plate [Fig. $1(f)$]. The slit length and distance from the aperture were chosen according to the separation conditions preventing the overlapping of sample autocorrelation and sample-reference cross correlation at the reconstruction [\[66\]](#page-11-0).

To prevent specimen damage by $Ga⁺$ ions in the aperture and reference milling process the thin plate was fixed to the

FIG. 1. Sketch of the (a) XMCD, (b) RSXS and (c) HERALDO experiments. (d) SEM image of the RSXS sample aperture. (e) Thin plate of the Co₈Zn₈Mn₄ fixed onto the membrane. (f) SEM image of the sample aperture and the reference slit drilled in the 1- μ m-thick gold plate for the HERALDO experiment. (g) $\cos Zn_8Mn_4$ thin plate fixed onto a gold plate by tungsten contact. The strong white/black contrast in the top/bottom parts of (d)–(g) corresponds to the Au wires sputtered onto the membrane [\[49\]](#page-11-0), which is irrelevant for this study.

membrane by means of a tungsten contact after the mask treatment [Fig. $1(e)$]. For both RSXS and HERALDO the thin plates were attached to the corresponding membranes by the single tungsten contacts [Fig. $1(g)$] to avoid the possible strain [\[50,67\]](#page-11-0).

The RSXS setup at Photon Factory, Japan was equipped with a high-vacuum chamber with a background pressure of 10[−]⁸ Torr [\[48\]](#page-11-0). The scattered intensity was collected by an in-vacuum charge-coupled device (CCD) area x-ray detector of 2048×2048 pixels (Princeton Instruments, Trenton, New Jersey, USA). The RSXS endstation MARES was used at ALBA synchrotron [\[64\]](#page-11-0). Resonant diffraction intensity was collected by a custom-designed CCD detector of 2148×2052 pixels (XCAM Co, Ltd., UK). Since small-angle scattering intensity is distributed near the transmitted direct beam, a tungsten beamstop was introduced to protect the detector for the RSXS experiment, while in the case of HERALDO the smaller aperture size allowed us to measure the holograms without using any beamstop. The magnetic field was applied parallel to the incident x-ray beam and perpendicular to the thin plate. A He-flow-type cryostat was used to control the sample temperature in a range from \sim 15 to \sim 320 K, as measured by the Si diode thermometers attached next to the sample holder and cryostat head [\[68\]](#page-11-0). A radiation shield was wrapped around the sample holder to reduce the heating of the specimen from the warm environment.

III. RESULTS AND DISCUSSIONS

The XAS signals averaged between RCP and LCP spectra near the $L_{2,3}$ edges of Mn and Co at $T = 130$ K are shown in Fig. $2(a)$. Surprisingly, despite the metallic nature of the $Co_8Zn_8Mn_4$ alloy, the Mn absorption shows a multiplet structure at the L_3 and L_2 edges. Well-resolved peaks at 1.3

and 3.5 eV above the absorption maxima at $E = 641.0$ eV and a doublet structure at the L_2 edge, which splits into two maximums at $E = 651.7$ eV and $E = 653.2$ eV, are clearly observable. Meanwhile, the Co L_3 and L_2 peak shapes at $E =$ 780.0 eV and $E = 794.1$ eV are similar to the broad spectrum of metallic Co [\[69–71\]](#page-11-0). This suggests that the fine structures of Mn $2p \rightarrow 3d$ transition should result from the localization of Mn 3*d* electrons rather than the oxidation of the surface [\[72–74\]](#page-11-0): otherwise, the multiplet structure of Co $L_{2,3}$ edges should be observed [\[75,76\]](#page-11-0). To qualitatively illustrate the features of the measured XAS, simulations were performed using the small cluster approach in the Xclaim software tool [\[77\]](#page-11-0). Mn $3d^5$ was assumed as an initial state and Mn $2p^5$ $3d^6$ was the final configuration. The simulated XAS lines were broadened with a Lorentz function with the full width Γ at half maximum (FWHM) of 1 eV. The spectrum for Co was calculated using initial $3d^8$ and final $2p^5$ $3d^9$ configurations, which reproduces the measured data despite some broadening of the latter. The observed fine structure of the XAS spectrum of Mn also nicely corresponds to the one calculated from the multiplet effects [Fig. $2(a)$].

XMCD signal measured in a magnetic field of $B = 0.5$ T well above the saturation is shown in Fig. $2(b)$ for both elements. Despite that the magnitude of the XMCD measured at the $L_{2,3}$ edges of Mn is about $1/5$ times smaller than the XMCD measured at Co $L_{2,3}$ edges, it is clear that the signs of the dichroic signals are the same. The XMCD signal at the Mn *L*² edge is notably suppressed, indicating quenching of the orbital moment. The sum rule analysis [\[78\]](#page-11-0) allows us to estimate the orbital to spin moment ratio for Mn and Co as $\mu_l(\text{Mn})/\mu_s(\text{Mn}) = 0.03$ and $\mu_l(\text{Co})/\mu_s(\text{Co}) = 0.0025$. Magnetic field dependence of the element-selective XMCD signals measured at 135 K can be found in the Supplemental Material [\[79\]](#page-11-0). The signs and magnitudes of the XMCD signals

FIG. 2. (a) XAS spectra of the bulk $Co_8Zn_8Mn_4$ near the Co and Mn *L*²,³ absorption edges: measured points are shown as the solid lines; calculated spectra are shown as the dashed lines. (b) XMCD signal. The measurements were performed at temperature $T = 130$ K at applied magnetic field $B = 0.5$ T.

indicate a ferromagnetic coupling of Co and Mn moments and a partial cancellation of Mn magnetization. This is in good agreement with the magnetization measurements, which have shown the reduction of magnetization and critical temperature with an increment of Mn concentration in $β$ -Mn-type Co-Zn-Mn compounds [\[24\]](#page-10-0). One can assume that while the Co-Co and Co-Mn couplings are ferromagnetic, the Mn-Mn interaction should be antiferromagnetic. This scenario is also suggested by a recent neutron diffraction study of the Co-Zn-Mn alloys in a wide composition range accompanied by the density functional theory (DFT) calculations [\[80\]](#page-11-0). This is in contrast to the case of Co-Mn alloys where Mn moments tend to align antiparralel to the host Co magnetization [\[81](#page-11-0)[–83\]](#page-12-0). On the other hand, the parent β -Mn compound shows strongly antiferromagnetic nearest-neighbor correlations in the 12*d* Mn sublattice, while strong ferromagnetic correlations between next-nearest neighbors were found [\[84\]](#page-12-0). Furthermore, recent DFT calculations of the β -Mn-type Co-Zn-Mn suggested larger localization of the Mn atoms at the 12*d* site than at 8*c*, being consistent with our observations [\[80\]](#page-11-0). For further quantitative discussions on the XAS and XMCD features, such as origin of multiplet spectrum of Mn, and determination

of spin and orbital contributions, additional measurements of different Co-Zn-Mn concentrations and spectra calculations from *ab initio* theory are highly desired.

Complex scattering factor *f* for resonant magnetic x-ray scattering can be described as

$$
f = (\mathbf{s} \cdot \mathbf{s}')f_c + i(\mathbf{s} \times \mathbf{s}') \cdot \mathbf{M}f_m^1 + (\mathbf{s} \cdot \mathbf{M})(\mathbf{s}' \cdot \mathbf{M})f_m^2, \quad (1)
$$

where *s* and *s'* are polarizations of the incident and scattered x rays, respectively, f_c is the charge scattering factor, M is the local magnetization, and f_m^1 and f_m^2 are factors attributed to the magnetic scattering maximized at the resonant condition. The last term in Eq. (1) containing scattering factor f_m^2 is quadratic in *M* and generally smaller than the other two terms [\[85\]](#page-12-0). Therefore, the scattering patterns $(|f|^2)$ measured at resonant conditions mainly consist of the squares of charge and magnetic scattering factors $|f_c|^2$ and $|f_m^1|^2$, and their interference $(|f_c^* f_m^1|)$. For linear polarization the measured intensities are dominated by the pure charge and magnetic scattering terms, while those measured using circular polarization contain a charge-magnetic cross term [\[86,87\]](#page-12-0). Consequently, the intensity of magnetic scattering can be simply distinguished from the charge scattering by subtracting the diffraction pattern measured (1) at off-resonant condition and appropriately normalized, (2) in the field-induced polarized state of the sample, or (3) above the critical temperature.

RSXS experiments were carried out at Mn $(E = 640.5 \text{ eV})$ and Co $(E = 779 \text{ eV})$ L_3 edges, where the magnetic scattering intensity was maximized. Far-field RSXS patterns were acquired with an exposition time of 1 s (excluding readout time ∼0.4 s) and a total of 200 expositions were averaged. In the case of the off-resonant conditions, the absorption was reduced and the measurement time was consequently reduced by a factor of 2. RSXS was measured in the temperature range between $T = 25$ K and $T = 270$ K during a field warming after field cooling (FWAFC) in $B = 70$ mT. The sample position was realigned after each temperature change to compensate for the effect of the thermal contraction (expansion) of the sample manipulator with a buffer time between the measurements (on average \approx 40 min) for stabilization. In the present experiment the linear polarization was used to minimize the influence of the charge-magnetic scattering interference term. In order to isolate the charge scattering contribution we have measured the scattering intensity at offresonance ($E = 645$ eV and $E = 785$ eV) conditions.

Typical coherent RSXS speckle patterns measured at $E =$ 779 eV at temperatures $T = 25$, 120, and 150 K are shown in Fig. [3.](#page-4-0) The missing area in the left part of each panel is due to the beamstop shadow. Tuning the energy to the Mn *L*³ edge results in the scaling of the scattering pattern due to the difference in the photons wavelength. Magnetic scattering intensity is weaker at the Mn edge, which can be explained by the lower Mn concentration in this compound, stronger absorption, and magnetization reduction, as expected from XMCD experiment. At room temperature and $T = 150$ K the small-angle scattering patterns demonstrate sixfold symmetry indicating a single-domain triangular skyrmion lattice. As it has been already shown by SANS and LTEM experiments, the skyrmion lattice phase can be supercooled by a field cooling process down to the low temperatures [\[61](#page-11-0)[,88,89\]](#page-12-0). According to the previous LTEM observations on a $\text{Co}_8\text{Zn}_8\text{Mn}_4$ thin

FIG. 3. (a) Schematic phase diagram of $\text{Co}_8\text{Zn}_8\text{Mn}_4$ and the procedure of the resonant soft x-ray scattering (RSXS) measurements. The sample is field cooled (FC) in the applied magnetic field $B = 70$ mT from room temperature down to 25 K (indicated by the blue arrow) and then warmed back to 300 K in the same field (indicated by the red arrow). Coherent RSXS speckle patterns measured for the $\rm{Co}_{8}\rm{Zn}_{8}\rm{Mn}_{4}$ sample at $E = 779$ eV corresponding to the L_3 absorption edge of Co at different temperatures (b) $T = 25$ K, (c) $T = 120$ K, and (d) $T = 150$ K and applied field *B* = 70 mT. White scale bar corresponds to 0.05 nm⁻¹.

plate, a hexagonal skyrmion lattice to amorphous state transition is reversible and accompanied by the elongation of individual skyrmions along one of the principal crystallographic axes, while the skyrmion density is conserved [\[89\]](#page-12-0).

Upon the field cooling (or FWAFC), the RSXS transforms to a homogeneous ringlike pattern corresponding to the intermediate "amorphous" phase at the temperature $T \approx 100$ K. Below $T \approx 100$ K four wide peaks appear at the coherent diffraction speckle pattern. Moreover, below the transition temperature, the skyrmion lattice parameter gradually decreases from $a_{Sk} = 112$ nm ($T = 100$ K) to $a_{Sk} = 76$ nm $(T = 25 \text{ K})$. Temperature dependence of the magnetic modulation q_{Sk} is shown in Fig. 4. q_{Sk} vector is also dependent on the magnetic field similar to the bulk case [\[61\]](#page-11-0) (inset in Fig. 4). Elongation of the q_{Sk} vector is, presumably, caused by increasing antiferromagnetic interaction between Mn ions superimposed on the helical order. Upon cooling from $T =$ 100 K to $T = 25$ K a coherent RSXS speckle pattern with fourfold symmetry can be observed. According to the previous results, the hexagonal skyrmion lattice [\[61\]](#page-11-0) is recovered by applying a stronger magnetic field of 300–500 mT at low

FIG. 4. Temperature dependence of the modulation vector of skyrmion lattice q_{Sk} . Inset shows magnetic field dependence of q_{Sk} measured at $T = 25$ K and $T = 50$ K.

temperature. Indeed, the magnitude of the q_{Sk} vector tends to shrink upon increment of the magnetic field (inset in Fig. 4). However, in the present experiment the magnetic scattering intensity arising from isolated skyrmions is hardly distinguishable in the presence of the background even when the charge scattering is subtracted. Corresponding RSXS patterns can be found in the Supplemental Material [\[79\]](#page-11-0).

The integrated speckle pattern intensities are demonstrating similar features for Co and Mn except for the difference in signal-to-noise ratio, which is better at $E =$ 779 eV. Radially integrated azimuthal profiles of the scattering patterns measured using soft x rays with energies $E =$ 779 eV and $E = 640.5$ eV can be found in the Supplemental Material [\[79\]](#page-11-0).

Fourier transform holography with an extended reference allows us to reconstruct the real-space image by single Fourier transform of the measured pattern multiplied by the linear differential operator in the direction parallel to the reference slit. Circularly polarized soft x rays with energies matching to Co and Mn *L*³ edges were used for the HERALDO experiment. Far-field diffraction patterns were collected with RCP and LCP to produce the interference pattern and reconstruct the magnetic texture. In this case information about the magnetic texture was simply encoded in the interference term $f_c^* \bar{f}_m^1$ between charge scattering arising from the reference slit and magnetic scattering from the $Co_8Zn_8Mn_4$ sample. The difference between the holograms taken with the two opposite x-ray helicities provided the isolated interference term which could be inverted to the real-space image via single Fourier transform and linear differential operator [\[12\]](#page-10-0). HERALDO patterns were acquired with an exposition time of 6 s (excluding readout time ∼0.4 s), and a total of 200 expositions were averaged, resulting in a total acquisition time of 20 min for each hologram. The sample was cooled down to the temperature $T = 20$ K, while the magnetic field of $B = 70$ mT was applied along the [001] direction parallel to the incident soft x-ray beam propagation vector.

Differences between the diffraction patterns taken with the two opposite x-ray polarizations at photon energies $E =$ 779 eV and $E = 640.5$ eV are shown in Figs. $5(a)$ and [5\(b\),](#page-5-0) respectively. The highest harmonics of the interference pattern can be found at $q_{\text{max}} = 0.2 \text{ nm}^{-1}$ which corresponds to the real-space resolution of 32 nm. The difference between

FIG. 5. Differential scattering patterns between LCP and RCP x rays of energies (a) $E = 779 \text{ eV}$ and (b) $E = 640.5 \text{ eV}$ taken at $T = 20 \text{ K}$. The color bar is given for both patterns in arbitrary units. Note that the strong linear patterns inclined upward right should be due to the incomplete subtraction of the charge scattering from the reference slit. (c) Fourier transform of (a) after applying linear differential filter and rotation of the image by 39◦. (d) and (e) Magnetic texture recorded at Co and Mn *L*³ edges, respectively. The color bar is given for both images in arbitrary units. The region outside the field of view is filled with the black background manually for clarity. (f) Fourier ring correlation analysis of the reconstructed images for Co and Mn. The orange line represents a calculated half-bit threshold and the dashed line indicates a irreversible cross point between the FRC function and half-bit threshold.

Fourier transform images taken with RCP and LCP at the Co *L*³ edge after applying the differential filter in the slit direction is shown in Fig. 5(c). The reconstructed real-space image shows the sample autocorrelation and two objectreference cross correlations delivered by both ends of the slits, as well as their complex conjugates. Magnetic contrast is inverted between the reconstructed object and conjugate. By taking into account the widths of magnetic domains where the local magnetization points antiparallel (parallel) to the applied field $B = 70$ mT at different temperatures (Fig. 6), we

FIG. 6. Real-space magnetic textures imaged at the Co L_3 edge ($E = 779$ eV) at temperatures (a) $T = 20$ K, (b) $T = 100$ K, and (c) $T = 120$ K and applied magnetic field $B = 70$ mT.

conclude that the regions with the negative (positive) values in Figs. $5(d)$ and $5(e)$ correspond to magnetization pointing antiparallel (parallel) to the field.

Figures $5(d)$ and $5(e)$ show magnification of the samplereference cross correlations taken from reconstructions of holograms taken for Co and Mn, respectively. Due to the difference of the wavelengths, the real-space image corresponding to magnetic texture of Mn ions was scaled by factor $E_1/E_2 = 779/640.5 \approx 1.22$. The magnetic structures exhibited by magnetic elements Mn and Co are similar to each other within the resolution limit. The signal-to-noise ratio is worse in the case of the measurement at the Mn L_3 edge due to the higher absorption of soft x rays at $E = 640.5$ eV and smaller concentration of Mn atoms compared to Co. Different magnitude of the magnetic moment between Co and Mn is also a reason for this. Therefore, the real-space image reconstructed for Mn atoms is slightly blurred [Fig. $5(e)$]. Additionally the reconstruction resolution is estimated from the Fourier ring correlation (FRC) [\[90\]](#page-12-0). By using FRC the spatial frequency dependence of the cross correlation of two real-space magnetic textures obtained from the reconstructions of Co and Mn HERALDO patterns [Fig. $5(f)$] is analyzed. The resolution was calculated at the point where the FRC curves irreversibly cross the half-bit threshold [\[91\]](#page-12-0). The real-space resolution that was found according to this criteria is similar to the resolution of 32 nm determined from the highest interference harmonics observed at the Fourier-space pattern.

Elongated skyrmions can be observed at the holograms taken both at $E = 779$ eV and $E = 640.5$ eV indicating a similar magnetic texture of Co and Mn sublattices, which coincides with the XMCD and RSXS data. Magnetic holograms were recorded during a FWAFC procedure up to 120 K at which the magnetic scattering is highly reduced due to the thermal shrinkage of the magnetic moments. Corresponding evolution of the magnetic texture measured at the Co absorption edge is shown in Fig. [6.](#page-5-0) The transformation of the elongated skyrmions to a more conventional shape with corresponding expansion takes place at $T = 120$ K [Fig. [6\(c\)\]](#page-5-0). Unfortunately, due to a thermal shrinkage of magnetic moments, the real-space reconstruction of magnetic texture at higher temperatures can be hardly distinguished from the background fluctuations.

Since the sample area probed by HERALDO was limited by the aperture size, we additionally performed a real-space inversion off the measured RSXS patterns by the iterative phase retrieval. Square root of the measured magnetic scattering intensity $\sqrt{I_m}$ shown in Fig. [3](#page-4-0) isolated from the charge scattering as described in Ref. [\[4\]](#page-10-0), was used as real part of the Fourier-space constraint. Fixed sample aperture size, shape, and orientation were used as real-space support. Pixels that were missing at the center of the diffraction pattern due to the beamstop shadow and subtracted charge scattering were substituted by the Fourier transform of the support. This approach is similar to the substitution used in the guided hybrid inputoutput (HIO) algorithm [\[92\]](#page-12-0). We used the implementation of the HIO algorithm [\[93\]](#page-12-0) with a feedback parameter $\beta = 0.9$ with assuming positivity and reality constraints for the realspace pattern. The phase-retrieval algorithm tended to stagnate to the local minima after 500 iterations. For each coherent scattering pattern the final real-space image was calculated

from an average of 500 algorithm trials with random initial phases. Reliability of the reconstructed real-space images was qualitatively and quantitatively examined by comparing the reconstructions performed for the data measured at Co and Mn *L*³ edges, respectively. Indeed, based on the results of the XMCD and HERALDO experiments the same orientation of the local magnetic moments of Co and Mn atoms was expected.

To quantitatively estimate the reliability of the reconstructions performed for Co and Mn we have introduced the twodimensional function

$$
R(i, j) = 1 - \frac{|\psi_{E_1}^{*(i,j)} - \psi_{E_2}^{*(i,j)}|}{|\psi_{E_1}^{*(i,j)}|},
$$

where $\psi_{E_1}^{*(i,j)}$ and $\psi_{E_2}^{*(i,j)}$ are values of (i, j) pixels of the reconstructed real-space patterns normalized by maximum value for corresponding image at $Co(E_1)$ and $Mn(E_2)$ L_3 edges. This normalization is needed to compensate for the difference in total magnetic contrast for Mn and Co patterns and transmission coefficient for the photons with different energies. Figure [7](#page-7-0) shows the real-space magnetic patterns reconstructed from data taken at the Co absorption edge at different temperatures. At $T = 25$ K and $T = 55$ K [Figs. [7\(a\)](#page-7-0) and [7\(b\)\]](#page-7-0) an array of magnetic skyrmions elongated along one of the crystallographic axes can be observed. As the temperature increases, some of the elongated skyrmions transform to the conventional circular shaped [Fig. $7(c)$], denoted as amorphous state. Unfortunately, similarly to the HERALDO experiment, the magnetic scattering intensity gradually decreases upon warming due to the thermal shrinkage of the magnetic moments. As a result, the signal-to-noise ratio is insufficient to reliably reconstruct the real-space image of the single-domain triangular skyrmion lattice at *T* > 120 K. In the Supplemental Material [\[79\]](#page-11-0) we show a few examples of the reconstructions performed for the skyrmion lattice and multidomain helical phases at higher temperatures. However, the real-space images obtained for these conditions correspond to the local minima in the solution space and vary sufficiently for the different starting phases. Two-dimensional reliability maps $R(i, j)$ for successful reconstructions can be found in the Supplemental Material [\[79\]](#page-11-0). Here we just note that the averaged reliability value $R = \frac{1}{N} \sum R(i, j)$, where *N* is the sample area in pixels, is not less than 89% for the data sets measured at 25 K.

From this aspect, the results of lensless imaging by iterative phase retrieval and HERALDO are consistent with each other. Both techniques provide a spatial resolution of 30 nm and sensitive to the signal-to-noise ratio. Data analysis in the case of coherent diffraction imaging is more sophisticated compared to HERALDO, but the latter requires advanced sample fabrication. Further attempts to improve the nanofabrication routine are required to obtain better real-space resolution.

The experimental results for the temperature- and fielddependent transformation of the skyrmion lattice obtained by soft x-ray scattering and imaging agree well with the results of Landau-Lifshitz-Gilbert (LLG) simulations calculated with realistic parameters for $\text{Co}_8\text{Zn}_8\text{Mn}_4$ thin plate by using mumax³ package $[94]$ (Fig. [8\)](#page-8-0). As hinted by present

FIG. 7. Reconstruction of coherent RSXS patterns at (a) $T = 25$ K, (b) $T = 55$ K, and (c) $T = 120$ K collected at the Co L₃ edge. The color bar for intensity (*z* scale) is given in arbitrary units. (d) Magnification of the region highlighted by the blue square in (a) is shown for the the different temperatures.

XMCD experiment and previous magnetization measurements [\[24\]](#page-10-0), the effective ferromagnetic exchange interaction in the $Co_8Zn_8Mn_4$ system may decrease with temperature due to antiferromagnetic correlations of the Mn sublattice. Moreover, manifestation of the cubic anisotropy is considerable at low temperatures. For the simulation we used exchange stiffness $A_{ex} = 9.2$ pJ/m and DMI constant $D =$ 0.00053 J/m measured experimentally by microwave spin-wave spectroscopy [\[95\]](#page-12-0), and cubic anisotropy in the form of an effective field [\[94\]](#page-12-0):

$$
\mathbf{B}_{\text{anis}} = -2K_c/M_s\{[(\mathbf{c}_1 \cdot \mathbf{m})\mathbf{c}_1][(\mathbf{c}_2 \cdot \mathbf{m})^2 + (\mathbf{c}_3 \cdot \mathbf{m})^2] + [(\mathbf{c}_2 \cdot \mathbf{m})\mathbf{c}_2][(\mathbf{c}_1 \cdot \mathbf{m})^2 + (\mathbf{c}_3 \cdot \mathbf{m})^2] + [(\mathbf{c}_3 \cdot \mathbf{m})\mathbf{c}_3][((\mathbf{c}_1 \cdot \mathbf{m})^2 + (\mathbf{c}_2 \cdot \mathbf{m})^2]\},\
$$

where $K_c = 5000 \text{ J/m}^3$ is the first-order cubic anisotropy constant determined from electron spin resonance (ESR) experiment $[96]$, \mathbf{c}_1 , \mathbf{c}_2 , and \mathbf{c}_3 is a set of mutually perpendicular unit vectors indicating the anisotropy directions (cubic axes), and $M_s = 350 \text{ kA/m}$ is the saturation magnetization [\[95\]](#page-12-0). A two-dimensional 200-nm-thick plate with area of $10 \times$ $10 \ \mu m^2$ with open boundary conditions and elementary cell size $5 \times 5 \times 200$ nm³ was simulated. To follow the experimental field-cooling protocol, we started the simulation with random initial spin configuration in the applied field along [001] direction. The field needed for robust SkX formation was set to $B = 150$ mT. The discrepancy between the observed and calculated field values is, presumably, caused by demagnetization effects. The characteristic helical pitch $\lambda =$ $4\pi A_{\rm ex}/D$ and skyrmion size are determined by the ratio of the exchange stiffness *A*ex and the Dzyaloshinskii constant *D* [\[29\]](#page-10-0). Therefore it is reasonable to assume two possible scenarios: (1) the temperature-dependent *q*-vector variation is caused by changing exchange integral due to the antiferromagnetic correlations in Mn sublattice towards lower temperatures; and (2) the emergence of the *q*-vector elongation can be also caused by enhancement of antisymmetric DMI due to the local noncomplanar structure of frustrated Mn. At the moment, the experimental data does not allow us to unambiguously distinguish between variations of *A*ex and *D*. To mimic the first scenario, we reduced the exchange stiffness parameter *A*ex from the measured value 9.2 to 3.2 pJ/m in a linear fashion, while the magnetic field $B = 150$ mT remained constant. Relaxation time of 10 ns was introduced between each step for magnetic texture stabilization. Upon gradual decrease of the exchange interaction in the system, the skyrmion lattice exhibits deformation similar to the LTEM and soft x-ray experiments [Figs. $7(a)$ – $7(c)$]. Elongation of the skyrmions is manifested due to the overall reduction of the exchange interaction, while its directionality along the cubic axes is dictated by the cubic anisotropy. Notably, this deformation takes place via directional expansion of each topological vortex, but not by merging of the neighboring skyrmions—in the latter case the total topological charge of the system would decrease, which is opposite to the topological charge conservation scenario reported earlier [\[89\]](#page-12-0). Radial averages of the two-dimensional fast Fourier transformation (FFT) patterns of out-of-plane projection of magnetization distribution was used to calculate the *q*-vector magnitude dependence for the resultant magnetic textures. The simulated variation in q_{Sk} induced by change in the exchange stiffness parameter *A*ex is in very good qualitative agreement with the temperature dependence (Fig. [9\)](#page-8-0). The shoulder in Fig. [9](#page-8-0) at *A*ex between 5 and 6 pJ/m corresponds to the smooth transition from the

FIG. 8. Micromagnetic simulations of Co₈Zn₈Mn₄ 10 × 10 μ m² thin plate: *z* projection of the magnetic texture is shown for various exchange from stiffness parameters $A_{ex} = 9.2 \times 10^{-12}$ J/m (a), 5.8×10^{-12} J/m (b), and 3.2×10^{-12} J/m (c). Bottom panels show magnification of the rectangular regions highlighted by the corresponding red boxes.

disk shaped to elongated skyrmions. Therefore we assume that the linear decrease of the exchange interaction in $\text{Co}_8\text{Zn}_8\text{Mn}_4$

takes place with decreasing temperature. In general, this consideration is consistent with the previous studies of the β -Mn alloys that have shown the presence of antiferromagnetic correlation of moments in the 12*d* Mn sublattice [\[80](#page-11-0)[,84,97–99\]](#page-12-0). We plan to directly address this question in our future studies. For example, recently developed spin-wave sensitive SANS technique allows us to directly probe *A*ex of a bulk specimen as a function of temperature [\[100\]](#page-12-0).

LLG simulation was used to probe the magnetic field evolution of the elongated skyrmion texture. The "field-cooled" magnetic texture shown in Fig. [10\(c\)](#page-9-0) with $A_{\rm ex} = 3.2$ pJ/m was used. Except for the discrepancy between the field magnitude values, the result is consistent with present x-ray and previous LTEM experiments—restoration of the hexagonal lattice of circular skyrmions from the deformed state by ramping the magnetic field was successfully reproduced [Figs. $10(a)$ – $10(f)$].

IV. CONCLUSION

In conclusion, by means of element-selective soft x-ray circular magnetic dichroism we have revealed the ferromagnetic arrangement of Co and Mn ions in a

FIG. 9. Measured dependence of q_{Sk} over temperature (symbols) and calculated dependence of q_{Sk} over the exchange stiffness (A_{ex}) parameter.

FIG. 10. Magnetic field evolution of the *z* projection of magnetization derived from the micromagnetic simulation of Co₈Zn₈Mn₄ thin plate in magnetic field $B = 150$ mT (a), $B = 250$ mT (b), $B = 300$ mT (c), $B = 350$ mT (d), $B = 400$ mT (e), and $B = 450$ mT (f) showing the hexagonal SkX restoration.

room-temperature skyrmion-hosting compound Co₈Zn₈Mn₄. Moreover, by using the coherent resonant small-angle soft x-ray scattering and holography with extended reference we can conclude that the topological magnetic texture is the same for both types of atoms in whole temperature range above T_g that is reliable for real-space reconstruction. Our results are consistent with each other and with the previous neutron scattering and Lorentz microscopy experiments and shows the transition from hexagonal skyrmion crystal to elongated skyrmion state that is accompanied by deformation of the individual vortices. Micromagnetic simulation suggests that such transition is driven by decreasing exchange interaction in the system and effect of the cubic anisotropy. This effective decrease of the ratio of symmetric exchange interaction to antisymmetric Dzyaloshinskii-Moriya mimics low-temperature antiferromagnetic frustration of the Mn sublattice. At lower temperature, antiferromagnetic correlations of Mn atoms is superimposed onto the long-range helical (skyrmion) modulation, resulting in shortening of the helical pitch and deformation of skyrmions. However, this effect is reversible and the hexagonal skyrmion lattice from elongated skyrmion state can be restored by increasing magnetic field even when the exchange stiffness is reduced, as learned from the micromagnetic simulation and previous experiments [\[61](#page-11-0)[,89\]](#page-12-0).

We have demonstrated lensless soft x-ray imaging of the magnetic texture at cryogenic temperatures and applied magnetic fields. HERALDO imaging was used with the circularly polarized soft x rays, while coherent diffraction imaging was performed with a linearly polarized beam. Both methods did not require focusing x-ray optics to perform magnetic imaging with resolution of a few tens of nanometers. Practically, the signal-to-noise ratio sufficient for the successful reconstruction was achieved only in the temperature range from $T =$ 20 K to $T = 120$ K due to the overall decay of the intensity of the magnetic scattering and charge-magnetic interference with increasing temperature.

Soft x-ray imaging methods allow us to simultaneously obtain element-selective real-space information and will be useful for further investigations of nontrivial magnetic textures in thin plates of polar magnets, since Néel-type skyrmions produce no contrast in Lorentz transmission electron microscopy.

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