

Zero-Dimensional Spin Accumulation and Spin Dynamics in a Mesoscopic Metal Island

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We have measured electron spin accumulation at 4.2 K and at room temperature in an aluminum island with all dimensions ($400 \text{ nm} \times 400 \text{ nm} \times 30 \text{ nm}$) smaller than the spin relaxation length. For the first time, we obtain uniform spin accumulation in a four-terminal lateral device with a magnitude exceeding the Ohmic resistance in the island. By controlling the magnetization directions of the four magnetic electrodes that contact the island, we have performed a detailed study of the spin accumulation. Spin precession measurements confirm the uniformity of our system and provide an accurate method to extract the spin relaxation time.

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What happens if we inject spin-polarized carriers into a small nonmagnetic island? This is an outstanding question in the rapidly developing field of spintronics [1]. In metallic systems with submicrometer dimensions, still many (of the order of 10^7 – 10^8) electron spins are involved. They will behave uniformly if the dimensions of the island are smaller than the spin relaxation length, $\lambda_{sf} = \sqrt{D\tau_{sf}}$, where D is the diffusion constant and τ_{sf} the spin relaxation time. We will show that, in this regime, the spin accumulation dynamics depends only on τ_{sf} and becomes independent of the transport properties such as D . Previous studies on electrical spin injection and detection have focused on four-terminal devices larger than the spin relaxation length [2–4] or in two-terminal pillar structures [5].

Here we report the study of the injection of spins in an aluminum island fabricated with *all* dimensions smaller than the spin relaxation length λ_{sf} . The island is weakly coupled to the four cobalt leads by means of tunnel barriers [6]. We will show that, to first order, the system is zero dimensional with respect to the spin and the induced spin polarization in the island is uniform. These conditions correspond to the regime $\tau_{\text{diff}} < \tau_{sf} \ll \tau_{\text{esc}}$, where τ_{diff} is the time for the electron to diffuse in the island and τ_{esc} the time to escape into the cobalt leads. In addition, there are two other (position-dependent) contributions to the chemical potential of the island, smaller than the spin accumulation, arising from the charge current (Ohmic resistance) and the spin current.

In such a system, spin accumulation can be described in an elementary way and becomes the result of two competing processes: the injection of spins, and their dynamics and relaxation mechanisms. The injection into the aluminum island is obtained by driving a current in and out of two (e.g., electrode Co1 and Co2) of the four cobalt electrodes that contact the island (see Fig. 1). Each current electrode i carries a charge current I and a spin current $\mathbf{I}_{m,i} = PI\mu_B\mathbf{m}_i/e$, where \mathbf{m}_i is the magnetization direction of the electrode. $P = (G^\uparrow - G^\downarrow)/(G^\uparrow + G^\downarrow)$ is the spin injection/detection efficiency of the tunnel bar-

rier, with G^\uparrow, G^\downarrow the tunnel conductances for the up and down spins, where up means oriented in the same direction as \mathbf{m}_i [2]. The spin relaxation mechanisms drive the out-of-equilibrium magnetization inside the island back to equilibrium at a rate τ_{sf}^{-1} .

The electrical detection of the spin imbalance is performed by using the two remaining cobalt electrodes, in this case Co3 and Co4. The signal detected by the electrode i has a spin independent contribution $\mu_0(x) = \int f_0(\epsilon, x) d\epsilon$ and a spin contribution $P\mathbf{m}_i \cdot \boldsymbol{\mu}$, with $\boldsymbol{\mu} = \int \mathbf{f}(\epsilon) d\epsilon$, where f_0 and \mathbf{f} are, respectively, the spin independent and spin dependent distribution functions [8].

The spin accumulation contribution to the total signal in the 0D case depends on the injecting vector $\mathbf{m}_{\text{inj}} = \mathbf{m}_1 - \mathbf{m}_2$, the detecting vector $\mathbf{m}_{\text{det}} = \mathbf{m}_3 - \mathbf{m}_4$, and the volume of the island \hat{V} :

$$R_s = \frac{V}{I} = \frac{P^2\tau_{sf}}{e^2\nu_{\text{DOS}}\hat{V}}\mathbf{m}_{\text{inj}} \cdot \mathbf{m}_{\text{det}}, \quad (1)$$

where $\nu_{\text{DOS}} = 2.4 \times 10^{28} \text{ eV}^{-1} \text{ m}^{-3}$ is the aluminum density of states at the Fermi energy. For collinear electrodes,

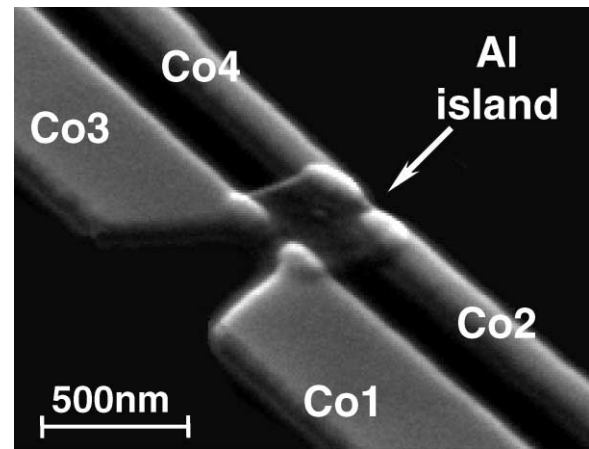


FIG. 1. Scanning electron microscope picture of a device. The square aluminum island in the middle is contacted by four cobalt electrodes of different widths [7].

this reduces to the formula obtained by Johnson [9]. Spin accumulation occurs *and* it is detected only if \mathbf{m}_1 and \mathbf{m}_2 are not parallel with each other and if \mathbf{m}_3 and \mathbf{m}_4 are also not parallel.

The devices are defined by electron beam lithography and a two-angle shadow mask technique. A square aluminum island $[(400 \times 400 \times 30) \text{ nm}^3]$ is deposited by *e*-beam evaporation at a base pressure of 1×10^{-6} mbar, through a suspended mask, followed by oxidization in pure oxygen ($2\text{--}10 \times 10^{-2}$ mbar for 1–5 min) to produce tunnel barriers with resistances in the range $R_{\text{TB}} = 1\text{--}40 \text{ k}\Omega$. Next, four cobalt electrodes 40 nm thick are deposited from a different angle to contact the island. The electrodes have different widths, one pair 500 nm wide and one pair 100 nm, with the widest one having the lowest coercive field. Owing to the magnetic shape anisotropy, the electrode's magnetization lies in the plane of the substrate, pointing in the positive or negative \hat{y} direction. By applying an in-plane external magnetic field, we can independently flip the magnetization of the electrodes. We identify a “parallel” and an “antiparallel” configuration [see Fig. 2(b)]. From the measurements, we conclude that a third, “anomalous” magnetic configuration also occurs, in which the two wide electrodes and one of the narrow ones are aligned, while the fourth one, probably due to a slightly different coercive field, is opposite.

The three possible electrical measuring configurations are depicted in Fig. 2(a): The current I is sent from I^+ to I^- , the detected voltage is $V = V^+ - V^-$, and the signal we plot is $R = V/I$ [10].

To describe spin transport and spin accumulation, we assume for the moment that magnetization direction in the island is collinear with the electrodes, with \uparrow directed in the positive \hat{y} direction. Then the island spin dependent chemical potentials are represented in terms of $\mu_{\uparrow,\downarrow}$ [8,11,12]. Figure 2(c) gives a schematic picture of the chemical potentials and the voltage contacts. In all three cases, the potential drop given by the charge current is $\Delta\mu_0 = eR_{\text{Ohm}}I$, where R_{Ohm} is the island four-terminal Ohmic resistance.

For the spin contribution, we analyze the three cases separately. In the antiparallel configuration, the spin accumulation signal is given by Eq. (1). In the parallel configuration, no net (average) spin accumulation occurs. However, spin-polarized carriers injected at Co1 and extracted at Co2 give rise to a the spin current of magnitude $|\mathbf{I}_m| = PI\mu_B/e$ that traverses the system, causing the spin polarization to be space dependent. It can be shown that this gives a contribution $\delta R = P^2R_{\text{Ohm}}$. This can be understood by considering that one of the two spin channels is partially used and the total conductance of the island decreases. In the limiting case of $P = 1$, the total conductance would halve. In the anomalous configuration, the above contribution to the resistance cancels, as can be seen in Fig. 2(c), and only the Ohmic resistance is detected.

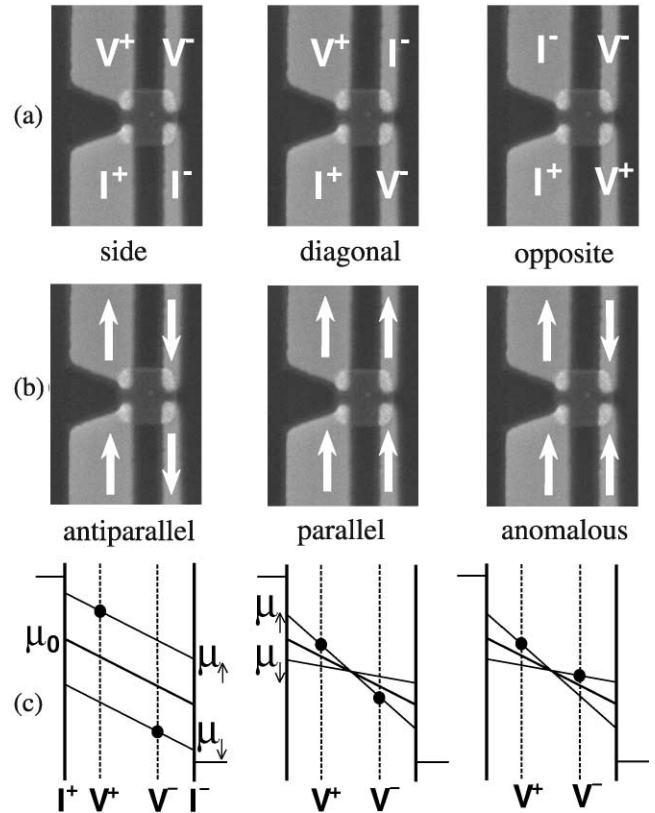


FIG. 2. (a) The three possible independent measuring configurations: Current is sent from I^+ to I^- , the measured voltage V is $V^+ - V^-$. (b) The possible magnetic configurations and (c) the corresponding chemical potentials for the spin up and down inside the island for the “side” configuration: The thick line is the average chemical potential μ_0 , the two thin lines $\mu_{\uparrow,\downarrow}$. The black dots indicate the potential measured by the V^+ and V^- probes for the case $P = 1$. In the parallel and anomalous configurations, an equal spin current density $|\mathbf{j}_m| = -\sigma_N\mu_B(\nabla\mu_{\uparrow} - \nabla\mu_{\downarrow})/(2e)$ flows through the island.

The measurements were performed by standard low frequency lock-in techniques with a modulation current of 10–100 μA . We have measured six devices in detail, one only at 4.2 K, two both at 4.2 K and at room temperature, and three at RT only. For the last three, we have also performed precession measurements (discussed later).

Figure 3 shows measurements for the three configurations at 4.2 K in device A, with all tunnel barriers having 20 $\text{k}\Omega$ resistance [13]. The magnetic field is applied in the \hat{y} direction. We start with the field at -100 mT , so that the electrode magnetizations are aligned in the negative \hat{y} direction. Ramping the field to positive values, we observe a sudden increase of the signal at $+30 \text{ mT}$, when the magnetization direction of the widest pair reverses. The magnetic configuration is now antiparallel, spin accumulation occurs, and the four-terminal resistance is enhanced. When the second pair of electrodes also switches at $+60 \text{ mT}$, the magnetization configuration is again parallel but with all

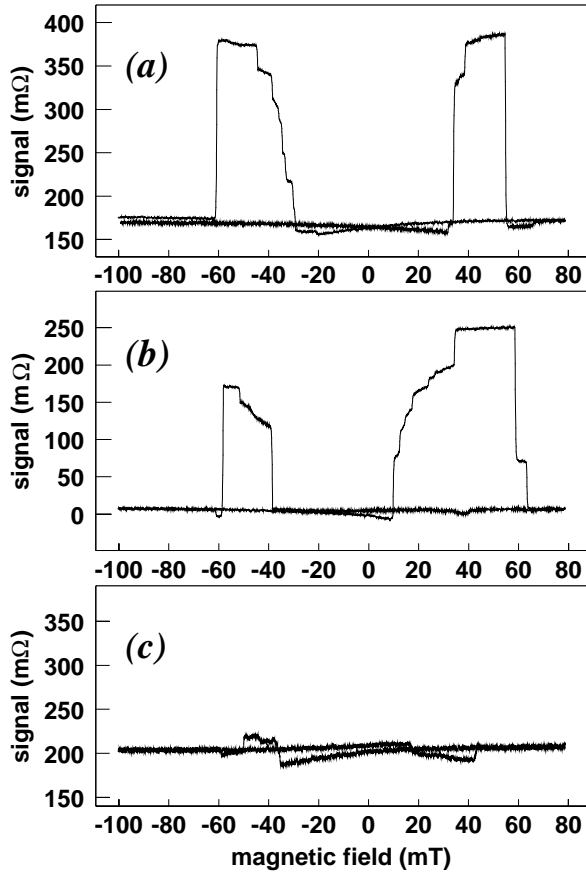


FIG. 3. Measurements of $R = (V^+ - V^-)/I$ as a function of the in-plane magnetic field at 4.2 K, in the (a) “side,” (b) “diagonal,” and (c) “opposite” configurations.

magnetizations pointing in the positive \hat{y} direction, and the signal drops again. For the “side” configuration, the spin accumulation signal is 220 mΩ and it is slightly larger, 250 mΩ, for the “diagonal” one.

The steps visible on the measurements at the switching of the larger electrodes are interpreted as the steplike reversal of the electrodes magnetization, and therefore as discrete changes of the injector and detector vectors, \mathbf{m}_{inj} and \mathbf{m}_{det} . Note that, in Fig. 3(b), the left peak does not reach full height. This is due to the incomplete reversal (of one) of the wide electrodes (probably due to domain formation). In this case, the fully antiparallel configuration is not reached. This mostly occurs at 4.2 K.

Graph 3(c) shows no spin signal for the “opposite” configuration: The two larger electrodes are always parallel as they flip at the same time. A similar behavior was also observed for a device with tunnel barrier resistances between 15 and 35 kΩ and for a device with tunnel barriers of 3–5 kΩ.

These results are consistent with the assumption of an almost uniform spin accumulation throughout the island [14]. Note also that the signal is about 20 times larger than the signal reported by Jedema *et al.* in a 1D geometry [4] and comparable with the two-terminal signal in

the pillar structures used to study the spin current induced magnetization reversal [5].

Figure 4(a) shows a measurement in the side configuration at room temperature for device B with tunnel barriers of 2 kΩ. Here the spin signal is $R_s = 60$ mΩ. The measurement presents a new feature around 70 mT: While switching from antiparallel to parallel, the signal dips $\delta R \approx 5$ mΩ below the signal in the parallel configuration. This can be explained by assuming an anomalous configuration, see Fig. 2(c), where only one narrow electrode has reversed. The detected signal is the lowest and equals the Ohmic resistance R_{Ohm} . At a higher field of 120 mT, the other narrow electrode also flips, returning to the parallel configuration, and the signal increases by $\delta R = P^2 R_{\text{Ohm}}$. We thus obtain a coarse estimation of $P \approx 16\%$. Another two devices with the same tunnel barrier resistances showed similar behavior and gave the same spin signal amplitude. A device with higher tunnel barriers showed a room temperature spin signal of 90 mΩ.

To accurately determine the spin relaxation time, we measured the precession of the injected spins under a magnetic field \mathbf{B} applied perpendicular to the device, in the positive \hat{z} direction. The component of the spins perpendicular to \mathbf{B} precesses with the Larmor angular frequency $\omega_L = g\mu_B|\mathbf{B}|/\hbar$, with $g \approx 2$ for aluminum

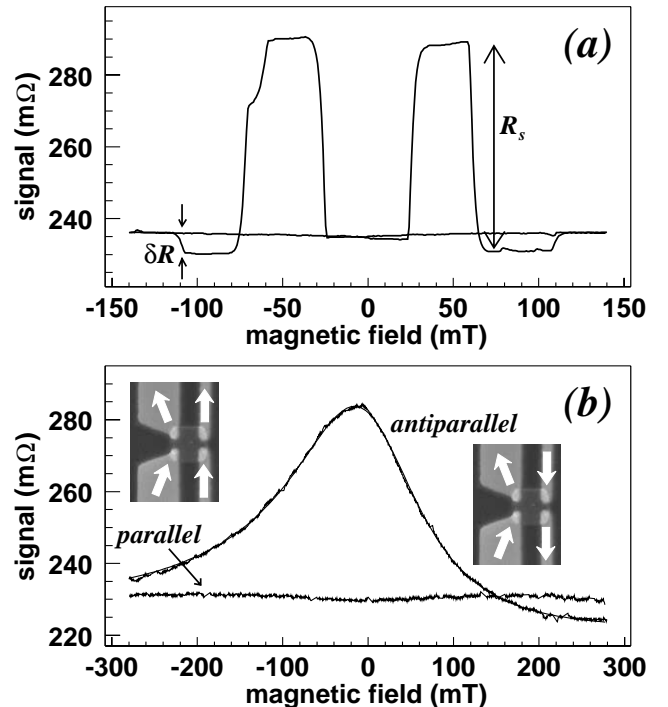


FIG. 4. (a) Spin signal at 300 K. The dip that appears at 70 mT is due to the “anomalous” magnetic configuration. (b) Spin precession for the same device in the parallel and antiparallel configurations, in the side configuration. The fitted curve fits the measurement closely. The insets represent the direction of the electrodes’ magnetization in the noncollinear case, i.e., assuming an angle ϕ between injector and detector.

[2,4]. To derive the spin signal, we use an approach similar to that of Johnson and Jedema, but now derived explicitly for the 0D geometry [11].

Let us assume \mathbf{m}_{inj} , \mathbf{m}_{det} perpendicular to \mathbf{B} and ϕ the angle between them. At $t = 0$, spins parallel to \mathbf{m}_{inj} are injected in the island and precess. The contribution to the detected signal at a later time t is proportional to $\exp(-t/\tau_{sf}) \cos(\omega_L t + \phi)$, where the exponential factor accounts for spin flip scattering. Integrating over the possible times $[0, +\infty)$, the spin accumulation signal becomes

$$R_s = \frac{P^2 \tau_{sf}}{e^2 \nu_{\text{DOS}} \hat{V}} \frac{\cos \phi - \omega_L \tau_{sf} \sin \phi}{1 + \omega_L^2 \tau_{sf}^2} |\mathbf{m}_{\text{inj}}| |\mathbf{m}_{\text{det}}|, \quad (2)$$

a linear combination of an even and an odd function in the field. This equation reduces to Eq. (1) for $|\mathbf{B}| = 0$.

In the experiment, we apply first a magnetic field in the $\hat{\mathbf{y}}$ direction to set the leads' magnetic configuration to either parallel or antiparallel. Then, with the $\hat{\mathbf{y}}$ field switched off, we measure the spin signal as a function of the perpendicular field. The resulting spin precession is shown in Fig. 4(b) for device B. In the parallel configuration, only a small dependence of the signal on the \mathbf{B} field is detected. In the antiparallel case, the spin accumulation reaches a maximum at -20 mT and decays asymmetrically. We fit the spin signal with Eq. (2) to which we have added a constant term for the background Ohmic resistance. The fitted curve, Fig. 4(b), is superimposed on the measurement, and agrees very well with the experimental data: We obtain $\tau_{sf} = 62$ ps at room temperature and $\phi = 0.13\pi$. The latter, we believe, reflects the fact that the tips of the larger electrodes have a triangular shape and the end domains are not exactly magnetized along the $\hat{\mathbf{y}}$ direction [see Fig. 4(b) insets]. Assuming that only the wide electrode magnetization is rotated by ϕ , $|\mathbf{m}_{\text{inj}}| = |\mathbf{m}_{\text{det}}| = 2 \cos(\phi/2)$, we find $P = 7\%$. These values agree with the results of Jedema *et al.* [4]. Note that, taking into account the detection efficiency P of the component of the signal, the spin accumulation is $(\mu_{\uparrow} - \mu_{\downarrow})/eI = R_s/P = 60 \text{ m}\Omega/7\% = 850 \text{ m}\Omega$, thus dominating the Ohmic resistance. Using the diffusion constant for aluminum at room temperature $D = 5 \times 10^{-3} \text{ m}^2/\text{s}$, the diffusion time $\tau_{\text{diff}} = L^2/D \approx 30$ ps [15], and the escape time $\tau_{\text{esc}} = R_{\text{TB}} e^2 \nu_{\text{DOS}} \hat{V} \approx 10^3 \tau_{sf}$. This shows that the relation $\tau_{\text{diff}} < \tau_{sf} \ll \tau_{\text{esc}}$ is satisfied.

In conclusion, we have measured zero-dimensional spin accumulation in a mesoscopic aluminum island at 4.2 K and at room temperature and also coherent spin precession. We have observed three contributions to the total detected signal: the overall spin accumulation (being the largest), the Ohmic resistance, and the effect of the spin current. From the precession measurements, we have determined the spin relaxation time and the orientation of the magnetic leads. The control of the spin accumulation

in the dc regime that we have demonstrated opens the way to the study of the island's magnetization dynamics with time dependent spin injection in the radio-frequency regime.

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 - [15] The spin relaxation length $\lambda_{sf} = \sqrt{D\tau_{sf}} = 550 \text{ nm}$.