Nonequilibrium phenomena in multiple normal-superconducting tunnel heterostructures

J. Voutilainen,¹ T. T. Heikkilä,^{1,2} and N. B. Kopnin^{1,3}

¹Low Temperature Laboratory, Helsinki University of Technology, P.O. Box 2200, FIN-02015 HUT, Finland

²Department of Physics and Astronomy, University of Basel, Klingelbergstr. 82, CH-4056 Basel, Switzerland

³L. D. Landau Institute for Theoretical Physics, 117940 Moscow, Russia

(Received 8 March 2005; revised manuscript received 28 April 2005; published 3 August 2005)

Using the nonequilibrium theory of superconductivity with the tunnel Hamiltonian, we consider a mesoscopic NISINISIN heterostructure, i.e., a structure consisting of five intermittent normal-metal (N) and superconducting (S) islands separated by insulating tunnel barriers (I). Applying the bias voltage between the outer normal electrodes one can drive the central N island very far from equilibrium. Depending on the resistance ratio of outer and inner tunnel junctions, one can realize either effective electron cooling in the central N island or create highly nonequilibrium energy distributions of electrons in both S and N islands. These distributions exhibit multiple peaks at a distance of integer multiples of the superconducting chemical potential. In the latter case the superconducting gap in the S islands is strongly suppressed as compared to its equilibrium value.

DOI: 10.1103/PhysRevB.72.054505

PACS number(s): 74.45.+c, 73.23.-b, 74.78.-w

I. INTRODUCTION

Mesoscopic electronic applications typically rely on phenomena which show best when the electrons in small wires are cooled to very low temperatures, ideally to the range of 10 mK. In this regime the crystal lattice is very weakly coupled to the electron system, and electron cooling via the lattice becomes difficult. An alternative approach is then to directly cool the electrons. This can be achieved by placing superconducting (S) contacts via insulating (I) barriers to the normal-metal (N) or superconducting (S') wire whose electrons are to be cooled.¹⁻³ By applying a voltage over such SINIS/SIS'IS coolers, it has been shown that one can cool electrons well below 100 mK with these structures,⁴ even when the lattice remains at a few hundred mK, or to enhance the superconductivity in the middle S' island.^{5–7} Optimally, such cooling should take the electron temperature to a few mK, a limit which is hardly reached in mesoscopic systems via other known means.

With this type of nonequilibrium cooling, the concept of the electron temperature is not always well defined,⁴ but one has to rather describe the full electron energy distribution function.^{8,9} In this case cooling corresponds to the sharpening of this distribution function, essentially removing the high-energy excitations.

One of the features limiting the performance of such SINIS coolers is the fact that the poor heat conductivity of the superconductors makes them inefficient reservoirs.¹⁰ An additional pair of normal-metal electrodes attached to superconducting electrodes of a SINIS cooler can improve the relaxation and enhance the cooling characteristics of the device.³ In this paper, we consider the effects of extra N electrodes of this type attached to a generic SINIS structure. The superconducting pieces are now assumed small enough so that they can be driven out of equilibrium by applying a bias voltage between the two normal electrodes. It is this arrangement of a NISIN'ISIN multiple heterostructure which is the subject of the present study. Its novel feature as compared to the traditional SINIS structure with bulk S elec-

trodes is that nonequilibrium is now induced in all inner islands of the structure, which, in turn, strongly enhances nonequilibrium effects both in the central N and in the adjacent S islands. The resulting distributions in each island can be inspected by transverse probe tunnel junctions.^{4,8}

The microscopic nonequilibrium theory of double-barrier SINIS junctions is based on time-dependent Keldysh Green function formalism (see Ref. 11 and references therein). In the present paper we extend the theory such that nonequilibrium distributions in superconducting islands are also allowed. The major modification is that the chemical potentials of the two superconducting islands cannot be chosen zero simultaneously since these islands have different potentials determined by the bias voltage. This is equivalent to a time dependence of the order parameters imposed by different time-dependent order-parameter phases. We restrict ourselves to a tunneling Hamiltonian approximation which effectively makes the problem spatially independent within each superconducting or normal island.

II. MODEL

The system we study is shown in Fig. 1. We use the tunnel Hamiltonian approach and neglect the proximity effects in both normal and superconducting islands. In nonequilibrium, each of the islands has a separate energy distribution. According to the tunnel Hamiltonian model, they should be



FIG. 1. NISINISIN structure: The central normal island (N) is placed between two superconducting islands (S) which are connected to two outer N electrodes serving as external leads. Each connection is realized through a tunnel contact with a resistance R_{ik} .

independent of both coordinates and directions of the momenta. This implies that each island is in a diffusive regime when the momentum direction dependence is averaged out within the first approximation in ℓ/L where ℓ is the impurity mean free path and *L* is the length of the contacting island. In addition, one has to assume that the intrinsic normal-state resistance *r* of each island is much smaller than any of the tunnel resistances *R* to satisfy the condition that the potential variation inside each island is smaller than its drop across the tunnel barrier.

A characteristic rate for tunneling from island 2 into island 1 to be defined later is $\eta_{12} = (4\nu_1 e^2 \Omega_1 R)^{-1}$, where ν_1 is the normal-state density of states (DOS) at the Fermi level in island 1 and Ω_1 is the volume of conductor 1. Competing with the inelastic relaxation, this rate determines whether the injection of new quasiparticles into the system is fast enough to drive the system out of equilibrium. The inelastic processes which uphold the thermal Fermi distribution can be neglected if $\eta \ge 1/\tau_{inel}$ which is equivalent to

$$\nu e^2 \Omega R / \tau_{\text{inel}} \sim (L \ell / \ell_{\text{inel}}^2) \Gamma \ll 1$$
 (1)

where $\Gamma = (N_m R/R_Q)(\overline{A}/A)$. Here \overline{A} stands for the average cross-section area of the conductor, L is its length such that $\Omega = \overline{A}L$, A is the area of the junction, $\ell_{\text{inel}} = \sqrt{D\tau_{\text{inel}}}$, and $D = v_F \ell/3$ is the diffusion coefficient. We also introduce the quantum resistance $R_Q = \pi \hbar/2e^2$, and the number of conducting modes $N_m \sim p_F^2 A/\hbar^2$ in the area of the junction.

In our work, we assume that the electron energy distribution within each island is homogeneous, which also implies a homogeneous potential and a constant effective temperature. Analysis of kinetic equations shows that this is achieved if $r \ll R$ or $\sigma \overline{A}/L \sim \nu D e^2 \overline{A}/L \gg 1/R$, which gives

$$DL^{-2} \gg (\nu e^2 R\Omega)^{-1} \sim \eta.$$

This puts a restriction $L \ll \ell \Gamma$. The limit $\ell \ll L$ holds if $\Gamma \gg 1$. Our model is thus applicable provided $1 \ll L/\ell \ll \Gamma$. Condition (1) of small inelastic interaction reads $\ell_{inel}^2 \gg \Gamma L \ell$.

III. FORMALISM

A. Transport equation

We use the standard Keldysh Green function formalism (see for example Refs. 12 and 13). We denote the Nambuspace matrix operator

$$\hat{G}^{-1} = \hat{\tau}_3 \frac{\partial}{\partial \tau} - \frac{\nabla^2}{2m} - \mu + \hat{H}, \quad \hat{H} = \begin{pmatrix} 0 & -\Delta \\ \Delta^* & 0 \end{pmatrix}.$$

It acts on matrix Green functions

$$\hat{G} = \begin{pmatrix} G & F \\ -F^{\dagger} & \overline{G} \end{pmatrix},$$

where \hat{G} stands for either retarded (advanced), $\hat{G}^{R(A)}$, or Keldysh, \hat{G}^{K} , Green function.

Introducing the total self-energy that contains tunneling self-energy together with both elastic and inelastic processes we can write equations for the retarded, advanced, and Keldysh Green functions in each island:

$$\begin{split} (\hat{G}^{-1} - \hat{\Sigma}^{R(A)}) \circ \hat{G}^{R(A)} &= \hat{1} \,\delta(x_1 - x_2), \\ (\hat{G}^{-1} - \hat{\Sigma}^R) \circ \hat{G}^K - \hat{\Sigma}^K \circ \hat{G}^A &= 0. \end{split}$$

The product $\hat{\Sigma}^{K_{\circ}}\hat{G}^{A}$ is a convolution over frequencies and momenta of the type

$$\hat{\Sigma}^{K} \circ \hat{G}^{A} = \int \hat{\Sigma}^{K}_{\epsilon_{1},\epsilon} \hat{G}^{A}_{\epsilon,\epsilon_{2}} \frac{d\epsilon}{2\pi}$$

The transport-like equations can be obtained in a standard way by applying the inverse operator to $\hat{G}^{R(A)}$ and \hat{G}^{K} from the right and subtracting the obtained equations from the equations above. Next we average the transport equations for the Green functions $\hat{G}(\mathbf{p}+\mathbf{k}/2,\mathbf{p}-\mathbf{k}/2)$ over the directions of the momentum \mathbf{p} keeping in mind that, in the tunnel approximation discussed in Sec. II, the Green functions are isotropic with respect to \mathbf{p} and are independent of the centerof-mass coordinate, i.e., they only have a component with $\mathbf{k}=0$. Therefore, the transport-like equations become

$$-\epsilon_{1}\hat{\tau}_{3}\hat{G}^{K}_{\epsilon_{1},\epsilon_{2}}+\hat{G}^{K}_{\epsilon_{1},\epsilon_{2}}\epsilon_{2}\hat{\tau}_{3}+\hat{H}\circ\hat{G}^{K}-\hat{G}^{K}\circ\hat{H}=\hat{\mathcal{I}}^{K}_{\epsilon_{1},\epsilon_{2}}, \quad (2)$$

$$-\epsilon_{1}\hat{\tau}_{3}\hat{G}^{R(A)}_{\epsilon_{1},\epsilon_{2}}+\hat{G}^{R(A)}_{\epsilon_{1},\epsilon_{2}}\epsilon_{2}\hat{\tau}_{3}+\hat{H}\circ\hat{G}^{R(A)}-\hat{G}^{R(A)}\circ\hat{H}=\hat{\mathcal{I}}^{R(A)}_{\epsilon_{1},\epsilon_{2}}. \quad (3)$$

Now \hat{G} stands for the function averaged over directions of **p**. The collision integrals in each island are

$$\begin{split} \hat{\mathcal{I}}_{\epsilon_1,\epsilon_2}^K &= \hat{\Sigma}^R \circ \hat{G}^K - \hat{G}^K \circ \hat{\Sigma}^A - \hat{G}^R \circ \hat{\Sigma}^K + \hat{\Sigma}^K \circ \hat{G}^A, \\ \\ \hat{\mathcal{I}}_{\epsilon_1,\epsilon_2}^{R(A)} &= \hat{\Sigma}^{R(A)} \circ \hat{G}^{R(A)} - \hat{G}^{R(A)} \circ \hat{\Sigma}^{R(A)}. \end{split}$$

The elastic processes drop out after averaging over momentum directions.

For tunneling between two superconductors (or normal metals) 1 and 2, the self-energy in island 1 is $\hat{\Sigma}_T(1) = i\eta_{12}\hat{g}(2)$ where the quasiclassical Green functions integrated over the energy $\xi_p = p^2/2m - E_F$ are

$$\hat{g} = \int \frac{d\xi_p}{\pi i} \hat{G}.$$

The quasiclassical functions satisfy the Eliashberg and Usadel equations that can be obtained from Eqs. (2) and (3), respectively, after integration over $d\xi_p$. The tunneling rate is defined as

$$\eta_{12} = \pi \nu_2 \Omega_1^{-1} \langle |T_{\mathbf{p},\mathbf{q}}|^2 \rangle = (4 \nu_1 e^2 \Omega_1 R_{12})^{-1}.$$
 (4)

Here $T_{\mathbf{p},\mathbf{q}}$ is the tunnel matrix element and R_{12} is the tunnel resistance of the contact between islands 1 and 2. Note that the self-energy in island 2 is $\hat{\Sigma}_T(2) = i \eta_{21} \hat{g}(1)$ where $\eta_{21} = \pi \nu_1 \Omega_2^{-1} \langle |T_{\mathbf{p},\mathbf{q}}|^2 \rangle = (4 \nu_2 e^2 \Omega_2 R_{12})^{-1}$. If island 1 has tunnel contacts to two other islands 2 and 3, the self-energy is a sum

$$\begin{split} \bar{\Sigma}_{T}(1) &= i \, \eta_{12} \hat{g}(2) + i \, \eta_{13} \hat{g}(3) \\ \eta_{12} &= (4 \nu_1 e^2 \Omega_1 R_{12})^{-1} \text{ and } \eta_{13} &= (4 \nu_1 e^2 \Omega_1 R_{13})^{-1}. \end{split}$$

where

We define

$$\hat{I}_{\epsilon_1,\epsilon_2}^{K} \equiv \begin{pmatrix} I_1 & I_2 \\ -I_2^{\dagger} & \overline{I}_1 \end{pmatrix} = \int \frac{d\xi_p}{\pi i} \hat{\mathcal{I}}_{\epsilon_1,\epsilon_2}^{K}.$$

The tunnel collision integral for island 1 in contact with island 2 takes the form

$$\begin{split} \tilde{I}_{T}^{k}(1) &= i \,\eta_{12} [\hat{g}^{R}(2) \circ \hat{g}^{K}(1) - \hat{g}^{R}(1) \circ \hat{g}^{K}(2) + \hat{g}^{K}(2) \circ \hat{g}^{A}(1) \\ &- \hat{g}^{K}(1) \circ \hat{g}^{A}(2)]. \end{split}$$
(5)

If the distribution function is time independent, the Keldysh Green function has the form

$$\hat{g}_{\epsilon_1,\epsilon_2}^K = \hat{g}_{\epsilon_1,\epsilon_2}^R (f_{1,\epsilon_2} + \hat{\tau}_3 f_{2,\epsilon_2}) - (f_{1,\epsilon_1} + \hat{\tau}_3 f_{2,\epsilon_1}) \hat{g}_{\epsilon_1,\epsilon_2}^A \tag{6}$$

where the odd- and even-in- ϵ distribution functions f_1 and f_2 are defined such that

$$f_{1,\epsilon} = -n(\epsilon) + n(-\epsilon), \quad f_{2,\epsilon} = 1 - n(-\epsilon) - n(\epsilon).$$

The actual quasiparticle energy distribution function *n* satisfies $f_{1,\epsilon}+f_{2,\epsilon}=1-2n(\epsilon)$.

After integrating Eq. (2) over $d\xi_p$ we obtain two kinetic equations for diagonal in ϵ components

$$\operatorname{Tr}(\hat{M}_{\epsilon,\epsilon}) = \operatorname{Tr}(\hat{I}^{K}_{\epsilon,\epsilon} - (\hat{I}^{R}_{\epsilon,\epsilon} - \hat{I}^{A}_{\epsilon,\epsilon})f_{1,\epsilon}),$$
(7)

$$\operatorname{Tr}(\hat{\tau}_{3}\hat{M}_{\epsilon,\epsilon}) = \operatorname{Tr}(\hat{\tau}_{3}[\hat{I}_{\epsilon,\epsilon}^{K} - (\hat{I}_{\epsilon,\epsilon}^{R} - \hat{I}_{\epsilon,\epsilon}^{A})f_{1,\epsilon}]).$$
(8)

The collision integrals here account for tunnel and inelastic processes $\hat{I} = \hat{I}_T + \hat{I}_{inel}$, where inelastic relaxation is due to electron-phonon and electron-electron interactions, $\hat{I}_{inel} = \hat{I}_{e-ph} + \hat{I}_{e-e}$. The matrix \hat{M} is

$$\begin{split} \hat{M} &= (\hat{H} \circ \hat{g}^K - \hat{g}^K \circ \hat{H}) - (\hat{H} \circ \hat{g}^R - \hat{g}^R \circ \hat{H}) f_1 \\ &+ f_1 (\hat{H} \circ \hat{g}^A - \hat{g}^A \circ \hat{H}). \end{split}$$

B. Charge and energy tunnel currents

After taking the trace and integrating over energy, Eq. (2) in island 1 yields

$$i\omega\left(\nu_{1}\int\frac{d\epsilon}{2}\int\frac{d\xi_{1}}{\pi i}\mathrm{Tr}\hat{G}_{\epsilon_{+},\epsilon_{-}}^{K}\right) = -i\nu_{1}\int\frac{d\epsilon}{2}\mathrm{Tr}(\hat{\tau}_{3}\hat{I}_{\epsilon_{+},\epsilon_{-}}^{K})$$

where $\epsilon_{\pm} = \epsilon \pm \omega/2$. This is the conservation of charge $Q(1) = eN(1)\Omega_1$ in the volume Ω_1 of island 1:

$$\frac{\partial Q(1)}{\partial t} = I(1). \tag{9}$$

Here N(1) is the particle density in island 1. The current that flows into island 1 is

$$I(1) = -ie\nu_1\Omega_1 \int \frac{d\epsilon}{4} \operatorname{Tr}(\hat{\tau}_3 \hat{I}_T^K(1)).$$
(10)

The inelastic collision integral drops out because it conserves the particle number. Multiplying Eq. (2) by ϵ and taking the trace of it one can obtain the balance of the energy $\mathcal{E}=\Omega E$ in the form

$$\frac{\partial \mathcal{E}(1)}{\partial t} = I_{\mathcal{E}}(1).$$

Here the energy density is

$$E = -\int \epsilon \operatorname{Tr}(\hat{\tau}_3 \hat{G}^K_{\epsilon_+, \epsilon_-}) \frac{d\epsilon}{4\pi i} \frac{d^3 p}{(2\pi)^3} + \frac{|\Delta|^2}{|g|} - Ne\varphi$$

where φ is the electric potential of the island. The energy current into island 1 is

$$I_{\mathcal{E}}(1) = -i\nu_1\Omega_1 \int \frac{d\epsilon}{4} \operatorname{Tr}((\epsilon - e\varphi_1\hat{\tau}_3)\hat{I}^K(1)).$$
(11)

The collision integral in Eq. (11) contains only tunnel and electron-phonon contributions, $I^{K}(1) = I_{e-ph}^{K}(1) + I_{T}^{K}(1)$. The electron-electron interactions drop out because of the energy conservation. The energy flow into island 1 can thus be separated into two parts. One part containing I_{e-ph}^{K} is the energy exchange with the heat bath (phonons). The other part contains the tunnel contribution $I_{T}^{K}(1)$ and is the energy current into island 1 through the tunnel contact.

IV. KINETIC EQUATIONS IN HYBRID STRUCTURES

A. Nonzero superconducting chemical potential

If a hybrid structure containing more than one superconducting island is voltage biased, at least one of the superconductors will have a nonzero chemical potential μ_s with a time-dependent phase of the order parameter. If the external voltage is constant and the resulting state is stationary, the phase is linear in time, $\chi = 2\mu_s t$, while the magnitude of the order parameter is time independent. Consider two sets of regular Green functions. One set of functions $\hat{g}^{R(A)}$ is taken in the basis where the order parameter phase varies in time. The other set belongs to the same order parameter magnitude but has a phase constant in time (zero for simplicity). However, the energies of the Green functions $g^{R(A)}$, $\overline{g}^{R(A)}$, $f^{R(A)}$, and $f^{\dagger R(A)}$ are shifted by different amounts proportional to the chemical potential in the previous representation. The relations between these two sets, the Green functions $g_{\epsilon,\epsilon'}^{R(A)}$ and $f_{\epsilon,\epsilon'}^{R(A)}$ and the steady-state functions with shifted energies, can be established from the spatially homogeneous Usadel equations, i.e., Eq. (3) integrated over $d\xi_p$. Since $\Delta = |\Delta| e^{+i2\mu_S t}$, i.e.,

$$\Delta_{\omega} = |\Delta| 2\pi \delta(\omega + 2\mu_{S}), \quad \Delta_{\omega}^{*} = |\Delta| 2\pi \delta(\omega - 2\mu_{S}),$$

one observes that the Usadel equations are satisfied by

$$g_{\epsilon,\epsilon'}^{R(A)} = g_{\epsilon+\mu_S}^{R(A)} 2\pi \delta(\epsilon - \epsilon'), \quad \overline{g}_{\epsilon,\epsilon'}^{R(A)} = \overline{g}_{\epsilon-\mu_S}^{R(A)} 2\pi \delta(\epsilon - \epsilon')$$
(12)

and

$$f_{\epsilon,\epsilon'}^{\dagger R(A)} = f_{\epsilon-\mu_S}^{\dagger R(A)} 2\pi \delta(\epsilon - \epsilon' - 2\mu_S)$$

$$f_{\epsilon,\epsilon'}^{R(A)} = f_{\epsilon+\mu_S}^{R(A)} 2\pi \delta(\epsilon - \epsilon' + 2\mu_S).$$
(13)

The functions $g_{\epsilon}^{R(A)} = -\overline{g}_{\epsilon}^{R(A)}$ and $f_{\epsilon}^{R(A)} = f_{\epsilon}^{\dagger R(A)}$ satisfy the steady-state Usadel equation

$$-2\epsilon f_{\epsilon}^{R(A)} + 2|\Delta|g_{\epsilon}^{R(A)} = I_{2,\epsilon}^{R(A)}$$
(14)

supplemented with the normalization

$$g_{\epsilon}^{R(A)})^2 - (f_{\epsilon}^{R(A)})^2 = 1.$$
 (15)

Therefore, we find from Eq. (6) for a nonzero chemical potential

$$g_{\epsilon,\epsilon'}^{K} = (g_{\epsilon+\mu_s}^{R} - g_{\epsilon+\mu_s}^{A})(f_{1,\epsilon} + f_{2,\epsilon})2\pi\delta(\epsilon - \epsilon'), \quad (16)$$

$$-\overline{g}_{\epsilon,\epsilon'}^{K} = (g_{\epsilon-\mu_{s}}^{R} - g_{\epsilon-\mu_{s}}^{A})(f_{1,\epsilon} - f_{2,\epsilon})2\pi\delta(\epsilon - \epsilon').$$
(17)

The off-diagonal Keldysh Green functions in Eq. (6) become

$$f_{\epsilon,\epsilon-\omega}^{K} = \left[f_{\epsilon+\mu_{S}}^{R}(f_{1,\epsilon+2\mu_{S}} - f_{2,\epsilon+2\mu_{S}}) - (f_{1,\epsilon} + f_{2,\epsilon})f_{\epsilon+\mu_{S}}^{A}\right]$$
$$\times 2\pi\delta(\omega + 2\mu_{S}). \tag{18}$$

$$f_{\epsilon,\epsilon-\omega}^{\dagger K} = \left[f_{\epsilon-\mu_{S}}^{\dagger R} (f_{1,\epsilon-2\mu_{S}} + f_{2,\epsilon-2\mu_{S}}) - (f_{1,\epsilon} - f_{2,\epsilon}) f_{\epsilon-\mu_{S}}^{\dagger A} \right] \\ \times 2\pi \delta(\omega - 2\mu_{s}).$$
(19)

B. Tunnel collision integrals

Assume that island 1 is a superconductor and island 2 is a normal metal. With Eqs. (12), (16), and (17), the matrix elements of the tunnel collision integral Eq. (5) in the superconducting island S give

$$Tr(\hat{I}_{T}^{K}(S)) = 4i \eta_{SN} \{ [g_{\epsilon+\mu_{S}}(S) + g_{\epsilon-\mu_{S}}(S)] [f_{1,\epsilon}(S) - f_{1,\epsilon}(N)] \\ + [g_{\epsilon+\mu_{S}}(S) - g_{\epsilon-\mu_{S}}(S)] [f_{2,\epsilon}(S) - f_{2,\epsilon}(N)] \}, \quad (20)$$
$$Tr(\hat{\tau}_{3}\hat{I}_{T}^{K}(S)) = 4i \eta_{SN} \{ [g_{\epsilon+\mu_{S}}(S) - g_{\epsilon-\mu_{S}}(S)] [f_{1,\epsilon}(S) - f_{1,\epsilon}(N)] \\ + [g_{\epsilon+\mu_{S}}(S) + g_{\epsilon-\mu_{S}}(S)] [f_{2,\epsilon}(S) - f_{2,\epsilon}(N)] \}. \quad (21)$$

Components of the retarded and advanced tunnel collision integrals $\hat{I}^{R(A)}$ are

$$\begin{split} I_2^{R(A)}(S) &= \pm i \, \eta_{SN} f^{R(A)}(S) \,, \\ &- I_2^{\dagger R(A)}(S) = \pm i \, \eta_{SN} f^{\dagger R(A)}(S) \,, \\ &I_1^{R(A)}(S) = \overline{I}_1^{R(A)}(S) = 0 \,. \end{split}$$

Both Keldysh and R(A) tunnel collision integrals in the normal island N are coupled to those in the island S by $\eta_{SN}I(N) = -\eta_{NS}I(S)$.

Using Eqs. (13) and (18), we find

$$\operatorname{Tr}(\hat{M}_{\epsilon,\epsilon}) = 2|\Delta|F_{\epsilon+\mu_{S}}(f_{1,\epsilon} - f_{1,\epsilon+2\mu_{S}} + f_{2,\epsilon} + f_{2,\epsilon+2\mu_{S}}) - 2|\Delta|F_{\epsilon-\mu_{S}}(f_{1,\epsilon-2\mu_{S}} - f_{1,\epsilon} + f_{2,\epsilon} + f_{2,\epsilon-2\mu_{S}}),$$

$$(22)$$

$$\begin{aligned} \operatorname{Tr}(\hat{\tau}_{3}M_{\epsilon,\epsilon}) &= 2|\Delta|F_{\epsilon+\mu_{S}}(f_{1,\epsilon} - f_{1,\epsilon+2\mu_{S}} + f_{2,\epsilon} + f_{2,\epsilon+2\mu_{S}}) \\ &+ 2|\Delta|F_{\epsilon-\mu_{S}}(f_{1,\epsilon-2\mu_{S}} - f_{1,\epsilon} + f_{2,\epsilon} + f_{2,\epsilon-2\mu_{S}}). \end{aligned}$$

$$\end{aligned} \tag{23}$$

After inserting Eqs. (20)–(23) into Eqs. (7) and (8) they yield the final kinetic equations to be solved for the distribution functions in the absence of inelastic scattering.

In the equations above, we introduce $g_{\epsilon} = (g_{\epsilon}^{R} - g_{\epsilon}^{A})/2$ $\equiv \operatorname{Re} g_{\epsilon}^{R}$ and also $F_{\epsilon} = (f_{\epsilon}^{R} + f_{\epsilon}^{A})/2 \equiv i \operatorname{Im} f_{\epsilon}^{R}$. The functions g_{ϵ} and F_{ϵ} are even in ϵ . With the account of tunnel and electron-phonon interactions, they can be found from the steady-state Usadel equations (14) and (15). These equations can be more easily solved when the inelastic interaction is small. If island 2 is normal, one finds

$$g_{\epsilon}^{R(A)} = \pm \frac{\epsilon \pm i\gamma}{\sqrt{(\epsilon \pm i\gamma)^2 - |\Delta|^2}}, \quad f_{\epsilon}^{R(A)} = \frac{|\Delta|}{\epsilon \pm i\gamma} g_{\epsilon}^{R(A)}, \quad (24)$$

where $\gamma = \eta_{SN}$.

For a N₁IS₁INIS₂IN₂ structure, each conductor S₁, N, or S₂ has tunnel contacts with two other conductors, thus the tunnel integrals are sums of the contributions from two islands. For example, if the inelastic interaction is small, the depairing rate in Eq. (24) is $\gamma(S_i) = \eta_{S_iN} + \eta_{S_iN_i}$.

C. Current conservation

For a NS junction, the tunnel current Eq. (10) into the superconductor S from a normal island N or $N_{1,2}$ is

$$I(N \to S) = \frac{1}{4eR_{SN}} \int_{-\infty}^{\infty} d\epsilon g_{\epsilon}(S) [f_{1,\epsilon-\mu_S}(S) - f_{1,\epsilon+\mu_S}(S) - f_{1,\epsilon-\mu_S}(S) + f_{1,\epsilon-\mu_S}(N) + f_{1,\epsilon+\mu_S}(N) + f_{2,\epsilon-\mu_S}(S) + f_{2,\epsilon+\mu_S}(S) - f_{2,\epsilon-\mu_S}(N) - f_{2,\epsilon+\mu_S}(N)].$$
(25)

Electric currents through both junctions are balanced when $I(N \rightarrow S_i) + I(N_i \rightarrow S_i) = 0$.

The energy current flowing into the superconducting island through each junction has the form

$$I_{\mathcal{E}}(N \to S) = \frac{1}{4e^2 R_{NS}} \int_{-\infty}^{\infty} d\epsilon \epsilon g_{\epsilon}(S) ([f_{1,\epsilon-\mu_S}(S) + f_{1,\epsilon+\mu_S}(S) - f_{1,\epsilon-\mu_S}(N) - f_{1,\epsilon+\mu_S}(N)] + [f_{2,\epsilon-\mu_S}(S) - f_{2,\epsilon+\mu_S}(S) - f_{2,\epsilon-\mu_S}(N) + f_{2,\epsilon+\mu_S}(N)]) - I(N \to S) [\mu_S/e + \varphi_S].$$
(26)

The energy conservation follows from the kinetic equations and is therefore an abundant condition. However, in the quasi-equilibrium limit, when the electron-electron interaction in each island dominates over the tunnel injection, the distribution functions are nearly thermal with certain temperatures specific for each island. In this case, the kinetic equations do not need to be solved explicitly, but the temperatures are found by requiring the conservation of the energy current. The energy current $I_{\mathcal{E}}(S \to N)$ flowing into the central normal island through each junction is obtained from Eq. (26) by changing the sign and replacing φ_S with φ_N so that the difference between the energy currents from N to S and from S to N is $I_{\mathcal{E}}(N \to S) + I_{\mathcal{E}}(S \to N) = I(N \to S)V_{NS}$, where V_{NS} $= \varphi_N - \varphi_S$ is the voltage across the junction. This difference of energy currents is compensated by the work produced by the voltage source and does not lead to the change in the energy of the superconductor.

D. Self-consistency equation and charge neutrality

The magnitude of the order parameter is found from Eq. (18). It is a real quantity, therefore,

$$\frac{|\Delta|}{\lambda} = \int_{-E_c}^{E_c} (\operatorname{Re} f_{\epsilon}^{R}) (f_{1,\epsilon+\mu_S} + f_{1,\epsilon-\mu_S} - f_{2,\epsilon+\mu_S} + f_{2,\epsilon-\mu_S}) \frac{d\epsilon}{4},$$
(27)

where E_c is the BCS cutoff energy. The self-consistency of the order parameter requires vanishing of its imaginary part:

$$\int_{-\infty}^{\infty} F_{\epsilon}(f_{1,\epsilon+\mu_{S}} - f_{1,\epsilon-\mu_{S}} - f_{2,\epsilon+\mu_{S}} - f_{2,\epsilon-\mu_{S}})d\epsilon = 0, \quad (28)$$

which results in

$$\int \mathrm{Tr}\hat{M}d\boldsymbol{\epsilon} = \int \mathrm{Tr}(\hat{\tau}_{3}\hat{M})d\boldsymbol{\epsilon} = 0.$$
 (29)

Equation (28) or the second condition in Eq. (29) together with the kinetic equation Eq. (8) is equivalent to $\int \text{Tr}(\hat{\tau}_3 \hat{I}^K) d\epsilon = 0$. Since the integrals over the inelastic electron-phonon and electron-electron collision integrals vanish, this is the condition of current conservation:

$$\int \mathrm{Tr}(\hat{\tau}_3 \hat{t}_T^K) d\epsilon = 0, \qquad (30)$$

which implies through Eq. (10) that the sum of currents flowing into an island is zero.

Charge density can be obtained by calculating Tr \hat{g}^{K} from Eqs. (16) and (17). Making shift of ϵ in the integral we obtain

$$\begin{split} N &= N_0 - 2\nu(e\varphi + \mu_S) \\ &+ \nu \int_{-\infty}^{\infty} g_{\epsilon}(f_{1,\epsilon+\mu_S} - f_{1,\epsilon-\mu_S} - f_{2,\epsilon+\mu_S} - f_{2,\epsilon-\mu_S}) \frac{d\epsilon}{2}. \end{split}$$

Here we take into account the divergence of the integral for large ϵ . Charge neutrality requires that $N=N_0$, i.e., the charge density in the superconducting state equals that in the normal state. The electric potential φ_S of a superconductor is thus coupled to the chemical potential through

$$e\Phi = \int_{-\infty}^{\infty} g_{\epsilon} (f_{1,\epsilon+\mu_S} - f_{1,\epsilon-\mu_S} - f_{2,\epsilon+\mu_S} - f_{2,\epsilon-\mu_S}) \frac{d\epsilon}{4}, \quad (31)$$

where $e\Phi = \mu_S + e\varphi_S$.

To summarize, for a system of *n* superconducting and m = n+1 normal pieces we have n+m pairs of the distribution



FIG. 2. (a) Chemical and electrostatic potentials of the superconducting island as a function of bias voltage. (b) *I-V* curves for different ρ . For clarity, in the case $\rho = 1000$ the potentials have been multiplied by the factor 100 and the currents by the factor 1000.

functions f_1 and f_2 and n parameters μ_S . For these unknowns, we have 2(n+m) functional kinetic equations [Eqs. (7) and (8)] and n conditions [Eq. (30)] of current conservation for each superconducting island. Note that the current balance of the type of Eq. (30) in each normal island is satisfied automatically due to the kinetic equations that have $\hat{M} \equiv 0$ in any normal conductor.

V. RESULTS

To further simplify our model, we limit the discussion to a case symmetric with respect to the central normal island, i.e., $R_{N_1S_1} = R_{N_2S_2} \equiv R_{out}$, $R_{NS_1} = R_{NS_2} \equiv R_{in}$, $|\Delta_1| = |\Delta_2| \equiv \Delta$, and $\varphi_{N_1} = -\varphi_{N_2}$, $\mu_{S_1} = -\mu_{S_2} \equiv \mu_S$. Therefore $f_1(N_1) = f_1(N_2)$ and $f_2(N_1) = -f_2(N_2)$. Moreover, when applied to kinetic equations in the normal island N, the symmetry yields $f_2(S_1) = -f_2(S_2) \equiv f_2(S), f_1(S_1) = f_1(S_2) \equiv f_1(S)$, and $f_2(N) = 0$. The system is biased by voltage V applied between the outermost normal electrodes N₁ and N₂.

A. Nonequilibrium state

1. Distribution functions

Under the conditions formulated in Sec. II when the inelastic relaxation can be neglected, $\eta \ge \tau_{e-e}^{-1}$, the kinetic equations (7) and (8) were solved numerically for nonequilibrium distributions in both normal and superconducting islands. Throughout numerical computations we use the value of the pair-breaking parameter $\Delta/\gamma = 1000$. For nonequilibrium states, the bath temperature was fixed at a value much lower than the gap, $T_{\text{bath}} = (0.1/1.764)\Delta$. This corresponds to *T*



FIG. 3. Nonequilibrium normal-island distributions for $\rho = 1000$. We use $v \equiv eV/\Delta$ here and in Figs. 4–7 below.



FIG. 4. Nonequilibrium normal metal distributions for $\rho = 1$.

=0.1 T_c if the gap coincides with the zero-temperature BCS value $\Delta_0 = 1.764T_c$.

The distribution functions in both N and S islands are in turn determined by the chemical potential of superconducting islands μ_S which was calculated self-consistently using the current conservation as discussed in the previous section. Chemical potential and the current-voltage curves are shown in Fig. 2.

We find three qualitatively distinct cases characterized by the ratio $\rho \equiv R_{out}/R_{in}$. For the ratio as large as $\rho = 1000$, the distribution in the normal island, n_N , is shown in Fig. 3 for several values of V. Under biasing V, the distribution is driven into nonequilibrium. At first, this is seen only as a slight deviation from the Fermi function n_F but for voltages $eV/\Delta \gtrsim 1$, the distribution n_N assumes the characteristic steplike profile. This behavior of n_N can be explained as follows. For very high ratios ρ , the superconducting chemical potential is small: $\mu_s/eV \ll 1$ [compare with Fig. 2(a)]. The kinetic equations (7) and (8) yield then $f_1(N) \approx f_1(S) \approx f_1(N_1)$. Since $f_2(N) = 0$, one has $2n_N = 1 - f_1(N) = 1 - f_1(N_1)$. The external leads N_1 and N_2 are in equilibrium such that $f_1(N_1) = -n_F(\epsilon)$ -eV/2)+ $n_F(-\epsilon - eV/2)$. Thus, $n_N = [n_F(\epsilon - eV/2) + n_F(\epsilon)]$ +eV/2]/2. Since $f_2(S)$ is small due to the high resistivity ratio, one has also $n_S \approx n_N$ for the superconducting island.

The distributions for the ratios $\rho = 1$ and $\rho = 1/1000$ are shown in Figs. 4 and 5. They also change their shape strongly above a certain voltage. The changes correspond to the sharp rise in the electric current through the junction [Fig. 2(b)]. The origin of this behavior is discussed in connection with the charge imbalance (see next subsection). The distribution functions of the N island show a cooling behavior: The distribution becomes very steep, thus corresponding to a low effective electron temperature at such bias voltages



FIG. 6. Effective temperature T_{eff} of Eq. (32).

when the chemical potential difference across the superconductor/normal-island junction is $|\mu_S| \approx \Delta$. One can define an effective temperature through⁴

$$k_B T_{\text{eff}} = \frac{\sqrt{6}}{\pi} \sqrt{\int_0^\infty [1 - f_{1,\epsilon}(N)] \epsilon d\epsilon}, \qquad (32)$$

keeping in mind that $f_2(N)=0$ for the center N island. This definition gives the actual electron temperature in (quasi)-equilibrium. The effective temperatures have minima for $\rho \leq 1$ as seen in Fig. 6.

For a good contact between the outer normal electrodes and the superconductors S, i.e., when $\rho = 1/1000$, there is essentially no difference between the superconductor distribution function n_S and that in the normal reservoir, a Fermi function with $\varphi_{N_1} = V/2$. This suggests that, for small ρ , the considered structure is similar to a SINIS system with superconducting reservoirs. The distribution in the central N island for low ρ is shown in Fig. 5. It resembles the distribution found for a SINIS structure.⁹

When the ratio ρ increases, the distribution in the superconductors, n_S , deviates from the Fermi function as shown in Fig. 7. Nonequilibrium distribution in the superconducting islands on both sides of the central normal island drives the state in the N island yet further from equilibrium (see Fig. 4). For larger ρ the cooling behavior becomes less pronounced and finally disappears (see Fig. 6).

For low and intermediate values of ρ , we observe novel features of highly nonequilibrium distributions both in the central normal and in the side superconducting islands. For small $\rho = 1/1000$, peaks in the energy distribution appear at



FIG. 5. Nonequilibrium normal metal distributions for $\rho = 1/1000$.



FIG. 7. Nonequilibrium superconductor distributions for various bias voltages when $\rho = 1$.

energies $\pm \epsilon = \Delta + \mu_S$ (see Fig. 5). In addition to these, new peaks appear in Fig. 4 for larger $\rho = 1$ at energies $\pm \epsilon = -\Delta + \mu_S$, for voltages considerably exceeding Δ/e . Both sets of peaks come as a result of recursion from singularities in g_{ϵ} and F_{ϵ} at $\epsilon = \pm \Delta$ in the kinetic equations. The new peaks at $\pm \epsilon = -\Delta + \mu_S$ are present as long as the distribution functions of the two reservoirs differ at the corresponding energies and appear as a result of suppression of the distribution function due to a large factor $(\Delta/\gamma)F_{\epsilon\pm\mu_S}$ in the subgap region in Eqs. (22) and (23). Thus, the requirement of the new peaks is roughly $\Delta - \mu_S < eV/2$ which can be fulfilled if $\mu_S \neq -eV/2$ as is the case for intermediate values of ρ .

2. Charge imbalance

As mentioned above, the distribution function of the center N island suffers a drastic change above a certain voltage. This change coincides with the upturn of the current as a function of the applied voltage in Fig. 2(b) and is accompanied by a deviation of the chemical potential μ_S from the electric potential $-e\varphi_S$ in the adjacent superconductor as determined by Eq. (31). In equilibrium, their difference $\Phi=0$. In nonequilibrium, a difference between μ_S and $-e\varphi_S$ appears according to Fig. 2(a). The singularities appear when the chemical potential difference between the superconductor and one of the contacting normal conductors approaches Δ . This corresponds to $eV \sim 2\Delta$ for a large mismatch between $R_{\rm in}$ and $R_{\rm out}$ or to $eV/2 \sim 2\Delta$ for $R_{\rm in} = R_{\rm out}$. To measure potentials φ_S , a capacitive connection would be required, in addition to the usual resistive connection only capable of detecting μ_{s} .

For a good contact between the outer normal electrodes and the superconductors S, i.e., when $\rho = 1/1000$, there is no deviation between $-e\varphi_S$ and μ_S . This can be understood by considering the function n_S , which coincides essentially with a Fermi function shifted by $\mu_S \approx -e\varphi_{N_1} = -eV/2$. For a Fermi function, $n_{\epsilon} = 1 - n_{-\epsilon}$, and the term in the brackets in Eq. (31) vanishes.

3. Gap instability

In a nonequilibrium state, the gap function Δ has to be calculated self-consistently using Eq. (27). Employing the equation for the critical temperature T_c ,

$$\frac{1}{\lambda} = \int_{0}^{E_{c}} \tanh \frac{\epsilon}{2T_{c}} \frac{d\epsilon}{\epsilon}, \qquad (33)$$

one can exclude the interaction constant in favor of T_c . For a low resistance ratio $\rho = 1/1000$, the gap does not change considerably, $\Delta \approx \Delta_0$ for all V, $\Delta_0 = 1.764T_c$ being the zerotemperature BCS gap. However, for a higher resistance ratio, the nonequilibrium energy gap is modified dramatically as shown in Fig. 8(a). One observes a drastic change in the gap for voltages coinciding with those where the change in the distribution is seen. The energy gap becomes a multi-valued function which implies hysteretic behavior accompanied by jumps of Δ at the corresponding voltages. For a very poor contact between the superconducting islands and the outer normal reservoirs, i.e., for high ρ when deviation from equi-



FIG. 8. (a) Gap Δ as a function of bias voltage for two resistance ratios $\rho = 1000$ (dashed line) and $\rho = 1$ (solid line). Both quantities are normalized to zero-temperature BCS gap Δ_0 . (b) Conversion between eV/Δ and eV/Δ_0 for $\rho = 1000$ (dashed line) and $\rho = 1$ (solid line).

librium is the largest, the gap function jumps down to very small values and superconductivity is nearly destroyed. For a lower tunnel resistance ratio ρ , the gap decrease is not so huge and superconductivity is less suppressed. Note that with respect to the order parameter magnitude, the bath temperature chosen for our calculations can be considered as zero. Indeed, the temperature was set much lower than Δ while the relevant energy scales for the distribution function are determined by the applied voltage and by Δ itself. Thus the thermal effects on the gap are negligible.

The predicted suppression of superconductivity in a nonequilibrium superconductor placed into a tunnel contact with a *nonequilibrium normal-metal electrode* contrasts to the superconductivity enhancement observed in tunnel SISIS structures^{5–7} where the nonequilibrium superconductor is in contact with *equilibrium superconducting* electrodes.

Note that since in our calculations we normalize all the values with the dimensions of energy (like voltage, excitation energy, and temperature) to the real gap magnitude Δ , we need to rescale the real voltage to its relative magnitude. Conversion between the relative, eV/Δ , and the real voltage normalized to the BCS gap, eV/Δ_0 , is provided by Fig. 8(b) where the graphs are given for ρ =1000 and ρ =1. As seen from Fig. 8(b), for ρ =1 high relative voltages eV/Δ can be achieved for comparatively low absolute voltage values eV/Δ_0 .

4. Visualization

The peaks in the distribution of N and S islands can be monitored by measuring the differential conductance of a



FIG. 9. Differential conductance for (a) the probe junction and (b) the corresponding quasiparticle energy distribution in the normal-metal island when $\rho = 1$, $eV/\Delta = 10$.



FIG. 10. Temperature in the normal-metal island as a function of bias voltage for bath temperatures (a) 0.05Δ and (b) 0.2Δ when the system is in quasiequilibrium.

probe tunnel SINIS or SIS'IS junction attached to the island in question. Let us consider a SINIS probe junction attached to the central N island as in Ref. 4. The distribution function in the N island is not modified by the measuring current if the tunnel resistance of the probe junction satisfies $R_{S_{PN}} \gg R_{in} + R_{out}$. The current through the probe junction is

$$I(N \to S_P) = -\frac{1}{eR_{S_PN}} \int_{-\infty}^{\infty} g_{\epsilon}(S_P) \\ \times [n_F(\epsilon) - n_N(\epsilon + eV_P/2)] d\epsilon,$$

where the bulk superconducting probe electrode S_P is assumed to be in equilibrium with a potential φ_P so that $\mu_{S_P} = -e\varphi_P = -eV_P/2$ where V_P is the voltage between the two probe electrodes. The energy gap Δ_P in the probe electrodes is assumed to have the magnitude corresponding to the BCS value for $T=T_{\text{bath}}$. As $T_{\text{bath}} \approx 0.1T_c$, the gap Δ_P is very close to Δ_0 . In addition, we set $\gamma_P = \gamma$ for the depairing rate in the probe electrodes. The differential conductance becomes

$$\frac{dI}{dV_P} = \frac{1}{2R_{S_PN}} \int_{-\infty}^{\infty} g_{\epsilon}(S_P) \frac{dn_N(\epsilon + eV_P/2)}{d\epsilon} d\epsilon.$$
(34)

Due to the peaks in $g_{\epsilon}(S_P)$ at $\epsilon = \pm \Delta_P$, the differential conductance should reproduce the peaks in the distribution function n_N at the probe voltages satisfying $\epsilon \pm \Delta_P = eV_P/2$.

The differential conductance $dI(N \rightarrow S_P)/dV_P$ together with the corresponding distributions in the central normal island for $eV/\Delta=10$ are shown in Fig. 9 for the ratio $\rho=1$. The peaks in the distribution in Fig. 9(b) are located at ϵ $=\pm\Delta-\mu_S$, i.e., $\epsilon=2.89\Delta$ and $\epsilon=4.89\Delta$. Two more are located at $\epsilon=\pm\Delta-3\mu_S$ or $\epsilon=10.7\Delta$ and $\epsilon=12.7\Delta$ [out of scale in Fig 9(b)]. Comparing these to the locations of the larger peaks in



FIG. 11. Temperature in the superconductor as a function of bias voltage for bath temperatures (a) 0.05Δ and (b) 0.2Δ when the system is in quasiequilibrium.



FIG. 12. Chemical potential of the superconductor as a function of bias voltage for bath temperatures (a) 0.05Δ and (b) 0.2Δ when the system is in quasiequilibrium.

Fig. 9(a) and using the Δ -vs- Δ_0 conversion curve of Fig. 8(a) we see that the peaks in the distribution are indeed reproduced at the probe voltage $eV_P/\Delta = 2\epsilon/\Delta \pm 2\Delta_P/\Delta$. The smaller peaks in Fig. 9(a) refer to the less pronounced structure in the distribution not well resolved in Fig. 9(b).

B. Quasi-equilibrium

As discussed in Sec. II, the condition for full nonequilibrium is $\eta \ge \tau_{\text{inel}}^{-1}$, where η is defined according to Eq. (4). The inelastic relaxation may be further separated to relaxation caused by electron-electron and electron-phonon interactions with collision rates τ_{e-e}^{-1} and τ_{e-ph}^{-1} , respectively. The experimental situation in nanoscale heterostructures⁴ corresponds often to the case $\tau_{e-e}^{-1} > \tau_{e-ph}^{-1}$. In particular, the limit of low coupling to the heat bath in S and/or N islands is frequently realized when the tunnel injection rate is intermediate between the electron-phonon and electron-electron relaxation rates, $\tau_{e-ph}^{-1} \ll \eta \ll \tau_{e-e}^{-1}$. We refer to this case as quasi-equilibrium. While the near absence of electron-phonon interactions prohibits the quasiparticles from coupling to the lattice, the rate of electron-electron scattering is high enough for the quasiparticles to assume a Fermi distribution with certain electron temperature. We have studied the cooling performance of our NISINISIN heterostructure in the quasiequilibrium limit looking at the electron temperature of the central N island.

We note that the simple expressions for the regular Green functions of the form of Eq. (24) are not applicable in the strict sense when the inelastic relaxation dominates. To find the exact expressions for the regular Green functions one has to solve Eqs. (14) and (15) with the proper inelastic collision integrals. However, to simplify our problem, we model the pair-breaking effects of inelastic relaxation by an effective pair-breaking rate γ in Eq. (24) in the same way as for the tunnel limit described in the previous sections. This approximation is frequently used in practical calculations. Here we put $\Delta/\gamma = 1000$ as above.

Applying the tunneling model shows that depending on the configuration of the quasiparticle traps, i.e., the ratio of outer and inner junction resistances ρ , effective cooling of the normal-metal island can be achieved as demonstrated also in Ref. 4. The temperatures of the central N island and of the contacting S island are shown in Figs. 10 and 11, respectively. The temperature of the N island indeed displays a minimum below bath temperature when the ratio ρ is small.



FIG. 13. Electric current between S and N islands as a function of bias voltage for bath temperatures (a) 0.05Δ and (b) 0.2Δ when the system is in quasiequilibrium.

However, for large ρ , the temperature monotonously rises above the bath temperature with an increasing voltage. As can be deduced from Figs. 10 and 11, the cooling effect is not attributed to the presence of two additional normal metal reservoirs. On the contrary, smallest ratio ρ , which corresponds to the strongest cooling, is seen to lead to almost constant $T_S = T_{\text{bath}}$ as it would be the case for pure superconducting reservoirs. Another remark concerns the cooling efficiency of a NISINISIN configuration as T_{bath} is lowered. Indeed, for $T_{\text{hath}}=0.2\Delta$ the temperature minimum for the depairing parameter $\gamma = \Delta/1000$ used for our calculations is roughly 0.063 Δ while the minimum is $T_{\min}=0.024\Delta$ for $T_{\text{bath}}=0.15\Delta$, and it is $T_{\min}\approx 0.013\Delta$ for $T_{\text{bath}} \leq 0.1\Delta$. From the numerical results in the case $1 \ge \rho \ge \gamma/\Delta$, we find that the minimum achieved temperature follows $T_{\rm min}/T_c \approx 0.24 \rho^{\zeta}$ where $\zeta \approx 0.5$. For smaller R_{out} , the minimum temperature T_{\min} is determined by the inverse proximity effect described by the depairing parameter γ (see Ref. 4). Combining both the superconductor heating due to a finite R_{out} and the effects of depairing, we can write an approximate formula for the minimum temperature for relatively low bath temperatures,

$$T_{\min}/T_c = 2.5(\gamma/\Delta)^{2/3} + 0.24\rho^{1/2}.$$
 (35)

For $\rho \leq 1$, the depairing rate γ is limited by R_{out} , i.e., $\gamma = 1/(4\nu_S e^2 \Omega_S R_{out}) = (r_S/2R_{out})E_T$, where $E_T = D/L^2$ is the Thouless energy in the superconducting island with length *L* and r_S is its resistance. Substituting this into Eq. (35), we find that the minimum temperature is optimized with

$$\rho \approx 6.4 [(E_T/\Delta)(r_S/R_{\rm in})]^{4/7}.$$
 (36)

Note, however, that Eq. (36) is valid provided $r_S/R_{in} \ll \rho \ll 1$.

The sharp rise in T_N and T_S occurs generally around $eV = 2\Delta$ but for larger values of ρ , when also $T_{\text{bath}} \rightarrow T_{\text{min}}$, the upturn shifts towards higher voltages V. This is because the upturn is determined by the condition $-\mu_S \approx \Delta$ rather than $eV/2 = \Delta$ as can be seen by comparing Figs. 10 and 11 to Fig. 12. For increasing bath temperature, though, this trend is smeared and disappears. The voltages corresponding to the temperature rise are also seen in the *IV* curves in Fig. 13 and in the differential conductance (Fig. 14).



FIG. 14. Differential conductance as a function of bias voltage for bath temperatures (a) 0.05Δ and (b) 0.2Δ when the system is in quasiequilibrium.

VI. DISCUSSION

A. Electron cooling

According to our results, the electron cooling is the most effective when the outer resistance is low, $\rho \ll 1$. In fact, both the effective temperature and the distribution function in the superconducting island almost coincide with those in the bath for such voltages that yield $|\mu_s| < \Delta$ especially for very low ratios of ρ . When the ratio ρ is low, a good contact between the inner superconducting island and the outer electrode makes the distributions in these two islands not so much different from each other, thus decreasing the role of the extra junction. This conclusion is valid only within the tunneling approximation. When the contact between the outer electrodes and superconducting islands are more transparent, cooling properties of the device are affected by the inverse proximity effect from the external normal leads.

For larger ratios ρ , the extra junction prevents the state of the superconducting island from reaching equilibrium, thus reducing the cooling power of the entire structure. Moreover, this limit has another disadvantage as far as the cooling performance is concerned: For larger voltages when eV approaches 2Δ , one expects a suppression of superconductivity in the *S* islands down to lower values of Δ and thus the cooler would become even less effective. This suggests that the cooling performance of an NISINISIN structure cannot be improved essentially by an extra tunnel junction as compared to that of a simple SINIS structure. However, the presence of the quasiparticle traps helps to practically realize the superconducting reservoirs by thermalizing them quickly to an object with a high thermal conductance.

B. Nonequilibrium distribution

Nonequilibrium distribution formed in the superconducting islands for high values of ρ results in yet stronger deviation from equilibrium in the central normal island. As seen from Figs. 4 and 5, the distribution function in the N island is characterized by peaks at energies $\pm \epsilon = \Delta \pm \mu_s$, and also $\pm \epsilon$ $= \Delta \pm 3\mu_s$, etc., for voltages considerably exceeding Δ/e . These peaks are clearly visible in the differential conductance of a probe tunnel SINIS junction made at the central normal island (Fig. 9). Simultaneously, nonequilibrium states in the superconducting island follow the gap which is strongly reduced as compared to its equilibrium BCS value Δ_0 . The transition into a nonequilibrium state is accompanied by a jump in the gap magnitude which leads to the jump in the relative voltage: high values of eV/Δ can be reached already for not very large absolute values of voltage eV/Δ_0 . This makes observation of the nonequilibrium states in N and S islands more easily accessible in experiments.

ACKNOWLEDGMENTS

We are thankful to J. P. Pekola for stimulating discussions. T.T.H. acknowledges funding by the Academy of Finland and the NCCR Nanoscience.

- ¹M. Nahum, T. M. Eiles, and J. M. Martinis, Appl. Phys. Lett. **65**, 3123 (1994).
- ²M. M. Leivo, J. P. Pekola, and D. V. Averin, Appl. Phys. Lett. 68, 1996 (1996).
- ³D. Golubev and A. Vasenko, in *International Workshop on Super-conducting Nano-electronics Devices*, edited by J. Pekola, B. Ruggiero, and P. Silvestrini (Kluwer Academic/Plenum Publishers, New York, 2002), p. 165.
- ⁴J. P. Pekola, T. T. Heikkilä, A. M. Savin, J. T. Flyktman, F. Giazotto, and F. W. J. Hekking, Phys. Rev. Lett. **92**, 056804 (2004).
- ⁵D. R. Heslinga and T. M. Klapwijk, Phys. Rev. B **47**, 5157 (1993).
- ⁶V. M. Dmitriev, V. N. Gubankov, and F. Y. Nad', in *Nonequilibrium Superconductivity*, edited by D. N. Langenberg and A. I. Larkin (North Holland, Amsterdam, 1986), p. 163; G. M. Eliashberg and B. I. Ivlev, in *ibid*.211.

- ⁷M. G. Blamire, E. C. G. Kirk, J. E. Evetts, and T. M. Klapwijk, Phys. Rev. Lett. **66**, 220 (1991).
- ⁸H. Pothier, S. Guéron, N. O. Birge, D. Esteve, and M. H. Devoret, Phys. Rev. Lett. **79**, 3490 (1997).
- ⁹F. Giazotto, T. T. Heikkilä, F. Taddei, R. Fazio, J. P. Pekola, and F. Beltram, Phys. Rev. Lett. **92**, 137001 (2004).
- ¹⁰J. P. Pekola, D. V. Anghel, T. I. Suppula, J. K. Suoknuuti, A. J. Manninen, and M. Manninen, Appl. Phys. Lett. **76**, 2782 (2000).
- ¹¹A. Brinkman, A. A. Golubov, H. Rogalla, F. K. Wilhelm, and M. Yu. Kupriyanov, Phys. Rev. B 68, 224513 (2003).
- ¹²A. I. Larkin and Yu. N. Ovchinnikov, in *Nonequilibrium Superconductivity*, edited by D. N. Langenberg and A. I. Larkin (North Holland, Amsterdam, 1986), p. 493.
- ¹³N. B. Kopnin, *Theory of Nonequilibrium Superconductivity* (Clarendon, Oxford, 2001).