

## Slow Lévy flights

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Among Markovian processes, the hallmark of Lévy flights is superdiffusion, or faster-than-Brownian dynamics. Here we show that Lévy laws, as well as Gaussian distributions, can also be the limit distributions of processes with long-range memory that exhibit very slow diffusion, logarithmic in time. These processes are path dependent and anomalous motion emerges from frequent relocations to already visited sites. We show how the central limit theorem is modified in this context, keeping the usual distinction between analytic and nonanalytic characteristic functions. A fluctuation-dissipation relation is also derived. Our results may have important applications in the study of animal and human displacements.

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### I. INTRODUCTION

Lévy flights (LFs) represent one of the most important extensions of the central limit theorem (CLT), a cornerstone of probability theory [1,2]. Lévy flights are sums of independent and identically distributed random variables that admit non-Gaussian limit laws due to their very large fluctuations. They find physical applications in laser cooling [3], optics [4], and chaotic transport [5]. Lévy flights are also paradigmatic of superdiffusive processes, i.e., anomalous types of transport where the characteristic diffusive length scale  $l(t)$  of an individual particle grows with time as  $t^\alpha$  with  $\alpha > 1/2$ , that is, faster than in classical Brownian motion (BM) [6–9].

In recent years, LFs (as well as the related Lévy walks [10]) have become prominent for modeling diffusion in a variety of complex systems. Power-law distributions of step lengths with diverging variance, a key feature of Lévy processes, are found to describe well the trajectories of immune cells in the brain [11], the displacements of animals [12–15], and hunter-gatherers [16,17] in their environments, and the travels of modern humans within and between cities [18–21]. However, the assumption of independence between steps does limit the applicability of genuine Lévy processes for modeling real systems, where non-Markovian effects and correlations can be strong. Deeper analysis of empirical data actually reveals that the diffusion of humans and animals (even those exhibiting Lévy patterns) is in general *subdiffusive* at large times, i.e., with  $l(t) \ll t^{1/2}$  [21–25]. Furthermore,  $l(t)$  commonly grows more slowly than a power law of time, namely, in a logarithmic way [21,24,25]: This behavior is even in sharper contrast to the superdiffusion of simple LFs.

Logarithmic diffusion can be generated in several ways, for instance, by continuous-time random-walk models with superheavy-tailed distributions of waiting times [26] or by certain iterated maps [27,28]. In the context of animal and human mobility, an important but little explored mechanism that may lead to very slow subdiffusion is spatial memory: Many living organisms actually keep revisiting familiar places [22–25,29,30]. Here we seek to understand, with the help of

a solvable model, how this type of memory can act as a self-attracting force that drastically constrains diffusion towards limited areas, giving rise to “home ranges,” and how this property can still be compatible with power-law-distributed step lengths.

The dynamics and limit distributions of constrained LFs are not well understood, except mainly for processes subjected to long waiting times or in external potentials [8,31]. Several limit theorems also exist for specific problems of sums of correlated random variables [32] and a few random walks with infinite memory of their previous displacements have exactly solvable first moments [33–35]. Yet very little is known about LFs composed of nonindependent steps, in particular processes with self-attraction. Self-attracting random walks are path-dependent processes where a walker tends to return to previously visited sites [36,37]. Numerical simulations and scaling arguments clearly show that self-attracting walks can exhibit subdiffusion [38–40]. These mathematically challenging processes cannot be readily analyzed with better known frameworks for subdiffusive phenomena, such as fractional Fokker-Planck equations [8] or scaled Brownian motions [41,42]. They are more related to diffusion in quenched disordered media [6], where some rigorous connections have been made with the Sinai model [43].

In this study we heuristically modify the CLT for processes that exhibit very slow diffusion and show that such modification exactly describes a class of self-attracting LFs and self-attracting random walks. Since the characteristic functions have a structure similar to that in the ordinary CLT, Gaussian and Lévy distributions emerge asymptotically in space, although the dynamics is strongly subdiffusive. We also derive a fluctuation-dissipation relation in the Gaussian case.

### II. GENERAL FORMULATION

Let  $P(n,t)$  be the probability that the position  $X_t$  of a particle at time  $t$  is  $n$  (where  $n$  and  $t$  are discrete), given that the particle is located at the origin  $n = 0$  at  $t = 0$ . We consider discrete one-dimensional walks, keeping in mind that discreteness is not relevant in the asymptotic limit. The results can also be extended straightforwardly to higher dimensions.

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We recall that for a standard random walk composed of  $t$  independent and identically distributed displacements  $\ell_i$  with distribution  $p(\ell)$ , the characteristic function of  $X_t$ , defined as  $\tilde{P}(k, t) \equiv \sum_{n=-\infty}^{\infty} e^{-ikn} P(n, t) = \langle e^{-ikX_t} \rangle$ , takes the form [9]

$$\tilde{P}(k, t) = \tilde{p}(k)^t = e^{\ln[\tilde{p}(k)]t}, \quad (1)$$

where  $\tilde{p}(k)$  is the characteristic function of  $\ell$ . Since  $\tilde{p}(0) = 1$  by normalization, in the unbiased ( $\langle \ell \rangle = 0$ ) and symmetric case, an expansion near  $k = 0$  gives

$$\tilde{p}(k) = 1 - C|k|^\mu + \dots \quad (2)$$

Two basic situations emerge: the analytic case  $\mu = 2$ , corresponding to  $\langle \ell^2 \rangle < \infty$  (and  $C = \langle \ell^2 \rangle / 2$ ), and the nonanalytic case  $0 < \mu < 2$  when  $\langle \ell^2 \rangle$  does not exist, due to a power-law decay of  $p(\ell)$ ,

$$p(\ell) \sim 1/|\ell|^{1+\mu}, \quad (3)$$

at large  $\ell$  [9]. Combining (1) and (2) yields the celebrated Gaussian-Lévy CLT

$$\tilde{P}(k, t) \rightarrow e^{-C|k|^\mu t}. \quad (4)$$

Equation (4) implies a scaling law  $P(n, t) \rightarrow t^{-1/\mu} f(n/t^{1/\mu})$  where the scaling function  $f(x)$  is a Gaussian distribution or a symmetric Lévy law  $L_{\mu,0}(x)$  for  $\mu = 2$  and  $0 < \mu < 2$ , respectively. The latter case is superdiffusive as the typical diffusion length is proportional to  $t^{1/\mu} \gg t^{1/2}$ .

Consider now a simple modification of Eq. (1): Suppose that for certain diffusion processes with memory or sums of correlated random variables (we do not need to specify a model at this point),  $\tilde{P}$  is not an exponential function of  $t$  but a power law

$$\tilde{P}(k, t) \simeq t^{-a(k)} = e^{-a(k)\ln t} \quad (5)$$

at large  $t$  and small  $k$ . The function  $a(k)$  satisfies  $a(0) = 0$ , owing to the normalization  $\tilde{P}(k=0, t) = 1$ . Again,  $a(k)$  can be generically analytic or nonanalytic near  $k = 0$ . In the first case, since  $\tilde{P}(k, t)^* = \tilde{P}(-k, t)$  and  $|\tilde{P}(k, t)| \leq 1$ , the Taylor expansion of the exponent must be of the form  $a(k) \simeq ia_1k + a_2k^2 + \dots$ , with  $a_1$  and  $a_2$  two real constants and  $a_2 > 0$ . For simplicity, we first consider  $a_1 = 0$ , or motion without bias.

In the nonanalytic case, the same arguments lead to  $a(k) \simeq a_\mu |k|^\mu$  with  $0 < \mu < 2$  a priori and  $a_\mu > 0$ . Inserting into (5), we see that the main difference from (4) is that the variable  $t$  is substituted by  $\ln t$ . Hence

$$P(n, t) \rightarrow \frac{1}{(\ln t)^{1/\mu}} f_\mu \left( \frac{n}{(\ln t)^{1/\mu}} \right), \quad (6)$$

where the limit laws  $f_\mu(x)$  are the same as in the ordinary CLT. If  $\mu = 2$ , diffusion is Gaussian but very slow,  $\langle X_t^2 \rangle = 2a_2 \ln t$ , in sharp contrast to BM, where  $\langle X_t^2 \rangle = 2Dt$ . [In this case, Eq. (6) should not be confused with the log-normal distribution, where the logarithm applies to the space variable, not the temporal one.] A basic Markovian example is, by construction, scaled Brownian motion, which is BM where the time  $T$  is rescaled as  $t = e^T$ . Such a process is also equivalent to BM with a time-dependent diffusion coefficient  $D(t)$  decaying as  $1/t$  at large  $t$  [42].

In the nonanalytic case, the situation looks paradoxical at first sight. The ensemble average  $\langle X_t^2 \rangle = \infty$  like in ordinary

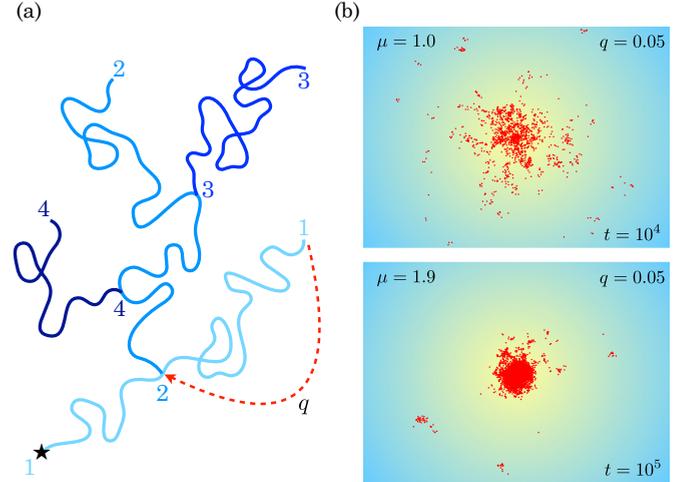


FIG. 1. (a) Schematic view of a process relocating at a constant rate  $q$  to sites occupied at previous times, these times being chosen stochastically. The numbers label the beginning and end of each excursion. Each end is followed by the beginning of the next excursion (arrow). (b) Two simulated trajectories corresponding to Lévy excursions with  $p(\ell) \sim 1/|\ell|^{1+\mu}$ , relocation rate  $q = 0.05$ , and relocation kernel given by Eq. (8) (panels at the same scale).

Lévy processes due to the broad tails of  $L_{\mu,0}(x)$  [or due to the fact that  $\partial^2 P(k, t)/\partial k^2$  does not exist at  $k = 0$ , from Eq. (5)]. Yet Eq. (6) also defines a typical diffusion length  $l(t) \propto (\ln t)^{1/\mu}$ , which grows extremely slowly. Therefore, based on this scaling length  $l(t)$ , motion is strongly subdiffusive and all the finite moments  $\langle |X_t|^v \rangle$  with  $v < \mu$  also evolve very slowly, as  $(\ln t)^{v/\mu}$ . Still, the process keeps superdiffusive features through the divergence of the second moment. This situation is reminiscent of scaling violation, which also arises in continuous-time random walks [44] or Lévy walks [10,45].

### III. RANDOM WALKS WITH RELOCATIONS

We now consider a concrete class of non-Markovian walks for which the above ideas apply. The processes of interest are self-attracting, namely, they tend to revisit locations visited in the past. Particular examples were studied numerically in [22,23] as animal movement models and theoretically in [25,46]. We present here a unified view of this class of processes.

Let  $q$  be a parameter ( $0 < q < 1$ ). At any time  $t$ , the walker chooses its next position according to the following rules.

(i) With probability  $1 - q$ , it performs a random displacement  $\ell$  drawn from a given distribution  $p(\ell)$  like in standard random walks or Lévy flights.

(ii) With the complementary probability  $q$ , it jumps (or resets) directly to the site occupied at some previous time  $t' \leq t$ . The time  $t'$  is chosen according to a given probability  $\pi_t(t')$ , or memory function, with  $\sum_{t'=0}^t \pi_t(t') = 1$  by normalization.

The rules are depicted in Fig. 1(a), with two simulated examples in Fig. 1(b). Note that in (ii), the next target site is chosen independently of its distance to the location  $X_t$  of the walker. If  $\pi_t(t') = \delta_{t',0}$ , the site chosen for revisit is unique (the origin), a case that corresponds to the well-studied random walk with resetting to the origin [47–50]. For more general

kernels, the walk is strongly path dependent but still described by a master equation

$$P(n, t+1) = (1-q) \sum_{\ell=-\infty}^{\infty} p(\ell) P(n-\ell, t) + q \sum_{t'=0}^t \pi_{t'} P(n, t'). \quad (7)$$

Standard random walks or Lévy flights are recovered for  $q = 0$ . If  $q \neq 0$ , the last term indicates that site  $n$  can be chosen to be occupied at time  $t+1$ , provided it was visited at the earlier time  $t'$ .

We first consider a uniform memory function, that is, independent of  $t'$ ,

$$\pi_{t'}(t') = \frac{1}{t+1}. \quad (8)$$

We call this case the preferential visit model (PVM): With such a kernel, rule (ii) is simply equivalent to choosing a given site  $n$  (among all visited sites) with probability proportional to the number of visits received by  $n$  since  $t=0$ . Therefore, the walker is prone to revisit familiar sites, at the expense of rarely visited ones. The moments  $\langle X_t^{2p} \rangle$  were calculated in [25] for the PVM with nearest-neighbor (NN) steps ( $\ell_i = \pm 1$ ) in rule (i). To solve Eq. (7) more generally, we define the Laplace transform of  $\tilde{P}(k, t)$ :

$$\hat{P}(k, \lambda) = \sum_{t=0}^{\infty} \lambda^t \sum_{n=-\infty}^{\infty} e^{-ikn} P(n, t). \quad (9)$$

By taking the double transform of Eq. (7) with the kernel (8) and writing  $\lambda^t/(t+1) = \lambda^{-1} \int_0^\lambda u^t du$ , we obtain

$$\hat{P}(k, \lambda) - 1 = (1-q) \tilde{p}(k) \lambda \hat{P}(k, \lambda) + q \int_0^\lambda du \frac{\hat{P}(k, u)}{1-u}. \quad (10)$$

Taking the derivative of Eq. (10), one obtains a first-order ordinary differential equation in the variable  $\lambda$ . As  $P(n, t=0) = \delta_{0,n}$ , the condition  $\hat{P}(k, 0) = 1$  must be enforced, leading to the exact solution

$$\hat{P}(k, \lambda) = (1-\lambda)^{-[1-a(k)]} [1 - (1-q) \tilde{p}(k) \lambda]^{-a(k)}, \quad (11)$$

with

$$a(k) = (1-q) \frac{1 - \tilde{p}(k)}{1 - (1-q) \tilde{p}(k)}. \quad (12)$$

We can infer the large- $t$  behavior of  $\tilde{P}(k, t)$  by studying the divergence of  $\hat{P}(k, \lambda)$  near  $\lambda = 1$ , with  $k$  fixed but small. Noting that  $a(k) \ll 1$ , Eq. (11) yields  $\hat{P}(k, \lambda) \simeq (1-\lambda)^{-[1-a(k)]}$ . This expression is simply inverted as

$$\tilde{P}(k, t) \simeq t^{-a(k)}, \quad (13)$$

as announced in (5). In the absence of bias, one can use Eq. (2), which, combined with (12), gives the exponent

$$a(k) \simeq \frac{1-q}{q} C |k|^\mu, \quad (14)$$

implying the limit law (6). We conclude that this random walk always diffuses logarithmically, unlike other reinforced walks

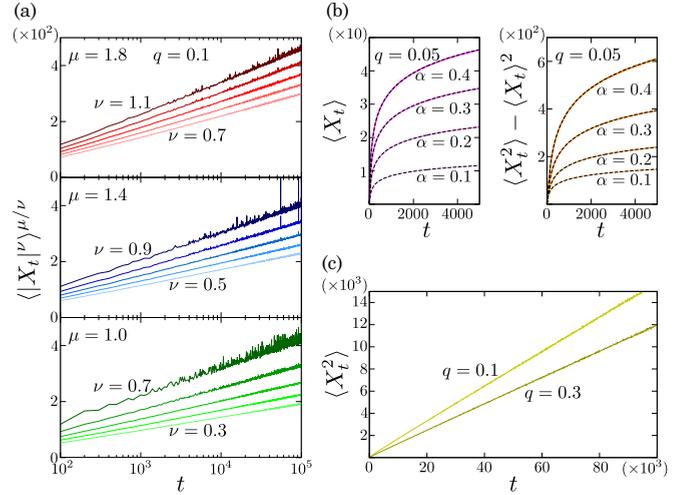


FIG. 2. Preferential visit model in one dimension. (a) Here  $\langle |X_t|^\nu \rangle^{\mu/\nu}$ , obtained from simulations with different  $\mu$  and  $\nu$  (averages over  $5 \times 10^5$  runs), is proportional to  $\ln t$  as expected. (b) Mean and variance of  $X_t$  for a NN walk with bias  $\alpha$  in rule (i). Colored solid lines are simulations and dark dashed lines, theory. (c) Normal diffusion for spatially uniform relocations.

that exhibit transitions to localized states [36,39]. Numerical simulations confirm the very slow dynamics, even when the excursions between relocations are Lévy flights, or  $\mu < 2$ : Perfect agreement with the prediction  $\langle |X_t|^\nu \rangle \sim (\ln t)^{\nu/\mu}$  for  $\nu < \mu$  is observed in Fig. 2(a). We emphasize again that the scaling function  $f(x)$  in this non-Markovian process is the same as for the underlying Markovian process between relocations (or with  $q = 0$ ). This property stems from the fact that the cumulant characteristic function  $\ln \tilde{p}(k)$  [Eq. (1)] and the function  $a(k)$  [Eq. (12)] have the same leading behavior at small  $k$ , except for a multiplicative constant. In other words, the analyticity or nonanalyticity of  $\tilde{P}(k, t)$  is preserved when  $q$  is set different from zero.

#### IV. GENERALIZATIONS

We now show that several extensions of the PVM also admit a propagator of the form given by Eq. (5).

##### A. Decaying memory

The results of the previous section do not change qualitatively by considering memory kernels other than a pure preferential one. For instance, the time in the past  $t'$  may be chosen not uniformly like in Eq. (8) but with a probability decaying with  $t-t'$ , the interval of time between a remembered occupation and the present time. Consider, for instance, a power-law memory decay

$$\pi_{t'}(t') = \frac{(t-t'+1)^{-\beta}}{\sum_{t''=0}^t (t-t''+1)^{-\beta}}, \quad (15)$$

with  $\beta > 0$  an exponent. Here the visits are still preferential, but with a tendency towards more recent sites (an effect actually observed in human mobility [51]). If  $\beta < 1$  the sum in (15) diverges at large  $t$  and can be substituted by an integral; by taking the Fourier transform of (7) and making the ansatz

$\tilde{P}(k, t) \simeq t^{-a(k)}$ , one obtains an integral equation for  $a(k)$ :

$$1 - (1 - q)\tilde{p}(k) = q(1 - \beta) \int_0^1 du (1 - u)^{-\beta} u^{-a(k)}. \quad (16)$$

Combining Eqs. (16) and (2) gives, at small  $k$ ,

$$a(k) \simeq \frac{1 - q}{q} \mathcal{F}(\beta) C |k|^\mu, \quad (17)$$

with

$$\mathcal{F}(\beta) = \left[ (1 - \beta) \int_0^1 du (1 - u)^{-\beta} \ln(1/u) \right]^{-1}.$$

Equation (17) shows that the scaling law (6) applies to more general processes than the PVM. [Equation (14) is recovered for  $\beta = 0$ .] Interestingly,  $\mathcal{F}(1) = \infty$ , which indicates that the scaling form (5) breaks down for  $\beta \geq 1$ . Actually, a calculation similar to the one above shows that, for  $\beta > 2$ , memory decays too fast to be relevant and the usual CLT (4) is recovered. Of course, these results do not mean that the aforementioned preservation property holds for arbitrary  $\pi_t(t')$ . For instance, for memory walks with  $1 < \beta < 2$  and steps  $\ell_i$  of finite variance, the process is non-Gaussian [46]. Likewise, Brownian random walks and Lévy flights subjected to stochastic resetting to the origin have asymptotic probability densities that are non-Gaussian [47] and non-Lévy [50], respectively.

### B. Model with bias

We now study the response of the non-Markovian walks (at fixed  $q$ ) to the presence of a constant forcing, namely, a bias  $\alpha \equiv \langle \ell \rangle = \sum_{\ell=-\infty}^{\infty} \ell p(\ell) \neq 0$ . Here we assume  $\langle \ell^2 \rangle < \infty$  or  $\mu = 2$ . By taking the first moment of Eq. (7), an equation for the average position  $\langle X_t \rangle \equiv \sum_{n=-\infty}^{\infty} n P(n, t)$  is obtained:

$$\langle X_{t+1} \rangle = (1 - q)[\langle X_t \rangle + \alpha] + q \sum_{t'=0}^t \pi_t(t') \langle X_{t'} \rangle \quad (18)$$

for any kernel  $\pi_t(t')$ . We now denote by  $\langle X_t^2 \rangle_0$  the mean-square displacement of the walker at zero bias. It is easy to show that  $\langle X_t^2 \rangle_0$  obeys exactly the same equation as (18), where  $\alpha$  has to be replaced by  $\langle \ell^2 \rangle_0 = \sum_{\ell=-\infty}^{\infty} \ell^2 p_0(\ell)$ , with  $p_0(\ell)$  unbiased. We deduce an Einstein fluctuation-dissipation relation (FDR)

$$\langle X_t \rangle = \frac{\alpha}{\langle \ell^2 \rangle_0} \langle X_t^2 \rangle_0. \quad (19)$$

The exact equality (19) is general: It is valid at all  $t$  and for any kernel  $\pi_t(t')$  (allowing one to recover results on the resetting to the origin with bias [48]). Despite being out of equilibrium, the FDR with constant bias in this system is the same as for ordinary random walks, where the response  $\langle X_t \rangle$  is entirely determined by the fluctuations at zero bias. With the kernel (15) and  $\beta < 1$ , the drift is thus logarithmic:  $\langle X_t \rangle \simeq \alpha \frac{1-q}{q} \mathcal{F}(\beta) \ln t$ , from Eqs. (19) and (17) with  $\mu = 2$ . The time evolution of the first moment  $\langle X_t \rangle$  is displayed on the left side of Fig. 2(b) for different parameter values.

In other words, the effective friction coefficient of the walker (proportional to  $\alpha \langle \dot{X}_t \rangle^{-1}$ ) grows linearly with  $t$ . This illustrates the nonstationarity emerging from long-range

memory and the increasingly sluggish dynamics caused by frequent relocations to the same preferred sites.

We further show that the combination of memory and bias has a drastic impact on the fluctuations of  $X_t$  around  $\langle X_t \rangle$ . We take, for example, the PVM with NN steps in rule (i) and expand Eq. (12), which is valid for any  $p(\ell)$ , near  $k = 0$ . Now using  $\tilde{p}(k) = 1 - i\alpha k - \frac{1}{2}k^2 + \dots$ , we obtain  $\tilde{P}(k, t) \simeq \exp[-i\mu_t k - \frac{1}{2}\sigma_t k^2]$ , which corresponds for  $P(n, t)$  to a Gaussian distribution of mean  $\mu_t$  and variance  $\sigma_t$ . We recover  $\mu_t = \alpha \frac{1-q}{q} \ln t$  [see (19)] and obtain, for  $\sigma_t$ ,

$$\sigma_t = \left[ \frac{1 - q}{q} + 2 \left( \frac{1 - q}{q} \right)^2 \alpha^2 \right] \ln t. \quad (20)$$

If  $q$  is small, the presence of a bias therefore strongly amplifies the fluctuations of  $X_t$ , as the second term in (20) is greater than zero and dominant. This effect is displayed on the right side of Fig. 2(b). For ordinary NN random walks, on the contrary, the bias decreases the fluctuations: In that case  $\sigma_t = (1 - \alpha^2)t$  and motion becomes deterministic at  $\alpha = 1$  (see, e.g., [52]).

## V. DISCUSSION AND CONCLUSION

In summary, we have shown that Lévy and Gaussian distributions can emerge generically far from the domain of applicability of the CLT, namely, in strongly subdiffusive path-dependent processes. We emphasize that the processes studied here exhibit subdiffusion because the relocation sites are selected heterogeneously in space. This situation is also encountered in the resetting to the origin, an extreme case where only one site receives all relocations, causing the typical diffusion length  $l(t)$  to tend to a constant [47]. To illustrate the importance of uneven relocations, one may by contrast consider a NN random walk, which, in rule (ii) above, relocates to a site chosen randomly and uniformly among the visited sites. In this case,  $l(t)$  roughly obeys  $dl/dt \sim (2R/l)[(R/2)/(1/q)]$ , with  $R = \sqrt{2D/q}$  the characteristic diffusion scale between two relocations,  $2R/l$  being the probability of resetting near the edges of the territory covered by the walk. This leads to  $l(t) \sim \sqrt{4Dt}$ , a normal diffusive behavior, which is qualitatively confirmed by the numerical simulations of Fig. 2(c).

The emergence of logarithmic diffusion can be understood qualitatively by drawing, from Fig. 1(a), an analogy to a branching random walk (see, e.g., [53,54]). Consider an initial normal random walk with a constant branching rate  $q_b$ . At each branching event, a new random walk is created that starts from the current position of the parent walk. The walks are independent, do not disappear, and all branch at the same rate  $q_b$ . The process follows until it is stopped at some final time  $T$ . Let us then imagine a single walker starting at the origin and following the paths left by all the branches, from the oldest to most recent, relocating at the start of the next branch when reaching the end of a branch. The average number of branches at time  $T$  is  $N_b(T) = e^{q_b T}$  and the total number of steps needed for the single walker to walk along all of them is  $t \simeq \int_0^T d\tau N_b(\tau) \simeq e^{q_b T} / q_b$ . At time  $t$ , the single walker will be at a typical distance  $l(t)$  from the origin, with  $l(t)^2 \sim T \simeq \frac{1}{q_b} \ln t$ . This form is surprisingly similar to our

result  $\langle X_t^2 \rangle \simeq \frac{1}{q} \ln t$  for the PVM at small  $q$ . The argument above can be repeated with branching Lévy flights, where  $l(t) \propto T^{1/\mu}$ , leading to a similar correspondence between the two models.

Note that the above analogy is only qualitative, as the PVM differs quantitatively from a set of branching random walks. Setting  $q_b = q$ , numerical simulations (not shown) indicate that, due to the rule of preferential visits, the relocation points in the memory model are distributed much more heterogeneously in space (namely, closer to the origin) than the branching points of the branching walks.

We conclude by mentioning that the processes studied here can explain two properties very often observed in human and animal mobility [15, 19–21, 25]: (a) Power-law-distributed step lengths can coexist with a very slow diffusion in the

long term (i.e., home-range behavior) and (b) the occupation of space by an individual within its home range is very nonuniform. Lévy flights with relocations to visited places could be an efficient strategy for searching and exploiting renewable resources, a challenge faced by many living organisms [12, 55–57].

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