

Macroscopic Finite Size Effects in Relaxational Processes

Shlomo Havlin^{1,2}, Armin Bunde^{1,2}, and Joseph Klafter³

¹ Institut für Theoretische Physik III, Justus-Liebig-Universität Giessen, D-35392 Giessen, Germany

² Minerva Center and Department of Physics, Bar-Ilan University, Ramat Gan, Israel

³ School of Chemistry, Tel Aviv University, Tel Aviv, Israel

Abstract. We present results on dynamical processes that exhibit a stretched exponential relaxation. When the relaxation is a result of two competing exponential processes, the size of the system, although macroscopic, play a dominant role. There exist a crossover time t_x that depends *logarithmically* on the size of the system, above which, the relaxation changes from a stretched exponential to a simple exponential decay. The decay rate also depends *logarithmically* on the size of the system. The results are relevant to large-scale Monte-Carlo simulations and should be amenable to experiments in low-dimensional macroscopic systems and mesoscopic systems.

Many relaxational processes in macroscopic systems are characterized by a relaxation function $Q(t)$ that exhibits a stretched exponential behavior,

$$Q(t) \sim Q(0) \exp[-(t/\tau)^\beta], \quad (1)$$

where $0 < \beta < 1$. Examples include viscoelastic relaxation [1], dielectric relaxation [2], glassy relaxations [3–5], relaxation in polymers [6,7] and long-time decay in trapping processes [8]. Many more examples [9–15] suggest that (1) is common to a very wide range of phenomena and macroscopic materials.

The origin of the stretched exponential is not always clear. In many cases it is assumed to be the result of a competition between two exponential processes. In some cases, e.g., trapping processes at long times, this assumption is well established, while in others, such as relaxation in glassy materials, this assumption has been controversially discussed [16,17] and alternative models have been also suggested [10,18–20].

We have recently investigated the occurrence of stretched exponential behavior in finite systems, in cases where the relaxation arises due to two competing exponential processes [21]

We have found that: (a) the size of the system, although macroscopic, plays a dominant role in the relaxation time pattern, leading to an exponential decay sufficiently at long times; (b) the crossover time, t_x , to the

exponential depends logarithmically on the system size; (c) the rate of the exponential decay also depends logarithmically on the system size, and (d) in the special examples of the trapping and the hierarchically constrained dynamics models the exponential relaxation may enter before the stretched exponential is reached. These results are of relevance to experiments in confined systems, mesoscopic systems and to Monte-Carlo simulations. Our theoretical predictions on the finite size effects can serve as an experimental test for identifying the origin of the mechanism leading to stretched exponential decay.

We assume that the relaxation function of the whole system can be represented by an integration over all possible states n , namely,

$$Q(t) = \int_0^\infty \Phi(n)Q(n,t)dn. \quad (2)$$

Here, $\Phi(n)$ is the probability that state n is occupied and $Q(n,t)$ is the dynamic relaxation of the n -th state.

Usually, in the case of a stretched exponential behavior, $\Phi(n)$ is assumed to behave as $\Phi(n) \sim \exp(-an^\alpha)$, while $Q(n,t)$ decays exponentially with time as $Q(n,t) \sim \exp(-bt/n^\gamma)$. A number of dynamical models that yield a stretched exponential decay can be formulated in terms of Eq.(2). These include the long-time behavior in the trapping problem [8], the target problem [20], direct energy transfer [20], trapping of nonidentical interacting particles [23], hierarchically constrained dynamics [16], models for relaxation in microemulsions and molecular glasses [24] and others.

We now concentrate on three examples: (i) A particle diffusing in a d -dimensional system with randomly distributed static traps, where we are interested in the survival probability $Q(t)$ of a particle. Here the state n represents a particle in a trap-free region of linear size n ; $\Phi(n)$ is the probability for the occurrence of a size n trap-free region, and $Q(n,t)$ is the survival probability of the particle in this region [8]. The exponent α is the dimension d of the system, and $\gamma = 2$ due to the diffusional motion. (ii) A linear system (chain) along which two types of particles (A and B) are diffusing and interacting via hard core interaction. However, only type A can be trapped by static traps which are randomly distributed along the chain. Here, $Q(t)$ is the survival probability of particles of type A , $\Phi(n)$ is the probability that a free trap region of size n occurs, and $Q(n,t)$ is the survival probability of a type A particle to survive in this region. The exponent α is the dimension of the system $\alpha = 1$ and $\gamma = 4$ is due to diffusion in the presence of hardcore interactions [25]. (iii) Hierarchically constrained dynamics, a model that has been proposed to account for glassy relaxation [16]. This model assumes that the relaxation of level n populated by spins, occurs in stages, and the constraint imposed by a faster degree of freedom must relax before a slower degree of freedom can relax. This implies that the time scale of relaxation in one level is subordinated to the relaxation below. A possible realization considered in [16] and here is a system with a discrete series of levels where the relaxation

time of level n is $\tau_n \sim n^\gamma$ (corresponding to the exponential form of $Q(n, t)$ in (2)), and the weight factor of level n , is $\Phi(n) \sim e^{-an}$ [12], corresponding to $\alpha = 1$. The first exponential in (2) is accordingly the probability to occupy level n and the second exponential represents the decay of that level.

The evaluation of the long time behavior of the integral in (2) is performed using the method of steepest descent. The main contribution to the integral arises from the maximum of the integrand in (2), which is obtained from the minimum of the function, $-an^\alpha - bt/n^\gamma$, appearing in the exponent. This yields that the main contribution to (2) comes from

$$n^* \cong (\gamma bt/\alpha a)^{1/(\alpha+\gamma)}, \quad (3)$$

leading to 1 with $\beta = \alpha/(\alpha + \gamma) < 1$, and $\tau = (\alpha/b\gamma)a^{-\gamma/\alpha}(\gamma/(\gamma + \alpha))^{1+\gamma/\alpha}$.

However, as shown below, these arguments are valid only in the thermodynamic limit where the system size is infinite. For a finite number N of traps (in the trapping system) or a *finite* system with a finite number N of spins (in the hierarchical constraint system) the relaxation function depends explicitly on N . Since our discussion is quite general for systems described by (2), in what follows we refer below to traps and spins in the above examples as elements.

For a single finite system consisting of N elements, the relaxation function $Q(t)$ represents an average quantity over the N elements,

$$Q(t) = \frac{1}{N} \sum_{\{n\}} m(n)Q(n, t), \quad (4)$$

where the sum is over all possible states n and $m(n)$ is the number of elements at state n , with $\sum_{\{n\}} m(n) = N$. Since the sum in (4) is over exponential functions, the value of $Q(t)$ will fluctuate for different sets of N . There will be a distribution of $Q(t)$, and we are interested in the typical $Q(t)$, which is around the peak of this distribution.

In the thermodynamic limit $N \rightarrow \infty$, all states n are occupied, $m(n)/N$ can be identified with $\Phi(n)$ and (2) follows. For N finite, in contrast, there exists a characteristic "maximum" state $n = n_{\max}(N)$, and this n_{\max} should replace the upper limit (∞) in (2),

$$Q(t) = \int_0^{n_{\max}} \Phi(n)Q(n, t) dn. \quad (5)$$

To estimate how n_{\max} depends on N , we note that the typical number of states n in a sample of N elements is $Z(n) \cong N\Phi(n) \cong N \exp(-an^\alpha)$. States with $Z(n) \ll 1$ will not occur in a typical system of N elements, and this yields

$$n_{\max} \cong \left(\frac{\ln N}{a} \right)^{1/\alpha}. \quad (6)$$

If $n^* \ll n_{\max}$, the upper limit in (2) can be approximated by infinity and thus leads to (1). However, if $n^* \gg n_{\max}$ the main contribution to (5) will not be from the maximum of the integrand, which is outside the range of integration, but from n_{\max} . Thus, for $n^* \gg n_{\max}$ we expect

$$Q(t) \cong Q(0)e^{-bt/n_{\max}^\gamma} \quad (7)$$

where the time constant of the relaxation, n_{\max}^γ , scales as $(\ln N)^{\gamma/\alpha}$. The crossover time from a stretched exponential (1) to an exponential (7) can be estimated from the condition $n^* = n_{\max}$, from which follows

$$t_x \cong \frac{\alpha a}{\gamma b} \left(\frac{\ln N}{a} \right)^{1+\gamma/\alpha}. \quad (8)$$

The striking point in (8) is the logarithmic dependence on N , which puts t_x in the range of observable time scales measurable in mesoscopic and even macroscopic systems. Indeed, the corresponding relaxation value $Q(t_x)$ scales as

$$Q(t_x) \sim N^{-\alpha/\gamma}, \quad (9)$$

independent of the microscopic parameters a and b . For the above three cases we find: (i) In the case of the trapping relaxation mechanism where $\alpha = d$ and $\gamma = 2$ we obtain,

$$Q(t_x)/Q(0) \sim N^{-d/2}. \quad (10)$$

(ii) In the non identical case $\alpha = 1$ and $\gamma = 1$ and thus

$$Q(t_x)/Q(0) \sim N^{-1/4}. \quad (11)$$

(iii) In the hierarchical constraint dynamics

$$Q(t_x)/Q(0) \sim N^{-1/\gamma}. \quad (12)$$

It is known [8e,23] that in both examples, for an infinite system, the stretched exponential behavior of (1) sets in only at very long times. Thus we expect that in the finite system, the crossover will mask the stretched-exponential pattern.

To test our analytical approach, we performed new Monte Carlo simulations on two cases (i) and (iii), the trapping model (case (i)) and the hierarchical constraint model (case (iii)). In the trapping model, we consider one and two dimensional systems with a fixed concentration $c = 0.5$ of randomly distributed traps, and vary the size N/c of the system. We calculated numerically the survival probability $Q(t)$ of a particle as a function of t and N . In the hierarchical model we have chosen $\tau_n \sim n$ i.e., $\gamma = 1$. We calculated the relaxation function for system sizes varying from $N = 10^2$ to $N = 10^5$.

As mentioned earlier, the relaxation function fluctuates for different sets of N . For obtaining the typical behavior of $Q(t)$, we have considered therefore

the "typical" average $Q(t)_{\text{typ}} \equiv \exp(\langle \ln Q(t) \rangle)$, where the brackets denote an average over many sets of N elements. Note that an arithmetic average over M sets of N elements can not be employed here, since it leads to a result identical for a larger system with $M \times N$ elements (see [4]). For a discussion of typical averages see [26]. For simplicity, we shall drop the index "typ" in the following.

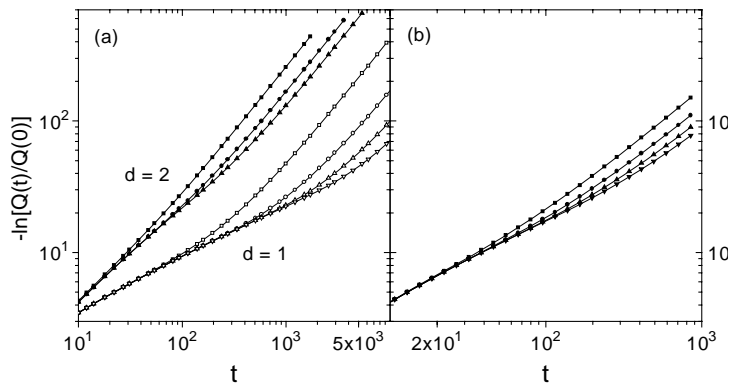


Fig. 1. Plot of $-\ln[Q(t)/Q(0)]$ as a function of t in a double logarithmic presentation for (a) the trapping model in $d = 1$ and $d = 2$, and (b) the hierarchical constraint model, for several system sizes. For the trapping model, the system sizes are $N = 2 \cdot 10^3$ (open square), $2 \cdot 10^5$ (open circle), $2 \cdot 10^7$ (open up triangle), $2 \cdot 10^9$ (open down triangle) in $d = 1$, and $N = 9 \cdot 10^2$ (full square), $9 \cdot 10^4$ (full circle), $9 \cdot 10^6$ (full up triangle) in $d = 2$. For the hierarchical model, the system sizes are $N = 10^2$ (full square), 10^3 (full circle), 10^4 (full up triangle), 10^5 (full down triangle).

Figure 1 shows $-\ln[Q(t)/Q(0)]$ as a function of t in a double logarithmic plot for (i) the trapping model in $d = 1$ and $d = 2$, and (iii) the hierarchical constraint model, both for several system sizes. In all cases, a crossover from an exponent $\beta < 1$ (at small t) towards $\beta = 1$ (at large t) can be easily recognized. The crossover time t_x shifts towards larger values when N increases.

To study the crossover behavior in a more quantitative manner, we have plotted in Fig. 2 the local exponents β obtained from the local slopes of Fig. 1, as a function of t . In both systems, for a fixed system size N , β first decreases with t , reaches a minimum value at a certain time that can be identified with t_x , and then increases monotonically with time towards $\beta = 1$. The figure shows that the minimum value of β has not yet reached its asymptotic value predicted for infinite systems, i.e., $\beta = 1/3$ ($d = 1$) and $\beta = 1/2$ ($d = 2$) for the trapping system and $\beta = 1/2$ for the hierarchical system.

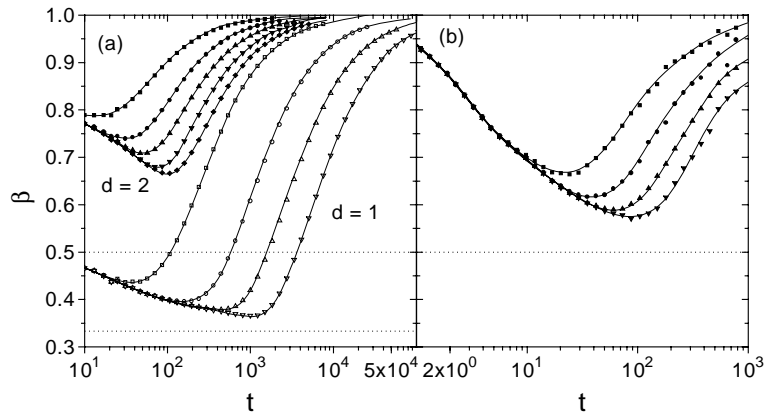


Fig. 2. Plot of the local exponents β calculated from the successive slopes of the corresponding curves in (1), (a) for the trapping model and (b) for the hierarchical model. The horizontal dashed lines represent the corresponding asymptotic ($N \rightarrow \infty$, $t \rightarrow \infty$) values of β .

To show the dependence of the crossover time t_x on the system size N we have plotted, in Fig. 3, the values of $t_x^{\alpha/(\alpha+\gamma)}$ as a function of $\ln N$. The crossover time was obtained numerically from the position of the minima of the curves in Fig. 2. The resulting straight lines are in full agreement with the prediction of (8), supporting our analytical approach.

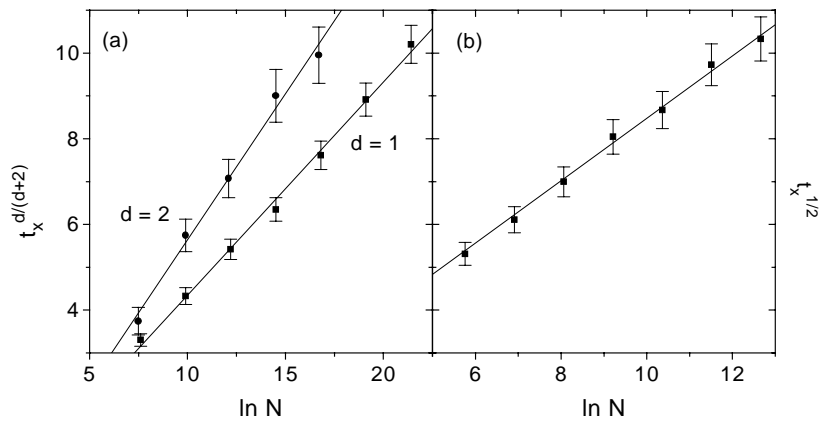


Fig. 3. Plot of $t_x^{\alpha/(\alpha+\gamma)}$ as a function of $\ln N$, for (a) the trapping model and (b) the hierarchical model. The straight line supports (8). The crossover times t_x were obtained from the positions of the minima of Fig. 2

In the following we discuss the relevance of our results to Monte-Carlo simulations and experiments. There exists a long standing puzzle in Monte-Carlo simulations of the trapping problem in $d = 2$ and 3 , that the predicted stretched exponential could not be observed [8], even for survival probabilities $Q(t)/Q(0)$ down to 10^{-21} in $d = 2$ [8b] and 10^{-67} in $d = 3$ [8g].

The finding of the logarithmic dependence of $Q(t)$ on the system size N explains this puzzle. The Monte-Carlo simulations in $d = 2$ and 3 were typically performed on 10^3 configurations with about 10^4 traps, which is equivalent to having a single system with $N \sim 10^7$ traps. Using (10), we expect for $N = 10^7$ traps $Q(t_x)/Q(0) \cong 10^{-7}$ in $d = 2$. Indeed, for times above t_x the exponent β approaches unity as predicted by our theory and seen clearly in Fig. 2a. Moreover, for this system size β never reaches the predicted thermodynamic value $\beta = 0.5$, the minimum value of β is about 0.65 . For $d = 3$, $Q(t_x)/Q(0) \cong 10^{-11}$ thus for smaller survival values ($t > t_x$) one again expects increasing values of β approaching unity. This explains the exponential decay found in the early Monte-Carlo simulations. Our results show that this is not an artefact but due to the finite size of the system. Moreover, they clearly indicate that the thermodynamic limit can not even be reached in one-dimensional macroscopic systems.

It would be of interest to test the above prediction experimentally by preparing experimental realizations where size effects can be controlled. Equations (8) and (10) suggest that the behavior around the crossover can be measured experimentally. For the trapping problem in linear systems, which has been studied experimentally [27,28], we expect for 10^8 sites and concentrations of traps c between 10^{-4} and 10^{-2} , that $Q(t_x)/Q(0) \sim 10^{-2} \div 10^{-3}$, which is a survival range that can be detected experimentally. For the non identical particles (case (iii)), we expect for 10^8 sites and concentration of traps c between 10^{-4} and 1 that $Q(t_x)/Q(0) \sim 10^{-1} \div 10^{-2}$ which is a survival range that can be well detected experimentally. The same arguments are valid for the target problem and therefore a similar crossover from stretched exponential to exponential decay is expected in relaxation experiments in low dimensional geometries [29]. Mesoscopic systems such as quantum dots, are also promising candidates for experiments where the crossover can be relevant. Identifying the logarithmic size dependence in experiments may provide support to the theories claiming that the observed stretched exponential is due to competing exponential processes, represented by (2).

This work was supported by the German Israeli Foundation (GIF).

References

1. R. Kohlrausch, Ann Phys. (Leipzig) **12**, 393 (1847)
2. G. Williams and D. C. Watts, Trans. Faraday Soc. **66**, 80 (1970)
3. V. Chamberlin, G. Mozurkewich and R. Orbach, Phys. Rev. Lett. **52**, 867 (1984)
4. F. Mezei and A. P. Murani, J. Magn. Mater. **14**, 211 (1979)

5. A. Plonka, *The Dependent Reactivity of Species in Condensed Matter* (Springer-Verlag, New York 1986)
6. A. A. Jones et al., *Macromolecules* **16**, 658 (1983)
7. K. L. Li et al., *Macromolecules* **21**, 2940 (1983)
8. (a) N. D. Donsker and S. R. S. Varadhan, *Commun. Pure Appl. Math.* **32**, 721 (1979); (b) P. Grassberger and Procaccia, *J. Chem. Phys.* **77**, 6281 (1982); (c) J. Klafter, G. Zumofen, A. Blumen, *J. Phys. Lett.* **45**, L49 (1984); (d) I. Webman, *Phys. Rev. Lett.* **52**, 220 (1984); (e) S. Havlin, M. Dishon, J. E. Kiefer, G. H. Weiss, *Phys. Rev. Lett.* **53**, 407 (1984); (f) M. Fixman, *Phys. Rev. Lett.* **52**, 791 (1984); (g) J.K. Anlauf, *Phys. Rev. Lett.* **52**, 1845 (1984)
9. A. K. Jonscher, *Nature* **267**, 673 (1977)
10. K. L. Ngai, *Comments Solid State Phys.* **9**, 127 (1979); **9**, 141 (1980)
11. K. Funke, *Prog. Solid St. Chem.* **22**, 11 (1993)
12. J. Klafter, M. F. Shlesinger, *Proc. Natl. Acad. Sci. U.S.A.* **83**, 848 (1986)
13. H. Scher, D. Bendler and M. Shlesinger, *Physics Today*, January 1991
14. J. C. Phillips, *Rep. Prog. Phys.* **59**, 1133 (1996)
15. J. C. Rasaiah, J. Zhu, J. B. Hubbard, and R. J. Rubin, *J. Chem. Phys.* **93**, 5768 (1990);
16. R. G. Palmer, D. L. Stein, E. Abrahams and P. W. Anderson, *Phys. Rev. Lett.* **53**, 958 (1984)
17. W. Götze and L. Sjögren, *Rep. Prog. Phys.* **55**, 241 (1992)
18. M. H. Cohen and G. S. Grest, *Phys. Rev. B* **24**, 4091 (1981)
19. M. F. Shlesinger and E. W. Montroll, *Proc. Natl. Acad. Sci. U.S.A.* **81**, 1280 (1984)
20. A. Blumen, J. Klafter, G. Zumofen, in *Optical Spectroscopy of Glasses*, ed. I. Zchokke (Reidel, Dordrecht 1986)
21. A. Bunde, S. Havlin, J. Klafter, G. Graff and A. Shehter, *Phys. Rev. Lett.* **78**, 3338 (1997)
22. J. C. Phillips, J. C. Rasaiah and J. B. Hubbard, *Phys. Rev. Lett.* , **80** 5453 (1998); A. Bunde, S. Havlin and J. Klafter, *Phys. Rev. Lett.* , **80** 5454 (1998)
23. A. Bunde, L. L. Mosely, H. E. Stanley, D. Ben-Avraham and S. Havlin, *Phys. Rev. A* **34**, 2575 (1986)
24. F. Sciortino and P. Tartaglia, *Physica A* **231**, 191 (1996)
25. T. E. Harris, *J. Appl. Pro.* **2**, 323 (1965); P. M. Richards, *Phys. Rev. B* **16**, 1393 (1977); S. Alexander and P. Pincus, *Phys. Rev. B* **18**, 2011 (1978)
26. A. Bunde nad J. Dräger, *Phys. Rev. E* **52**, 52 (1995); J. Dräger and A. Bunde, *ibid*, **54**, 4596 (1996)
27. R. A. Auerbach and G. L. McPherson, *Phys. Rev. B* **33**, 6815 (1986)
28. R. Knockenmuss and H. U. Gudel, *J. Chem. Phys.* **86**, 1104 (1987)
29. J. M. Drake et al., *Phys. Rev. Lett.* **61**, 865 (1988)