Diffusion laws in dendritic spines

David Holcman · Zeev Schuss

Received: 1 August 2011 / Accepted: 27 October 2011 / Published online: 27 October 2011
© 2011 Holcman, Schuss; licensee Springer. This is an Open Access article distributed under the terms of the Creative Commons Attribution License

Abstract Dendritic spines are small protrusions on a neuronal dendrite that are the main locus of excitatory synaptic connections. Although their geometry is variable over time and along the dendrite, they typically consist of a relatively large head connected to the dendritic shaft by a narrow cylindrical neck. The surface of the head is connected smoothly by a funnel or non-smoothly to the narrow neck, whose end absorbs the particles at the dendrite. We demonstrate here how the geometry of the neuronal spine can control diffusion and ultimately synaptic processes. We show that the mean residence time of a Brownian particle, such as an ion or molecule inside the spine, and of a receptor on its membrane, prior to absorption at the dendritic shaft depends strongly on the curvature of the connection of the spine head to the neck and on the neck’s length. The analytical results solve the narrow escape problem for domains with long narrow necks.

1 Introduction

Recognized more than 100 hundreds years ago by Ramón y Cajal, dendritic spines are small terminal protrusions on neuronal dendrites, and are considered to be the
main locus of excitatory synaptic connections. The general spine geometry consists of a relatively narrow cylindrical neck connected to a bulky head (the round part in Figure 1). Their geometrical shape correlates with their physiological function [1–6]. More than three decades ago, the spine-dendrite communication associated with the particle transfer was already anticipated [7] to be mediated not only by pure diffusion but it was hypothesized to involve other mechanisms such as twitching. This idea was reinforced by the findings [8] that inside the spine, the cytoplasmic actin is organized in filaments, involved in various forms of experimentally induced synaptic plasticity by changing the shape or volume of the pre- and postsynaptic side and by retracting and sprouting synapses. The fast dendritic spine contraction was finally confirmed in cultured hippocampal neurons [9] and consequences were studied theoretically in [10–12]. Interestingly, a serial electron microscopy and three-dimensional reconstructions of dendritic spines from Purkinje spiny branchlets of normal adult rats allowed to relate spine geometry to synaptic efficacy [1]. This image reconstruction approach leads to the conclusion that the cerebellar spine necks are unlikely to reduce transfer of synaptic charge by more than 5-20%, even if their smooth endoplasmic reticulum were to completely block passage of current through the portion of the neck that it occupies. The constricted spine neck diameter was proposed to isolate metabolic events in the vicinity of activated synapses by reducing diffusion to neighboring synapses, without significantly influencing the transfer of synaptic charge to the postsynaptic dendrite [1].

Change of spine morphology can be induced by synaptic potentiation protocols [13–15] and indeed intracellular signaling such as calcium released from stores alters the morphology of dendritic spines in cultured hippocampal neurons. These changes in geometry can affect the spine-dendrite communication. One of the first quantitative assessment of geometry was obtained by a direct measurement [16] of diffusion though the spine neck. Using photobleaching and photorelease of fluorescein dextran, by generating concentration gradients between spines and shafts in rat CA1 pyramidal neurons, the time course of re-equilibration was well approximated by a single exponential decay, with a time constant in the range of 20 to 100 ms. The role of the spine neck was further investigated using flash photolysis of caged calcium [3, 17]
and theory [18], and the main conclusion was that geometrical changes in the spine neck such as the length or the radius are key modulator of the spine-dendrite communication [12, 19, 20], affecting calcium dynamics. However, in all these studies, the nature of the connection between the neck and head was not considered. The theoretical studies [19, 21] considered non-smooth connection only of the head to the narrow cylindrical neck (Figure 1) and did not account for any effect of curvature. This is precisely the goal of this article to investigate the consequences of this connection.

The connection between the head and the neck is not only relevant for the three-dimensional diffusion, but also essential to the analysis of other synaptic properties. Indeed, synaptic transmission and plasticity involve the trafficking of receptors [22–27] such as AMPA or NMDA receptors (AMPARs or NMDARs) that mediate the glutamatergic-induced synaptic current. Single particle approaches have further [28, 29] revealed the heterogeneity of two-dimensional trajectories occurring on the neuron surface, suggesting that there are several biophysical processes involved in regulating the receptor motion. In addition, the number and type of receptors that shape the synaptic current [23] could be regulated by the spine geometry. This question was further explored theoretically [30, 31], using asymptotic expressions for the residence time and experimentally [32] by monitoring the movements of AMPARs on the surface of mature neurons using FRAP. Employing a combination of confocal microscopy and photobleaching techniques in living hippocampal CA1 pyramidal neurons, a correlation between spine shape parameters and the diffusion and compartmentalization of membrane-associated proteins was recently confirmed [33]. Lateral diffusion seems to be a constitutive process of AMPAR trafficking; it depends on spine morphology and is restricted by the spine geometry [34].

In this article, we develop a method for computing the NET from composite spine-like structures that consist of a relatively large compartment (head) $\Omega_1$ and a narrow cylindrical neck of cross section $|\partial \Omega_a|$ and length $L$ (see Figure 2). Our connection formula is given as

$$\bar{\tau}_{x \to \partial \Omega_a} = \bar{\tau}_{x \to \partial \Omega_1} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{|\partial \Omega_a|D}. \quad (1)$$

The connection between the two parts in the context of the NET problem was attempted in [21, 35] for the oversimplified model of a discontinuous connection. Here, we study a large class of connections and reveal the role of curvature in the spine-neck connection in regulating diffusion flux through narrow necks. More specifically, we study here the residence time of a Brownian particle from the spine head to the absorbing end of the spine neck moving either on the surface or inside the spine. We use the results of [36, 37] for the mean first passage time (MFPT) to an absorbing boundary at the end of a cusp-shaped protrusion in the head. They account for the effects of curvature generated by the neck-head connection in the spine. The reciprocal of the MFPT is the rate of arrival (probability flux) of Brownian particles from the head to the dendrite [38]. We calculate the narrow escape time (NET) from spine-shaped domains with heads connected smoothly and non-smoothly to the neck.
The composite domain consists of a head $\Omega_1$ connected by an interface $\partial\Omega_i$ to narrow neck $\Omega_2$. The entire boundary is reflecting, except for a small absorbing part $\partial\Omega_a$.

2 The NET from a domain with a bottleneck

We consider two- and three-dimensional composite domains $\Omega$ that consist of a head $\Omega_1$ connected through a small interface $\partial\Omega_i$ to a narrow cylindrical neck $\Omega_2$. The boundary of $\Omega$ is assumed reflecting to Brownian particles, except the far end of $\Omega_2$, denoted $\partial\Omega_a$, which is absorbing. For example, in Figure 2, the interface $\partial\Omega_i$ is the black segment AB and the absorbing boundary $\partial\Omega_a$ is the segment CD at the bottom of the strip. The NET from such a composite domain cannot be calculated by the methods of [39–42], because the contribution of the singular part of Neumann’s function to the MFPT in a composite domain with a funnel or another bottleneck is not necessarily dominant. The method of matched asymptotic expansions requires different boundary or internal layers at a cusp-like absorbing window than at an absorbing window which is cut from a smooth reflecting boundary (see [43–46]). The methods used in [21, 35] for constructing the MFPT in a composite domain of the type shown in Figure 1d are made precise here and the new method extends to the domains of the type shown in Figure 1c.

First, we recount some basic facts about the NET [35, 39–41, 43–45, 47, 48]. The NET is the MFPT of a Brownian trajectory to a small absorbing part of the boundary of a domain, whose remaining boundary reflects Brownian trajectories. Refined asymptotic formulas for the NET were derived in [42, 46, 49, 50], and were used to estimated chemical reaction rates.

Consider Brownian motion $x(t)$ in a sufficiently regular bounded domain $\Omega$, whose boundary $\partial\Omega$ consists of a reflecting part $\partial\Omega_r$ and an absorbing part $\partial\Omega_a$. The expected lifetime of $x(t)$ in $\Omega$, given $x(0) = x \in \Omega$, is the MFPT $v(x)$ of $x(t)$ from $x$ to $\partial\Omega_a$ is the solution of the mixed boundary value problem [38]

$$\Delta v(x) = -\frac{1}{D} \quad \text{for } x \in \Omega \quad (2)$$

$$v(x) = 0 \quad \text{for } x \in \partial\Omega_a \quad (3)$$

$$\frac{\partial v(x)}{\partial n} = 0 \quad \text{for } x \in \partial\Omega_r, \quad (4)$$

where $\partial v(x)/\partial n$ is the normal derivative at the boundary point $x$. If the size of the absorbing part $\partial\Omega_a$ of the boundary is much smaller than the reflecting part $\partial\Omega_r$, the
MFPT $\bar{\tau} = v(x)$ is to leading order independent of $x \in \Omega_a$ and can be represented by the Neumann function $N(x, y)$ as

$$\bar{\tau} = v(y) = -\frac{1}{D} \int_{\Omega} N(x, y) \, dx - \int_{\partial \Omega_a} N(x, y) \frac{\partial v(x)}{\partial n} \, dS_x. \quad (5)$$

The sum of the integrals is independent of $y \in \Omega_a$ outside a boundary layer near $\partial \Omega_a$. The Neumann function is a solution of the boundary value problem

$$\Delta_x N(x, y) = -\delta(x - y) + \frac{1}{|\Omega|} \quad \text{for } x, y \in \Omega \quad (6)$$

and

$$\frac{\partial N(x, y)}{\partial n_x} = 0 \quad \text{for } x \in \partial \Omega, y \in \Omega \quad (7)$$

and is defined up to an additive constant [39, 47].

### 2.1 The MFPT from the head to the interface

In the two-dimensional case considered in [40] the interface $\partial \Omega_i$ is an absorbing window cut from the smooth reflecting boundary of $\Omega_1$, as in Figure 1d. The MFPT $\bar{\tau}_{x \rightarrow \partial \Omega_i}$ is the NET from the reflecting domain $\Omega_1$ to the small interface $\partial \Omega_i$ (of length $a$), such that $\epsilon = \pi |\partial \Omega_i| / |\partial \Omega_1| = \pi a / |\partial \Omega_1| \ll 1$ (this corrects the definition in [40]). It is given by

$$\bar{\tau}_{x \rightarrow \partial \Omega_i} = \frac{|\Omega_1|}{\pi D} \ln \frac{|\partial \Omega_1|}{\pi |\partial \Omega_i|} + O(1) \quad (8)$$

for $x \in \Omega_1$ outside a boundary layer near $\partial \Omega_i$.

In particular, if $\Omega_1$ is a disk of radius $R$, then for $x =$ the center of the disk,

$$\bar{\tau}_{x \rightarrow \partial \Omega_i} = \frac{R^2}{D} \left[ \log \frac{R}{a} + 2 \log 2 + \frac{1}{4} + O(\epsilon) \right], \quad (9)$$

averaging with respect to a uniform distribution of $x$ the disk

$$\bar{\tau}_{x \rightarrow \partial \Omega_i} = \frac{R^2}{D} \left[ \log \frac{R}{a} + 2 \log 2 + \frac{1}{8} + O(\epsilon) \right]. \quad (10)$$

When the interface $\partial \Omega_i$ (of length $a$) is located at an algebraic cusp with radius of curvature $R_c$ (see Figures 1c and 2), the MFPT is given in [36, 37] as

$$\bar{\tau} = \frac{|\Omega_1|}{4D \sqrt{2a / R_c}} (1 + O(1)) \quad \text{for } \epsilon \ll |\partial \Omega|. \quad (11)$$

In the case of Brownian motion on a spherical head of the surface of revolution obtained by rotating the curve in Figure 1d about its axis, $\Omega_1$ is a sphere of radius $R$ centered at the origin, connected to $\Omega_2$ by a circle $\partial \Omega_i$ centered on the north-south axis near the south pole, with small radius $a = R \sin \delta / 2$. The domain $\Omega_2$ is a right
Fig. 3 A surface of revolution with a funnel. The $z$-axis points down.

cylinder of radius $a$ connected to $\Omega_1$ at $\partial\Omega_i$, and the absorbing boundary $\partial\Omega_a$ is the circle of radius $a$ at the bottom of the cylinder. The MFPT from $\Omega_1$ to $\partial\Omega_i$ is given in [37, 41, 51–53] as

$$\bar{\tau}_{x\to\partial\Omega_i} = \frac{2R^2}{D} \log \frac{\sin \frac{\theta}{2}}{\sin \frac{\delta}{2}},$$

where $\theta$ is the angle between $x$ and the south-north axis of the sphere.

A surface of revolution generated by rotating a curve about an axis, as in Figure 3, with a funnel of diameter $\epsilon$ can be represented parametrically as

$$x = r(z) \cos \theta, \quad y = r(z) \sin \theta, \quad 0 \leq \theta < 2\pi, 0 \leq z < \Lambda,$$

where the axis of symmetry is the $z$-axis with $z = 0$ at the top of the surface and $z = \Lambda$ at the end of the funnel, $r$ is distance to the $z$-axis, and $r = r(z)$ is the equation of the generating curve. We have

$$r(z) = O(\sqrt{z}) \quad \text{near } z = 0$$

$$r(z) = a + \ell^{-\alpha}(\Lambda - z)^{1+\alpha}(1 + o(1)) \quad \text{for } \alpha > 0 \text{ near } z = \Lambda,$$

where $\ell$ has dimension of length. For $\alpha = 1$ the parameter $\ell$ is related to the radius of curvature $R_c$ at $z = \Lambda$ by $\ell = 2R_c$. For $\alpha > 0$ [37]

$$\bar{\tau}_{x\to\partial\Omega_i} \sim \frac{|\Omega_1|}{2D} \left( \frac{1}{\pi} \right)^{1+\alpha} \frac{\ell^{\alpha/1+\alpha}}{\sin \frac{\pi}{1+\alpha}},$$

where $|\Omega_1|$ is the entire area of the surface. In particular, for $\alpha = 1$ we get the MFPT

$$\bar{\tau}_{x\to\partial\Omega_i} \sim \frac{|\Omega_1|}{4D \sqrt{2a/R_c}}.$$

The case $\alpha = 0$ corresponds to a circular cap of a small radius $a$ cut from a closed surface.

The MFPT of Brownian motion from a solid ball $\Omega_1$ to a disk $\partial\Omega_i$ of small radius $a$ near the south pole is given by [42]

$$\bar{\tau}_{x\to\partial\Omega_i} = \frac{|\Omega_1|}{4aD} \left[ 1 + \frac{a}{\pi R} \log \frac{R}{a} + o \left( \frac{a}{R} \log \frac{R}{a} \right) \right].$$
(note that the MFPT is independent of $x$ up to second order, see [46]). For a general three-dimensional domain, $\Omega_1$ the MFPT to a circular cap $\partial\Omega_i$ cut from a smooth boundary is given by [42]

$$\bar{\tau}_{x \to \partial\Omega_i} = \frac{|\Omega_1|}{4aD} \left[ 1 + \frac{L_x + R_x}{2\pi} \frac{|\partial\Omega_i|}{|\partial\Omega_1|} \right]^{1/2} \log \sqrt{\frac{|\partial\Omega_1|}{|\partial\Omega_i|}} + o \left( \frac{|\partial\Omega_i|}{|\partial\Omega_1|} \log \sqrt{\frac{|\partial\Omega_1|}{|\partial\Omega_i|}} \right) + O(1)D,$$

(18)

where $L_x, R_x$ are the principal curvatures at a point $x$, and $|\partial\Omega_i| = \pi a^2$ is the area of the circular cap.

When the interface $\partial\Omega_i$ is a circular disk of radius $a$ at the end of an axisymmetric solid funnel, the MFPT is drastically affected and changes to

$$\bar{\tau}_{x \to \partial\Omega_i} = \frac{1}{\sqrt{2}} \left( \frac{R_c}{a} \right)^{3/2} \frac{|\Omega_1|}{R_cD} (1 + o(1)) \text{ for } a \ll R_c,$$

(19)

where $R_c$ is the radius of curvature at the end of the funnel [37].

### 3 Connecting a head to a narrow neck

We consider Brownian motion in a domain $\Omega$ that consists of a head, which is a regular bounded domain $\Omega_1$, and a narrow neck $\Omega_2$, which is a right circular or planar cylinder of length $L$, perpendicular to the boundary $\partial\Omega_1$, and of radius $a$ (see Figure 2). Thus, the interface $\partial\Omega_i$ between the head and the neck is a line segment, a circle, or a circular disk, depending on the dimension. We assume that $\partial\Omega_1 - \partial\Omega_i$ is reflecting and that the other basis of the neck, $\partial\Omega_a \subset \partial\Omega_2$, is absorbing for the Brownian motion. The length (or area) $|\partial\Omega_i|$ is given by

$$|\partial\Omega_i| = \begin{cases} a & \text{for a line segment} \\ 2\pi a & \text{for a circle} \\ \pi a^2 & \text{for a disk}. \end{cases}$$

(20)

The MFPT $\bar{\tau}_{x \to \partial\Omega_a}$ can be represented as [54], [38, Lemma 10.3.1, p. 388]

$$\bar{\tau}_{x \to \partial\Omega_a} = \bar{\tau}_{x \to \partial\Omega_i} + \bar{\tau}_{\partial\Omega_i \to \partial\Omega_a},$$

(21)

where the MFPT $\bar{\tau}_{\partial\Omega_i \to \partial\Omega_a}$ is $\bar{\tau}_{x \to \partial\Omega_a}$, averaged over $\partial\Omega_i$ with respect to the flux density of Brownian trajectories in $\Omega_1$ into an absorbing boundary at $\partial\Omega_i$ (see [38] for further details).

First, we calculate $\bar{\tau}_{\partial\Omega_i \to \partial\Omega_a}$ and the absorption flux at the interface. In the narrow neck $\Omega_2$ the boundary value problem (2)–(4) can be approximated by the one-dimensional boundary value problem

$$Du_{zz} = -1$$
where the value at the interface \( u(L) = u_H \) is yet unknown. The solution is given by

\[
  u(z) = -\frac{z^2}{2D} + Bz, \tag{22}
\]

so that

\[
  u(L) = u_H = -\frac{L^2}{2D} + BL, \tag{23}
\]

which relates the unknown constants \( B \) and \( u_H \). The constant \( B \) is found by multiplying Equation 2 by the Neumann function \( N(x, y) \), integrating over \( \Omega_1 \), applying Green’s formula, and using the boundary conditions (3) and (4). Specifically, we obtain for all \( y \in \partial \Omega_i \)

\[
  v(y) = -\frac{1}{D} \int_{\Omega_1} N(x, y) \, dx - \int_{\partial \Omega_i} N(x, y) \frac{\partial v(x)}{\partial n} \, dS_x + \frac{1}{|\Omega_1|} \int_{\Omega_1} v(x) \, dx. \tag{24}
\]

Approximating, as we may, \( v(y) \approx u(L) \) and using (23), we obtain

\[
  -\frac{L^2}{2D} + BL = -\frac{1}{D} \int_{\Omega_1} N(x, y) \, dx - \int_{\partial \Omega_i} N(x, y) \frac{\partial v(x)}{\partial n} \, dS_x + \frac{1}{|\Omega_1|} \int_{\Omega_1} v(x) \, dx. \tag{25}
\]

Because \( v(x) \) is the solution of the boundary value problem (2)–(4) in the entire domain \( \Omega = \Omega_1 \cup \Omega_2 \), the meaning of (25) is the connecting rule (21), where

\[
  \bar{\tau}_{x \to \partial \Omega_a} = \frac{1}{|\Omega_1|} \int_{\Omega_1} v(x) \, dx \tag{26}
\]

\[
  \bar{\tau}_{\partial \Omega_i \to \partial \Omega_a} = u(L) \tag{27}
\]

\[
  \bar{\tau}_{x \to \partial \Omega_c} = -\frac{1}{D} \int_{\Omega} N(x, y) \, dx - \int_{\partial \Omega_i} N(x, y) \frac{\partial v(x)}{\partial n} \, dS_x. \tag{28}
\]

Equation 26 gives the MFPT, averaged over \( \Omega_1 \). The averaging is a valid approximation, because the MFPT to \( \partial \Omega_i \) is constant to begin with (except in a negligible boundary layer). Equation 27 is the MFPT from the interface to the absorbing end \( \partial \Omega_a \) of the strip, and (28) follows from (5).

Matching the solutions in \( \Omega_1 \) and \( \Omega_2 \) continuously across \( \partial \Omega_i \), we obtain the total flux on \( \partial \Omega_i \) as

\[
  J = D \int_{\partial \Omega_i} \frac{\partial v(x)}{\partial n} \, dS_x = -(|\Omega_1| + |\Omega_2|), \tag{29}
\]
Noting that \( \partial v(x)/\partial n = -u'(0) = -B \), we get from (20) and (29) that

\[
B = \begin{cases} 
\frac{|\Omega_1|}{aD} + \frac{L}{D} & \text{for a line segment} \\
\frac{|\Omega_1|}{2\pi aD} + \frac{L}{D} & \text{for a circle} \\
\frac{|\Omega_1|}{\pi a^2 D} + \frac{L}{D} & \text{for a circular disk.}
\end{cases}
\]  
(30)

Finally, we put (21)–(30) together to obtain

\[
\bar{\tau}_{x \to \partial \Omega_a} = \bar{\tau}_{x \to \partial \Omega_i} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{|\partial \Omega_a D|}.
\]  
(31)

The MFPT \( \bar{\tau}_{x \to \partial \Omega_i} \) is given by (8)–(19) above.

3.1 The NET from two- and three-dimensional domains with bottlenecks

The expression (31) for the NET from a domain with a bottleneck in the form of a one-dimensional neck, such as a dendritic spine, can be summarized as follows. Consider a domain \( \Omega \) with head \( \Omega_1 \) and a narrow cylindrical neck \( \Omega_2 \) of length \( L \) and radius \( a \). The radius of curvature at the bottleneck in smooth connecting funnel is \( R_c \). In the two-dimensional case

\[
\bar{\tau}_{x \to \partial \Omega_a} = \begin{cases} 
\frac{|\Omega_1|}{\pi D} \ln \frac{|\partial \Omega_1|}{a} + \frac{O(1)}{D} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{aD} & \text{planar spine connected to the neck at a right angle} \\
\frac{\pi |\Omega_1|}{D} \sqrt{\frac{R_c}{a}} (1 + o(1)) + \frac{L^2}{2D} + \frac{|\Omega_1|L}{2\pi aD} & \text{planar spine with a smooth connecting funnel} \\
\frac{|\Omega_1|}{2\pi D} \log \frac{\sin \frac{\theta}{2}}{\sin \frac{\theta}{2}} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{2\pi aD} & \text{spherical spine connected to the neck at a right angle} \\
\frac{|\Omega_1|}{2D} \left( \frac{1}{(1 + \alpha)} \sin \frac{\pi}{1 + \alpha} \right) + \frac{L^2}{2D} + \frac{|\Omega_1|L}{2\pi aD} & \text{spherical spine with a smooth connecting funnel},
\end{cases}
\]  
(32)

where \( R \) is the radius of the sphere, \( a = R \sin \delta/2 \), and \( \theta \) is the initial elevation angle on the sphere. If \( |\Omega_1| \gg aL \) and \( L \gg a \), the last term in (32) is dominant, which is the manifestation of the many returns of Brownian motion from the neck to the head prior to absorption at \( \partial \Omega_a \) (see an estimate in [19]). The last line of (32) agrees with the explicit calculation in [37].

The NET of a Brownian motion from a three-dimensional domain \( \Omega \) with a bottleneck in the form of a narrow circular cylinder of cross-sectional area \( \pi a^2 \) is given
Fig. 4 Left: The NET of Brownian motion on a sphere with a bottleneck connected by a smooth funnel to the neck (dashed line), and with a non-smooth connection (continuous line). Right: The NET of Brownian motion in a ball with a bottleneck connected by a smooth funnel to the neck (dashed line), and with a non-smooth connection (continuous line).

by

\[ \tilde{\tau}_{x \to \Omega_a} = \begin{cases} 
\frac{1}{4aD} \left[ |\Omega_1| \left[ 1 + \frac{a}{\pi R} \log \frac{R}{a} \right] + \frac{O(1)}{D} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{\pi a^2 D} \right] \\
\text{solid spherical head of radius } R \text{ connected to the neck at a right angle} \\
\frac{1}{4aD} \left[ 1 + \frac{(Lx + Rx)}{2\pi} \left| \frac{\partial \Omega_a}{\partial x} \right|^{1/2} \right] \\
\times \log \left( \frac{|\partial \Omega_a|}{|\partial \Omega_1|} \right) + o \left( \sqrt{\frac{|\partial \Omega_a|}{|\partial \Omega_1|}} \log \sqrt{\frac{|\partial \Omega_a|}{|\partial \Omega_1|}} \right) \\
+ \frac{O(1)}{D} + \frac{L^2}{2D} + \frac{|\Omega_1|L}{\pi a^2 D} \\
\text{a general head connected to the neck at a right angle} \\
\frac{1}{\sqrt{2}} \left( \frac{R_c}{a} \right)^{3/2} \frac{|\Omega_1|}{R_c D} (1 + o(1)) + \frac{L^2}{2D} + \frac{|\Omega_1|L}{\pi a^2 D} \\
\text{a general head connected smoothly to the neck by a funnel}, 
\end{cases} \tag{33} \]

where \( R_c \) is the curvature at the cusp. The order 1 term can be computed for the sphere using the explicit expression of the Neumann-Green function [46]. Figures 4 and 5 show the NET for various parameters, such as the neck length and radius.

Finally, the influence of the neck length on the residence time is shown in Figure 5: changing the neck length modulates dramatically the residence time. Interestingly, the geometry of the connection affects much significantly the dimension two rather than that the three dimensional Brownian motion.
4 Discussion and conclusion

We have shown that the mean residence time (or flux) of Brownian particles inside a spine-like structure or on its surface depends strongly on the geometrical properties of both head and neck. Surprisingly, it also depends strongly on the smoothness of the connection between the two.

The application to a freely diffusing AMPA receptor, which is responsible for the excitatory synaptic current, shows that its motion on the spine membrane is strongly restricted by dendritic spine geometry. Our results can be used to estimate the residence time of the receptor on the membrane if interactions with any scaffolding molecules are neglected (the latter are mostly concentrated in a local microdomain called the PSD). Using Equation 32 for non-smooth geometry (Figure 1b) and for a spherical head of radius $R = 1 \mu m$, a neck length $L = 1 \mu m$, a neck radius $a = 0.1 \mu m$, and a diffusion coefficient $D = 0.1 \mu m^2/s$, we obtain from the third line of Equation 32 that the residence time is $\tau_{2D} \approx 260 s$, while for a smooth connecting geometry (Figure 1a) with a curvature radius of $R_c = 1 \mu m$ at the connecting neck-head $\tau_{2D} \approx 1150 s$ (line 2 in Equation 32). Evidently, the residence time is more than doubled, leading to the conclusion that there is a significant difference between the function of spines with smoothly and non-smoothly connected necks. We conclude from this analysis that an AMPA receptors that do not interact with the PSD stay on a typical dendritic spine between one and a half to 5 min on average and this residence time is controlled mostly by the geometrical properties of the spine.

We now consider the residence time for freely diffusing particles such as molecules, mRNA, and ions (e.g., calcium) inside a dendritic spine. For a calcium ion, the diffusion constant is about $D_{ca} = 400 \mu m^2/s$ [12]. Calcium ions that exit the spine only to the dendrite shaft at the end of the neck, but not through exchangers, give the following residence time estimates. Using formula 33 for a non-smooth connection between the spine head and the neck, we obtain that $\tau_{3D} \approx 195 ms$ (line 1), while for a smooth connection with radius of curvature $1 \mu m$, the residence time is $\tau_{3D} \approx 820 ms$ (line 3 in Equation 33). Interestingly, the mean residence time is tripled from the non-smooth to a smooth connection.
A remaining open question is to extend the present analysis to the case where many binding molecules can trap receptors. This effect should be expected to significantly increase the residence time inside a dendritic spine, as has already been observed in [55] for the case of a receptor inside the PSD. The present mathematical analysis of the residence time provides a solution to the narrow escape problem for domains with bottlenecks [21, 35]. Other generalizations of this study is to include the dynamics of many receptors [30, 56] or/and to study dendritic trafficking [57].

There are many other factors that affect the spine-dendrite communication with respect to calcium. This includes calcium pumps, endogenous buffers, calcium stores, the number and rates of exchangers. These mechanisms affect the residence of calcium in spines [58–60] and it would be interesting to add them in the present analysis.

**Competing interests**

The authors declare that they have no competing interests.

**Authors’ contributions**

DH and ZS contributed equally to the manuscript. All authors read and approved the final manuscript.

**Acknowledgements**

This research was supported by an ERC starting Grant.

**References**

1. Harris, K.M., Stevens, J.K.: Dendritic spines of rat cerebellar Purkinje cells: serial electron microscopy with reference to their biophysical characteristics. J. Neurosci. 12, 4455–4469 (1988)
2. Bourne, J.N., Harris, K.M.: Balancing structure and function at hippocampal dendritic spines. Annu. Rev. Neurosci. 31, 47–67 (2008)
3. Korkotian, E., Holcman, D., Segal, M.: Dynamic regulation of spine-dendrite coupling in cultured hippocampal neurons. Eur. J. Neurosci. 20(10), 2649–2663 (2004)
4. Hotulainen, P., Hoogenraad, C.C.: Actin in dendritic spines: connecting dynamics to function. J. Cell Biol. 189(4), 619–629 (2010)
5. Newpher, T.M., Ehlers, M.D.: Spine microdomains for postsynaptic signaling and plasticity. Trends Cell Biol. 5, 218–227 (2009)
6. Sheng, M., Hoogenraad, C.C.: The postsynaptic architecture of excitatory synapses: a more quantitative view. Annu. Rev. Biochem. 76, 823–847 (2007)
7. Crick, F.: Do dendritic spines twitch? Trends Neurosci. 5, 44–46 (1982)
8. Filipova, E., Delay, R.J.: Cytoplasmic actin in dendritic spines as a possible mediator of synaptic plasticity. J. Cell Biol. 95, 350–365 (1982)
9. Korkotian, E., Segal, M.: Spike-associated fast contraction of dendritic spines in cultured hippocampal neurons. Neuron 30(3), 751–758 (2001)
10. Holcman, D., Schuss, Z., Korkotian, E.: Calcium dynamic in dendritic spines and spine motility. Biophys. J. 87, 81–91 (2004)
11. Holcman, D., Schuss, Z.: Modeling calcium dynamics in dendritic spines. SIAM J. Appl. Math. 65(3), 1006–1026 (2005)
12. Holcman, D., Korkotian, E., Segal, M.: Calcium dynamics in dendritic spines, modeling and experiments. Cell Calcium 37(5), 467–475 (2005)
13. Papa, M., Segal, M.: Morphological plasticity in dendritic spines of cultured hippocampal neurons. Neuroscience 71(4), 1005–1011 (1996)
14. Collin, C., Miyaguchi, K., Segal, M.: Dendritic spine density and LTP induction in cultured hippocampal slices. J. Neurophysiol. 77(3), 1614–1623 (1997)

15. Korkotian, E., Segal, M.: Release of calcium from stores alters the morphology of dendritic spines in cultured hippocampal neurons. Proc. Natl. Acad. Sci. USA 96(21), 12068–12072 (1999)

16. Svoboda, K., Tank, D.W., Denk, W.: Direct measurement of coupling between dendritic spines and shafts. Science 272(5262), 716–719 (1996)

17. Bloodgood, B.L., Sabatini, B.L.: Neuronal activity regulates diffusion across the neck of dendritic spines. Science 310(5749), 866–869 (2005)

18. Holcman, D., Marchewka, A., Schuss, Z.: Survival probability of diffusion with trapping in cellular neurobiology. Phys. Rev. E, Stat. Nonlinear Soft Matter Phys. 72(3), 031910 (2005)

19. Biess, A., Korkotian, E., Holcman, D.: Diffusion in a dendritic spine: the role of geometry. Phys. Rev. E, Stat. Nonlinear Soft Matter Phys. 76, 021922 (2007)

20. Holcman, D., Kupka, I.: Some questions in computational cellular biology. Fixed Point Theory Appl. 7(1), 67–83 (2010). doi:10.1007/s11784-010-0012-1

21. Berezhkovskii, A.M., Barzykin, A.V., Zitserman, V.Y.: Escape from cavity through narrow tunnel. J. Chem. Phys. 130(24), 245104 (2009)

22. Chen, L., Chetkovich, D.M., Petralia, R.S., Sweeney, N.T., Kawasaki, Y., Wenthold, R.J., Bredt, D.S., Nicoll, R.A.: Stargazin regulates synaptic targeting of AMPA receptors by two distinct mechanisms. Nature 408, 936 (2000)

23. Bredt, D.S., Nicoll, R.A.: AMPA receptor trafficking at excitatory synapses. Neuron 40, 361 (2003)

24. Adesnik, H., Nicoll, R.A., England, P.M.: Photoactivation of native AMPA receptors reveals their real-time trafficking. Neuron 48(6), 977–985 (2005)

25. Shi, S.H., Hayashi, Y., Petralia, R.S., Zaman, S.H., Wenthold, R.J., Svoboda, K., Malinow, R.: Rapid spine delivery and redistribution of AMPA receptors after synaptic NMDA receptor activation. Science 284(5421), 1811–1816 (1999)

26. Malinow, R., Malenka, R.C.: AMPA receptor trafficking and synaptic plasticity. Annu. Rev. Neurosci. 25, 103–126 (2002)

27. Holcman, D., Triller, A.: Modeling synaptic dynamics and receptor trafficking. Biophys. J. 91(7), 2405–2415 (2006)

28. Earnshaw, B.A., Bressloff, P.C.: A biophysical model of AMPA receptor trafficking and its regulation during LTP/LTD. J. Neurosci. 26, 12362–12373 (2006)

29. Ashby, M.C., Maier, S.R., Nishimune, A., Henley, J.M.: Lateral diffusion drives constitutive exchange of AMPA receptors at dendritic spines and is regulated by spine morphology. J. Neurosci. 26, 7046 (2006)

30. Hugel, S., Abegg, M., de Paola, N., Caroni, P., Gähwiler, B.H., McKinney, R.A.: Dendritic spine morphology determines membrane-associated protein exchange between dendritic shafts and spine heads. Cereb. Cortex 19(3), 697–702 (2009)

31. Holcman, D., Triller, A.: Modeling synaptic dynamics and receptor trafficking. Biophys. J. 91(7), 2405–2415 (2006)

32. Hugel, S., Abegg, M., de Paola, N., Caroni, P., Gähwiler, B.H., McKinney, R.A.: Dendritic spine morphology determines membrane-associated protein exchange between dendritic shafts and spine heads. Cereb. Cortex 19(3), 697–702 (2009)

33. Holcman, D., Triller, A.: Modeling synaptic dynamics and receptor trafficking. Biophys. J. 91(7), 2405–2415 (2006)

34. Schuss, Z., Singer, A., Holcman, D.: The narrow escape problem for diffusion in cellular microdomains. Proc. Natl. Acad. Sci. USA 104(41), 16098–16103 (2007)

35. Holcman, D., Hoze, N., Schuss, Z.: Narrow escape through a funnel and effective diffusion on a crowded membrane. Phys. Rev. E, Stat. Nonlinear Soft Matter Phys. 84, 021906 (2011)

36. Holcman, D., Schuss, Z.: Brownian motion in dire straits: Part I. Preprint (2011)

37. Schuss, Z.: Diffusion and Stochastic Processes: An Analytical Approach. Springer, New York (2010)

38. Singer, A., Schuss, Z., Holcman, D., Eisenberg, B.: Narrow escape I. J. Stat. Phys. 122(3), 437–463 (2006)

39. Singer, A., Schuss, Z., Holcman, D., Eisenberg, B.: Narrow escape II. J. Stat. Phys. 122(3), 465–489 (2006)

40. Singer, A., Schuss, Z., Holcman, D.: Narrow escape III. J. Stat. Phys. 122(3), 491–509 (2006)

41. Singer, A., Schuss, Z., Holcman, D.: Narrow escape and leakage of Brownian particles. Phys. Rev. E, Stat. Nonlinear Soft Matter Phys. 78, 051111 (2008)

42. Ward, M.J., Keller, J.B.: Strong localized perturbations of eigenvalue problems. SIAM J. Appl. Math. 53, 770–798 (1993)
44. Ward, M.J., Henshaw, W.D., Keller, J.B.: Summing logarithmic expansions for singularly perturbed eigenvalue problems. SIAM J. Appl. Math. 53, 799–828 (1993)
45. Ward, M.J., Van De Velde, E.: The onset of thermal runaway in partially insulated or cooled reactors. IMA J. Appl. Math. 48, 53–85 (1992)
46. Cheviakov, A., Ward, M.J., Straube, R.: An Asymptotic analysis of the mean first passage time for narrow escape problems: Part II: the sphere. Multiscale Model. Simul. 8(3), 836–870 (2010)
47. Holcman, D., Schuss, Z.: Escape through a small opening: receptor trafficking in a synaptic membrane. J. Stat. Phys. 117(5–6), 191–230 (2004)
48. Kolokolnikov, T., Titcombe, M., Ward, M.J.: Optimizing the fundamental Neumann eigenvalue for the Laplacian in a domain with small traps. Eur. J. Appl. Math. 16, 161–200 (2005)
49. Holcman, D., Schuss, Z.: Diffusion through a cluster of small windows and flux regulation in microdomains. Phys. Lett. A 372, 3768–3772 (2008)
50. Holcman, D., Schuss, Z.: Diffusion escape through a cluster of small absorbing windows. J. Phys. A, Math. Theor. 41, 155001 (2008)
51. Gandolfi, A., Gerardi, A., Marchetti, F.: Diffusion-controlled reactions in two dimensions. Acta Appl. Math. 4, 139–159 (1985)
52. Linderman, J., Laufengerger, D.: Analysis of intracellular receptor/Ligand sorting: calculation of mean surface and bulk diffusion times within a sphere. Biophys. J. 50, 295–305 (1986)
53. Coombs, D., Straube, R., Ward, M.: Diffusion on a sphere with localized traps: mean first passage time, eigenvalue asymptotics, and Fekete points. SIAM J. Appl. Math. 70(1), 302–332 (2009)
54. Matkowsky, B.J., Schuss, Z., Tier, C.: Uniform expansions of the transition rate in Kramers’ problem. J. Stat. Phys. 35(3–4), 443–456 (1984)
55. Taflia, A., Holcman, D.: Dwell time of a Brownian molecule in a microdomain with traps and a small hole on the boundary. J. Chem. Phys. 126(23), 234107 (2007)
56. Bressloff, P.C., Earnshaw, B.A.: A dynamical corral model of protein trafficking in spines. Biophys. J. 96, 1786–1802 (2009)
57. Bressloff, P.C.: Cable theory of protein receptor trafficking in dendritic trees. Phys. Rev. E, Stat. Nonlinear Soft Matter Phys. 79, 041904 (2009)
58. Majewska, A., Brown, E., Ross, J., Yuste, R.: Mechanisms of calcium decay kinetics in hippocampal spines: role of spine calcium pumps and calcium diffusion through the spine neck in biochemical compartmentalization. J. Neurosci. 20(5), 1722–1734 (2000)
59. Majewska, A., Tashiro, A., Yuste, R.: Regulation of spine calcium dynamics by rapid spine motility. J. Neurosci. 20(22), 8262–8268 (2000)
60. Yuste, R., Majewska, A., Holthoff, K.: From form to function: calcium compartmentalization in dendritic spines. Nat. Neurosci. 7, 653–659 (2000)