

Statistical physics of biological motion: Crawling cells and foraging bumblebees

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Outline

two parts:

- 1 **cell migration**
- 2 **bumblebee foraging**

in both cases:

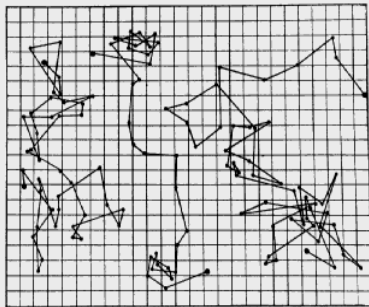
- **motivation and experiment**
- **experimental results and statistical analysis**
- **theoretical modeling and summary**

Part 1:

Cell Migration

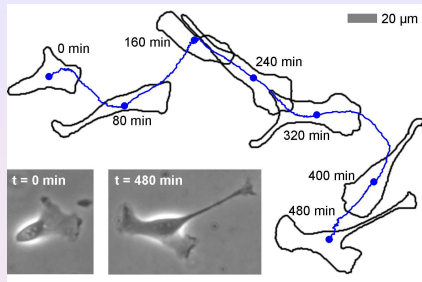
Brownian motion of migrating cells?

Brownian motion



Perrin (1913)

three colloidal particles,
positions joined by straight
lines



Dieterich et al. (2008)

single biological cell crawling on
a substrate

Brownian motion?

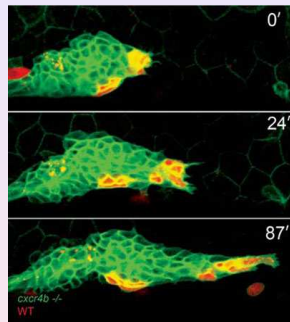
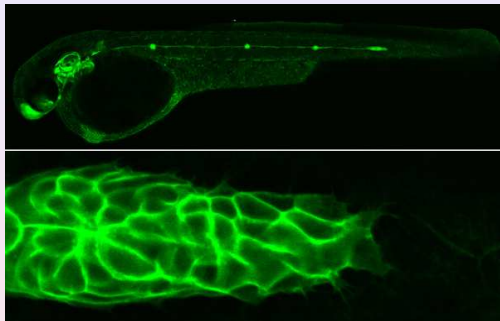
conflicting results:

yes: Dunn, Brown (1987)

no: Hartmann et al. (1994)

Why cell migration?

motion of the *primordium* in developing zebrafish:



Gilmour (2008)

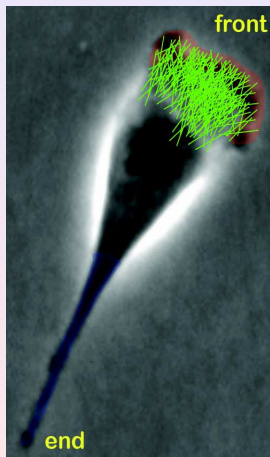
positive aspects:

- morphogenesis
- immune defense

negative aspects:

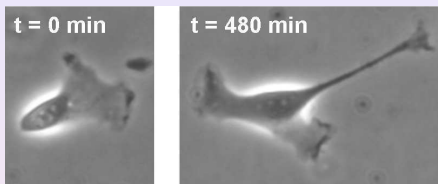
- tumor metastases
- inflammation reactions

How do cells migrate?



- **membrane protrusions and retractions** ~ force generation:
 - lamellipodia (front)
 - uropod (end)
 - actin-myosin network
- formation of a **polarized state**
front/end
- cell-substrate **adhesion**

Our cell types and some typical scales



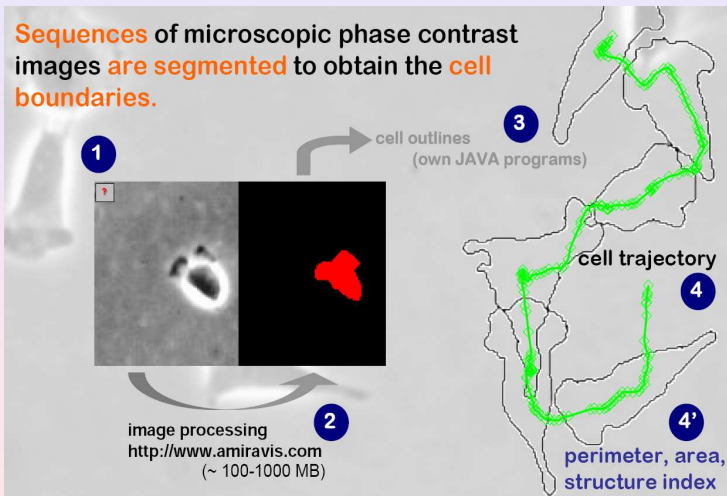
- **renal epithelial MDCK-F (Madin-Darby canine kidney) cells**; two types: wildtype (NHE^+) and NHE -deficient (NHE^-)
- observed up to **1000 minutes**: here *no* limit $t \rightarrow \infty$!
- cell diameter **$20-50\mu\text{m}$** ; mean velocity $\sim 1\mu\text{m}/\text{min}$; lamellipodial dynamics \sim **seconds**

movies: NHE+: t=210min, dt=3min

NHE-: t=171min, dt=1min

Measuring cell migration

Sequences of microscopic phase contrast images **are segmented** to obtain the **cell boundaries**.



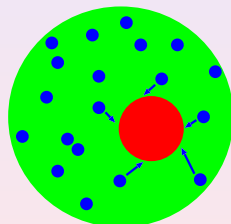
Theoretical modeling of Brownian motion

‘Newton’s law of stochastic physics’:

$$\dot{\mathbf{v}} = -\kappa\mathbf{v} + \sqrt{\zeta} \boldsymbol{\xi}(t) \quad \text{Langevin equation (1908)}$$

for a **tracer particle of velocity \mathbf{v}** immersed in a fluid

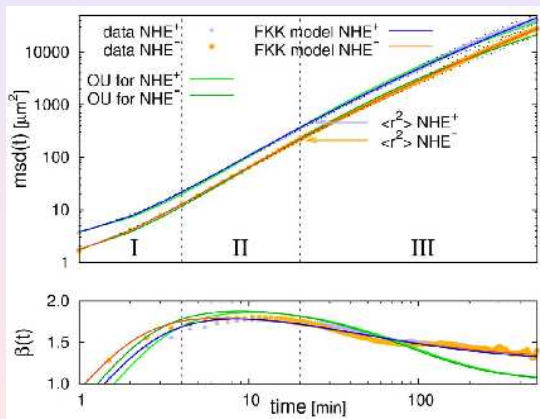
force decomposed into **viscous damping** and **random kicks of surrounding particles**



Application to cell migration?

Mean square displacement

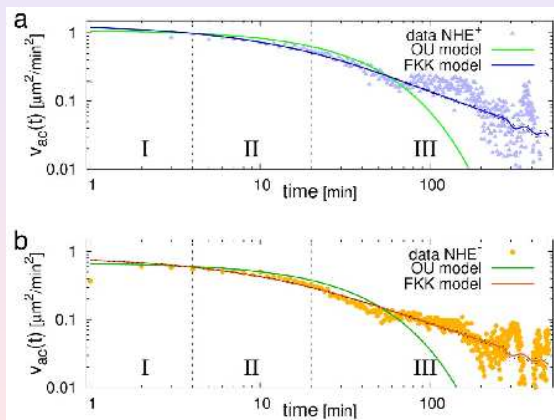
- $msd(t) := \langle [\mathbf{x}(t) - \mathbf{x}(0)]^2 \rangle \sim t^\beta$ with $\beta \rightarrow 2$ ($t \rightarrow 0$) and $\beta \rightarrow 1$ ($t \rightarrow \infty$) for Brownian motion; $\beta(t) = d \ln msd(t) / d \ln t$



anomalous diffusion if $\beta \neq 1$ ($t \rightarrow \infty$); here: **superdiffusion**

Velocity autocorrelation function

- $v_{ac}(t) := \langle \mathbf{v}(t) \cdot \mathbf{v}(0) \rangle \sim \exp(-\kappa t)$ for Brownian motion
- fits with same parameter values as $msd(t)$



crossover from **stretched exponential to power law**

Position distribution function

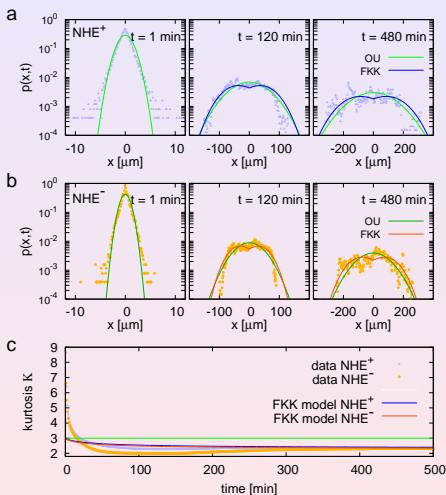
- $P(x, t) \rightarrow$ Gaussian ($t \rightarrow \infty$) and kurtosis

$$\kappa(t) := \frac{\langle x^4(t) \rangle}{\langle x^2(t) \rangle^2} \rightarrow 3 \quad (t \rightarrow \infty)$$

for Brownian motion (green lines, in 1d)

- *other solid lines*: fits from our model; parameter values as before

note: model needs to be amended to explain short-time distributions



crossover from peaked to broad **non-Gaussian distributions**

The model

Fractional Klein-Kramers equation (Barkai, Silbey, 2000):

$$\frac{\partial P}{\partial t} = -\frac{\partial}{\partial x} [vP] + \frac{\partial^{1-\alpha}}{\partial t^{1-\alpha}} \kappa \left[\frac{\partial}{\partial v} v + v_{th}^2 \frac{\partial^2}{\partial v^2} \right] P$$

with probability distribution $P = P(x, v, t)$, damping term κ , thermal velocity v_{th} and **Riemann-Liouville fractional derivative of order $1 - \alpha$** defined by

$$\frac{\partial^\gamma P}{\partial t^\gamma} := \begin{cases} \frac{\partial^m P}{\partial t^m} & , \quad \gamma = m \\ \frac{\partial^m}{\partial t^m} \left[\frac{1}{\Gamma(m-\gamma)} \int_0^t dt' \frac{P(t')}{(t-t')^{\gamma+1-m}} \right] & , \quad m-1 < \gamma < m \end{cases}$$

with $m \in \mathbb{N}$; for $\alpha = 1$ ordinary Klein-Kramers equation recovered

4 fit parameters v_{th}, α, κ (plus another one for ‘biological noise’ on short time scales)

Solutions for this model

analytical solutions (Barkai, Silbey, 2000):

- **mean square displacement:**

$$msd(t) = 2v_{th}^2 t^2 E_{\alpha,3}(-\kappa t^\alpha) \rightarrow 2 \frac{D_\alpha t^{2-\alpha}}{\Gamma(3-\alpha)} \quad (t \rightarrow \infty)$$

with $D_\alpha = v_{th}^2 / \kappa$ and *generalized Mittag-Leffler function*

$$E_{\alpha,\beta}(z) = \sum_{k=0}^{\infty} \frac{z^k}{\Gamma(\alpha k + \beta)}, \quad \alpha, \beta > 0, \quad z \in \mathbb{C};$$

note that $E_{1,1}(z) = \exp(z)$: $E_{\alpha,\beta}(z)$ is a generalized exponential function

- **velocity autocorrelation function:**

$$v_{ac}(t) = v_{th}^2 E_{\alpha,1}(-\kappa t^\alpha) \rightarrow \frac{1}{\kappa \Gamma(1-\alpha) t^\alpha} \quad (t \rightarrow \infty)$$

- for $\kappa \rightarrow \infty$ fractional Klein-Kramers reduces to a *fractional diffusion equation* yielding $P(x, t)$ in terms of a Fox function (Schneider, Wyss, 1989)

Possible physical interpretation

Physical meaning of the fractional derivative?

fractional Klein-Kramers equation is *approximately* related to the generalized Langevin equation

$$\dot{v} + \int_0^t dt' \kappa(t-t')v(t') = \sqrt{\zeta} \xi(t)$$

e.g., Mori, Kubo (1965/66)

with **time-dependent friction coefficient** $\kappa(t) \sim t^{-\alpha}$

cell anomalies might originate from **glassy behavior** of the cytoskeleton gel, where power law exponents are conjectured to be universal (Fabry et al., 2003; Kroy et al., 2008)

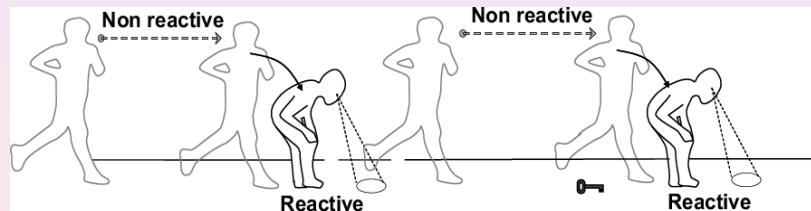
nb: anomalous dynamics observed for *many different cell types*

Possible biological interpretation

Biological meaning of the anomalous cell migration?

experimental data and theoretical modeling suggest *slower diffusion for small times* while *long-time motion is faster*

compare with **intermittent optimal search strategies** of foraging animals (Bénichou et al., 2006)



note: controversy about **modeling the migration of foraging animals** (albatros, **bumblebees**, fruitflies,...)

Summary: Anomalous cells

- different **cell dynamics** on different **time scales**
(cp. with **Lévy hypothesis**, which suggests scale-freeness)
- for long times cells crawl **superdiffusively** with **power law decay of velocity correlations and non-Gaussian position pdfs**
- **stochastic modeling** of experimental data by a **generalized Klein-Kramers equation**

Part 2:

Bumblebee Foraging

Motivation

bumblebee foraging – two very practical problems:

1. find food (nectar, pollen) in complex landscapes



2. try to avoid predators

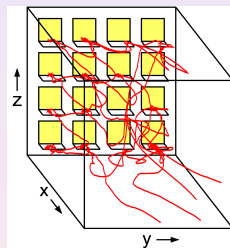
What type of motion?

Study bumblebee foraging in a *laboratory experiment*.

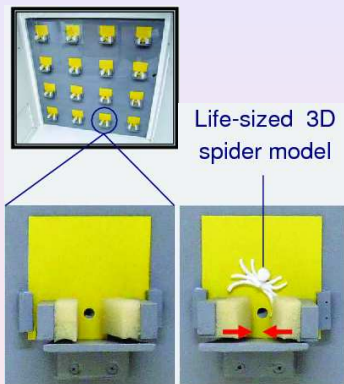
The bumblebee experiment

Ings, Chittka, *Current Biology* **18**, 1520 (2008):
bumblebee foraging in a cube of $\simeq 75\text{cm}$ side length

- artificial yellow flowers: 4x4 grid on one wall
- two cameras track the position (50fps) of a single bumblebee (*Bombus terrestris*)
- **advantages:** systematic **variation of the environment**;
easier than tracking bumblebees on large scales
- **disadvantage:** no 'free flight' of bumblebees



Variation of the environmental conditions



Life-sized 3D spider model

movie

three experimental stages:

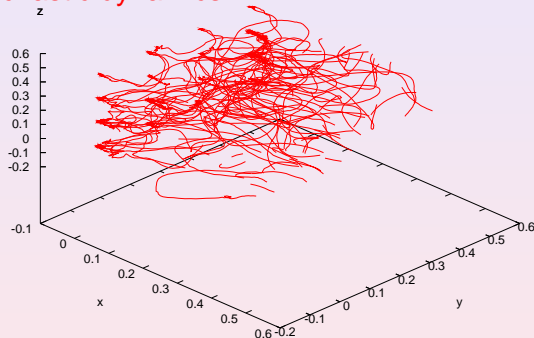
- 1 spider-free foraging
- 2 foraging under predation risk
- 3 memory test 1 day later

safe and **dangerous**
flowers

#bumblebees=30 , #data per bumblebee for each stage \approx 7000

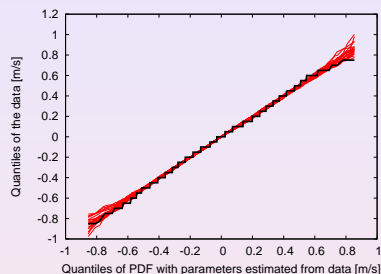
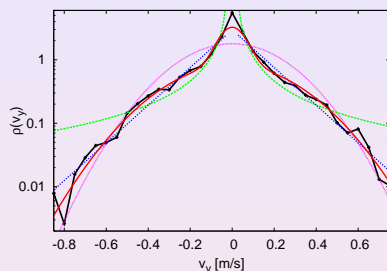
Bumblebee experiment: two main questions

- 1 What **type of motion** do the bumblebees perform in terms of **stochastic dynamics**?



- 2 Are there **changes of the dynamics** under **variation of the environmental conditions**?

Velocity distributions: analysis



left: experimental **pdf of v_y -velocities** of a single bumblebee in the spider-free stage (black crosses) with max. likelihood fits of **mixture of 2 Gaussians**; **exponential**; **power law**; **single Gaussian**

right: **quantile-quantile plot** of a Gaussian mixture against the experimental data (black) plus **surrogate data**

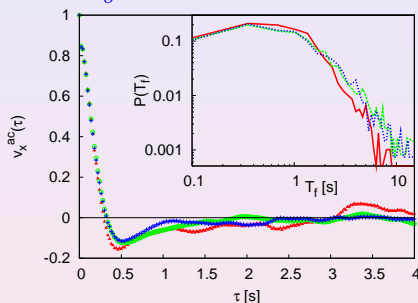
Velocity distributions: interpretation

- **best fit** to the data by a **mixture of two Gaussians** with different variances (quantified by information criteria with resp. weights)
- **biological explanation:** models **spatially different flight modes** near the flower vs. far away, cf. intermittent dynamics

big surprise: no difference in pdf's between different stages under variation of environmental conditions!

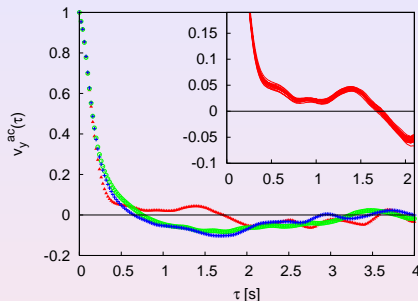
Velocity autocorrelation function \perp to the wall

$$V_x^{AC}(\tau) = \frac{\langle (v_x(t) - \mu)(v_x(t + \tau) - \mu) \rangle}{\sigma^2} \quad \text{with average over all bees}$$



- plot: spider-free stage, predation thread, memory test
- \exists **anti-correlations** for $\tau \simeq 0.5$: bees return to flowers
- only small **quantitative changes** under predation thread, cf. shift of minimum in $V_x^{AC}(\tau)$ and changes in pdf of flight times (inset): more flights with long durations

Velocity autocorrelation function \parallel to the wall

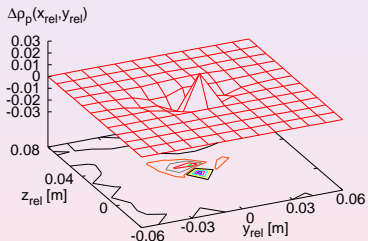


- plot: spider-free stage, predation thread, memory test
- \exists **profound qualitative change** of correlations from positive for spider-free to negative in case of spiders
- resampling of data (inset) confirms existence of positive correlations

\Rightarrow all **changes** are in the **velocity correlations**, *not* the pdf's!

Predator avoidance and a simple model

predator avoidance as
difference in position pdfs
spider / no spider from data:



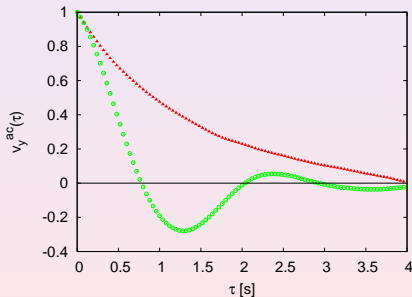
positive spike: *hovering*;
negative region: *avoidance*

modeled by Langevin equation

$$\frac{dv_y}{dt}(t) = -\eta v_y(t) - \frac{\partial U}{\partial y}(y(t)) + \xi(t)$$

η : friction coefficient,

ξ : Gaussian white noise



simulated velocity correlations with
repulsive interaction potential U
bumblebee - spider **off** / **on**

Summary: Clever bumblebees

- mixture of **two Gaussian velocity distributions** reflects **spatial adjustment** of bumblebee dynamics to flower carpet
- all changes to predation threat are contained in the **velocity autocorrelation functions**, which exhibit highly **non-trivial temporal behaviour**

(nb: **Lévy hypothesis** suggests that all relevant foraging information is contained in pdf's)
- **change of correlation decay** in the presence of spiders due to **experimentally extracted repulsive force** as reproduced by generalized Langevin dynamics

Collaborators and literature

work performed with:

1. cells: P.Dieterich, R.K., R.Preuss, A.Schwab,
Anomalous Dynamics of Cell Migration, PNAS **105**, 459 (2008)

2. bees: F.Lenz, T.Ings, A.V.Chechkin, L.Chittka, R.K.,
*Spatio-temporal dynamics of bumblebees foraging under
predation risk*, Phys. Rev. Lett. **108**, 098103 (2012)

