Effects of tubular inhomogeneities on the tubuloglomerular feedback (TGF) system
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Introduction
An inhomogeneous model (IM) of rat thick ascending limb (TAL) is used to predict dynamic behaviors of the tubuloglomerular feedback (TGF) loop and to explain the experimental data from hypertensive rats [5], which have shown that TGF system may exhibit irregular oscillations with characteristics of deterministic chaos (Fig. 1, B and C).

Results
To understand the effect of inhomogeneities on the model behavior we incrementally modified the homogeneous base model, and considered two spatially homogeneous models (IM zero backleak and HM non-zero backleak) and one spatially inhomogeneous model (IM), built on experimental observations [1,2].

Bifurcation diagrams
Figure 2. Schematic of model components.
Figure 3. Bifurcation diagrams based on characteristic equations for linearized model equations (right column). These diagrams are regions of the parameter plane that predict the steady-state and oscillatory behaviors based on the real parts of the eigenvalues λ1, λ2, and λ3. Confirmation of solution behaviors depends on numerical simulations using the full model equations (left column). Points A, B, and C correspond to panels A, B, and C of Fig. 4; their parameters are (0.135, 8). More model simulations for the IM are shown in Fig. 5, corresponding to points BH(0.1, 10) and D2(0.075, 10).

In the Steady State region, the only stable solution is the time-independent steady state. Above the curve Re λ1 > 0, 1 stable oscillations are possible; above the curve Re λ2 > 0, 2 stable oscillations are possible, i.e., the stable oscillations have a frequency about twice that found in the previous region 1f LCO, especially above the Re λ2 > Re λ1, where Re λ2 > Re λ1. The region labeled 2f-LCO exhibits bistability with respect to model solutions, i.e., two stable LCOs, of differing frequencies, can be elicited, depending on initial conditions.

Effects of backleak and inhomogeneities
Effects of backleak and inhomogeneities on the base case model are illustrated for the points marked in Fig. 3.

First, points A, B, and C with the same pair of parameters are considered (Fig. 4). As expected from Fig. 3, point A corresponds to a periodic, non-sinusoidal oscillation at the fundamental frequency, while points B and C corresponds to oscillations twice the fundamental frequency.

More irregular are the spectra for other selected points, D1 and D2, in the parameter plane of the IM (Fig. 5).

Methods
Figure 2. Schematic of model components.

Figure 4. Right panels: Oscillations in chloride concentration at MD and in SNGFR computed from the mathematical model at the chosen points of Fig. 3. All resulting concentration waveforms are shifted upwards, the average values in each case being higher than the steady-state concentration values approximated by 32.0 mM. Left panels: Corresponding power spectra of tubular fluid chloride concentration at the MD (solid lines) and of SNGFR (gray shades), with both linear and logarithmic vertical axes. The general trend of decreasing amplitude as a function of frequency indicates that the TAL operates as a low-pass filter. It also reveals the nodal structure of the chloride concentration arising at the MD. The first nodal frequency for the MD concentration appears indeed at the primary fundamental frequency, while all the rest nodal frequencies are multiples of the first one.

Figure 5. Numerical simulations for chloride concentration arising at the MD, corresponding to points D1 (0.1, 10) and D2 (0.075, 10). For Fig. 3, for IM (solid lines) and HM non-zero backleak (gray shades).

Summary
1. We used an inhomogeneous model (IM) of rat TAL to investigate the effects of non-constant TAL radius and non-constant chloride transport rate on tubular concentration undergoing a transient oscillatory perturbation, and on TGF-mediated oscillations.
2. We show that the physiologic parameter space has regions that correspond to four qualitatively different model behaviors:
   • a time-independent steady state,
   • a sustained oscillation with frequency f,
   • a sustained oscillation of frequency ~2f, and
   • two sustained oscillations, differing frequencies, that depend on initial conditions.
3. The waveforms and power spectra arising in our model are similar to complex waveforms and corresponding power spectra from hypertensive rats. Thus we hypothesize that irregular oscillations in hypertensive rats are attributable, at least in part, to the TAL inhomogeneities.
4. Power spectra from hypertensive rats contain other frequency signature that remain to be explained.

References

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