

# Mutual inhibition and bistability: two key elements of the endochondral ossification process



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## INTRODUCTION

### Background

- *in vitro* engineering of tissues : may be achieved by mimicking *in vivo* tissue development
- Endochondral ossification : chondrocytes pass through a succession of states (proliferative and hypertrophic state)

### Sox9-Runx2 switch and bistability

- Sox9 program is characteristic of the **proliferative state** and Runx2 program of the **hypertrophic state**
- Runx2 and Sox9 inhibit each other
- $\beta$ -catenin, downstream of Wnt and BMP, is a key factor in this mechanism

### Aim of this study

- model the switch between Sox9 (proliferative) program and Runx2 (hypertrophic) program
- experimental validation

## MATERIALS & METHODS

### MODEL 1

- a literature-based mathematical model describing BMP and Wnt pathways and various cross-talks [1]
- **mutual inhibition** between BMP and Wnt
- regulation of the amount of  $\beta$ -catenin in the nucleus
- **parameter values** derived from previous models [1] and experiments reported in literature [3]
- 19 variables, 49 parameters

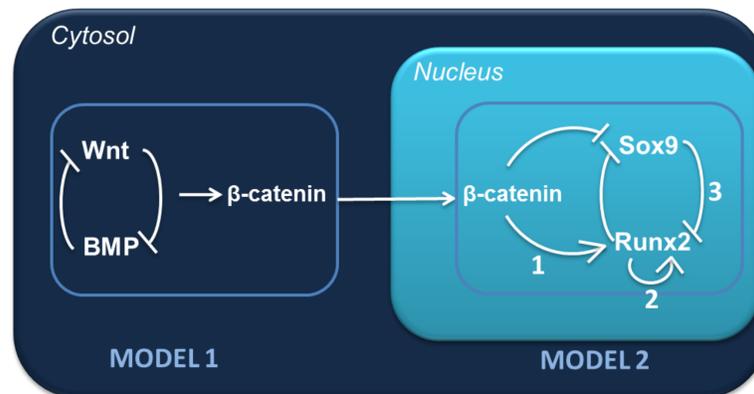


Figure 1 : Schematic representation of chondrocyte with indication of both models

### MODEL 2

- a literature-based mathematical model describing the switch between Sox9 and Runx2
- **mutual inhibition** between Sox9 and Runx2
- **auto-activation** of Runx2
- **parameter values** by screening of parameter space to find parameter sets generating **bistable behavior** [2]
- 3 variables, 16 parameters

**Ordinary Differential Equations (ODEs)** describe the temporal evolution of the various model constituents (numbers refer to interactions in Figure 1)

$$\frac{d[Runx2]}{dt} = \left( \frac{[\beta_{cat}]^{n1}}{K_1^{n1} + [\beta_{cat}]^{n1}} \right) \xrightarrow{1} \left( \alpha_{Runx2} \frac{[Runx2]^{n2}}{K_2^{n2} + [Runx2]^{n2}} \right) \xrightarrow{2} \left( \frac{K_3^{n3}}{K_3^{n3} + [Sox9]^{n3}} \right) \xrightarrow{3} [Runx2] \xrightarrow{\text{degradation}}$$

## RESULTS

- the model predicts that :
  - ✓ activation of Wnt upregulates  $\beta$ -catenin and provokes the switch between the Sox9 state and the Runx2 state
  - ✓ activation of BMP inhibits the transition of  $\beta$ -catenin to the nucleus but cannot provoke a switch from the Runx2 state towards the Sox9 state
  - ✓ two stable states (bistability) are obtained for appropriate parameter sets

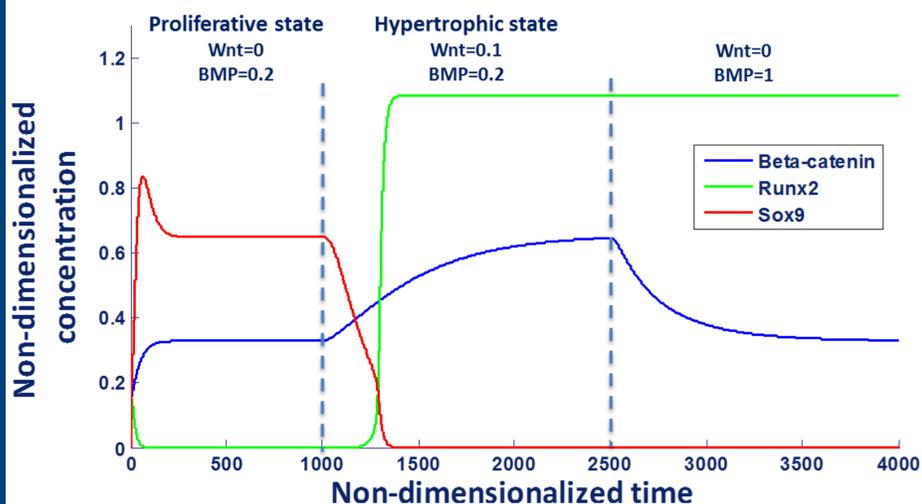


Figure 2 : Influence of Wnt and BMP on  $\beta$ -catenin, Sox9 and Runx2

## DISCUSSION

- in absence of quantitative parameter information, the ODE model presented here provides qualitative predictions on changes in the concentrations of all modelled components
- the model is able to reproduce the switch between the Sox9 program and the Runx2 program for specific parameter sets
- the model behavior is in concordance with experimental results present in the literature [4]
- mathematical models can be used to enhance our understanding of signaling cascades and their interactions

## REFERENCES

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