Algebraic Systems Biology: A Case Study for the Wnt Pathway

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Abstract
Steady-state analysis of dynamical systems for biological networks gives rise to algebraic varieties in high-dimensional spaces whose study is of interest in their own right. We demonstrate this for the shuttle model of the Wnt signaling pathway. Here, the variety is described by a polynomial system in 19 unknowns and 36 parameters. It has degree 9 over the parameter space. This case study explores multistationarity, model comparison, dynamics within regions of the state space, identifiability, and parameter estimation, from a geometric point of view. We employ current methods from computational algebraic geometry, polyhedral geometry, and combinatorics.

Keywords
Biochemical reaction networks · Nonlinear algebra · β-catenin/Wnt signaling · Steady-state variety · Polyhedra · Algebraic matroids

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1 Introduction

The theory of biochemical reaction networks is fundamental for systems biology (Klipp et al. 2009; Voit 2012). It is based on a wide range of mathematical fields, including dynamical systems, numerical analysis, optimization, combinatorics, probability, and, last but not least, algebraic geometry. There are numerous articles that use algebraic geometry in the study of biochemical reaction networks, especially those arising from mass action kinetics. A tiny selection is Craciun and Feinberg (2005), Feliu and Wiuf (2012), Karp et al. (2012), Pérez Millán et al. (2012), and Shiu and Sturmfels (2010).

We here perform a detailed analysis of one specific system, namely the shuttle model for the Wnt signaling pathway, introduced recently by MacLean et al. (2015). Our aim is twofold: to demonstrate how biology can lead to interesting questions in algebraic geometry and to apply state-of-the-art techniques from computational algebra to biology.

The dynamical system we study consists of the following 19 ordinary differential equations. Their derivation and the relevant background from biology will be presented in Sect. 2.

\[
\begin{align*}
\dot{x}_1 &= -k_1 x_1 + k_2 x_2 \\
\dot{x}_2 &= k_1 x_1 - (k_2 + k_{26}) x_2 + k_{27} x_3 - k_3 x_2 x_4 + (k_4 + k_5) x_{14} \\
\dot{x}_3 &= k_{26} x_2 - k_{27} x_3 - k_{14} x_3 x_6 + (k_{15} + k_{16}) x_{15} \\
\dot{x}_4 &= -k_3 x_2 x_4 - k_9 x_4 x_{10} + k_4 x_{14} + k_8 x_{16} + (k_{10} + k_{11}) x_{18} \\
\dot{x}_5 &= -k_{28} x_5 + k_{29} x_7 - k_6 x_5 x_8 + k_5 x_{14} + k_7 x_{16} \\
\dot{x}_6 &= -k_{14} x_3 x_6 - k_{20} x_6 x_{11} + k_{15} x_{15} + k_{19} x_{17} + (k_{21} + k_{22}) x_{19} \\
\dot{x}_7 &= k_{28} x_5 - k_{29} x_7 - k_{17} x_7 x_9 + k_{16} x_{15} + k_{18} x_{17} \\
\dot{x}_8 &= -\dot{x}_{16} = -k_6 x_5 x_8 + (k_7 + k_8) x_{16} \\
\dot{x}_9 &= -\dot{x}_{17} = -k_{17} x_7 x_9 + (k_{18} + k_{19}) x_{17} \\
\dot{x}_{10} &= k_{12} - (k_{13} + k_{30}) x_{10} - k_9 x_4 x_{10} + k_{31} x_{11} + k_{10} x_{18} \\
\dot{x}_{11} &= -k_{23} x_{11} + k_{30} x_{10} - k_3 x_{11} - k_{20} x_6 x_{11} - k_{24} x_{11} x_{12} + k_{25} x_{13} + k_{21} x_{19} \\
\dot{x}_{12} &= -\dot{x}_{13} = -k_{24} x_{11} x_{12} + k_{25} x_{13} \\
\dot{x}_{14} &= k_3 x_2 x_4 - (k_4 + k_5) x_{14} \\
\dot{x}_{15} &= k_{14} x_3 x_6 - (k_{15} + k_{16}) x_{15} \\
\dot{x}_{16} &= k_9 x_4 x_{10} - (k_{10} + k_{11}) x_{18} \\
\dot{x}_{19} &= k_{20} x_6 x_{11} - (k_{21} + k_{22}) x_{19}
\end{align*}
\]

The quantity \( x_i \) is a differentiable function of an unknown \( t \), representing time, and \( \dot{x}_i(t) \) is the derivative of that function. This dynamical system has five linear conservation laws:

\[
\begin{align*}
0 &= (x_1 + x_2 + x_3 + x_{14} + x_{15}) - c_1 \\
0 &= (x_4 + x_5 + x_6 + x_7 + x_{14} + x_{15} + x_{16} + x_{17} + x_{18} + x_{19}) - c_2 \\
0 &= (x_8 + x_{16}) - c_3 \\
0 &= (x_9 + x_{17}) - c_4 \\
0 &= (x_{12} + x_{13}) - c_5
\end{align*}
\]
The 31 quantities $k_i$ are the rate constants of the chemical reactions, and the five $c_i$ are the conserved quantities. Both of these are regarded as parameters, so we have 36 parameters in total. Our object of interest is the steady-state variety, which is the common zero set of the right-hand sides of (1) and (2), and hence contains all steady states of the system (real and complex). This variety lives in $\mathcal{K}_1^{19}$, where $\mathcal{K}$ is an algebraically closed field that contains the rational numbers $\mathbb{Q}$ as well as the 36 parameters $k_i$ and $c_i$. If these parameters are fixed to be particular real numbers, then we can take $\mathcal{K} = \mathbb{C}$, the field of complex numbers. If it is preferable to regard $k = (k_1, \ldots, k_{31})$ and $c = (c_1, \ldots, c_5)$ as vectors of unknowns, then $\mathcal{K} = \mathbb{Q}(k, c)$ is the algebraic closure of the rational function field. In this latter setting, when all parameters are generic, we shall derive the following result:

**Theorem 1.1** The polynomials in (1)–(2) have nine distinct zeros in $\mathcal{K}_{19}$ when $\mathcal{K} = \mathbb{Q}(k, c)$.

By analyzing the steady-state variety, we can better understand the model and thus the biological system. Note that the model is nonlinear. Depending on the application, sometimes it is preferable to work over the complex numbers $\mathbb{C}$, while other times we restrict our analysis to a real positive orthant. For example, when using tools from algebraic geometry, we work over an algebraically closed field, e.g., $\mathbb{C}$ or $\mathcal{K} = \mathbb{Q}(k, c)$. Regardless of the choice of field, the aim is to predict the system’s behavior, offer biological insight, and determine what data are required to verify or reject the model. Here is a list of questions one might ask about our model from the perspective of systems biology.

1.1 Biological Problems

These are numbered according to the section that addresses them.

4. *For what real positive rate parameters and conserved quantities does the system exhibit multistationarity?* This question is commonly asked when using a dynamical system for modeling a real-world phenomenon. When modeling a process that experimentally appears to have more than one stable equilibrium, multistationary models are preferred.

5. *Suppose we can measure only a subset of the species concentrations. Which subsets can lead to model rejection?* If all species are measurable at steady state, then we can substitute data into the system (1) and check that all expressions $\dot{x}_i$ are close to zero. If only some $x_i$ are known, we still want to be able to evaluate models with the available data.

6. *Give a complete description of the stoichiometric compatibility classes for the chemical reaction network.* A stoichiometric compatibility class is the set of all points accessible from a given state via the reactions in the system. This question relates more closely to the dynamics of the system, but also has ramifications for the set of all steady states.

7. *What information does species concentration data give us for parameter estimation?* In particular, are the parameters identifiable? Identifiability means that having many measurements of the concentrations $x$ can determine the reaction rate
constants \( k \). If not identifiable, we will explore algebraic constraints imposed by the species concentration data. This question is relevant for complete and partial steady-state data (usually noisy).

These questions are open challenges for medium to large models in systems biology and medicine (Klipp et al. 2009; Voit 2012). The book chapter (MacLean et al. 2015) illustrates standard mathematical and statistical methods for addressing these questions for small-to-medium Wnt signaling models. Here, we examine these questions for a medium-to-large Wnt model from the perspective of algebraic geometry. The aim is to provide insight into global behavior by applying tools from nonlinear algebra to systems biology. Below are the algebraic problems underlying the four biological problems listed above.

### 1.2 Algebraic Problems

These are numbered to match the corresponding biological problems, so the number refers to the section that addresses the problem.

1. Describe the set of points \((k, c) \in \mathbb{R}_{>0}^{31} \times \mathbb{R}_{>0}^5\) such that the polynomials (1)–(2) have two or more positive zeros \(x \in \mathbb{R}_{>0}^{19}\). When is there only one? Identify the discriminant.

2. Which projections of the variety defined by (1) into coordinate subspaces of \(K^{19}\) are surjective? Equivalently, describe the algebraic matroid on the ground set \(\{x_1, \ldots, x_{19}\}\).

3. The conservation relations (2) specify a linear map \(\chi : \mathbb{R}^{19} \to \mathbb{R}^5\), \(x \mapsto c\). Describe all the convex polyhedra \(\chi^{-1}(c) \cap \mathbb{R}_{>0}^{19}\) where \(c\) runs over the points in the open orthant \(\mathbb{R}_{>0}^5\). Specifically, what is the full set of feasible \(x\)-values (not just at steady state) given a fixed value for \(c\)?

4. **Complete data:** Describe the matroid on the ground set \(\{k_1, k_2, \ldots, k_{31}\}\) that is defined by the linear forms on the right-hand sides of (1), for fixed steady-state concentrations. Note that these linear forms live in \(\mathbb{Q}(x)[k]\).

5. **Partial steady-state data without noise:** Repeat the analysis after eliminating some of the \(x\)-coordinates.

6. **Partial steady-state data with noise:** For the remaining \(x\)-coordinates, suppose that we have data which are approximately on the projected steady-state variety. Determine a parameter vector \((k, c)\) that best fits the data.

In this paper, we shall address these questions and several related ones, after explaining the various ingredients. A particular focus is the exchange between the algebraic formulation and its biological counterpart. Our presentation is organized as follows.

In Sect. 2, we review the basics on the Wnt signaling pathway, we recall the shuttle model of MacLean et al. (2015), and we derive the dynamical system (1)–(2). In Sect. 3, we establish Theorem 1.1, and we examine the set of all steady states. This is here regarded as a complex algebraic variety in an affine space of dimension \(55 = 19 + 31 + 5\) with coordinates \((x, k, c)\).
In Sects. 4, 5, 6, and 7, we address the four problems stated above. The numbers of the problems refer to the respective sections. Each section starts out with an explanation of how the biological problem and the algebraic problem are related. The rationale behind Sect. 4 is likely to be familiar to most of our readers, given that multistationarity has been discussed widely in the literature; see, e.g., Craciun and Feinberg (2005) and Pérez Millán et al. (2012). On the other hand, in Sect. 5, we employ the language of matroid theory. This may be unfamiliar to many readers, especially when it comes to the algebraic matroid associated with an irreducible algebraic variety. Section 6 characterizes the polyhedral geometry encoded in the conservation relations (2). This is a case study in the spirit of (Shiu and Sturmfels 2010, Figure 1). Section 7 addresses the problems of parameter identifiability and parameter estimation. Finally, in Sect. 8, we return to the biology, and we discuss what our findings might imply for the study of Wnt signaling and other systems.

2 From Biology to Algebra

Cellular decisions such as cell division, specialization, and cell death are governed by a rich repertoire of complex signals that are produced by other cells and/or stimuli. In order for a cell to come to an appropriate decision, it must sense its external environment, communicate this information to the nucleus, and respond by regulating genes and producing relevant proteins. Signaling molecules called ligands, external to the cell, can bind to proteins called receptors, initializing the propagation of information within the cell by molecular interactions and modifications (e.g., phosphorylation). This signal may be relayed from the cytoplasm into the nucleus via molecules, and the cell responds by activation or deactivation of gene(s) that control, for example, cell fate. The complex interplay of molecules involved in this information transmission is called a signaling transduction pathway. Although many signaling pathways have been defined biochemically, much is still not understood about them or how a signal results in a particular cellular response. Mathematical models constructed at different scales of molecular complexity may help unravel the central mechanisms that govern cellular decisions, and their analysis may inform and guide testable hypotheses and therapies.

In this paper, we focus on the canonical Wnt signaling pathway, which is involved in cellular processes, both during development and in adult tissues. This includes stem cells. Dysfunction of this pathway has been linked to neurodegenerative diseases and cancer. Consequently, Wnt signaling has been widely studied in various organisms, including amphibians and mammals. Researchers are interested in how the extracellular ligand Wnt affects the protein β-catenin, which plays a pivotal role in turning genes on and off in the nucleus.

The molecular interactions within the Wnt signaling pathway are not yet fully understood. This has led to the development and analysis of many mathematical models. The Wnt shuttle model (MacLean et al. 2015) includes an abstraction of the signal transduction pathway (via activation/inactivation of molecules) described above. The model also takes into account molecules that exist, interact, and move between different compartments in the cell (e.g., cytoplasm and nucleus). Biologists understand the
Wnt system as either Wnt off or Wnt on. However, such a scenario is rarely binary (i.e., different concentration levels of Wnt may exist) and inherently depends on spatial movement of molecules. The Wnt shuttle model includes complex interactions with nonlinearities arising in the equations. In particular, it includes both the Wnt off and Wnt on scenarios, by adjusting initial conditions or parameter values. The biology needed to understand the model can be described as follows. See also Table 1.

Wnt off: When cells do not sense the extracellular ligand Wnt, β-catenin is degraded (broken down). The degradation of β-catenin is partially dependent on a group of molecules (Axin, APC, and GSK-3) that form the destruction complex. Crucially, the break down of β-catenin occurs when the destruction complex is in an active state; modification to the destruction complex by proteins, called phosphatases, changes it from inactive to active. Additionally, β-catenin can degrade independent of the destruction complex. Synthesis of β-catenin occurs at a constant rate.

Wnt on: When receptors on the surface of a cell bind to Wnt, the Wnt signaling transduction pathway is initiated. This enables β-catenin to move into the nucleus where it binds with transcription factors that regulate genes. This signal propagation is mediated by the following molecular interactions. After Wnt stimulus, the protein Dishevelled is activated near the membrane. This in turn inactivates the destruction complex, thereby preventing the destruction of β-catenin, allowing it to accumulate in the cytoplasm through natural synthesis. Throughout the molecular interactions in the signaling pathway, intermediate complexes can form (e.g., β-catenin bound with Dishevelled).

Space: The location of molecules plays a pivotal role: β-catenin moves between the cytoplasm and the nucleus (to reach target genes and regulate them). Dishevelled and molecules that form the destruction complex shuttle between the nucleus and the cytoplasm. However, it is assumed that only the inactive destruction complex can shuttle (since in the cytoplasm it would be bound to β-catenin). Phosphatases exist in both the nucleus and the cytoplasm, but the movement across compartments is not included in the model. Symmetry of reactions is assumed if the species exist in both compartments. Intermediate complexes are assumed to be short-lived, or not small enough for movement across compartments.

The Wnt shuttle model of MacLean et al. (2015) has 19 species whose interactions can be framed as biochemical reactions. These species correspond to variables $x_1, \ldots, x_{19}$ in our dynamical system (1). Namely, $x_i$ represents the concentration of the species that is listed in the $i$th row in Table 1. The second column in Table 1 indicates the biological meaning of the 19 species. The symbols in the last column are those used in the presentation of the Wnt shuttle model in MacLean et al. (2015).

The 19 species in the model interact according to the 31 reactions given in Table 2. Each reaction comes with a rate constant $k_i$. These are the coordinates of our parameter vector $k$. The 31 reactions in Table 2 translate into a dynamical system $\dot{x} = \Psi(x; k)$. Here, $\Psi$ is a vector-valued function of the vectors of species concentrations $x$ and rate constants $k$. The choice of $\Psi$ is up to the modeler. In this paper, we assume that $\Psi$ represents the law of mass action (Klipp et al. 2009, §2.1.1). This is precisely what is used in MacLean et al. (2015) for the Wnt shuttle model. The resulting dynamical system is (1). We refer to Craciun and Feinberg (2005), Feliu and Wiuf (2012), Karp
Table 1 Nineteen species in the Wnt shuttle model

| Variable | Species                        | Symbol |
|----------|--------------------------------|--------|
| $x_1$    | Dishevelled in cytoplasm (inactive) | $D_1$  |
| $x_2$    | Dishevelled in cytoplasm (active) | $D_a$  |
| $x_3$    | Dishevelled in nucleus (active)  | $D_{an}$ |
| $x_4$    | Destruction complex in cytoplasm (active) | $Y_a$  |
| $x_5$    | Destruction complex in cytoplasm (inactive) | $Y_i$  |
| $x_6$    | Destruction complex in nucleus (active) | $Y_{an}$ |
| $x_7$    | Destruction complex in nucleus (inactive) | $Y_{in}$ |
| $x_8$    | Phosphatase in cytoplasm         | $P$    |
| $x_9$    | Phosphatase in nucleus           | $P_n$  |
| $x_{10}$ | $\beta$-catenin in cytoplasm    | $x$    |
| $x_{11}$ | $\beta$-catenin in nucleus      | $x_n$  |
| $x_{12}$ | TCF (gene transcription in nucleus) | $T$    |
| $x_{13}$ | Intermediate complex             | $C$    |
| $x_{14}$ | Intermediate complex, $\beta$-catenin: TCF in nucleus | $C_{xT}$ |
| $x_{15}$ | Intermediate complex, $\beta$-catenin: dishevelled in cytoplasm | $C_{YT}$ |
| $x_{16}$ | Intermediate complex, destruction complex: dishevelled in nucleus | $C_{YD}$ |
| $x_{17}$ | Intermediate complex, destruction complex: phosphatase in cytoplasm | $C_{YP}$ |
| $x_{18}$ | Intermediate complex, destruction complex: phosphatase in nucleus | $C_{YP_n}$ |
| $x_{19}$ | Intermediate complex, $\beta$-catenin: destruction complex in cytoplasm | $C_{xY}$ |
|          | Intermediate complex, $\beta$-catenin: destruction complex in nucleus | $C_{xYn}$ |

et al. (2012), Pérez Millán et al. (2012), and Shiu and Sturmfels (2010) and their many references for mass action kinetics and its variants. In summary, Table 2 can be translated into the dynamical system (1) under the law of mass action. The five relations in (2) constitute a basis for the linear space of conservation relations of the model in Table 2 assuming mass action kinetics.

We refer to $x_1, \ldots, x_{19}$ as the species concentrations, $k_1, \ldots, k_{31}$ as the rate parameters, and $c_1, \ldots, c_5$ as the conserved quantities. We write $\mathbf{x}$, $\mathbf{k}$, and $\mathbf{c}$ for the vectors with these coordinates. As is customary in algebraic geometry, we take the coordinates in the complex numbers $\mathbb{C}$, or possibly in some other algebraically closed field $K$ containing the rationals $\mathbb{Q}$.

Our aim is to understand the relationships between $\mathbf{x}$, $\mathbf{k}$, and $\mathbf{c}$ in the Wnt shuttle model. To this end, we introduce the steady-state variety $S \subset \mathbb{C}^{55}$. This is the set of all points $(\mathbf{x}, \mathbf{k}, \mathbf{c})$ that satisfy the equations $\dot{x}_1 = \cdots = \dot{x}_{19} = 0$ in (1) along with the five conservation laws in (2). We write our ambient affine space as $\mathbb{C}^{55} = \mathbb{C}_x^{19} \times \mathbb{C}_k^{31} \times \mathbb{C}_c^5$. 
Table 2  Thirty-one reactions in the Wnt shuttle model

| Reaction | Explanation |
|----------|-------------|
| \(x_1 \xrightarrow{k_1} x_2\) | (In)activation of dishevelled, depends on Wnt |
| \(x_2 + x_4 \xleftrightarrow{k_3}{k_4} x_{14} \xrightarrow{k_5} x_2 + x_5\) | Destruction complex active \(\rightarrow\) inactive |
| \(x_5 + x_8 \xrightarrow{k_6}{k_7} x_{16} \xrightarrow{k_8} x_4 + x_8\) | Destruction complex inactive \(\rightarrow\) active |
| \(x_4 + x_{10} \xleftrightarrow{k_9}{k_{10}} x_{18} \xrightarrow{k_{11}} x_4 + \emptyset\) | Destruction complex-dependent \(\beta\)-catenin degradation |
| \(\emptyset \xrightarrow{k_{12}} x_{10}\) | \(\beta\)-catenin production |
| \(x_{10} \xrightarrow{k_{13}} \emptyset\) | Destruction complex-independent \(\beta\)-catenin degradation |
| \(x_3 + x_6 \xleftrightarrow{k_{14}}{k_{15}} x_{15} \xrightarrow{k_{16}} x_3 + x_7\) | Destruction complex active \(\rightarrow\) inactive (nucleus) |
| \(x_7 + x_9 \xleftrightarrow{k_{17}}{k_{18}} x_{17} \xrightarrow{k_{19}} x_6 + x_9\) | Destruction complex inactive \(\rightarrow\) active (nucleus) |
| \(x_6 + x_{11} \xleftrightarrow{k_{20}}{k_{21}} x_{19} \xrightarrow{k_{22}} x_6 + \emptyset\) | Destruction complex-dependent \(\beta\)-catenin degradation (nucleus) |
| \(x_{11} \xrightarrow{k_{23}} \emptyset\) | Destruction complex-independent \(\beta\)-catenin degradation (nucleus) |
| \(x_{11} + x_{12} \xleftrightarrow{k_{24}}{k_{25}} x_{13}\) | \(\beta\)-catenin binding to TCF (nucleus) |
| \(x_2 \xrightarrow{k_{26}}{k_{27}} x_3\) | Shuttling of active dishevelled |
| \(x_5 \xrightarrow{k_{28}}{k_{29}} x_7\) | Shuttling of inactive-form destruction complex |
| \(x_{10} \xleftrightarrow{k_{30}}{k_{31}} x_{11}\) | Shuttling of \(\beta\)-catenin |

This emphasizes the distinction between the species concentrations, rate parameters, and conserved quantities.

3 Ideals, Varieties, and Nine Points

We begin here with an initial investigation of the varieties of interest in our model analysis. This section is written in the language of algebraic geometry. For a nonalgebraic
Lemma 3.1 The ideal $I$ admits the nontrivial decomposition $I = I_m \cap I_e$, where $I_e = I : (x_1)$ and $I_m = I + (x_1)$, both of these components have codimension 14, and $I_e$ is a prime ideal.

The ideal $I_m$ is called the main component, while $I_e$ is called the extinction component, since it reflects those steady states where a number of the reactants “run out.” Both of these ideals live in $\mathbb{Q}[x, k]$, and we now present explicit generators. The extinction component equals

$$
I_e = \langle x_1, x_2, x_3, x_5, x_7, x_{14}, x_{15}, x_{16}, x_{17}, k_{30} x_{10} - (k_{23} + k_{31}) x_{11} - k_{22} x_9, k_{13} x_{10} + k_{23} x_{11} + k_{11} x_{18} + k_{22} x_9 - k_{12}, k_{24} x_{11} x_{12} - k_{25} x_{13}, k_{20} x_6 x_{11} - (k_{21} + k_{22}) x_9, k_9 x_4 x_{10} - (k_{10} + k_{11}) x_{18} \rangle.
$$

The ideal $I_e$ is found to be prime in $\mathbb{Q}[x, k]$. The main component equals

$$
I_m = \langle k_{16} x_{15} - k_{19} x_{17}, k_5 x_{14} - k_8 x_6, k_{30} x_{10} - (k_{23} + k_{31}) x_{11} - k_{22} x_9, k_{13} x_{10} + k_{23} x_{11} + k_{11} x_{18} + k_{22} x_9 - k_{12}, k_{28} x_5 - k_{29} x_7, k_{26} x_2 - k_{27} x_3, k_1 x_{11} - k_{22} x_6, k_{24} x_{11} x_{12} - k_{25} x_{13}, k_{20} x_6 x_{11} - (k_{21} + k_{22}) x_9, k_9 x_4 x_{10} - (k_{10} + k_{11}) x_{18}, (k_{4} k_6 k_{8} k_{14} k_{16} k_{18} k_{26} k_{29} + k_{4} k_6 k_{8} k_{14} k_{16} k_{19} k_{26} k_{29} + k_5 k_6 k_{8} k_{14} k_{16} k_{19} k_{26} k_{29} + k_3 k_5 k_7 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_8 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_7 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_8 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_7 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_8 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_7 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_8 k_{16} k_{17} k_{19} k_{27} k_{28} + k_3 k_5 k_7 k_{16} k_{17} k_{19} k_{27} k_{28}) x_{14} x_{9} \rangle.
$$

This ideal is not prime in $\mathbb{Q}[x, k]$. For instance, the variable $k_1$ is a zerodivisor modulo $I_m$, as seen from the last generator. Removing the factor $k_1$ from the last
generator yields the quotient ideal $I_m : \langle k_1 \rangle$. However, even that ideal has several associated primes. All of these prime ideals, except for one, contain some of the rate constants $k_i$.

That special component is characterized in the following proposition. Given any ideal $J \subset \mathbb{Q}[x, k]$, we write $\tilde{J} = \mathbb{Q}(k)[x]J$ for its extension to the polynomial ring $\mathbb{Q}(k)[x]$ in the unknowns $x_1, \ldots, x_{19}$ over the field of rational functions in the parameters $k_1, \ldots, k_{31}$.

**Proposition 3.2** The ideal $J_m = \tilde{I}_m \cap \mathbb{Q}[x, k]$ is prime. Its irreducible variety $V(J_m) \subset \mathbb{C}^{50}$ has dimension 36; it is the unique component of $V(I_m)$ that maps dominantly onto $\mathbb{C}^{31}_k$.

**Proof** The ideal $\tilde{I}_m$ has the same generators as $I_m$ but now regarded as polynomials in $x$ with coefficients in $\mathbb{Q}(k)$. Symbolic computation in the ring $\mathbb{Q}(k)[x]$ reveals that $\tilde{I}_m$ is a prime ideal. This implies that $J_m$ is a prime ideal in $\mathbb{Q}[x, k]$, and hence, $V(J_m)$ is irreducible. The dimension statement follows from the result of Lemma 3.1 that $I_m$ is a complete intersection. This ensures that $V(I_m)$ has no lower-dimensional components, by Krull’s Principal Ideal Theorem. Finally, $V(J_m)$ maps dominantly onto $\mathbb{C}^{31}_k$ because $J_m \cap \mathbb{Q}[k] = \{0\}$.  

**Corollary 3.3** The ideal $\tilde{I}$ is radical, and it is the intersection of two primes in $\mathbb{Q}(k)[x]$:

$$\tilde{I} = \tilde{I}_e \cap \tilde{I}_m.$$  

**Proof** This follows from Proposition 3.2 and the primality of $I_e$ in Lemma 3.1.  

The decomposition has the following geometric interpretation. We now work over the field $K = \mathbb{Q}(k)$. All rate constants are taken to be generic (random rational values). Then, $V(\tilde{I})$ is the five-dimensional variety of all steady states in $K^{19}$. This variety is the union of two irreducible components,

$$V(\tilde{I}) = V(\tilde{I}_e) \cup V(\tilde{I}_m),$$

where each component is five-dimensional. The first component lies inside the 10-dimensional coordinate subspace $V(x_1, x_2, x_3, x_5, x_7, x_{14}, x_{15}, x_{16}, x_{17})$. Hence, it is disjoint from the hyperplane defined by the first conservation relation $x_1 + x_2 + x_3 + x_{14} + x_{15} = c_1$. In other words, $V(\tilde{I}_e)$ is mapped into a coordinate hyperplane under the map $\chi : K^{19} \rightarrow K^5, x \mapsto c$.

On the other hand, the second component $V(\tilde{I}_m)$ maps dominantly onto $K^5$ under $\chi$. Theorem 1.1 states that the generic fiber of this map consists of nine reduced points. Equivalently,

$$\chi^{-1}(c) \cap V(\tilde{I}) = \chi^{-1}(c) \cap V(\tilde{I}_m)$$

is a set of nine points in $K^{19}$. We are now prepared to argue that this is indeed the case.
**Computational Proof of Theorem 1.1** We consider the ideal of the variety (4) in the polynomial ring \( \mathbb{Q}(k, c)[x] \). This polynomial ring has 19 variables, and all 36 parameters are now scalars in the coefficient field. This ideal is generated by the right-hand sides of (1) and (2). Performing a Gröbner basis computation in this polynomial ring verifies that our ideal is zero-dimensional and has length 9 (i.e., the model has nine steady states that are complex or real values). Hence, (4) is a reduced affine scheme of length 9 in \( K^{19} \).

Fast numerical verification of this result is obtained by replacing the coordinates of \( k \) and \( c \) with generic values. In Macaulay2, one finds, with probability 1, that the resulting ideals in \( \mathbb{Q}[x] \) are radical of length 9. We also verified this result via numerical algebraic geometry, using the two software packages Bertini (Bates et al. 2013) and PHCpack (Verschelde 1999).

To summarize the main point of this section, we have now proven that the Wnt shuttle model can have a maximum of nine isolated steady states (over \( \mathbb{C} \)).

### 4 Multistationarity and its Discriminant

This section centers around Question 4 from the Introduction: *For what real positive rate parameters and conserved quantities does the system exhibit multistationarity?* This is commonly asked about biochemical reaction networks and about dynamical systems in general, and it is a question for which computational algebraic geometry has had an active role. For example, it is shown in Pérez Millán et al. (2012) that if the steady-state ideal of the network is binomial, then there are simple sign conditions to test for multistationarity.

Mathematically, this is a problem of *real* algebraic geometry. Writing \( S \) for the steady-state variety in \( \mathbb{C}^{55} \), we are interested in the fibers of the map \( \pi_{k,c} : S \cap \mathbb{R}_{>0}^{55} \rightarrow \mathbb{R}_{>0}^{31} \times \mathbb{R}_{>0}^{5} \). The fiber of \( (k, c) \in \mathbb{R}_{>0}^{31} \times \mathbb{R}_{>0}^{5} \) is the preimage \( \pi_{k,c}^{-1}(k, c) \) and contains the steady states of the system for the rate constants and conserved quantities \( (k, c) \). According to Theorem 1.1, the general fiber consists of nine complex points \( x \in \mathbb{C}^{19} \), when the map \( \pi_{k,c} \) is taken over \( \mathbb{C} \). But here we take it over the reals \( \mathbb{R} \) or over the positive reals \( \mathbb{R}_{>0} \).

In our application to biology, we only care about concentration vectors \( x \) whose coordinates are real and positive. Thus, we wish to stratify \( \mathbb{R}_{>0}^{31} \times \mathbb{R}_{>0}^{5} \) according to the cardinality of

\[
\pi_{k,c}^{-1}(k, c) = \{ (x, k', c') \in S \cap \mathbb{R}_{>0}^{55} : k' = k \text{ and } c' = c \}. \tag{5}
\]

This stratification comes from a decomposition of the 36-dimensionalorthant \( \mathbb{R}_{>0}^{31} \times \mathbb{R}_{>0}^{5} \) into connected open semialgebraic subsets. The walls in this decomposition are given by the *discriminant* \( \Delta \), a giant polynomial in the 36 unknowns \( (k, c) \) that is to be defined later.

We begin with the following result on what is possible with regard to real positive solutions.
Theorem 4.1 Consider the polynomial system in (1)–(2) where all parameters \( k_i \) and \( c_j \) are positive real numbers. The set (5) of positive real solutions can have 1, 2, or 3 elements.

Proof For random choices of \((k, c) = (k_1, \ldots, k_{31}, c_1, \ldots, c_5)\) in the orthant \( \mathbb{R}^{36}_{>0} \), our polynomial system has nine complex solutions, by Theorem 1.1. For the following two special choices of the 36 parameter values, all nine solutions are real. First, take \((k, c)\) to be the vector

\[
(1.7182818, 53.2659, 3.4134082, 0.61409879, 0.61409879, 3.4134082, 0.98168436, 0.98168436, \\
92.331732, 0.86466471, 79.9512906, 97.932525, 1, 3.2654672, 0.61699064, 0.61699064, \\
37.913879, 0.86466471, 0.86466471, 4.7267833, 0.17182818, 0.68292191, 1, 0.55950727, \\
1.0117639, 1.7182818, 1.7182818, 0.99326205, 0.99326205, 5.9744464, 1, 4.9951026, \\
16.4733784, 1.6006340000000001, 1.2089126, 2.7756596399999998).
\]

The resulting system has three positive solutions \( x \in \mathbb{R}^{19}_{>0} \). Next, let \((k', c')\) be the vector

\[
(0.948166, 7.45086, 5.72974, 3.96947, 7.21145, 7.8761, 1.87614, 8.11372, 6.21862, 5.24801, \\
3.10707, 1.08146, 5.22133, 5.84158, 911392, 4.28788, 4.81201, 9.67849, 1.34452, 7.38597, \\
6.64451, 7.10229, 8.57942, 5.79076, 6.33244, 1.53916, 1.39658, 0.81673, 5.8434, 3.86223, \\
7.22696, 1.45438, 3.36482, 6.06453, 4.82045, 3.6014).
\]

Here, one solution to our system is positive. By connecting the two parameter points above with a general curve in \( \mathbb{R}^{36}_{>0} \) and by examining in-between points \((k'', c'')\), we can construct a system with two positive solutions. All computations were carried out using \texttt{Bertini} (Bates et al. 2013). \qed

Remark 4.2 At present, we do not know whether the number of real positive solutions can be larger than three. We suspect that this is impossible, but we currently cannot prove it.

The difficulty lies in the fact that the stratification of \( \mathbb{R}^{36}_{>0} \) is extremely complicated. In computer algebra, the derivation of such stratifications is known as the problem of real root classification. For a sample of recent studies in this direction, see Chen et al. (2013), Faugère et al. (2008), and Rodriguez and Tang (2015). Real root classification is challenging even when the number of parameters is 3 or 4; clearly, 36 parameters are out of the question. The stratification of \( \mathbb{R}^{36}_{>0} \) by behavior of (5) has too many cells.

While symbolic techniques for real root classification are infeasible for our system, we can use numerical algebraic geometry to gain insight into the stratification of \( \mathbb{R}^{36}_{>0} \). Coefficient-parameter homotopies (Morgan and Sommese 1989) can solve the steady-state polynomial system (1)–(2) for multiple choices of \((k, c)\) quickly. For our computations, we use \texttt{Bertini.m2}. This is the \texttt{Bertini} interface for \texttt{Macaulay2}, as described in Bates et al. (2013). Each system has 19 equations in 19 unknowns, and for random \((k, c)\), each system has nine complex solutions. Such a system can be solved in less than one second using the \texttt{bertiniParameterHomotopy} function from \texttt{Bertini.m2}.

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Table 3  Frequencies for the sampling schemes

| # of real solutions | 9    | 7    | 5    | 3    |
|---------------------|------|------|------|------|
| Freq. for uniform sampling | 5760 | 3675 | 544  | 13   |
| Freq. for integer sampling    | 2138 | 5181 | 2522 | 122  |

Table 4  Frequencies for testing robustness scheme

| # of real solutions | Freq. | # of pos. solutions | Freq. |
|---------------------|-------|---------------------|-------|
| 9                   | 9879  | 3                   | 9879  |
| 7                   | 121   | 1                   | 121   |

Below, we describe the following experiment. We sample 10,000 parameter vectors \((k, c)\) from two different probability distributions on \(\mathbb{R}^{36}_{>0}\). In each case, we report the observed frequencies for the number of real solutions and number of positive solutions. We then follow these experiments with a specialized sampling scheme for testing numerical robustness.

**Uniform sampling scheme:** Here, we choose \((k, c)\) uniformly from the cube \((0.0, 100.0)^{36}\). Sampling 10,000 parameter vectors from this scheme and solving the steady-state system for each of these parameter vectors in Bertini, we obtained 9,992 solution sets that contained nine complex points. Solution sets with less than nine points occur when some paths in the coefficient-parameter homotopy fail. We call solution sets with nine solutions good.

**Integer sampling scheme:** Here, we select \((k, c)\) uniformly from \(\{1, 2, 3\}^{36}\). Sampling 10,000 parameter vectors according to this scheme and solving the corresponding steady-state system returned 9,963 good solution sets. Table 3 records how many of the good solution sets had 9, 7, 5, 3 real solutions; all solution sets had one positive real solution.

These computations indicate that for most parameter vectors in \((0, 100)^{36}\), we will see only one positive solution to the steady-state system. But while the set of parameter vectors that result in multiple steady states is not very large, we can give evidence that multistationarity is preserved under small perturbations. This is our next point.

**Testing robustness:** Let \((k^*, c^*)\) be the first point in the proof of Theorem 4.1. For each index \(i \in \{1, \ldots, 19\}\), we choose \(y_i\) uniformly from \((-0.03 \cdot k^*_i, 0.03 \cdot k^*_i)\) and then set \(k_i = k^*_i + y_i\). We ran the same process for the \(c_i\). Sampling 10,000 parameter vectors this way and solving the corresponding steady-state systems returned 10,000 good solution sets, and they are as shown in Table 4.

In the remainder of this section, we properly define the discriminant \(\Delta\) that separates the various strata (regions in the parameter space corresponding to a fixed number of steady states) in \(\mathbb{R}^{36}_{>0}\). Let \(\Delta_{\text{int}}\) denote the Zariski closure in \(\mathbb{C}^{31}_k \times \mathbb{C}^5_c\) of all parameter vectors \((k, c)\) for which (1)–(2) does not have nine isolated complex solutions, and there are no solutions with \(x_i = 0\) for some \(i\). It can be shown that \(\Delta_{\text{int}}\) is a hypersurface that is defined over \(\mathbb{Q}\), so it is given by a unique (up to sign) irreducible squarefree polynomial in \(\mathbb{Z}[k, c]\). We use the symbol \(\Delta_{\text{int}}\) also for that polynomial. To be precise,
\( \Delta_{\text{int}} \) is the discriminant of a number field \( L \) with \( K \supset L \supset \mathbb{Q} \), namely \( L = \mathbb{Q}(k, c) \) is the field of definition of the finite \( K \)-scheme (4).

Next, for any \( i \in \{1, 2, \ldots, 19\} \), consider the intersection of the steady-state variety \( S \) with the hyperplane \( \{x_i = 0\} \). The Zariski closure of the image of \( S \cap \{x_i = 0\} \) under the map \( \pi_{k, c} \) is a hypersurface in \( \mathbb{C}^19 \times \mathbb{C}^3_{\mathcal{L}} \), defined over \( \mathbb{Q} \), and we write \( \Delta_{x_i=0} \) for the unique (up to sign) irreducible polynomial in \( \mathbb{Z}[k, c] \) that vanishes on that hypersurface. We now define

\[
\Delta := \Delta_{\text{int}} \cdot \text{lcm}(\Delta_{x_1=0}, \Delta_{x_2=0}, \ldots, \Delta_{x_{19}=0}).
\]

This product with a least common multiple (lcm) is the discriminant for our problem.

**Example 4.3** The degree of \( \Delta_{\text{int}} \) as a polynomial only in \( c = (c_1, c_2, c_3, c_4, c_5) \) equals 34. To illustrate this, we set \( c = (5, 16+C, \frac{8}{5}−C, \frac{6}{5}+C, 3−C) \) where \( C \) is a parameter, and

\[
k = \left( \frac{9}{5}, \frac{9}{5}, 3, \frac{2}{3}, \frac{2}{3}, 3, 1, 1, 100, \frac{4}{5}, 80, 100, 1, 3, \frac{2}{3}, \frac{2}{3}, 38, \frac{4}{5}, 5, \frac{4}{8}, \frac{3}{5}, 1, \frac{1}{2}, 19, \frac{7}{4}, \frac{7}{4}, 1, 1, 5, 1 \right).
\]

Under this specialization, the polynomial \( \Delta_{\text{int}} \) becomes an irreducible polynomial of degree 34 in the parameter \( C \). Its coefficients are enormously large integers. It has 14 real roots.

For the other factors \( \Delta_{x_i=0} \) of the discriminant, we find the following specializations:

\[
\begin{align*}
x_1 &\to 0, x_2 \to 0, x_3 \to 0, x_4 \to (C+16)(5C−8), x_5 \to C+16, x_6 \to (C+16)(5C+6), \\
x_7 &\to C+16, x_8 \to 5C−8, x_9 \to 5C+6, x_{10} \to \text{a quartic } q(C), x_{11} \to 0, x_{12} \to C−3, \\
x_{13} &\to C−3, x_{14} \to (C+16)(5C−8), x_{15} \to (C+16)(5C+6), x_{16} \to (C+16)(5C−8), \\
x_{17} &\to (C+16)(5C+6), x_{18} \to (C+16)(5C−8)q(C), x_{19} \to (C+16)(5C+6).
\end{align*}
\]

These polynomials have eight distinct real roots in total, so the total number of real roots of the discriminant is \( 14 + 8 = 22 \). These are the break points where real root behavior changes:

\[
\begin{array}{cccccccc}
(9, 0) & -77.2388 & (9, 0) & -16.0000 & (9, 0) & -5.28669 & (7, 0) & -1.57472 \\
(9, 0) & 1.46506 & (9, 0) & -1.34899 & (7, 0) & -1.29581 & (9, 0) & 1.20000 \\
(9, 1) & -1.19215 & (9, 1) & -1.18389 & (7, 1) & -0.584325 & (9, 3) & -0.361808 \\
(7, 3) & 0.191039 & (5, 1) & 1.30812 & (7, 1) & 1.33197 & (5, 1) & 1.60000 \\
(5, 0) & 1.60161 & (3, 0) & 3.0000 & (3, 0) & 4.26306 & (5, 0) & 11.1174 \\
(7, 0) & 21.4165 & (9, 0) & 310.141 & (9, 0) & & & \\
\end{array}
\]

In this table, we list all 22 roots of the specialized discriminant \( \Delta(C) \). The eight boldface values of \( C \) are the roots of (6): Here, one of the coordinates of \( x \) becomes zero. At the other 14 values of \( C \), the number of real roots changes. Between any two
roots, we list the pair \((r, p)\), where \(r\) is the number of real roots and \(p\) is the number of positive real roots. For instance, for \(-0.361808 < C < 0.191039\), there are seven real roots of which three are positive.

We have now proven that three positive real roots can exist for a choice of parameters. Moreover, we characterized the discriminant that stratifies the parameter space into regions of different real root behavior as \(e\) varies.

5 Algebraic Matroids and Parametrizations

Question 5 asks: Suppose we can measure only a subset of the species concentrations. Which subsets can lead to model rejection? This issue is important for the Wnt shuttle model because, in the laboratory, only some of the species have been measured such as \(x_{10}\) and \(x_{11}\), i.e., \(\beta\)-catenin in the cytoplasm and nucleus as well as other members of the destruction complex (Tan et al. 2012).

We shall address Question 5 using algebraic matroids. Matroid theory allows us to analyze the structure of relationships among the 19 species in Table 1. This first appeared in MacLean et al. (2015). We here present an in-depth study of the matroids that govern the Wnt shuttle model.

An introduction to (algebraic) matroids can be found in Oxley (2011); they have been applied in Király et al. (2013, 2015) to problems involving the completion of partial information. General algorithms for computing algebraic matroids are derived in Rosen (2014). We briefly review basic notions.

**Definition 5.1** A matroid is an ordered pair \((X, I)\), where \(X\) is a finite set, here regarded as unknowns, and \(I\) is a subset of the power set of \(X\). These satisfy certain independence axioms. For an algebraic matroid, we are given a prime ideal \(P\) in the polynomial ring \(K[X]\) generated by \(X\), and \(I\) consists of subsets of \(X\) whose images in \(K[X]/P\) are algebraically independent over \(K\). Thus, the collection of independent sets is \(I = \{Y \subseteq X : P \cap K[Y] = \{0\}\}\).

1. **Bases** are maximal independent sets, i.e., subsets in \(I\) with maximal cardinality.
2. **Rank** is a function \(\rho\) from the power set of \(X\) to the natural numbers, which takes as input a set \(Y \subseteq X\) and returns the cardinality of the largest subset of \(Y\) in \(I\).
3. **Closure** is a function from the power set \(2^X\) to itself. The input is a set \(Y\), and the output is the largest set containing \(Y\) with the same rank.
4. **Flats** are the elements in \(2^X\) that lie in the image of the closure map.
5. **Circuits** are the sets of minimal cardinality not contained in \(I\).

We are here interested in the matroid \(M\) that is defined by the prime ideal \(P = \tilde{I_m}\) in \(\mathbb{Q}(k)[x]\). Recall that we use tildes to indicate that we are considering the \(k_i\)s as coefficients as opposed to variables. The ground set \(X\) of the matroid \(M\) the set of species concentrations \(\{x_1, \ldots, x_{19}\}\). Its independent sets are the subsets of \(X\), whose elements are algebraically independent in the quotient ring \(K[X]/\tilde{I_m}\). Since \(V(\tilde{I_m})\) is five-dimensional, each basis consists of five elements in \(X\). In our application,
bases are the maximal subsets of $X$ that can be specified independently at steady state; they are also the minimal-cardinality sets that can be measured to learn all species concentrations. The rank of a set $Y \subseteq \{x_1, \ldots, x_{19}\}$ indicates the number of measurements required to learn the concentrations for every element of $Y$. Choosing an arbitrary set of species concentrations $Y \subseteq \{x_1, \ldots, x_{19}\}$ to measure, the values of the elements in the closure of $Y$, which is a flat, are determined up to finite choice.

Circuits furnish our answer to Question 5: They are minimal sets of species that can be used to test compatibility of the data with the model. For each circuit $Y$, there is a unique-up-to-scalar relation in $\overline{T_m} \cap \mathbb{Q}(k)[Y]$, called the circuit polynomial of $Y$. If the measurements indicate that this relation is not satisfied, then the model and data are not compatible.

Proposition 5.2 The algebraic matroid of $\overline{T_m}$ has rank 5. It has 951 circuits, summarized in Table 5. Of the 11,628 subsets of $X$ of size 5, precisely 2389 are bases. The 2092 bases summarized in Table 6 have base degree 1, while the remaining 297 have base degree 2.

Given a basis $Y \subseteq \{x_1, \ldots, x_{19}\}$ of the algebraic matroid $M$, its base degree is the length of the generic fiber of the projection of $V(\overline{T_m})$ onto the $Y$-coordinates. The computation of the matroid $M$ was carried out using the methods described in Rosen (2014). It was first reported in MacLean et al. (2015), along with the matroids of alternative models for the Wnt pathway. The idea there was to find subsets of variables that were dependent for different models.

Our matroid analysis here goes beyond (MacLean et al. 2015) in several ways:

1. Instead of using generic values for the parameters $k$, we leave the parameters in symbolic form. Specifically, we take our circuit polynomials to have (relatively prime) coefficients in $\mathbb{Z}[k]$. This gives us a new tool for model rejection, e.g., in situations where only one data point is known but some parameter values are available.

2. We show how circuits can be used in parameter estimation; this will be done in Sect. 8.

3. We use the degree-1 bases to derive rational parametrizations of the variety $V(\overline{T_m})$. Such parameterizations can be helpful when analyzing theoretical properties of the model; see, for example, the proof of Proposition 6.2.

We now explain Table 5. A circuit polynomial has type $(i, j)$ if it contains $i$ rate parameters ($k$-variables) and $j$ species concentrations ($x$-variables). The entry in row $i$ and column $j$ in Table 5 is the number of circuits of type $(i, j)$. Zero values are omitted for clarity.

Example 5.3 There are five circuits of type $(2, 2)$. One of them is $\dot{x}_1 = -k_1 x_1 + k_2 x_2$. Most of the 951 circuit polynomials in $\overline{T_m}$ are more complicated. In particular, they are nonlinear in both $x$ and $k$. For instance, the unique circuit polynomial of type $(6, 11)$ equals

$$(-k_{15}k_{17}k_{19}k_{20}k_{25} - k_{16}k_{17}k_{19}k_{20}k_{25})x_7x_9x_{13} + (k_{14}k_{16}k_{18}k_{21}k_{24}$$
$$+ k_{14}k_{16}k_{19}k_{21}k_{24} + k_{14}k_{16}k_{18}k_{22}k_{24} + k_{14}k_{16}k_{19}k_{22}k_{24})x_3x_12x_{19}.$$
In Sect. 7, we will consider the role of these nonlinear functions in parameter estimation.

Because they give rise to rational parameterizations of $V(\tilde{I}_m)$, the most desirable bases of our algebraic matroid are those of base degree 1.

**Proposition 5.4** Let $P \subset K[X]$ be a prime ideal, $Y$ a basis of its algebraic matroid, $|X| = n$ and $|Y| = r$. If $Y$ has base degree 1, then $V(P)$ is a rational variety, and the basic circuits of $Y$ specify a birational map $\varphi_Y : K^r \dashrightarrow K^n$ whose image is Zariski dense in $V(P)$.
Proof For each coordinate $x_i$ in $X \setminus Y$, there exists a circuit containing $Y \cup \{x_i\}$; this is the basic circuit of $(Y, x_i)$. Since $Y$ has base degree 1, the generic fiber of the map $V(P) \to K^r$ consists of a unique point. Therefore, the circuit polynomial is linear in $x_i$. It has the form

$$p_i(Y) \cdot x_i + q_i(Y), \quad \text{where } p_i, q_i \in K[Y].$$

The $i$-coordinate of the rational map $\varphi_Y$ equals $x_i$ if $x_i \in Y$ and $-q_i(Y)/p_i(Y)$ if $x_i \notin Y$. \qed

From Propositions 5.2 and 5.4, we obtain 2092 rational parametrizations of the variety $V(\tilde{\mathcal{I}}_m)$. These are the maps $\varphi_Y : K^5 \to K^9$, where $Y$ runs over all bases of base degree 1. Using these $\varphi_Y$, we obtain 2092 representations of the steady-state variety (4) as a subset of $K^5$, where now $K = \mathbb{Q}(k, c)$. Namely, we consider the preimages of the five hyperplanes defined by (2). These are hypersurfaces in $K^5$ whose intersection represents the nine points in (4). We performed the following computation for all 2092 bases $Y = \{x_1, \ldots, x_9\}$ of base degree 1:

1. Substitute $x = \varphi_Y(x_1, \ldots, x_9)$ into the five linear equations (2).
2. Clear the denominators $d_1, \ldots, d_5$ in each equation to get polynomials $h_1, \ldots, h_5$ in $K[Y]$.
3. The saturation ideal $J_Y = \langle h_1, \ldots, h_5 \rangle : \langle d_1d_2 \cdots d_5 \rangle^\infty$ represents the preimage of (4) under $\varphi_Y$.

Given such a large choice of parametrizations, we seek one where $J_Y$ has desirable properties, for example where the structure of the generating polynomials coincides with known properties of $V(\tilde{\mathcal{I}}_m)$. We use the following criterion: consider subsets of five of the generators of $J_Y$, compute the mixed volume of their Newton polytopes, and fix a subset minimizing that mixed volume. In the census of 2092 bases in Table 6, that minimum is referred to as the mixed volume of $Y$.

By Bernstein’s Theorem, the mixed volume is the number of solutions to a generic system with the five given Newton polytopes. We seek bases $Y$ where this matches the number nine from Theorem 1.1. We see that the mixed volume is nine for 416 of the bases in Table 6.

Example 5.5 The basis $Y = \{x_1, x_4, x_6, x_8, x_{13}\}$ has base degree 1 and mixed volume 9. The other variables can be expressed in terms of $Y$ as follows. For brevity, we set

$$r(x_4, x_6) = k_9k_{11}k_{20}k_{22}x_4x_6 + k_9k_{11}(k_{21} + k_{22})(k_{23} + k_{31})x_4$$

$$+ k_{20}k_{22}(k_{10} + k_{11})(k_{13} + k_{30})x_6 + (k_{10} + k_{11})(k_{21} + k_{22})(k_{13}k_{23} + k_{23}k_{30} + k_{13}k_{31}).$$

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x_2 = \frac{k_1}{k_2} x_1

x_3 = \frac{k_1 k_{26}}{k_{27}} x_1

x_5 = \frac{k_1 k_5 (k_7 + k_8)}{k_2 k_6 k_8 (k_4 + k_5)} x_8

x_7 = \frac{k_1 k_5 k_9 (k_7 + k_8)}{k_2 k_5 k_9 k_29 (k_4 + k_5)} x_8

x_9 = \frac{k_1 k_4 k_{16} k_{26} k_{29} (k_4 + k_5) (k_{18} + k_{19})}{k_2 k_4 k_5 k_{17} k_{19} k_{27} (k_7 + k_8) (k_15 + k_16)} x_6 x_8

x_{10} = \frac{k_1 k_{26} (k_{10} + k_{11}) (k_{30} + k_{31} + k_{10} + k_{22}) (k_{23} + k_{31})}{r(x_4, x_6)}

x_{11} = \frac{k_1 k_{30} (k_{10} + k_{11}) (k_{21} + k_{22})}{r(x_4, x_6)}

x_{12} = \frac{k_1 k_{30} (k_{10} + k_{11}) (k_{21} + k_{22})}{r(x_4, x_6)} \frac{k_{25}}{k_{24} x_3}

x_{14} = \frac{k_1 k_3}{k_2 (k_4 + k_5)} x_1 x_4

x_{15} = \frac{k_1 k_{14} k_{26}}{k_{27} (k_{15} + k_{16})} x_1 x_6

x_{16} = \frac{k_1 k_3 k_5}{k_2 k_8 (k_4 + k_5)} x_1 x_4

x_{17} = \frac{k_1 k_{14} k_{16} k_{26}}{k_{27} (k_{15} + k_{16})} x_1 x_6

x_{18} = \frac{k_1 k_{14} k_{16} k_{26} (k_{20} + k_{22} + k_{23} + k_{31})}{r(x_4, x_6)} x_4

x_{19} = \frac{k_1 k_{20} k_{30} (k_{10} + k_{11})}{r(x_4, x_6)} x_6

This map \( \varphi_Y \) is substituted into (2), and then we saturate. The resulting ideal \( J_Y \) equals

\[
(\alpha_1 x_6 x_8 + \alpha_2 x_4 + \alpha_3 x_6, \quad \alpha_4 x_1 x_6 + \alpha_5 x_1 + \alpha_6 x_8 + \alpha_7,
\]

\[
\alpha_8 x_1 x_4 + \alpha_9 x_8 + \alpha_{10}, \quad \alpha_{11} x_4 x_6 x_1 + \alpha_{12} x_4 x_1 + \alpha_{13} x_6 x_1 + \alpha_{14} x_1 + \alpha_{15},
\]

\[
\alpha_{16} x_4 x_6^2 + \alpha_{17} x_6^3 + \alpha_{18} x_4 x_6 + \alpha_{19} x_6^2 + \alpha_{20} x_8^2 + \alpha_{21} x_1 + \alpha_{22} x_4 + \alpha_{23} x_6 + \alpha_{24} x_8 + \alpha_{25},
\]

where the \( \alpha_1, \ldots, \alpha_{25} \) are certain explicit rational functions in the \( k \)-parameters.

The outcome of the preceding analysis is a collection of parametrizations which describe the variety \( V(\tilde{J}_m) \) in terms of fewer variables, all of which were among our original set. Now, instead of a large polynomial system which acts in a highly special way, we have a smaller polynomial system, whose corresponding variety acts more in line with the defining polynomials, i.e., the mixed volume of the polynomial system is equal to the number of solutions for generic coefficients. As a result, numerical techniques for estimation will be better conditioned.

### 6 Polyhedral Geometry

Dynamics of the system while not at steady state cannot typically be studied with algebraic methods. One exception is the set of all possible states accessible from a given set of initial values via the chemical reactions in the model. This set is called a stoichiometric compatibility class in the biochemistry literature. Mathematically, these classes are convex polyhedra. We determine them all for the Wnt shuttle model. This resolves Problem 6 from the Introduction.

The conservation relations (2) define a linear map \( \chi \) from the orthant of concentrations \( \mathbb{R}_{\geq 0}^{19} \) to the orthant of conserved quantities \( \mathbb{R}_{\geq 0}^5 \). We express this projection as a \( 5 \times 19 \)-matrix:
\[
\begin{pmatrix}
  c_1 \\
  c_2 \\
  c_3 \\
  c_4 \\
  c_5
\end{pmatrix} =
\begin{pmatrix}
  1 & 1 & 1 & \ldots & \ldots & 1 & 1 & \ldots \\
  \ldots & 1 & 1 & 1 & 1 & 1 & 1 & 1 & 1 \\
  \ldots & \ldots & 1 & \ldots & \ldots & 1 & 1 & \ldots \\
  \ldots & \ldots & \ldots & \ldots & \ldots & 1 & 1 & \ldots \\
  \ldots & \ldots & \ldots & \ldots & \ldots & \ldots & \ldots & \ldots
\end{pmatrix}
\begin{pmatrix}
  x_1 \\
  x_2 \\
  x_3 \\
  \vdots \\
  x_{18} \\
  x_{19}
\end{pmatrix}
\] (7)

Let \( P_c \) denote the fiber of the map \( \chi \) for \( c \in \mathbb{R}^5_{\geq 0} \). This is known in the biochemical literature as the invariant polyhedron or the stoichiometric compatibility class of the given \( x \); see, e.g., [Shiu and Sturmfels 2010, (3)]. The fiber over the origin is \( P_0 = \mathbb{R}^5_{\geq 0} \{ e_{10}, e_{11} \} \), the two-dimensional orthant formed by all positive linear combinations of the basis vectors \( e_{10} \) and \( e_{11} \). If \( c \in \mathbb{R}^5_{\geq 0} \) is an interior point, then \( P_c \) is a 14-dimensional convex polyhedron of the form \( P_0 \times \tilde{P}_c \) where \( \tilde{P}_c \) is a 12-dimensional (compact) polytope. Two vectors \( c \) and \( c' \) are considered equivalent if their invariant polyhedra \( P_c \) and \( P_{c'} \) have the same normal fan. This property is much stronger than being combinatorially isomorphic. The equivalence classes are relatively open polyhedral cones, and they define a partition of \( \mathbb{R}^5_{\geq 0} \). This partition is the chamber complex of the matrix (7). For a low-dimensional illustration, see (Shiu and Sturmfels 2010, Figure 1). Informally speaking, the chamber complex classifies the possible boundary behaviors of our dynamical system.

**Proposition 6.1** The chamber complex of our \( 5 \times 19 \)-matrix divides \( \mathbb{R}^5_{\geq 0} \) into 19 maximal cones. It is the product of a ray, \( \mathbb{R}_{\geq 0} \), and the cone over a subdivision of the tetrahedron. That subdivision consists of 18 smaller tetrahedra and 1 bipyramid, described in detail below.

**Proof** The product structure arises because the matrix has two blocks after permuting columns, an upper left \( 4 \times 17 \) block and a lower right \( 1 \times 2 \) block (1 1). Our task was to compute the chamber decomposition of \( \mathbb{R}^4_{\geq 0} \) defined by the \( 4 \times 17 \)-block. After deleting zero columns and multiple columns, we are left with a \( 4 \times 7 \)-matrix, given by the seven left columns in

\[
M =
\begin{pmatrix}
  a & b & c & d & e & f & g \\
  0 & 1 & 0 & 0 & 1 & 0 & 0 \\
  1 & 1 & 1 & 0 & 0 & 0 & 1 \\
  0 & 0 & 1 & 0 & 0 & 1 & 0 \\
  0 & 0 & 0 & 1 & 0 & 0 & 1
\end{pmatrix}
\]

The correspondence between the seven left columns of \( M \) and the columns of (7) is as follows:

\[
a = \{ x_4, x_5, x_6, x_7, x_{18}, x_{19} \}, \quad b = \{ x_{14}, x_{15} \}, \quad c = \{ x_{16} \},
\]

\[
d = \{ x_{17} \}, \quad e = \{ x_1, x_2, x_3 \}, \quad f = \{ x_8 \}, \quad g = \{ x_9 \}.
\]

The remaining columns of \( M \) are additional vertices in the subdivision.
The following table lists the 19 maximal chambers. For each chamber, we list the extreme rays and the facet-defining inequalities. For instance, the chamber in $\mathbb{R}^5_{\geq 0}$ denoted by $efjk$ is the orthant spanned by the columns $e, f, j, k$ and of the matrix $M$ times the ray $(0, 0, 0, 0, 1)^T$. It is defined by $c_5 \geq 0$ together with the four listed inequalities: $c_4 \geq 0$, $\min(c_1, c_3) \geq c_2 \geq c_4$.

| Chamber | Inequalities |
|---------|-------------|
| $abcd$  | $\{c_4, c_3, c_1, c_2 - c_4 - c_3 - c_1\}$ |
| $bcdl$  | $\{c_2 - c_3 - c_1, c_2 - c_4 - c_1, c_2 - c_4 - c_3, -c_2 + c_4 + c_3 + c_1\}$ |
| $efjk$  | $\{c_2 - c_2 + c_4, -c_2 + c_3, -c_2 + c_1\}$ |
| $bcjl$  | $\{c_4, -c_2 + c_1 + c_3, c_2 - c_3 - c_4, c_2 - c_1 - c_4\}$ |
| $bdil$  | $\{c_3, -c_2 + c_4 + c_1, c_2 - c_4 - c_3, c_2 - c_3 - c_1\}$ |
| $beij$  | $\{c_3, c_4, c_1 - c_2, c_2 - c_3 - c_4\}$ |
| $cdhl$  | $\{c_1, -c_2 + c_3 + c_4, c_2 - c_4 - c_3, c_2 - c_4 - c_1\}$ |
| $cfhh$  | $\{c_4, c_1, -c_2 + c_3, c_2 - c_4 - c_1\}$ |
| $dghi$  | $\{c_1, c_3, -c_2 + c_4, c_2 - c_1 - c_3\}$ |
| $egik$  | $\{c_3, -c_2 + c_4, c_2 - c_3, -c_2 + c_1\}$ |
| $fghk$  | $\{c_1, -c_1 + c_2, -c_2 + c_3, -c_2 + c_4\}$ |
| $efjk$  | $\{c_4, c_1 - c_2, c_2 - c_4, -c_2 + c_3\}$ |
| $bijl$  | $\{c_2 - c_1, -c_2 + c_1 + c_3, -c_2 + c_4 + c_1, c_2 - c_3 - c_4\}$ |
| $chjl$  | $\{c_2 - c_3, -c_2 + c_4 + c_3, -c_2 + c_3 + c_1, c_2 - c_4 - c_1\}$ |
| $dhl$   | $\{c_2 - c_4, -c_2 + c_4 + c_1, -c_2 + c_3 + c_4, c_2 - c_1 - c_3\}$ |
| $ghik$  | $\{c_4 - c_2, c_2 - c_3, c_2 - c_1, -c_2 + c_1 + c_3\}$ |
| $eijk$  | $\{c_2 - c_4, c_2 - c_3, c_1 - c_2, -c_2 + c_3 + c_4\}$ |
| $fhjk$  | $\{c_2 - c_4, c_2 - c_1, -c_2 + c_3, -c_2 + c_4 + c_1\}$ |
| $hijkl$ | $\{c_2 - c_4, c_2 - c_3, c_2 - c_1, -c_2 + c_4 + c_3, -c_2 + c_4 + c_1, -c_2 + c_3 + c_1\}$ |

Interpreting the columns of $M$ as homogeneous coordinates, the table describes a subdivision of the standard tetrahedron into 18 tetrahedra and one bipyramid $hijkl$. These cells use the 12 vertices $a, b, \ldots, l$. The reader is invited to check that this subdivision has precisely 39 edges and 47 triangles, so the Euler characteristic is correct: $12 - 39 + 47 - 19 = 1$.

We shall prove the following result about the Wnt shuttle model.

**Proposition 6.2** Suppose that the rate constants $k_i$ and the conserved quantities $c_j$ are all strictly positive. Then, no steady states exist on the boundary of the invariant polyhedron $P_e$.

**Proof** Consider the two components $I_m$ and $I_e$ of the steady-state ideal $I$ given in Lemma 3.1. We intersect each of the two varieties with the affine-linear space defined by the conservation relations (2) for some $e \in \mathbb{R}^5_{\geq 0}$. We claim that all solutions $x$ satisfy $x_i \neq 0$ for $i = 1, 2, \ldots, 19$.

For the main component $V(I_m)$, we prove this assertion with the help of the parametrization $\varphi_Y$ from Example 5.5. If the values of $x_1, x_4, x_6, x_8, x_{13}$ and of the expression $r(x_4, x_6)$ are nonzero, then each coordinate of $\varphi_Y$ is nonzero. We next observe that $r(x_4, x_6) > 0$ for any $k > 0$ and $x \geq 0$. A case analysis, using binomial relations in the ideal $I_m$, reveals that if any of $x_1, x_4, x_6, x_8, x_{13}$ are zero, some coordinate of $e$ is forced to zero as well.
It remains to consider the extinction component. Its ideal $I_e$ contains the set $b \cup e = \{x_1, x_2, x_3, x_{14}, x_{15}\}$. The corresponding columns of the matrix in (7) are the only columns with a nonzero entry in the fourth row. This implies that $c_4 = 0$ holds for every steady state in $V(I_e)$. We conclude that there are no steady states on the boundary of the polyhedron $P_c$. □

Remark 6.3 In this proof, we did not need the detailed description of the chamber complex, because of the special combinatorial structure in the Wnt shuttle model. In general, when studying chemical reaction networks that arise in systems biology, an analysis like Proposition 6.1 is requisite for gaining information about possible zero coordinates in the steady states.

Algebraically, our proof shows that the extinction component $I_e$ is not relevant when all conserved quantities $c_i$ are strictly positive. Indeed, at all points $x$ in the variety $V(I_e)$, at least one of the corresponding linear forms evaluates to zero.

To summarize this section, we have now fully characterized each set of equivalent stoichiometric compatibility classes for the Wnt shuttle model. Furthermore, we showed that if the rate constants and conserved quantities are strictly positive, then all steady states have strictly positive species concentrations.

7 Parameter Estimation

Question 7 asks: What information does species concentration data give us for parameter estimation? This question is of particular importance to experimentalists, as species concentrations depend on initial conditions, whereas parameter values are intrinsic to the biological process being modeled. Identifiability of parameters has been studied in many contexts, notably in statistics (Garcia-Puente et al. 2013) and in biological modeling (Meshkat and Sullivant 2014). Sometimes, as in Meshkat and Sullivant (2014), parameters are determined from complete time course data of the dynamical system, making a differential algebra approach desirable. In the present paper, we focus on the steady-state variety, so we consider data collection only at steady state. We assume that there is a true but unknown parameter vector $k^* \in \mathbb{R}^{31}$ of rate constants, and our data are sampled from the positive real points $x$ on the variety in $\mathbb{R}^{19}$ that is defined by the 19 polynomials in (1).
7.1 Complete Species Information

The first algebraic question we answer: To what extent is the true parameter vector $k^*$ determined by points on its steady-state variety?

To address this question, we form the polynomial matrix $F(x)$ of format $19 \times 31$ whose entries are the coefficients of the right-hand sides of (1) regarded as linear forms in $k$, i.e., the entries of $F(x)$ are polynomials in $x$. With this notation, our dynamical system (1) can be written in matrix-vector product form as

$$\dot{x} = F(x) \cdot k.$$ 

Our noiseless data points are sampled from

$$\{ x \in \mathbb{R}_{\geq 0}^{19} : F(x) \cdot k^* = 0 \}. \quad (8)$$

Let $x_1, x_2, x_3, \ldots$ denote generic data points in (8). The set of all parameter vectors $k$ that are compatible with these data is a linear subspace of $\mathbb{R}^{31}$, namely it is the intersection

$$\text{kernel}(F(x_1)) \cap \text{kernel}(F(x_2)) \cap \text{kernel}(F(x_3)) \cap \cdots \quad (9)$$

The best we can hope to recover from sampling data is the following subspace containing $k^*$:

$$\bigcap_{x \text{ in } (8)} \text{kernel}(F(x)) \subset \mathbb{R}^{31}. \quad (10)$$

We refer to (10) as the space of parameters compatible with $k^*$. A direct computation reveals:

**Proposition 7.1** The space of all parameters compatible with $k^*$ is a 14-dimensional subspace of $\mathbb{R}^{31}$. If $x$ is generic, then the kernel of $F(x)$ is a 17-dimensional subspace of $\mathbb{R}^{31}$.

This has the following noteworthy consequence for our biological application:

**Corollary 7.2** The parameters of the Wnt shuttle model are not identifiable from steady-state data, but there are 14 degrees of freedom in recovering the true parameter vector $k^*$.

Even though the parameters of the Wnt shuttle model are not identifiable, we would still like to gain a more precise understanding of the subspaces in Proposition 7.1. To do this, we shall return to the combinatorial setting of matroid theory. We introduce two matroids on the 31 reactions in Table 2. The common ground set is $K = \{ k_1, k_2, \ldots, k_{31} \}$. The one-point matroid $M_{\text{one}}$ is the rank 17 matroid on $K$ defined by the linear subspace kernel($F(x)$) of $\mathbb{R}^{31}$ where $x \in \mathbb{R}^{19}$ is generic. The parameter matroid $M_{\text{par}}$ is the rank 14 matroid on $K$ defined by the space (10) of all parameters compatible with a generic $k^*$. The following result, obtained by calculations, reflects the block structure of the matrix $F(x)$. 

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Proposition 7.3 The one-point matroid $\mathcal{M}_{\text{one}}$ is the graphic matroid of the graph shown in Fig. 1a. Its seven connected components are matroids of ranks 3, 3, 7, 1, 1, 1, 1. The rank 14 parameter matroid $\mathcal{M}_{\text{par}}$ is obtained from $\mathcal{M}_{\text{one}}$ by specializing the rank 7 component to the rank 4 matroid on 11 elements whose affine representation is shown in Fig. 1b.

This characterizes the combinatorial constraints imposed on the parameters $k$ by measuring the species concentrations at steady state. For a single measurement $x$, the result on $\mathcal{M}_{\text{one}}$ tells us that the $19 \times 31$-matrix $F(x)$ has rank $14 = 31 - \text{rank}(\mathcal{M}_{\text{one}})$. After row operations, it block-decomposes into two matrices of format $3 \times 6$, one matrix of format $4 \times 11$, and four matrices of format $1 \times 2$. Each of these seven matrices is row-equivalent to the node–edge cycle matrix of a directed graph, with underlying undirected graph as in Fig. 1a.

Consider the graph with edges 9, 10, 11, 12, 13, 20, 21, 22, 23, 30, 31, these numbers correspond to the indices on the rate parameters. The cycle $\{22, 23, 30, 31\}$ reveals that our measurement $x$ imposes one linear constraint on $k_{22}, k_{23}, k_{30}, k_{31}$. If we take further measurements, as in (9), then six of the seven blocks of $F(x)$ remain unchanged. Only the $4 \times 11$-block of $F(x)$ must be enlarged, to a $7 \times 11$-matrix. The rows of that new matrix specify the affine-linear dependencies among 11 points in $\mathbb{R}^3$. That point configuration is depicted in Fig. 1b. For instance, the points $\{9, 10, 11\}$ are collinear and the points $\{20, 21, 22\}$ are collinear, but these two lines are skew in $\mathbb{R}^3$. From the other line, we see that that repeated measurements at steady state impose two linear constraints on $k_{22}, k_{23}, k_{30}, k_{31}$. The matroid describing the relations among rate parameters indicates what information about parameters we can extract from thorough species concentration data. When data are plentiful, this tells us what constraints the $k$-variables satisfy.

7.2 Circuit Data

The second question we address in this section: Given partial species concentration data, is any information about parameters available? In Sect. 7.1, all 19 concentrations...
$x_i$ were available for a steady state. In what follows, we suppose that $x_i$ can only be measured for indices $i$ in a subset of the species, say $C \subset \{1, \ldots, 19\}$. In our analysis, it will be useful to take advantage of the rank 5 algebraic matroid in Proposition 5.2, since that matroid governs dependencies among the coordinates $x_1, \ldots, x_{19}$ at steady state.

We here focus on the special case when $C$ is one of the 951 circuits of the algebraic matroid of $\mathcal{I}_m$. Let $f_C$ be the corresponding circuit polynomial, as in Table 5. We regard $f_C$ as a polynomial in $x$ whose coefficients are polynomials in $\mathbb{Q}[k]$. Suppose that $f_C$ has $r$ monomials $x^{a_1}, \ldots, x^{a_r}$. We write $\Phi_C \in \mathbb{Q}[k]^r$ for the row vector of coefficients, so our circuit polynomial is the dot product $f_C(k, x) = \phi_C(k) \cdot (x^{a_1}, \ldots, x^{a_r})^T$. We write $\mathcal{V}_C \subset \mathbb{R}^r$ for the algebraic variety parametrized by $\Phi_C(k)$. Thus, $\mathcal{V}_C$ is the Zariski closure in $\mathbb{R}^r$ of the set $\{\Phi_C(k') : k' \in \mathbb{R}^{31}\}$.

Our idea for parameter recovery is this: Rather than looking for a $k$ compatible with the true parameter $k^*$, in the sense of Proposition 7.1, we seek a point $y = \Phi_C(k)$ in $\mathcal{V}_C$ that is compatible with $\Phi_C(k^*)$. And, only later do we compute a preimage of $y$ under the map $\mathbb{R}^{31} \to \mathbb{R}^r$ given by $\Phi_C$. Most interesting is the case when $\mathcal{V}_C$ is a proper subvariety of $\mathbb{R}^r$. Direct computations yield the following:

**Proposition 7.4** For precisely 288 of the 951 circuits $C$ of the algebraic matroid of the steady-state ideal $\mathcal{I}_m$, the coefficient variety $\mathcal{V}_C$ is a proper subvariety in its ambient space $\mathbb{R}^r$. In each of these cases, the defining ideal of $\mathcal{V}_C$ is of one of the following four types:

\begin{align*}
   &\langle y_2 y_6 - y_3 y_5 \rangle \\  &\langle y_5 y_6 - 2 y_3 y_7, y_5^2 - 4 y_2 y_7, y_3 y_5 - 2 y_2 y_6, y_2 y_6^2 - y_3^2 y_7 \rangle \\  &\langle y_3 y_5^2 - y_2 y_5 y_6 + y_1 y_6^2 \rangle \\  &\langle 2 y_3 y_4 - y_2 y_5, y_2 y_3 - 2 y_1 y_5, y_2^2 - 4 y_1 y_4 \rangle
\end{align*}

**Example 7.5** Consider the circuit $C = \{6, 10, 18\}$. The circuit polynomial $f_C$ equals

\[
(k_{13} k_{20} k_{22} + k_{20} k_{22} k_{30}) \cdot x_6 x_{10} + k_{11} k_{20} k_{22} \cdot x_6 x_{18} - k_{12} k_{20} k_{22} \cdot x_6 \\
+ (k_{13} k_{21} k_{23} + k_{13} k_{22} k_{23} + k_{21} k_{23} k_{30} + k_{22} k_{23} k_{30} + k_{13} k_{21} k_{31} + k_{13} k_{22} k_{31}) \cdot x_{10} \\
+ (k_{11} k_{21} k_{23} + k_{11} k_{22} k_{23} + k_{11} k_{21} k_{31} + k_{11} k_{22} k_{31}) \cdot x_{18} \\
- (k_{12} k_{21} k_{23} + k_{12} k_{22} k_{23} + k_{12} k_{21} k_{31} + k_{12} k_{22} k_{31}).
\]

Here, $r = 6$, and we write $\Phi_C(k) = (y_1, y_2, y_3, y_4, y_5, y_6)$ for the vector of coefficient polynomials, for example $y_1 = k_{13} k_{20} k_{22} + k_{20} k_{22} k_{30}$. The variety $\mathcal{V}_C$ is the hypersurface in $\mathbb{R}^5$ defined by the equation $y_2 y_6 = y_3 y_5$.

We now sample data points $x_i$ from the model with the true (but unknown) parameter vector $k^*$. Each such point defines a hyperplane $\{y \in \mathbb{R}^r : y \cdot (x_1^{a_1}, \ldots, x_r^{a_r}) = 0\}$. The parameter estimation problem is to find the intersection of these data hyperplanes with the variety $\mathcal{V}_C$. That intersection contains the point $y^* = \Phi_C(k^*)$, which is what we aimed to recover; all $k \in \Phi^{-1}(y^*)$ would be compatible with the model.
7.3 Noisy Circuit Data

The final question we consider in this section is: *Given partial species concentration data with noise, is any information about parameters available?*

As in Sect. 7.2, we fix a circuit $C$ of the algebraic matroid in Sect. 5, and we assume that we can only measure the concentrations $x_j$ where $j \in C$. Each measurement $x_i \in \mathbb{R}^C$ still defines a hyperplane $y \cdot (x_i^a, \ldots, x_i^r) = 0$ in the space $\mathbb{R}^r$. But now, the true vector $y^* = \Phi_C(k^a)$ is not exactly on that hyperplane, but only close to it. Hence, if we take $s$ repeated measurements, with $s > r$, the intersection of these hyperplanes should be empty.

We propose to find the best fit by solving the following least squares optimization problem:

\[
\text{Minimize } \sum_{i=1}^{s} \left( y \cdot (x_i^a, \ldots, x_i^r) \right)^2 \text{ subject to } y \in V_C \cap S^{r-1},
\]

where $S^{r-1} = \{ y \in \mathbb{R}^r : y_1^2 + y_2^2 + \cdots + y_r^2 = 1 \}$ denotes the unit sphere. When the variety $V_C$ is the full ambient space $\mathbb{R}^r$, this is a familiar regression problem, namely to find the hyperplane through the origin that best approximates $s$ given points in $\mathbb{R}^r$. Here “best” means that the sum of the squared distances of the $s$ points to the hyperplane is minimized. This happens for 663 of the 951 circuits $C$, and in that case, we can apply standard techniques.

However, for the 288 circuits $C$ identified in Proposition 7.4, the problem is more interesting. Here, the hyperplanes under consideration are constrained to live in a proper subvariety. In that case, we need some algebraic geometry to reliably find the global optimum in (15) (Gross et al. 2015).

Our problem is to minimize a quadratic function over the real affine variety $V_C \cap S^{r-1}$. The quadratic objective function is generic because the $x_i$ are sampled with noise. The intrinsic algebraic complexity of our optimization problem was studied by Draisma et al. (2015). That complexity measure is the *ED degree* of $V_C \cap S^{r-1}$, which is the number of solutions in $\mathbb{C}^r$ to the critical equations of (15). Here, by ED degree, we mean the ED degree of $V_C \cap S^{r-1}$, when considered in generic coordinates. This was called the *generic ED degree* in Ottaviani et al. (2014).

We illustrate our algebraic approach by working out the first instance (11) in Proposition 7.4.

*Example 7.6* Suppose we are given $s$ noisy measurements of the concentrations $x_6, x_{10}, x_{18}$. In order to find the best fit for the parameters $k$, we employ the circuit polynomial $f_C$ in Example 7.5. We compute $y \in \mathbb{R}^6$ by solving the corresponding optimization problem (16). This problem is to minimize a random quadratic form subject to two quadratic constraints

\[
y_2y_6 - y_3y_5 = y_1^2 + y_2^2 + y_3^2 + y_4^2 + y_5^2 + y_6^2 - 1 = 0.
\]

We solve this problem using the method of Lagrange multipliers. This leads to a system of polynomial equations in $y$. Using saturation, we remove the singular locus.
of (16), which is the circle \( \{ y \in \mathbb{R}^6 : y_1^2 + y_4^2 - 1 = y_2 = y_3 = y_5 = y_6 = 0 \} \). The resulting ideal has precisely 40 zeros in \( \mathbb{C}^6 \). In the language of Draisma et al. (2015) and Ottaviani et al. (2014), the generic ED degree of the variety (16) equals 40.

There are many ways to approach the problem of parameter estimation. Numerical algebraic geometry (Gross et al. 2015) is one of them. In this section, we used the Wnt shuttle model to demonstrate the breadth of geometric and combinatorial questions that can arise in parameter estimation, including and beyond understanding the geometry of the underlying quadratic optimization problem. Furthermore, we showed how matroid theory can be used to validate/invalidate models and provide relationships about parameters for parameter estimation.

8 From Algebra to Biology

The aims of this paper were (1) to demonstrate how biology can lead to interesting questions in algebraic geometry, and (2) to apply new techniques from computational algebra in biology. So far, our tour through (numerical) algebraic geometry, polyhedral geometry, and combinatorics has demonstrated a range of mathematical questions to explore, such as the number of steady states (over \( \mathbb{C} \)), the chamber complex that provides permissible regions of dynamics, as well as algebraic matroids for parameter identifiability, experimental design, and model rejection.

In this section, we will focus on translating our analysis into applicable considerations for the research cycle in systems biology, as illustrated in Fig. 2. In what follows, we discuss some concrete applications and results pertaining to the steps (a), (b), and (c) in Fig. 2.

**Analysis of the Model:** Before any experiments are performed, our techniques inform the modeler of the global steady-state properties of the model. The number of real solutions to system (1)–(2), stated in Theorem 1.1, governs the number of observable steady states. Various sampling schemes, as described in Sect. 4, demonstrated that most parameter values lead to only one observable steady state for the Wnt shuttle model. We produced a set of parameter values and conserved quantities with three real solutions, and two solutions are also attainable. If the “true” parameters \( k^* \) and...
If multiple states are observed experimentally, then the model must be capable of multistationarity. In the Wnt shuttle model, the system is capable of multiple steady states; however, based on parameter sampling, the frequency of this occurrence is low, and parameters in this regime are somewhat stable under perturbation. The discriminant of the system is a polynomial of degree 34 in $c$, and our analysis along a single line in $c$-space illustrates the high degree of complexity inherent in the full stratification of the 36-dimensional parameter space.

**Experimental Design:** In Sect. 6, the combinatorial structure of the various stoichiometric compatibility classes was fully characterized. As the conserved quantities $c = (c_1, \ldots, c_5)$ range over all positive real values, the set of all compatible species concentration vectors $x$ will take one of 19 polyhedral shapes $P_c$. This may find application in identifying multiple steady-state solutions for specific rate constants $k$. A possible choice for initial conditions when performing experiments is on or near the vertices of the 14-dimensional polyhedron $P_c$. The idea is that the primary steady state would lie deep in the interior of $P_c$, but one or more additional steady states could occur when rate concentrations are most extreme given fixed conserved quantities.

**Example 8.1** Suppose the conserved quantities vector lies in the bipyramid described in Proposition 6.1, e.g., $c = (1, 2, 2, 2, 3)$. The preimage of $c$ in $x$-space is a product of the orthant $\mathbb{R}_{\geq 0}\{e_{10}, e_{11}\}$ and a 12-dimensional polytope with 400 vertices:

$$(1, 0, 0, 2, 0, 0, 0, 2, 2, 0, 3, 0, 0, 0, 0, 0, 0, 0) ,$$

and 399 of its permutations. This product is the polyhedron $P_c$. If we have control over initial conditions, we might begin by exploring near the vertices. As suggested above, this might reveal interesting systems behavior, like multiple steady states.

In the laboratory, the experimentalist makes choices of what to measure and what not to measure. For instance, measuring a particular $x_i$ may be infeasible, or there may be a situation in which measuring concentration $x_i$ can preclude measuring concentration $x_j$.

For every strategy, we fix a cost vector, listing the costs of making each measurement. We use the symbol $N$ to indicate infeasible measurements. Suppose there are two different ways to run the experiment; then, we have a $2 \times 19$ cost matrix $P$, whose rows are cost vectors for each experiment. We multiply $P$ by the 0–1-incidence matrix for the 951 circuits of Proposition 5.2. That matrix has a 1 in row $i$ and column $j$ if circuit $j$ contains species $i$, and 0 otherwise. The product is a matrix of size $2 \times 951$. For $N \to \infty$, the $2 \times 951$ matrix has a finite entry in position $(i, j)$ precisely when the strategy $i$ can measure the circuit $j$. Minimizing over those finite cost entries selects the most cost-effective experiment to measure a circuit.

**Example 8.2** Suppose that none of the intermediate complexes $x_{13}, \ldots, x_{19}$ are measurable, and that we are able to measure only one phosphatase concentration ($x_4$ or $x_8$) in each experimental setup. A corresponding cost matrix might look like
Example 8.3 (Model rejection). Suppose that rate parameters are unknown, and that we have collected data for variables \( x_1, x_4, x_{14} \). The circuit polynomial is 
\[
k_1k_3x_1x_4 + (-k_2k_4 - k_5k_5)x_{14},
\]
which specializes to \( x_1x_4 - 2x_{14} \). If the evaluation of the positive quantity \( |x_1x_4 - 2x_{14}| \) lies above a threshold \( \epsilon \), then we can reject the model as not matching the data.

Every circuit polynomial of the matroid is a \textit{steady-state invariant}; depending on which experiment was performed, the collection of measured variables must contain some circuit. Even if one can measure all 19 species at steady state, it is not possible to recover all 31 kinetic rate constants, but we do have relationships that must be satisfied among parameters (MacLean et al. 2015).

Example 8.4 (Parameter Estimation). Suppose that rate parameters are unknown, and that we have collected data for \( x_6, x_{10}, x_{18} \). The corresponding circuit polynomial \( f_C \) is shown in Example 7.5. We know that the coefficients of \( f_C \) satisfy the constraint 
\[
y_2y_6 = y_3y_5.
\]
Suppose our experiments lead to the following 10 measurements for the vector \( (x_6, x_{10}, x_{18}) \):

\[
\begin{align*}
(0.390982, 4.83152, 6.08251), & (0.70659, 4.98107, 3.83617), \\
(1.4316, 4.30851, 12.5809), & (0.995583, 4.01222, 15), & (0.413817, 4.08114, 14.902), \\
(0.232206, 3.38274, 23.3162), & (0.219045, 5.06008, 3.67175), & (0.704106, 3.52804, 21.1037), \\
(0.648732, 3.6505, 19.7008) &
\end{align*}
\]

The data lead us to the following function to optimize in (15):

\[
\begin{align*}
57.2345y_1^2 + 376.181y_1y_2 + 801.672y_2^2 - 27.5625y_1y_3 & - 96.4429y_2y_3 \\
+ 3.36521y_3^2 + 179.49y_1y_4 + 564.034y_2y_4 & - 42.729y_3y_4 \\
+ 178.839y_4^2 + 564.034y_1y_5 & \\
+ 242.31y_2y_5 - 144.7y_3y_5 & + 1054.49y_4y_5 + 2263.2y_5^2 - 42.729y_1y_6 \\
- 144.7y_2y_6 + 10.339y_3y_6 & - 83.8072y_4y_6 - 269.749y_5y_6 + 10y_6^2
\end{align*}
\]
The global minimum of this quadratic form on the codimension 2 variety (16) has coordinates
\[ y_1 = 0.183472, \ y_2 = 0.152416, \ y_3 = 0.959232, \ y_4 = 0.038042, \]
\[ y_5 = 0.00335267, \ y_6 = 0.211. \]

Given these values, one now has three degrees of freedom in estimating the nine parameters \( k_i \) that appear in the circuit polynomial \( f_C \). The other 10 coordinates of \( k \) are unspecified.

9 Conclusion

Focusing on the Wnt shuttle model, we have looked at four common questions asked by experimentalists in systems biology. Each of these questions inspires a range of algebraic, combinatorial, and geometric problems, a small sample of which we have explored in this paper. Many current methods from dynamical systems to analyze models are limited to small models with a low-dimensional parameter space, or requires knowledge of parameter values and numerical studies. This often prevents the analysis of medium to large models. As we have seen, properties of the underlying algebraic varieties can give meaningful insights into the models. In particular, algebraic and geometric methods, as shown here and described in previous literature in this vein, can offer ways to circumvent current limitations on traditional tools. Additionally, there is new mathematics to be gained here. Systems biology has the potential to lead to considerable new advances in real algebraic geometry.

We have answered questions for only one single model; however, biochemical reaction networks describe a structured class of polynomial dynamical systems. The hope for the future would be to exploit this structure to make general statements regarding such systems, which requires novel approaches in nonlinear algebra, polyhedral geometry, and matroid theory.

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