

# Automated Conflict Detection Between Medical Care Pathways

Philip Weber, Bosco Filho, Mark Lee, Ian Litchfield, Ruth Backman

University of Birmingham, UK

School of Computer Science | Institute of Applied Health Research

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# What is the Problem?

In England > 15 million people have a long-term health condition.

Around 70% of the money spent on health and social care.

In the UK 2.9 million with three or more (Multi-morbidity) – by 2018.

- aging, smoking, diet, inactivity, ...
- cancer, heart disease, lung disease, diabetes, depression, ...

Complex processes for treatment (people, factors, clinical evidence ...)

UK National Institute for Care Excellence (NICE):

- Clinical Guidelines → Care Pathways.

A care pathway is essentially a process for treatment of a disease.

# Care Guidelines and Pathways

Secure | <https://pathways.nice.org.uk/pathways/chronic-obstructive-pul> ☆ ABP

Menu **NICE** Sign in

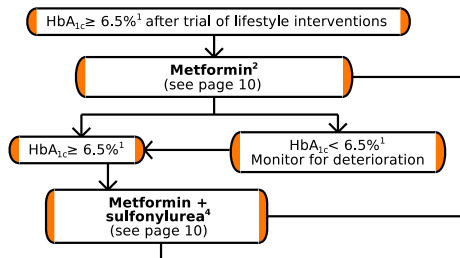
## COPD overview

```
graph TD; A[Person over 16 at risk of COPD] --> B[Diagnosis and assessment]; A --> C[Multidisciplinary team]; A --> D[See what NICE says on patient experience]; B --> E[Management]; B --> F[Managing exacerbations]; C --> E; C --> F; E --> G[Palliative care]; F --> G;
```

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# Two Problems

1. **Informal modelling** – potential for inconsistency,
2. Focus on **single conditions** – potential for conflict.



- **Implicit cycle** of retesting.
- What does **Metformin** conflict with?
- What does **HbA1c** interact with?

Our work:

- 1 define a **formal pathway model** to capture clinical pathways,
- 2 develop automated methods for **conflict detection**,
- 3 recommend minimal solutions for **conflict resolution**.

# 1. Modelling clinical guidelines

**Requirement:** formal modelling for analysis.

Many options (YAWL, Petri Nets, Computer Interpretable Guidelines, ...)

Business Process Model and Notation (BPMN):

- well-known *de facto* business process modelling language,
- increasingly prevalent for modelling clinical pathways,
- graphical, intuitive, flexible, 'subset-able'.

But

- no formal semantics,
- models can be unstructured,
- especially the semantics w.r.t. data are unspecified.

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# Data in Care Pathways

Routing may be **dependent on** and **modify** data, e.g.

- Is patient already **taking medication**  $m$ ?
- If test  $X > \text{value } v$ , refer for treatment  $X$  else retest in  $M$  months.
- If patient age  $A > a$  prescribe drug  $x$  else  $y$ .
- Record the fact of **prescription of drug**  $Z$ .

BPMN has the **Data Object** element, but

- **semantics open to interpretation,**
- **decoupled from the control-flow.**

Literature covering formalisation of

- BPMN integration with data objects, e.g. [Meyer et al., 2013];
- interaction between processes and databases [Sun et al., 2014];
- **seems more complex than we need;**
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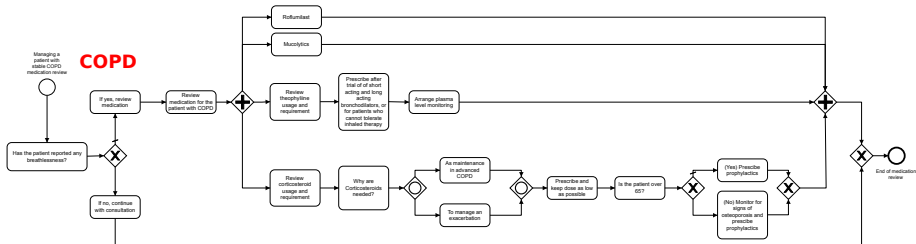
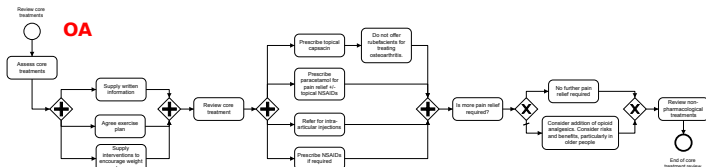
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# BPMN for Two Pathway Fragments

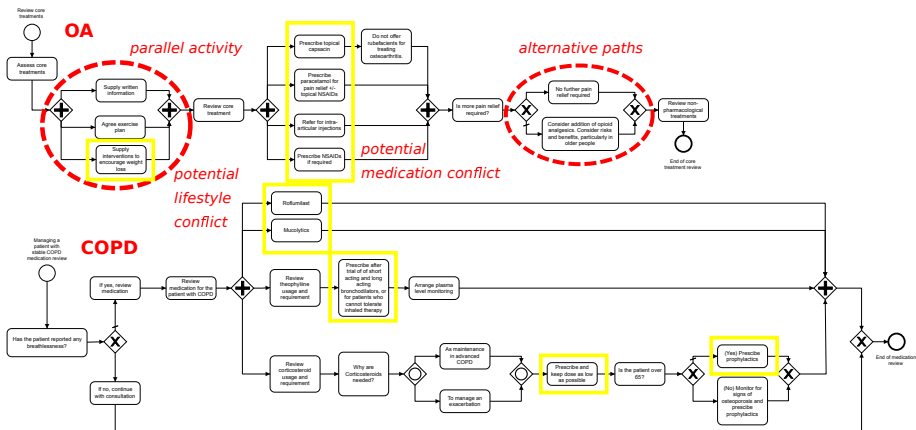


Fragments of doctors' appointments for review of

**OA** : Osteoarthritis.

**COPD** : Chronic Obstructive Pulmonary Disease (Lung Disease).

# BPMN for Two Pathway Fragments



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OA : Osteoarthritis.

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Formal execution semantics ...

- subset of BPMN notation – can be expanded,
- **formal semantics** of execution,
- based on Workflow Graphs ([Vanhatalo et al., 2007] formalism),

... and integration with data,

- semantics of dependence on – and modification of – **data**,
- based on Coloured Petri Nets.

# Workflow Graphs

Effectively a **subset of BPMN** allowing the main control-flow patterns and imposing some structure on the model [Vanhatalo et al., 2007].

- $G = (N, E)$ , nodes  $N$ , edges  $E$ , such that  $E \subseteq N \times N$ ,
- $N \in \{\text{START, STOP, ACTIVITY, FORK, JOIN, DECISION, MERGE}\}$ ,
- $G$  is **well-formed** by definition,
- allowing for **atomic** activities, parallel and alternative behaviours.

Semantics of  $G$  is a **token game** (cf Petri Nets).

- **State**  $s$  of  $G$  is a mapping  $s : E \rightarrow \mathbb{N}$  assigning **tokens** to edges in  $E$ .
- $s(e) = k \iff$  in state  $s$ , edge  $e$  carries  $k \in \mathbb{N}$  tokens.
- execution of node  $n$  changes the state to  $s' : s \xrightarrow{n} s'$ .

This says nothing about data.

## Flexible approach to model data:

- Fixed set of  $d$  variables  $X = \{x_1, \dots, x_d\}$  of types  $\mathcal{T}(x_i) \in \{\mathcal{T}_1, \dots, \mathcal{T}_m\}$ ,
- valuations  $V = (\nu_1, \dots, \nu_d)$  assigned to  $X$  as the process executes.
  - $V$  assigned to token ('colour').
- Activity may be guarded by pre- and post-conditions  $c(\cdot)$ :
  - $c(\cdot)$  is a first-order logic formula over  $X$ ,
  - $c(\cdot) \models V$  if the valuation  $V$  satisfies  $c(\cdot)$ ,
  - e.g.  $pre(a) := c(x_1, \dots, x_d) \triangleq (x_i > 55)$ .
- Activity may carry out data modifications:
  - statement  $f(\cdot) : V \rightarrow V'$  over variables in  $X$ ,
  - e.g.  $x := x + 1$  or  $x := False$ ,
- data may require synchronisation – managing the control-flow.

**Implicit data** is referenced but not modified.

- e.g. database of drug-drug-disease interactions (Stockley / BNF).

## Extension of Workflow Graphs:

- $G = (N, l, E, X, pre, post, mod)$ ,
- $l : N \rightarrow \{\text{START, END, ACTIVITY, EXCLUSIVE, INCLUSIVE, PARALLEL}\}$ ,
- $X = \{X_1, \dots, X_d\}$ ,
- $\{pre, post\} : N \rightarrow C$ ,
- $mod : N \rightarrow D$  (database),
- allowing for **atomic** activities, parallel, exclusive or inclusive choice.

## Semantics defined in terms of before- and after- conditions and states,

- $m : E \rightarrow \{T_1, T_2, \dots\}$  is a **marking** describing the **state**,
- mapping each edge  $e \in E$  to coloured tokens,  $T_i = (t_i, V_i)$ ,
- execution modifies the **state**  $m \xrightarrow{n} m'$  and (perhaps) **valuation**  $V \rightarrow V'$ .



# BPMN+V: Execution Semantics

e.g. For an **ACTIVITY**  $a$  in a well-formed BPMN+V model:

- one input and one output sequence flow  $e_{in}, e_{out}$ .
- $a$  consumes  $T = (t, V)$  from  $e_{in}$  and returns  $T' = (t, V')$  on  $e_{out}$ ,
- if  $\exists T = (t, V) \in m(e_{in}) \mid pre(a) \models V$ , //  $V$  satisfies any pre-condition.
- then  $m \xrightarrow{a} m'$ , where

1.  $post(a) \models V'$ , and

2.  $m'(e) =$

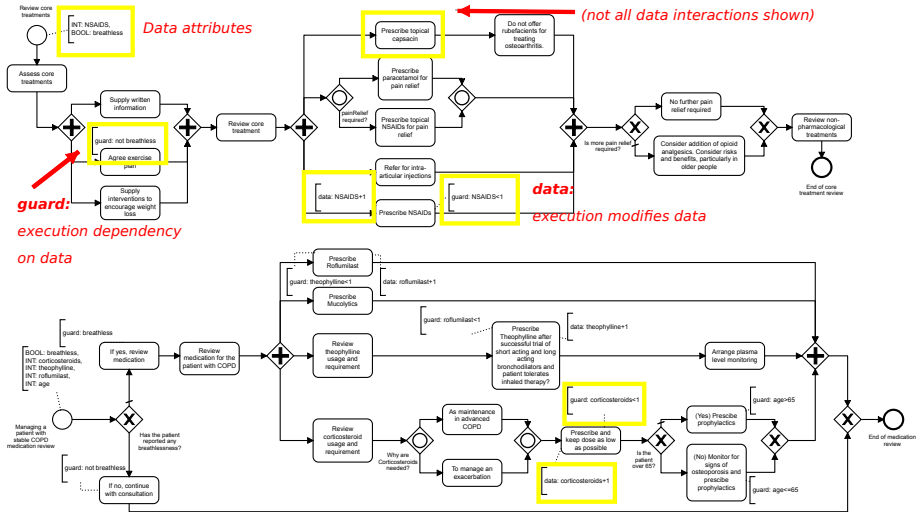
$$\begin{cases} m(e) \setminus \{T\} & \text{if } e = e_{in}, \quad // V' \text{ satisfies any post - condition.} \\ m(e) \cup \{T'\} & \text{if } e = e_{out}, \quad // \text{token is 'moved'}. \\ m(e) & \text{otherwise.} \end{cases}$$

e.g.  $pre(a) := \neg\text{NSAIDS}$ ,

$f(a) \triangleq \text{corticosteroids} := \text{corticosteroids} + 1$ .

– Similarly for all node types.

# BPMN+V: Data Annotation



## 2. Conflict Detection

# Conflict Detection

The **problem**: to identify conflicts between clinical care guidelines followed concurrently in treating patients with multiple morbidities.

Assume

- two BPMN+V models (care pathways)  $M_1, M_2$ ,
- interacting with database  $\mathcal{D}$ ,
- shared set of  $d$  variables  $X = X_1 \cup X_2$ ,
- set of  $k$  *constraints*  $C = \{C_1, \dots, C_k\}$ .

**Constraint**  $C_r$  is a logical formula over  $X$ , e.g.

- if  $x_i$  and  $x_j$  indicate prescription of two medications,
- which must not be taken together,
- then  $C_r(x_1, \dots, x_d) \triangleq \neg(x_i \wedge x_j)$ .

The **problem**:

*identify all pairs of execution paths through  $M_1, M_2$  which will modify the variables in  $X$  so that at least one of the  $C$  is violated.*

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# State Space Approach to Conflict Detection

Current process for conflict detection (evaluate BPMN+V):

- 1 Model clinical pathways in BPMN+V,
- 2 simple parallel composition,

Assumes patient (potentially) starts both pathways concurrently.  
Assumes no common activities.

Future: intelligent model composition.

# State Space Approach to Conflict Detection

Current process for conflict detection (evaluate BPMN+V):

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- 3 annotate with constraints (potential conflicts, e.g. meds. dependencies),
- 4 identify data combinations for which to explore the model,

- Identify  $d$  variables  $X$  involved in conditions,
- values  $V$  checked/assigned.
- Create a 'covering set' of  $2^d$  initial data settings for validation:
  - e.g.  $\{y > 1, y \leq 1\}$  for a condition  $y > 1$ .

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- 4 identify data combinations for which to explore the model,
- 5 state space exploration for each data combination (via CPN),

- Transform to CPN (take advantage of existing methods).
- Construct  $2^d$  reachability graphs  $R_i$  (explore state space).
- Conflicting Activities indicated by non-final dead markings linked by common variables.
- Repeat for individual models (detect data-related inconsistencies)
- and composed models (detect conflicts).

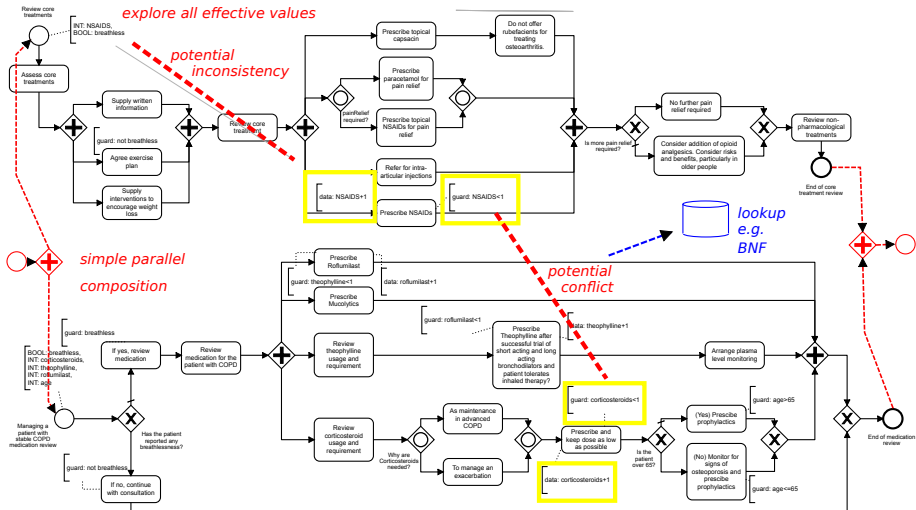


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- 5 state space exploration for each data combination (via CPN),
- 6 identify 'non-final dead markings',
- 7 visualise and interpret the conflicting activities and data combinations.

# Composed OA and COPD Model

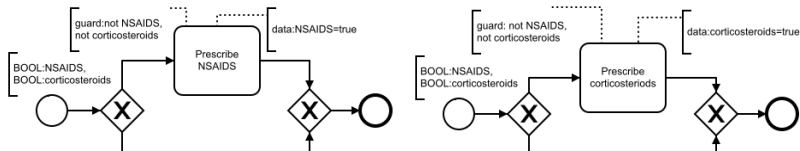


(Possibly invalid assumption that the patient starts following both models at the same time.)

# 3. Evaluation

## 3-stage evaluation:

### 1 Artificial process fragments, e.g.



### 2 Randomly-generated models,

- block-structured expansion,
- controlled block probabilities and number of conflicts.

### 3 Running example – Osteoarthritis (OA) and COPD pathways,

- 14 activities,
- 3 variables,
- up to 11,000 states in the composed model.

# Report and Visualise Results

Model	Activity	Data	Initial Data	Conflict Model	Conflict Activity	Conflict Data
OA	Agree exercise plan [-breathless]	breathless=True	breathless=True			
OA	Prescribe NSAIDs	NSAIDs=1	NSAIDs=1			
COPD	Prescribe cortico. and keep ...	cortico.=1	cortico.=1			
COPD	Prescribe roflumilast [th.lline < 1]	th.lline=1	th.lline=0	COPD	Prescribe th.lline after ...	rofl.
COPD	Prescribe th.lline ... [rofl. < 1]	rofl.=1	rofl.=0	COPD	Prescribe roflumilast	th.lline
OA	Prescribe NSAIDs	cortico.=1	cortico.=0	COPD	Prescribe and keep ...	NSAIDs
COPD	Prescribe cortico. and keep ...	NSAIDs=1	NSAIDs=0	OA	Prescribe NSAIDs	cortico.

data inconsistencies for OA (top),

- cannot proceed with exercise plan if patient presents with breathlessness,
- must not over-prescribe NSAIDs.

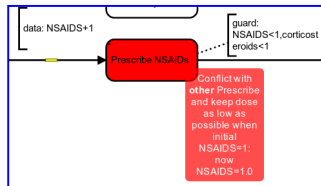
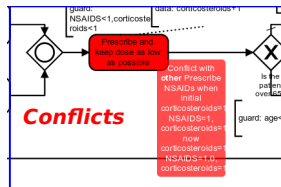
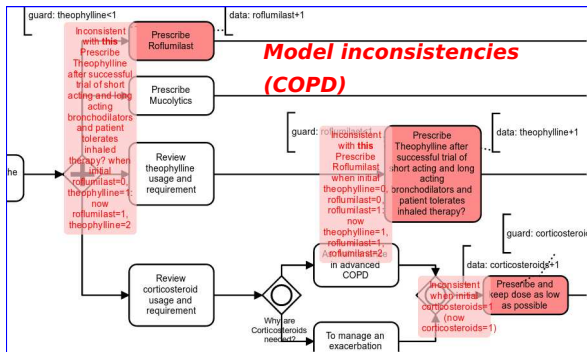
COPD (centre),

- must not over-prescribe corticosteroids,
- check/prescribe Roflumilast and Theophylline are mutually exclusive,
- but parallel structure allows both to be executed.

conflicts between the models (bottom).

- corticosteroids and NSAIDs are mutually exclusive across both pathways.

# Report and Visualise Results



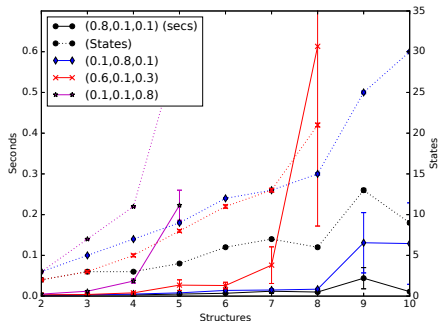
“**Inconsistent with this** {activity} when {data settings}”.

“**Conflict with other** {activity} when {data settings}”.

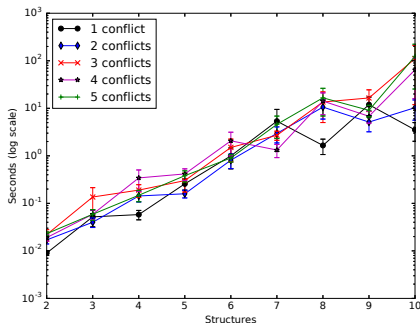
Annotation using <http://bpmn.io/> and/or Camunda Modeller.

# Conflict Detection: Performance of State Space Method

Bespoke BPMN+V and CPN implementation.  
Averages over 30 randomly generated models.



(a) Varying complexity models [ $p(seq)$ ,  $p(xor)$ ,  $p(pll)$ ].



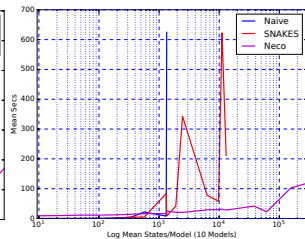
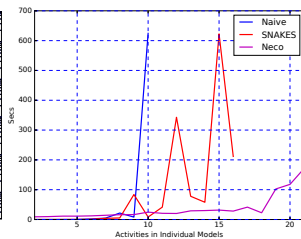
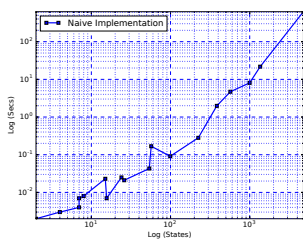
(b) Composed models, varying numbers of conflicts.

- (a) #states vs time (seconds) to run the conflict detection process,  
– models generated with varying probability of sequence, alternate or parallelism.
- (b) increasing #conflicts, in models with low probability of concurrent activity.

# Conflict Detection: Performance of State Space Method

Bespoke vs **SNAKES** [1] vs **Neco** [2].

Averages over 30 randomly generated models.



[1] **SNAKES**: <https://snakes.ibisc.univ-evry.fr/>

"SNAKES is a Python library that provides all the necessary to define and execute many sorts of Petri nets",

[2] **Neco**: <https://github.com/Lvyn/neco-net-compiler>

"Neco . . . takes a Petri net and builds a **library** that has all the primitives to explore the state space . . . optimised in many ways."



## Modelling

- user interface, software tool and case study,
- data integration with sources of data and conflict.

## Conflict Detection

- model **composition** – adequacy of simplistic approach,
- conflict detection using **logical specification and constraint solvers**,
- scheduling constraints.

## Conflict Resolution

- recommendation of minimal changes for conflict resolution,
- e.g. bypass activities,
- e.g. reschedule.

# Thank you

*Phil Weber*

`http://www.birmingham.ac.uk/mitcon/`

`http://www.cs.bham.ac.uk/~weberpy/  
p.weber.1@cs.bham.ac.uk`