





Structural Causal Models are Credal Networks?

joint work with Marco Zaffalon (IDSIA) and Rafael Cabañas (UAL)

Alessandro Antonucci (<u>alessandro@idsia.ch</u>) Senior Lecturer Researcher IDSIA USI-SUPSI

SIPTA Seminar, Virtual Event, Nov 28, 2023



slides



Istituto Dalle Molle di Studi sull'Intelligenza Artificiale (IDSIA)





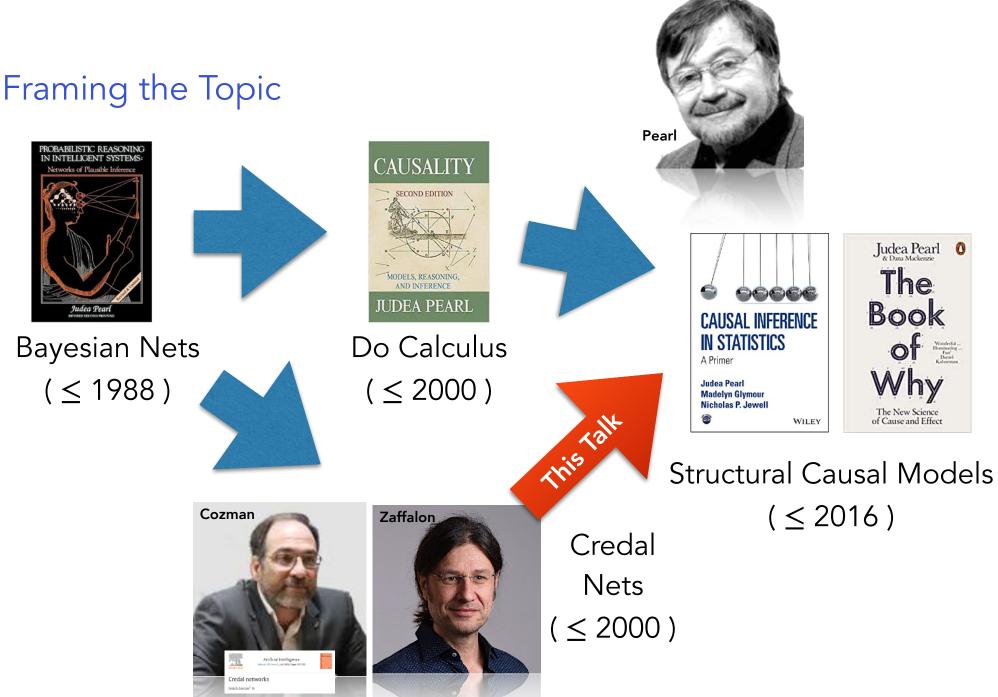


- World-class Research Institute on AI founded in 1988 in Lugano
- Affiliated with both University of Lugano (USI) and University of Applied Sciences of Southern Switzerland (SUPSI)
- Staff ~100 people + 50 PhD



Angelo Dalle Molle (1908 - 2002)





(Science >) AI > Deep Learning

🔊 Andrej Karpathy blog

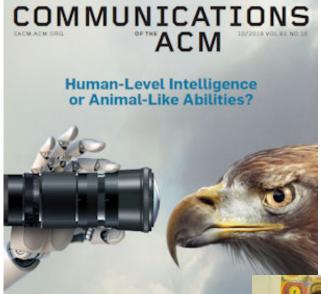
The Unreasonable Effectiveness of Recurrent Neural Networks May 21, 2015

RESEARCH ARTICLE | BIOLOGICAL SCIENCES |

f 🌶 in 🖾 🤮 The unreasonable effectiveness of deep learning in artificial intelligence

Terrence J. Sejnowski 💷 🗠 Authors Info & Affiliations

Edited by David L. Donoho, Stanford University, Stanford, CA, and approved November 22, 2019 (received for review September 17, 2019) January 28, 2020 117 (48) 30033-30038 https://doi.org/10.1073/pnas.1907373117



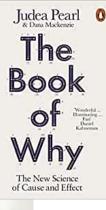
Computing within Limits Transient Electronics Take Shape Q&A with Dina Katabi Formally Verified Software in the Real World



"Deep learning has instead given us machines with truly impressive abilities but no intelligence.

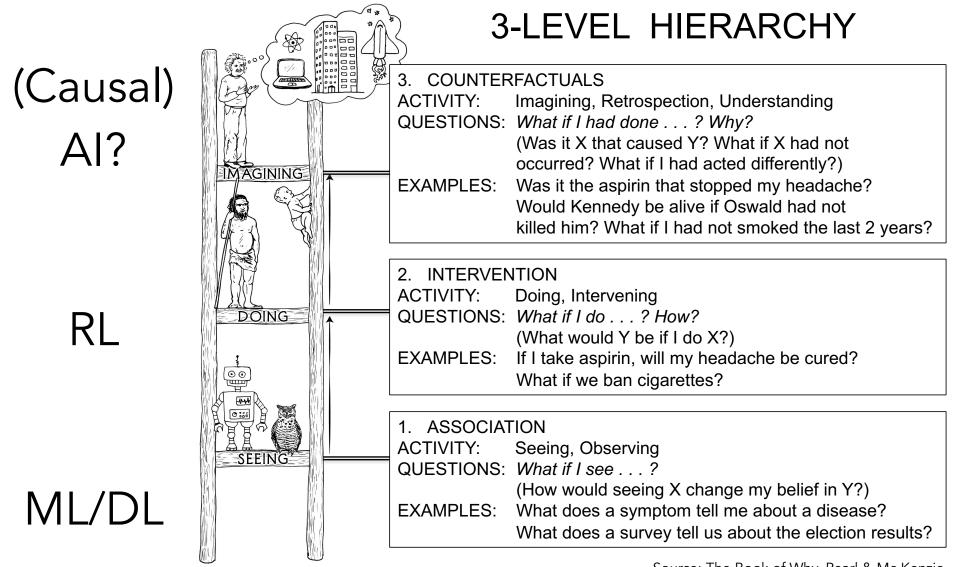
The difference is profound and lies in the **absence of a** model of reality."







Pearl's Ladder of Causation and the Need for a Causal Al



Source: The Book of Why, Pearl & Mc Kenzie



Pearl's Ladder of Causation and the Need for a Causal Al credal nets (Cau as a tool to climb the top s? (= counterfactuals) of the ladder ML/D EXAMPLES: What does a symptom tell me about a disease? What does a survey tell us about the election results?

Source: The Book of Why, Pearl & Mc Kenzie



- Manifest **endogenous** variable X
- Observations \mathscr{D} available
- From \mathscr{D} statistical learning of P(X)



Boolean XP(X = 0) = p



- Manifest **endogenous** variable X
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- A latent **exogenous** variable U

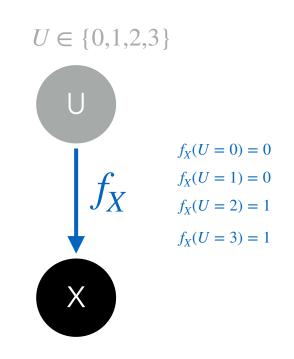




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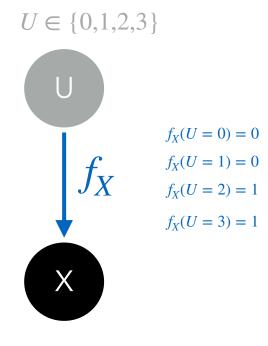
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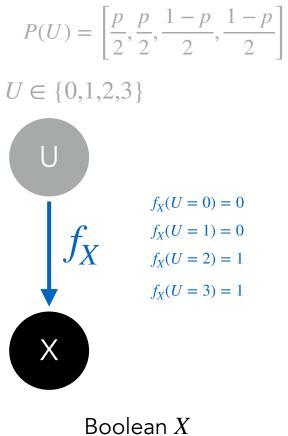
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- A P(U) giving P(X)? More than one!



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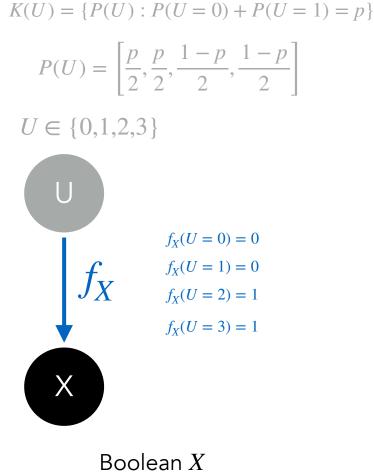


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- Causal inference to be based on the credal set K(U) compatible with P(X)

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P(X=0) = p



- Manifest **endogenous** variable X
- Observations \mathcal{D} available

From \mathcal{D} statistical learning of P(X)

 $K(U) = \{P(U) : P(U = 0) + P(U = 1) = p\}$

SUPSI

$$P(U) = \left[\frac{p}{2}, \frac{p}{2}, \frac{1-p}{2}, \frac{1-p}{2}\right]$$

 $U \in \{0, 1, 2, 3\}$

A lat This is a (minimalistic) State throu structural causal model f_X su

U

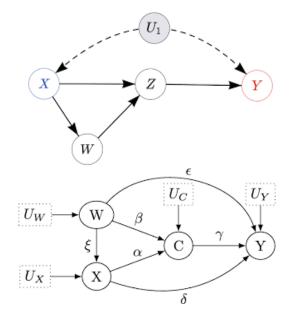
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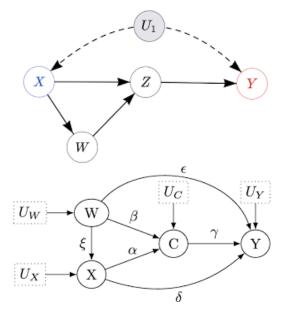


- $\mathbf{X} := (X_1, \dots, X_n)$ (endogenous variables)
- $\mathbf{U} := (U_1, \dots, U_m)$ (exogenous variables)
- Directed graph \mathcal{G} assumed to be semi-Markovian = root in **U**, non-root in **X**
- Equation $X = f_X(Pa_X)$ for each $X \in \mathbf{X}$
- Marginal P(U) for $U \in \mathbf{U}$ (assessed if possible)



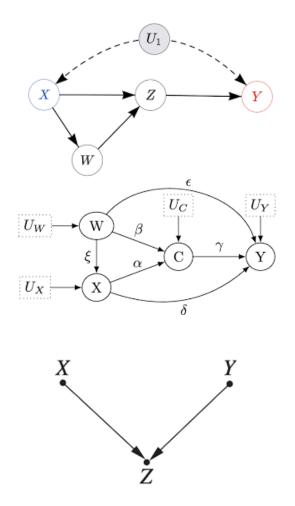


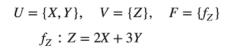
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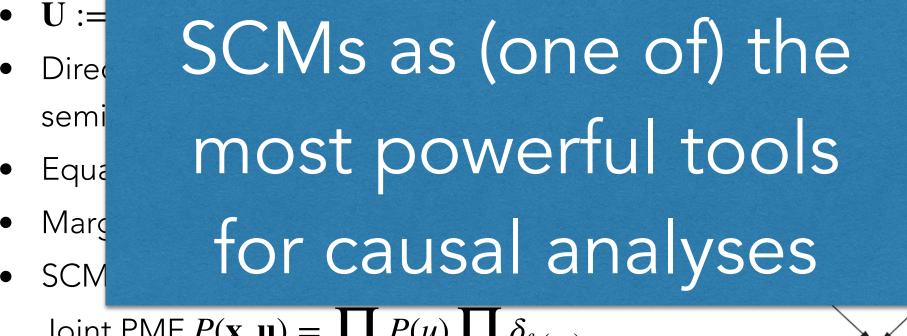
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 $U = \{X, Y\}, \quad V = \{Z\}, \quad F = \{f_Z\}$ $f_Z : Z = 2X + 3Y$

 U_Y



Headache Example (Staying on the First Rung)

- You take aspirin (X = 1) and headache vanishes (Y = 1)
- Probability that this has been due to aspirin?
- Observational data ${\mathscr D}$ about the two variables available
- From \mathscr{D} , P(Y=0 | X=0) = 0.5 > P(Y=0 | X=1) = 0.1

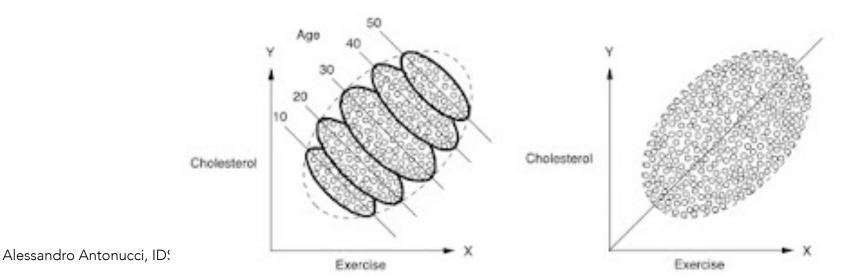
$$X \bullet \longrightarrow \bullet Y$$

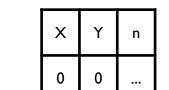
x	Y	n
0	0	
0	I	
I	0	
I	I	



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- Not genuine causal analysis: adding further covariates might give contradictory results (Simpson's paradox)





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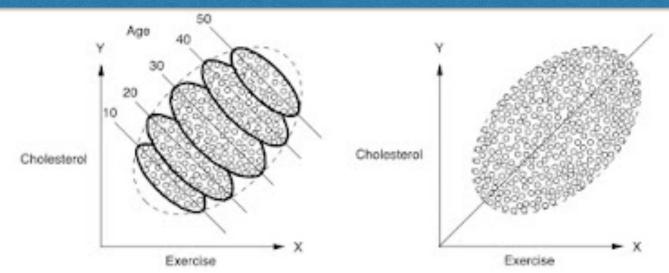
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X



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 Time to climb up
 the ladder



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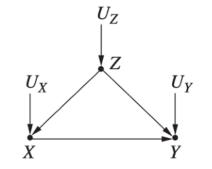
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n



Take the Aspirin! (Interventions = Second Rung)

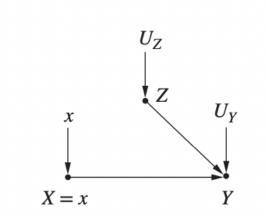
- Gender Z as an additional (endogenous) variable
- Markovian \mathscr{G} (one exo parent for each endo)
- Force people to take aspirin = **intervention** do(X = 1)





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 U_{Y}

 U_Z

Ζ

 U_X

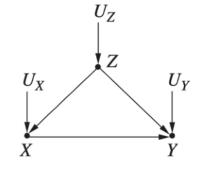


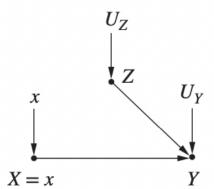
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- Pearl's **do calculus** allows to reduce interventional queries to observational ones (solved by BN inference)

E.g., backdoor $P(y \text{ do}(X = x)) = \sum P(y x, z) \cdot P(z)$

• Do calculus only needs ${\mathscr G}$ (and not the SCM)!







Identifiability of Causal Queries

- Do calculus reduces interventional to observational queries by exploiting d-separation in SCMs
- Sound and complete (graph-theoretic) algorithm
 + inference in the empirical joint PMF
- Alternatively: surgery and inference in the SCM ...

v a	ind analy	ze causal	diagrams
vironment for creating, editing, and o known as directed acyclic graphs or e focus is on the use of causal diagrams studies in epidemiology and other ormation, see the "learn" page.			Versions The following versions of DAG available: • Development version Recent development ana
d	Learn	Code	May contain new features, could also contain new bu • Experimental version
	Learn more about DAGs and DAGitty.	The R package "depitty" is available on CRNN or	Maat recort development snapshot, May not over w 3.0; Released 2019-01-00 2.2; Released 2019-01-02 2.2; Released 2014-13-3; 2.1; Released 2014-02-00 2.0; Released 2014-02-00 1.1; Released 2017-01-12

DAGitty — drav

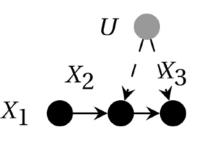


Identifiability of Causal Queries

- Do calculus reduces interventional to observational queries by exploiting d-separation in SCMs
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 + inference in the empirical joint PMF
- Alternatively: surgery and inference in the SCM ...
- Not all queries can be computed by do calculus.
 If not we call the query **unidentifiable**
- Emerging idea: unidentifiable queries are only partially identifiable (bounds can be estimated!)
- Recent works in this field by various groups: sampling (Bareinboim), poly programming (Shpitser)

Gitty — draw and analyze causal diagrams							
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 $P(x_3 \ \operatorname{do}(x_2) \in [l, u]$





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Back to Headache (Moving to the Third Rung)

- What if I had not taken the aspirin, would have headache stayed?
- An intervention contrasting the current observation ...
- This is a **counterfactual** query $P(Y_{X=0} = 0 \ X = 1, Y = 1)^U$ \bullet (called probability of necessity, PN, sub denote do)

SUPSI

X



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- An intervention contrasting the current observation ...
- This is a **counterfactual** query $P(Y_{X=0} = 0 \ X = 1, Y = 1)^U$ (called probability of necessity, PN, sub denote do)
- We need the complete SCM: $\mathcal{G} + \{f_X\}_{X \in \mathbf{X}} + \{P(U)\}_{U \in \mathbf{U}}$
- With complete SCM, an augmented model called twin network with duplicated endogenous variables is used X' for counterfactual analysis after surgery
- (Non-trivial) counterfactuals are unidentifiable!

SUPSI

X





To Compute Counterfactuals ...

- We need a fully specified SCM, i.e.,
 - 1. Graph ${\mathscr G}$ over $({\mathbf X}, {\mathbf U})$
 - (often available by domain expert or Markovian assumption)
 - 2. Endogenous equations $\{f_X\}_{X \in \mathbf{X}}$ (available or obtained by complete enumeration)
 - 3. Exogenous marginals $\{P(U)\}_{U \in \mathbf{U}}$ (rarely available)



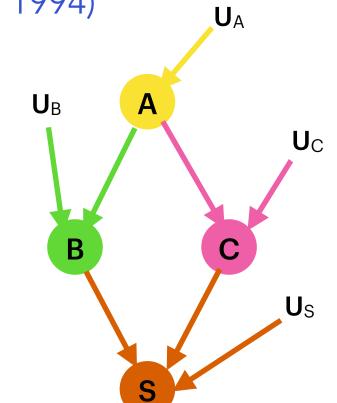
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- Latent $P(\mathbf{U}) = \prod P(U)$ unavailable? We have data \mathcal{D} about \mathbf{X}
- Compute counterfactual = Compute $\{P(U)\}_{U \in U}$ from \mathcal{D}
- Not a new problem: LP approach for special cases already in Balke and Pearl (1994), but do-calculus reduced attention to CFs



Causal Analysis at the Party (Balke & Pearl 1994)

Ann sometimes goes to parties Bob is not a party guy, but he likes Ann and he might be there Carl broke up with Ann, he tries to avoid Ann, but he likes parties Carl and Bob hate each other, they might have a Scuffle if both at the party



besides such knowledge assume we have observations \mathcal{D} corresponding to a joint mass function P(A, B, C, S)(e.g., in the form of a BN)



Causal Analysis at the Party (Balke & Pearl 1994)

Ann sometimes goes CAUSAL GOSSIP UB A Bob is not a party guy, INTERVENTIONAL COUNTERFACTUAL

and he might be there Carl Ann must not be he trie at the party, or Bob would be there Carl and Bob by by bother, they might have a Scuffle if $boP(B^{th}do(\overline{a})) = ?$



"If Bob were at the party, Us then Bob and Carl would surely Scuffle"

besides such knop $S_{b} = P(S_{b} = b) = ?$ we have observations \mathcal{D} corresponding

a (fully specified) SCM can answer these questions

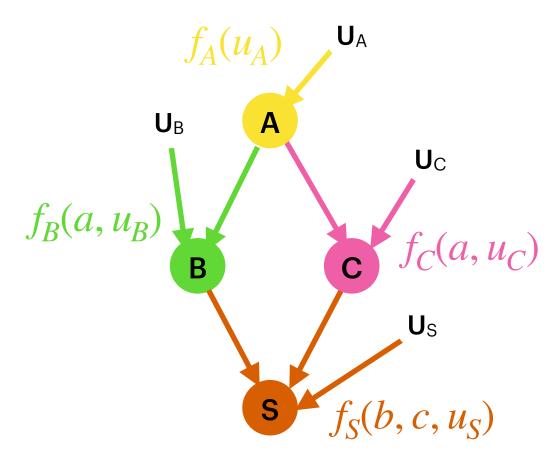
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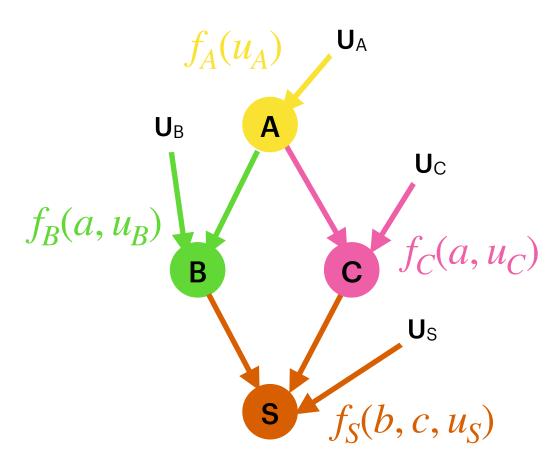


• Find the exogenous marginals? $P(U_A)P(U_B)P(U_C)P(U_S)$





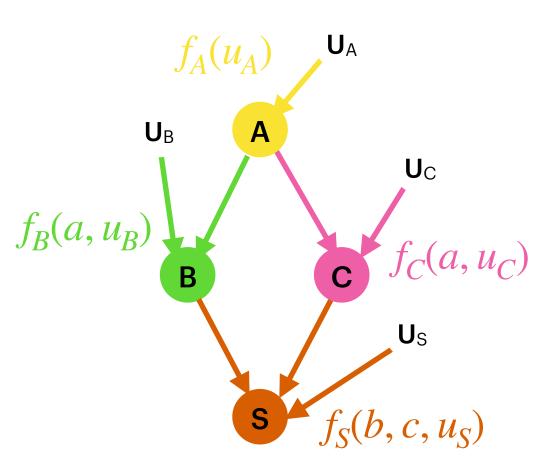
- Find the exogenous marginals? $P(U_A)P(U_B)P(U_C)P(U_S)$
- Endogenous (= with D) consistency



$$\sum_{u_A, u_B, u_C, u_D} \left[p(u_A) \cdot \delta_{a, f_A(u_A)} \cdot p(u_B) \cdot \delta_{b, f_B(a, u_B)} \cdot p(u_C) \cdot \delta_{c, f_C(a, u_C)} \cdot p(u_S) \cdot \delta_{s, f_S(b, c, u_S)} \right] = \tilde{p}(a, b, c, s)$$



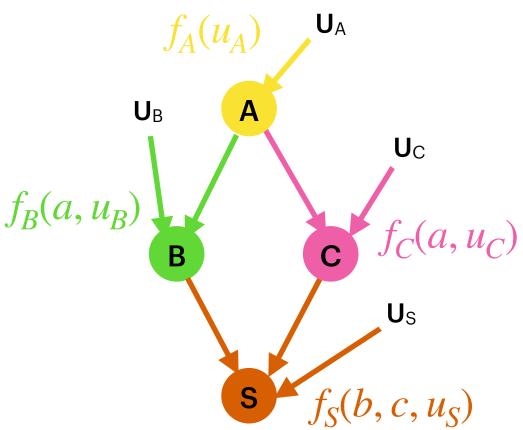
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- This induces global non-linear (so-called Verma) constraints



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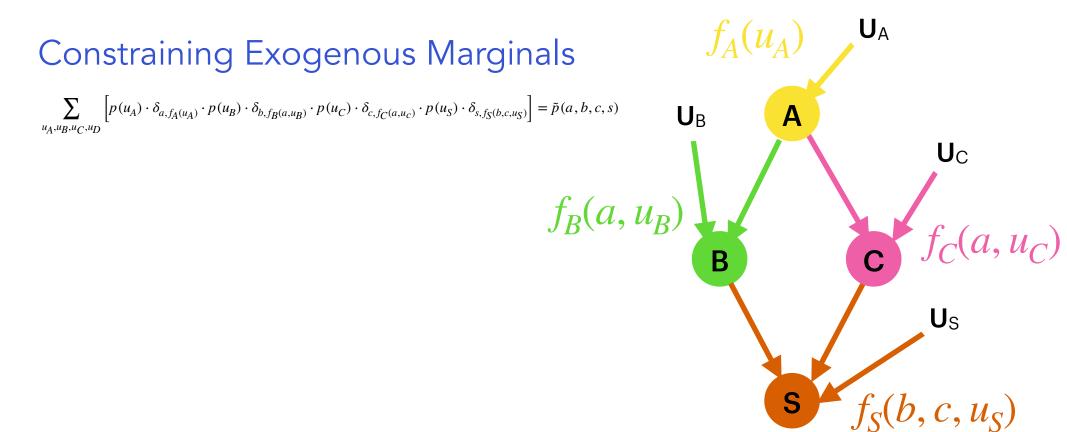
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- Constraints became local and linear ones by marginalisation and conditioning (Zaffalon et al., 2020)



$$\sum_{u_A, u_B, u_C, u_D} \left[p(u_A) \cdot \delta_{a, f_A(u_A)} \cdot p(u_B) \cdot \delta_{b, f_B(a, u_B)} \cdot p(u_C) \cdot \delta_{c, f_C(a, u_c)} \cdot p(u_S) \cdot \delta_{s, f_S(b, c, u_S)} \right] = \tilde{p}(a, b, c, s)$$

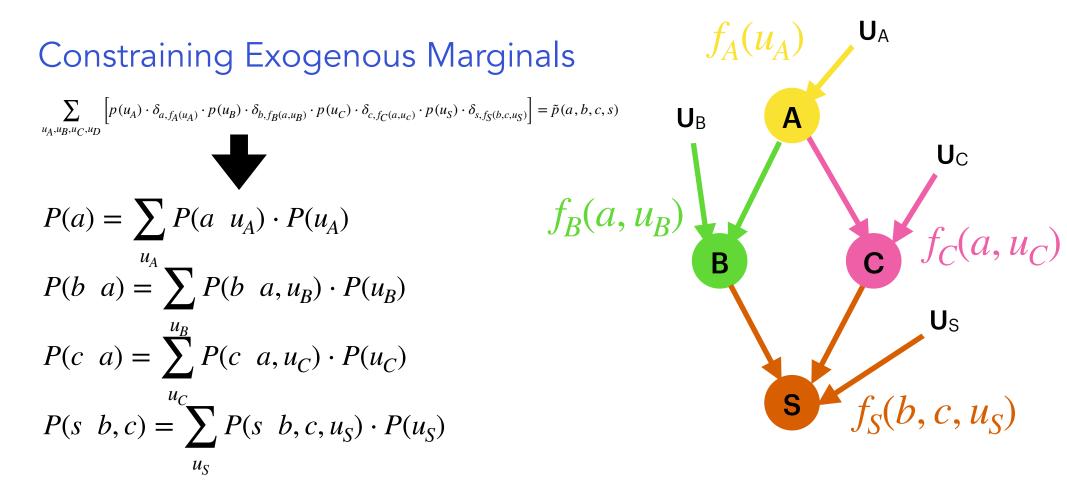




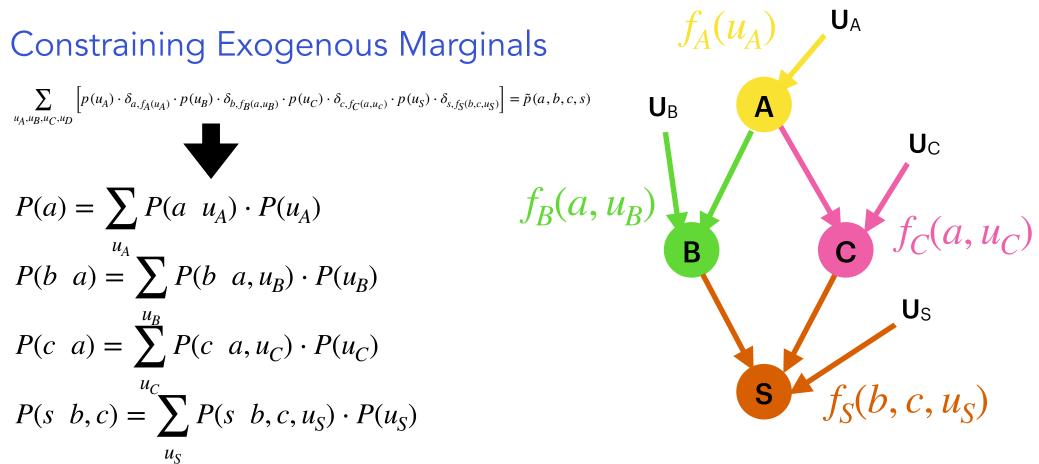






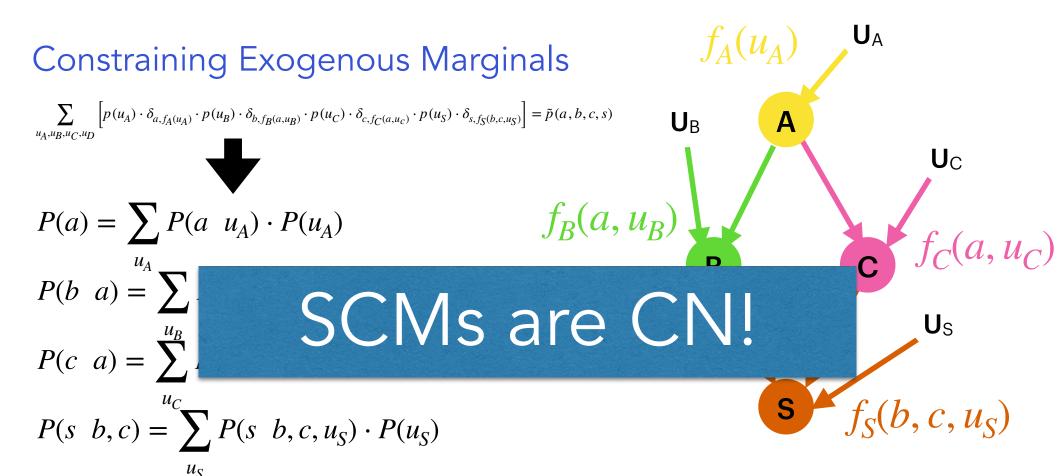






- Linear constraints on marginal exogenous probabilities leading to the credal sets specification $K(U_A)$, $K(U_B)$, $K(U_C)$, $K(U_S)$
- Structural equations (= endogenous CPTS) remain unaffected

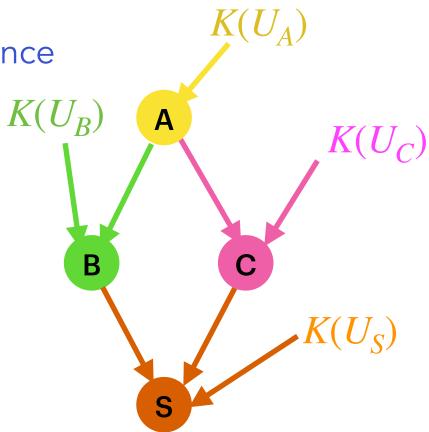




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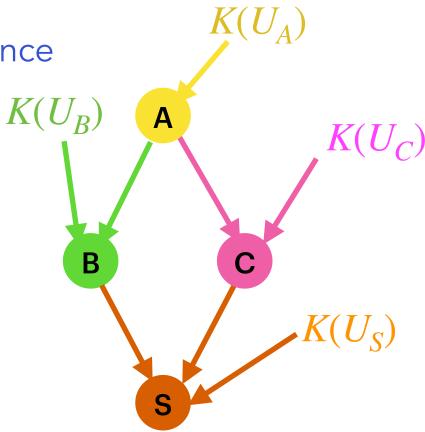


• Consistent SCMs as a single CN



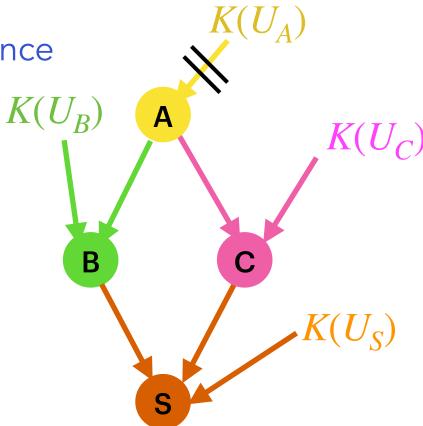


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- CN algs to compute bounds!



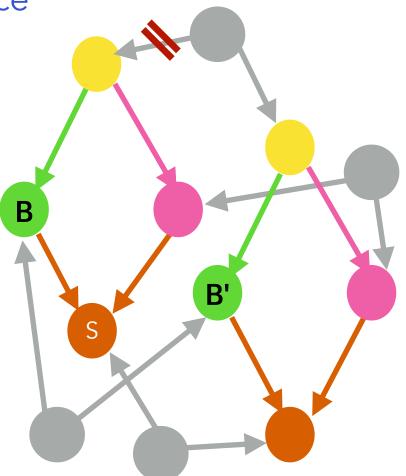


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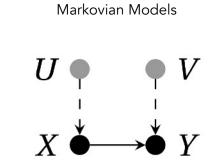
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- Interventions are straightforward $P(B \ do(\overline{a})) \in [\underline{P}'(B \ \overline{a}), \overline{P}'(B \ \overline{a})]$
- Counterfactuals require twin nets $P(S_b \ \overline{b}) \in [\underline{P}(S \ b, \overline{b}'), \overline{P}(S \ b, \overline{b}')]$
- Identifiable? $\underline{P} = \overline{P}$

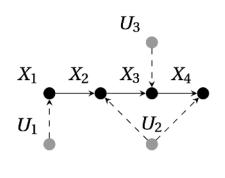




Markovian and Quasi-Markovian SCMs as CNs

1: f	for $X \in \mathbf{X}$ do		
2:	$U \leftarrow \operatorname{Pa}(X) \cap \boldsymbol{U}$	// U as the unique exogenous	parent of X
3:	$\underline{\operatorname{Pa}}(X) \leftarrow \operatorname{Pa}(X) \setminus \{U\}$	// Endogenous p	arents of X
4:	if $\underline{Pa}(X) = \emptyset$ then		
5:	$K(U) \leftarrow \{P'(U) : \sum_{u \in f_v^{-1}} P'(u) = 1$	$\tilde{P}(x)$, $\forall x \in \Omega_X$ }	// Eq. (4)
6:	else		
7:	$K(U) \leftarrow \{P'(U) : \sum_{u \in f_{X \operatorname{ing}(X)}^{-1}(x)} P'(u)\}$	$(u) = \tilde{P}(x \underline{pa}(X)), \forall x \in \Omega_X, \forall \underline{pa}(X) \in \Omega_{\underline{Pa}_X}$	// Eq. (6)
8:	end if		
9: e	end for		





Algo	Algorithm 2 Given an SCM <i>M</i> and a PMF $\tilde{P}(\mathbf{X})$, return CSs $\{K(U)\}_{U \in \mathbf{U}}$						
1: f	or $U \in \boldsymbol{U}$ do						
2:	$\{X_U^k\}_{k=1}^{n_U} \leftarrow \operatorname{Sort}[X \in X : U \in \operatorname{Pa}(X)]$	// Children of U in topological order					
3:	$\gamma \leftarrow \phi$						
4:	for $(x_U^1, \ldots, x_U^{n_U}) \in \times_{k=1}^{n_U} \Omega_{\mathbf{X}_U^k}$ do						
5:	for $(\underline{\text{pa}}(X_U^1), \dots, \underline{\text{pa}}(X_U^{n_U})) \in \times_{k=1}^{n_U} \Omega_{\underline{\text{pa}}}$	(X_{II}^k) do					
6:	$\Omega'_U \leftarrow \bigcap_{k=1}^{n_U} f_{X^k_U \underline{\mathrm{pa}}(X^k_U)}^{-1}(x^k_U)$						
7:	$\gamma \leftarrow \gamma \cup \left\{ \sum_{u \in \Omega'_U} P(u) = \prod_{k=1}^{n_U} \tilde{P}(x_U^k) \right\}$	$ x_U^1,\ldots,x_U^{k-1},\underline{\mathrm{pa}}(X_U^1)),\ldots,\underline{\mathrm{pa}}(X_U^k))\Big\}$					
8:	end for						
9:	end for						
10:	$K(U) \leftarrow \{P(U): \gamma\}$	// CS by linear constraints on $P(U)$					
11: e	and for						

Quasi-Markovian Models







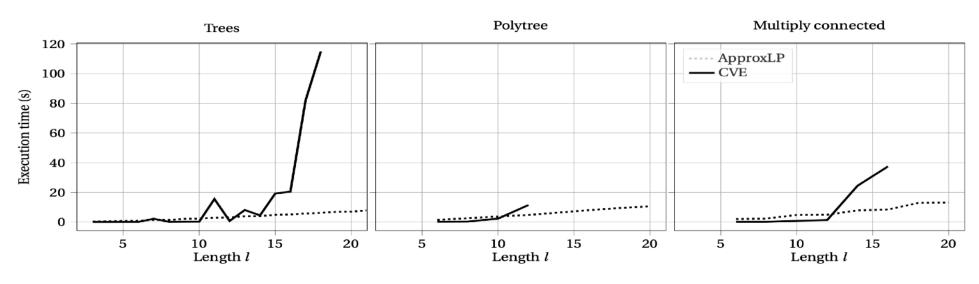
Software and Experiments Crede Models Agentine Java library for CNs







Software and Experiments Create Accelerations Lava library for CNs Control of Create Acceleration Lava Library for CNs Create Acceleration Lava Library for CNs

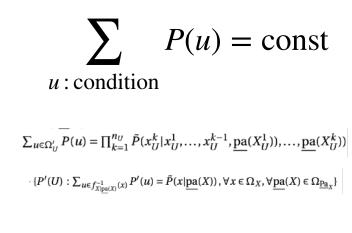


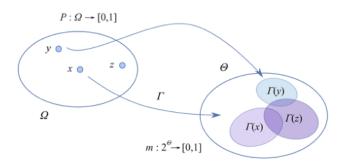
Exact inference by credal variable elimination only for small models ApproxLP (Antonucci et al., 2014) allows to process larger models RMSE always <0.7%



Intermezzo: Belief Functions (as Credal Sets)

- Linear constraints for CN induced by SCM have a peculiar form
- These are CS corresponding to **belief functions** (Dempster '68, Shafer '76)
- Class of generalised probabilistic models
- PMF distributes mass over the singletons, BF over (poss. overlapping) sets
- Dempster's multi-valued mapping, in SCMs $\mathbf{U} = f^{-1}(\mathbf{X})$, BF(\mathbf{U}) := $f^{-1}[P(\mathbf{X})]$
- Dedicated conditioning/combination rules



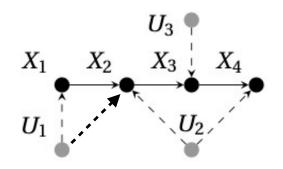


Credits: Fabio Cuzzolin



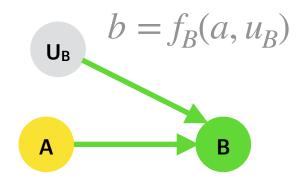
Back to SCM2CN: Non Quasi-Markovian Case

- Non Quasi-Markovian? Non-Linear constraint
- E.g., $\sum P(u_1) \cdot P(u_2) = \dots$
- Merge exogenous variables $U := (U_1, U_2)$
- Independence constraints can be disregarded (but higher exogenous dimensionality)
- Again CN approximate inference to solve causal queries
- State space dimensionality affects complexity
- We might have very large latent spaces ...



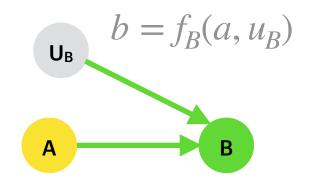


- Finding the equations given \mathscr{G} only
- P(B | A) should be a deterministic CPT





- Finding the equations given ${\mathscr G}$ only
- P(B | A) should be a deterministic CPT

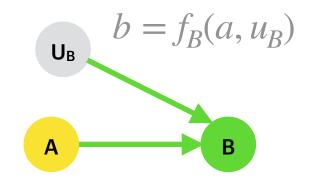


 $P(B \ A)$

						A=1	A=O	A=1
B=0	1	1	1	0	0	1	0	0
B=1	0	0	0	1	1	0	1	1
	<i>B</i> =	= 0	<i>B</i> :	=A	<i>B</i> =	: ¬A	B	= 1



- Finding the equations given $\mathcal G$ only
- P(B | A) should be a deterministic CPT
- U_B indexing all these deterministic CPTs

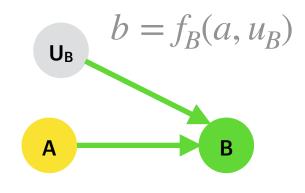


 $P(B \ A, U)$

	A=0	A=1	A=O	A=1	A=O	A=1	A=0	A=1
B=0	1	1	1	0	0	1	0	0
B=1	0	0	0	1	1	0	1	1
	U	=0	U	=1	U	=2	U	=3
	<i>B</i> =	= 0	<i>B</i> =	= A	<i>B</i> =	= ¬A	B	= 1



- Finding the equations given $\mathcal G$ only
- P(B | A) should be a deterministic CPT
- U_B indexing all these deterministic CPTs
- Knowledge might discard some states (ex., Bob goes to the party if Ann does)



 $P(B \ A, U)$

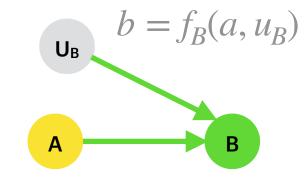




- Finding the equations given ${\mathscr G}$ only
- P(B | A) should be a deterministic CPT
- U_B indexing all these deterministic CPTs
- Knowledge might discard some states (ex., Bob goes to the party if Ann does)
- With Boolean parent & child) U = 4 in general (exp size) :

$$U = X \prod_{Y \in Pa_Y} Y$$

even more challenging with multiple exogenous parents



 $P(B \ A, U)$





CFs based on

B=O

U=0

B = 0

- Finding the equations given \mathcal{G} only
- P(B | A) should be a deterministic CPT
- U_{R} indexing
- Knowledge r (ex., Bob goe
- \mathcal{G} and \mathcal{D} only • With Boolea in general (exp size) :

$$U = X \prod_{Y \in Pa_Y} Y$$

even more challenging with multiple exogenous parents U=3

 $b = f_R(a, u_R)$

A, U)

U=2

B = A $B = \neg A$ B = 1

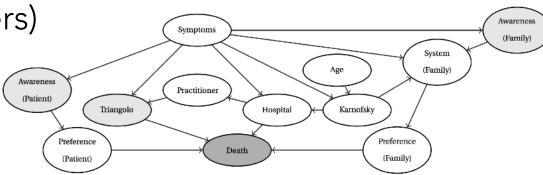
UΒ

U=1



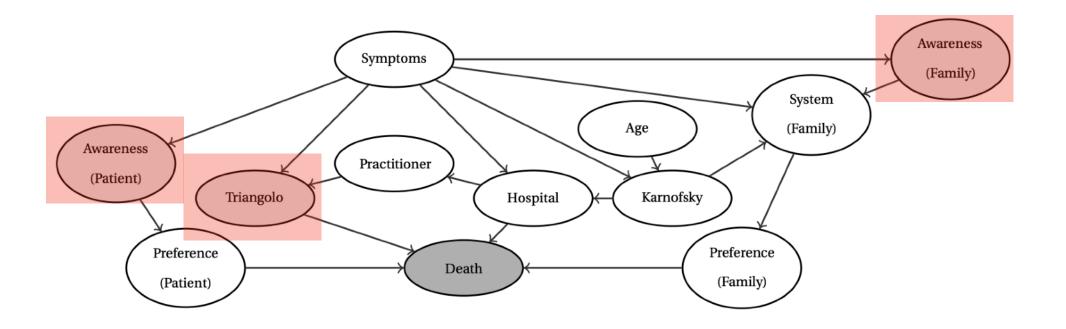
- Study of terminally ill cancer patients' preferences wrt their place of death (home or hospital)
- *G* obtained by expert knowledge and data
- Exogenous variables?
- Markovian assumption (= no confounders)







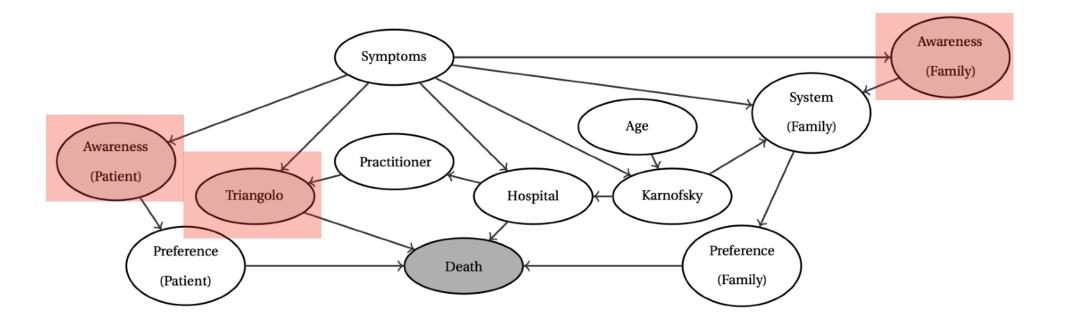
- Most patients prefer to die at home
- But a majority actually die in institutional settings
- Interventions by health care professionals can facilitate dying at home?



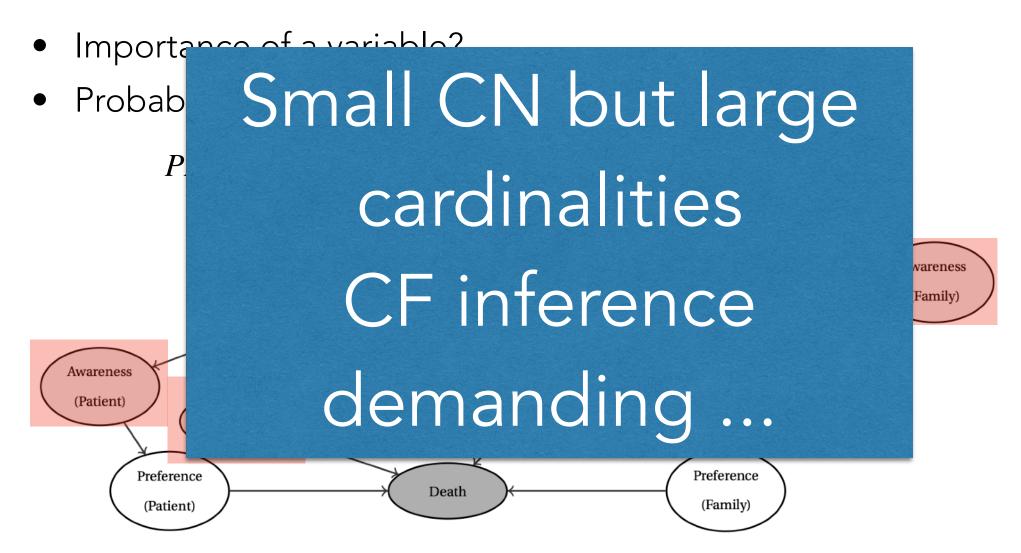


- Importance of a variable?
- Probability of necessity and sufficiency

 $PNS := P(Y_{X=1} = 1, Y_{X=0} = 0)$









Causal Expectation Maximisation (Zaffalon et al., 2021)

- Exogenous variables are always missing (MAR, asystematic, way)
- Expectation Maximisation (Dempster 1977)
 - Random initialisation of P(U)
 - E-step: Missing data completion by expected (fractional) counts
 - M-step: "completed" data to retrain P(U)
 - Iterate until convergence
- EM goes to a (local/global) max of $\log P(\mathcal{D})$



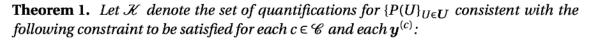
U1	U2	X1	X2	n
*	*	0	0	•••
*	*	0	1	
*	*	1	0	
*	*	1	1	•••





Causal EM: Likelihood Unimodality

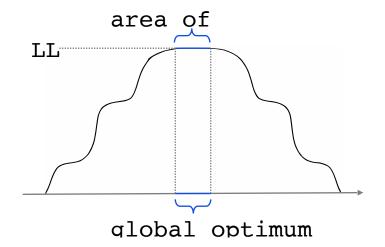
- Causal EM reduce should converge to global maxima only the corresponding P(U) belongs to credal set K(U)
- Sampling initialisations = sampling of K(U)
- For each sample we obtain an inner point



(8)

$$\sum_{\substack{\boldsymbol{u}^{(c)}: f_X(\mathrm{pa}_X) = x \\ \forall X \in \boldsymbol{X}^{(c)}}} \prod_{U \in \boldsymbol{U}^c} P(u) = \prod_{X \in \boldsymbol{X}^{(c)}} \hat{P}(x | \boldsymbol{y}_X^{(c)}),$$

where the values of u, x and $y_X^{(c)}$ are those consistent with $u^{(c)}$ and $y^{(c)}$. If $\mathcal{K} \neq \phi$, the log-likelihood in Eq. (7) achieves its global maximum if and only if $\{P(U)\}_{U \in U} \in \mathcal{K}$. If $\mathcal{K} = \phi$, the marginal log-likelihood in Eq. (7) can only take values strictly lower than the global maximum.







Causal EM: Guarantees?

- We first reduced causal queries to CN inference
- Causal EM reduces CN inference to (iterated) BN inference
- Identifiable queries? Each sample gives the same values (a numerical alternative to do-calculus)
- Unidentifiable? Each sample as an inner point
- Credible intervals can be derived

Theorem 5. Let $[a^*, b^*]$ denote the exact probability bounds of a causal query. Say that $\rho := \{r_i\}_{i=1}^n$ are the outputs of n EMCC iterations, while [a, b] is the interval induced by ρ , i.e., $a := \min_{i=1}^n r_i$ and $b := \max_{i=1}^n r_i$. By construction $a^* \le a \le b \le b^*$. The following inequality holds:

$$P\left(a - \varepsilon L \le a^* \le b^* \le b + \varepsilon L \, \middle| \, \rho\right) = \frac{1 + (1 + 2\varepsilon)^{2-n} - 2(1 + \varepsilon)^{2-n}}{(1 - L^{n-2}) - (n-2)(1 - L)L^{n-2}},\tag{13}$$

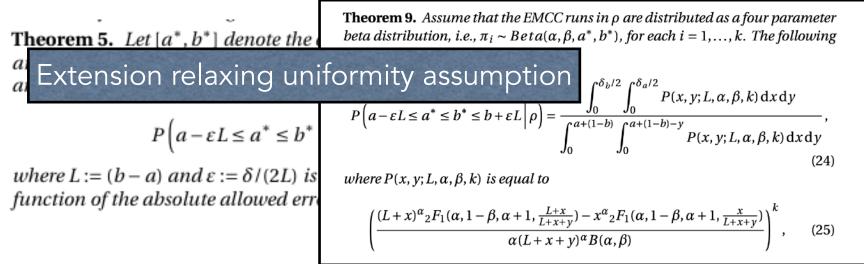
where L := (b - a) and $\varepsilon := \delta/(2L)$ is the relative error at each extreme of the interval obtained as a function of the absolute allowed error $\delta \in (0, L)$.





Causal EM: Guarantees?

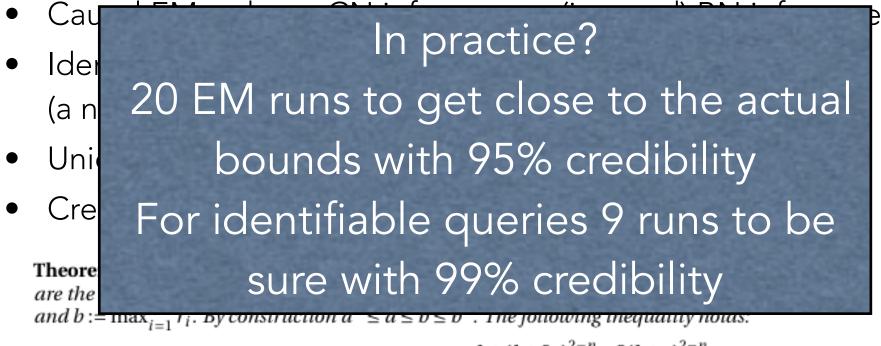
- We first reduced causal queries to CN inference
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- Unidentifiable? Each sample as an inner point
- Credible intervals can be derived





Casual EM: Guarantees?

• We first reduced causal queries to CN inference



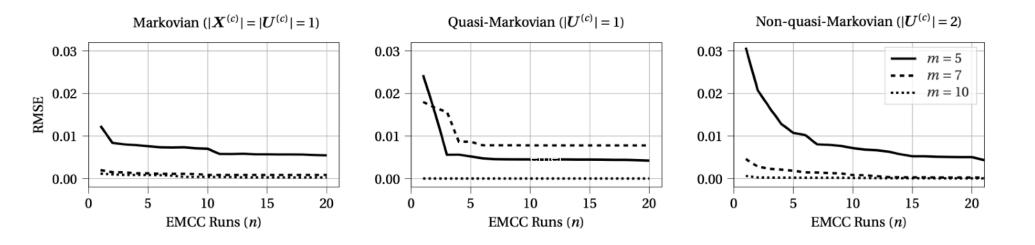
$$P\left(a - \varepsilon L \le a^* \le b^* \le b + \varepsilon L \,\middle|\, \rho\right) = \frac{1 + (1 + 2\varepsilon)^{2-n} - 2(1 + \varepsilon)^{2-n}}{(1 - L^{n-2}) - (n-2)(1 - L)L^{n-2}},\tag{13}$$

where L := (b - a) and $\varepsilon := \delta/(2L)$ is the relative error at each extreme of the interval obtained as a function of the absolute allowed error $\delta \in (0, L)$.





Causal EM: Experiments



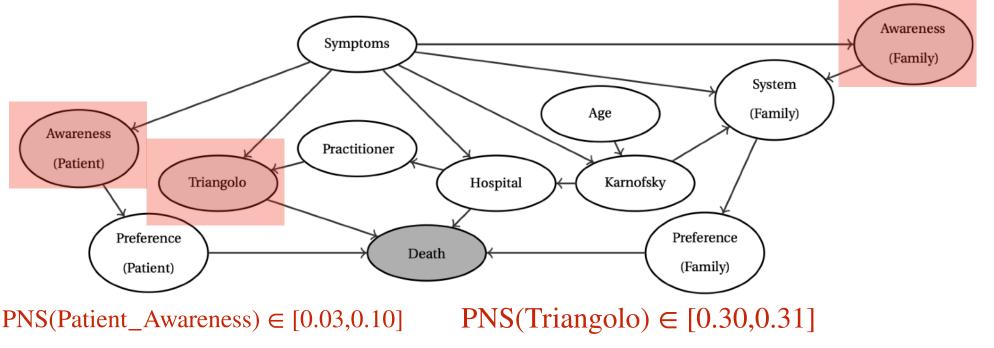
PNS for artificial SMCs: quick convergence (= much faster than direct CN approach)





Counterfactual Analysis in Palliative Cares by Causal EM

- Importance of a variable?
- Probability of necessity and sufficiency $PNS := P(Y_{X=1} = 1, Y_{X=0} = 0)$
- 15 EM runs before convergence PNS(Family_Awareness) ∈ [0.06,0.10]



Alessandro Antonucci, IDSIA



One should act on Triangolo first: for instance,
 In by making Triangolo available to all patients, we
 Pr should expect a reduction of people at the hospital by 30%

This would save money too, and would allow politicians to do economic considerations as to which amount it is even economically profitable to fund Triangolo, and have patients die at home, rather than spending more to have patients die at the hospital

 $PNS(Patient_Awareness) \in [0.03, 0.10] \qquad PNS(Triangolo) \in [0.30, 0.31]$

).06,0.10]



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SUPSI

Causal Analysis from **Biased** Data

• Selective data acquisition (untreated M and treated F missing)

Treatment X	Recovery Y	Gender Z	counts
0	0	0	2
I	0	0	41
0	I.	0	114
I	Ι	0	313
0	0	I	107
l.	0	I.	109
0	I	I	13
I.	I.	I.	I.



SUPSI

Causal Analysis from **Biased** Data

- Selective data acquisition (untreated M and treated F missing)
- A (Boolean) **selector** variable $S \equiv (X \neq Z)$

Treat, X	Recover y	Gender Z	Selector S	counts
*	*	*	0	2
I	0	0	Ι	41
*	*	*	0	114
I	Ι	0	I	313
0	0	Ι	Ι	107
*	*	*	0	109
0	Ι	Ι	Ι	13
*	*	*	0	I





Causal Analysis from **Biased** Data

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- A (Boolean) **selector** variable $S \equiv (X \neq Z)$
- Assume we know $n(S = 0) \propto P(S = 0)$

Treat, X	Recover y Y	Gender Z	Selector S	counts
Ι	0	0	I	41
I	I	0	I	313
0	0	I	I	107
0	I	I	I	13
*	*	*	0	226





Causal Analysis from **Biased** Data

- Selective data acquisition (untreated M and treated F missing)
- A (Boolean) **selector** variable $S \equiv (X \neq Z)$
- Assume we know $n(S = 0) \propto P(S = 0)$
- Interventional queries with bias?
- Do calculus for selection bias Barenboim & Tian (AAAI, 2015)

Treat, X	Recover y Y	Gender Z	Selector S	counts
I	0	0	I	41
I	Ι	0	I	313
0	0	I	I	107
0	I	I	I	13
*	*	*	0	226

Recovering Causal Effects from Selection Bias					
Elias Bareinboim*	Jin Tian*				
Computer Science Department	Department of Computer Science				
University of California, Los Angeles	Iowa State University				
Los Angeles, CA. 90095	Ames, IA. 50011				
eb@cs.ucla.edu	jtian@iastate.edu				

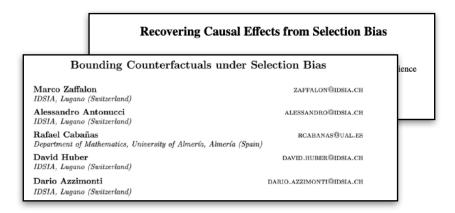


SUPSI

Causal Analysis from **Biased** Data

- Selective data acquisition (untreated M and treated F missing)
- A (Boolean) **selector** variable $S \equiv (X \neq Z)$
- Assume we know $n(S = 0) \propto P(S = 0)$
- Interventional queries with bias?
- Do calculus for selection bias Barenboim & Tian (AAAI, 2015)
- Unidentifiable queries?
- Our EM(CC) can be used for that!

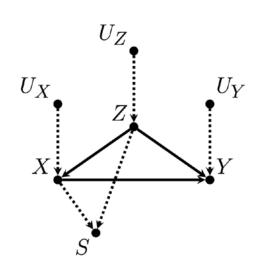
Treat, X	Recover y Y	Gender Z	Selector S	counts
I	0	0	I	41
I	I	0	I	313
0	0	I	I	107
0	I	I	I	13
*	*	*	0	226





• *S* determined by an equation, a SCM!

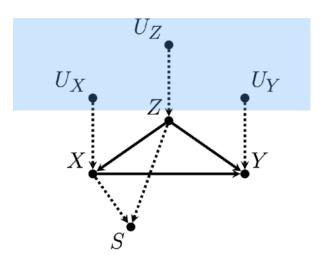
UX	UY	UZ	х	Y	Z	S	n
*	*	*	I	0	0	I	41
*	*	*	Ι	Ι	0	Ι	313
*	*	*	0	0	I	Ι	107
*	*	*	0	I	I	I	13
*	*	*	*	*	*	0	226





- *S* determined by an equation, a SCM!
- CN approach? No, S = 1 induces relations between P(U)'s in the CN

	UX	UY	UZ	x	Y	Z	S	n
ľ	*	*	*	I	0	0	I	41
	*	*	*	I	I	0	I	313
	*	*	*	0	0	I	ļ	107
	*	*	*	0	I	I	I	13
	*	*	*	*	*	*	0	226

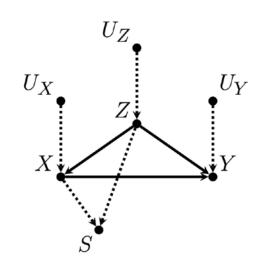






- *S* determined by an equation, a SCM!
- CN approach? No, S = 1 induces relations between P(U)'s in the CN
- EM? Maybe, but "non-rectangular" missingness, might kill unimodality ...
- Convergence to max preserved? (hence inner points of $[\underline{P}, \overline{P}]$)

UX	UY	UZ	х	Y	Z	S	n
*	*	*	I	0	0	I	41
*	*	*	I	I	0	I	313
*	*	*	0	0	I	I	107
*	*	*	0	I	I	I	13
*	*	*	*	*	*	0	226



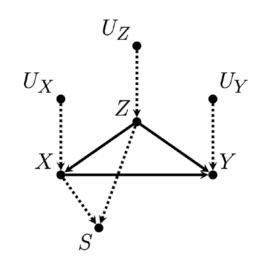




- *S* determined by an equation, a SCM!
- CN approach? No, S = 1 induces relations between P(U)'s in the CN
- EM? Maybe, but "non-rectangular" missingness, might kill unimodality ...
- Convergence to max preserved? (hence inner points of $[\underline{P}, \overline{P}]$) Yes!

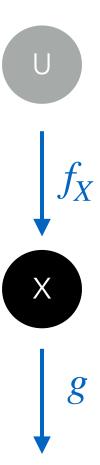
Theorem 4 As a function of $\{P(U)\}_{U \in U}$, the log-likelihood in Eq. (7) has no local maxima and a global maximum equal to the value LL^* in Eq. (6). Such a maximum is achieved if and only if the M-compatibility constraints in Eqs. (8) and (9) are satisfied.

UX	UY	UZ	х	Y	Z	S	n
*	*	*	Ι	0	0	Ι	41
*	*	*	I	I	0	I	313
*	*	*	0	0	I	I	107
*	*	*	0	I	I	I	13
*	*	*	*	*	*	0	226





S	Sketch of the proof										
	UX	UY	UZ	х	Y	Z	S	n			
	*	*	*	Ι	0	0	I	41			
	*	*	*	I	I	0	I	313			
	*	*	*	0	0	I	I	107			
	*	*	*	0	I	I	I	13			
	*	*	*	*	*	*	0	226			



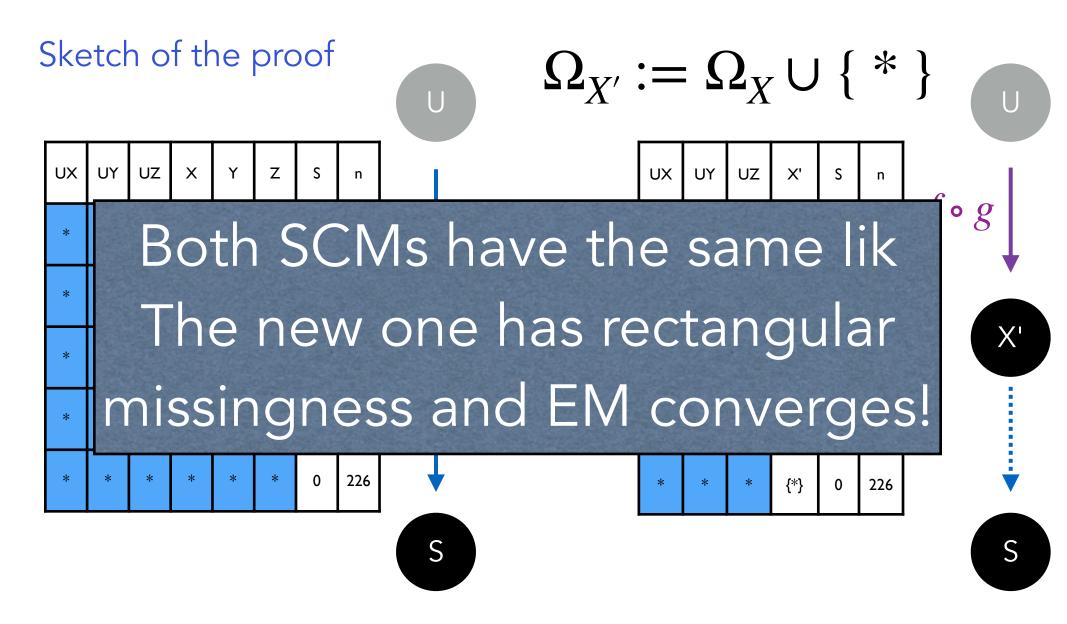
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SUPSI



Sketch of the proof						U		$\Omega_{X'}$:=	: (\mathbf{D}_X	U	{	*	}			
	UX	UY	υz	х	Y	z	S	n		0		UX	UY	UZ	X'	S	n	C
	*	*	*	I	0	0	I	41	f_2	X		*	*	*	{100}	I	41	$f \circ g$
	*	*	*	I	I	0	I	313				*	*	*	{110}	I	313	
	*	*	*	0	0	I	I	107	X			*	*	*	{001}	I	107	
	*	*	*	0	I	I	I	13	8	8		*	*	*	{011}	I	13	
	*	*	*	*	*	*	0	226	↓			*	*	*	{*}	0	226	▼
•					-	-			S									S







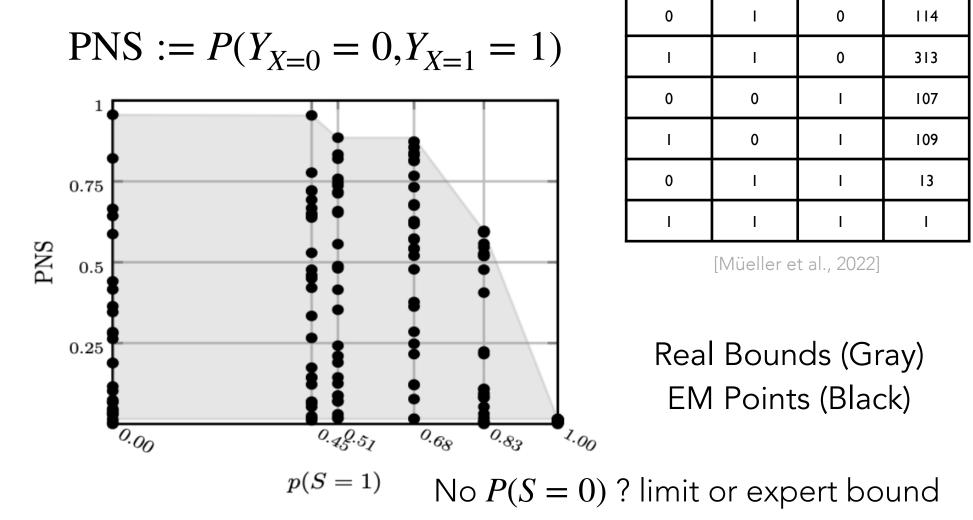
counts

2

41

Counterfactual Bounds for Biased Data

Probability of Necessity and Sufficiency



Treatment

Х

0

I

Recovery

0

0

Gender

0

0



Current Work: Hybrid Data

Learning to Bound Counterfactual Inference from Observational, Biased and Randomised Data

Marco Zaffalon^a, Alessandro Antonucci^{a,*}, Rafael Cabañas^b, David Huber^a

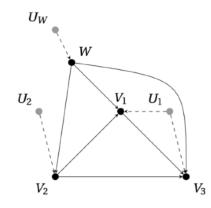
^aIDSIA, Lugano (Switzerland) ^bDepartment of Mathematics, University of Almería, Almería (Spain)

Study	Treatment	Gender	Survival	Counts
	do(drug)	female	survived	489
	do(drug)	female	dead	511
	do(drug)	male	survived	490
interventional	do(drug)	male	dead	510
Interventional	do(no drug)	female	survived	210
	do(no drug)	female	dead	790
	do(no drug)	male	survived	210
	do(no drug)	male	dead	790
	drug	female	survived	378
	drug	female	dead	1022
	drug	male	survived	980
observational	drug	male	dead	420
observational	no drug	female	survived	420
	no drug	female	dead	180
	no drug	male	survived	420
	no drug	male	dead	180

Table 1: Data from interventional and observational studies on the potential effects of a drug on patients affected by a deadly disease.

Treatment	Gender	Survival	W	Counts
drug	female	survived	drug	489
drug	female	dead	drug	511
drug	male	survived	drug	490
drug	male	dead	drug	510
no drug	female	survived	no drug	210
no drug	female	dead	no drug	790
no drug	male	survived	no drug	210
no drug	male	dead	no drug	790
drug	female	survived	w_{ϕ}	378
drug	female	dead	w_{ϕ}	1022
drug	male	survived	w_{ϕ}	980
drug	male	dead	w_{ϕ}	420
no drug	female	survived	w_{arphi}	420
no drug	female	dead	w_{ϕ}	180
no drug	male	survived	w_{ϕ}	420
no drug	male	dead	w_{ϕ}	180

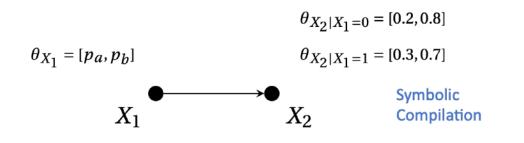
Table 2: A merged version of the two datasets in Table 1 with the index variable *W*.

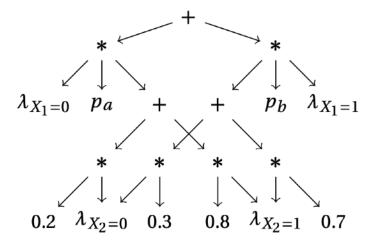




Current Work: Symbolic Knowledge Compilation (TPM 2023)

- Joint work with Adnan Darwiche and Hizuo Chen
- Our EM requires many (BN) queries
- Equations remain constant
- Compile BN once, use many times
- Symbolic compilation

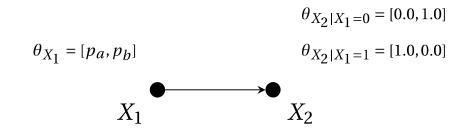


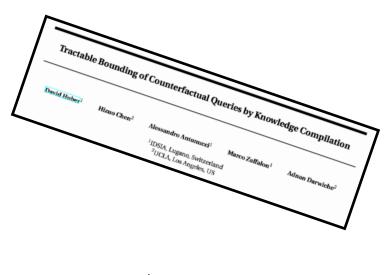


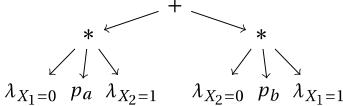


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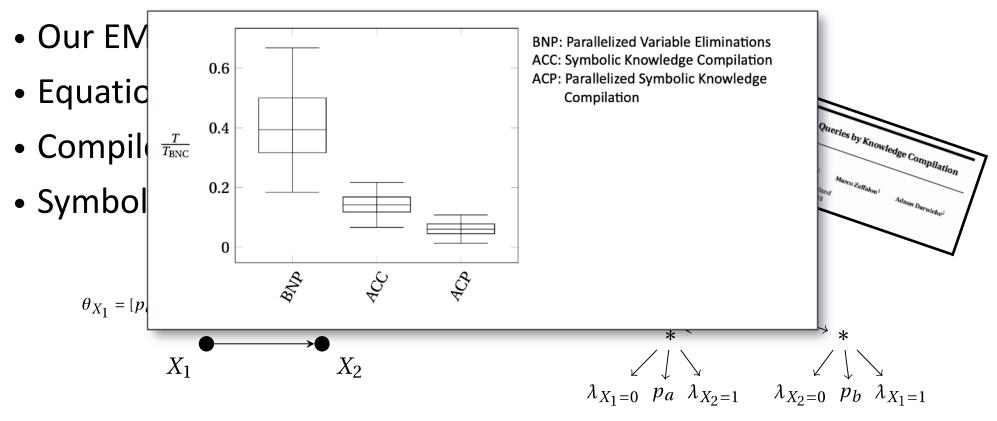


83



Current Work: Symbolic Knowledge Compilation (TPM 2023)

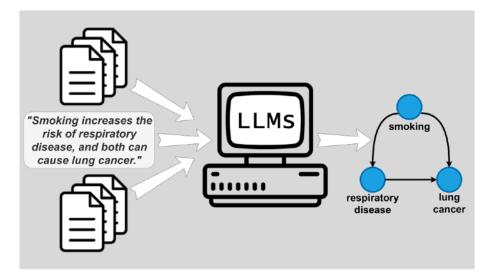
Joint work with Adnan Darwiche and Hizuo Chen







• GPT parsing causal statements in natural language



- Link with IPs? Multiple causal graphs might be returned!
- Many recent papers on bounding counterfactual wrt ignorance about the causal structure (credal structures?)





Conclusions

- Causality theories have an intimate connection with credal models
- Past research about CNs might offer new tools for causal analysis
- CNs offer formalism for a deeper SCMs understanding
- Lot of work to be done, causal ML/RL just at the beginning!
- Our current directions are:
 - Canonical models
 - Continuous Variables
 - XAI (Counterfactual Explanations)
 - "Credal EM" propagating credal initialisations?



models

alysis

Conclusions

- Causality theories have an intimate connection
- Past research about CNs might offer new
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- Our current dire Saint
 - Canop
 - Parallelisation Со
 - Continuous Variables
 - Learning Causal Graphs (GPT)



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