Causal Transformer for Estimating Counterfactual Outcomes

Valentyn Melnychuk, Dennis Frauen, Stefan Feuerriegel

LMU Munich, Munich, Germany

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Institute of AI for Management @ LMU Munich

Who are we?



Valentyn Melnychuk, PhD candidate



Dennis Frauen, PhD candidate



Prof. Dr. Stefan Feuerriegel

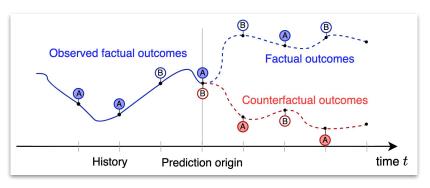
Main research topics: Causal machine learning, Treatment effect estimation, Causal representation learning

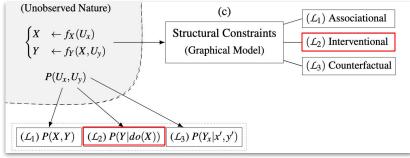
www.ai.bwl.uni-muenchen.de

Introduction: Estimating counterfactual outcomes over time

Why this is important?

- Counterfactual prediction allows to answer individualized "what if" questions: what will happen to the patient, if I apply an alternative sequence of treatments, counterfactual to a standard treatment policy
- Here, **potential outcomes** are meant, which correspond to the interventional level of valuation in Pearl's Hierarchy of Causal Inference¹
- Growing opportunity to employ **observational data**:
 - randomized controlled trials (RCTs) are costly and/or unethical
 - abundance of large-scale observational data, e.g., electronic health records







Introduction: Estimating counterfactual outcomes over time

Problem formulation

Given observational dataset of:

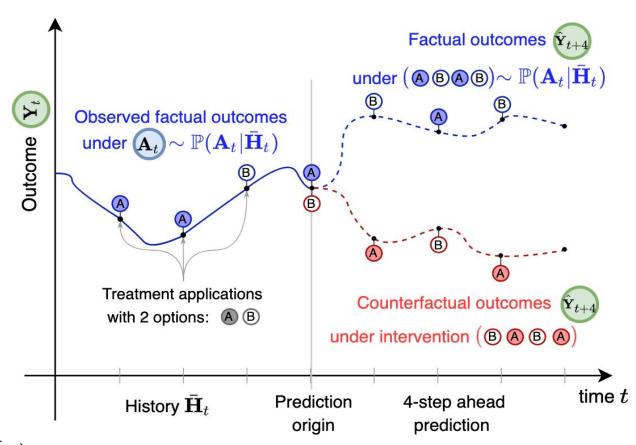
- x time-varying covariates (e.g., blood pressure)
- v static covariates (e.g., age)
- categorical treatments (e.g., ventilation)
- (factual*) outcomes (e.g., respiratory frequency)

we want to estimate expected **counterfactual outcomes over time** starting from prediction origin for a given sequence of treatment interventions:

$$\mathbb{E}\big(\mathbf{Y}_{t+\tau}\big[\bar{\mathbf{a}}_{t:t+\tau-1}\big] \mid \mathbf{H}_t\big)$$

$$\tau\text{-step-ahead} \quad \text{sequence of counterfactual outcome} \quad \text{treatment interventions} \quad \text{history before prediction origin}$$

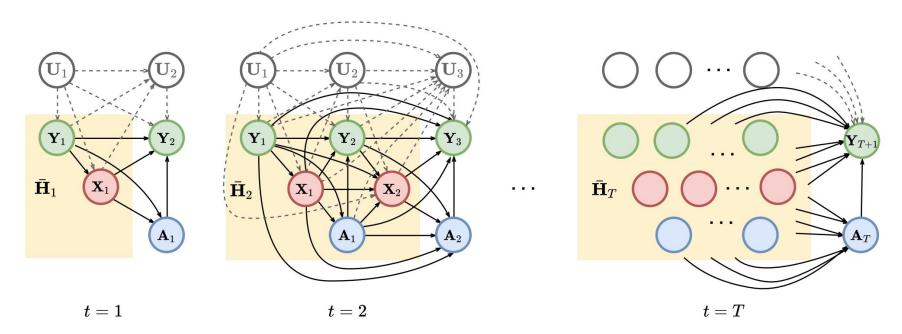
For that, we aim to learn a function $g(au, ar{\mathbf{a}}_{t:t+ au-1}, ar{\mathbf{H}}_t)$



Introduction: Assumptions

Identifiability assumptions

- ullet Consistency. If $ar{f A}_t=ar{f a}_t$ is a given sequence of treatments for some patient, then ${f Y}_{t+1}[ar{f a}_t]={f Y}_{t+1}$
- Sequential Overlap. There is always a non-zero probability of receiving/not receiving any treatment, conditioning on the previous history: $0 < \bar{\mathbb{P}}(\mathbf{A}_t = \mathbf{a}_t \mid \bar{\mathbf{H}}_t = \bar{\mathbf{h}}_t) < 1$
- Sequential Ignorability. Current treatment is independent of the potential outcome, conditioning on the observed history $\mathbf{A}_t \perp \mathbf{Y}_{t+1}[\mathbf{a}_t] \mid \bar{\mathbf{H}}_t$



Introduction: Task complexity

Why estimation is hard?

- Fundamental problem of causal inference:
 counterfactual outcomes are never directly observed in a real world
- Traditional machine learning to learn $g(\cdot)$ is either **sub-optimal** (one-step-ahead prediction) or **biased** (multiple-step-ahead prediction) in the presence of time-varying confounding
- Observed history grows with time:
 - existing reinforcement literature is non-applicable as this is a non-Markovian setting
 - existing literature for cross-sectional setting, e.g. individual treatment effect (ITE) / conditional average treatment effect (CATE), also falls short
- Although the causal effect is identifiable, i.e., with G-Computation formula, it is unclear, how to leverage a bias-variance tradeoff and computational complexity:

$$\mathbb{E}\left(\mathbf{Y}_{t+\tau}[\bar{\mathbf{a}}_{t:t+\tau-1}] \mid \bar{\mathbf{H}}_{t}\right) = \int_{\mathbb{R}^{d_{x}} \times \cdots \times \mathbb{R}^{d_{x}}} \mathbb{E}\left(\mathbf{Y}_{t+\tau} \mid \bar{\mathbf{H}}_{t}, \bar{\mathbf{x}}_{t+1:t+\tau-1}, \bar{\mathbf{y}}_{t+1:t+\tau-1}, \bar{\mathbf{a}}_{t:t+\tau-1}\right) \times \prod_{j=t+1}^{t+\tau-1} \mathbb{P}\left(\mathbf{x}_{j}\mathbf{y}_{j} \mid \bar{\mathbf{H}}_{t}, \bar{\mathbf{x}}_{t+1:j-1}, \bar{\mathbf{y}}_{t+1:j-1}, \bar{\mathbf{a}}_{t:j-1}\right) d\bar{\mathbf{x}}_{t+1:t+\tau-1} d\bar{\mathbf{y}}_{t+1:t+\tau-1}$$

Introduction: Related methods

Related methods

- Marginal Structural Models (MSMs) (Robins et al., 2000; Hernan et al., 2001)
 - Base models: linear models wrt. a fixed window taken from history
 - Estimation: (1) propensity score estimation; (2) pseudo-outcome regressions, with IPTW weighted trajectories
- Recurrent Marginal Structural Networks (RMSNs) (Lim et al., 2018)
 - Base models: 2 propensity LSTMs, encoder LSTM, decoder LSTM
 - Estimation: (1) propensity score estimation; (2) pseudo-outcome regressions, with IPTW weighted trajectories
- Counterfactual Recurrent Network (CRN) (Bica et al., 2020)
 - o Base models: encoder LSTM, decoder LSTM
 - Estimation: balanced representations via gradient reversal
- **G-Net** (Li et al., 2021)
 - Base models: time-varying covariates and outcome LSTM
 - Estimation: sampling-based G-computation

Introduction: Research gap – Our contributions

Research gap

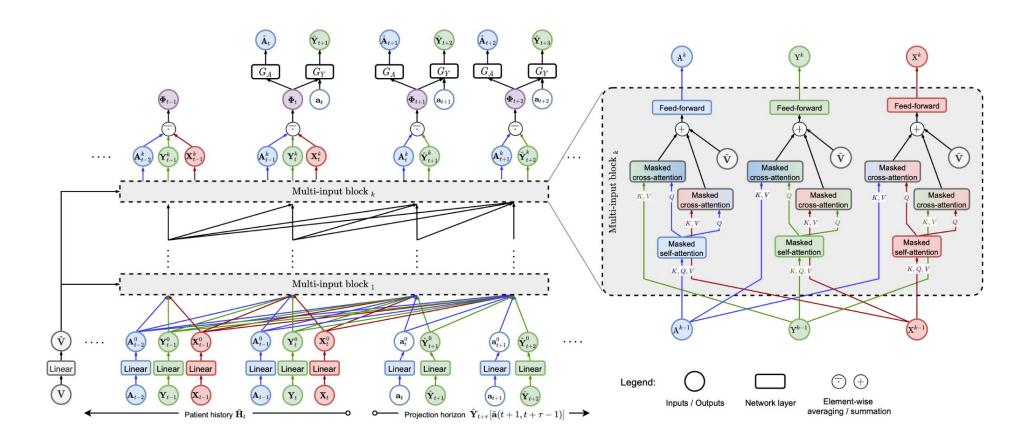
Current state-of-the-art methods are built on top of long short-term memory (LSTM), thus
rendering inferences for complex, long-range dependencies challenging

Our contributions

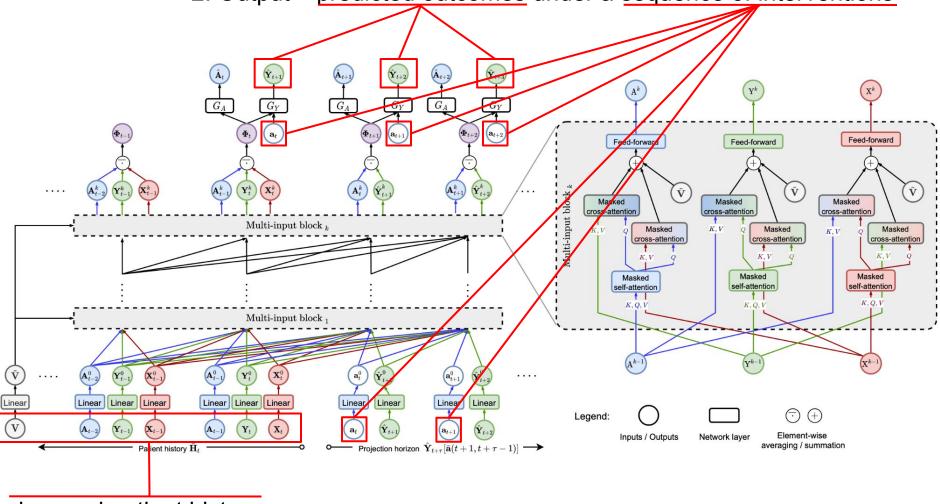
Causal Transformer (CT) is an end-to-end model, first tailoring of transformers to a counterfactual prediction task over time:

- CT captures complex, long-range dependencies between time-varying covariates, treatments and outcomes
- CT employs a novel adversarial counterfactual domain confusion (CDC) loss to address a time-varying confounding
- CT achieves **state-of-the-art performance** on synthetic, semi-synthetic & real benchmarks

CT is a single end-to-end model for **both one- and multiple-step-ahead prediction**



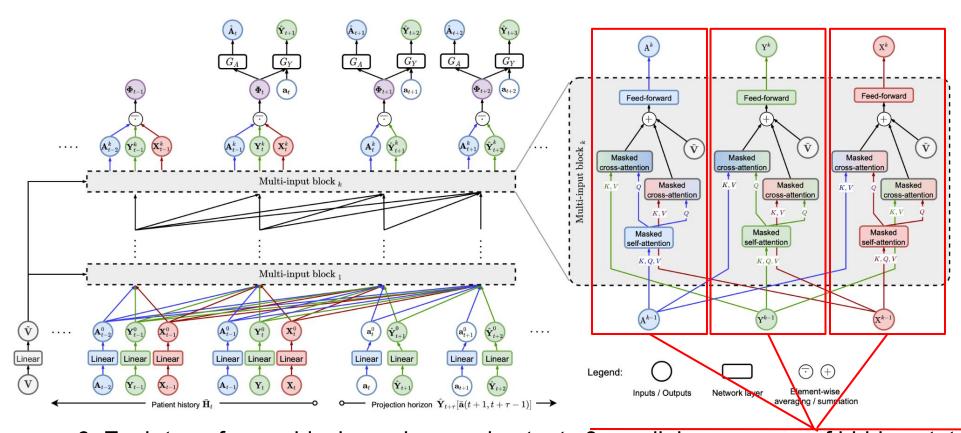
2. Output – <u>predicted outcomes</u> under a <u>sequence of interventions</u>



1. Input – observed patient history

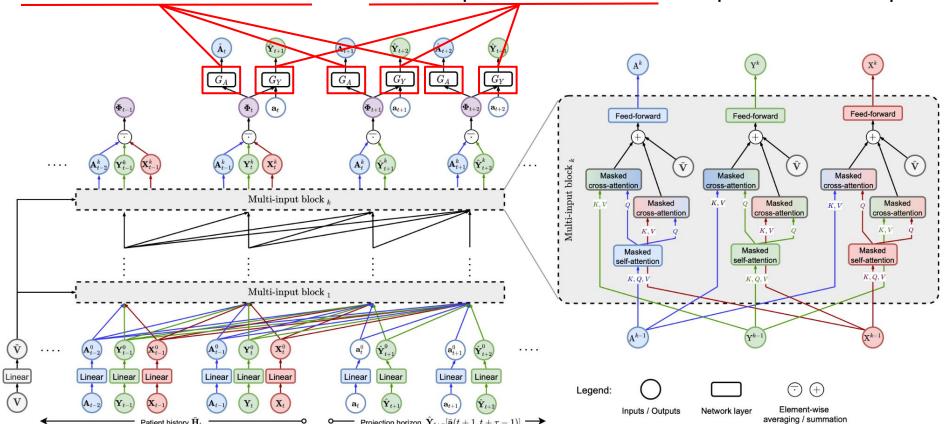
4. Outputs of the last block are averaged 3. Inputs are and form balanced representations transformed with a stack of multi-input blocks Feed-forward Feed-forward Feed-forward $\tilde{\mathbf{V}}$ Multi-input block Masked Masked cross-attention cross-attention cross-attention Multi-input block k Masked Masked Masked cross-attention cross-attention Masked self-attention self-attention self-attention Multi-input block Linear Linear Linear Linear Linear \odot Legend: Inputs / Outputs Network laver Projection horizon $\hat{\mathbf{Y}}_{t+\tau}[\bar{\mathbf{a}}(t+1,t+\tau-1)]$

5. Each block is equipped with self-attention, cross-attention and feed-forward layers



6. Each transformer block receives and outputs 3 parallel sequences of hidden states. I.e., there CT has 3 subnetworks, and the information between them is shared via cross-attentions

7. We place treatment classifier network and outcome prediction network on top of balanced representations



8. Both treatment classifier and outcome prediction networks are used for the novel counterfactual domain confusion loss (CDC) loss

Other details

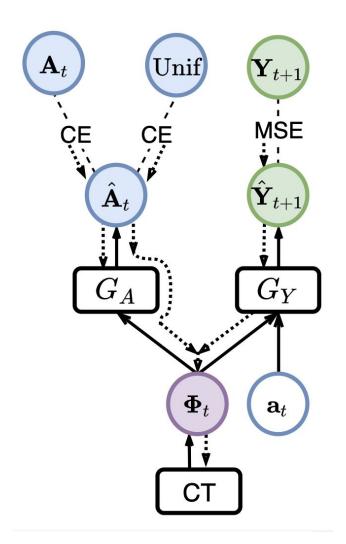
- Each transformer block is minimal¹ and combines
 - (i) multi-head self-/cross-attention with residual connections
 - (ii) feed-forward layer with residual connections
 - (iii) layer normalization
- We employed attentional dropout², analogously to the recurrent dropout in LSTMs.
- In every self- and cross-attention, we use trainable **relative positional encodings**³, which:
 - considers the order of treatments, outcomes and time-varying covariates relatively to the prediction origin. E.g., they allow us to distinguish sequences such as, e. g., <treatment A → side effect → treatment B > from <treatment A → treatment B → side-effect>
 - allow for better generalization to unseen sequence length by dropping the order information for the distant past
- Mini-batch augmentation with masking is used to enable multi-step-ahead prediction, where future time-varying covariates are unavailable

¹ Dong, Yihe, Jean-Baptiste Cordonnier, and Andreas Loukas. "Attention is not all you need: Pure attention loses rank doubly exponentially with depth." International Conference on Machine Learning. PMLR, 2021.

² Zehui, Lin, et al. "DropAttention: a regularization method for fully-connected self-attention networks." arXiv preprint arXiv:1907.11065 (2019).

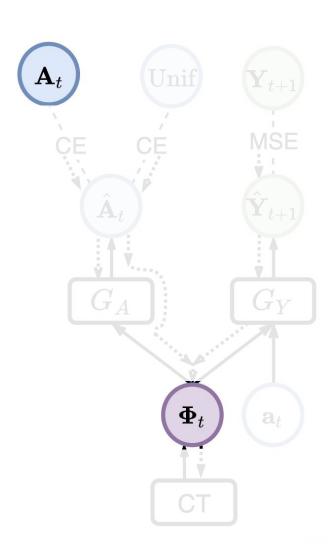
³ Shaw, Peter, Jakob Uszkoreit, and Ashish Vaswani. "Self-attention with relative position representations." arXiv preprint arXiv:1803.02155 (2018).

- Idea stems from the unsupervised domain adaptation¹
- CDC is an adversarial objective, which aims at same time to:



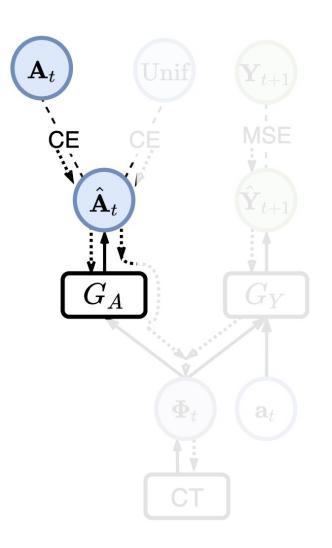
¹ Tzeng, Eric, et al. "Simultaneous deep transfer across domains and tasks." Proceedings of the IEEE international conference on computer vision (2015)

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 - (a) make **balanced representations** (Φ_t) **non-predictive** of the current treatment (A_t)



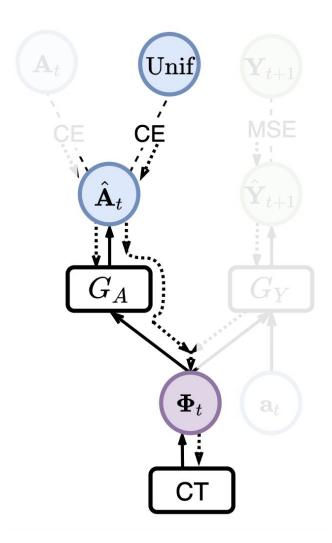
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 - by minimizing cross-entropy of current treatment wrt. G_A



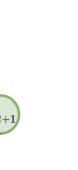
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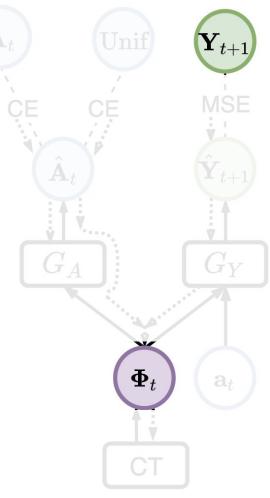
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 - (b) make balanced representations Φ_t predictive of the outcome Ψ_{t+1}

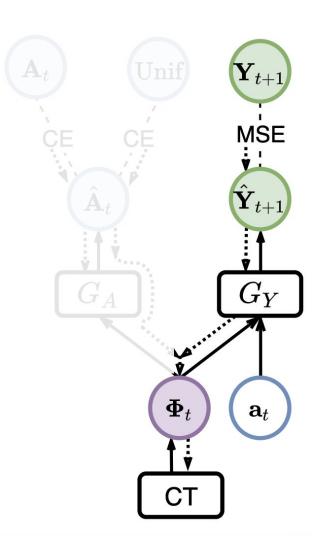




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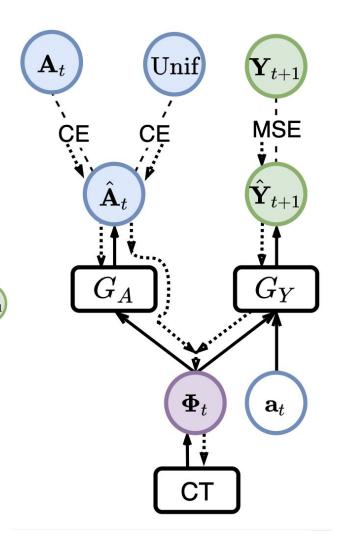
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 - by minimizing cross-entropy of current treatment wrt. G_A
 - by minimizing cross-entropy between uniform treatment and output of treatment classifier network wrt.
 - (b) make balanced representations (Φ_t) predictive of the outcome (\mathbf{Y}_{t+1})
 - ullet minimizing factual MSE wrt. lacktriangle and lacktriangle





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- Adversarial learning is further stabilized with exponential moving average (EMA) of model weights



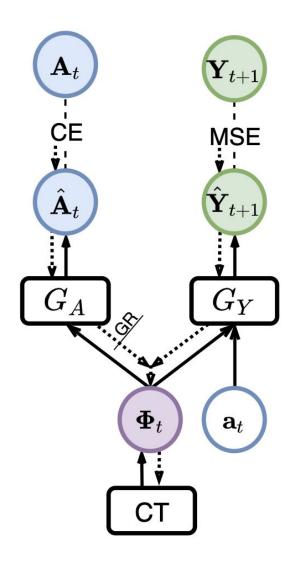
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Causal Transformer: Theoretical insights

- Previously proposed **gradient reversal**¹ (CRN, Bica et al., 2020) extends in two ways:
 - if badly chosen hyperparameter -> representation may be predictive of opposite treatment
 - gradients vanish, if treatment classifier network learns too fast
- We prove a theorem, similar to (CRN, Bica et al., 2020): finding a solution to an adversarial objective of CDC loss renders distributions of representations conditional on each treatment equal (= balanced)
- In our case, we minimize a reversed KL-divergence:

CDC loss (our paper)	Gradient reversal (CRN, Bica et al., 2020)				

where $P_j^{\Phi}(x')$ is a distribution of representation conditional on treatment j



Ganin, Yaroslav, and Victor Lempitsky. "Unsupervised domain adaptation by backpropagation." International conference on machine learning. PMLR, 2015

Experiments: Datasets – Baselines

We evaluate CT based on:

Datasets

o synthetic datasets based on pharmacokinetic-pharmacodynamic model of tumor growth

$$\mathbf{Y}_{t+1} = \left(1 + \rho \log \left(\frac{K}{\mathbf{Y}_t}\right) - \beta_c C_t - (\alpha_r d_t + \beta_r d_t^2) + \varepsilon_t\right) \mathbf{Y}_t \qquad \mathbf{A}_t^c, \mathbf{A}_t^r \sim \mathrm{Bernoulli}\left(\sigma \left(\frac{\gamma}{D_{\max}}(\bar{D}_{15}(\bar{\mathbf{Y}}_{t-1}) - D_{\max}/2)\right)\right)$$

o self-designed **semi-synthetic dataset** based on MIMIC-III dataset

$$\mathbf{Z}_{t}^{j,(i)} = \underbrace{\alpha_{S}^{j} \, \text{B-spline}(t) + \alpha_{g}^{j} \, g^{j,(i)}(t)}_{\text{endogenous}} + \underbrace{\alpha_{f}^{j} \, f_{Z}^{j}(\mathbf{X}_{t}^{(i)})}_{\text{exogenous}} + \underbrace{\varepsilon_{t}}_{\text{noise}} \qquad \begin{aligned} p_{\mathbf{A}_{t}^{l}} &= \sigma \Big(\gamma_{A}^{l} \bar{A}_{T_{l}}(\bar{\mathbf{Y}}_{t-1}) + \gamma_{X}^{l} f_{Y}^{l}(\mathbf{X}_{t}) + b_{l} \Big) \qquad \mathbf{Y}_{t}^{j} &= \mathbf{Z}_{t}^{j} \, + E^{j}(t) \\ \mathbf{A}_{t}^{l} \sim \text{Bernoulli}\left(p_{\mathbf{A}_{t}^{l}}\right), \end{aligned}$$

- real-world dataset (MIMIC-III)
- Only synthetic and semi-synthetic data have ground-truth counterfactuals; real-world evaluation is a proof of concept
- We compared root-mean-squared error (RMSE) of one and multiple-step-ahead predictions.
 For multiple-step-ahead we sampled a fixed number of random counterfactual trajectories

Baselines

- Marginal Structural Models (MSMs) (Robins et al., 2000; Hernan et al., 2001)
- Recurrent Marginal Structural Networks (RMSNs) (Lim et al., 2018)
- Counterfactual Recurrent Network (CRN) (Bica et al., 2020)
- G-Net (Li et al., 2021)

Experiments: Results

Results

• CT achieves **superior performance** over current baselines for benchmarks with long-range dependencies and long prediction horizons, e.g., for semi-synthetic benchmark:

	$ \tau = 1$	au = 2	$\tau = 3$	au=4	$\tau = 5$	$\tau = 6$	au=7	$\tau = 8$	$\tau = 9$	$\tau = 10$
MSMs (Robins et al., 2000) RMSNs (Lim et al., 2018) CRN (Bica et al., 2020) G-Net (Li et al., 2021)	$ \begin{vmatrix} 0.37 \pm 0.01 \\ 0.24 \pm 0.01 \\ 0.30 \pm 0.01 \\ 0.34 \pm 0.01 \end{vmatrix} $	$ \begin{vmatrix} 0.57 \pm 0.03 \\ 0.47 \pm 0.01 \\ 0.48 \pm 0.02 \\ 0.67 \pm 0.03 \end{vmatrix} $	0.74 ± 0.06 0.60 ± 0.01 0.59 ± 0.02 0.83 ± 0.04	0.88 ± 0.03 0.70 ± 0.02 0.65 ± 0.02 0.94 ± 0.04	1.14 ± 0.10 0.78 ± 0.04 0.68 ± 0.02 1.03 ± 0.05	1.95 ± 1.48 0.84 ± 0.05 0.71 ± 0.01 1.10 ± 0.05	3.44 ± 4.57 0.89 ± 0.06 0.72 ± 0.01 1.16 ± 0.05	> 10.0 0.94 ± 0.08 0.74 ± 0.01 1.21 ± 0.06	> 10.0 0.97 \pm 0.09 0.76 \pm 0.01 1.25 \pm 0.06	> 10.0 1.00 ± 0.11 0.78 ± 0.02 1.29 ± 0.06
EDCT w/ GR ($\lambda = 1$) (ours) CT ($\alpha = 0$) (ours) * CT (ours)	0.29 ± 0.01 0.20 ± 0.01 0.20 ± 0.01	0.46 ± 0.01 0.38 ± 0.01 0.38 ± 0.01	0.56 ± 0.01 0.45 ± 0.01 0.45 ± 0.01	0.62 ± 0.01 0.50 ± 0.02 0.49 ± 0.01	0.67 ± 0.01 0.52 ± 0.02 0.52 ± 0.02	0.70 ± 0.01 0.55 ± 0.02 0.53 ± 0.02	0.72 ± 0.01 0.56 ± 0.02 0.55 ± 0.02	0.74 ± 0.01 0.58 ± 0.02 0.56 ± 0.02	0.76 ± 0.01 0.60 ± 0.02 0.58 ± 0.02	0.78 ± 0.01 0.61 ± 0.02 0.59 ± 0.02

Lower = better (best in bold)

 Among all the neural models, our CT has the smallest runtime, due to single-stage training procedure with CDC loss and usage of self-attention:

	Main stages of training & inference	Total runtime (in min)			
MSMs	2 logistic regressions for IPTW & linear regression	3.5	±	0.3	
RMSNs	2 networks for IPTW & encoder & decoder	109.7	±	2.3	
CRN	encoder & decoder	75.3	±	17.5	
G-Net	single network & MC sampling for inference	118.0	±	2.0	
CT (ours)	single multi-input network	13.5	±	4.8	

Experiments: Ablation study

Ablation types

Based on synthetic datasets we evaluate different versions of CT with varying:

- (a) different components within the subnetworks (positional encodings, attentional dropout)
- (b) different losses (CDC vs Gradient reversal vs no balancing, w/ vs w/o EMA of weights)
- (c) single-subnetwork variant of CT (EDCT) vs original CT

Results

- Combination of end-to-end three subnetworks architecture and the novel CDC loss is crucial (neither work better alone)
- Simply switching the backbone from LSTM to transformer and using gradient reversal as in CRN (Bica et al., 2020) gives worse results
- CDC loss also improves the performance of CRN

		τ:	= 1	$\tau = 6$	
		$\gamma = 1$	$\gamma = 4$	$\gamma = 1$	$\gamma = 4$
CT	(proposed)	0.80	1.32	0.63	0.93
	w/ non-trainable PE*	± 0.00	-0.02	+0.01	-0.03
a	w/ absolute PE*	+0.04	+0.16	+0.15	+1.00
	w/o attentional dropout*	± 0.00	+0.07	+0.00	+0.09
	w/o cross-attention*	+0.03	+0.16	+0.06	+0.10
	w/o EMA $(\beta = 0)^*$	+0.03	+0.38	+0.03	+0.33
b	w/o balancing ($\alpha = 0$; $\beta = 0.99$)*	-0.01	-0.02	± 0.00	+0.07
	$w/GR (\lambda = 1)$	+0.02	+0.17	+0.08	+0.33
_	EDCT w/ GR ($\lambda = 1$)	+0.16	+0.08	+0.05	+0.23
С	EDCT w/ DC ($\alpha = 0.01$; $\beta = 0.99$)	-0.03	+0.10	-0.03	+0.23

Lower = better;



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Conclusion

We proposed a novel, state-of-the-art method: the **Causal Transformer** which is designed to capture complex, long-range patient trajectories

It combines a custom subnetwork architecture to process the input together with a new counterfactual domain confusion loss for end-to-end training



Source Code: github.com/Valentyn1997/ CausalTransformer



ArXiv Paper: arxiv.org/abs/2204.07258

Extended related work

Method	Setting	Model type (backbone)	Time	Treatments	Framework
HITR (Xu et al., 2016) CGP (Schulam & Saria, 2017) MOGP (Soleimani et al., 2017) SyncTwin (Qian et al., 2021) DCRN (Berrevoets et al., 2021)	DGM (X) C, SO, SI, CSI (X) DGM (X) DGM (X) C, SO, Cov (X)	NP (GP) NP (GP) SP (GP) SP (GRU-D, LSTM) P (3 LSTMs)	Disc & Cont Cont Disc & Cont Disc Disc	Seq, Cat Seq, Cat Seq, Cont Single-time, Bin Seq, Bin	G-computation G-computation G-computation Synthetic control Disentangled representation
* MSMs (Robins et al., 2000) * RMSNs (Lim et al., 2018) * CRN (Bica et al., 2020) * G-Net (Li et al., 2021)	C, SO, SI (✔) C, SO, SI (✔) C, SO, SI (✔) C, SO, SI (✔)	P (Logistic & linear regressions) P (LSTM) P (LSTM) P (LSTM)	Disc Disc Disc Disc	Seq, Cat Seq, Cat Seq, Cat Seq, Cat	IPTW weighted loss IPTW weighted loss BR (gradient reversal) G-computation
* Causal Transformer (this paper)	C, SO, SI	P (3 transformers)	Disc	Seq, Cat	BR (CDC)

^{* =} Methods with the same assumptions as ours (and thus included in our baselines)

Legend:

- Setting: consistency (C), sequential overlap (SO), sequential ignorability (SI), sequential ignorability but conditional on covariates (Cov), continuous sequential ignorability (CSI), assumed data generating model (DGM)
 - Model: parametric (P), semi-parametric (SP), and non-parametric (NP)
 - Time: discrete (Disc) or continuous (Cont) time steps
 - Treatments: sequential (Seq), binary (Bin), categorical (Cat), continuous (Cont).
 - Framework: inverse probability of treatment weights (IPTW), balanced representations (BR)

Attention primer

1. Linear transformations:

$$egin{aligned} Q^{(i)} &= Q^{(i)}(\mathrm{H}^b) = \mathrm{H}^b \, W_Q^{(i)} + \mathbf{1} b_Q^{(i) op}, \ K^{(i)} &= K^{(i)}(\mathrm{H}^b) = \mathrm{H}^b \, W_K^{(i)} + \mathbf{1} b_K^{(i) op}, \ V^{(i)} &= V^{(i)}(\mathrm{H}^b) = \mathrm{H}^b \, W_V^{(i)} + \mathbf{1} b_V^{(i) op}, \end{aligned}$$

- 2. Attention weights and scores:
 - w/o relative positional encoding

$$\operatorname{Attn}^{(i)}(Q^{(i)}, K^{(i)}, V^{(i)}) = \operatorname{softmax}\left(\frac{Q^{(i)}K^{(i)}}{\sqrt{d_{qkv}}}\right)V^{(i)}$$

w/ relative positional encoding

$$\operatorname{Attn}^{(i)}(Q^{(i)}, K^{(i)}, V^{(i)}) = \operatorname{softmax}\left(\frac{Q^{(i)}K^{(i)\top}}{\sqrt{d_{qkv}}}\right)V^{(i)}$$

$$(\operatorname{Attn}(Q, K, V))_{i} = \sum_{j=1}^{t} \alpha_{ij}(V_{j} + a_{ij}^{V}), \quad \alpha_{ij} = \operatorname{softmax}_{j}\left(\frac{Q_{i}^{\top}(K_{j} + a_{ij}^{K})}{\sqrt{d_{qkv}}}\right)$$

3. Multi-head attention:

$$MHA(Q, K, V) = Concat(Attn^{(1)}, \dots, Attn^{(n_h)})$$

Encoder-Decoder Causal Transformer: Architecture

Two separate transformers, i.e., encoder and decoder, for each task of one- and multiple step ahead predictions

