Sequential Pattern-based Anomaly Detection for Enhanced Interpretability



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 - Co-chair, AIME 2020
- Triathlete in the mornings ..











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- Collaborations: Columbia University, Maccabi Healthcare Services, AIIMS/IIT New Delhi, Peking University, UTHealth, UPenn/CHOP and more
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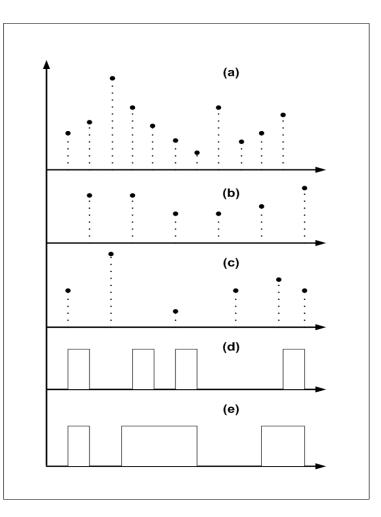


Funders and Collaborators

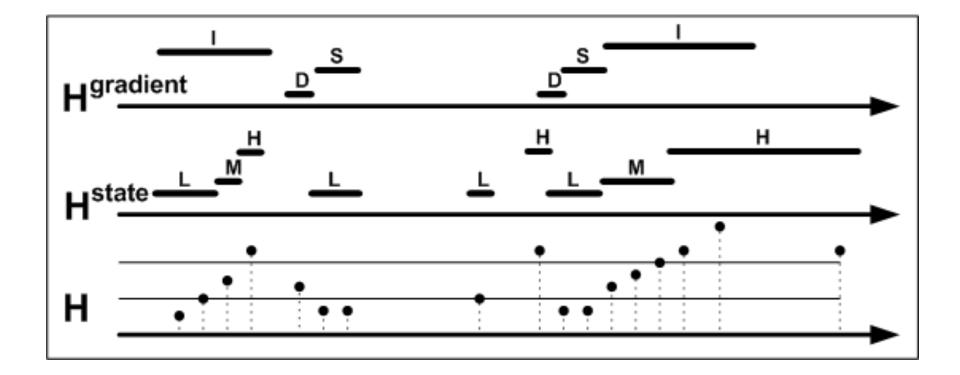




Temporal Raw Data



From Time Points to Time Intervals Series



Robert Moskovitch, Yuval Shahar, Classification Driven Temporal Discretization of Multivariate Time Series, *Data Mining and Knowledge Discovery*, 29, 4, 871-913, 2015.

Motivation

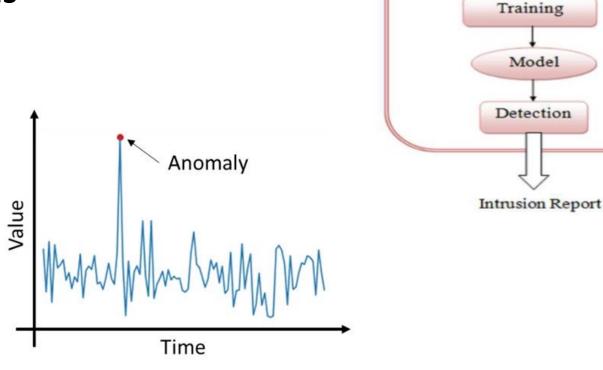
- Industrial Control Systems (ICS) are essential networks of hardware and software that manage industrial processes, such as water purification plants.
- Increased internet connectivity exposes ICS to cyber threats, highlighting the need for robust security measures and advanced anomaly detection techniques.



Anomaly Detection

Anomaly detection is a field of study that focuses on identifying unusual or abnormal behavior in data.

- Statistical methods
- Machine learning algorithms
- Data mining techniques
- Rule-based methods



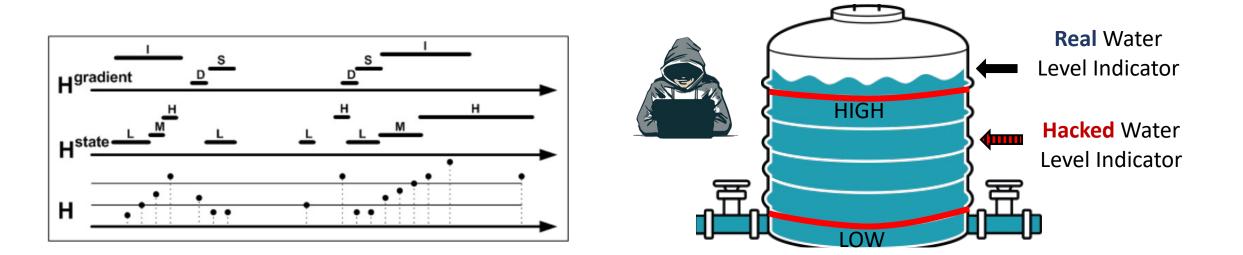
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Monitored Environment

Parameterization

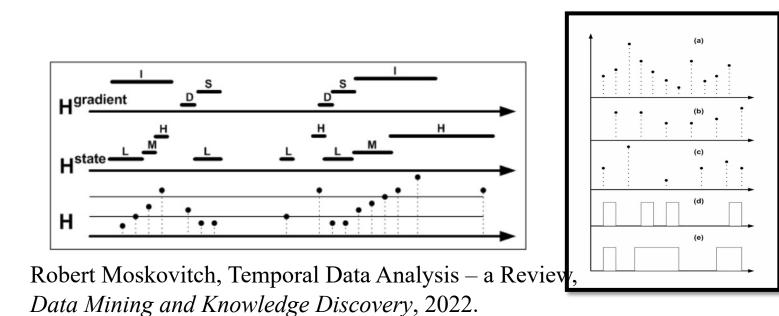
Research Goals

Objective 1: Develop an anomaly detection model using **sequential patterns**. **Objective 2**: Develop an **interpretable** algorithm for cyber security experts in ICS.



Pros of Temporal Abstraction

- Handle missing values robust to gaps in data.
- Handle irregular sampling unifies inconsistent data collection intervals.
- Generalization detects patterns across varying time intervals.
- Interpretability offer clearer interpretation than continuous data.



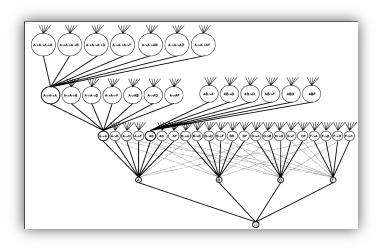
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 8	 	 	_	_
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 9	 	 		
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low	 	 		
Patient 10			10	_

Sequential Pattern Mining

- Database consists of ordered events, in which each event is 1 time-unit long.
- <u>Apriori property</u>: If a sequence is infrequent, then all its super-sequences must also be infrequent.

Example: < e a > is infrequent, then

< e a b > must be infrequent as well.



SID	Sequences		
10	<badc></badc>		
20	< (bd) a (de) c b >		
30	< a d c b >		
40	< a d (be) c >		
50	< c d b c a b >		

Sequential Pattern Mining - Example

SID	Sequences
10	< b a c >
20	< (bd) a (de) c b >
30	< a d c (bc) >
40	< a d (bc) c >
50	< c (bc) c a b >

minimum support =



Frequent Patterns	Support
< a >	5
< a b >	4
< a c >	4
< b >	5
< (bc) >	4
< b c >	4
< c >	5
< c c >	4

Sequential Pattern Mining

Real example of pattern:

"Water tank fills, then empties"

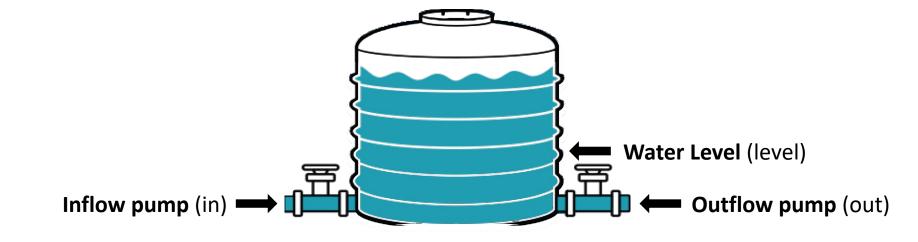
<u>3 sensors</u>: **inflow pump** - in, **water level** - level, **outflow pump** – out <u>Abstractions Levels</u>: **High** – **H**, **Low** - **L**

< (in_H, level_L, out_L) (in_H, level_H, out_L) (in_L, level_H, out_H) >

"Tank Filling Up"

"Tank Full"

"Tank Draining Down"



Sequential Pattern - definitions

Horizontal Support (HS)

HS of a pattern P in sequence S – The number of times pattern P is contained in S

SID	Sequences	
10	<bac></bac>	
20	< (b ,d) a (d,e) c b >	HS(< b >, 20) = 2
30	< a d c b >	HS(< a >, 30) = 1
40	< a d (b,e) c >	
50	<cbcabc></cbcabc>	HS(< c c >, 50) = 3
		< c b c a b c > < c b c a b c > < c b c a b c >

Sequential Pattern - definitions

Mean Duration (MD)

MD of a pattern P in sequence S – Sum of time spans for each P instance in S, divided by P's

occurrences.

SID	Sequences	
10	<bac></bac>	
20	< (b ,d) a (d,e) c b >	MD(< b >, 20) = $\frac{1+1}{2} = \frac{2}{2} = 1$
30	< a d c b >	
40	< a d (b,e) c >	
50	<cbcabc></cbcabc>	$ MD(, 50) = \frac{3+4+6}{3} = \frac{13}{3} = 4.33 $
		< c b c a b c >
		< c b c a b c >
		< c b c a b c >

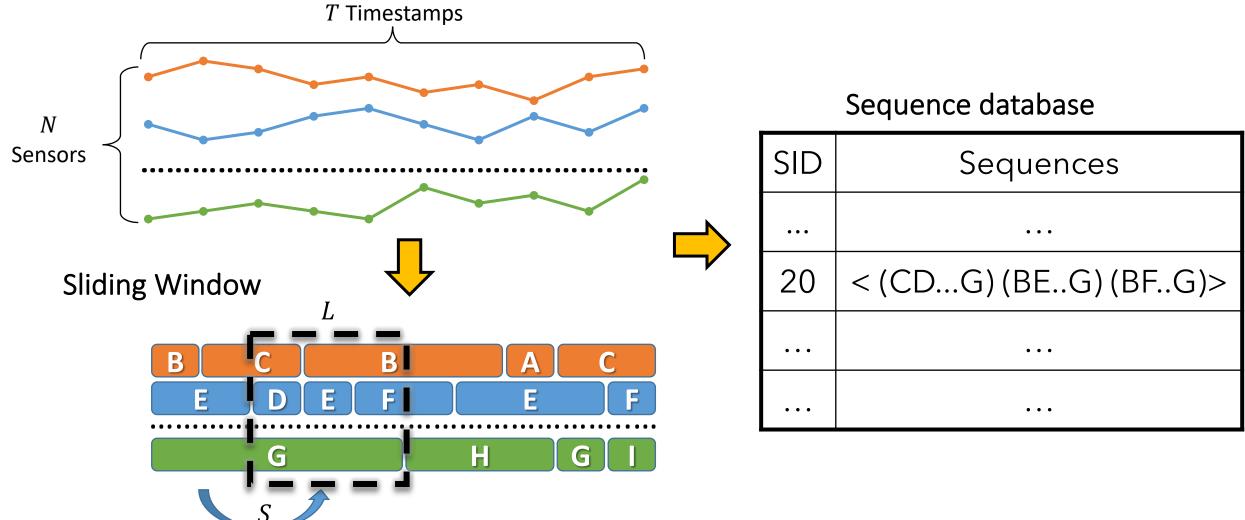
Attack Periods Anomaly Detection Hypothesis

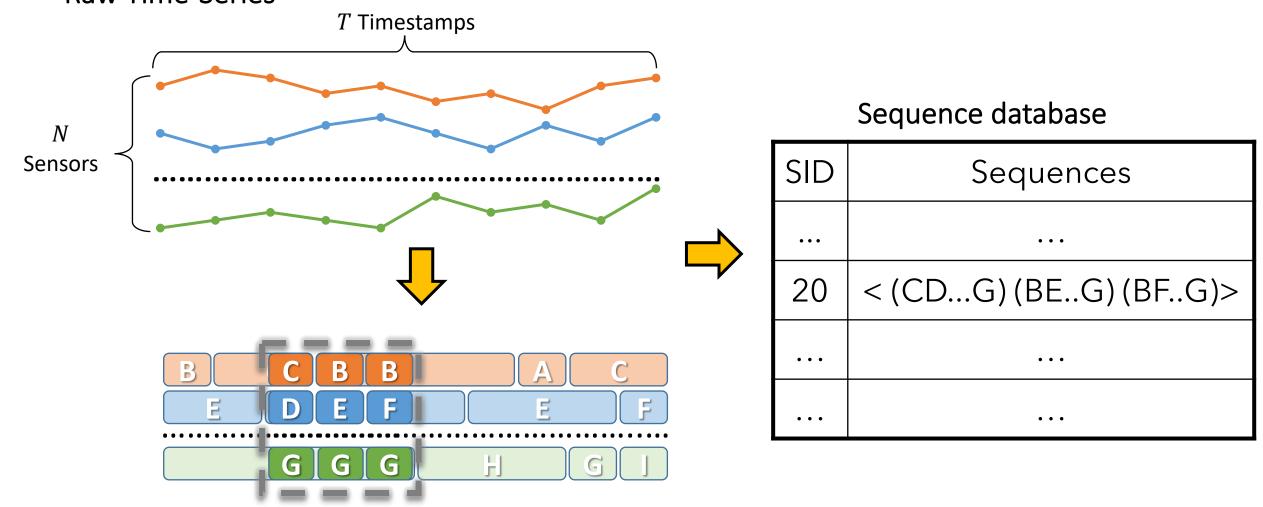
Since attacks can be seen as **anomalous periods** of sensor data, sequential patterns that are considered **frequent in mostly normal periods** will appear as **missing or have different properties**

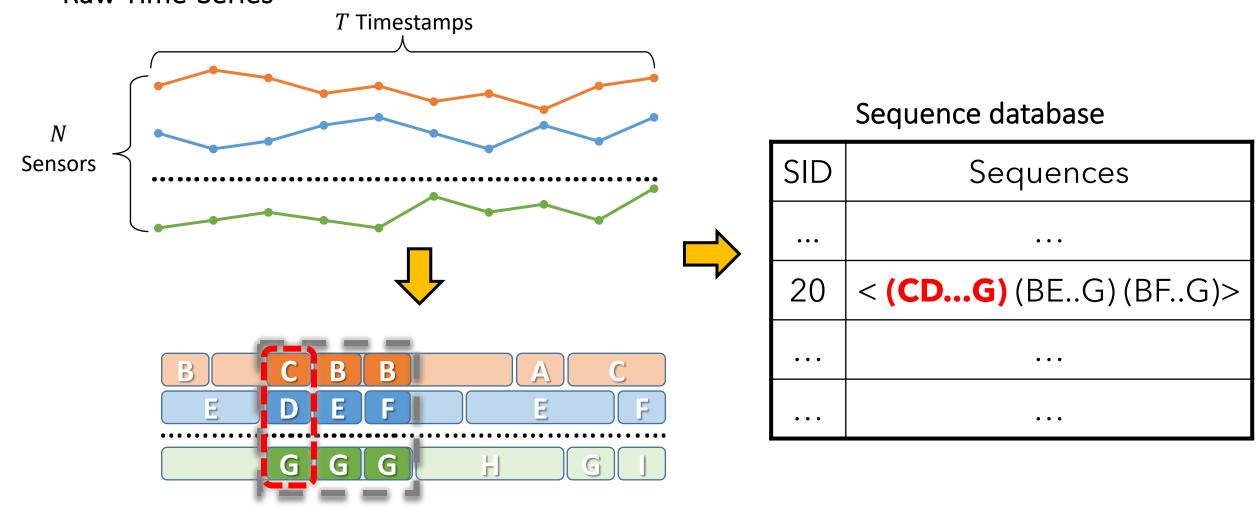
Motivation

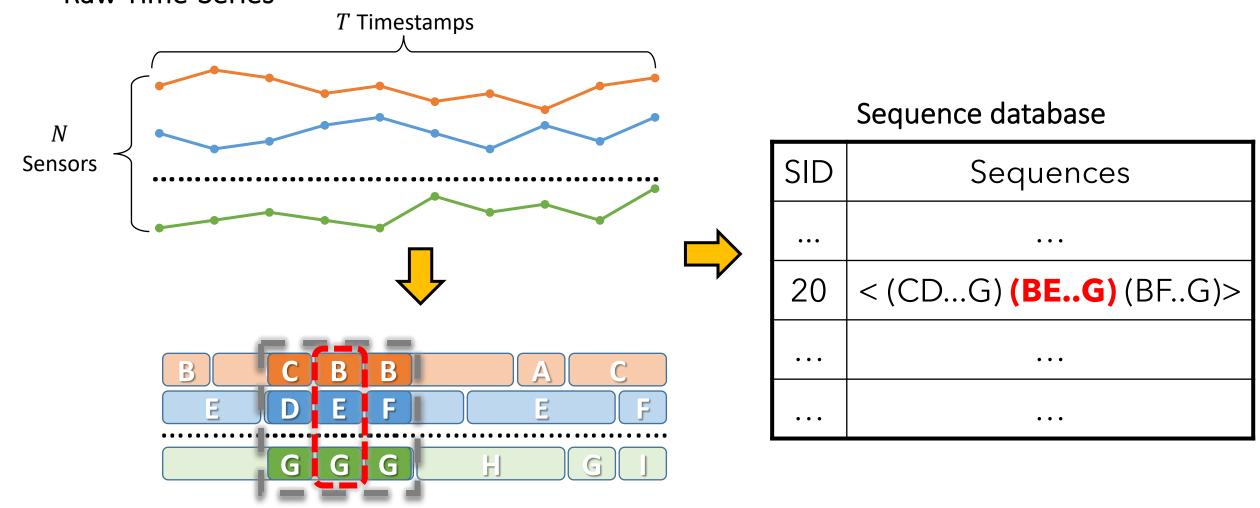
- ✓ Background
- Proposed Method

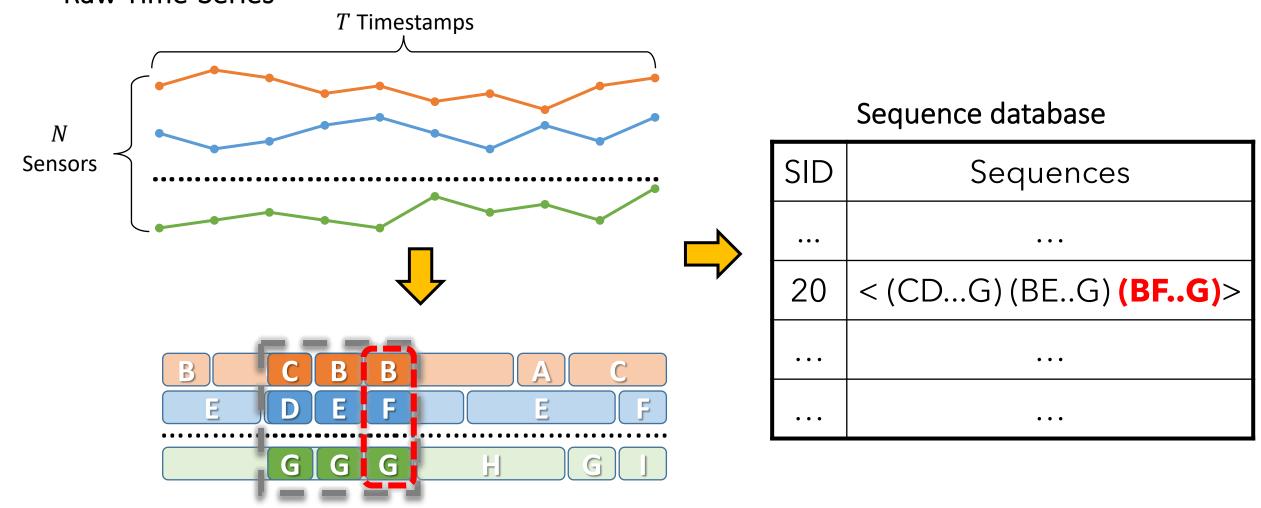
Proposed Method





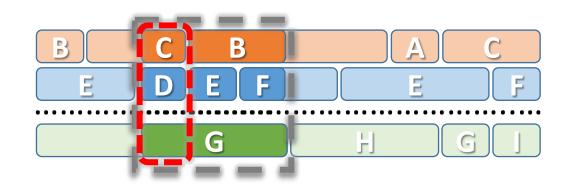






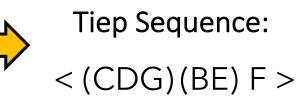
We also introduce and evaluate a more efficient approach for sequence extraction - "Tiep" (Time Interval Endpoint).

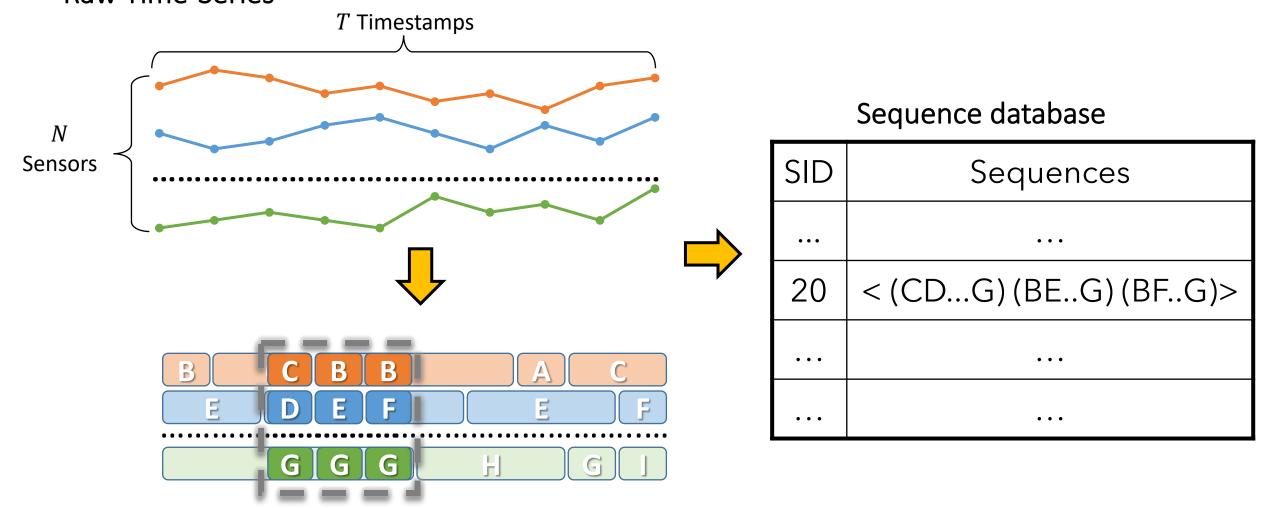
- Aims to mitigate scalability issues in high-resolution data.
- Unlike traditional methods, focuses only on starting time points.
- Significantly reducing computational overhead and memory usage.



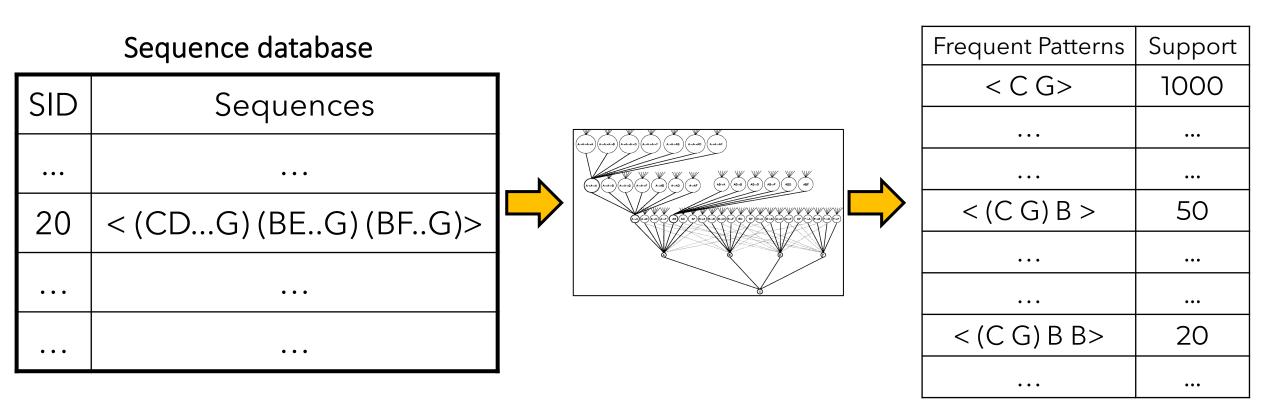
Full Original Sequence:

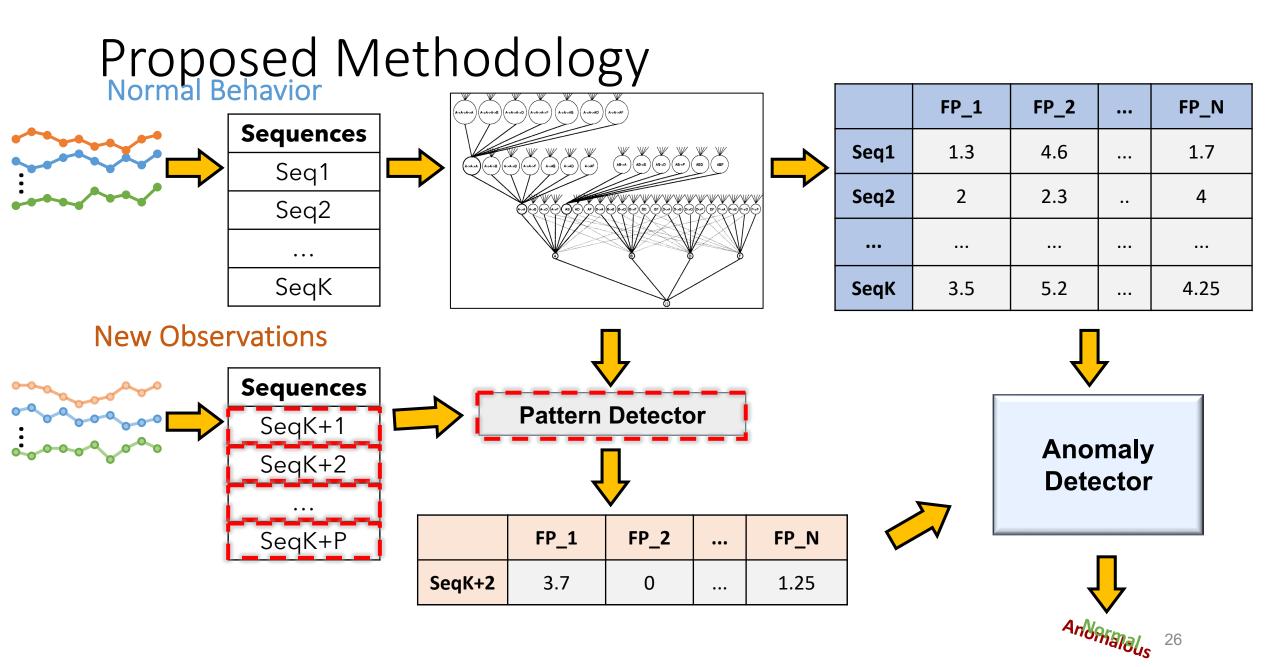
< (CDG) (BEG) (BFG)>





Sequential Pattern Mining





- ✓ Motivation
- ✓ Background
- ✓ Proposed Method
- Evaluation

Evaluation

Dataset

Secure Water Treatment (SWaT)

- Source: iTrust, Singapore University of Technology and Design
- URL: <u>https://itrust.sutd.edu.sg/itrust-labs_datasets/</u>
- **Objective**: Evaluate anomaly detection in industrial control systems

Divided into two parts: (1) Training set and (2) Test set

- Training set: 7 days of normal operation data
- **Test set**: 4 days of data with 36 injected cyber-physical attacks

Timestamp	FIT101	LIT101	AIT201	MV101	P101	P102	AIT202
22/12/2015 4:00:00 PM	2.470294	261.5804	244.3284	2	2	1	8.19008
22/12/2015 4:00:01 PM	2.457163	261.1879	244.3284	2	2	1	8.19008
22/12/2015 4:00:02 PM	2.439548	260.9131	244.3284	2	2	1	8.19008
22/12/2015 4:00:03 PM	2.428338	260.285	244.3284	2	2	1	8.19008
22/12/2015 4:00:04 PM	2.424815	259.8925	244.4245	2	2	1	8.19008
22/12/2015 4:00:05 PM	2.425456	260.0495	244.5847	2	2	1	8.19008
22/12/2015 4:00:06 PM	2.472857	260.2065	244.5847	2	2	1	8.19008
22/12/2015 4:00:07 PM	2.513532	260.5991	244.5847	2	2	1	8.19008
22/12/2015 4:00:08 PM	2.559972	261.0309	244.5847	2	2	1	8.19008
22/12/2015 4:00:09 PM	2.598085	261.1093	244.809	2	2	1	8.19008
22/12/2015 4:00:10 PM	2.630753	261.7766	244.809	2	2	1	8.19008
22/12/2015 4:00:11 PM	2.649329	261.7766	244.809	2	2	1	8.19008
22/12/2015 4:00:12 PM	2.654133	261.8944	244.8731	2	2	1	8.19008
22/12/2015 4:00:13 PM	2.646446	261.6589	244.8731	2	2	1	8.19008
22/12/2015 4:00:14 PM	2.625949	261.2664	245.0333	2	2	1	8.19008
22/12/2015 4:00:15 PM	2.61602	260.8346	245.0333	2	2	1	8.19008
22/12/2015 4:00:16 PM	2.609935	261.0309	245.0333	2	2	1	8.19008
22/12/2015 4:00:17 PM	2.602889	261.1093	245.0333	2	2	1	8.19008
22/12/2015 4:00:18 PM	2.587516	260.9916	245.0333	2	2	1	8.19008
22/12/2015 4:00:19 PM	2.573103	261.3056	245.0333	2	2	1	8.19008
22/12/2015 4:00:20 PM	2.556769	261.6589	245.4499	2	2	1	8.19008

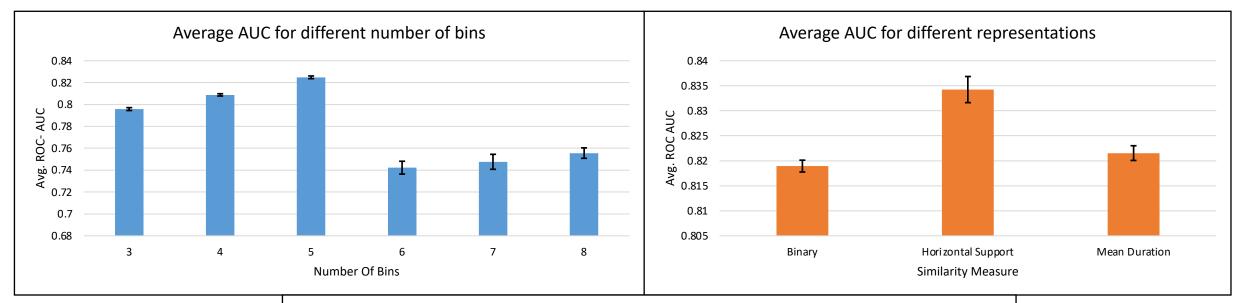
Research Questions

- 1. Which **number of bins** has the best performance in abstracting our data?
- 2. Which **metric** has the most effective performance between frequent patterns and transactions, such as Binary, Horizontal Support, and Mean Duration?
- 3. Which state-of-the-art **anomaly detector**, including OneClassSVM and SGDOneClassSVM, achieves optimal generalization on our data and has the best performance when utilizing them?

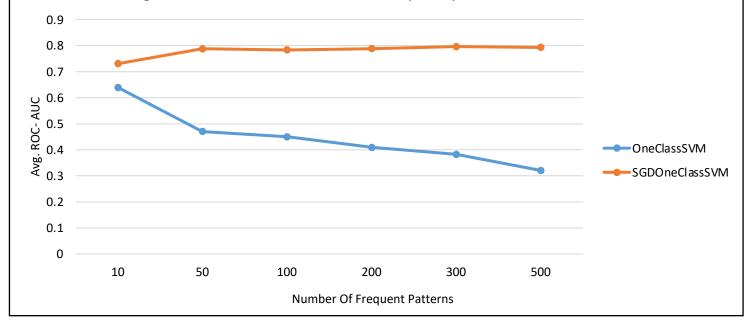
- ✓ Motivation
- ✓ Background
- ✓ Proposed Method
- ✓ Evaluation
- Initial Results

Initial Results

Key Results



Average AUC for different number of frequent patterns for each Classifier



Overall Results

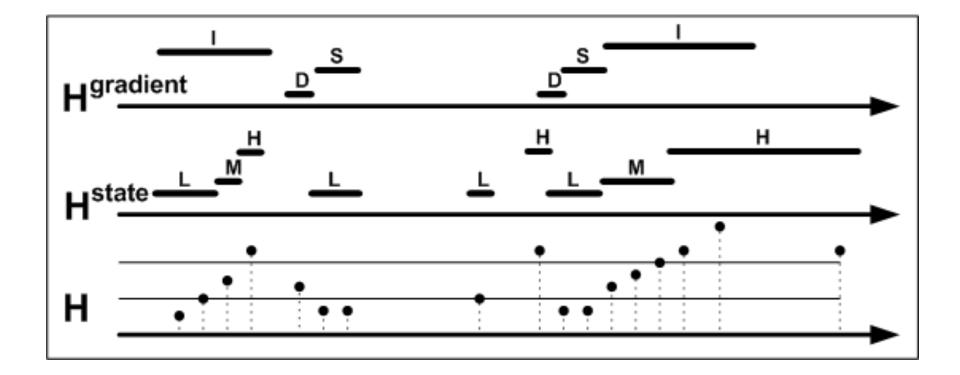
Method	SWaT		* P-Preci	* P-Precision, R-Recall		
	P *	R *	AUC	F1		
MERLIN	0.656	0.2547	0.6175	0.3669		
LSTM-NDT	0.7778	0.5109	0.714	0.6167		
DAGMM	0.9933	0.6879	0.8436	0.8128		
OmniAnomaly	0.9782	0.6957	0.8467	0.8131		
MAD-GAN	0.9593	0.6957	0.8463	0.8065		
USAD	0.9977	0.6879	0.846	0.8143		
MTAD-GAT	0.9718	0.6957	0.8464	0.8109		
CAE-M	0.9697	0.6957	0.8464	0.8101		
GDN	0.9591	0.6957	0.8462	0.8101		
GRN-50	0.9972	0.5921	0.8781	0.7389		
GRN-100	0.9986	0.5909	0.8845	0.7496		
Our best algorithm:	0.998	0.583	0.8646	0.736		

- Motivation
- ✓ Background
- ✓ Proposed Method
- ✓ Evaluation
- ✓ Initial Results
- Future Research Directions

Future Research Directions

More datasets Evaluation with TIEPs Explainability Using TIRPs Using TIRPs with Similarity

From Time Points to Time Intervals Series



Robert Moskovitch, Yuval Shahar, Classification Driven Temporal Discretization of Multivariate Time Series, *Data Mining and Knowledge Discovery*, 29, 4, 871-913, 2015.

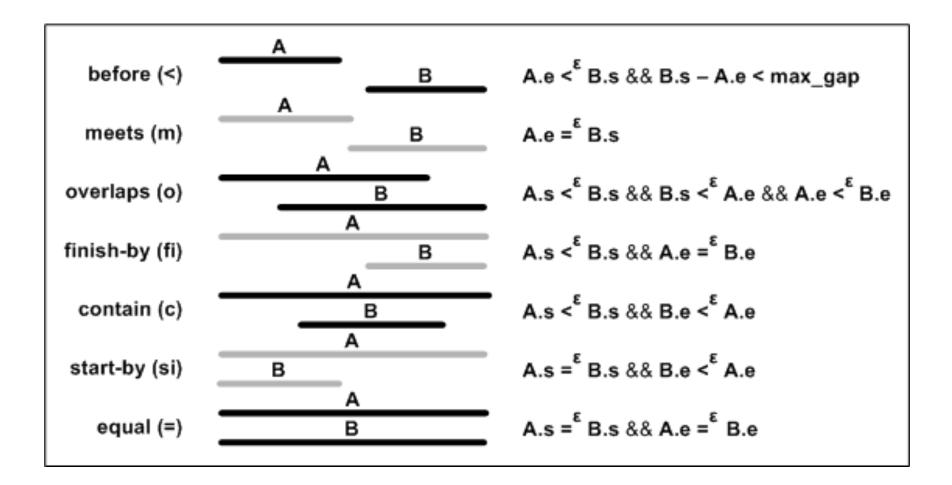
Time Intervals Related Patterns Discovery

HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 8	
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 9	
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 10	

Time Intervals Related Patterns Discovery

HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 8		
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 9		
HA1c.High Insulin.High Glucose.Norm Cholesterol.Low statines.Low Patient 10		

Allen's (1983) Temporal Logic



Time Intervals Related Pattern - TIRP

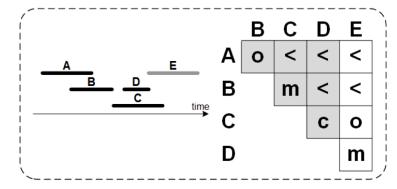
A TIRP is a conjunction of pairwise temporal relations

{A o B, A < C, As < D, A < E, B m C, B < D, B < E, C c D, C o E, D m E}

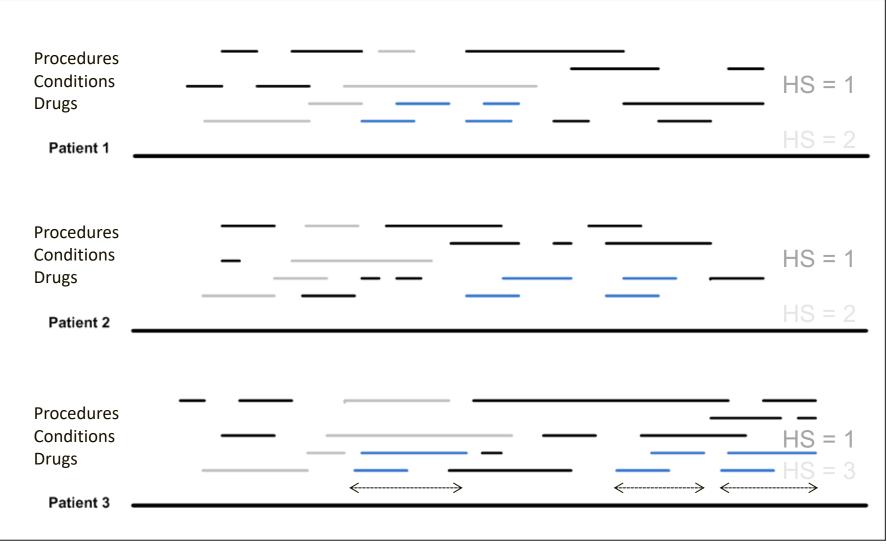
A k-sized TIRP includes $k(k-1)/2 = (k^2-k)/2$ temporal relations

TIRPs have several metrics, which can be predictive

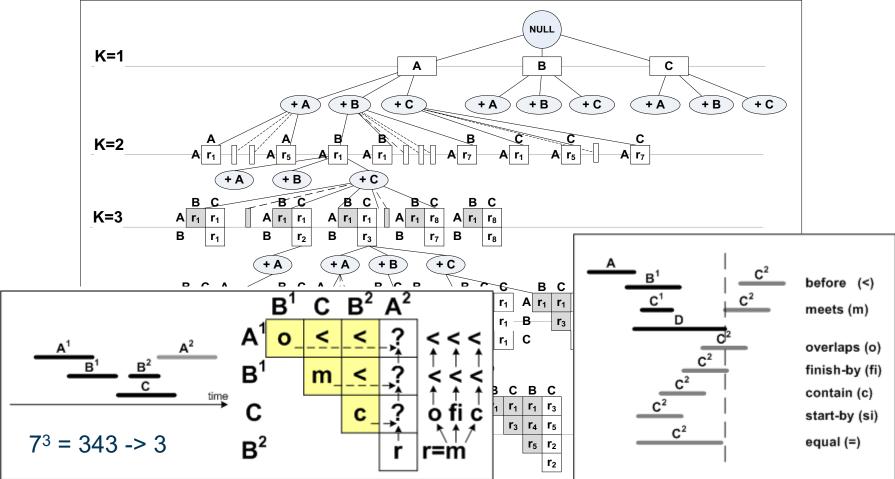
- Vertical Support how many patients have the TIRP in the database
- Horizontal Support how many instances (episodes) of the TIRP were in the past hours (or weeks)
- Mean Duration what is the average duration of these instances



Time Intervals Related Patterns Discovery – an illustration

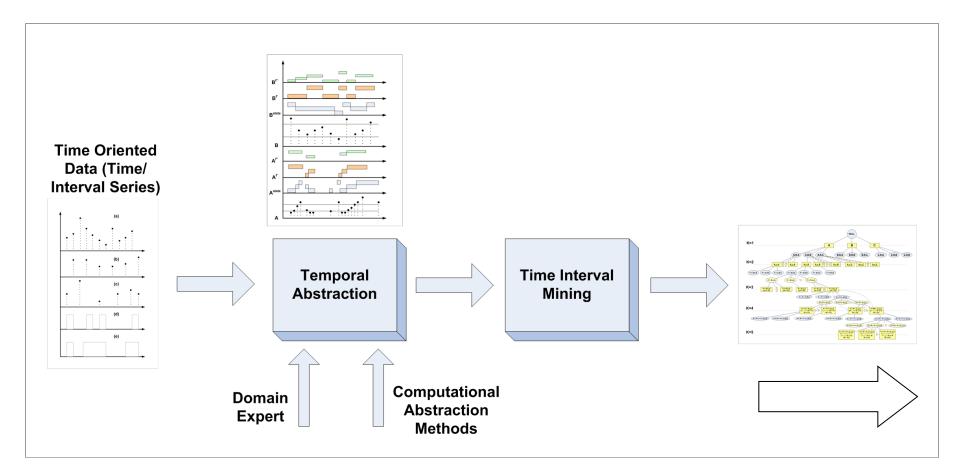


KarmaLego Fast Time Intervals Mining



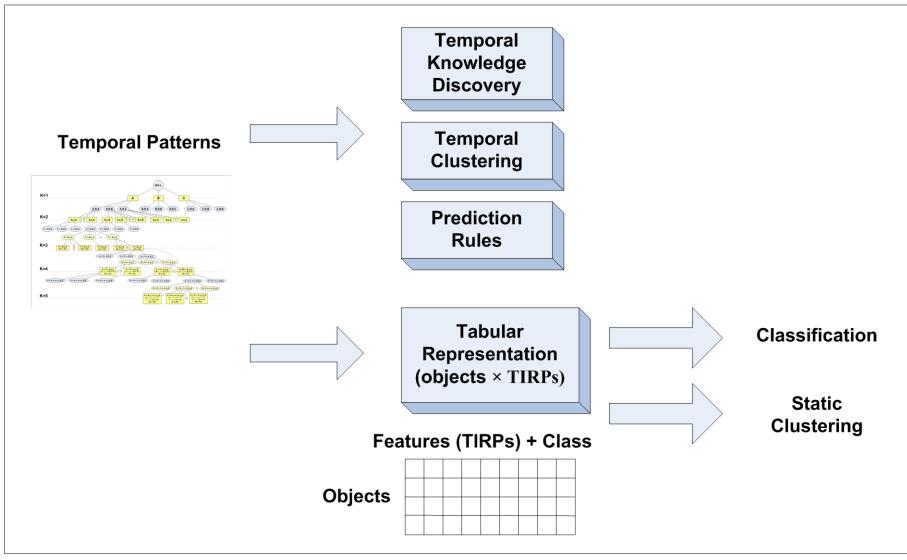
Moskovitch et al, Outcomes Prediction via Time Intervals Related Patterns, IEEE International Conference on Data Mining (ICDM), 2015

KarmaLego General Workflow



Moskovitch et al, Fast Time Intervals Mining by Exploiting the Transitivity of Temporal Relations, Knowledge and Information Systems, 2015.

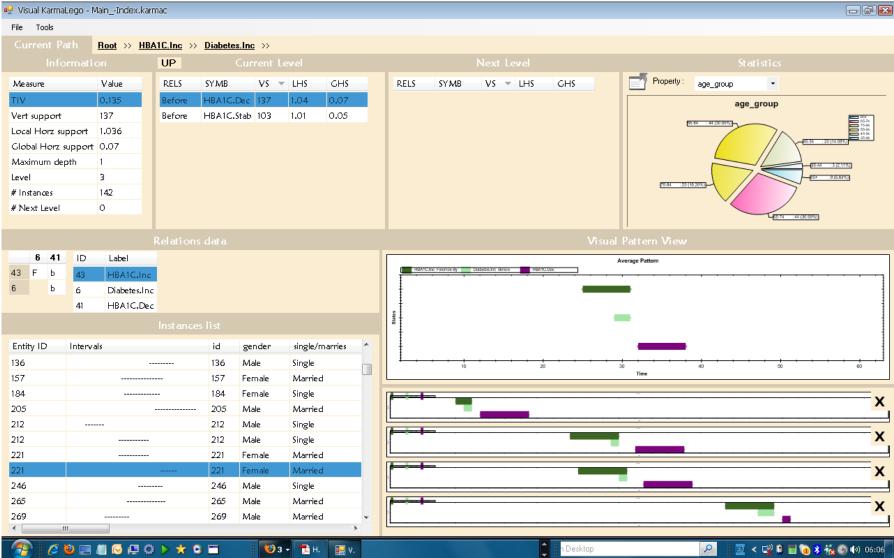
Use of TIRPs from KarmaLego



KarmaLegoV

🖳 Visual KarmaLeo	go - MainIndex.kar	rmac											
File Tools													
Current Path	h <u>Root</u> >> <u>HB</u>	A1C.Inc >>											
Inform						St at i st i cs							
Measure	Value	RELS	SYMB		VS LH	s ghs	•	RELS	SY MB	VS	LHS	GHS	Property : gender
TIV	0.098	Before	Diabetes.De	c 4	489 1.19	9 0.29		Before	HBA1C.Dec	: 137	1.04	0.07	gender
Vert support	221	Meets	Diabetes.De	ec 2	260 1.0	7 0.14		Before	HBA1C.Stat	5 103	1.01	0.05	T Grade
Local Horz supp	oort 1.054	Overlaps	Diabetes.Inc	c 2	285 1.0	5 0.15							Alale 124 (53 22'5)
Global Horz sup	oport 0.114	Contains	Diabetes.Inc	c 5	569 1.4	6 0.41							
Maximum depth	n 2	Meets	Diabetes.Inc	c 2	276 1.0	7 0.14							
Level	2	Before	Diabetes.Inc	c 5	504 1.18	8 0.29							
# Instances	233	Finishes By	Diabetes.Inc	c 2	221 1.0	5 0.11							Emain : 156 (46.78%)
# Next Level	2	Starts	Diabetes.Lo	w 3	319 1	0.16							
		Before	Diabetes.Lo	w 1	32 1.11	0.07	v						
												Visual Patt	tern View
43 F 43 6											Pattern		
Entity ID In	ntervals		id ge	nder	single/ma	rries 🔷							
205			205 Ma	ale	Married								-
212			212 Ma	ale	Single				10		20		30 40 50
212			212 Ma	ale	Single								
221			221 Fer	male	Married			-					X
221			221 Fer	male	Married		۱ <u>ــــــــــــــــــــــــــــــــــــ</u>						
246			246 Ma	ale	Single							-	X
265			265 Ma	ale	Married		1						
269			269 Ma	ale	Married			-					
286			286 Ma	ale	Married		-						X
291				male	Married								
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310			310 Fer	male	Single	• •			•				
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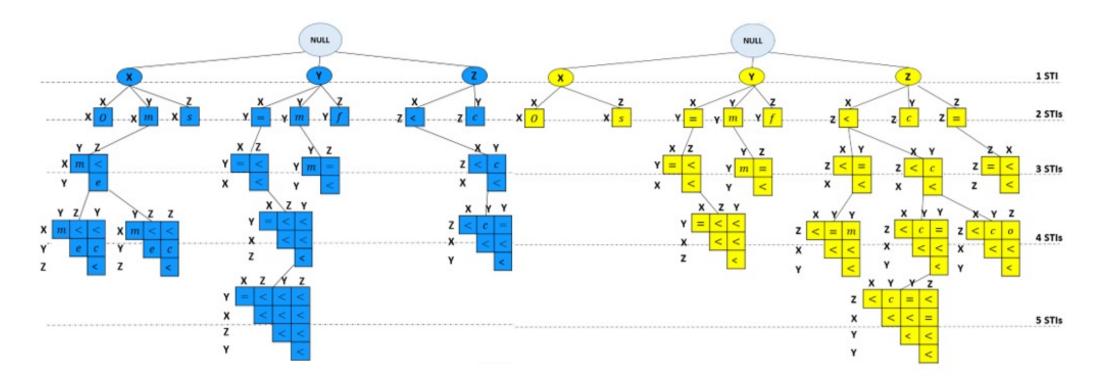
KarmaLegoV



KarmaLegoV – Search Results

ID 42 43 44 45 46 47 48	Propety HBA1C.Stab HBA1C.Inc LDL.Lev1 LDL.Lev2 LDL.Lev3 LDL.Lev4 LDL.Dec	1 C 2 C 4 C 5 C 6 C 7 C	Property A Diabetes.Low Diabetes.Med Diabetes.Dec Diabetes.Stab Diabetes.Low Diabet	ID Propert 33 HBA1C 40 HBA1C 41 HBA1C 42 HBA1C 43 HBA1C 44 LDLLe 45 LDLLe	Lev3 Lev4 Dec Stab	Limit V Supp From 0	To 0		Average Pattern Crowster Stat Superson to Growster Stat Superson Converses Stat Superson Con		Search Close elect Index ave Results
Level	Property #1	Property #2	Property #3	Property #4	Property #5	Property #6	Property #7	Property #8	Relations	H Support	V Supp
5	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Cholesterol.Stab	Glucose.Stab	HBA1C.Dec			e.b.b.b.b.e.b.b.e.e.b.b.b.b.b.b.	148	0.07
•	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Cholesterol.Stab	Glucose.Stab	Creatinine.Stab	HBA1C.Dec		e.b.b.b.b.e.b.b.e.e.b.b.e.e.e.b.b.b.b.b	118	0.06
i	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Cholesterol.Stab	Creatinine.Stab	HBA1C.Dec			e.b.b.b.b.e.b.b.e.e.b.b.b.b.b.	131	0.06
	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Glucose.Stab	HBA1C.Dec				e.b.b.b.b.e.b.b.b.b.	159	0.08
	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Glucose.Stab	Creatinine.Stab	HBA1C.Dec			e.b.b.b.b.e.b.b.e.e.b.b.b.b.b.	122	0.06
;	HBA1C.Inc	Cholesterol.Inc	HBA1C.Stab	Creatinine.Stab	HBA1C.Dec				e.b.b.b.b.e.b.b.b.b.	138	0.07
	HBA1C.Inc	Cholesterol.Inc	LDL.Stab	HBA1C.Dec					e.b.b.b.b.b.	133	0.06
;	HBA1C.Inc	Cholesterol.Inc	LDL.Stab	Cholesterol.Stab	HBA1C.Dec				e.b.b.b.b.e.b.b.b.b.	120	0.06
Ļ	HBA1C.Inc	Cholesterol.Inc	Glucose.Stab	HBA1C.Dec					e.b.b.b.b.b.	173	0.08
	HBA1C.Inc	Cholesterol.Inc	Glucose.Stab	Creatinine.Stab	HBA1C.Dec				e.b.b.b.b.e.b.b.b.b.	130	0.06
Ļ	HBA1C.Inc	Cholesterol.Inc	Glucoæ.Inc	HBA1C.Dec					e.e.e.b.b.b.	147	0.07
	HBA1C.Inc	Cholesterol.Inc	Glucoæ.Inc	HBA1C.Stab	Cholesterol.Stab	HBA1C.Dec			e.e.e.b.b.b.b.b.b.e.b.b.b.b.b.	122	0.06
	HBA1C.Inc	Cholesterol.Inc	Glucose.Inc	HBA1C.Stab	Cholesterol.Stab	Glucose.Stab	HBA1C.Dec		e.e.e.b.b.b.b.b.b.e.b.b.b.e.e.b.b.b.b.b	106	0.05
	HBA1C.Inc	Cholesterol.Inc	Glucoæ.Inc	HBA1C.Stab	Glucose.Stab	HBA1C.Dec			e.e.e.b.b.b.b.b.b.e.b.b.b.b.b.	113	0.05
	HBA1C.Inc	Cholesterol.Inc	Glucoæ.Inc	Cholesterol.Stab	HBA1C.Dec				e.e.e.b.b.b.b.b.b.b.	131	0.06
	HBA1C.Inc	Cholesterol.Inc	Glucoæ.Inc	Cholesterol.Stab	Glucose.Stab	HBA1C.Dec			e.e.e.b.b.b.b.b.b.e.b.b.b.b.b.	111	0.05
	HBA1C.Inc	Cholesterol.Inc	Glucose.Inc	Creatinine.Stab	HBA1C.Dec				e.e.e.b.b.b.b.b.b.b.	104	0.05
	HBA1C.Inc	Cholesterol.Inc	Creatinine.Stab	HBA1C.Dec					e.b.b.b.b.b.	163	0.08
	HBA1C.Inc	Glucose.Dec	HBA1C.Dec						b.b.s.	248	0.11
	HBA1C.Inc	Glucose.Dec	HBA1C.Dec						F.b.b.	106	0.05
	nd, 1 Selected					III					•

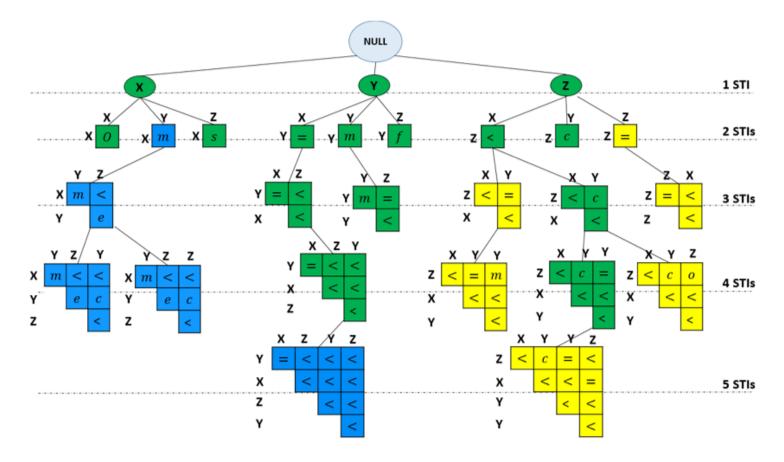
Visualization of Predictive TIRPs in Two Populations



a. Population 1 TIRPs Tree

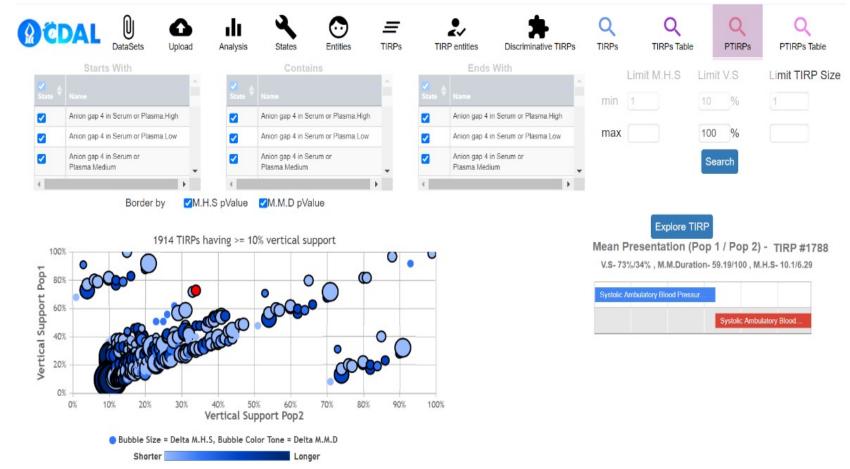
b. Population 2 TIRPs Tree

Visualization of Predictive TIRPs in Two Populations



c. The Populations Unified TIRPs Tree

Visualization of Predictive TIRPs in Two Populations

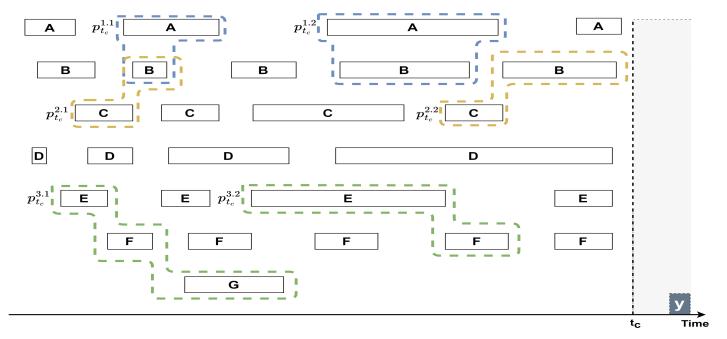


Guy Shitrit, Noam Tractinsky, Robert Moskovitch, Visualization of Frequent Temporal Patterns in Single or Two Populations, *Journal of Biomedical Informatics*, 2022.

Continuous event's prediction via TIRPs



Detected TIRPs Instances



Itzhak et al, Continuous Prediction of Temporal Pattern Completion, PAKDD 2023

Next Directions with TIRPs Similarity

- The idea is to develop a similarity function for a TIRP, based on their time intervals duration, and temporal relations
- Use that for Anomaly Detection
- Use that for COPE identification

Time Intervals Related Pattern - TIRP

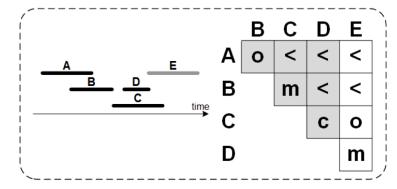
A TIRP is a conjunction of pairwise temporal relations

{A o B, A < C, As < D, A < E, B m C, B < D, B < E, C c D, C o E, D m E}

A k-sized TIRP includes $k(k-1)/2 = (k^2-k)/2$ temporal relations

TIRPs have several metrics, which can be predictive

- Vertical Support how many patients have the TIRP in the database
- Horizontal Support how many instances (episodes) of the TIRP were in the past hours (or weeks)
- Mean Duration what is the average duration of these instances

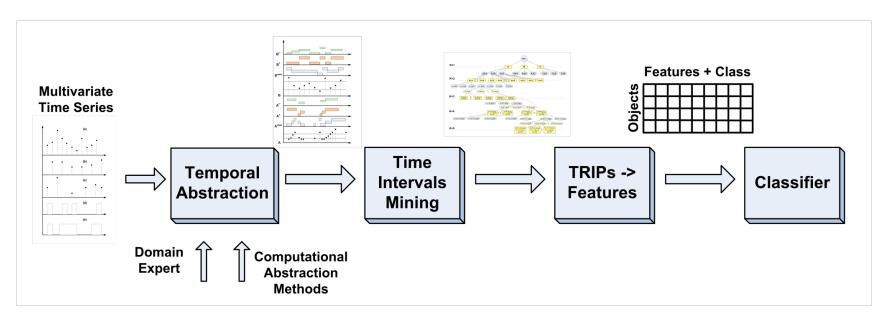


Using Similarity for Anaomaly Detection, or COPE ideantification

- Using the similarity function for Anomaly Detection
 - How the patterns in the data are similar or anomaluous?
 - Not similar -> anomalous, and how? Using a threshold
 - Include explainability which patterns, and how different?
- Using the similarity function for COPE Identification
 - After having a TIRP that idenitifies a COPE (system situation)
 - How similar is it?
 - Or how different than common appearance
 - – will give much more granular detection framework

KarmaLegoSification

- A major problem in multivariate time series classification is the various types of raw temporal data
- TIRPs can be useful here as well, as classification features

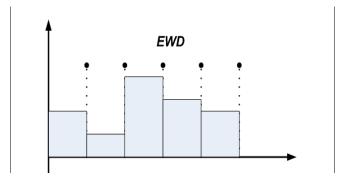


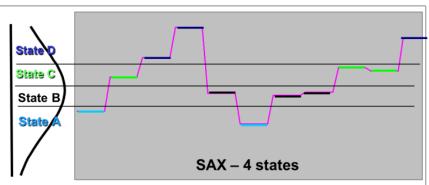
Moskovitch et al., Data Mining and Knowledge Discovery, 2015

Unsupervised Discretization: EWD and SAX

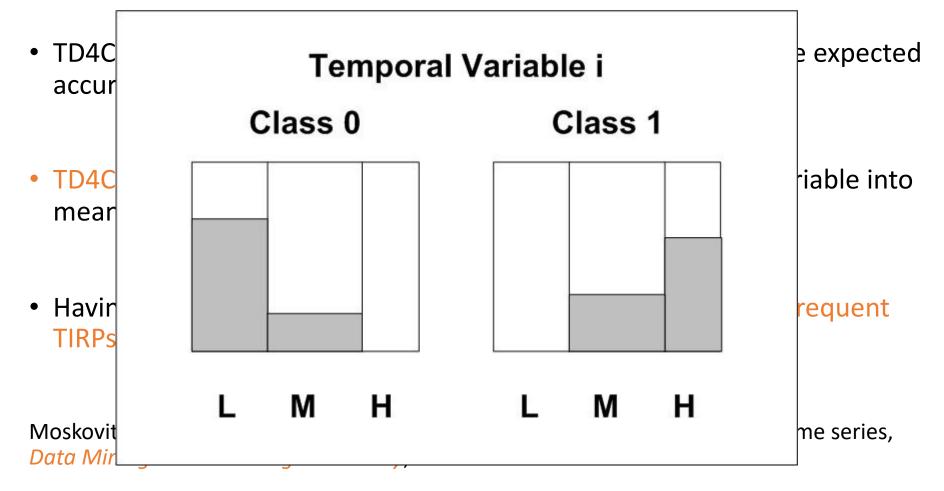
EWD the continuous values range is divided into k equal bins (states)

Symbolic Aggregate approXimation Based on PAA, Piecewise Aggregate Approximation, A time series segmenting algorithm (Keogh et al.,2003).





TD4C - Temporal Discretization for Classification



TD4C Formulation

- Given C = {c₁, c₂,.. c_n} classes, E entities divided into {E₁, E₂,.. E_c} sets of entities per class and T = {t₁, t₂,.. t_m} temporal variables, and A a TD4C abstraction method.
- The problem is to find the set of cutoffs for each temporal variable t_i that increases the difference in the dominant states in each class.
- Thus, we want to measure the distribution of the states in each class entities, and to measure when these are most different.
- > For that three measure were determined:

• Entropy
• Entropy

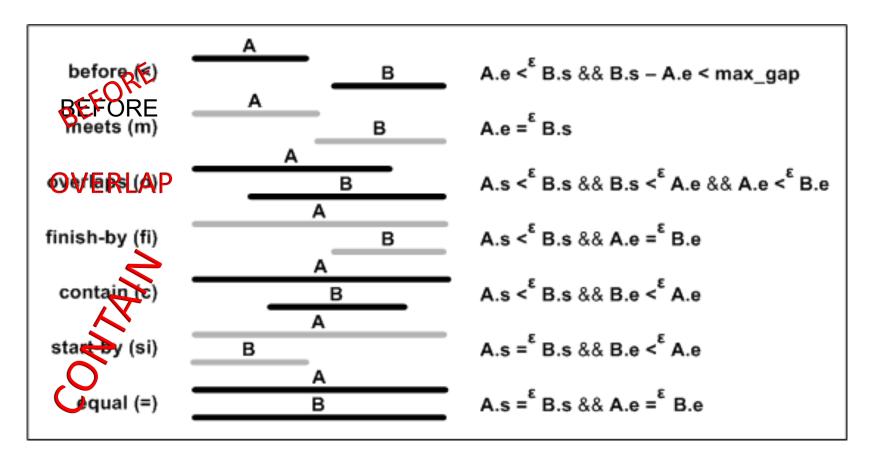
$$E(c) = -\sum_{i=1}^{k} p_i \cdot \log(p_i) \qquad D = \sum_{i=1}^{c} \sum_{j=i+1}^{c} |E(c_i) - E(c_j)|$$
• Cosinus

$$similarity(v, u) = \frac{v \cdot u}{\|v\|\|u\|} \qquad D = \sum_{i=1}^{c} \sum_{j=1}^{c} similarity(c_i, c_j)$$
• Kullback-Leibler

$$KL(P, Q) = \sum_{i=1}^{k} p_i \log\left(\frac{p_i}{q_i}\right) \qquad D = \sum_{i=1}^{c} \sum_{j=i+1}^{c} KL(c_i, c_j)$$

Generalization of Allen's Temporal Relations

KarmaLegoS enables to mine TIRPs with 7 (Allen's original) or 3 more general temporal relations



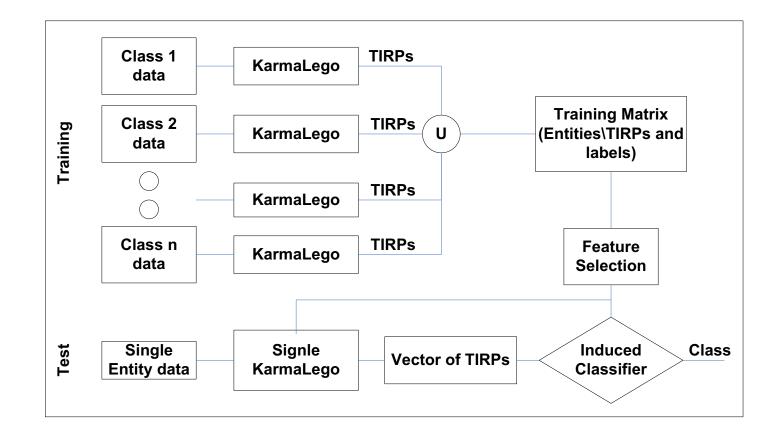


entity	Tirp ₁	Tirp ₂	Tirp ₃	Tirp ₄			Tirp n	Class
P ₁	0	2 .3	3.4	1.2	0	0	0	0
P <u>2</u>	0	0	1.23	3 .56	2 .54	1.23	3 .3	1
P ₃	0	0	2 .56	3 .34	7	0	0	0
₽4	1 .6	3 .34	1 .45	2 .7	0	0	0	0
P ₅	2 .2	2 .34	0	0	0	0	2 .4	1
P ₆	1.2	2 .5	0	1 .56	0	0	1 .34	1
Pm	1.8	0	0	2 .23	0	0	0	1

Datasets

- ICU Dataset 645 patients who underwent cardiac surgery at the AMC in Amsterdam (2002-2004). Includes over 12 hours of *High* and *Low frequency*. 196 patients were mechanically ventilated for more than 24 hours (70%), and the rest were 449.
- Diabetes- Contains 2038 diabetic patients data along 5 years (2002-2007) from Israeli HMO, measured monthly HbA1c, Glucose, Cholesterol values and medication purchased.
 992 males and 1012 females, having a quite balanced (~50%) dataset.
- Hepatitis Laboratory measurements of Hepatitis B and C patients, admitted in Japan. Eleven temporal variables, having the top vertical support.
 204 Hepatitis B patients and 294 Hepatitis C patients (~60%).

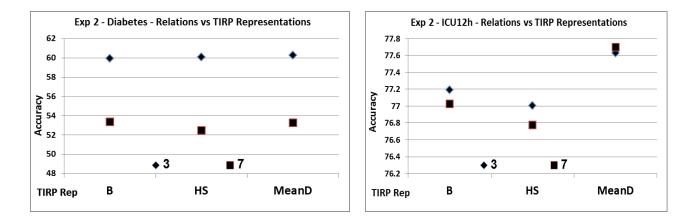
KarmaLegoS Evaluation Setup

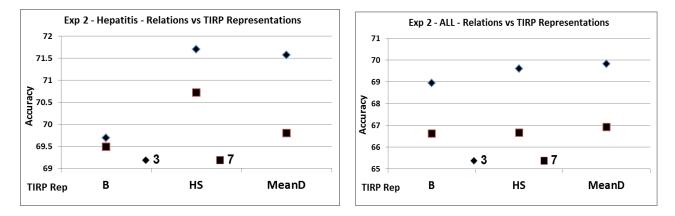


KarmaLegoS - Parameters

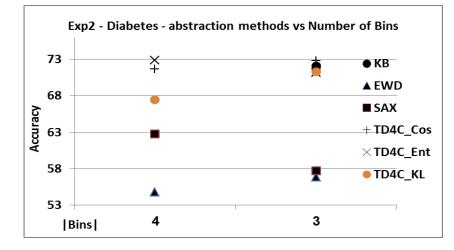
- To evaluate KarmaLegoS 5 experiments were designed for the various parameters in the KarmaLegoS framework, including:
 - Abstraction method (KB, EWD, SAX, TD4Cs)
 - Number of bins (3, 4)
 - Temporal relations set (3, 7)
 - TIRP Representation (Binary, HS, MeanD)
- For that experiments we designed and ran on three real datasets using 10 fold cross validation and RandomForest, compared according to the Accuracy measure.

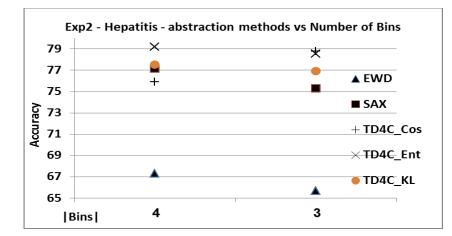
Temporal Relations and TIRP representation

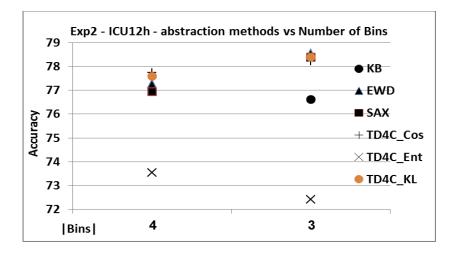


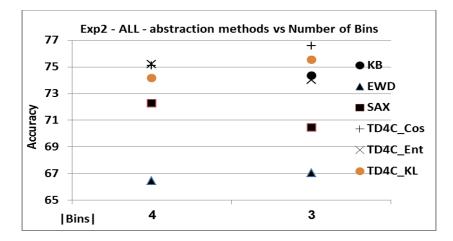


Abstraction Methods vs Bins Number ε=0, 3 temporal relations, MeanD and NoFS





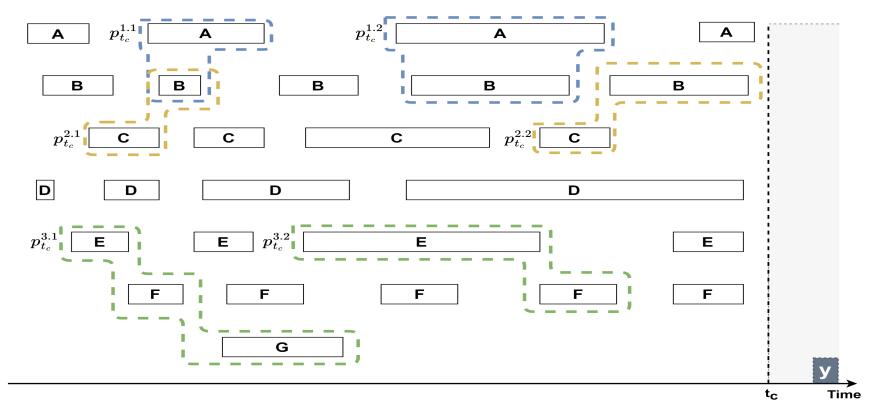




Continuous event's prediction via TIRPs



Detected TIRPs Instances



Continuous Prediction of TIRP's Completion

At any timepoint (e.g., t_c^1 , t_c^2 , t_c^3 , and t_c^4), we aim to continuously estimate the probability of the TIRP's completion.

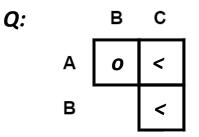
Nevo Itzhak, Szymon Jaroszewicz, Robert Moskovitch, Continuously Predicting a Time Intervals Based Pattern Completion Towards Event Prediction, PAKDD, **Osaka, Japan, 2023**.

Segmented CPM (SCPM)

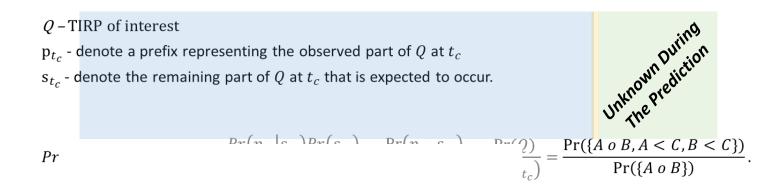
Q – TIRP of interest

 \mathbf{p}_{t_c} - denote a prefix representing the observed part of Q at t_c

 s_{t_c} - denote the remaining part of Q at t_c that is expected to occur.



$$Pr(Q|t_c) = Pr(\underline{s_{t_c}|p_{t_c}}) = \frac{Pr(p_{t_c}|s_{t_c})Pr(s_{t_c})}{Pr(p_{t_c})} = \frac{Pr(p_{t_c},s_{t_c})}{Pr(p_{t_c})} = \frac{Pr(Q)}{Pr(p_{t_c})} = \frac{Pr(A \circ B, A < C, B < C)}{Pr(A \circ B)}$$



Segmented CPM (SCPM)

Q – TIRP of interest

- \mathbf{p}_{t_c} denote a prefix representing the observed part of Q at t_c
- \mathbf{s}_{t_c} denote the remaining part of Q at t_c that is expected to occur.

$$Pr(Q|t_{c}) = Pr(\underline{s}_{t_{c}}|p_{t_{c}}) = \frac{Pr(p_{t_{c}}|s_{t_{c}})Pr(s_{t_{c}})}{Pr(p_{t_{c}})} = \frac{Pr(Q)}{Pr(p_{t_{c}})} = \frac{Pr(A \circ B, A < C, B < C)}{Pr(D_{t_{c}})}.$$

$$Methods$$

$$Evaluation$$

$$Results$$

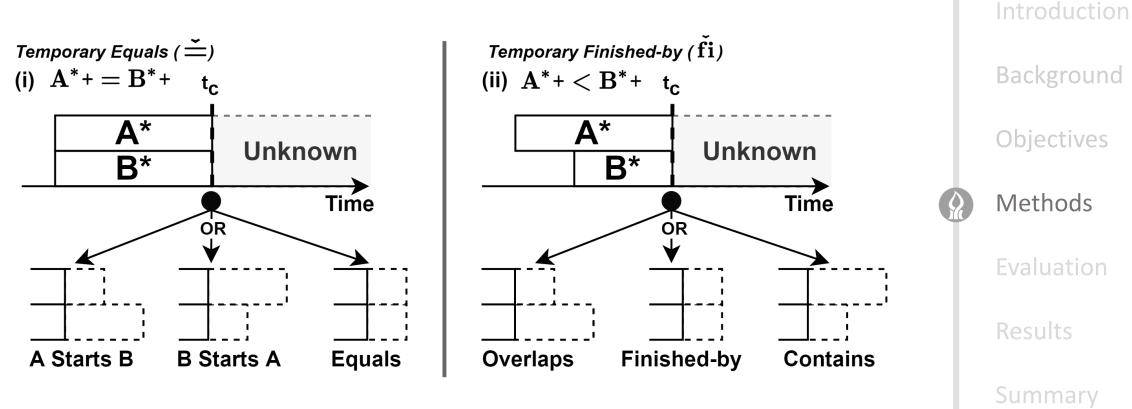
$$Summary$$

Introduction

Background

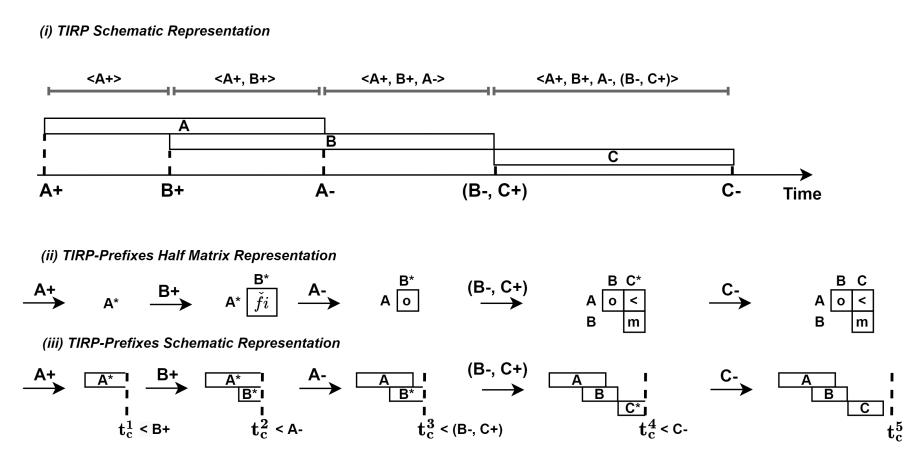
Objectives

Unfinished Coinciding STIs



Possible evolving temporal relations, given that the start times of a pair of unfinished STIs are known.

TIRP-Prefixes Representation

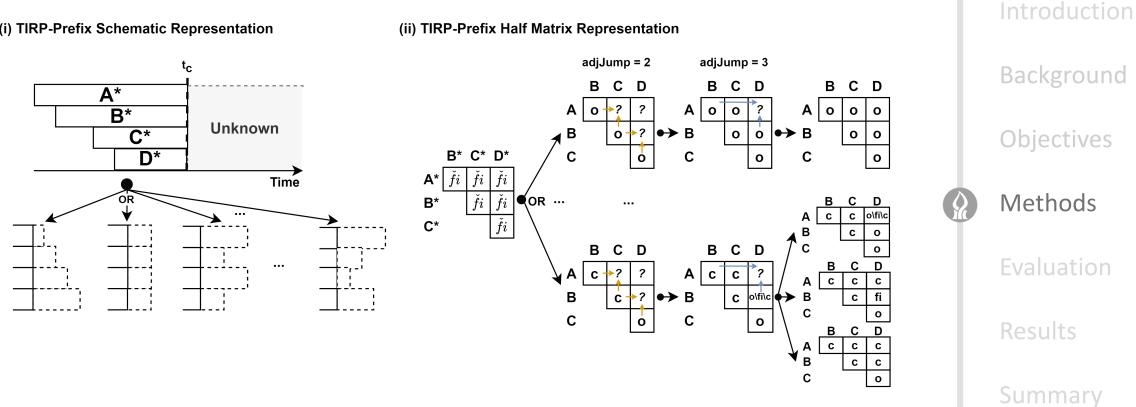


A TIRP of interest is divided into TIRP-prefixes that are part of the TIRP's evolving process

Slide 74 out of 25

Unfinished Coinciding STIs

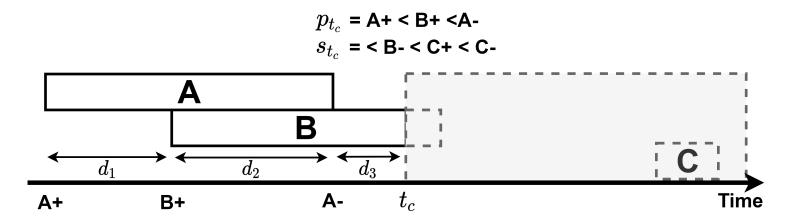
(i) TIRP-Prefix Schematic Representation



For four unfinished symbolic time intervals, a naive generation of all the suitable temporal relations among them requires generating up to $3^{(4^2-4)/2} = 3^6 = 729$ patterns

Slide 75 out of 25

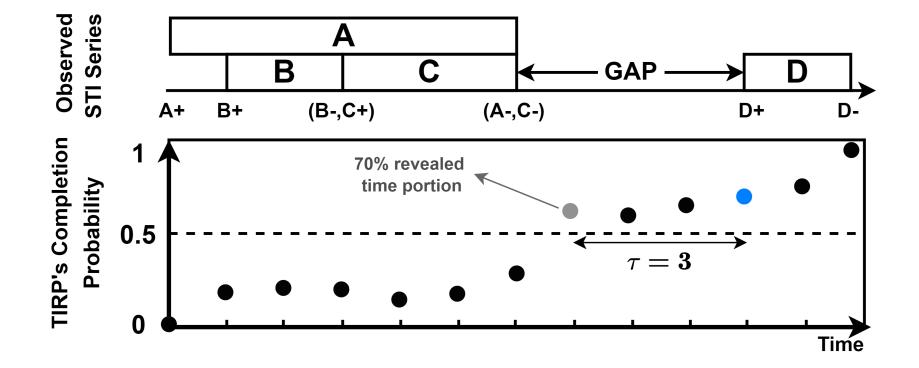
Machine-Learning-based CPM (CPML)



Time durations d1, d2, and d3 are based on tieps A+, B+, A-, and tc, which are used as *features for the classifiers* to perform the TIRP's completion prediction.

Instance	d1	d2	d3	 dk	Class
1	1	2	12	 0	1
2	9	12	1	 0	0
n	1	2	11	 0	1

Early Warning Strategies



An alert could be raised after the probability was consistently exceeded for some pre-defined decision time delay

Evaluation Research Questions

A. Which CPM performs better, in terms of prediction performance and earliness, in predicting the completion of a TIRP?

B. Which value of τ performs best, in terms of prediction performance and earliness, in predicting the completion of a TIRP?

Evaluation

Datasets

 Table 1. The evaluation datasets' parameters

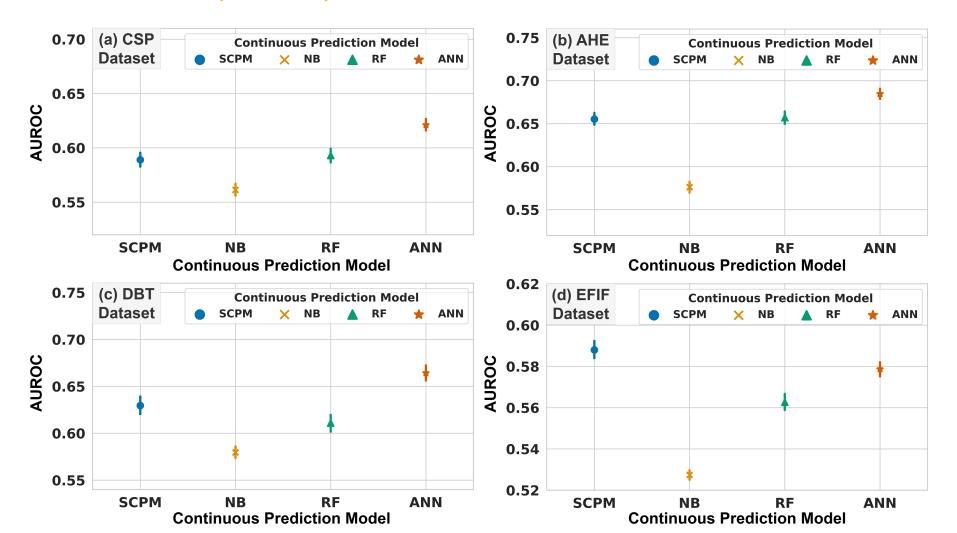
Name	#Ent	#Var	#Timestamps	Granularity	#EntEvent	#TIRPs
\mathbf{CSP}	329	13	720	minutes	115~(35%)	257
AHE	$1,\!000$	4	238	hours	500~(50%)	246
\mathbf{DBT}	1,710	12	24	months	239~(14%)	256
EFIF	823	15	144	weeks	121~(15%)	529

We evaluated the proposed models using four real-life datasets:

- 1. CSP dataset ICU patients, who underwent cardiac surgery Event: low cardiac index with values lower than 2.5 L/min/m
- 2. AHE dataset ICU patients from multiple ICUs, in which half of the patients had acute hypertensive episodes *Event: AHE onset*
- 3. DBT dataset Type II diabetes patients Event: high Hemoglobin A1C with values greater than 9%
- 4. EFIF dataset elderly first fall of residents Event: First fall

Results Preliminary Analysis

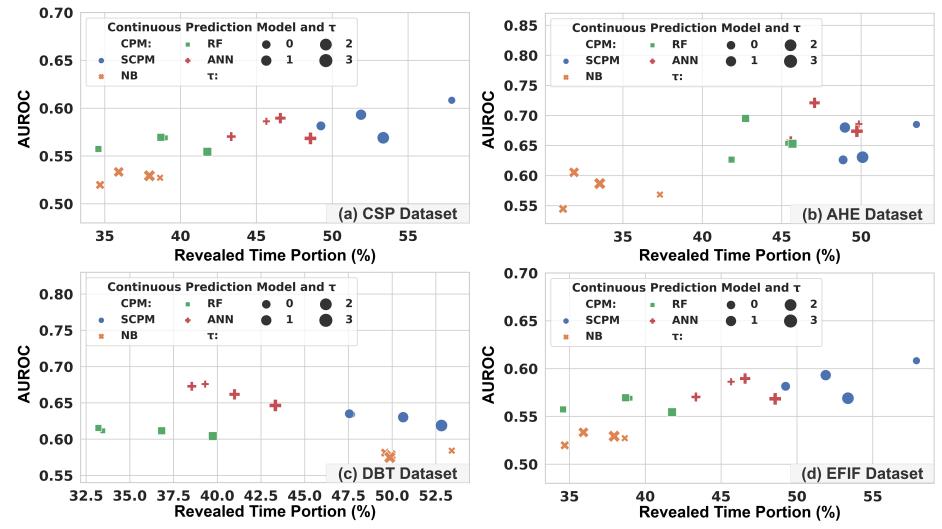
ANN performed better than the other models.



Results Preliminary Analysis

More accurate models need more time to make decisions.

SCPM provided the latest predictions NB and RF provided the earliest predictions.





- 1. Continuous TIRP's completion prediction
- 2. Uncertainty related to evolving temporal relations and our solution-- TIRP-prefix representation
- 3. CPML based on an ANN outperformed other models with 1.5% AUROC on average, while CPML based on NB or RF provided early TP predictions.

Thank You!

Prof. Robert Moskovitch Head, Complex Data Analytics Lab Software and Information Systems Engineering Ben Gurion University Israel



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Relevant References

- Omer Harel, Robert Moskovitch, Complete Closed Time Intervals-Related Patterns Mining, The 35th AAAI Conference on Artificial Intelligence (AAAI 2021), Vancouver, Canada, 2021. Rank A*
- Nevo Itzhak, Szymon Jaroszewicz, Robert Moskovitch, Continuously Predicting a Time Intervals Based Pattern Completion Towards Event Prediction, PAKDD, Osaka, Japan, 2023.
- Nevo Itzhak, Maya Schvetz, Itay Pesach, Robert Moskovitch, Acute Hypertensive Episodes Prediction, *Artificial Intelligence in Medicine*, 2023.
- Maya Shwetz, Lior Fuchs, Victor Novack, Robert Moskovitch, Outcomes Prediction in Longitudinal Data: Study Designs Evaluation, use case in ICU Acquired Sepsis, *Journal of Biomedical Informatics*, 2021.

Sepsis Acquired During ICU Admission

Patient with suspected infection

Identified by administration of antibiotics and sampling of body fluid

Patient with Sequential Organ Failure Assessment (SOFA) score >= 2

> SOFA determines the extent of a person's organ rate of failure

> ICU-acquired Sepsis is defined as one that started at least 48 hours after admission

Maya Shwetz, Lior Fuchs, Victor Novack, Robert Moskovitch, Outcomes Prediction in Longitudinal Data: Study Designs Evaluation, use case in ICU Acquired Sepsis, *Journal of Biomedical Informatics*, 2021.

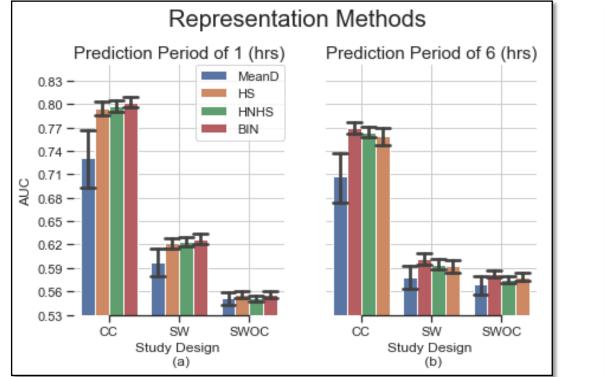
	ICU Acquired Sepsis
	↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓ ↓
t _o I Admission First 2 days in ICU	Time interval of 48 hours,
Start Time	SOFA>=2

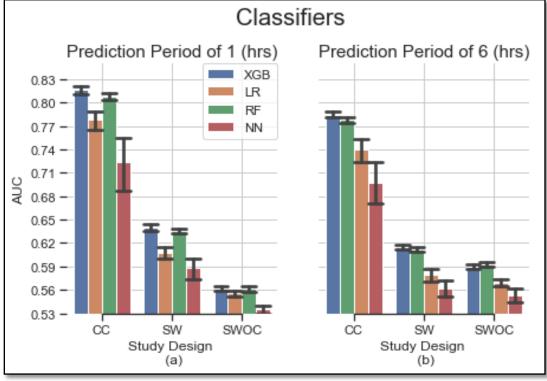
Exp 1- Predict the Occurrence of Sepsis Acquired in ICU Onset a Certain Time in Advance

- Case-Control design relative to the outcome
- Case-Crossover design with sliding window approach
- Sliding window (Case-Crossover-Control)

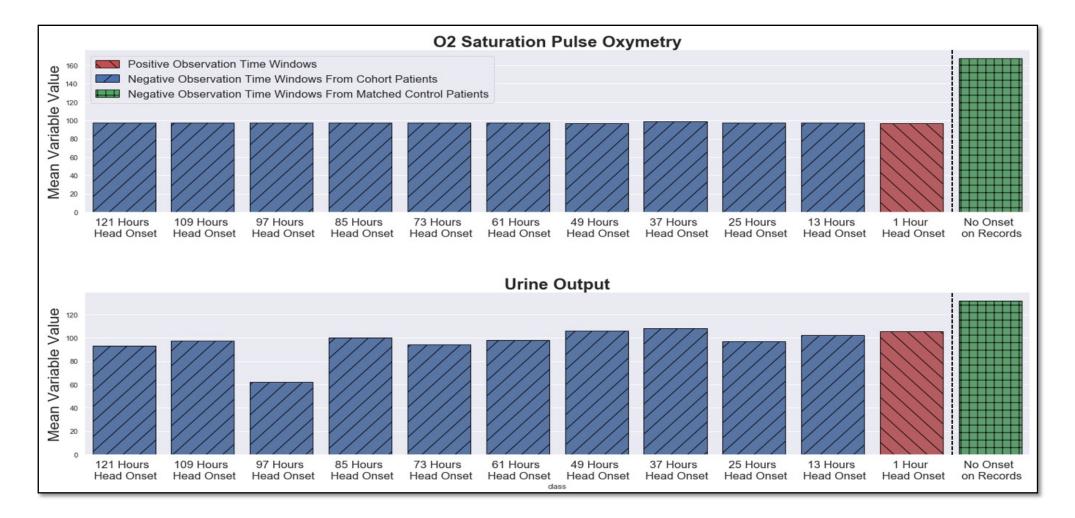
		С	ohort Subject		
					ICU Acquired Sep
	Negative Window 6/12 hours	Negative Window 6/12 hours	Negative Window 6/12 hours	Positive Window 6/12 hours	_
-					
t _o Admiss Start Ti		/	• t _x		Prediction Time (1/6 hours)
		N	/latched Cont	rol	
			<u> </u>	Negative wind 6/12 hours	
t _o Admiss Start TI			••••• t,	-p-o	t _x -p

Representation Methods and Classifiers





Low Performance Possible Reason

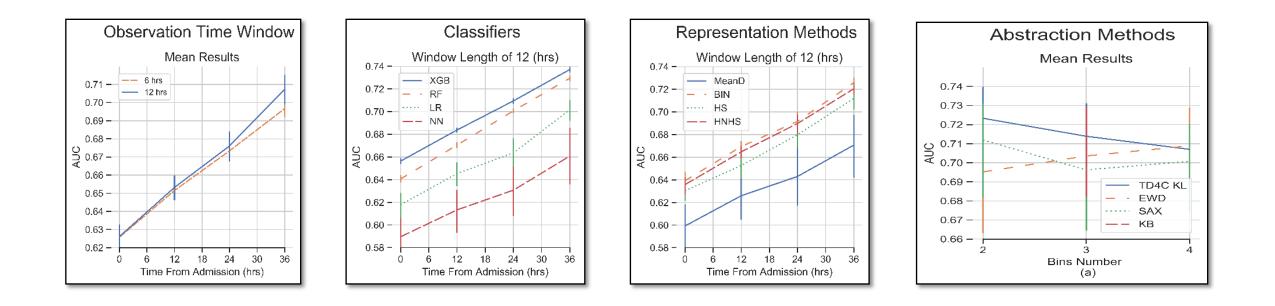


Exp 2 - Predict whether a patient will develop sepsis acquired in ICU at his stay

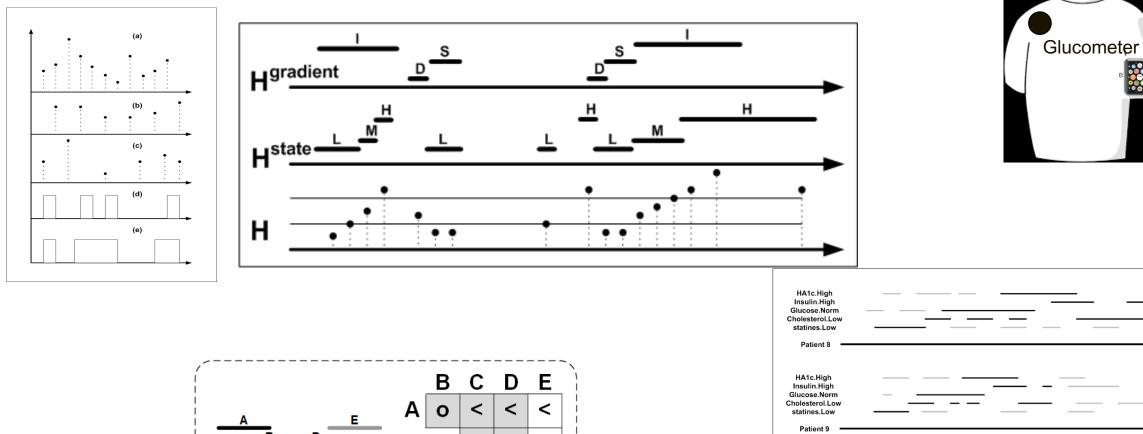
 Case-Control design relative to prior event, admission start time, to the outcome

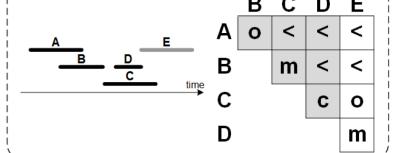
	Cohort Subject
	ICU Acquired Sepsis
	· •
Positive Observation Window (O ₁)	
<u></u>	<u></u>
t _o t Admission	t _x
Start TIME	
	Control Subject
Negative Observation	
Window (O ₀)	
t _o t Admission	n
Start TIME	

AUC results per hours from start ICU admission and observation period



Temporal Data Analytics (IoT)





insuin.nign	
Glucose.Norm	
Cholesterol.Low	
statines.Low	
statilles.Low	
Patient 8	
HA1c.High	
Insulin.High	
Glucose.Norm	
Cholesterol.Low	
statines.Low	
Statifies.LOW	
Patient 9	
HA1c.High	
Insulin.High	
Glucose.Norm	
Cholesterol.Low	
statines.Low	
Patient 10	
Patient 10	

Al in Medicine

- Artificial Intelligence in Medicine (AIM) one of the most important domains for human kind health
- Covers many research topics: text, image, temporal, and decision support
- In recent years with the breakthroughs in deep learning, there are tasks in medicine, such as image processing that benefited significantly. However, also longitudinal data, especially in Intense Care Units. AI was used in automating treatments, consisting on clinical guidelines, which were computerized.
- The speakers will cover ideally the following topics: Electronic Health Records Analytics Image Processing Longitudinal Data Analytics in medical data (outpatient, or inpatient data, or ICU).
 - Generative AI in Medicine.

Order of preference	Country	Full name	Institution	Email address	Research field	Expertise/reason for choice
1	Israel	Dr. Mor Peleg	Haifa University	peleg.mor@gmail.com	Al for medicine	Chief Editor, Journal of Biomedical Informatics
2	Israel	Dr. Erez Shmueli	Tel Aviv University	shmueli@tau.ac.il	Breathing Complications Diagnosis, BIG DATA in Medicine	Had made meaningful contributions in that field
3	Israel	Dr. Eran Segal	Weitzman Institute		Diabetes data analytics	Well known studies around the world
1	Japan	Dr. Takanori Hasegawa	M&D Data Science Center, Tokyo Medical and Dental University	tk.hasegawa@gmail.com https://www.hase62.jp/	bioinformatics	Human genome, transcriptome, epigenome, metagenome analysis, and clinical data for personalized and preventive medicine through modelling, prediction and inference
2	Japan	Dr. Shuhei Kurita	RIKEN-AIP	Shuhei.kurita@riken.jp https://shuheikurita.github.io/	Vision and Language	Vision and Language are two most important and popular fields in generative Al.
3	Japan	Dr. Ryoma Bise	Kyushu University	bise@ait.kyushu-u.ac.jp	Computer Vision for bioinformatics	 Pathological Image Segmentation/Classification, Cell Tracking Many papers published in top- tier conferences

Temporal Data Analytics

- While most of the works and methods developed in machine learning ignore time, dues to challenges and intention to simplify problems, longitudinal data is available more and more in recent years with the advancements of the Internet of Things.
- Thus, temporal data analytics is an important topic in many applicative domains, such as security in computers networks, intensive care unit and generally medicine, predictive maintenance and more.
- Other relevant tasks are forecasting in stocks, and other domains, in which sensory data are available. In recent decades there is a growing interest in discovery of frequent temporal patterns, such as sequential patterns, time intervals patterns and other, and their use for various tasks, such as classification, and more.

Order of preference	Country	Full name	Institution	Email address	Research field	Expertise/reason for choice
1	Israel	Mark Last	Ben Gurion University	mlast@bgu.ac.il	Data Mining	Had made several works in temporal data analysis
2	Israel	Assaf Shuster	Technion	assaf@cs.technion.ac.il	Data Mining	Is working in time series data
1	Japan	Koji Inoue	Kyoto University	Inoue.koji.3x@kyoto-ua.c.jp http://www.sap.ist.i.kyoto- u.ac.jp/members/inoue/	Spoken dialogue systems, Multimodal signal processing, Human-robot interaction	Highly evaluated young researcher in the field: http://www.sap.ist.i.kyoto- u.ac.jp/members/inoue/profile.html
2	Japan	Sho Yokoi	Tohoku University	yokoi@tohoku.ac.jp http://www.cl.ecei.tohoku.ac.jp/~yokoi/	Natural Language Processing	Active researcher who is also with RIKEN-AIP.