

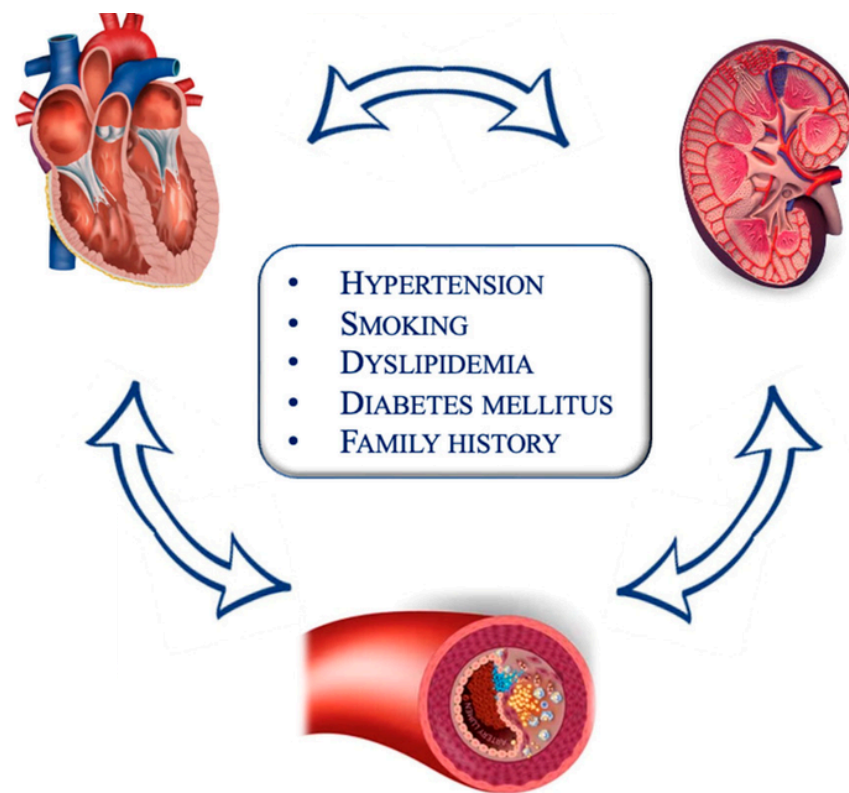
# End-to-End Reward Decomposition and Explainable Interaction: A Medical Inverse Reinforcement Learning Framework

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# Introduction



## Problem Statement:

- Patient Profile: 65 - year - old male admitted with diabetes, chronic kidney disease, and heart failure simultaneously.
- Treatment Dilemma
  - Doctors need to balance treatment effectiveness for all three diseases.
  - Must avoid drug conflicts and adverse reactions.
- Data and Interaction Issues
  - Incomplete records of disease progression.
  - Dynamic, complex interactions between the three conditions over time.

## Research Motivation:

- How can we **clearly** decompose and explain multiple clinical objectives in **complex medical decisions**?
- How can we robustly complete and utilize patient data when observations are **missing or incomplete**?
- How can we make every step of AI-driven medical decisions fully **transparent and traceable** for clinicians?

# Related Work

Research Direction	Recent Representative Works & References	Main Advances	Limitations & Challenges
<b>Inverse Reinforcement Learning (IRL) in Healthcare</b>	<ul style="list-style-type: none"><li>- Jayaraman &amp; Desman, 2024 (Nat. Digital Med.): Federated IRL</li><li>- Snoswell et al., 2024 (Front. Digital Health): Telemedicine preference IRL</li><li>- Fang et al., 2024 (arXiv): Safe clinical IRL</li></ul>	<ul style="list-style-type: none"><li>- Enables personalized medical recommendations</li><li>- Addresses privacy via federated learning</li><li>- Models patient preferences for telehealth</li></ul>	<ul style="list-style-type: none"><li>- Assumes complete data</li><li>- Struggles with complex multimorbidity interactions</li><li>- Limited reward interpretability</li></ul>
<b>Causal Modeling for Multimorbidity</b>	<ul style="list-style-type: none"><li>- Langenberg et al., 2023 (Nat. Medicine): Multimorbidity causal framework</li><li>- Yin et al., 2025 (arXiv): Multi-agent RL for disease interaction</li></ul>	<ul style="list-style-type: none"><li>- Reveals disease progression patterns</li><li>- Captures dynamic interactions between multiple diseases</li></ul>	<ul style="list-style-type: none"><li>- Difficult to disentangle temporal causal effects</li><li>- Lacks real-time, individualized decision support</li></ul>
<b>Learning from Partially Observed Data</b>	<ul style="list-style-type: none"><li>- Liu et al., 2024 (ICLR): POMDP and inverse weighting</li><li>- Xia et al., 2024 (OpenReview): Contrastive learning for EHR</li></ul>	<ul style="list-style-type: none"><li>- Adapts to incomplete medical records</li><li>- Utilizes partial data effectively through contrastive learning</li></ul>	<ul style="list-style-type: none"><li>- Uncertainty quantification issues</li><li>- Limited theoretical guarantees for convergence</li></ul>
<b>Optimization of Multimorbidity Treatment Decisions</b>	<ul style="list-style-type: none"><li>- Zhang et al., 2024 (AAAI): Multi-task RL for comorbidity</li><li>- Tan et al., 2024 (arXiv): Hierarchical multi-agent RL for multi-organ care</li></ul>		

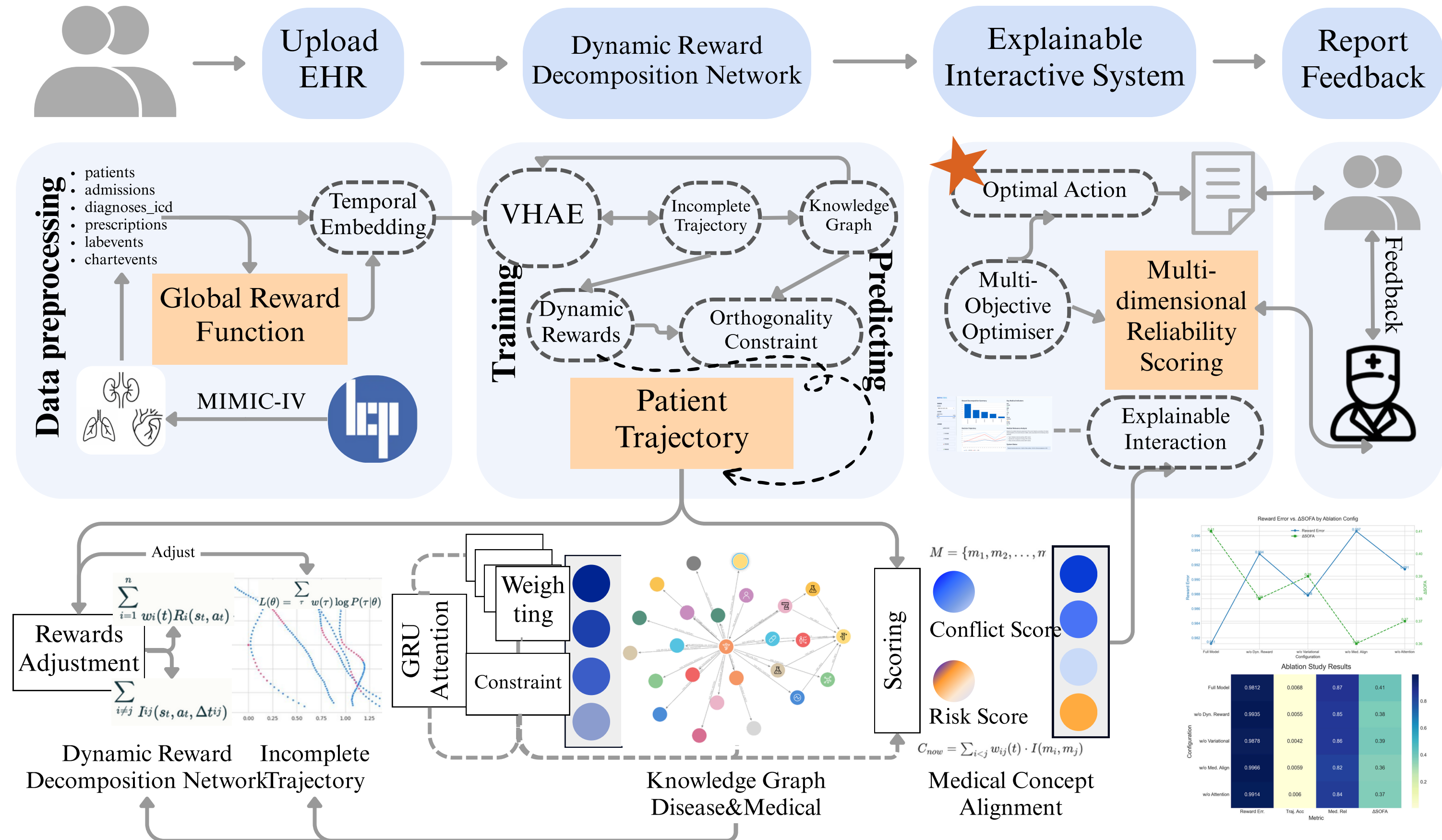
• IRL on incomplete, real-world multimorbidity data.

• Causal disentanglement of dynamic disease interactions.

• Robust learning under partial observation and uncertainty.

• Temporal conflict detection and multi-objective optimization.

# Methodology





# Core Technical Innovations

## Transparent Reward Decomposition

- Breaks complex clinical goals into clear, independent components.
- Orthogonality and attention ensure each reward is clinically meaningful and explainable.
- 

$$R(s, a) = \sum_{i=1}^K w_i(s) \cdot r_i(s, a), \quad L_{ortho} = \sum_{i \neq j} |r_i^T r_j|$$

## Robust Trajectory Completion

- Fills in missing patient data using a hybrid of deep learning and medical knowledge.
- Guarantees reliable, realistic recovery even with high data gaps.

$$L = \text{ELBO} + \alpha \cdot KG_{Loss} + \beta \cdot \text{Temporal\_Smoothness}$$

## Full-Path Explainability

- Visualizes every step of AI decision-making across time, features, and strategy.
- Empowers clinicians with dynamic, interactive, and traceable insights.



# Experimental Results

## 1 Method Comparison

Table 1: Comparison of Different Methods

Method	Components			
	Reward Decomp.	Trajectory Comp.	Medical Align.	Visual.
MERIT-IRL	✓	×	×	×
XAI-Med	✓	×	✓	✓
PO-IRL	×	✓	×	×
LRD-IIRL	✓	✓	✓	✓

## 2 Quantitative Results

Table 2: Quantitative Results on MIMIC-IV Dataset

Method	Reward Reconstruction			Traj. Acc	Med. Rel
	MSE	RMSE	Rel. Error		
MERIT-IRL	0.9951	0.9975	0.9951	0.0049	0.80
XAI-Med	0.9932	0.9966	0.9932	0.0068	0.85
PO-IRL	1.0023	1.0011	1.0023	-0.0023	0.75
LRD-IIRL	<b>0.9812</b>	<b>0.9905</b>	<b>0.9812</b>	<b>0.0068</b>	<b>0.87</b>

## 6 Evaluation Metrics

Table 6: Evaluation Metrics Definition

Metric	Calculation	Clinical Meaning
Reward Error	$\frac{1}{n} \sum_{i=1}^n (r_i - \hat{r}_i)^2$	Treatment effect prediction
Traj. Acc	$\frac{1}{n} \sum_{i=1}^n \mathbb{I}(s_i = \hat{s}_i)$	State prediction accuracy
Med. Rel	$\frac{1}{n} \sum_{i=1}^n \text{sim}(c_i, \hat{c}_i)$	Concept alignment degree
$\Delta$ SOFA	$\text{SOFA}_{t+1} - \text{SOFA}_t$	Safety indicator

## 3 Ablation Study

Table 3: Ablation Study Results

Config.	Performance Metrics			$\Delta$ SOFA
	Reward Err.	Traj. Acc	Med. Rel	
Full Model	0.9812	0.0068	0.87	0.41
w/o Dyn. Reward	0.9935	0.0055	0.85	0.38
w/o Variational	0.9878	0.0042	0.86	0.39
w/o Med. Align	0.9966	0.0059	0.82	0.36
w/o Attention	0.9914	0.0060	0.84	0.37

## 4 Clinical Evaluation

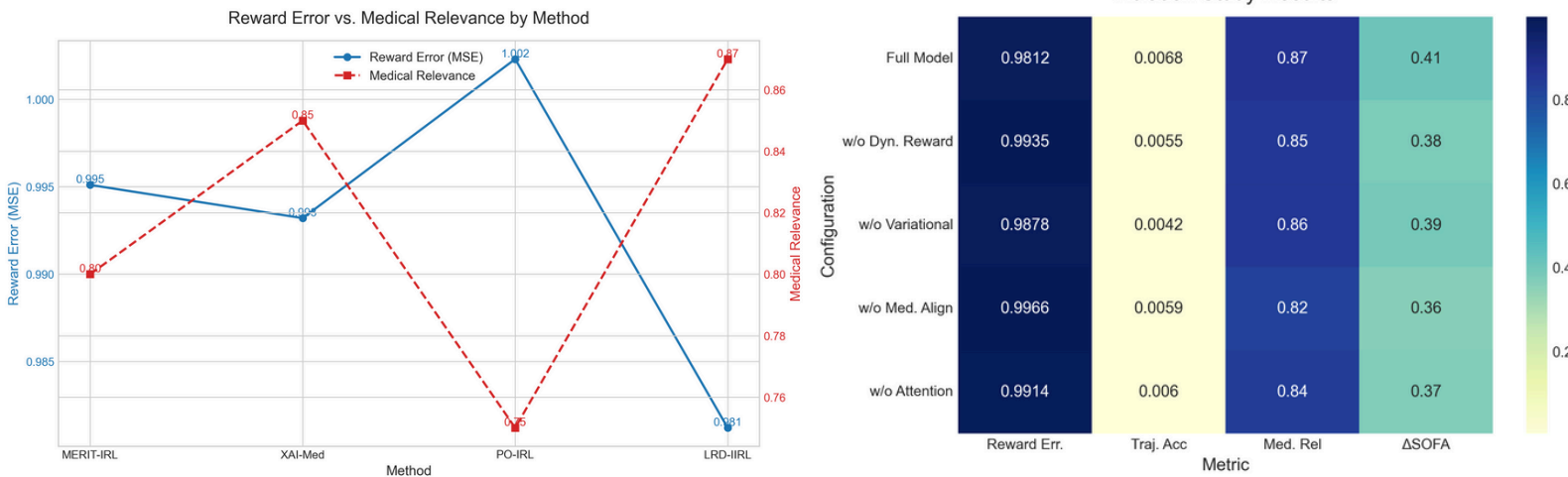
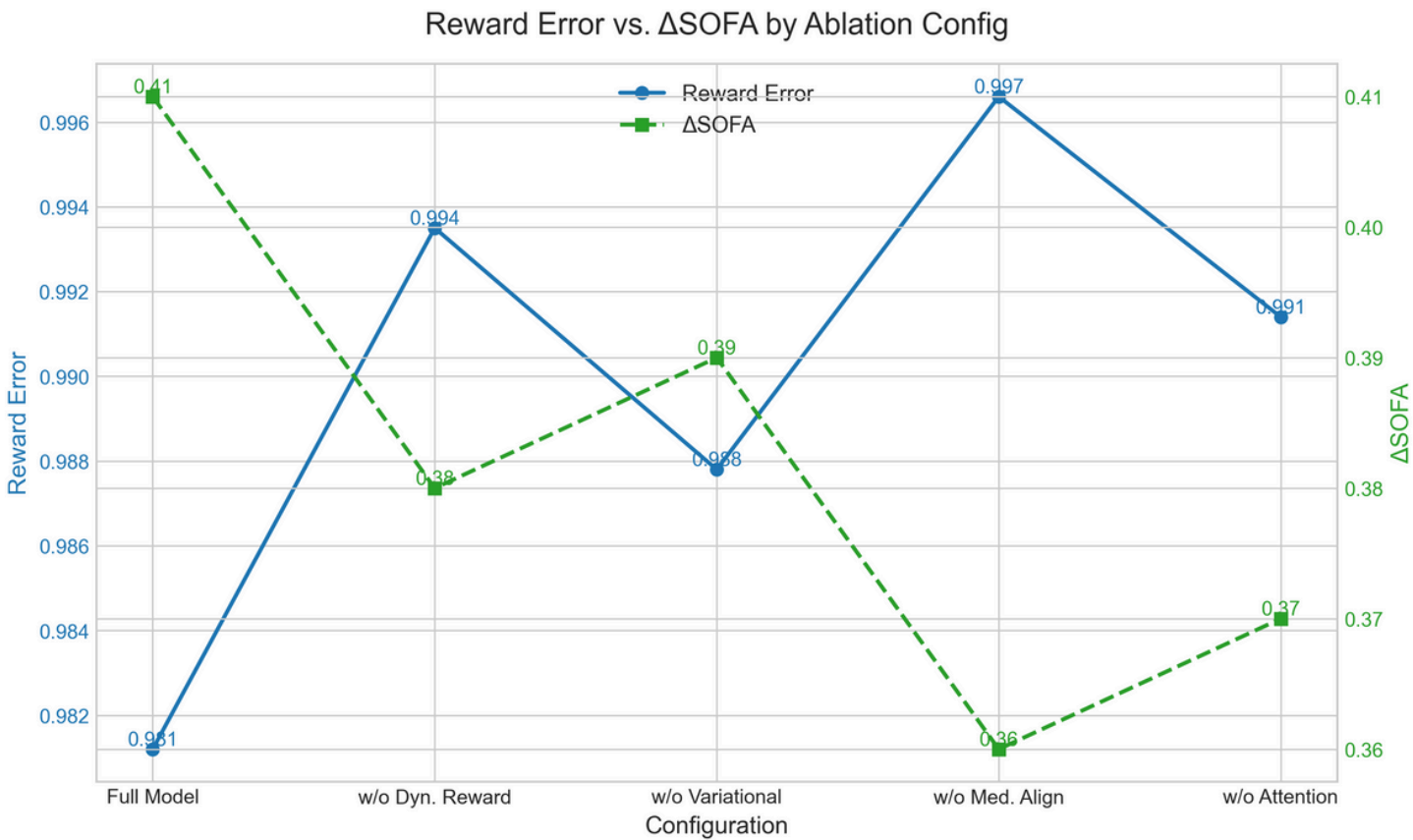
Table 4: Clinical Evaluation Results

Scenario	Expert Ratings			Acc. Rate
	Treatment	Interp.	Usability	
ICU Multi-morb.	$4.7 \pm 0.3$	$4.5 \pm 0.4$	$4.6 \pm 0.3$	92%
Oncology	$4.5 \pm 0.4$	$4.3 \pm 0.5$	$4.4 \pm 0.4$	88%
Emergency	$4.3 \pm 0.5$	$4.2 \pm 0.6$	$4.3 \pm 0.5$	85%

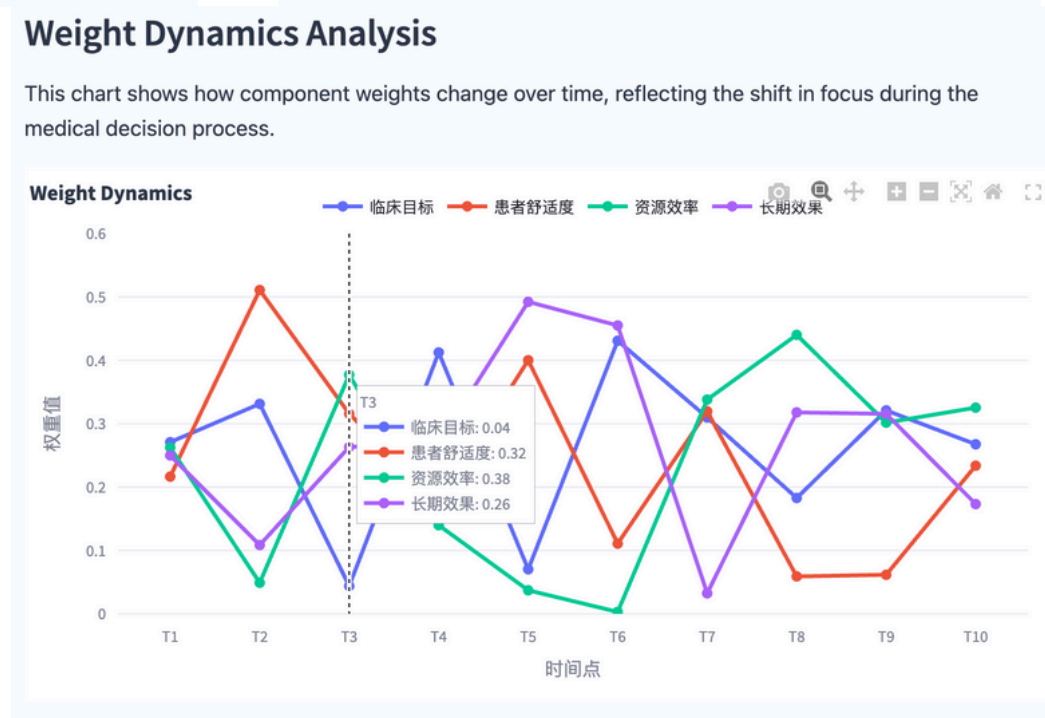
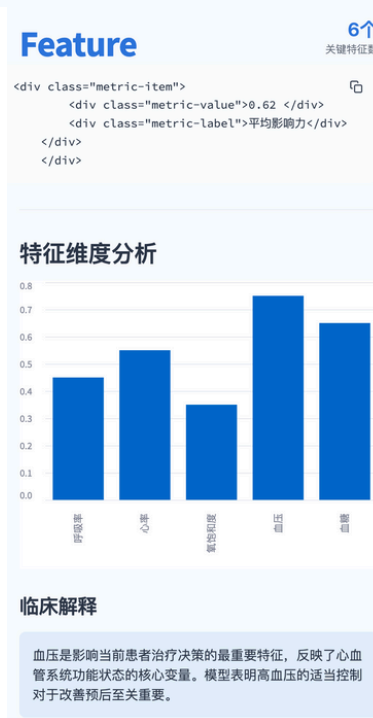
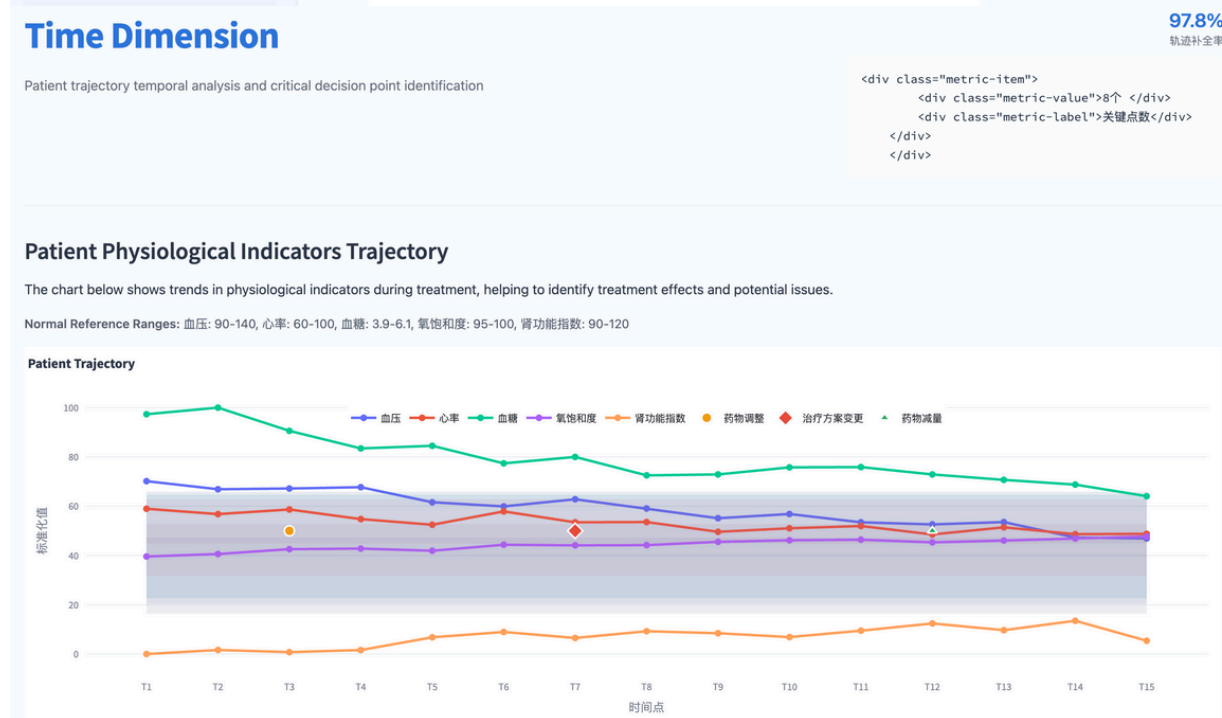
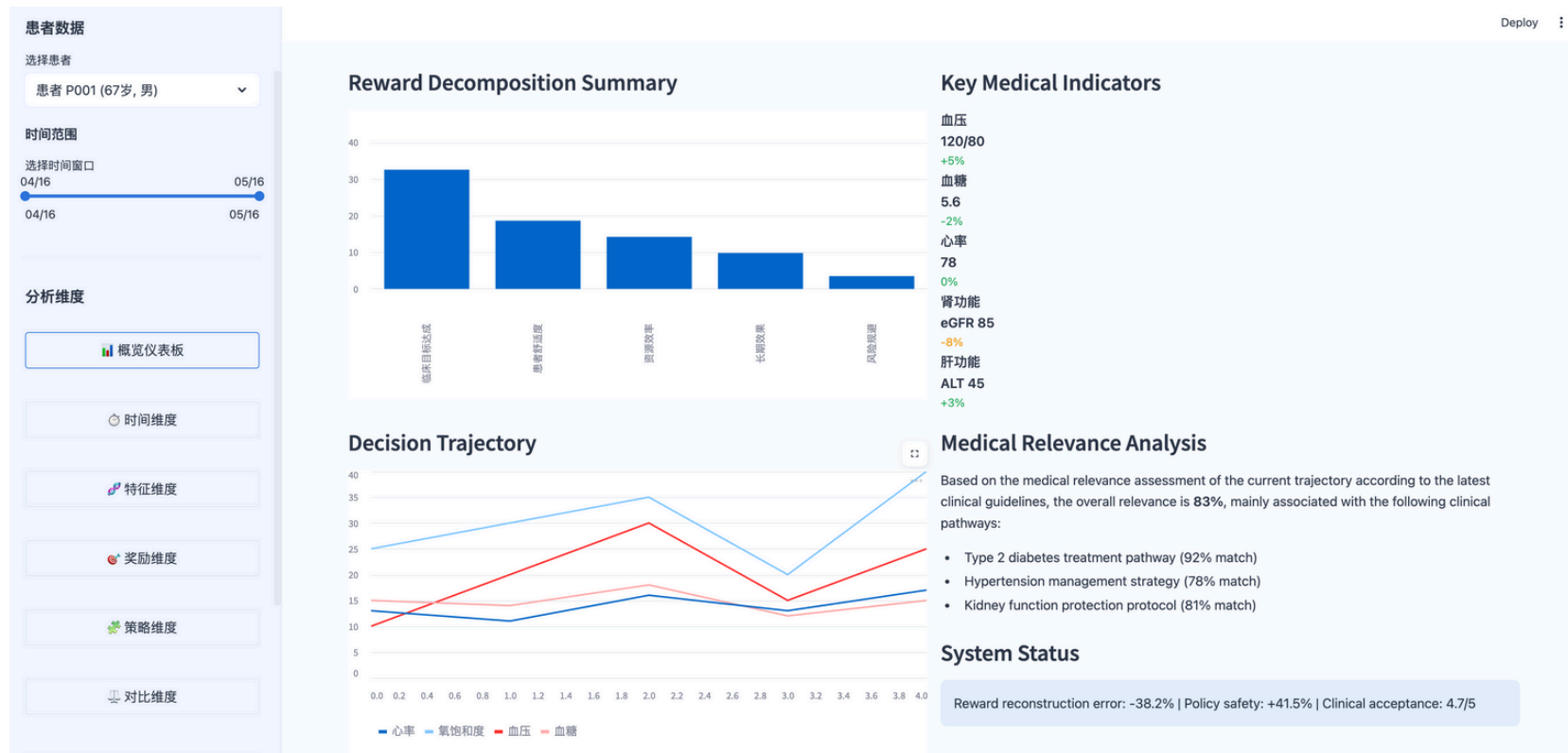
## 5 Dataset Statistics

Table 5: Dataset Statistics

Dataset	Sample Size			Miss. Rate
	Train	Val.	Test	
MIMIC-IV	15,000	3,000	2,000	5.2%
Synthetic	10,000	2,000	1,000	0.0%



# Interactive System Performance





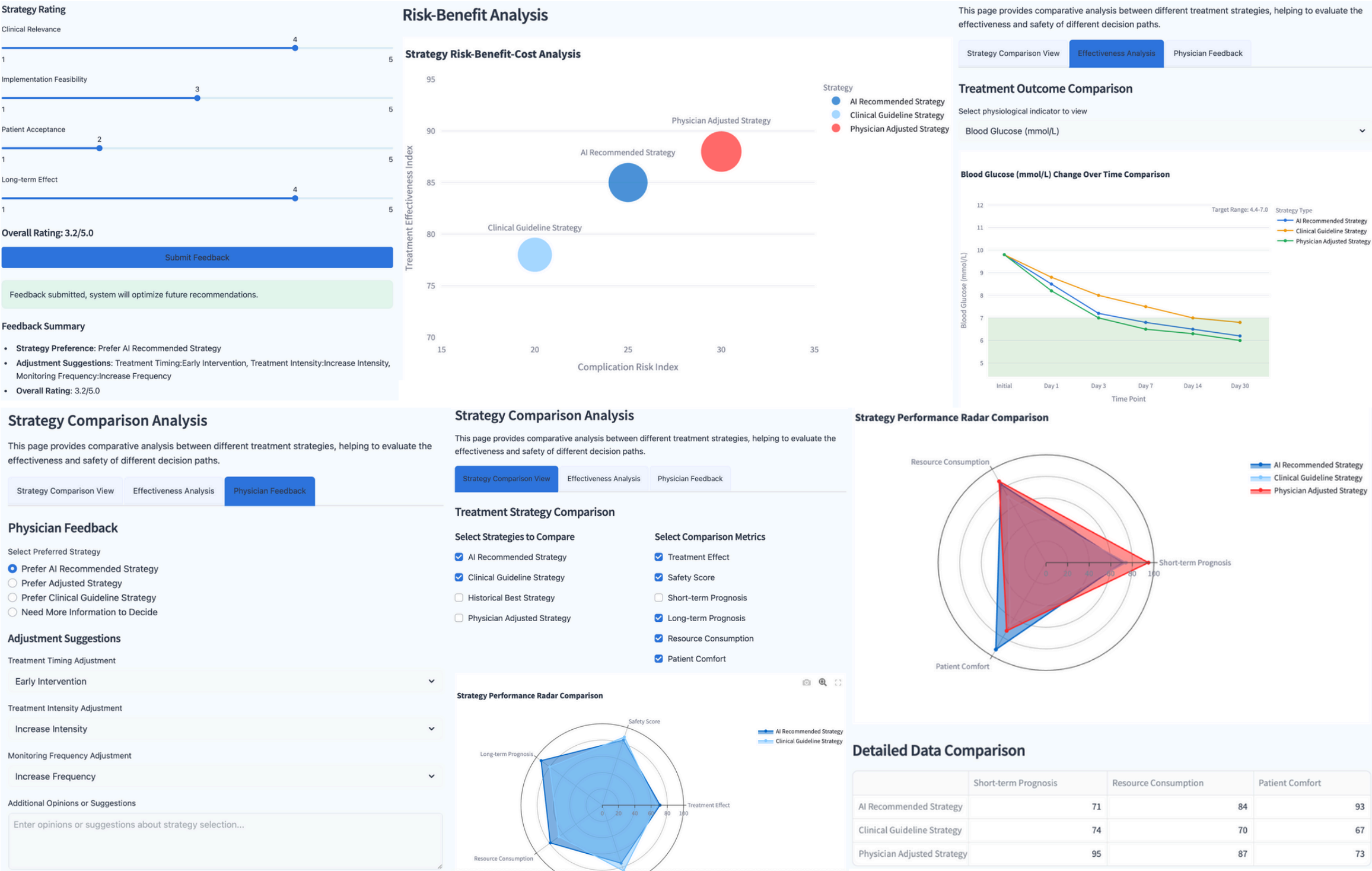
# Adaptation Scenario

**Clinical Decision Support:**  
Assists physicians in developing safe and effective treatment plans for patients with complex conditions like diabetes with heart failure.

**Preventive Medicine:**  
Identifies early warning signs of disease interactions, enabling proactive interventions before complications develop.

**Clinical Research:**  
Provides insights into disease interactions and treatment efficacy, supporting new therapeutic approach development.

**Personalized Medicine:**  
Adapts recommendations to patient-specific factors, including disease duration and severity patterns.





# Conclusion

## Technical Innovations

- We introduce LRD-IIRL, an end-to-end framework that unifies reward decomposition, robust trajectory completion, and full-path explainability for clinical decision-making.
- Our method leverages orthogonal constraints and medical knowledge to disentangle multi-objective rewards, ensuring clinical interpretability.
- A hybrid variational autoencoder and knowledge graph module enables reliable patient trajectory recovery, even with high missingness.

## Empirical Validation

- Extensive experiments on large-scale clinical datasets (e.g., MIMIC-IV) show LRD-IIRL outperforms state-of-the-art baselines in reward reconstruction, trajectory completion, and medical relevance.
- The framework achieves higher expert acceptance rates, demonstrating its value for trustworthy, real-world medical AI deployment.
- LRD-IIRL sets a new standard for interpretable and robust AI in healthcare.

## Clinical Trust and Transparency

- The five-dimensional XAI-Viz system delivers transparent, interactive visualizations across time, features, rewards, policy, and expert comparison.
- Every AI decision is traceable and clinically aligned, bridging the trust gap between clinicians and intelligent systems.
- Our approach empowers doctors to understand, validate, and refine AI recommendations in real-world scenarios.

# Q&A

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