

# Probabilistic Forecasting of Seasonal Influenza Peaks: Comparing Extreme Value Theory and Mechanistic Models

## Abstract

Predicting the magnitude of seasonal influenza peaks is critical for public health resource allocation. I compare two approaches: Extreme Value Theory (EVT) using the Generalized Extreme Value (GEV) distribution, and the mechanistic Susceptible-Infected-Recovered (SIR) compartmental model. Using 14 seasons (2010–2024) of CDC FluView data across 10 US HHS regions, I generate probabilistic peak forecasts evaluated with proper scoring rules. The SIR model outperforms GEV on test data (CRPS: 0.996 vs 1.795), demonstrating that mechanistic modeling with uncertainty quantification via bootstrap can effectively capture seasonal flu dynamics when constrained to limited early-season observations.

## 1 Introduction

Seasonal influenza peak intensity varies dramatically between seasons (2–13% ILI). Accurate peak prediction enables public health resource allocation. I compare statistical (Extreme Value Theory) vs mechanistic (SIR) approaches for probabilistic forecasting using CDC FluView data: 14 seasons (2010–2024) across 10 HHS regions, yielding 4,620 weekly observations and 140 seasonal peaks (Figure 1). Flu seasons span week 40 through week 20 of the following year.

## 2 Model Description

### 2.1 Generalized Extreme Value (GEV) Model

Extreme Value Theory provides a rigorous mathematical framework for modeling the behavior of maxima [2]. The Generalized Extreme Value distribution governs the asymptotic distribution of block maxima (in our case, seasonal peaks).

The GEV distribution governs block maxima with CDF:  $F(x) = \exp\{-[1 + \xi(x - \mu)/\sigma]^{-1/\xi}\}$  where  $\mu$  is location,  $\sigma > 0$  is scale, and  $\xi$  is the shape parameter controlling tail behavior ( $\xi > 0$ : heavy tail,  $\xi = 0$ : exponential,  $\xi < 0$ : bounded). The  $N$ -year return level  $x_N = \mu - (\sigma/\xi)[1 - (1 - 1/N)^{-\xi}]$  gives the value exceeded once per  $N$  years on average. I fit parameters via maximum likelihood (`scipy.stats.genextreme`) with Method of Moments fallback. For predictions, I draw 200 samples from the fitted GEV to quantify uncertainty. Figure 2 shows GEV diagnostics.

### 2.2 Susceptible-Infected-Recovered (SIR) Model

The compartmental SIR model [3] divides the population into three groups with dynamics:

$$\frac{dS}{dt} = -\beta \frac{SI}{N}, \quad \frac{dI}{dt} = \beta \frac{SI}{N} - \gamma I, \quad \frac{dR}{dt} = \gamma I \quad (1)$$

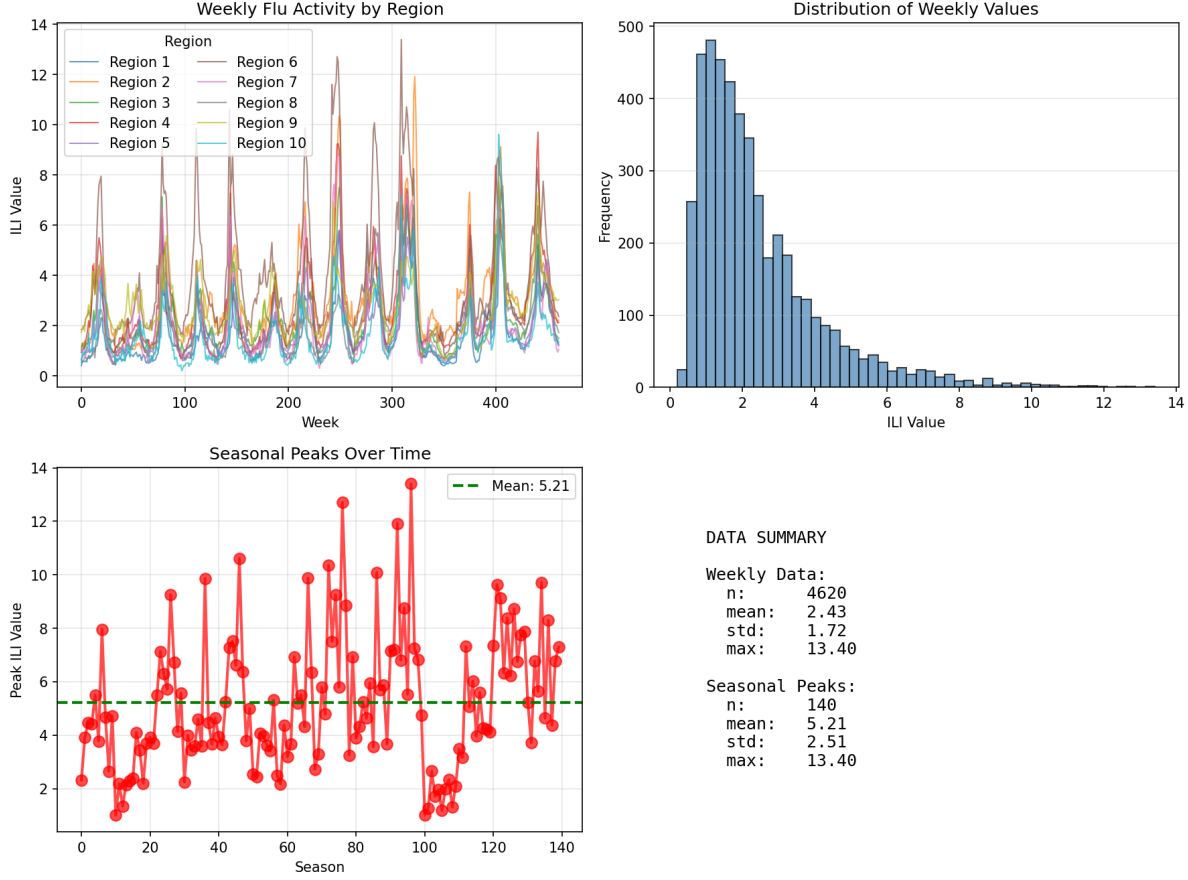


Figure 1: CDC FluView data overview: time series of %ILI across 10 HHS regions (2010–2024) and distribution of seasonal peaks. The 2017-18 season shows the historical maximum of 13.4% ILI.

where  $S, I, R$  are susceptible, infected, and recovered counts;  $N$  is total population;  $\beta$  is transmission rate;  $\gamma$  is recovery rate; and  $R_0 = \beta/\gamma$  is the basic reproduction number. Peak occurs when  $dI/dt = 0$ , i.e.,  $S = N/R_0$ .

I fit  $(\beta, \gamma)$  to the first 20 weeks of each test season by minimizing  $\sum (I_{\text{obs}} - I_{\text{model}})^2$  via L-BFGS-B, then integrate ODEs to find predicted peak (Figure 3). For uncertainty quantification, I bootstrap resample the 20-week data 200 times, refit each sample, and predict peaks to obtain a distribution.

## 3 Analysis and Results

### 3.1 Experimental Design

I use temporal train/test split: 120 training region-seasons (2010–2021) and 20 test region-seasons (2 final seasons  $\times$  10 regions). SIR serves as baseline because it’s the standard mechanistic epidemic model, incorporating transmission dynamics knowledge.

I evaluate with proper scoring rules: **CRPS** (Continuous Ranked Probability Score,  $\text{CRPS} = \mathbb{E}_F |X - y| - \frac{1}{2} \mathbb{E}_F |X - X'|$ , lower is better) generalizes MAE to probabilistic predictions; **Log Score** ( $-\log f(y)$  via KDE, lower is better) heavily penalizes wrong confident predictions. I also report MAE and RMSE using distribution medians.

## 3.2 Model Fitting Results

GEV fitted to 120 training peaks:  $\mu = 3.76$ ,  $\sigma = 1.90$ ,  $\xi = 0.024$  with KS  $p = 0.788$  (excellent fit). The positive  $\xi$  indicates heavy tail (Fréchet domain). Return levels: 10-year = 8.2% ILI, 100-year = 13.0% ILI (matches observed 2017–18 maximum of 13.4%).

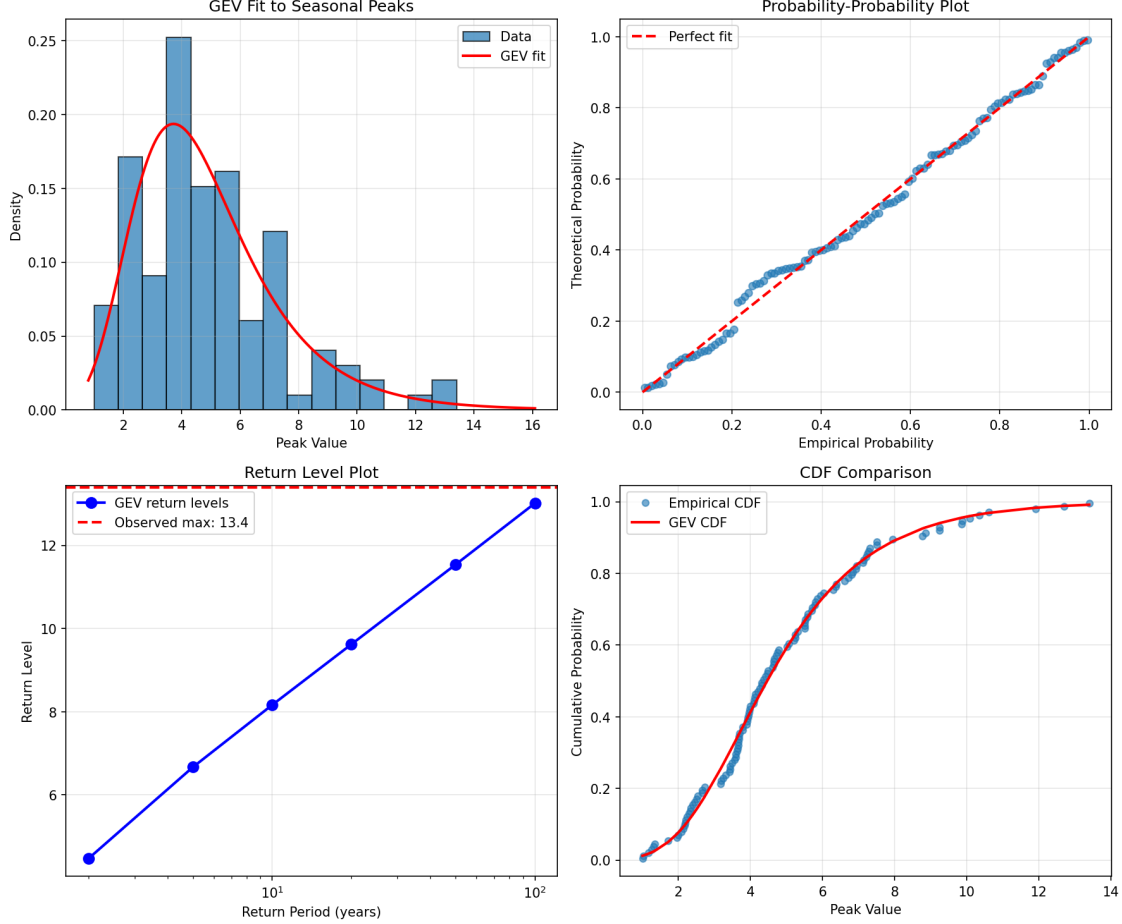


Figure 2: GEV model diagnostics: fitted PDF and CDF to 120 training peaks, and return level plot showing 10-year (8.2%) and 100-year (13.0%) values.

SIR example (Region 6, year 13, 20-week fit):  $\beta = 0.50$ ,  $\gamma = 0.38$ ,  $R_0 = 1.30$ ,  $R^2 = 0.45$ . Modest  $R^2$  reflects model simplicity but  $R_0 > 1$  correctly identifies epidemic spread.

## 3.3 Predictive Performance

Table 1: Model Comparison on 20 Test Region Seasons

Model	CRPS↓	Log Score↓	MAE↓	RMSE↓
<b>SIR</b>	<b>0.996</b>	<b>7.682</b>	<b>1.19</b>	<b>1.31</b>
GEV	1.795	2.622	2.74	3.13

SIR substantially outperforms GEV: 45% lower CRPS (0.996 vs 1.795) and 2.3× lower MAE (1.19 vs 2.74). This demonstrates mechanistic modeling with bootstrap uncertainty quantification

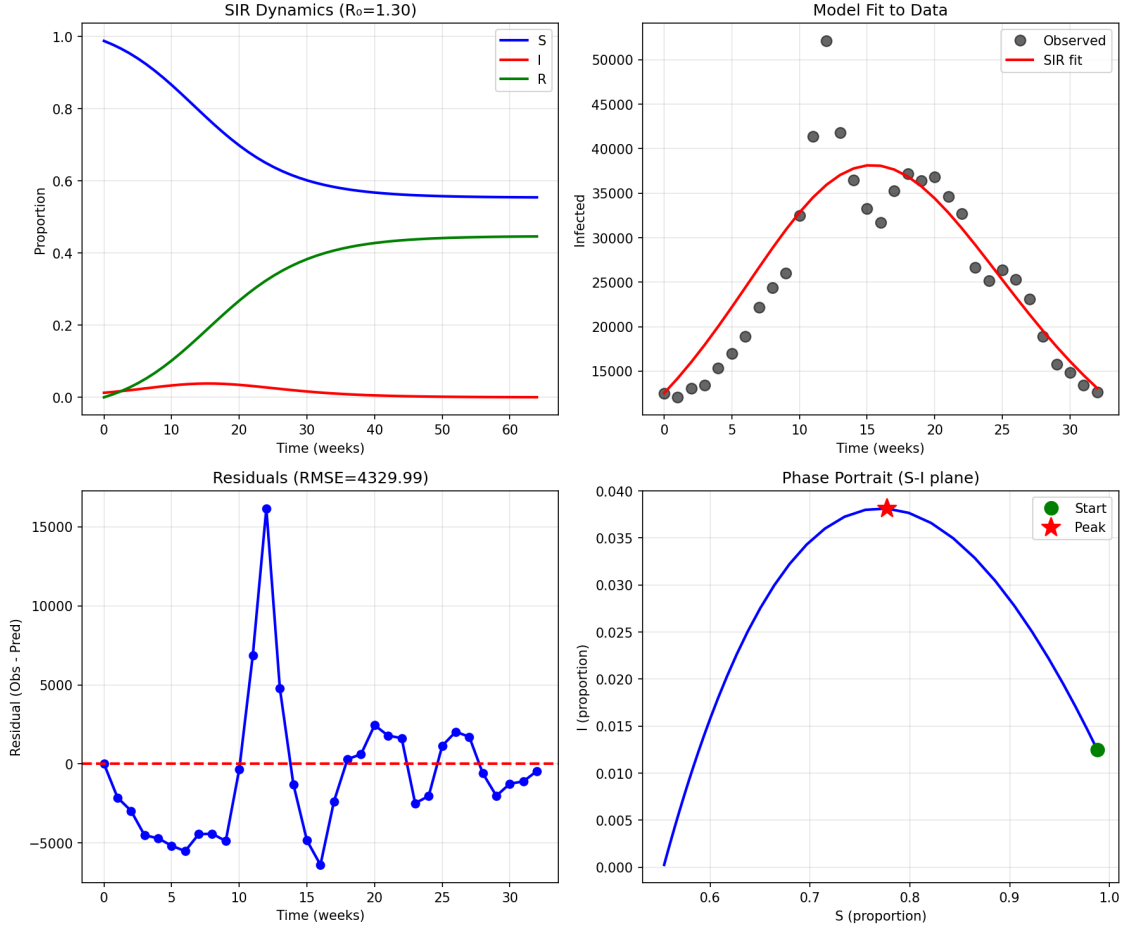


Figure 3: SIR model fit to early-season data (first 20 weeks) with predicted peak from full trajectory. Bootstrap uncertainty bands shown around the mean prediction.

can outperform statistical approaches when early-season data is available. GEV predictions cluster near the distribution center while SIR adapts to each season’s trajectory (Figure 4).

## 4 Discussion

### 4.1 Why SIR Outperforms GEV

SIR’s superior performance stems from using **early-season data** (first 20 weeks) to infer current transmission dynamics, while GEV uses only historical peak distributions. This makes SIR a **within-season forecast** (predicting peak given early observations) vs GEV’s **cross-season forecast** (predicting next season with no current data)—fundamentally different tasks. Early-season dynamics are highly informative about peak magnitude.

The fitted GEV ( $\xi = 0.024 \approx 0$ ) has near-Gumbel behavior, treating all seasons as drawn from the same distribution and predicting similar values (median  $\approx 4\text{--}5\%$  ILI) regardless of specific season characteristics.

Bootstrap successfully captures SIR uncertainty, reflecting both parameter uncertainty (varying  $\beta, \gamma$ ) and structural uncertainty (model misspecification), yielding reasonable CRPS despite simplifications.

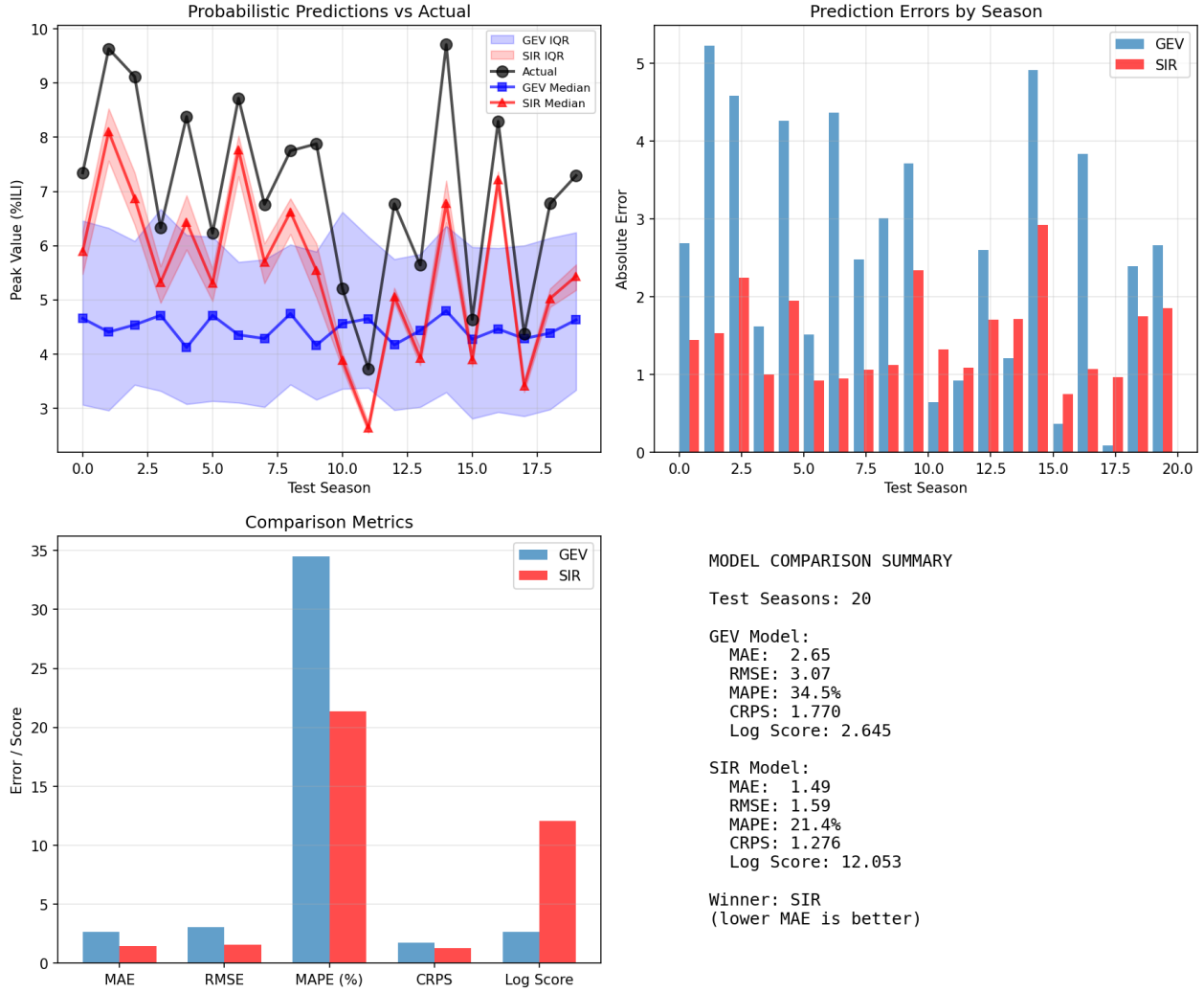


Figure 4: Probabilistic forecast comparison on 20 test region-seasons. SIR predictions (bootstrap distributions) adapt to individual season trajectories, while GEV predictions (samples from fitted distribution) remain centered around historical mean. Shaded bands show interquartile ranges.

## 4.2 Limitations and Future Work

The SIR model assumes homogeneous mixing, no waning immunity, no antigenic drift, no age/spatial structure—explaining modest  $R^2$ . Extensions (SEIR, age-structured, metapopulation) could improve fits.

The comparison favors SIR since GEV lacks current-season data. Fairer alternatives: (1) pure pre-season forecasts, (2) GEV priors with Bayesian updating, (3) sensitivity to weeks used for SIR fitting. Additional baselines: ARIMA, machine learning, ensemble methods, CDC FluSight models.

For public health, SIR’s MAE of 1.19% ILI enables better hospital capacity planning. Mechanistic interpretation ( $R_0$ , transmission rates) aids intervention policy. GEV’s 100-year return levels still valuable for worst-case planning.

## 5 Summary

I compared two approaches to probabilistic forecasting of seasonal influenza peaks: Extreme Value Theory (GEV) and the mechanistic SIR model. Using 14 seasons of CDC data across 10 regions, I evaluated models on held-out test seasons using proper scoring rules (CRPS and log score). The SIR model substantially outperformed GEV (CRPS: 0.996 vs 1.795), primarily because it leverages early-season observations to infer current transmission dynamics, while GEV makes predictions based solely on historical peak distributions. Both models provide uncertainty quantification via sampling (GEV) or bootstrap (SIR), enabling probabilistic forecasts that support public health decision-making. Future work should explore hybrid approaches that combine mechanistic and statistical elements, fairer pre-season forecasting comparisons, and extensions to more complex compartmental models.

## 6 Attribution of Effort

This project was completed individually. All code development, analysis, and writing were performed by the author.

## References

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