



Spatial Modeling of *P. ramorum* in California:

Strengths and weaknesses of rule-based, statistical, and cellular automata methods

**Ross K. Meentemeyer
University of North Carolina at Charlotte
Department of Geography & Earth Sciences**

I. Rule-Based Modeling

- incorporates epidemiological and biological factors driving establishment and spread in California plant communities
 - Host susceptibility
 - Host epidemiology
 - Moisture
 - Temperature

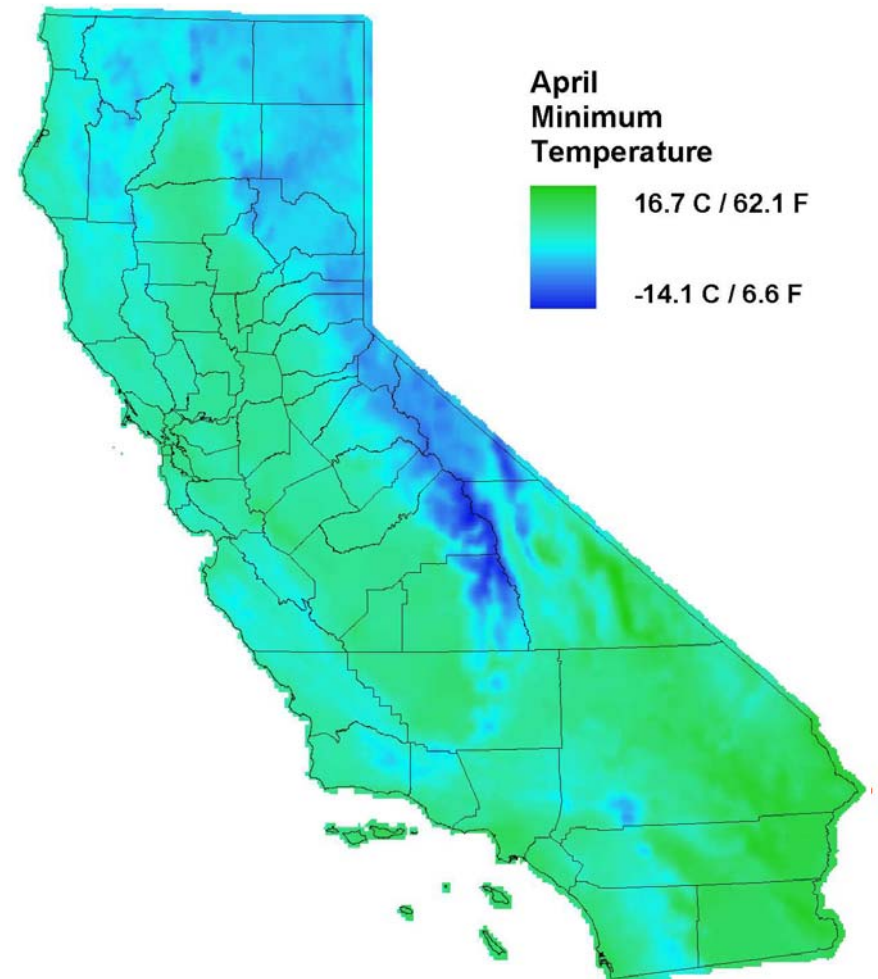


Parameterizing the Model

- Scored 5 variables to encode magnitude & direction of their effect on disease spread

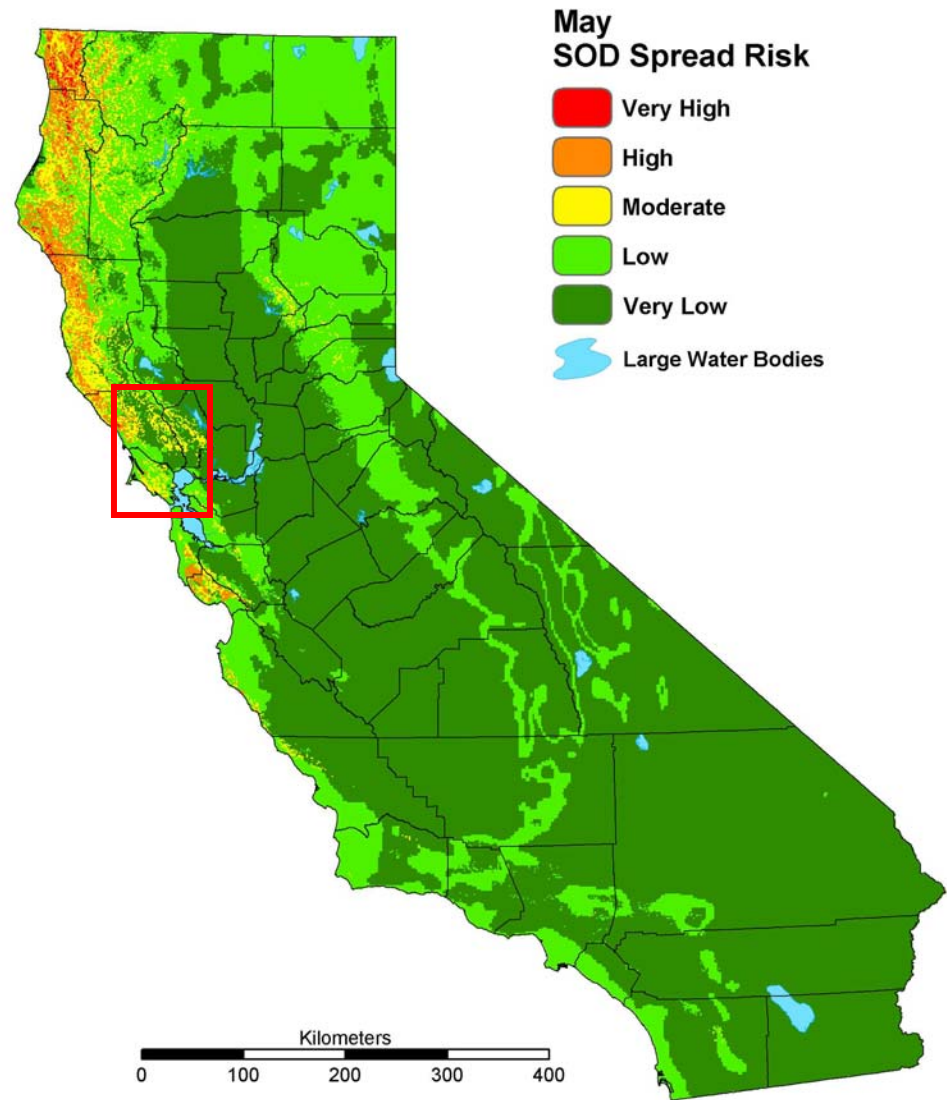
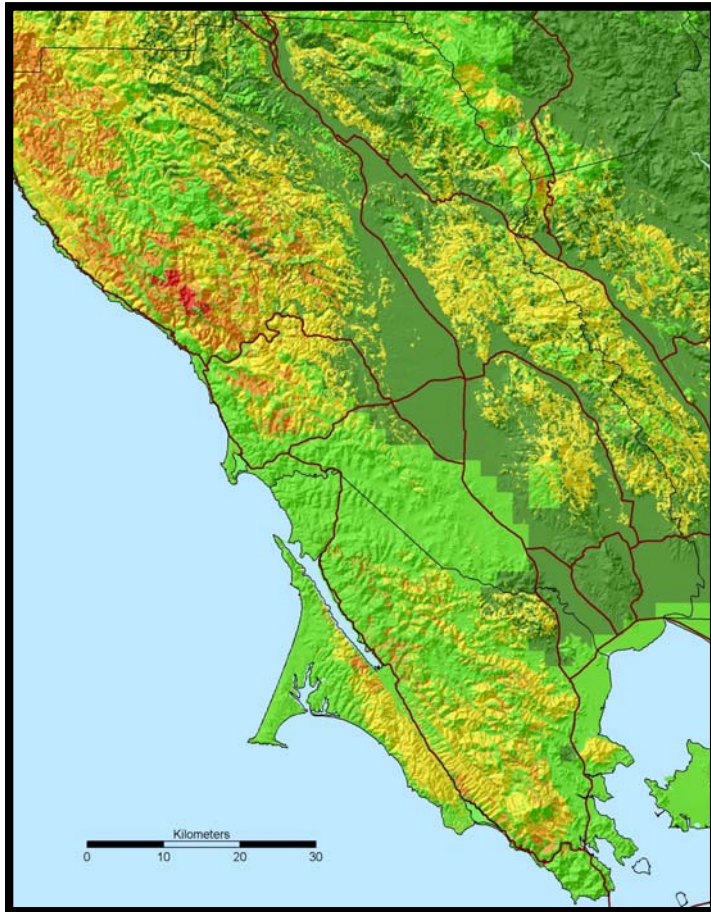
Variables

- Host species abundance
- Precipitation
- Relative Humidity
- Maximum Temperature
- Minimum Temperature



Model Results

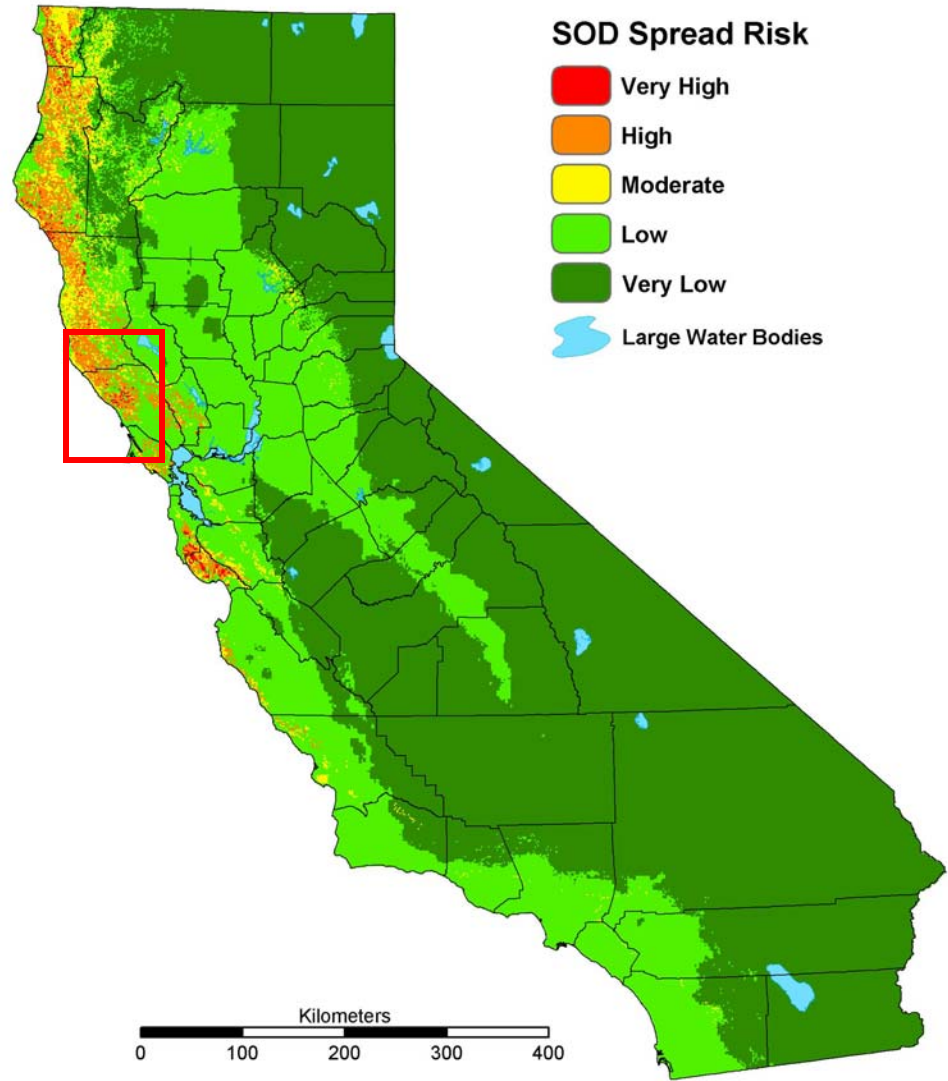
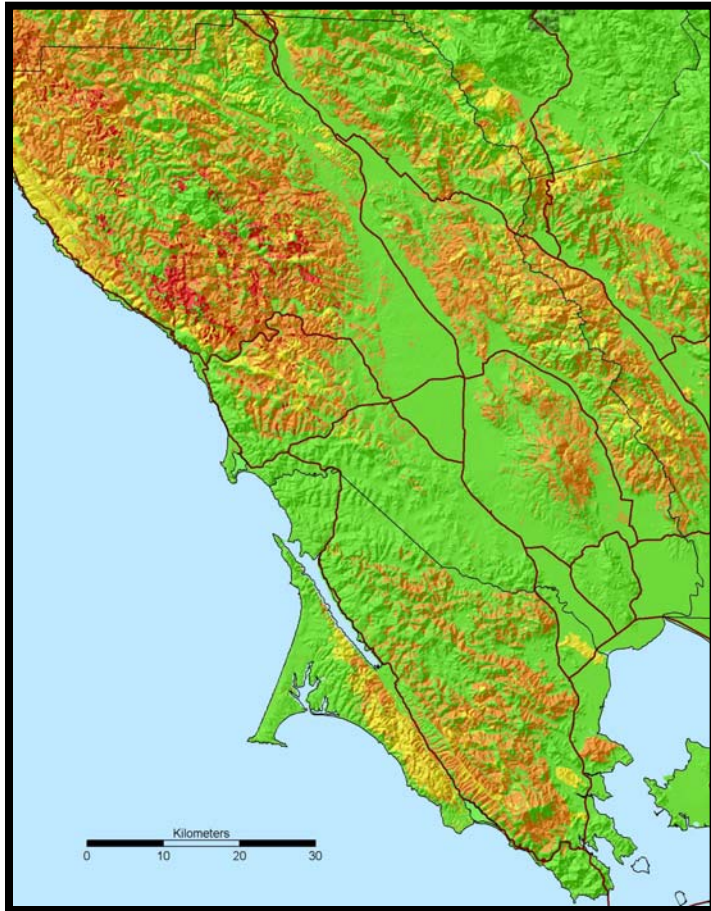
Seasonal Changes: December - May



Meentemeyer et al. 2004 (*Forest Ecology & Management*)

Model Results

Cumulative Risk: December - May



Meentemeyer et al. 2004 (*Forest Ecology & Management*)

Rule-Based Modeling

Strengths

- simple methods
- works well with limited data on pathogen survival and transmission

Limitations

- too simple
- may not reflect real field conditions
- risk-based (not probabilistic) and static

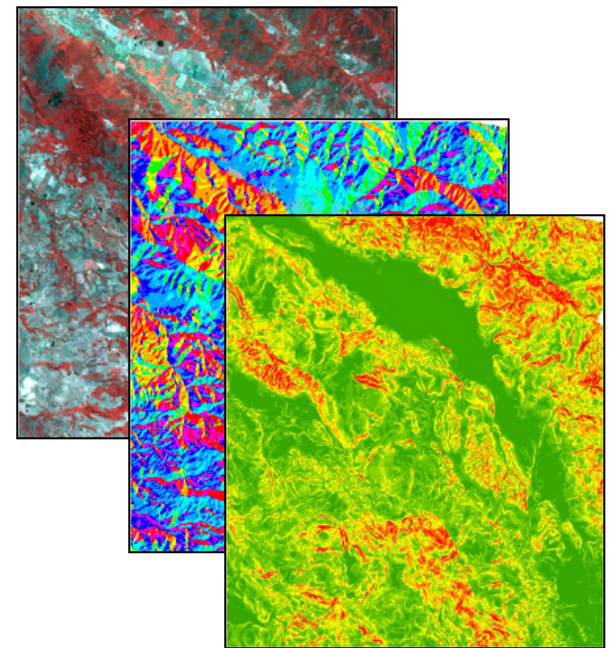
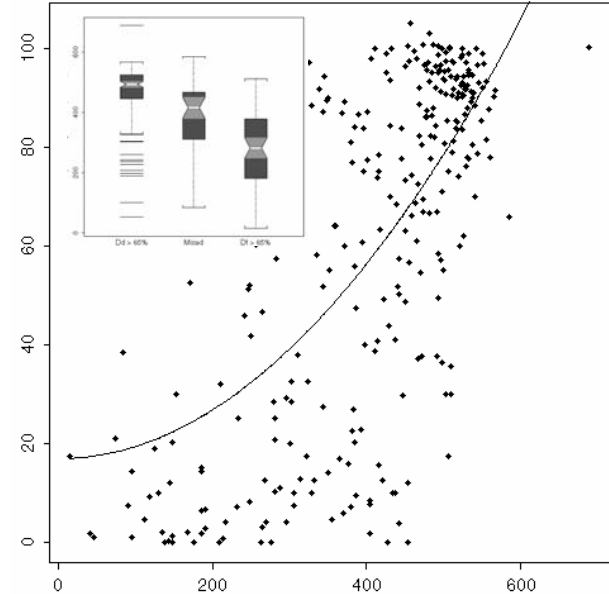
II. Statistical Modeling

- based on observed relationships between pathogen occurrence, host composition, and environmental conditions

Example Methods

- Regression
 - CART
 - Neural Networks
- require presence & absence data

- once a model is developed, can apply its equation across mapped variables in a GIS



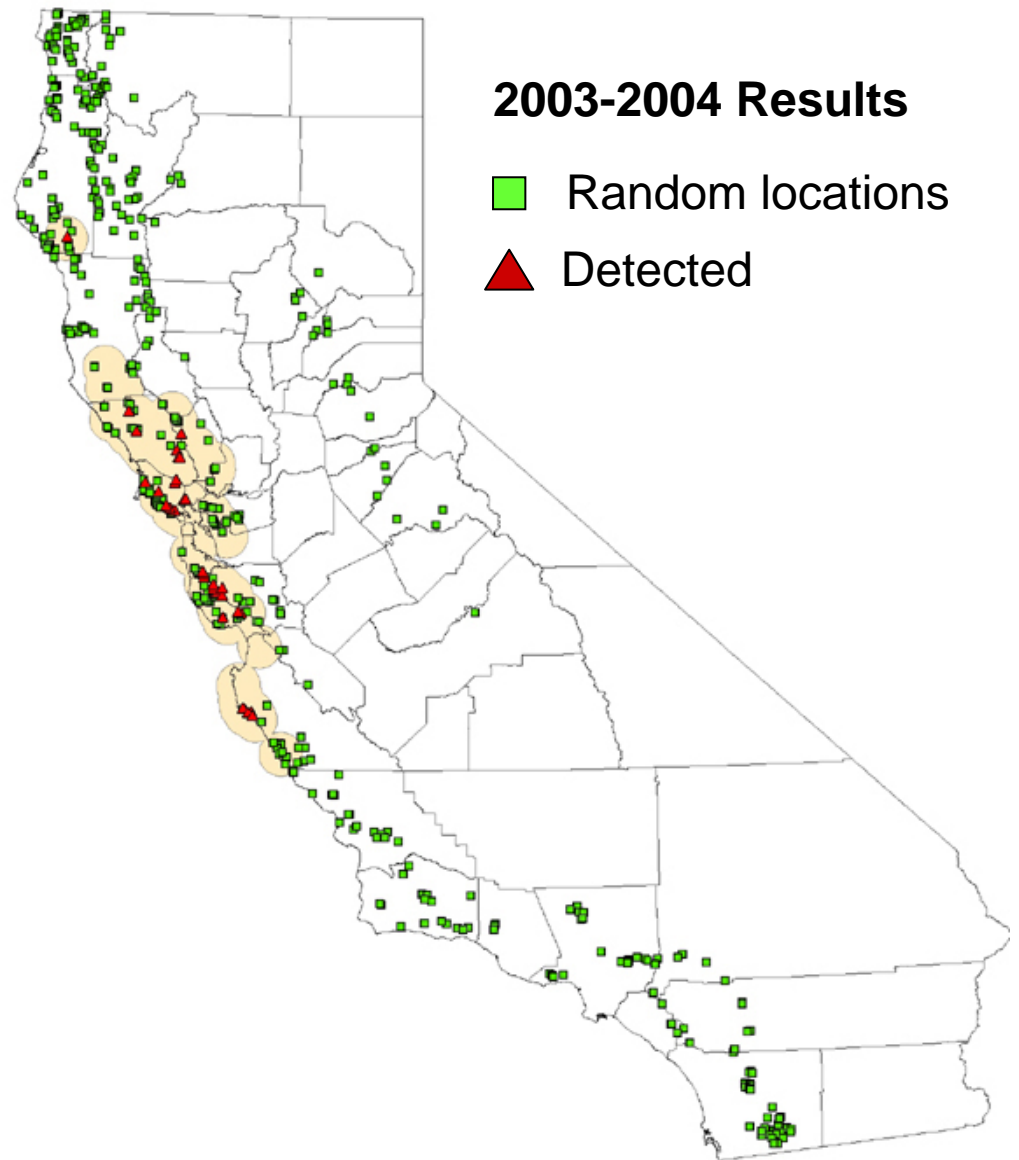
Response Variable

- extensive dataset on distribution *P. ramorum*
- within high-risk forests, we have surveyed 495 random locations (2003-2004)
- *P. ramorum* detected at 33 of the 495 sites.

n = 9 of 139 in 2003

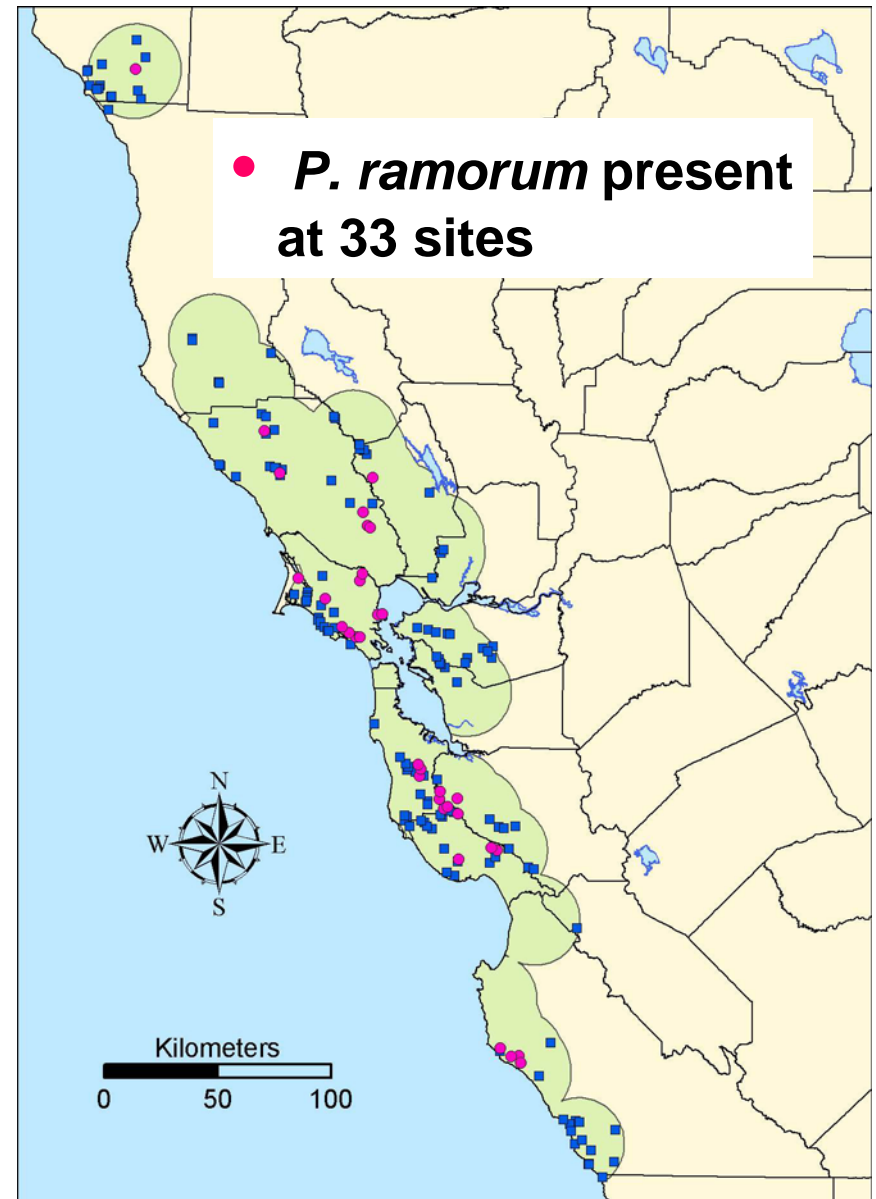
n = 24 of 347 in 2004

- 20 km was farthest detection



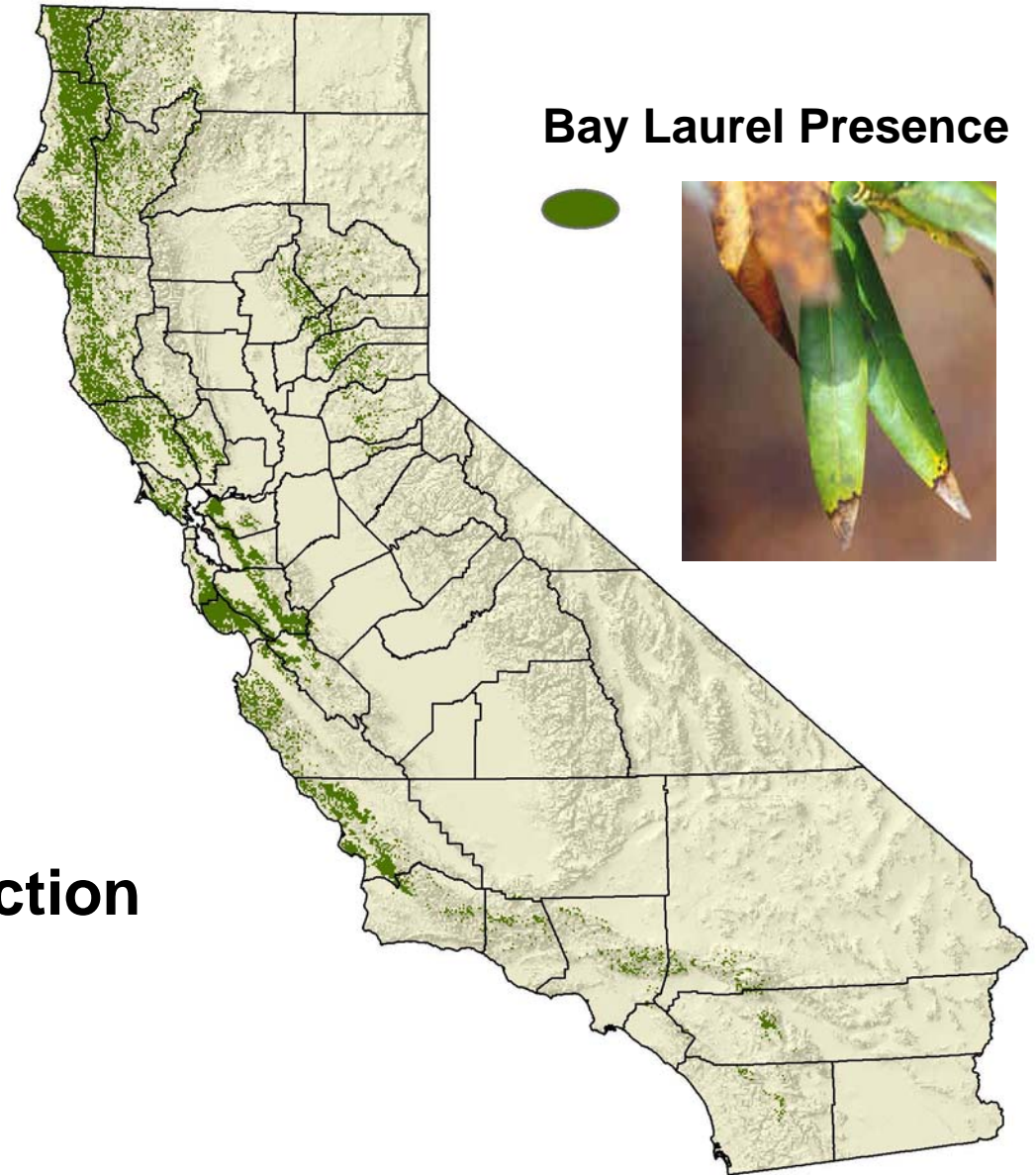
Defining the Analysis Region

- mapped COMTF sites
- limited to areas < 20 km of confirmed infection sites
- yielded **166** random sites from survey

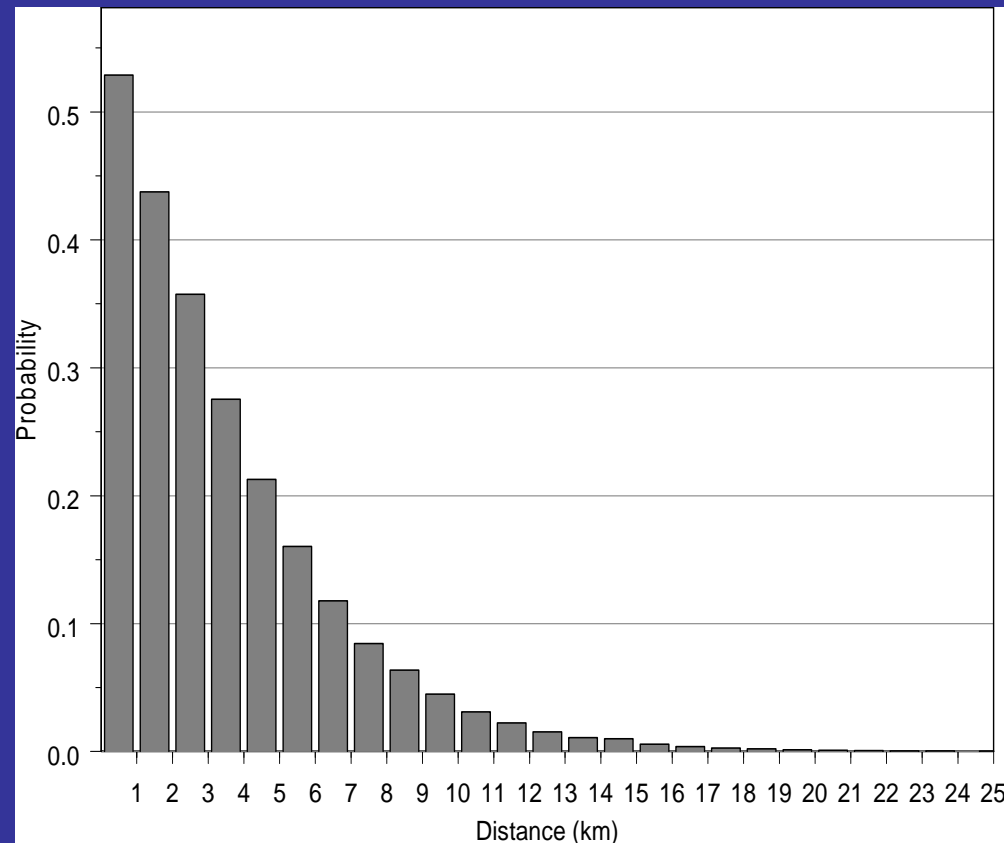


Predictor Variables

- Population pressure
- Climate conditions
- Fire history
- Presence of bay laurel & tanoak
- Distance to known infection



Statistically-Based Model



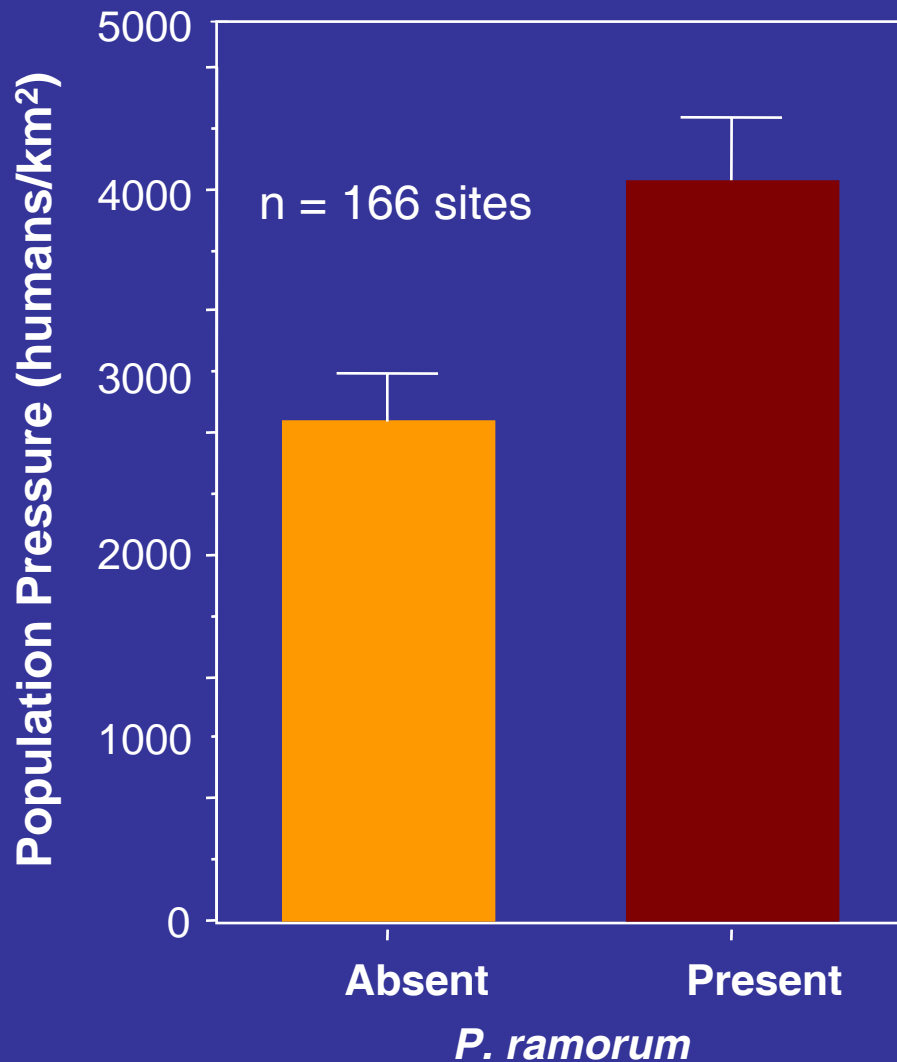
Logistic Regression ($r^2 = 0.51$)

Distance (-)	p=0.0000
Pop. pressure (+)	p=0.0001
Bay laurel (+)	p=0.0002
Precipitation (+)	p=0.0001
Pop. pressure x Precip.	p=0.0001

Eliminated Variables

Min. & Max. Temperature
Relative Humidity
Fire History
Tanoak presence

Statistically-Based Model



Logistic Regression ($r^2 = 0.51$)

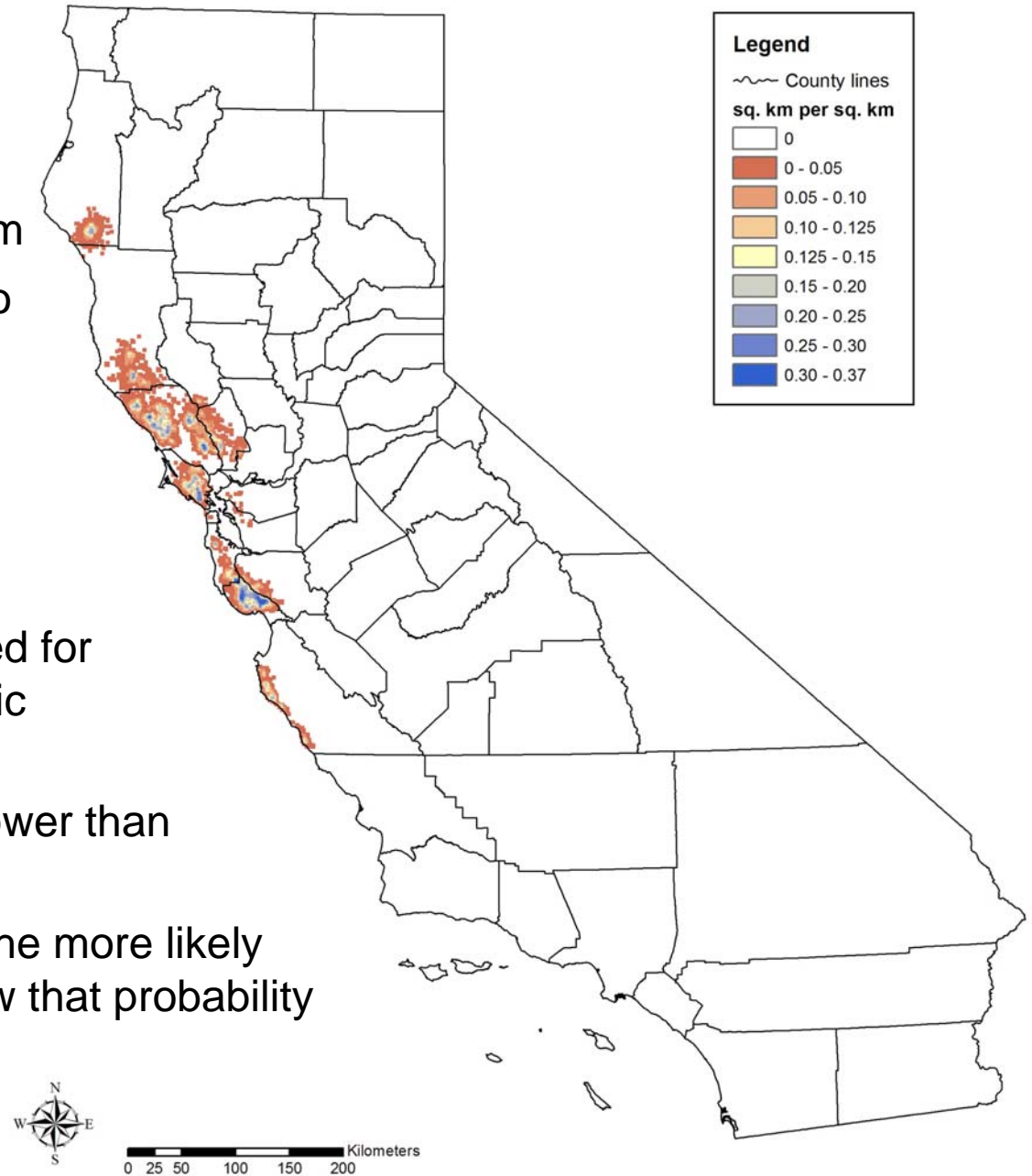
Distance (-)	p=0.0000
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Pop. pressure x Precip.	p=0.0001

Eliminated Variables

Min. & Max. Temperature
Relative Humidity
Fire History
Tanoak presence

Application and Simulation in GIS

- Mapped all variables at 50 x 50 m
- logistic regression was applied to each variable to produce a map of occurrence probability (0-1)
- used Monte Carlo simulation to predict distribution across unsampled regions
- a random number (0-1) generated for each cell and compared to logistic probability
- cell infected if random number lower than probability of infection
- higher the predicted probability the more likely the random number will fall below that probability



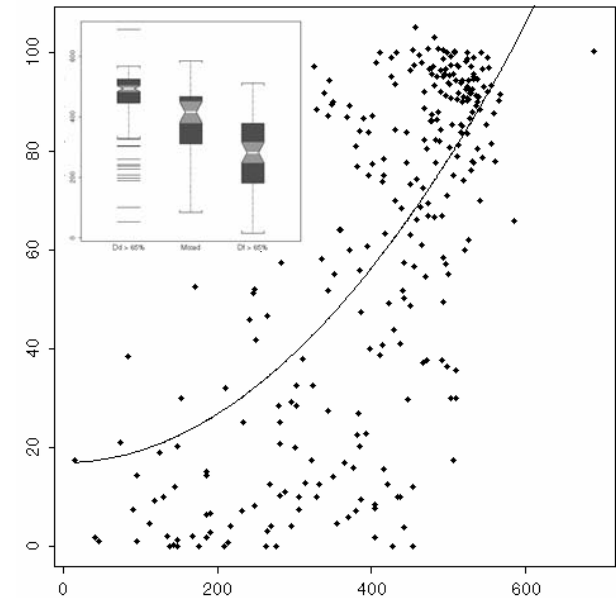
Statistical Modeling

Strengths

- established methods like regression
- empirical relationships across study system facilitate ecological and epidemiological understanding

Limitations

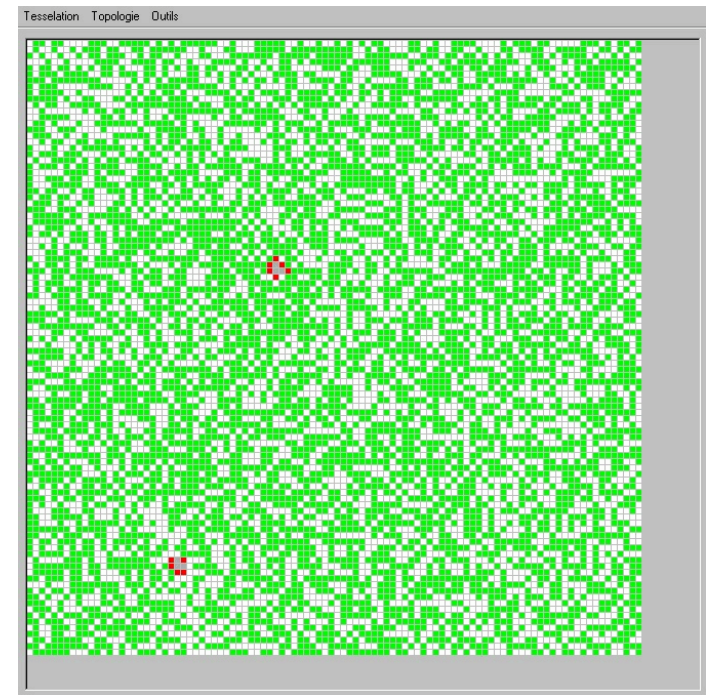
- susceptible sites have not yet been exposed (distance variables partially offset this)
- probabilistic, but still static (not dynamic)
- need dynamic models of establishment and spread driven by weather events and dispersal vectors



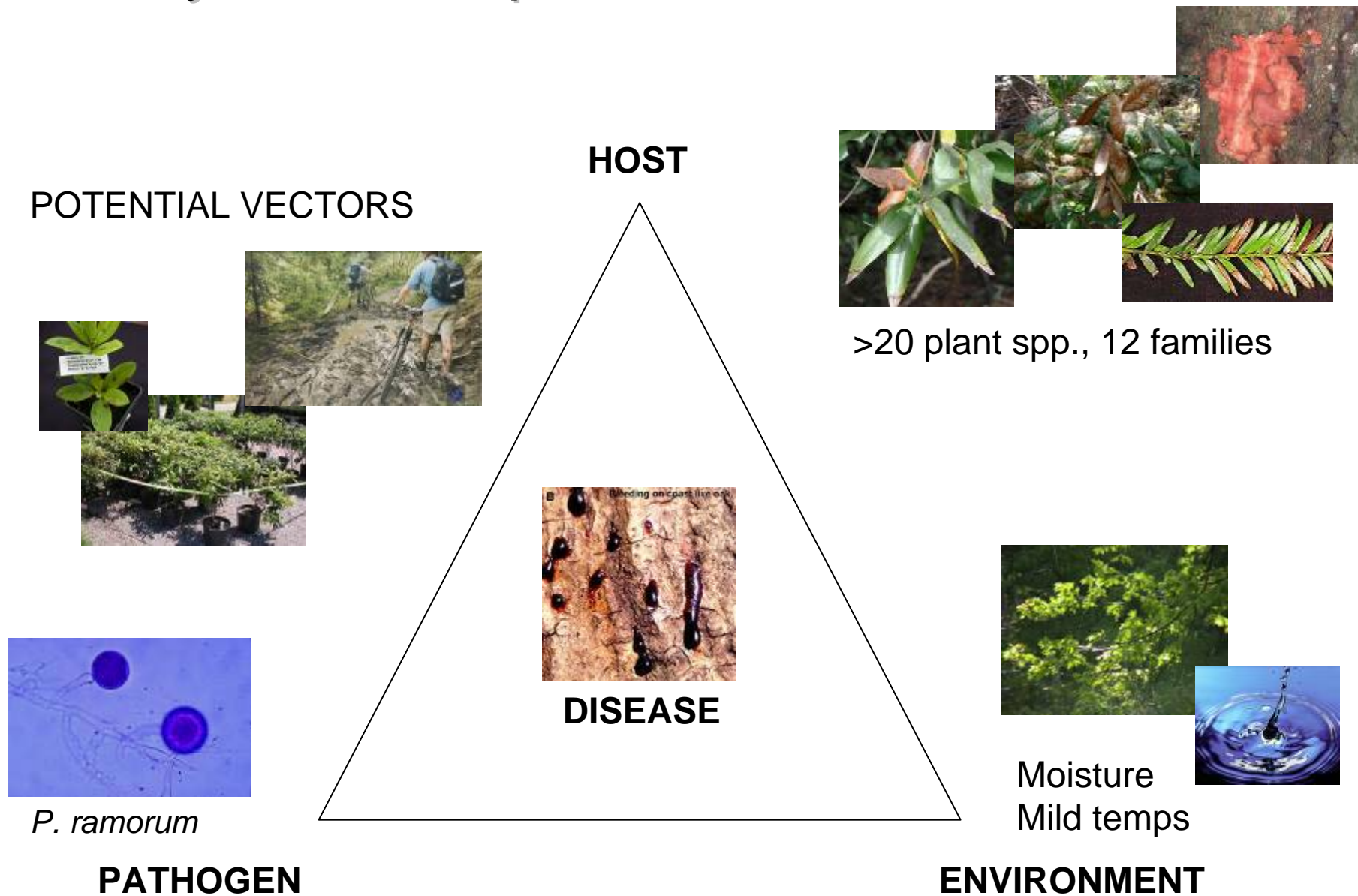
III. Cellular Automata Modeling

- powerful approach for modeling processes driving establishment & spread
- may conduct repeatable non-destructive experiments at large scales
 - effect of extreme weather
 - efficacy of management practices
- requires parameterization of system components across a grid and specified time steps (e.g. 1hr, 1day)

Spread of a Wildfire



System Components of *P. ramorum*



System Components

In the model

$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$

P = Probability of infection

S = Host susceptibility

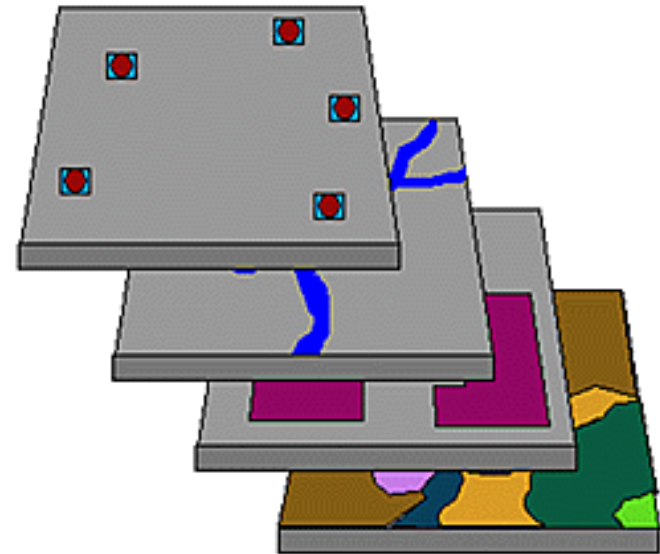
T = Temperature

P = Precipitation

I = Surrounding disease levels

D = Distance

H = Human vectors



Model Approach

- Probability of infection modeled for each cell based on system components
 - Number between zero and one

Model landscape
Time step 1

0.08	0.21	0.11	0	0
0	0.15	0.50	0.01	0
0.04	0.02	0.13	0.18	0
0.42	0	0.26	0.06	0.22
0	0.26	0	0.04	0

Model Approach

- Probability of infection modeled for each cell based on system components
 - Number between zero and one
- Weighted random process
 - Generate random numbers between zero and one

Model landscape
Time step 1

0.08	0.21	0.11	0	0
0.15	0.23	0.79		
0	0.15	0.50	0.01	0
	0.10	0.52	0.34	
0.04	0.02	0.13	0.18	0
0.07	0.86	0.25	0.97	
0.42	0	0.26	0.06	0.22
0.64		0.29	0.11	0.73
0	0.26	0	0.04	0
	0.84		0.12	

Model Approach

- Probability of infection modeled for each cell based on system components
 - Number between zero and one
- Weighted random process
 - Generate random numbers between zero and one
- Compare probability to random numbers
 - **Infected** – *Probability > Random*
 - **Uninfected** – *Probability < Random*
 - No host vegetation (0)

Model landscape
Time step 1

0.08 0.15	0.21 0.23	0.11 0.79	0	0
0	0.15 0.10	0.50 0.52	0.01 0.34	0
0.04 0.07	0.02 0.86	0.13 0.25	0.18 0.97	0
0.42 0.64	0	0.26 0.29	0.06 0.11	0.22 0.73
0	0.26 0.84	0	0.04 0.12	0

Model Approach

- Probability of infection modeled for each cell based on system components
 - Number between zero and one
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 - Generate random numbers between zero and one
- Compare probability to random numbers
 - **Infected** – $Probability > Random$
 - **Uninfected** – $Probability < Random$
 - No host vegetation (0)
- Outputs disease intensity as count of times cell is infected

Model landscape
Time step 1

0	0	0	0	0
0	1	0	0	0
0	0	0	0	0
0	0	0	0	0
0	0	0	0	0

Model Approach

- Probability of infection modeled for each cell based on system components
 - Number between zero and one
- Weighted random process
 - Generate random numbers between zero and one
- Compare probability to random numbers
 - **Infected** – $Probability > Random$
 - **Uninfected** – $Probability < Random$
 - No host vegetation (0)
- Outputs disease intensity as count of times cell is infected
- Iterates on a weekly time step

Model landscape
Time step 2

0	1	0	0	0
0	2	1	0	0
0	0	1	0	0
0	0	0	0	0
0	0	0	0	0

Model Approach

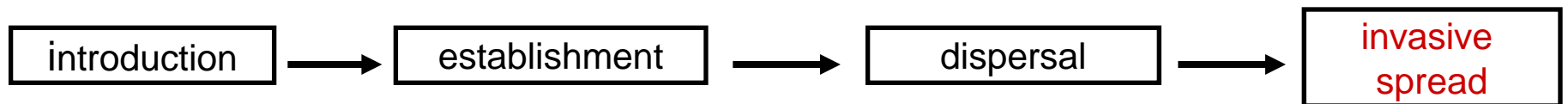
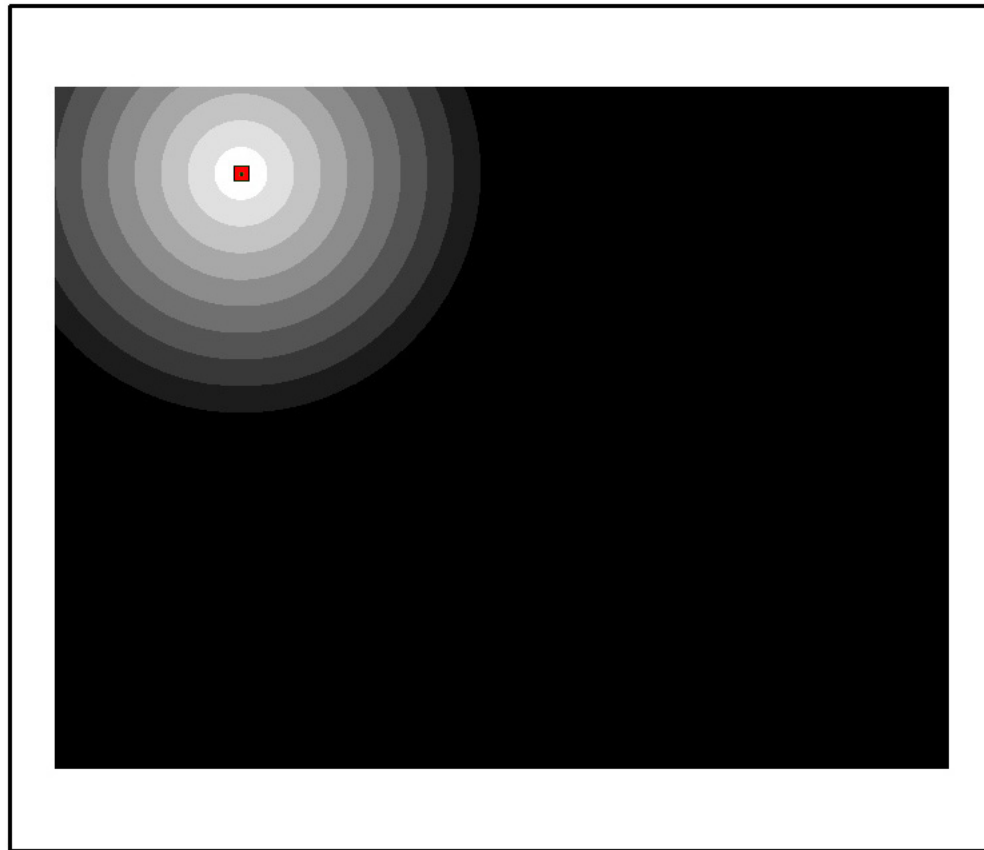
- Probability of infection modeled for each cell based on system components
 - Number between zero and one
- Weighted random process
 - Generate random numbers between zero and one
- Compare probability to random numbers
 - **Infected** – $Probability > Random$
 - **Uninfected** – $Probability < Random$
 - No host vegetation (0)
- Outputs disease intensity as count of times cell is infected
- Iterates on a weekly time step

Model landscape
Time step 3

0	1	0	0	0
0	3	2	0	0
0	0	1	0	0
0	0	0	0	0
0	0	0	1	0

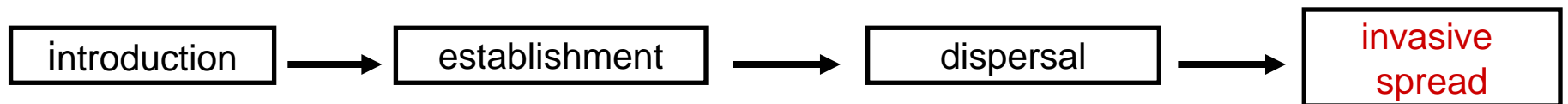
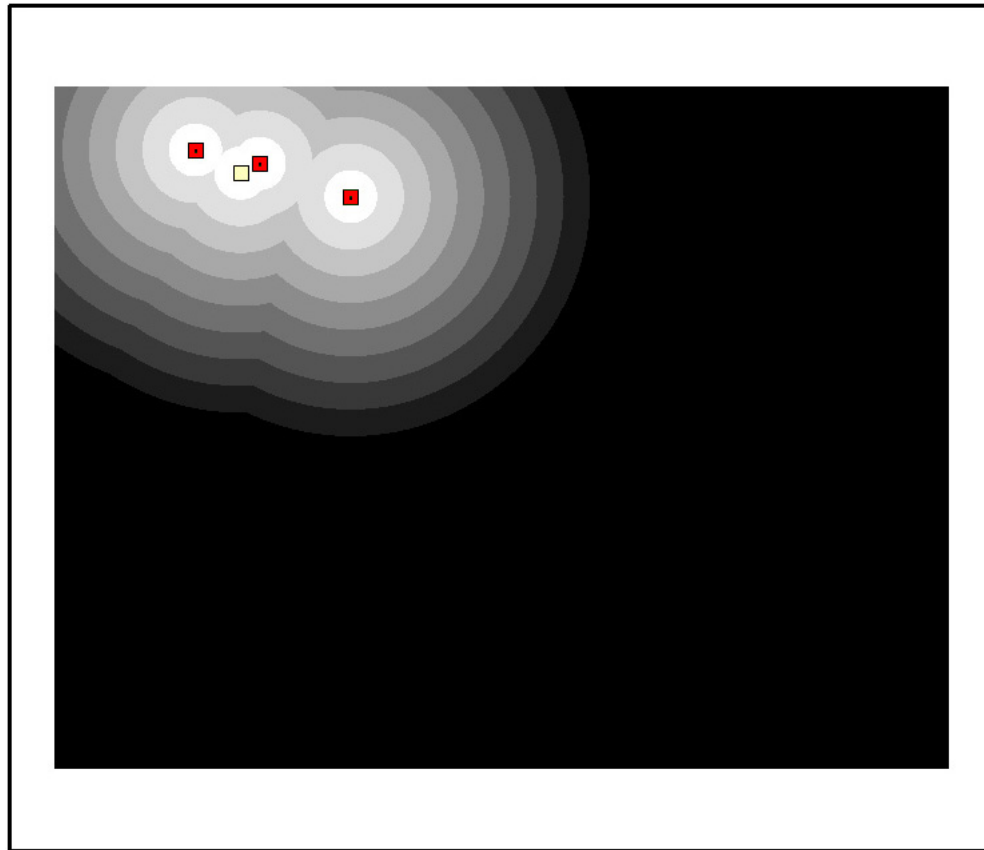
Model Approach

Example Simulation



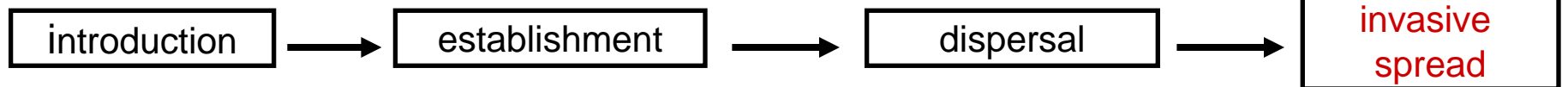
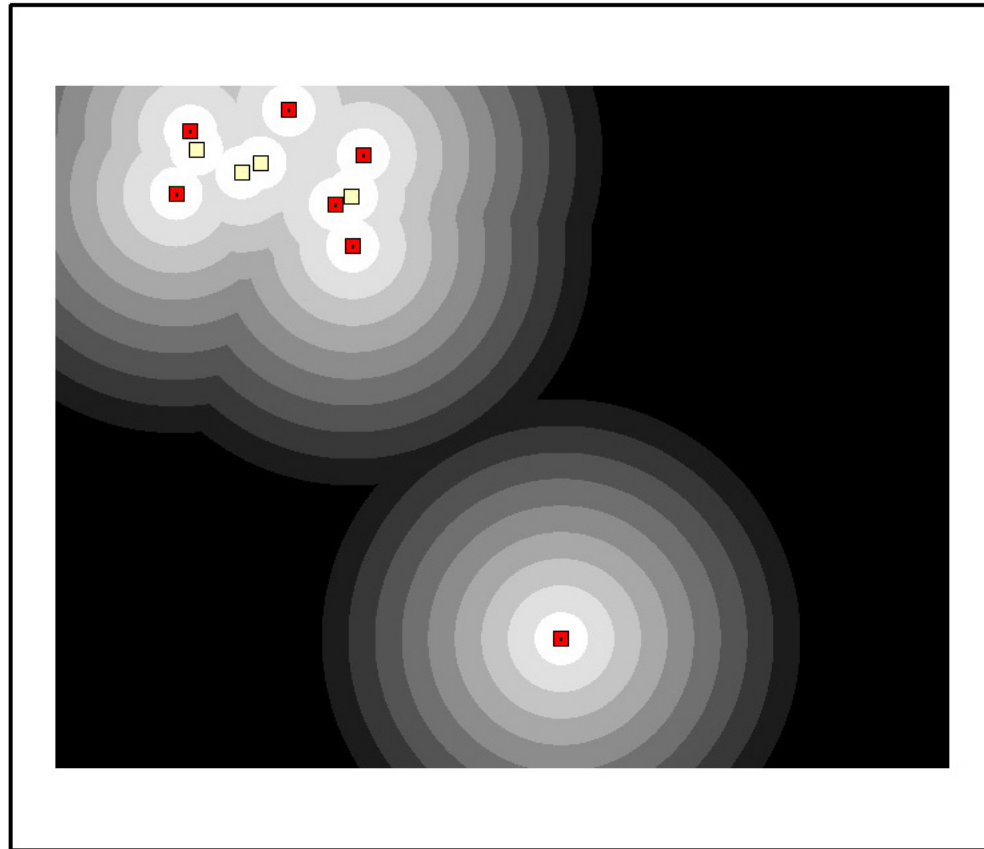
Model Approach

Example Simulation



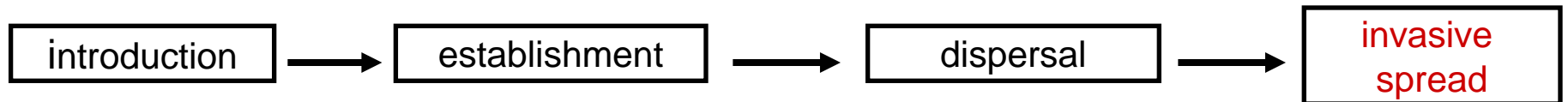
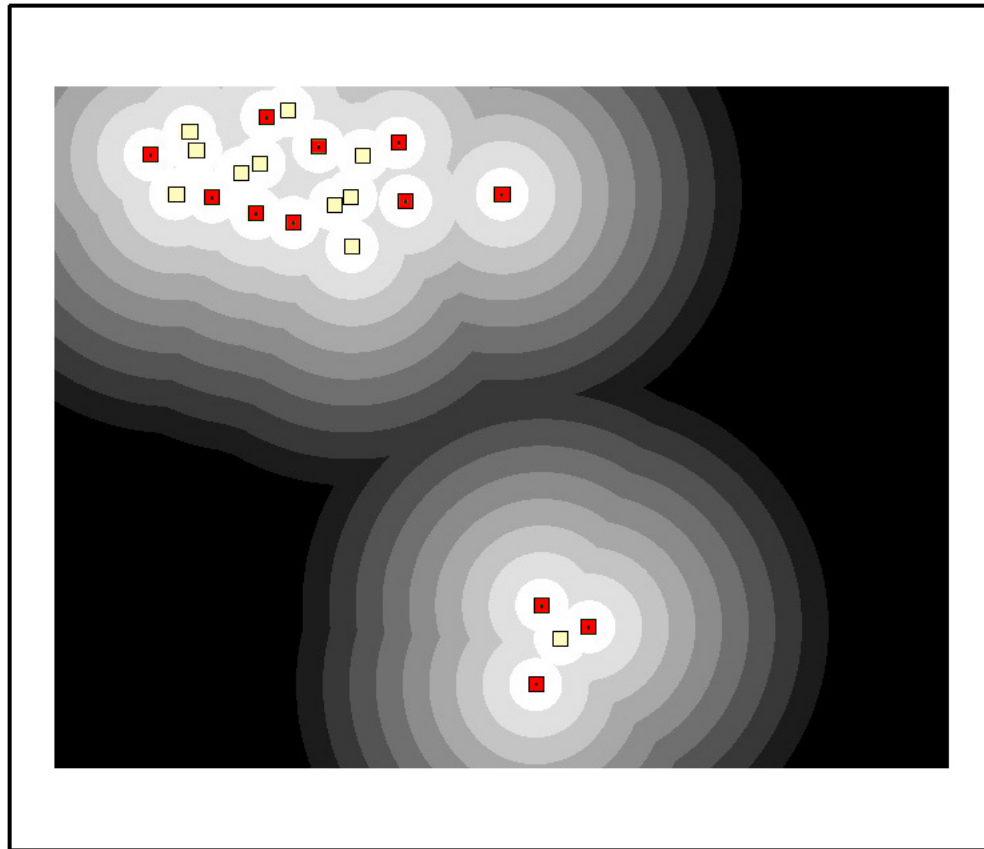
Model Approach

Example Simulation



Model Approach

Example Simulation



Mapping and Parameterizing System Components



Spatial Grain = 250 m cells

Time Step = 1 week

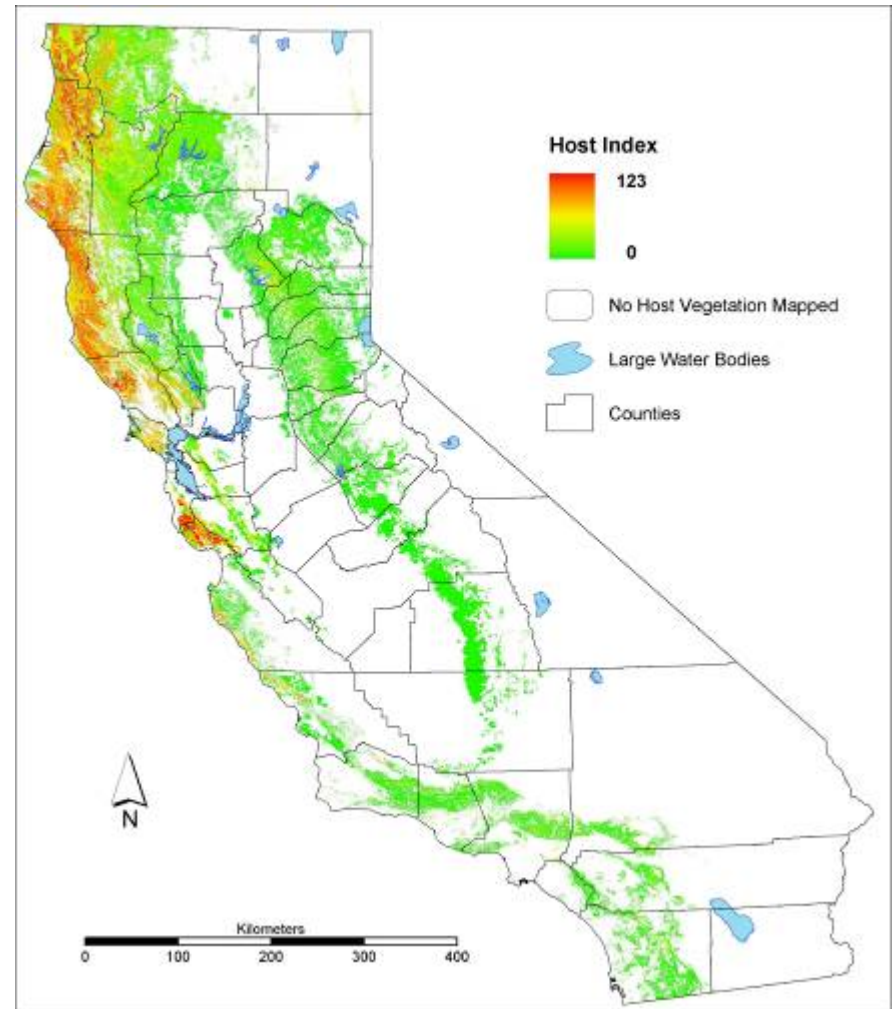
Host Susceptibility

Parameterization

Host Index (S)

- Meentemeyer *et al.* 2004
- Ranked susceptibility of each species
- Using CALVEG data, each cell received a composite score based on susceptibility rank and abundance of all species present
- Output scaled to 100 points

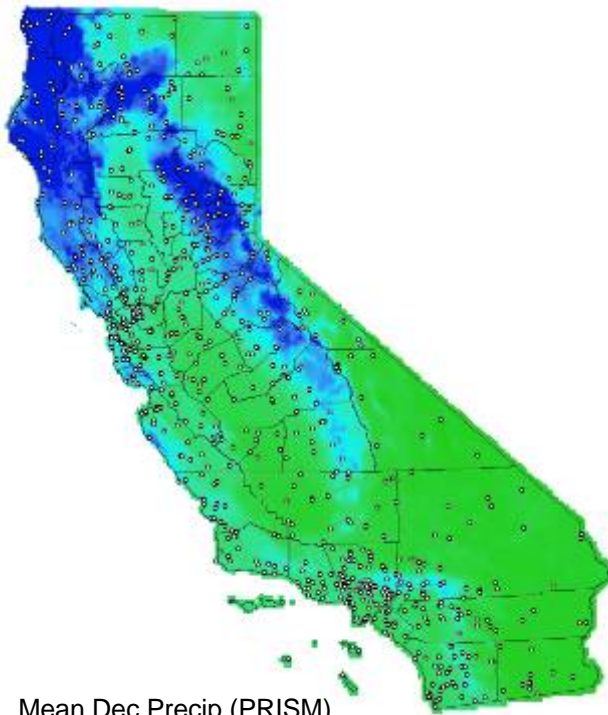
$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$



Weather Events

Mapping Methods

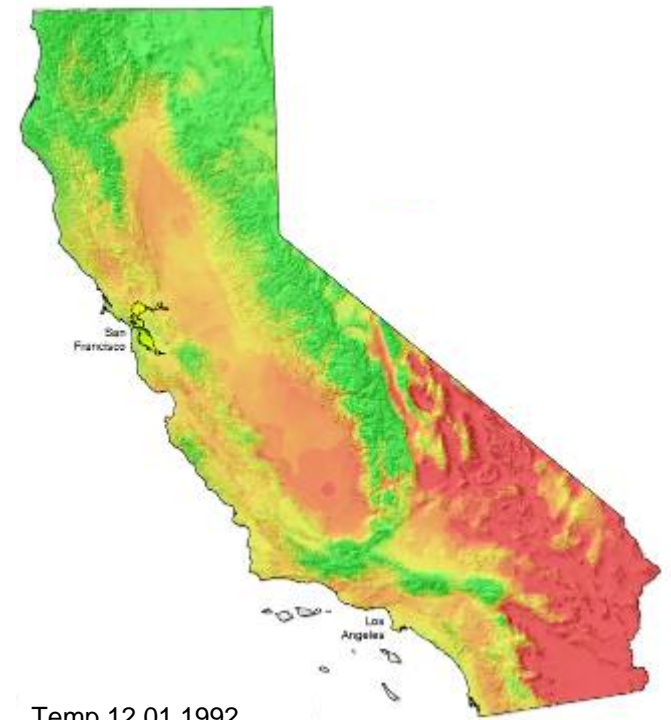
- Daily precip & temp at NCDC-NOAA point locations
- Integrate point data with long-term average grids (PRISM)



Mean Dec Precip (PRISM)
NCDC Base Stations



Precip 12.01.1992

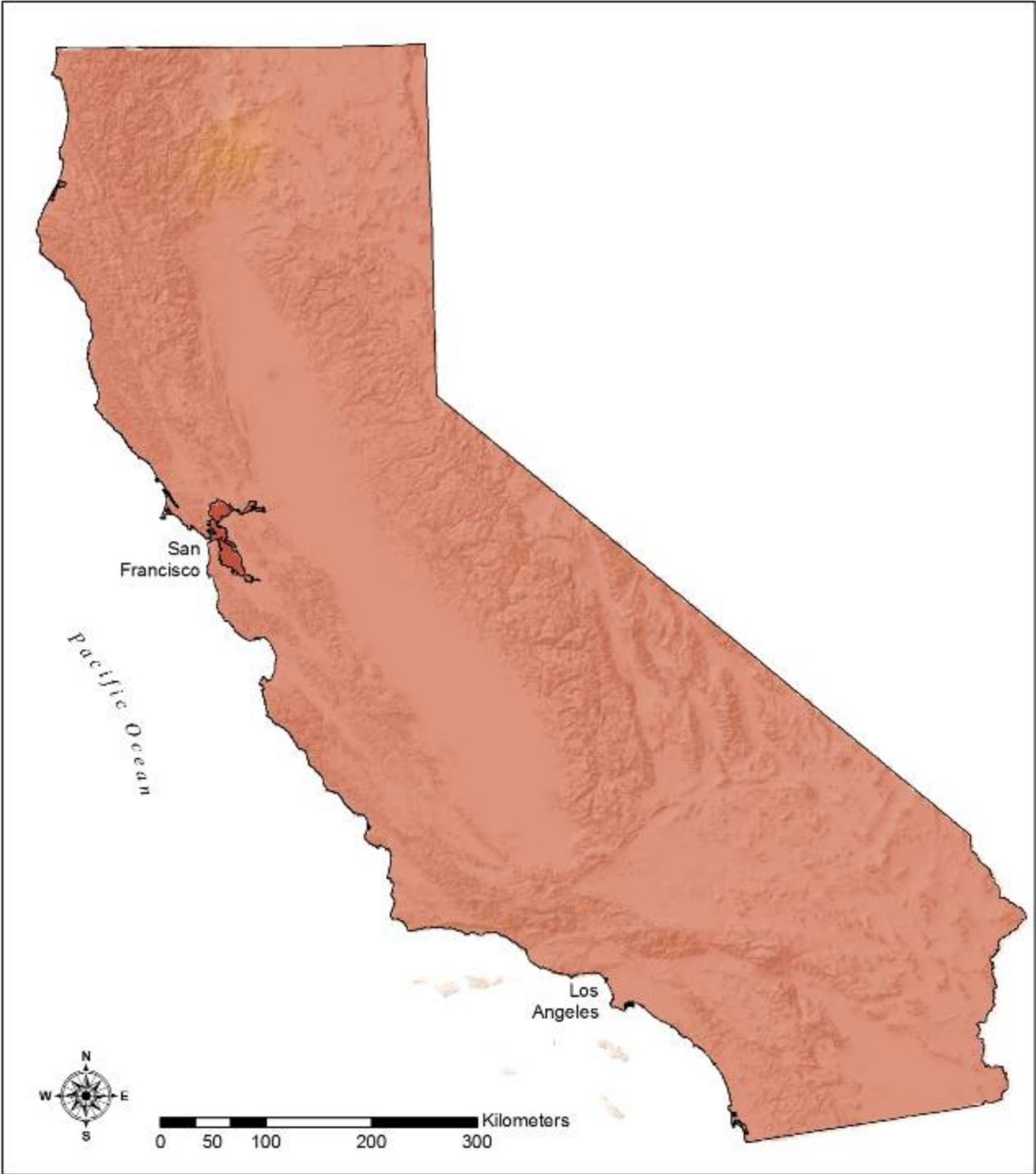


Temp 12.01.1992

Hunter & Meentemeyer (in press) *J. Appl. Meteor.*

PRECIPITATION

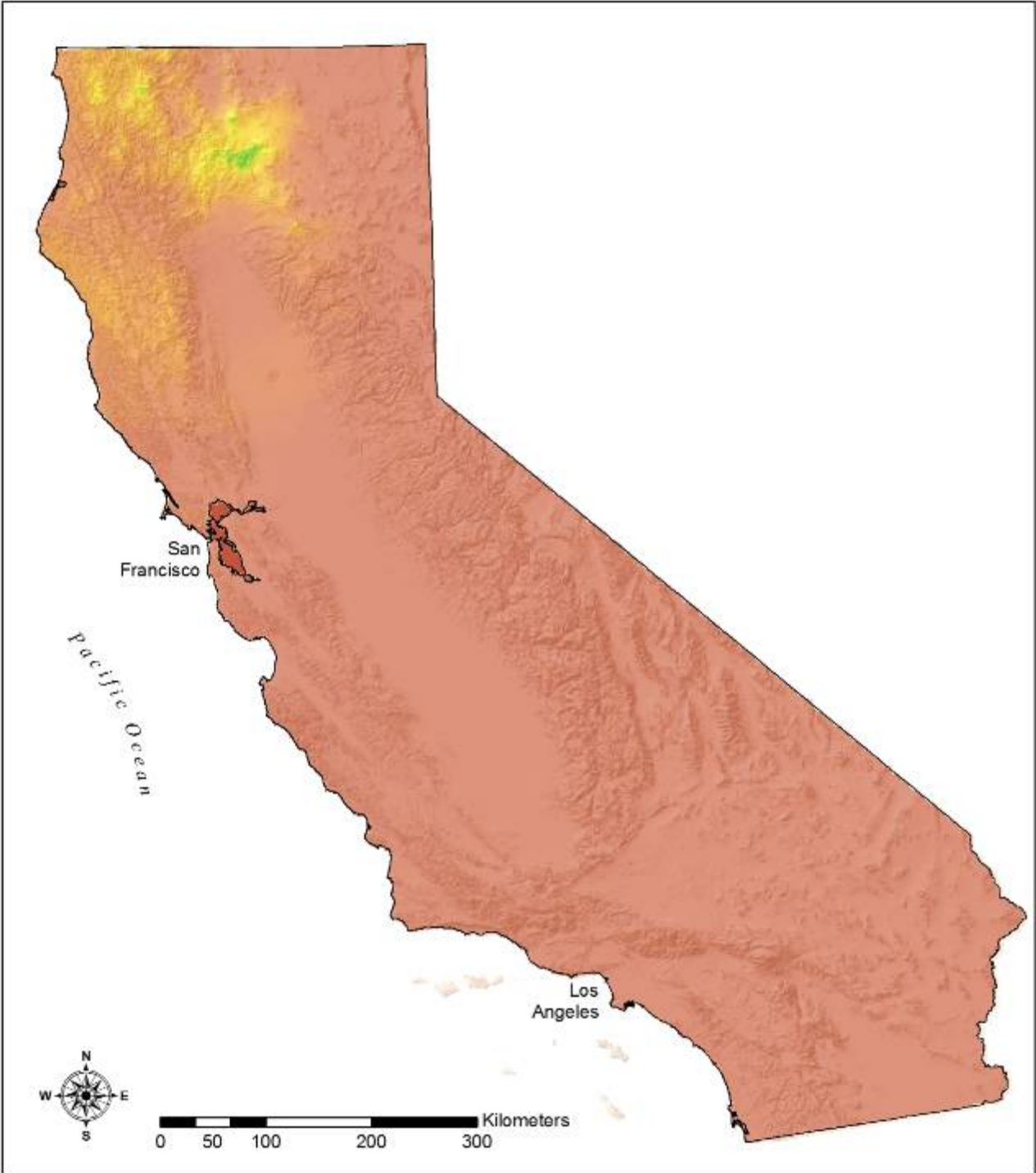
March 20, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

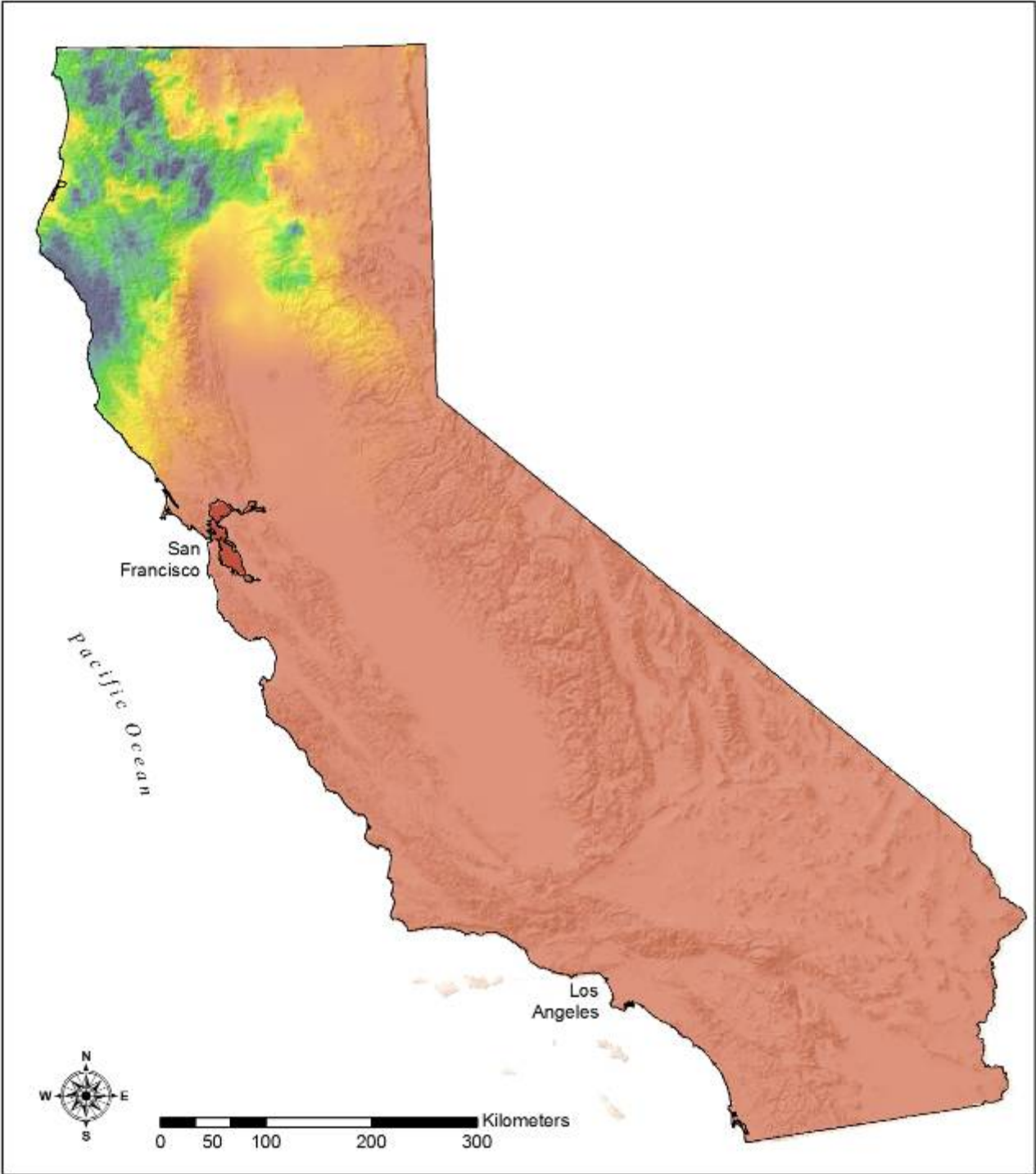
March 21, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

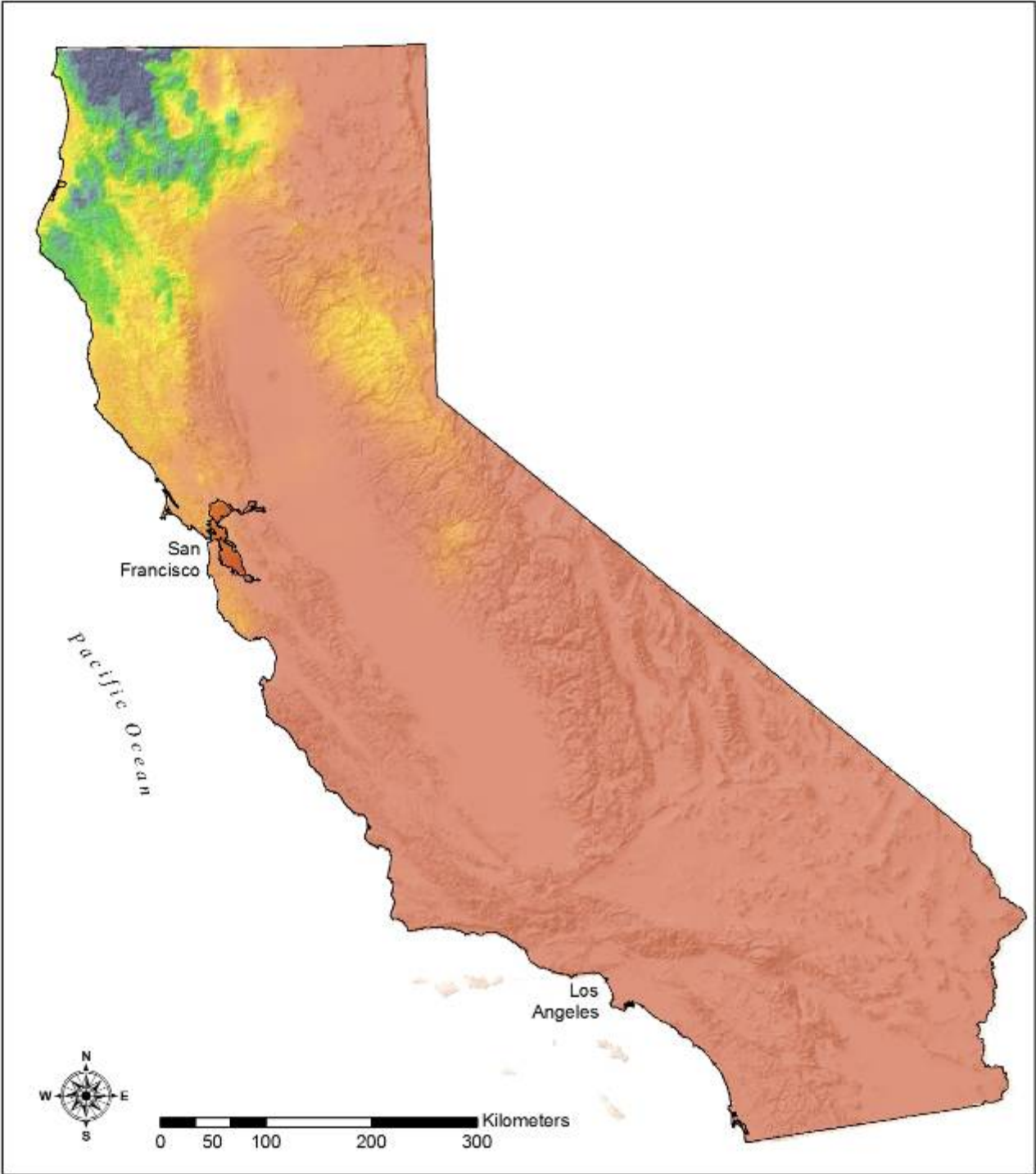
March 22, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

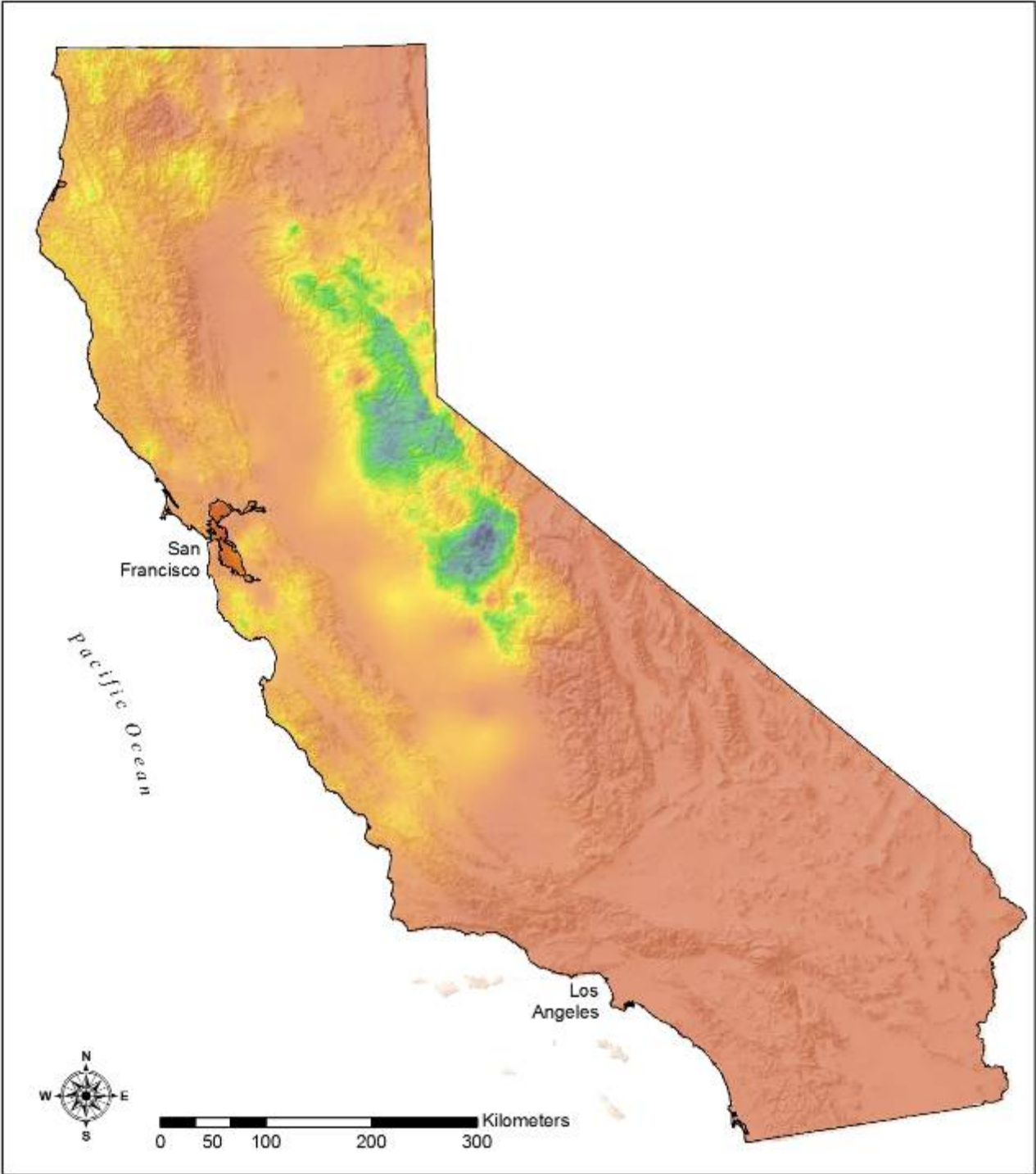
March 23, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

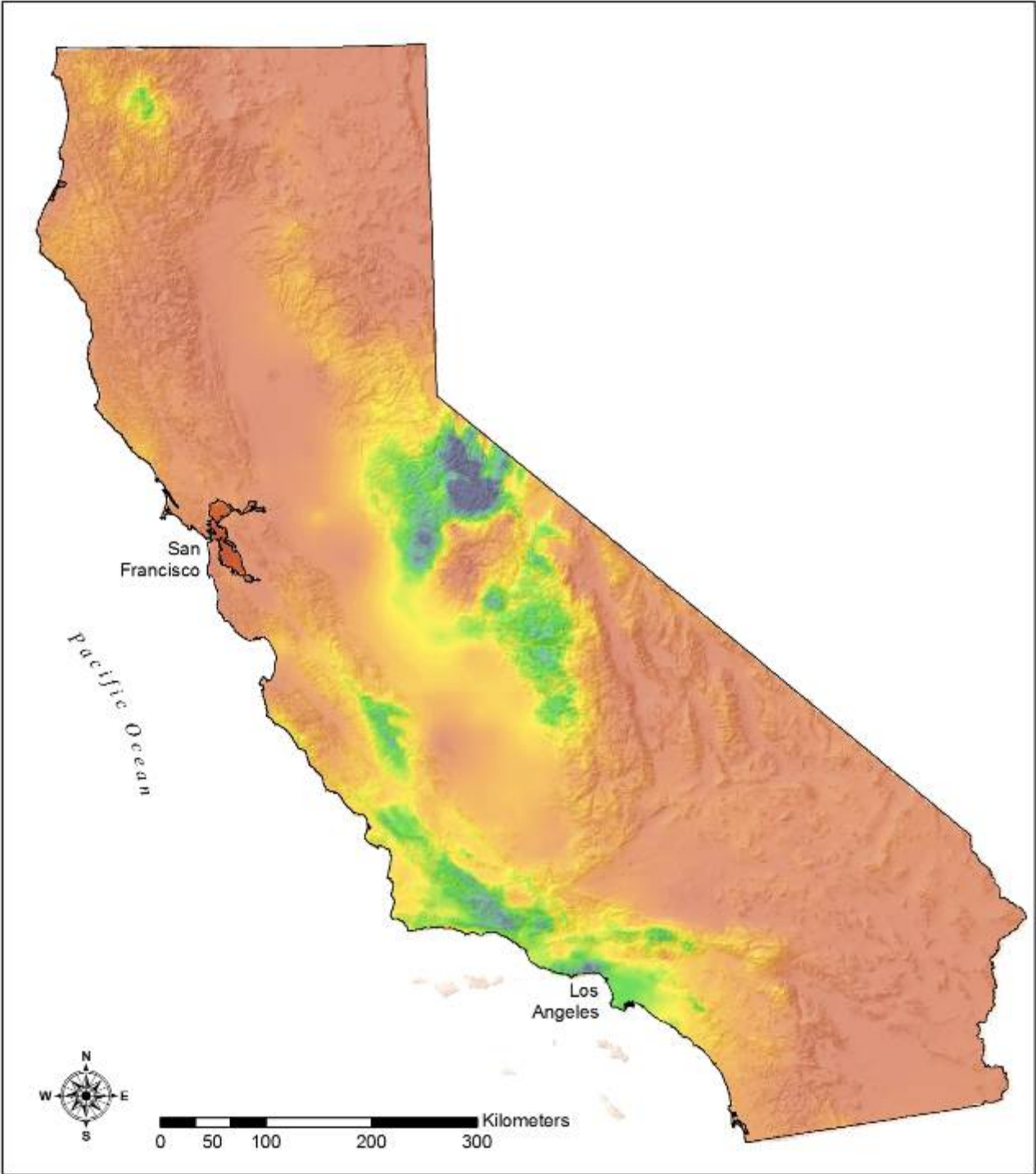
March 24, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

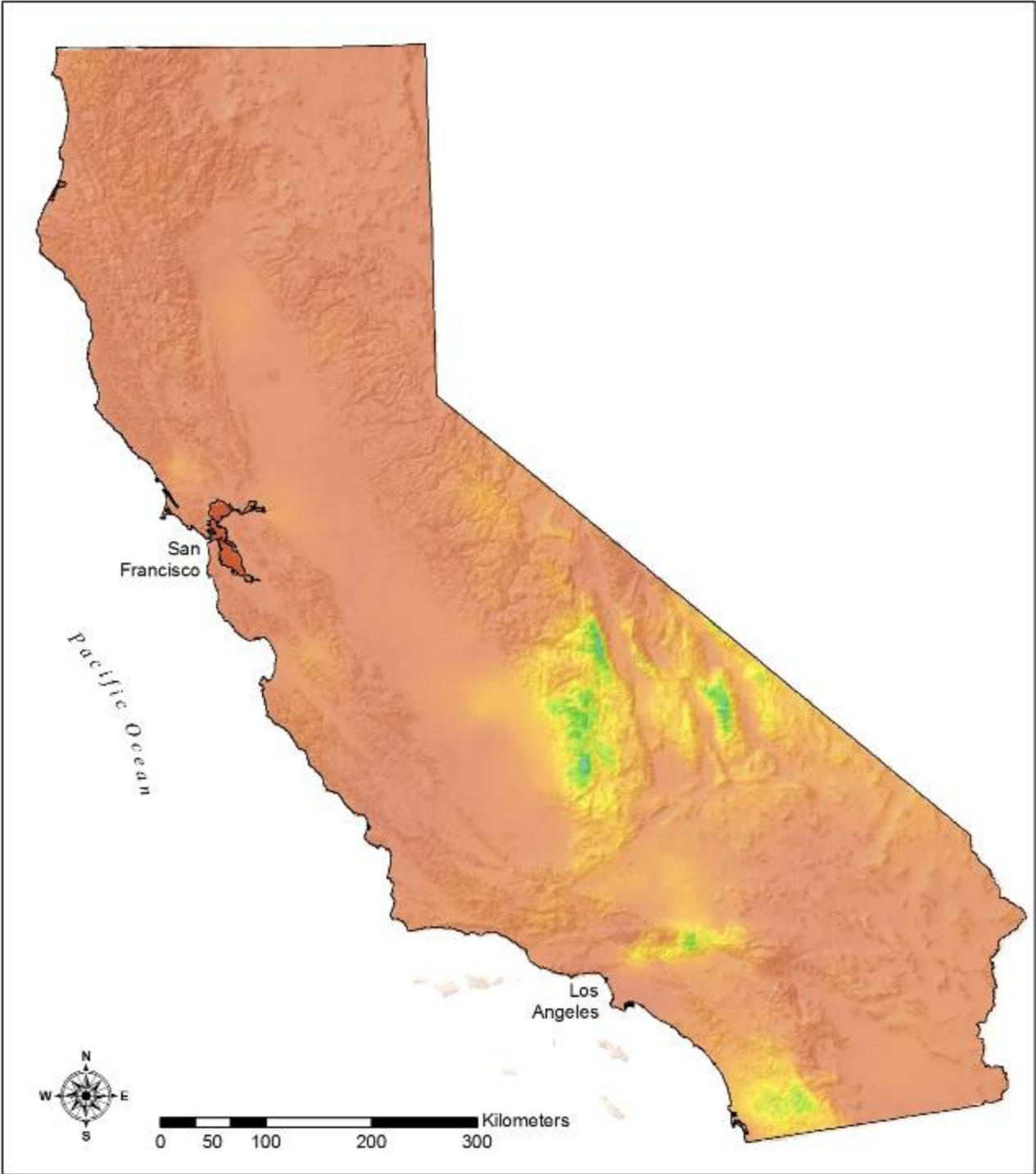
March 25, 1998



Hunter & Meentemeyer (in press)
J. Appl. Meteor.

PRECIPITATION

March 26, 1998

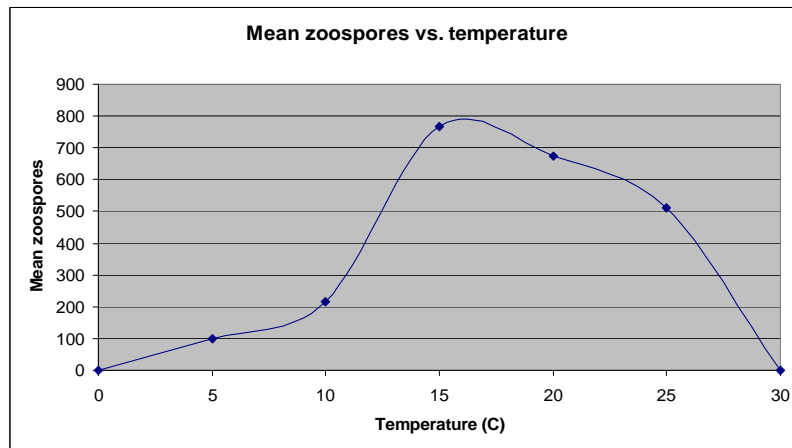


Hunter & Meentemeyer (in press)
J. Appl. Meteor.

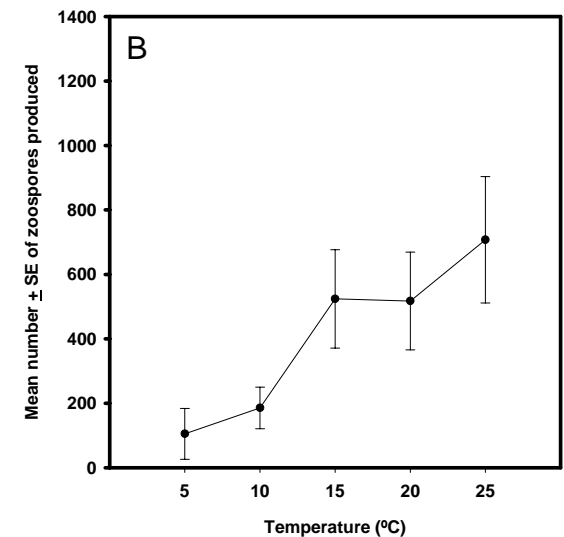
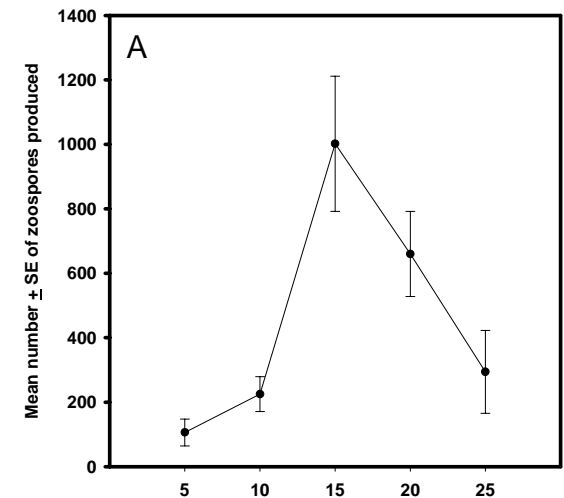
Weather Events

Parameterizing Temperature

- Davidson *et al.* (2005) lab studies
 - Measured zoospore production on bay
 - 7 set temps, 2 trials, 2 sets of 10 leaves each trial
- In the model
 - Average 2 trials
 - Fit curvilinear equation, scaled to 10 point scale
- Coded into model using weekly mean, max daily



$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$

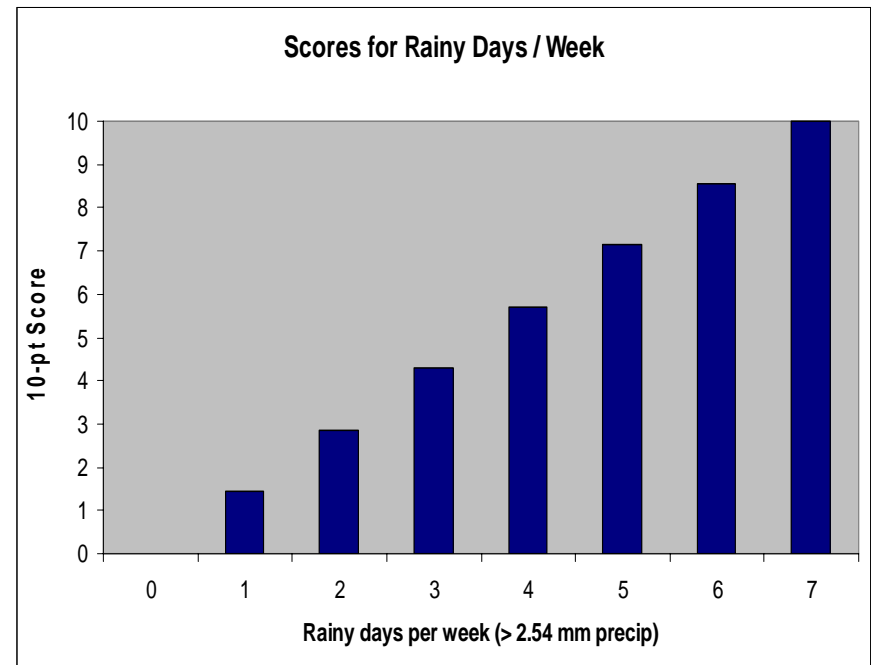


Davidson *et al.* 2005 *Phytopathology*

Weather Events

Parameterizing Precipitation

- Garbelotto / Davidson studies
 - 9-12 hrs of leaf moisture required for significant infection
- In the model
 - Rainy days are >2.5 mm precip
 - Assume linear relationship between potential sporulation and # of wet days per week
 - Scale # of wet days per week to 10 point scale

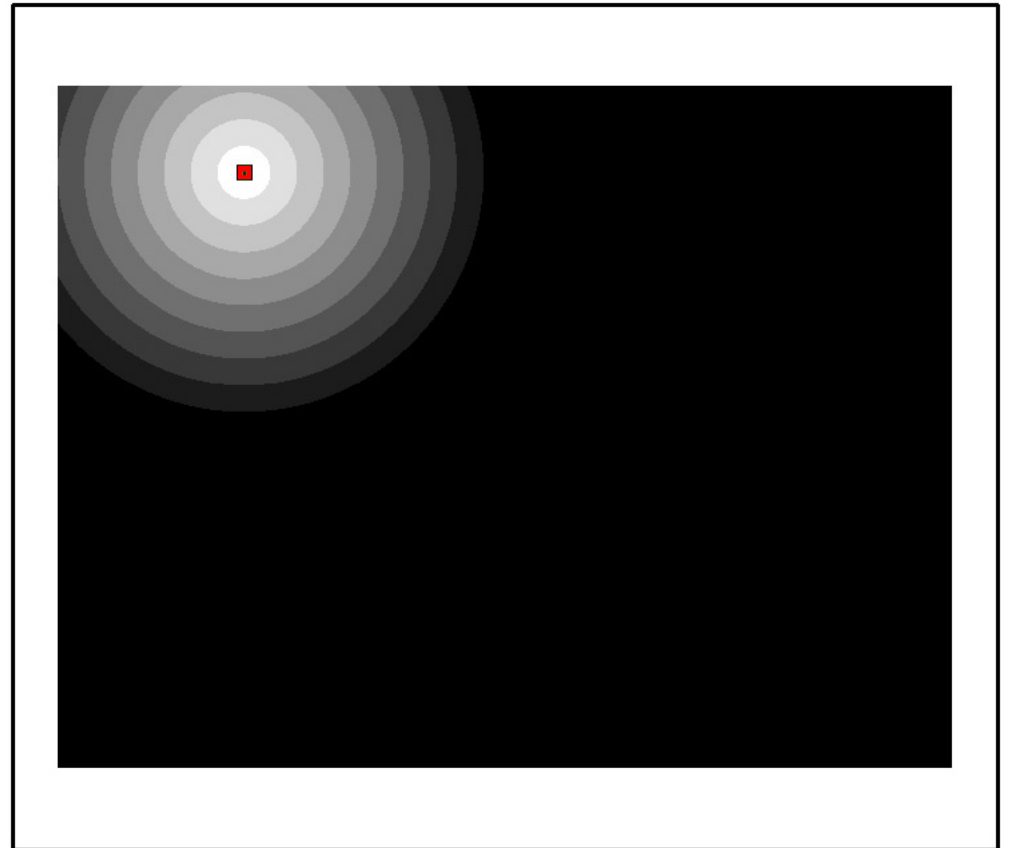


$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$

Dispersal distance

Mapping Methods

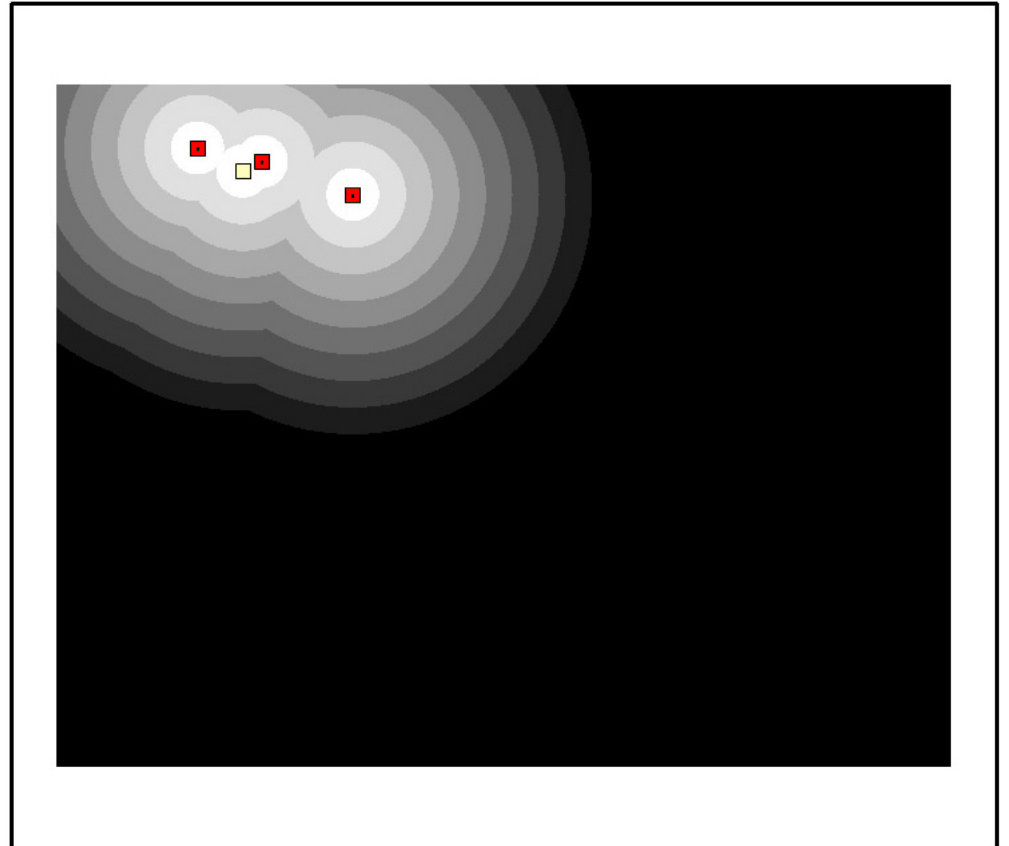
- Straight-line distance to nearest infected site
- Recalculated for each time step



Dispersal distance

Mapping Methods

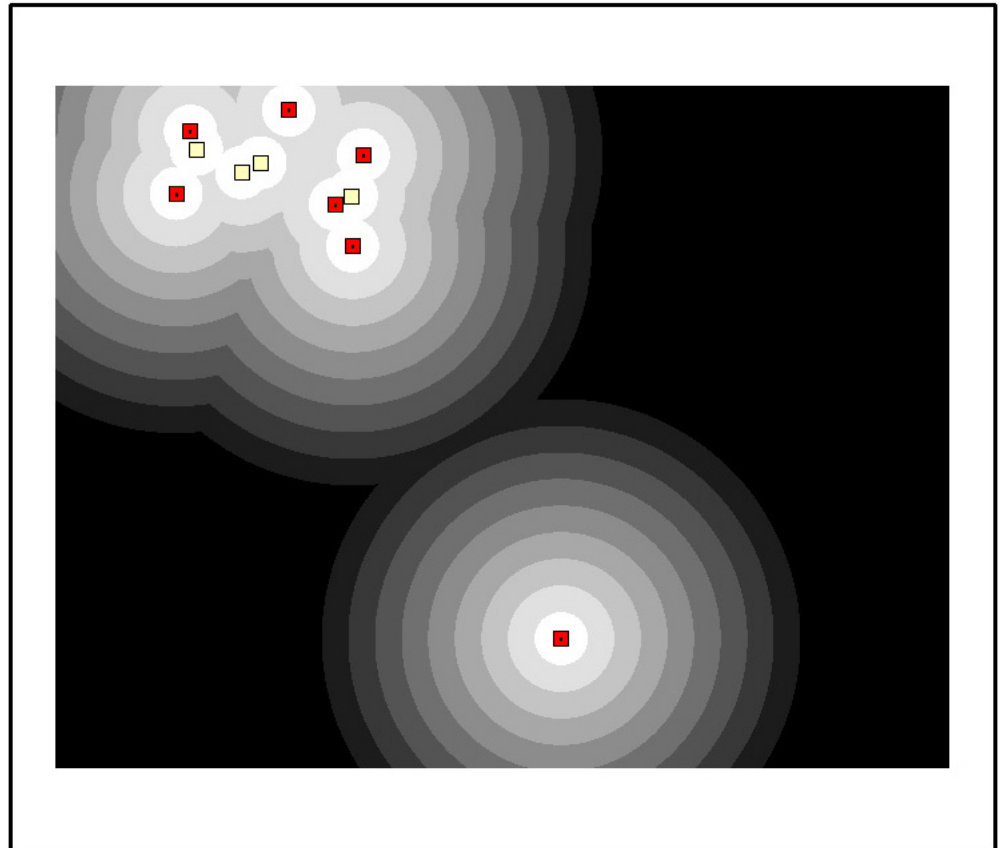
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Dispersal distance

Mapping Methods

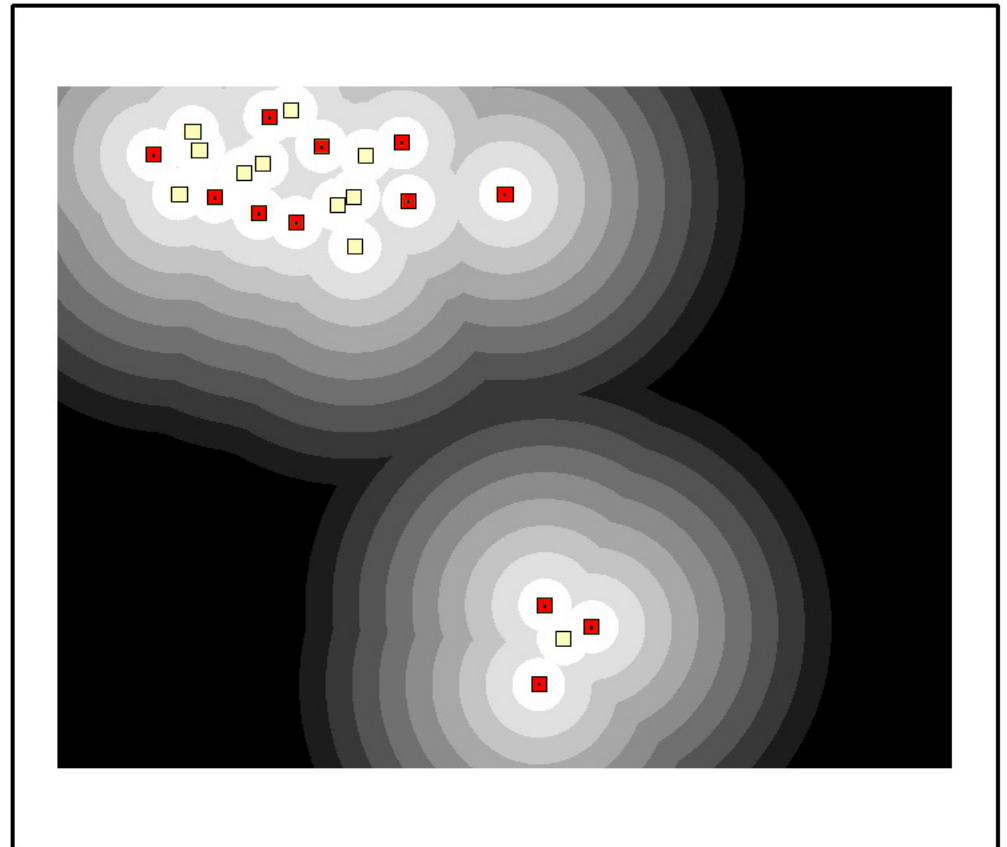
- Straight-line distance to nearest infected site
- Recalculated for each time step



Dispersal distance

Mapping Methods

- Straight-line distance to nearest infected site
- Recalculated for each time step

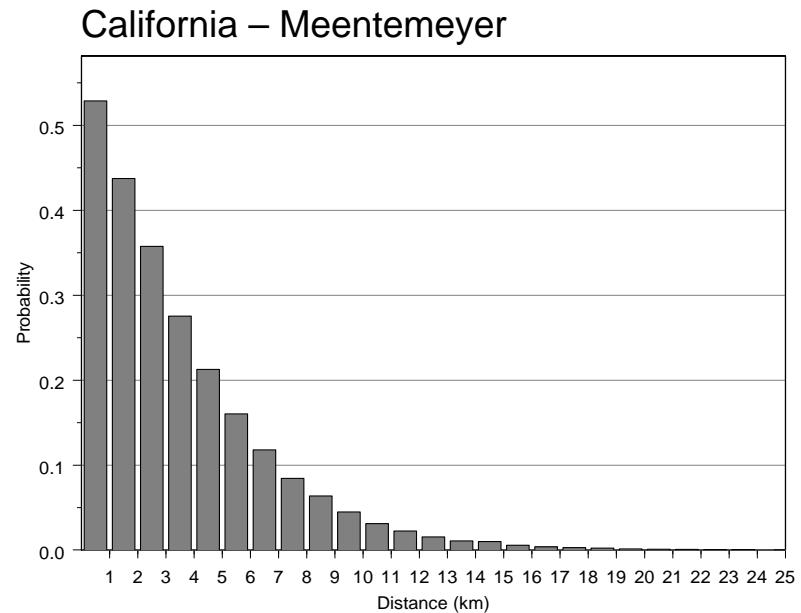
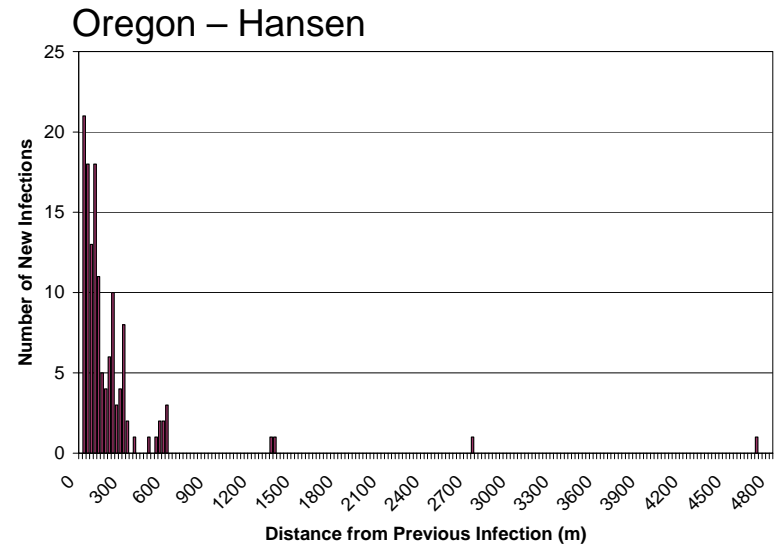


Dispersal distance

Parameterization

- 2 distance decay studies are being explored
- Model is programmed based on these curves and other potential curves

$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$



Surrounding disease levels (I)

- **Assumption:** higher pathogen abundance in surrounding cells increases chance of infection
- In the model
 - Output an index of pathogen abundance at each time step
i.e. the count that each cell has been infected
 - To account for surrounding disease levels, sum the counts within a 1250m rectangle and scale to 10 pts

250m

0	1	0	0	0
0	3	2	0	0
0	0	1	0	0
0	0	0	0	0
0	0	0	1	0

Output from previous example

=

-----1250m-----

		8		

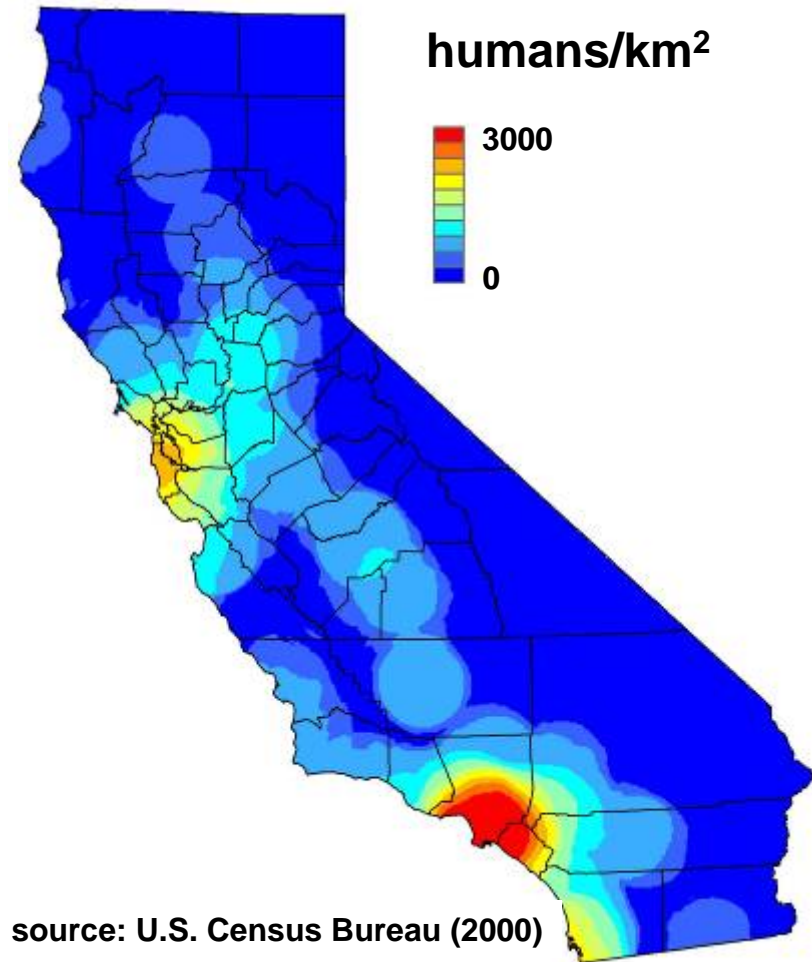
Output of surrounding pathogen abundance for an individual cell

$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$

Potential long-distance vectors

Mapping Methods

- *P. ramorum* disperses locally through several physical pathways
 - i.e. rain splash, wind-driven rain, stream water
- Human activity poses highest risk for long-distance dispersal
- Human population density within 50km radius areas

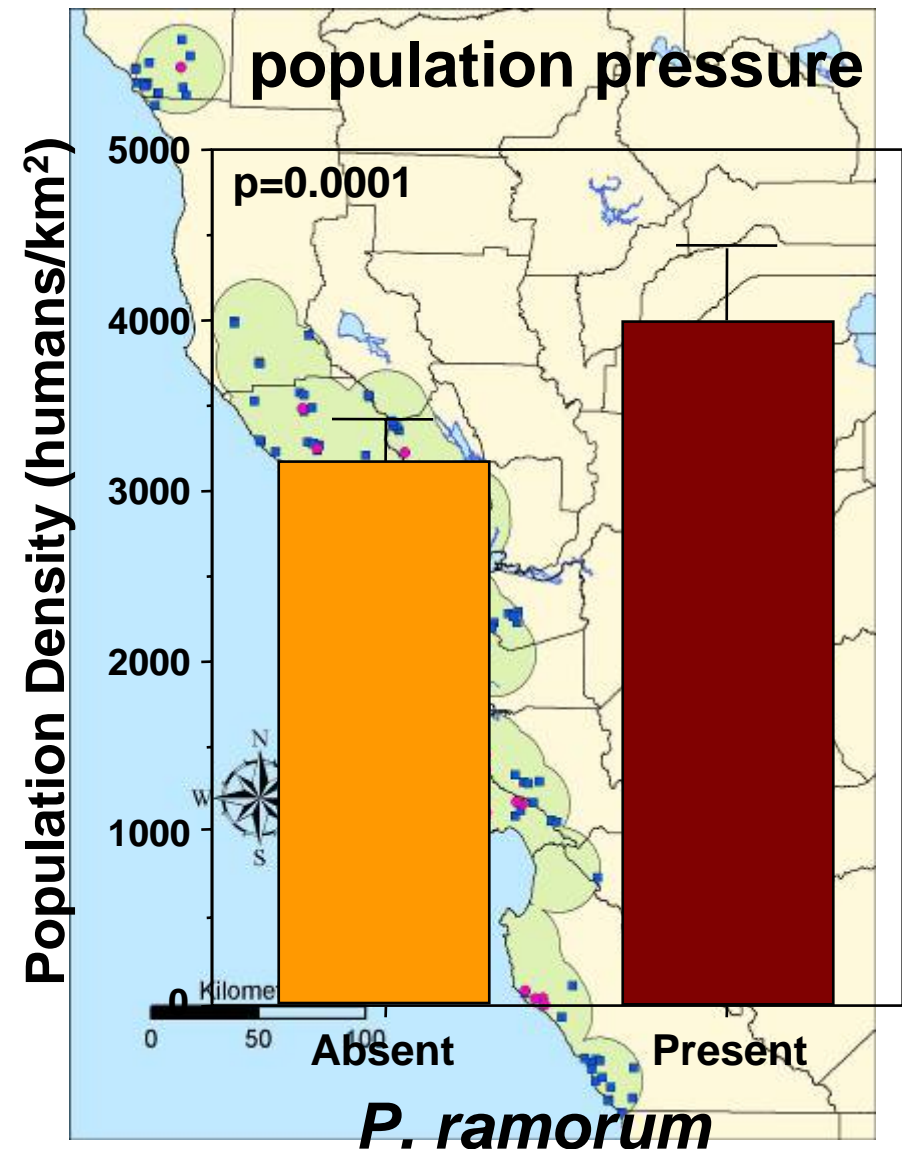


Potential long-distance vectors (H)

Parameterization

- Logistic regression
- Response variable
 - Presence/absence
 - Statewide foliar survey data (Meentemeyer) n=166
- Determine relationship between pop density and infection probability while controlling for host vegetation and climate
- Scale probabilities to 10 point scale for integration with other parameters

$$P_{t+1} \approx \frac{S * P * T * I * H}{D}$$



Applying the model

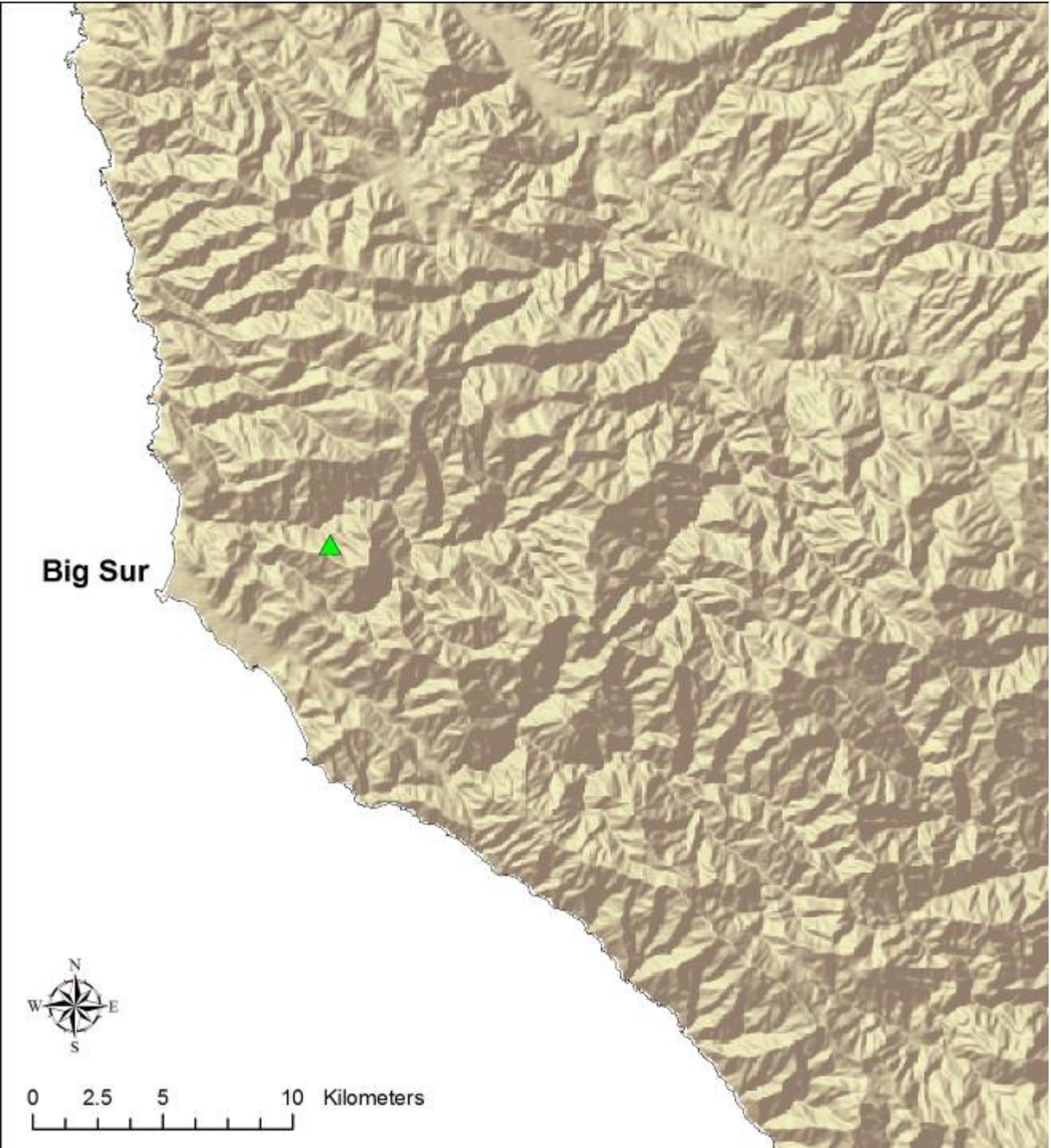


Example model run

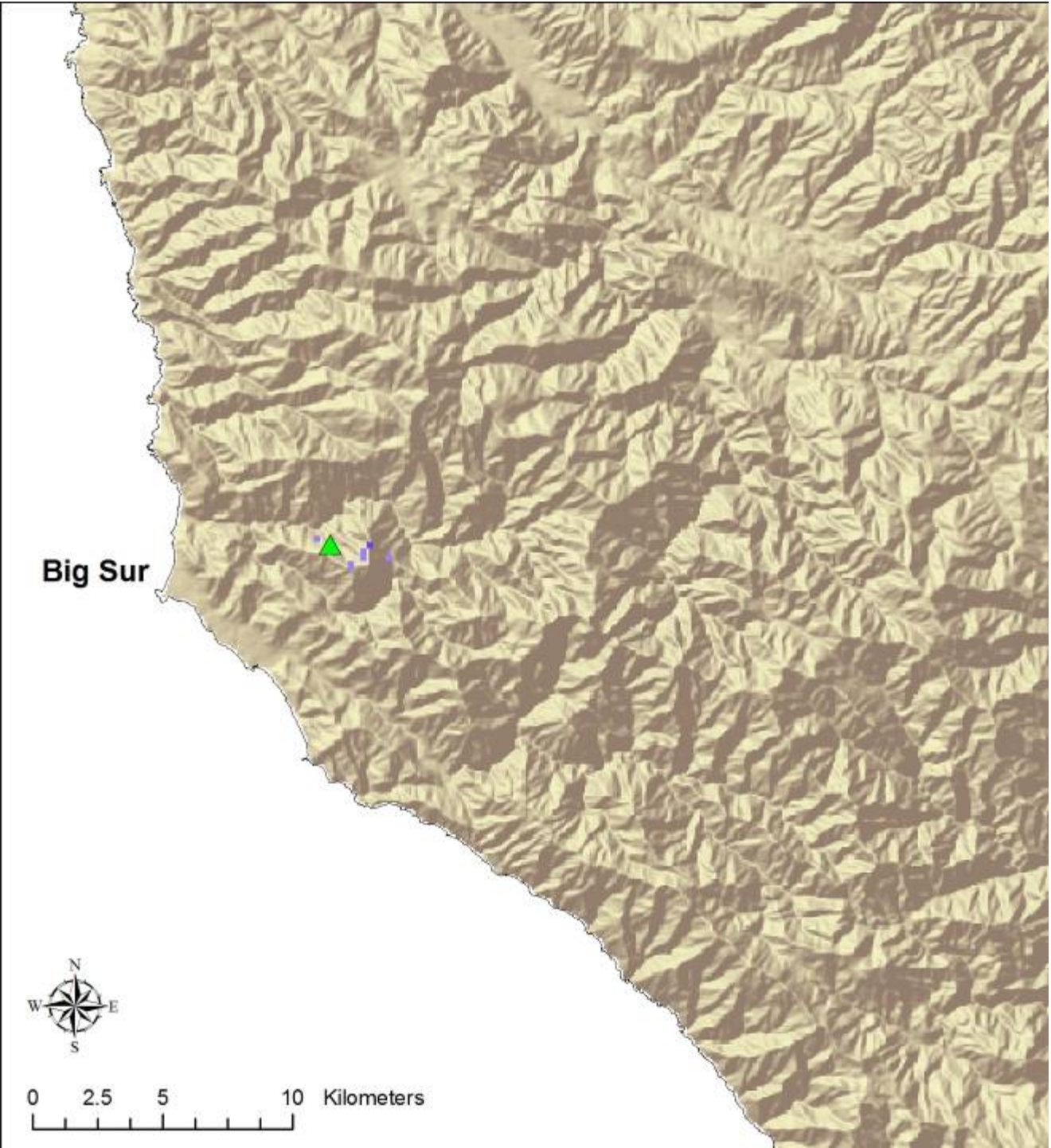
- Initiated spread with a single infected cell
 - Large patch of highly susceptible vegetation near Big Sur
- Initiated in 1985
 - Weekly time step through Dec 2003



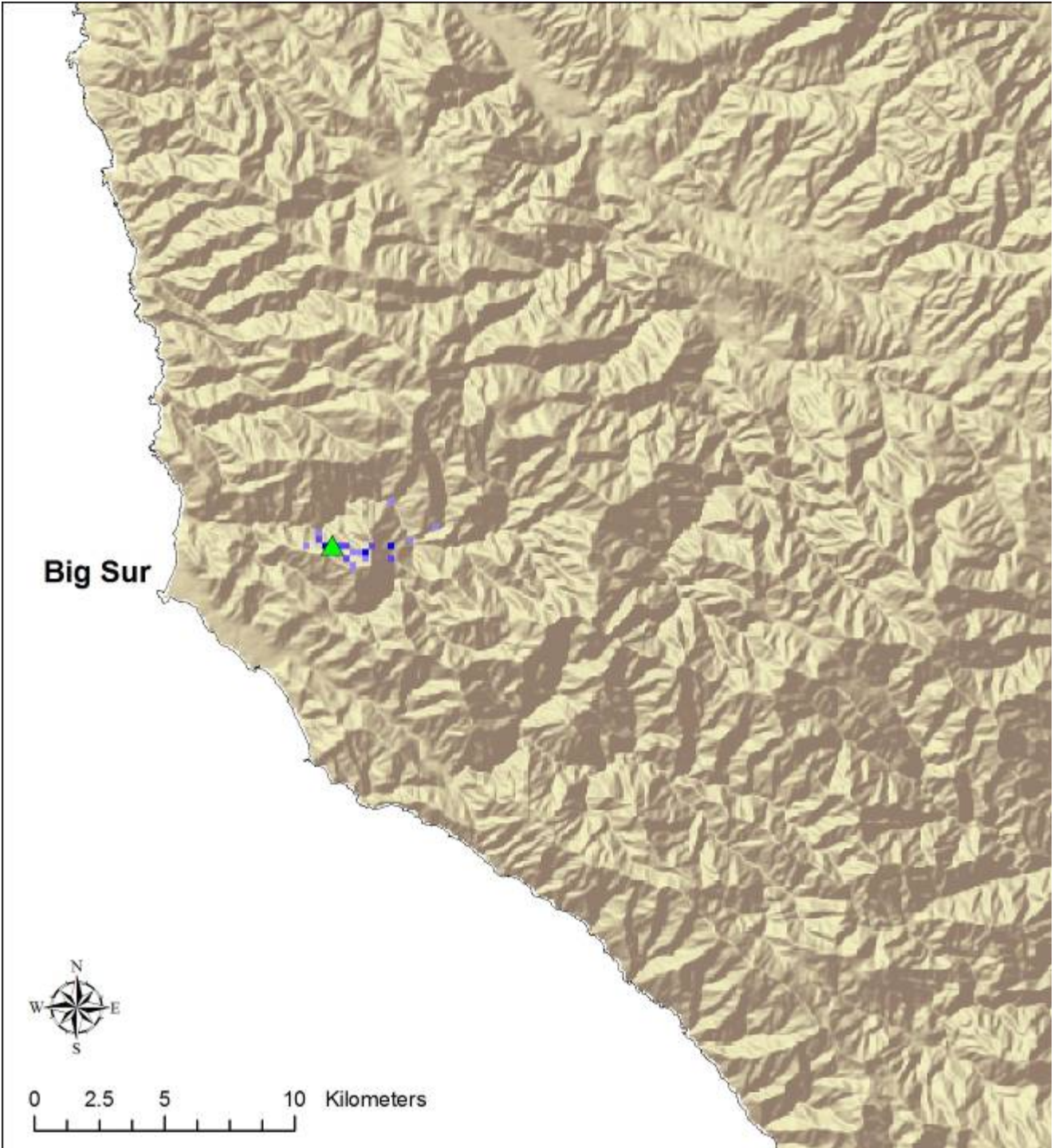
1985



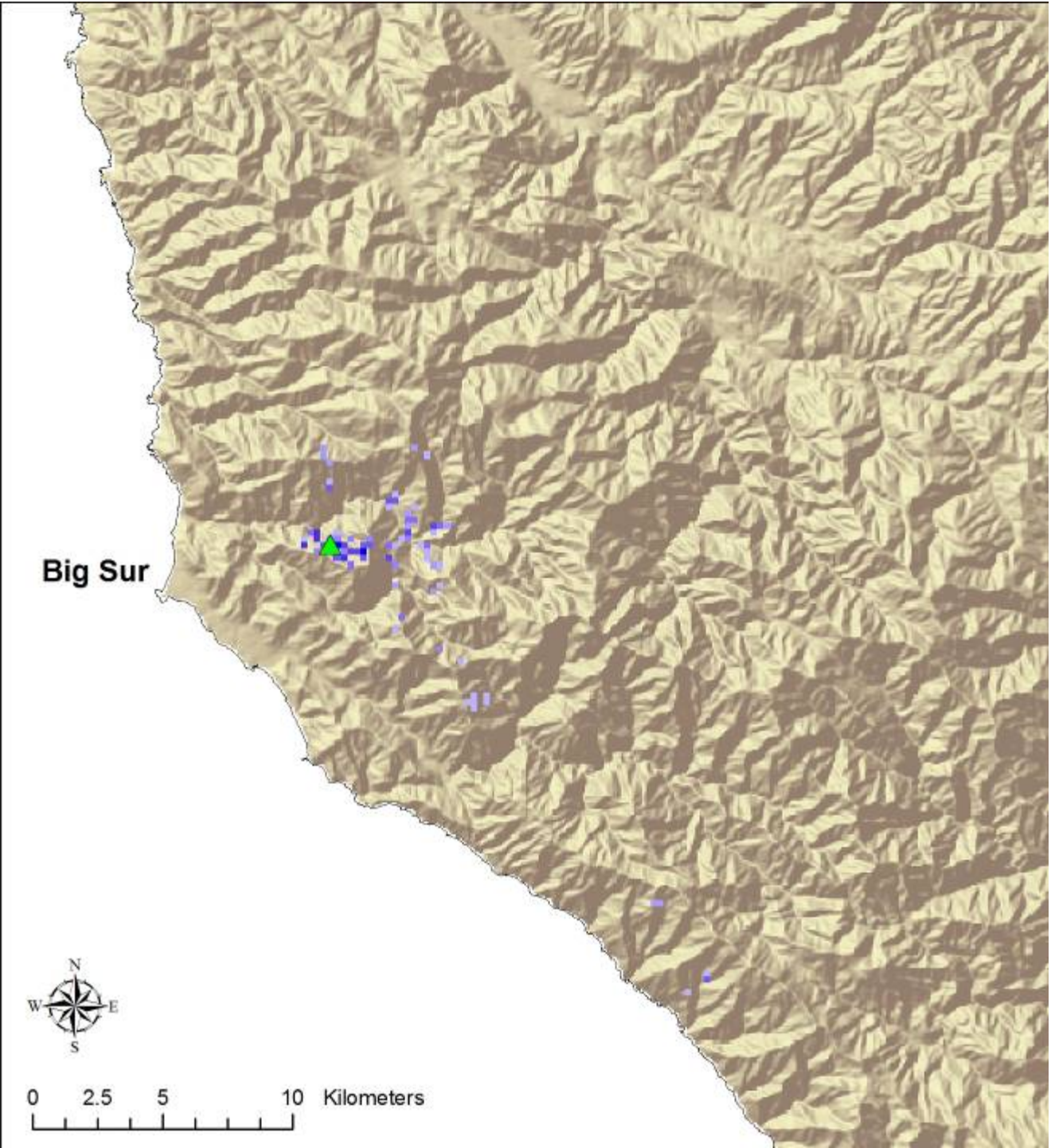
1988



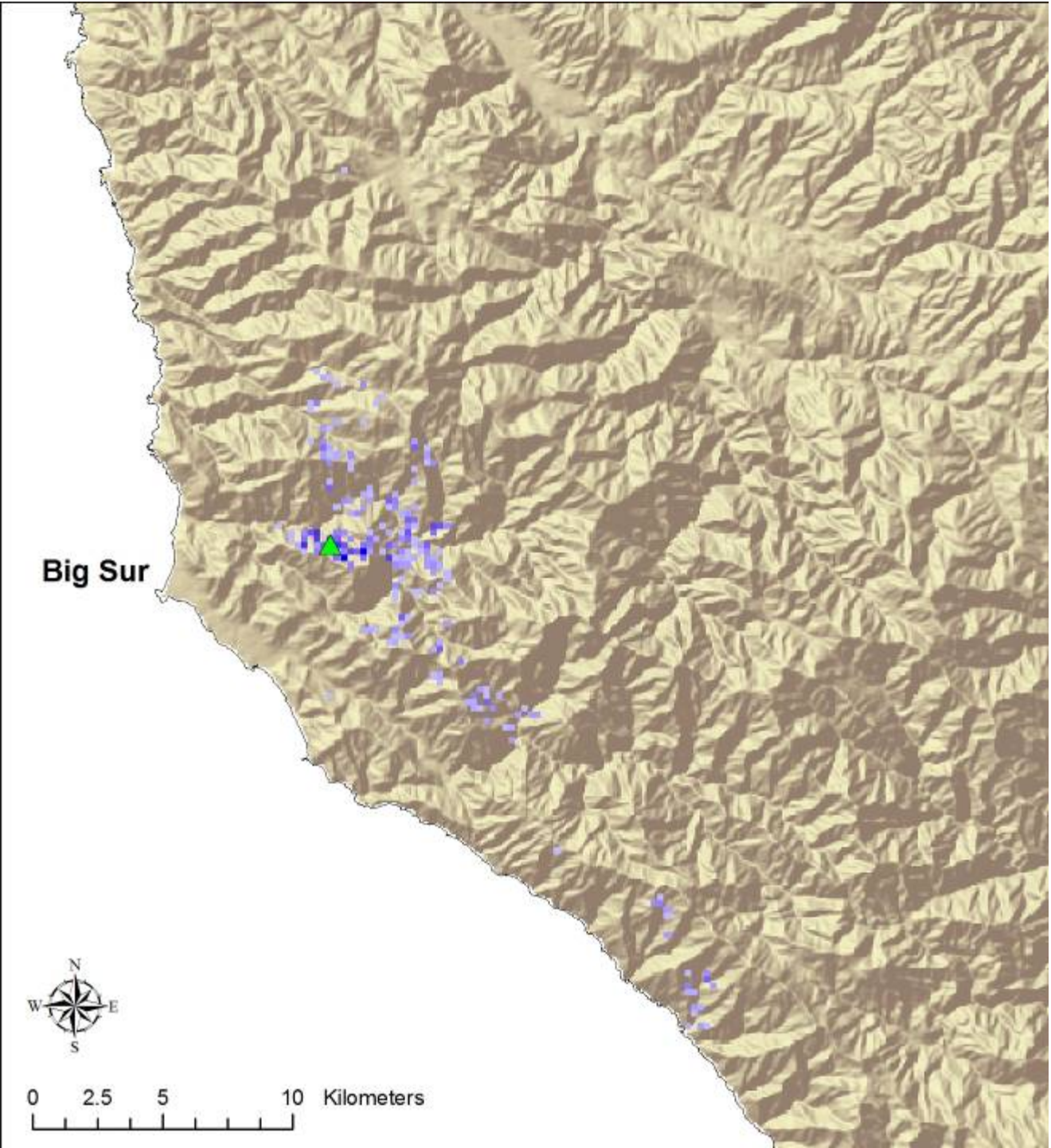
1991



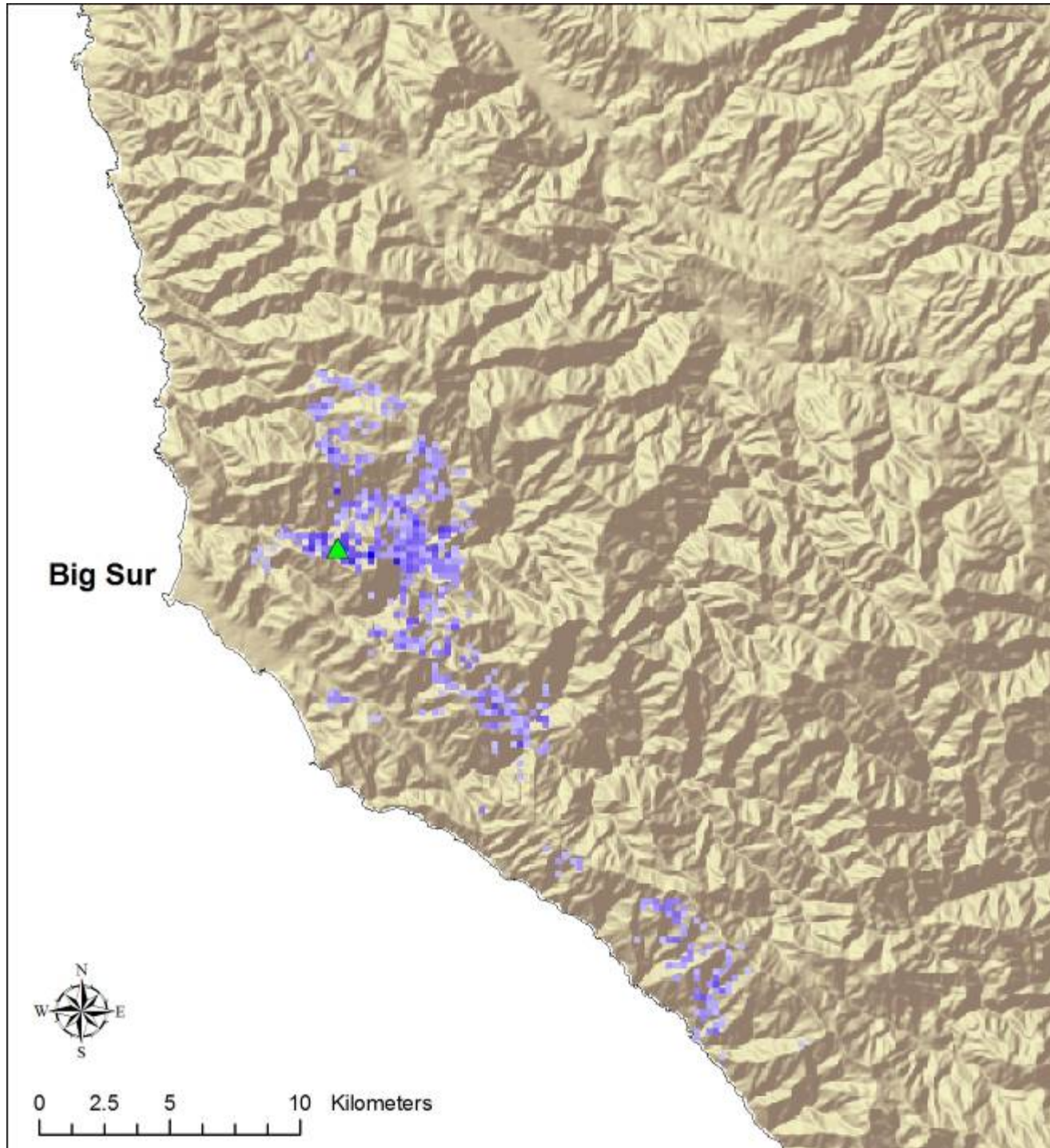
1994



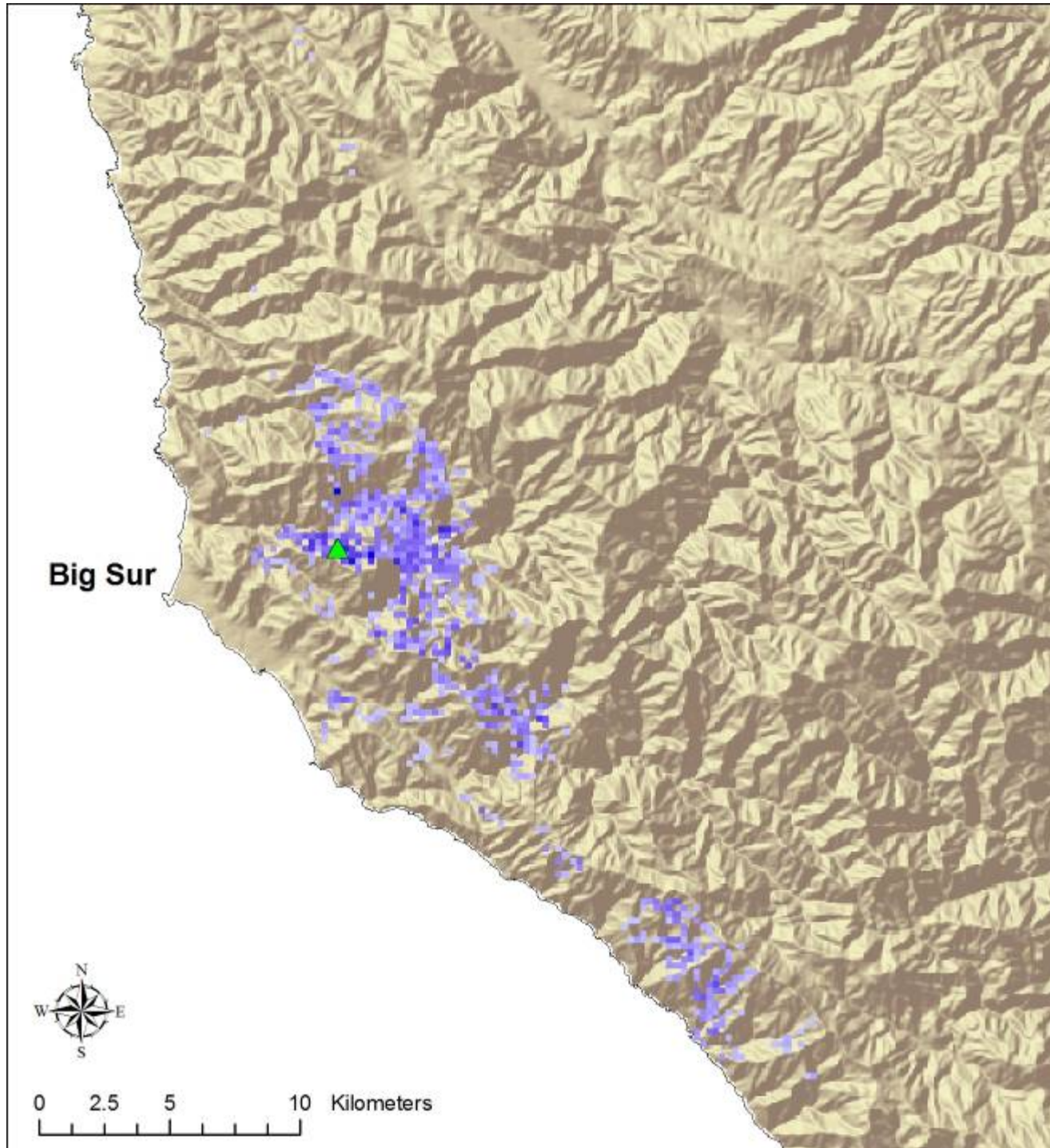
1997



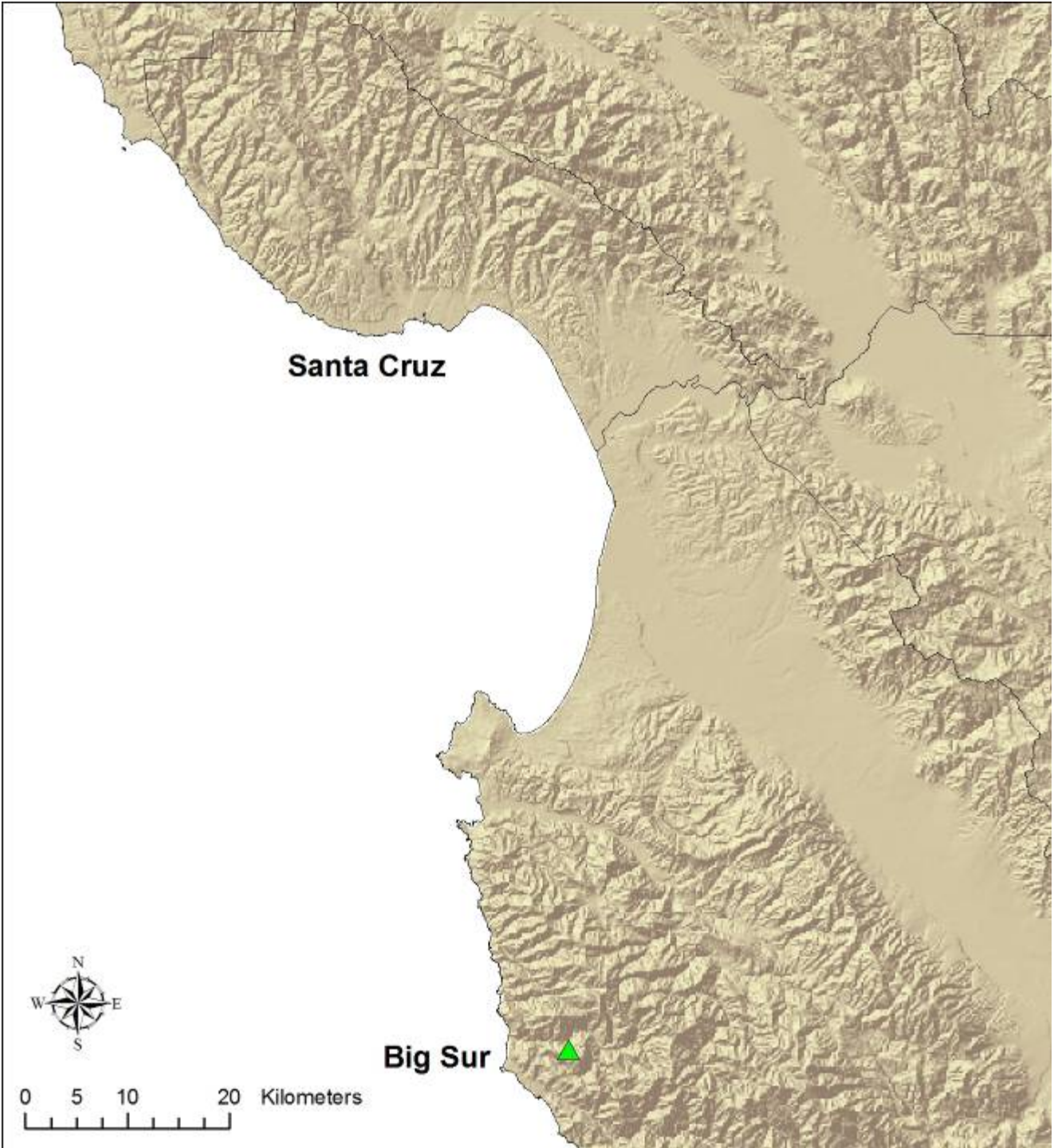
2000



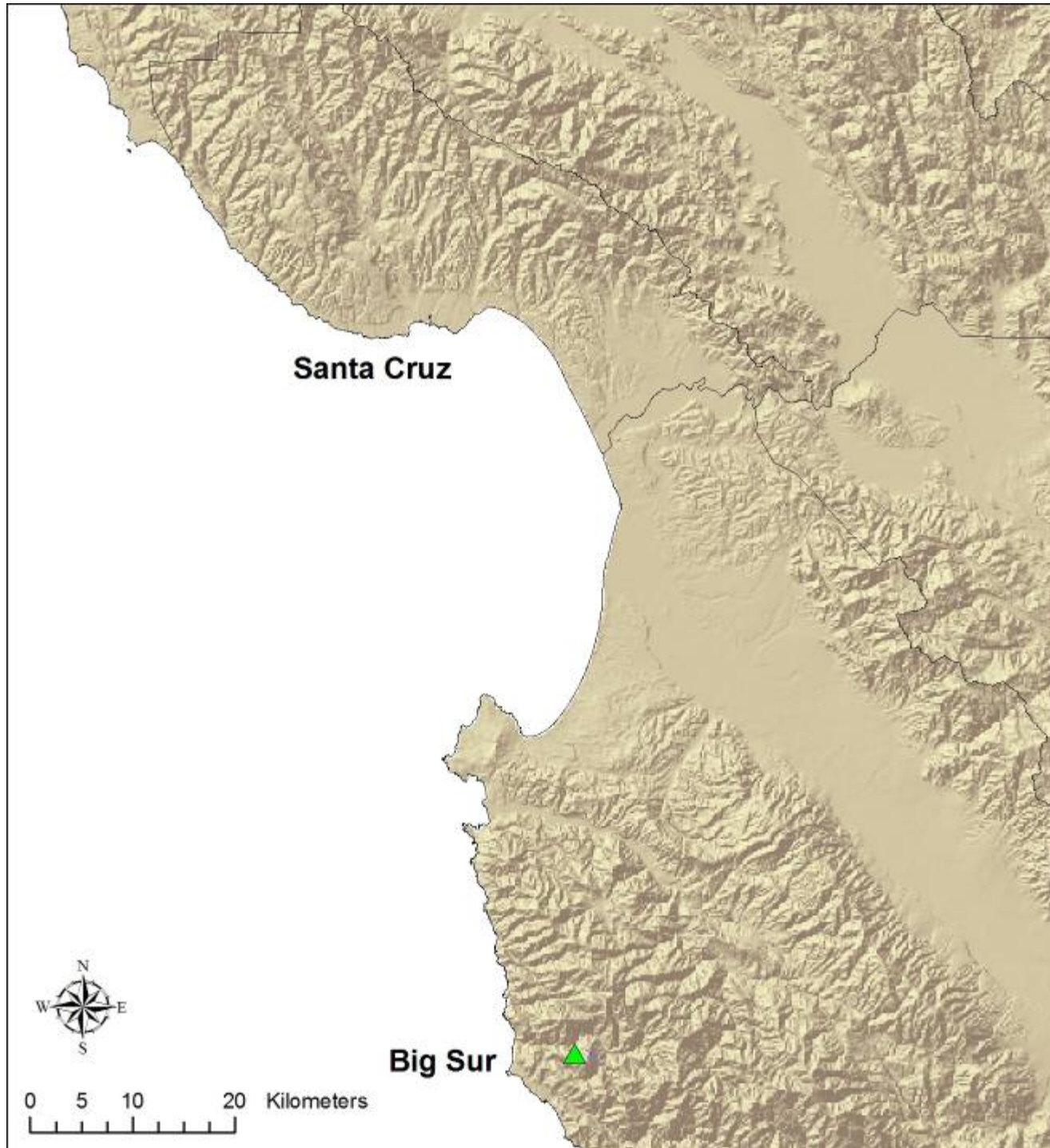
2003



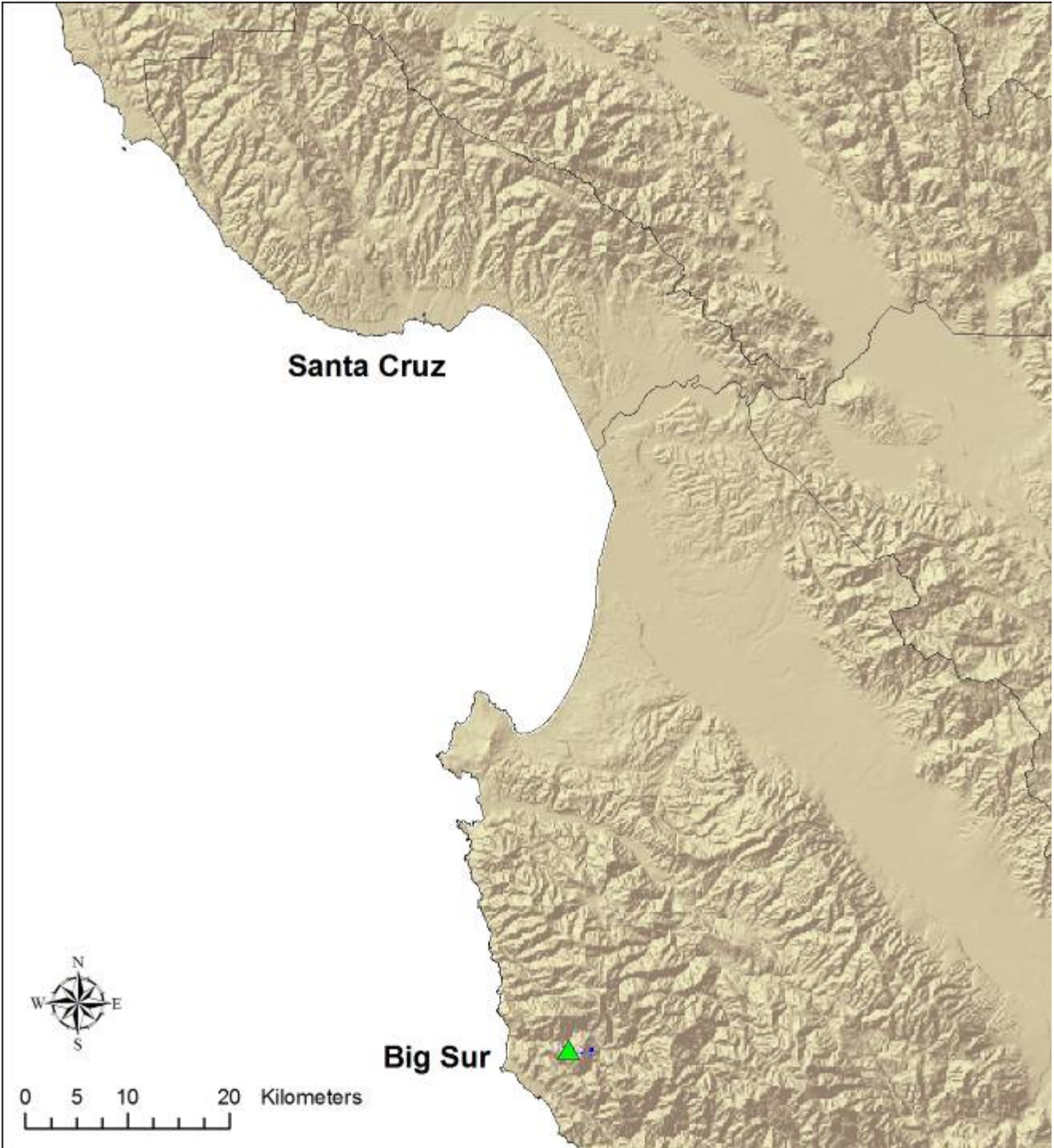
1985



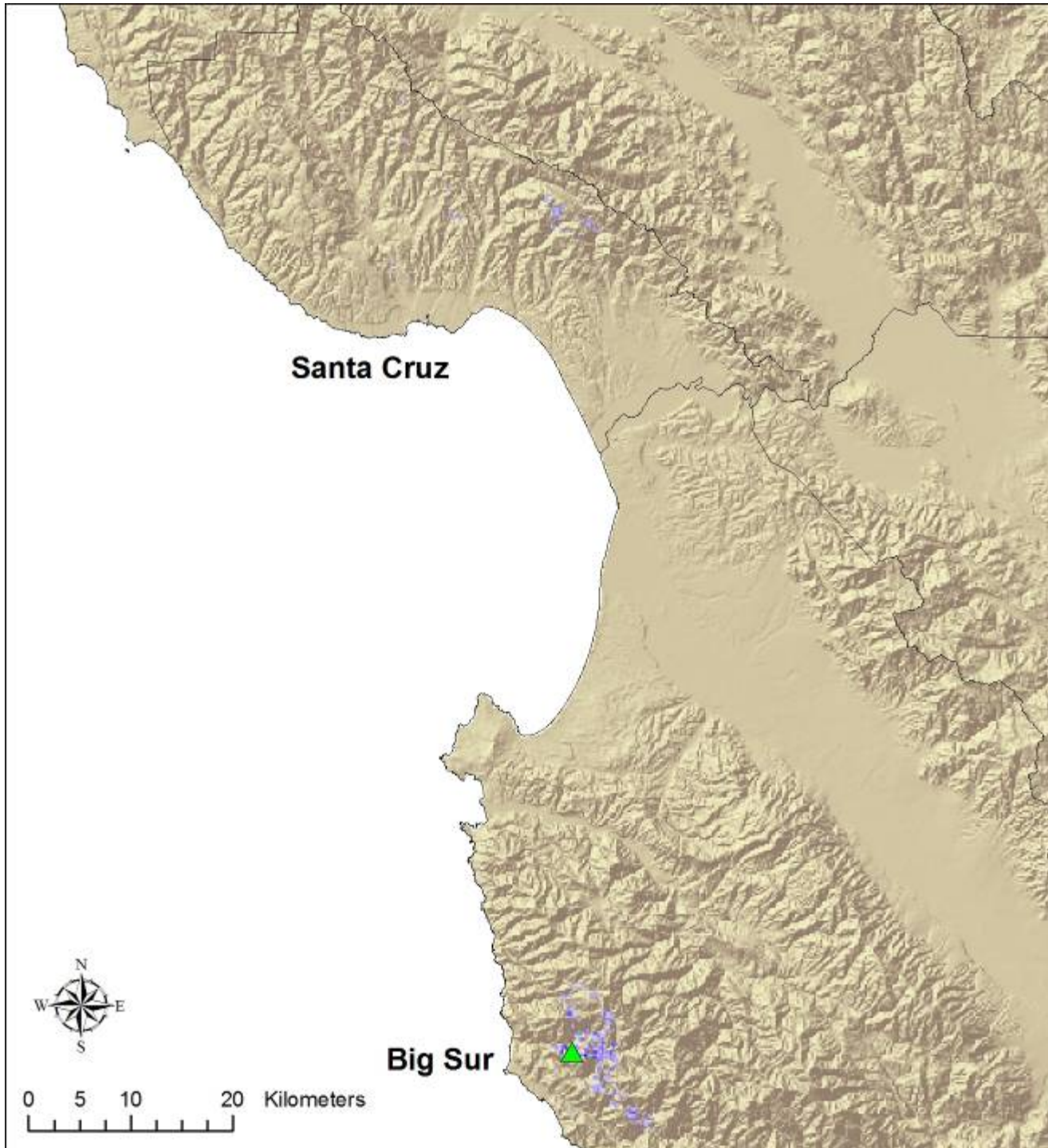
1988



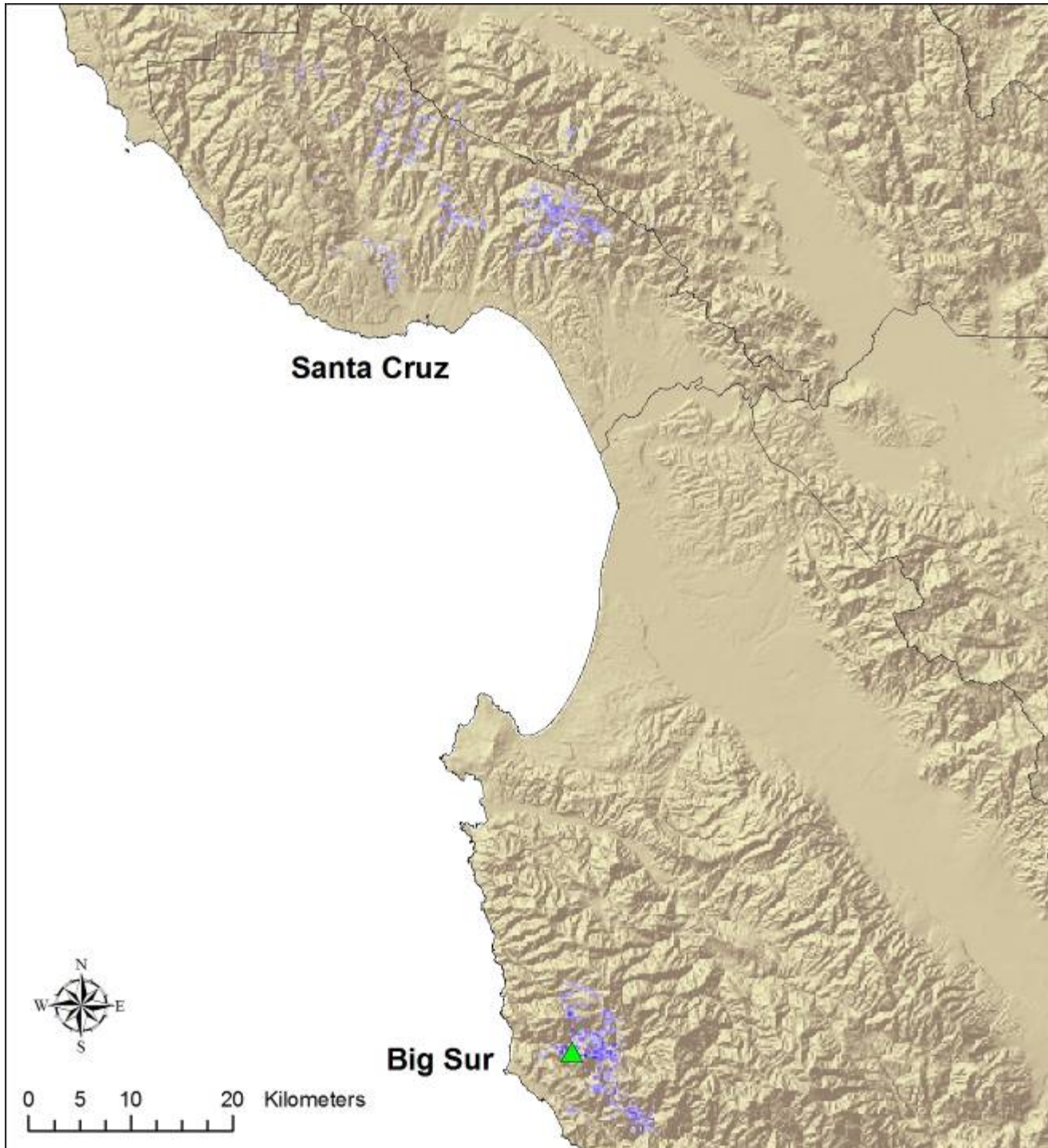
1991



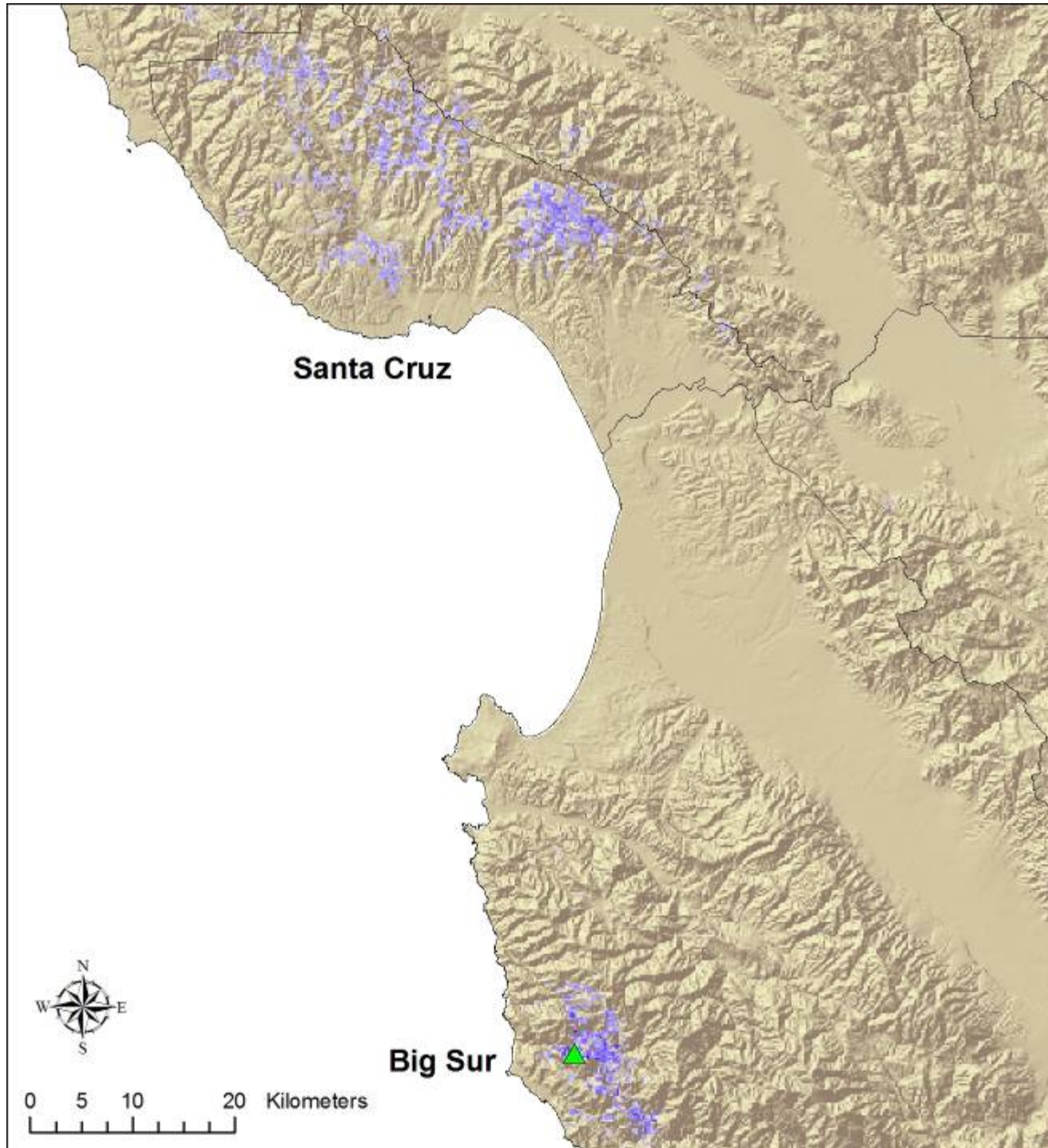
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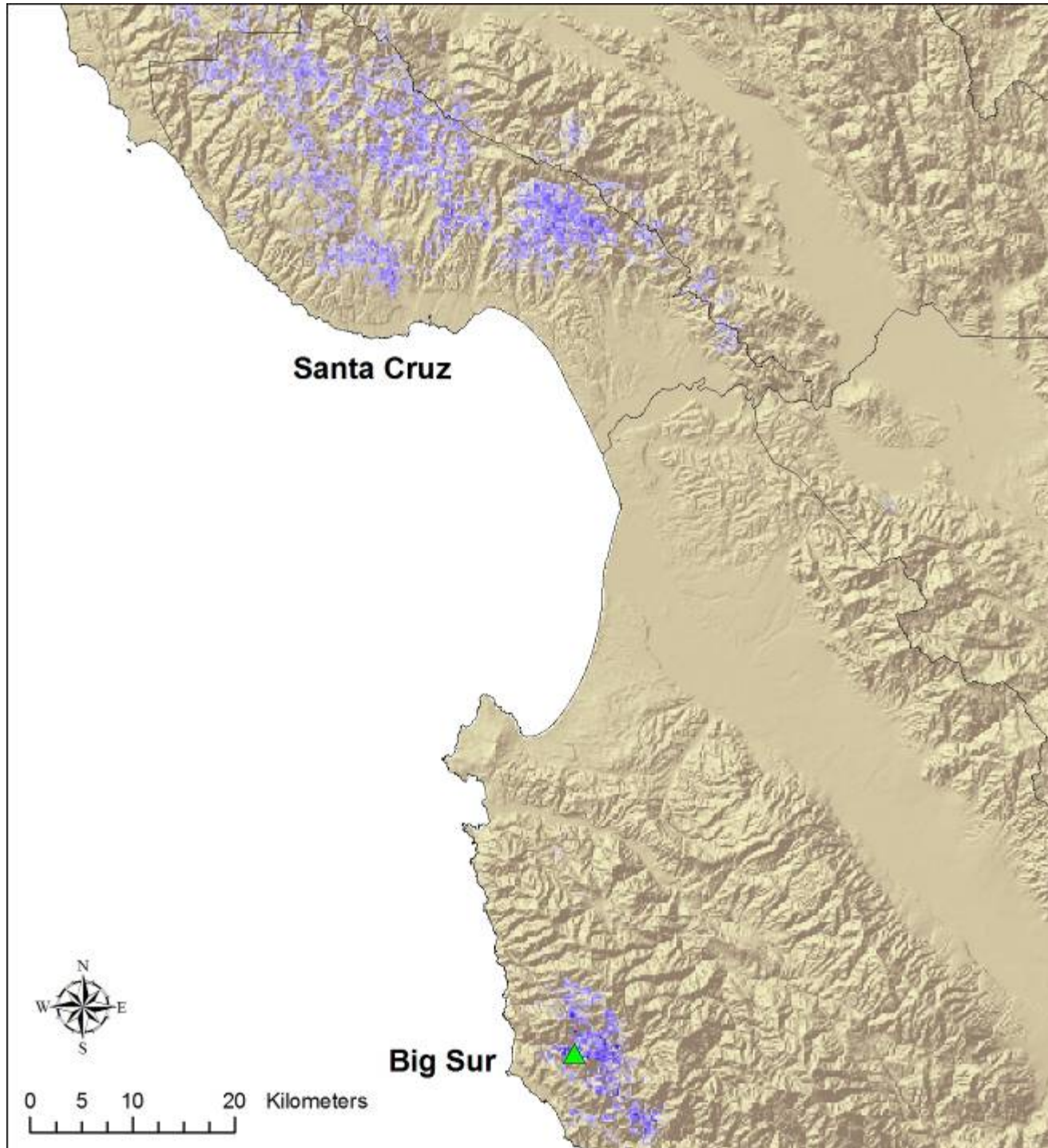
1997



2000



2003



Model Evaluation

- Compare output maps with observed disease distribution
 - Run model 10 times
 - Assess predicted vs. observed infected area within 100 5k blocks
- Evaluate magnitude and direction of error and determine predictive accuracy
- Conduct sensitivity analysis of parameters



Additional research questions

- Test hypotheses about dispersal and infection processes
 - Influence of weather events
 - Simulate management scenarios



Significance

- Critical for predicting and slowing the spread of *P. ramorum* and other invasive species
 - Incorporate spatial and temporal heterogeneity of environmental conditions into spread modeling
 - Focus management efforts on high-risk landscapes
 - Test alternate management scenarios

