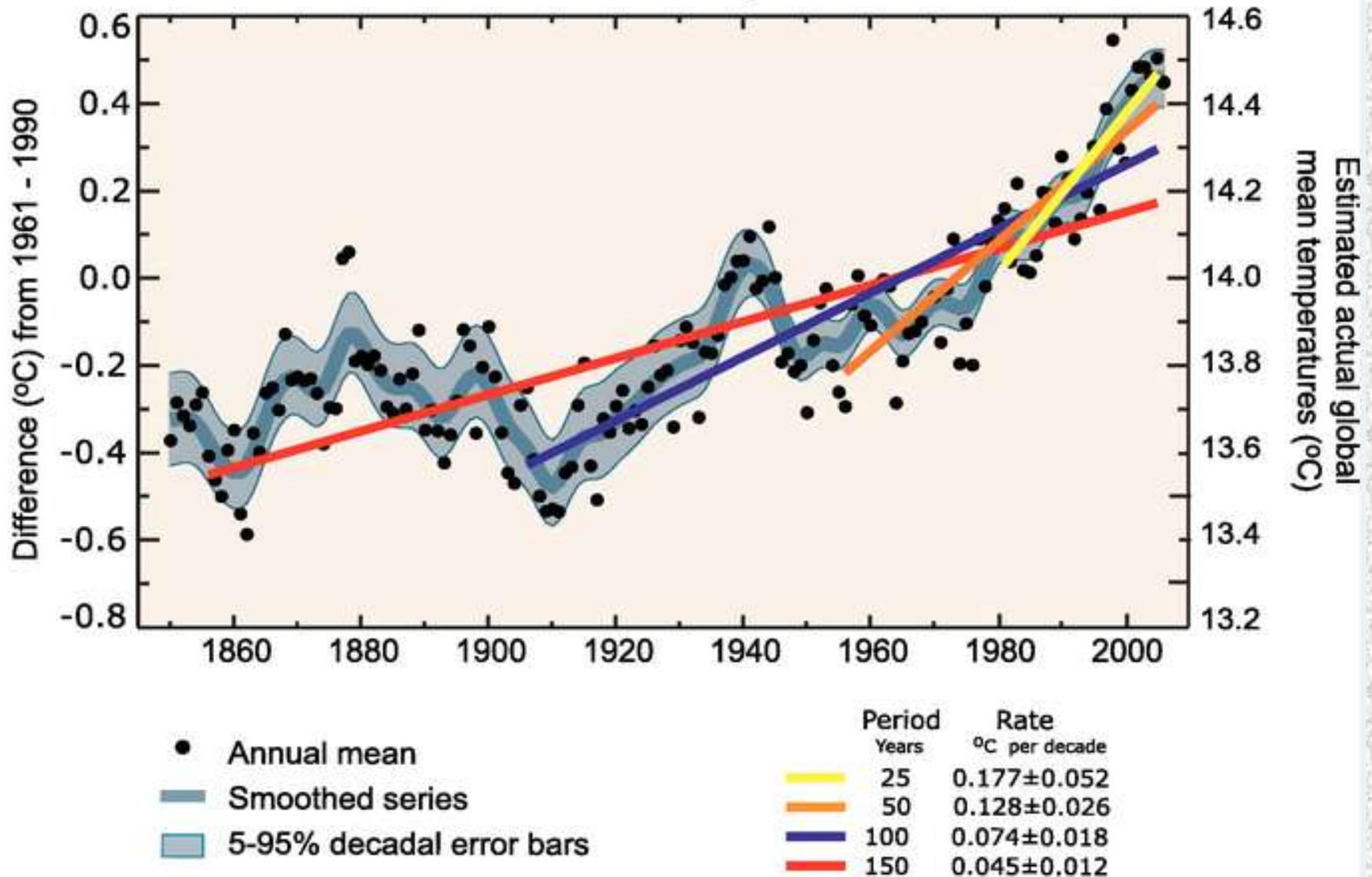


# Global Mean Temperature

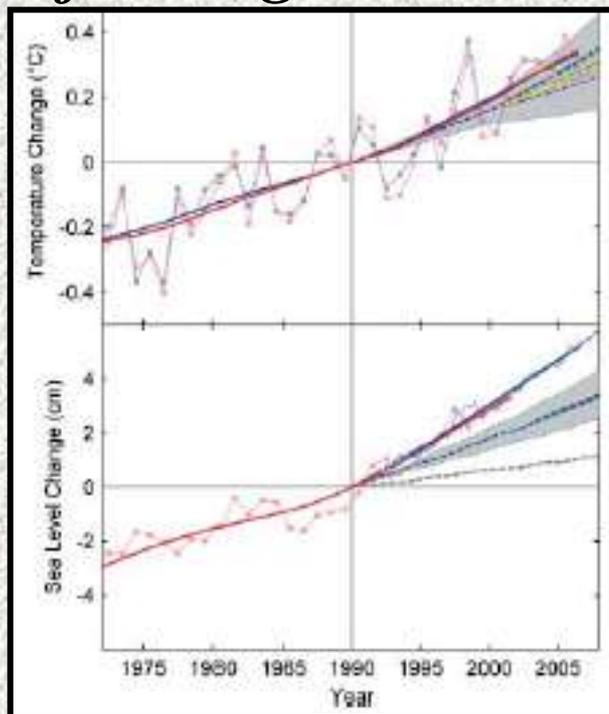


**Does 'velocity' of climate change matter for wildlife?**

# Increasing rates of change in abiotic parameters

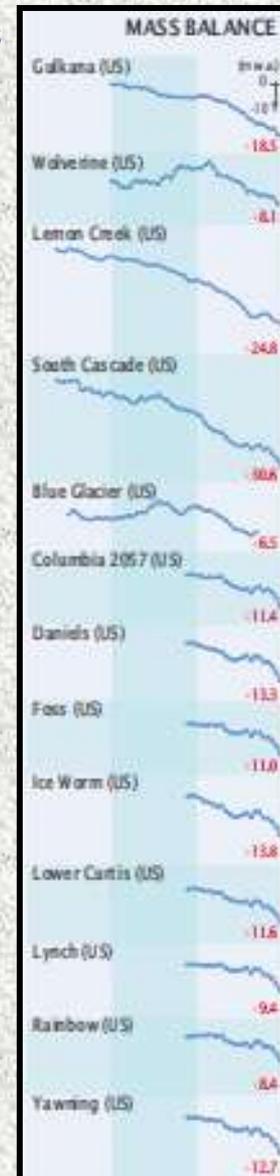
**Temperature change  
Sea-level rise**

Rahmstorf et al. 2006

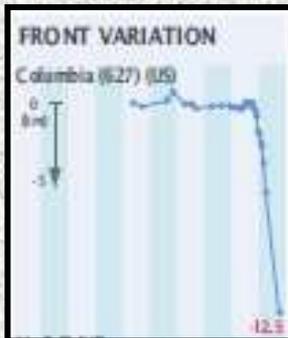


**Glacial mass balance**

UNEP-WGMS 2008



**Extent of glaciers**



Mote et al. 2003

**Changes in 1 Apr SWE**

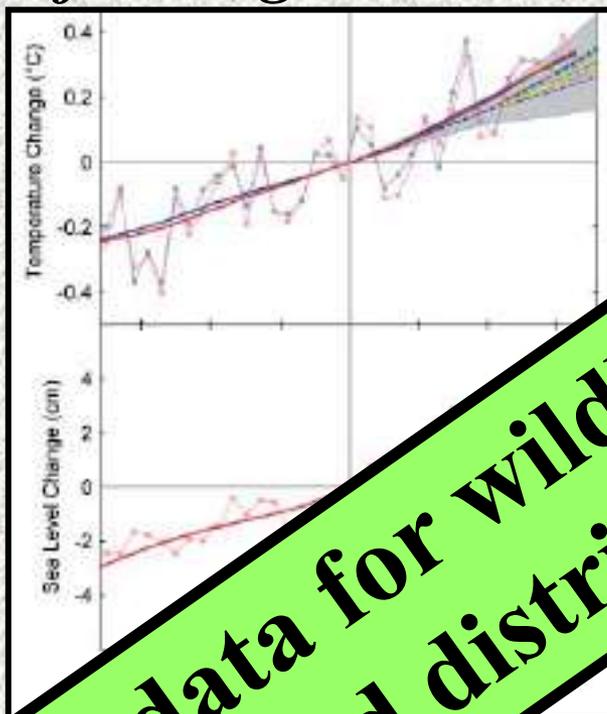
1930s to 1990s			1945-55 to 1990s		
Observed SWE	VIC SWE	Precip.	Observed SWE	VIC SWE	Precip.
-13.6%	+1.3%	+3.5%	-29.2%	-15.5%	-4.6%
+10.8%	+2.2%	+8.6%	-15.8%	-8.8%	+0.5%
+2.9%	-13.8%	+10.4%	-2.2%	-24.6%	-1.1%
+8.9%	-6.4%	+10.1%	-21.6%	-18.2%	+2.1%

UNEP-WGMS 2008

# Increasing rates of change in abiotic parameters

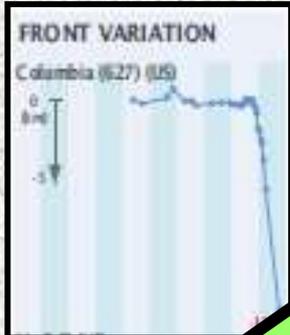
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Glaciers

Extent of glaciers

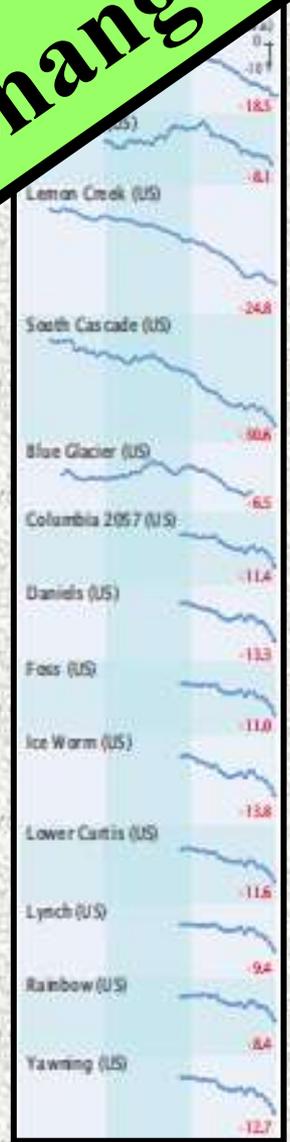


UNEP-WGMS 2003

**Much fewer data for wildlife, esp. for local extinction and distributional change**

Changes in 1 Apr SWE

	1945-55 to 1990s				
	Observed SWE	Precip.	Observed SWE	VIC SWE	Precip.
	+1.3%	+3.5%	-29.2%	-15.5%	-4.6%
	+10.8%	+8.6%	-15.8%	-8.8%	+0.5%
	+2.9%	+10.4%	-2.2%	-24.6%	-1.1%
	+8.9%	+10.1%	-21.6%	-18.2%	+2.1%



**Conserving wildlife in  
mountain ecosystems:  
*importance of a broad-scale perspective***

**Erik A. Beever**

**USGS Northern Rocky Mtn. Science Center**

# Overview: *The Big Picture*

- **Montane ecosystems and their wildlife**
- Critical importance of mechanisms
- Sage-obligate phenology: birds, bugs, flowers
- Am. pikas (change vs. current, changing pace)
- New technologies, approaches
- Context-dependency: *it all depends ...*

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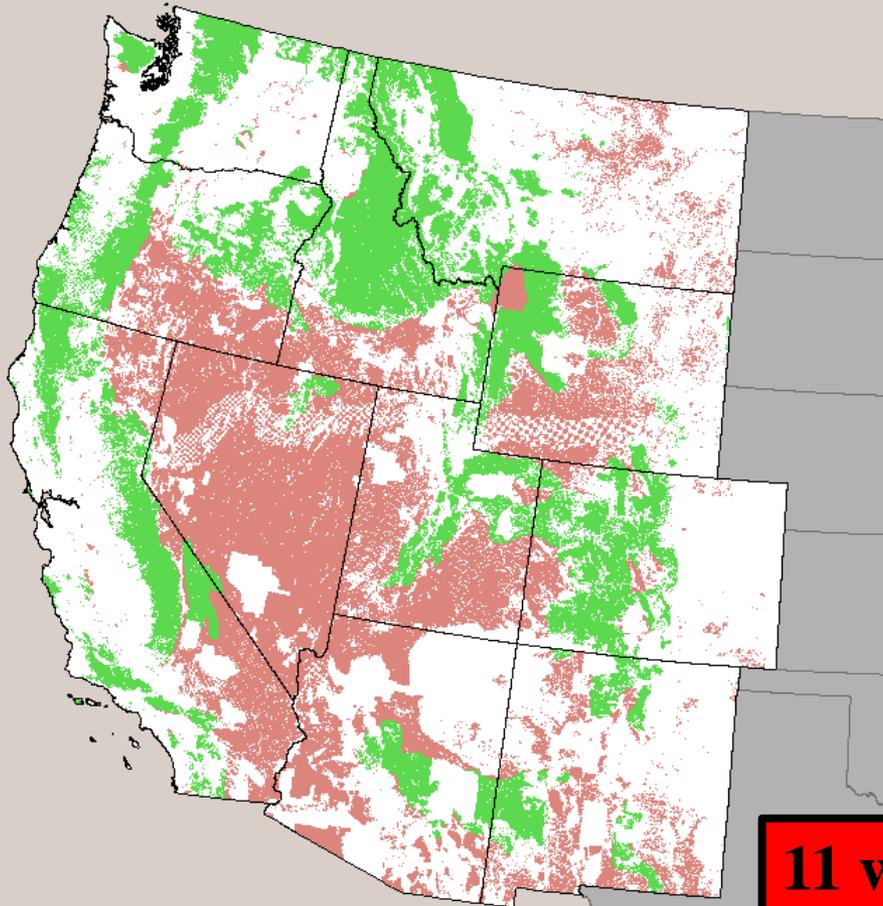
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# Montane ecosystems ...

- **Higher PPT, % as snow, UV radiation, wind**
  - **Greater variability in climate**
- **Lower ambient T, atmosph. P, O<sub>2</sub>**
- **Pronounced veg. zonation, fires**
- **Lower intensity & diversity of many human land uses**
- **Poorly developed soils, lower NPP, shorter growing season**
- **Higher proportion of wilderness, conservation areas**



# Mountain ecosystems are disproportionately federal



(PADUS) v1.1 data

Lt. green      Purple

- 56% of mountainous landscapes are federally managed
- 39% of non-mountainous landscapes are federally managed

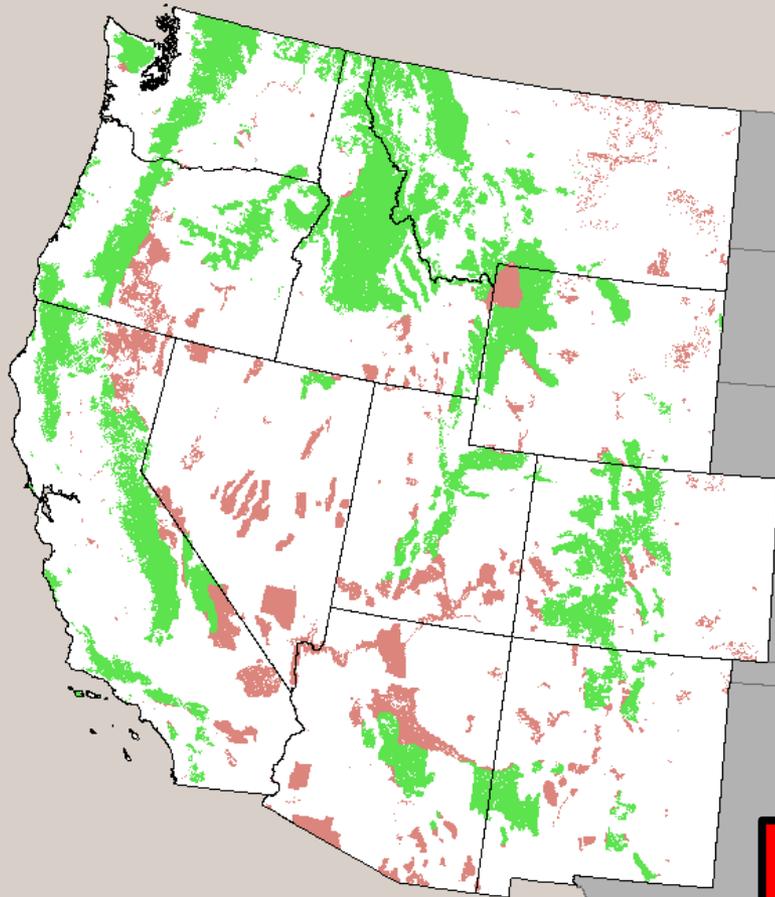
**11 western U.S. states**

**Fed** = USFS + DOI lands

**Non-fed** = all other lands

**Mountainous** = Categories C5, C6, & D3-6 from Hammond's (1970) classes of land-surface form

# Most-protected lands are mostly mountainous



Lt. green

Purple

- 72% of strictest-conservation federally managed lands are mountainous
- 9.9% of non-mountainous landscapes are federally managed under strictest conservation

**11 western U.S. states**

**Fed** = USFS + DOI lands

**Non-fed** = all other lands

**Mountainous** = Categories C5, C6, & D3-6 from Hammond's (1970) classes of land-surface form

# ... and their montane wildlife

- less biogeographic connection to other montane pop'ns
- high flexibility in reproduction, behavior, seasonality
- variability across latitudinal, elevational, and PPT clines
- **Diverse life-history responses to harshness:**
  - seasonal ranges, migration (elevational) 
  - subnivean activity, year-round 
  - torpor, hibernation, or estivation to over-winter 

# Potential mechanisms of GCC on montane spp.

- Altered food abundance



- Altered community structure



- Habitat fragmentation  species-area losses

- Exceeding (narrow) physiological tolerances

- Acute heat stress

- Chronic thermal stress

- Winter cold stress (melted snow, insulation)

# Potential mechanisms of GCC on montane spp.

- Increased vulnerability to crepuscular/nocturnal predators



- Increased susceptibility to disease, pests, parasites

- Altered snow/ice cover, precipitation, streamflow, humidity, soil moisture, insolation

  - Increased water stress

- **Mtn. ecosystems & their climates & wildlife are poorly understood**



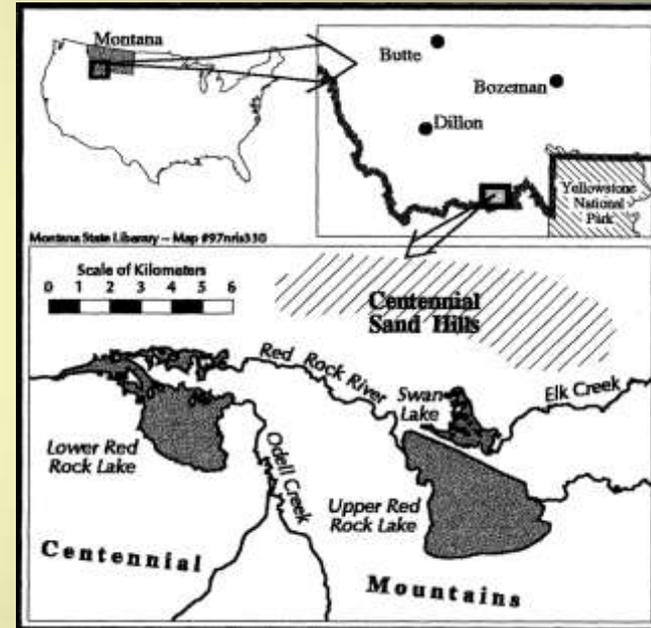
# Phenological mismatches for migr. birds

- **Red Rock Lakes NWR** – establ. 1935, for trumpeter swans and migratory birds
- on W edge of GYE; long history w/humans
- 66% wilderness; elevations 1,900 - 3,100 m



# Phenology: *the stage & actors*

- Migratory passerines eat plants, insects
- Encompass elevational gradients (space-t)
- Pre- and post-breeding
- Birds, bugs, plants, climate
- Energetics affect fecundity



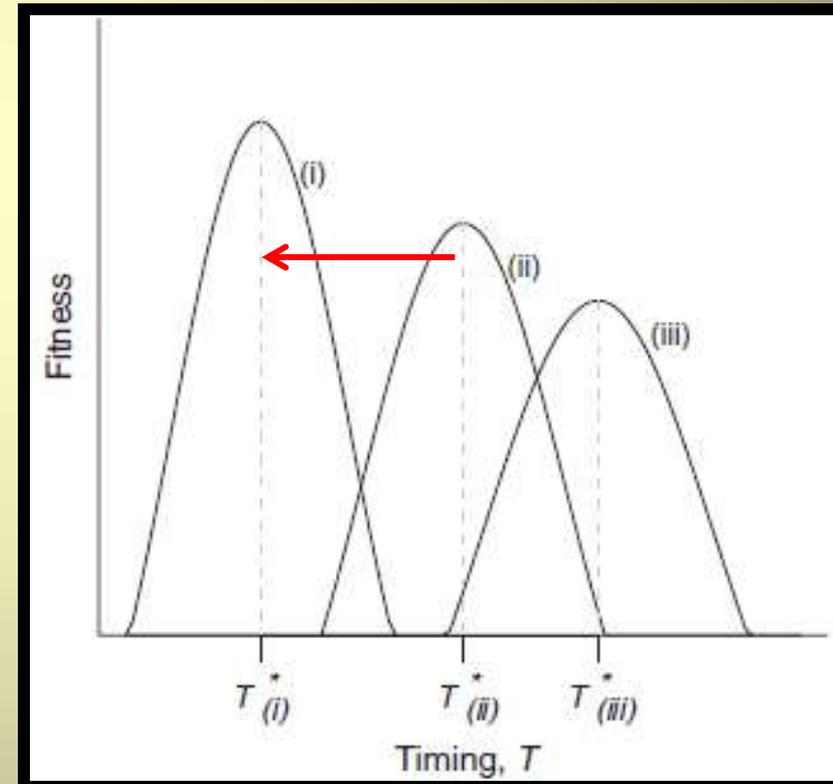
Vesper sparrow  
Broadly distributed  
<10% decline, last 10 yrs  
1 ssp. Of Concern

Brewer's sparrow  
Sage-obligate  
Significant ↓, last 10-20 yrs  
S2 – S5; of cons. interest



# What is phenological mismatch?

- ❶ Migration timing → day length
- ❷ Insect emergence → degree-days
- ❸ Birds arrive after insect emergence passed
- ❹ Climate has warmed



# Phenology: *objectives*

- **Identify absolute & relative timing of bird arrival, insect emergence, & plant flowering;**
- **Characterize the shape of the temporal change in these events throughout the snow-free season;**
- **Compare above across areas experiencing different magnitudes of climate change;**
- **Quantify fitness consequences of birds consuming plants vs. insects;**
- **Compare mismatch in specialist vs. a more-generalist bird species**

# Phenology: *results to date*

- Seasonal changes in body mass differed between spp.
- In Brewer's sparrows, mass declined on recently grazed areas, slightly increased on areas w/ 7 yrs rest



# Phenology: *results to date*

- **Seasonal changes in insect abundance differed between size classes**



# Phenology: *results to date*

- **Territorial males captured from arrival through pre-breeding & egg-laying periods**
- **Declines in insect contribution to diet throughout season**
- **Decline sharper in Brewer's sparrows**

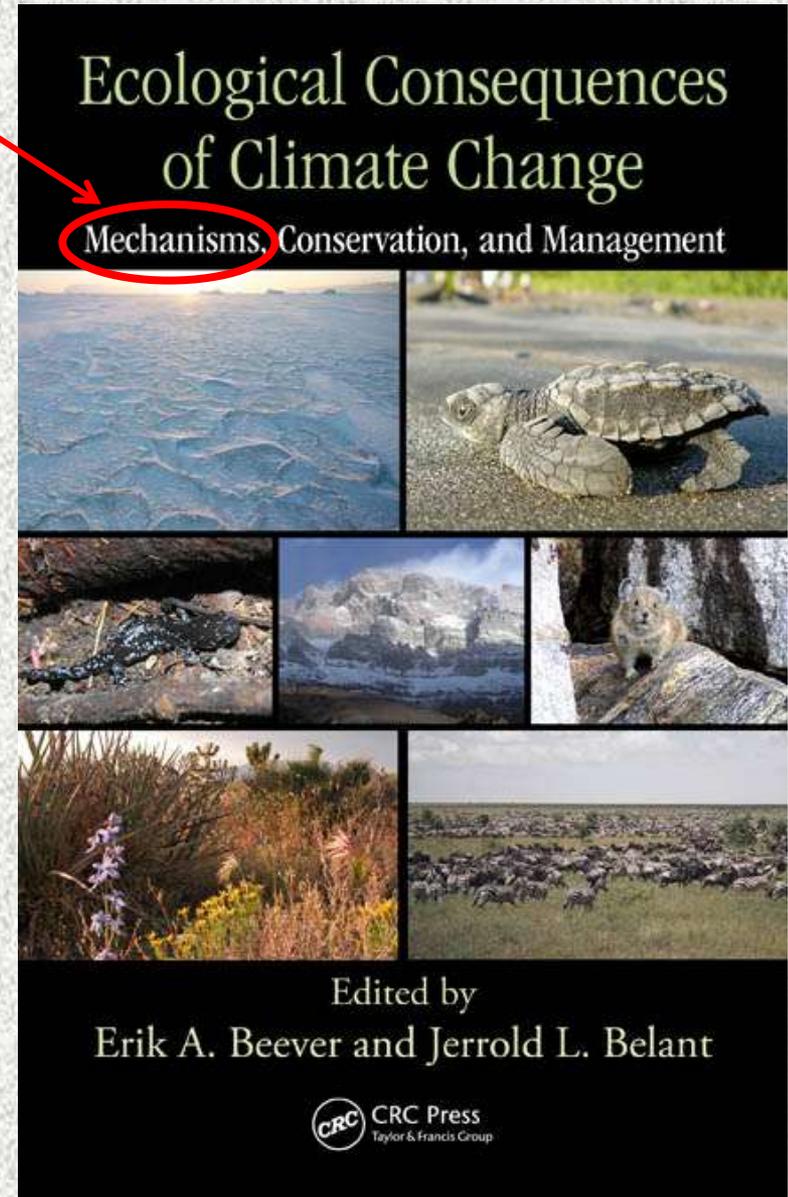
# Phenology: *collaborators*

- **Red Rock Lakes NWR**
- **The Nature Conservancy**
- **Montana Fish, Wildlife, and Parks**
- **University of California, Santa Cruz**
- **University of Montana-Western**
- **University of Rhode Island**
- **U.S. Geological Survey**



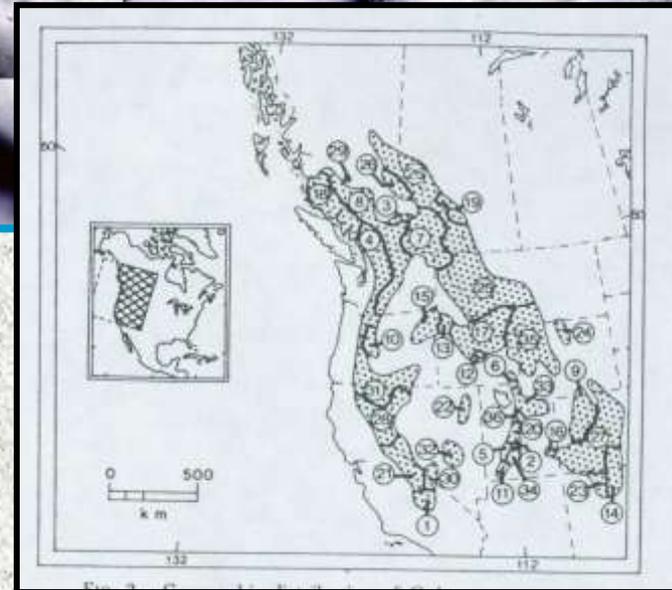
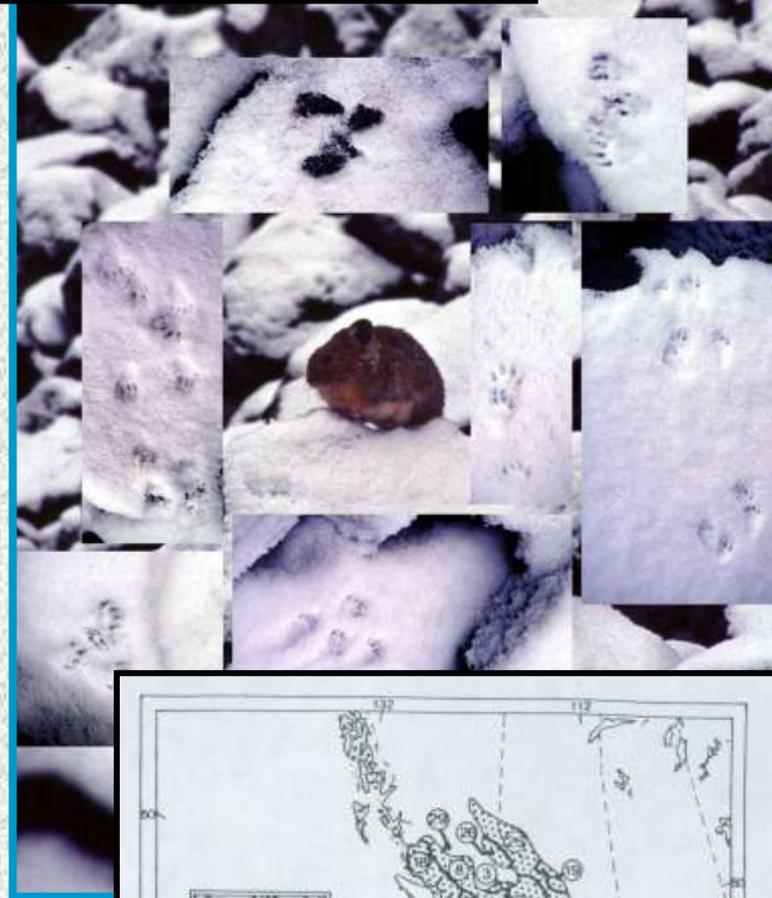
# Mechanisms are *very* important !

- Help understand *why* and *how* behind who, where, what
- Essential for adaptation, mitigation mgmt & conservation strategies



# *Ochotona princeps* Richardson

- **Coprophagous** (to conserve H<sub>2</sub>O)
- **Territorial** (cheek-gland rubbing)
- Broad repertoire of **vocalizations** (7)
- **Active** year-round (rare for montane)
- Inhabit **only talus** and talus-like areas
- **Don't move** very far → radiation



# *Pikas as model sp. for ecological-niche testing*

- Locally abundant; rare 4 mammals
- Relatively stable population sizes
- Highly detectable (haypiles, calls)
  - Monitoring, research are less expensive
- Easily defined habitat, NOT changing over time
  - losses *not* confounded by habitat change
- HSTal records & 15 yrs of recent data indicate a changing distribution



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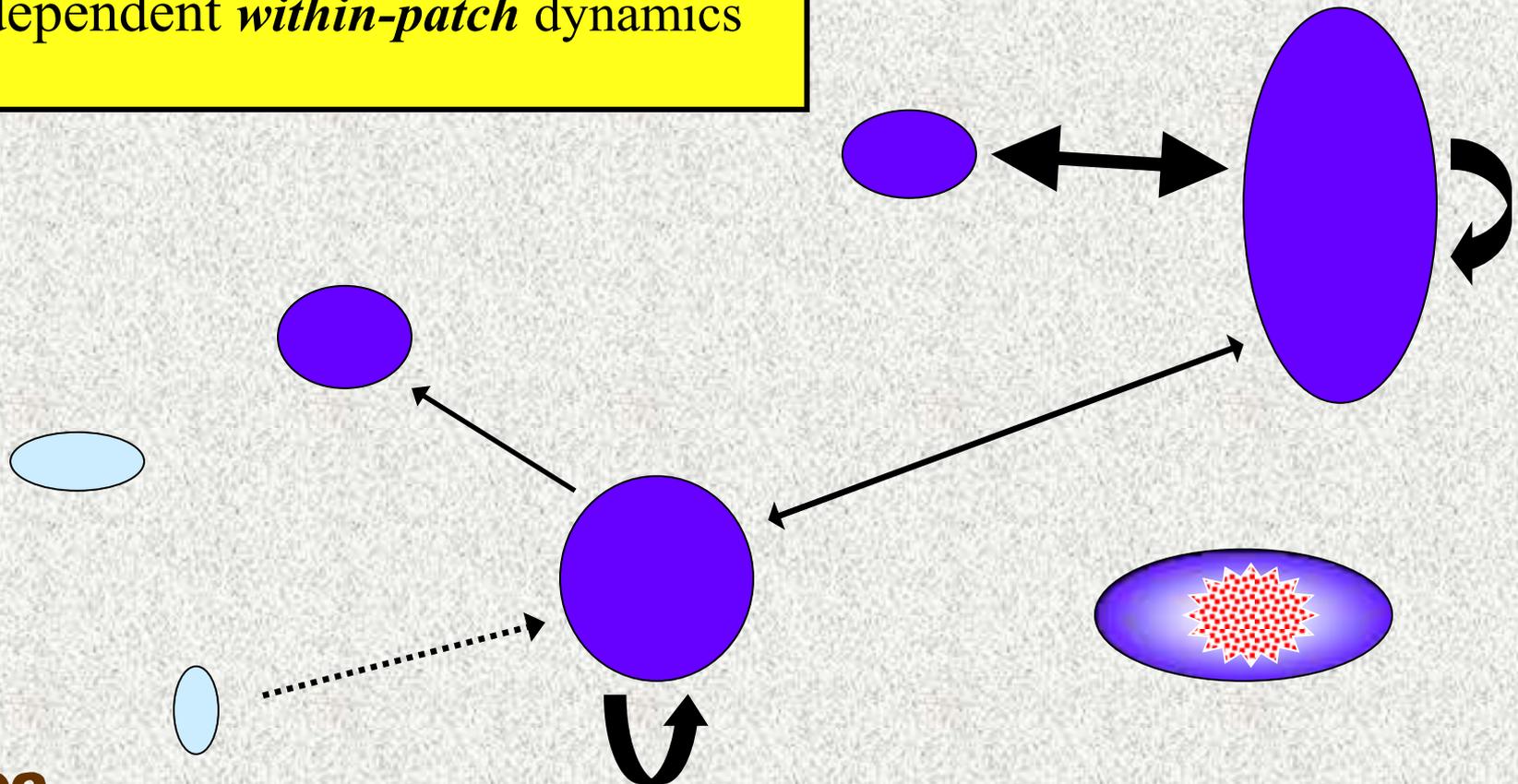
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# Dynamics of metapopulations

- Extinction-colonization dynamics
- Low dispersal frequency
- Independent *within-patch* dynamics



# Questions $\Rightarrow$ Hypotheses

- Have there been any distributional changes since historical specimen records?
- What combination of factors was responsible for changes (if any)?
- Did the pace and drivers of losses differ between 20<sup>th</sup> Century and last decade?
- Across broad domains, is pattern of site-level losses best predicted by magnitude of change in climatic attributes, or by relative status of climatic attributes?



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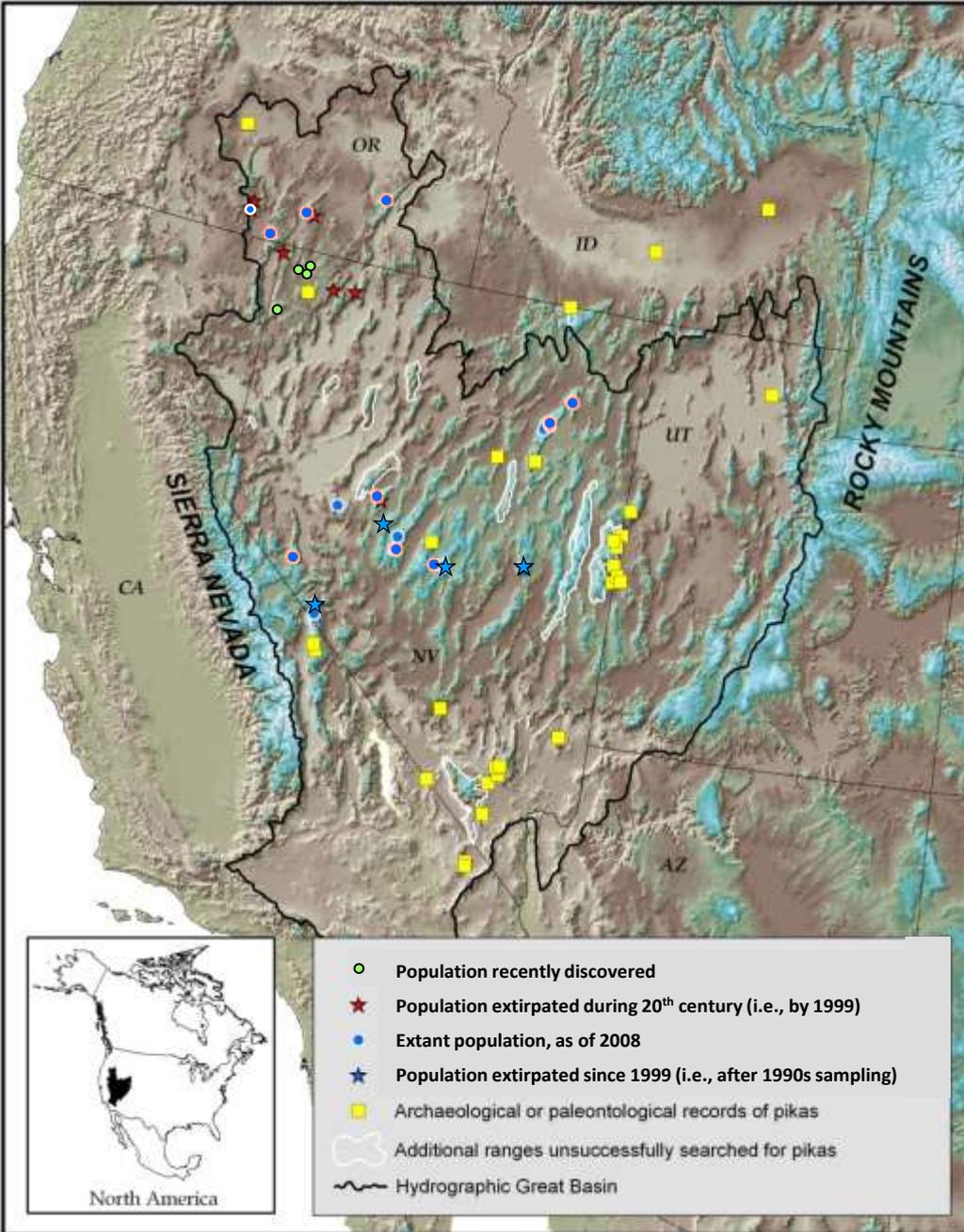


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**Study sites**  
 within the Great Basin  
 (blue areas: >2,286 m)



**3 periods of sampling**

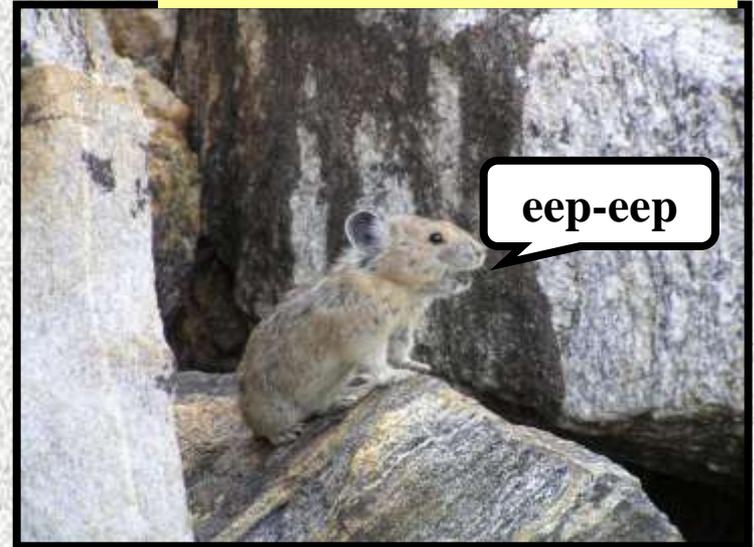
Historic	1898-1956
Recent_1	1994-1999
Recent_2	2003-2008

# Pika evidences

Sighting



Call (AKA 'vocalization')



Active haypile, sighting



# Pika industry



# Old evidences of pikas

Feces: dry



Old haypiles



Feces: moist



**Three Lks. (Lamoille Cnyn), Ruby Mtns., ne NV  
(classic pika habitat)**



**Long Cnyn, Ruby Mtns., ne NV**

**(pikas remain at site, but lowest taluses unoccupied)**





**Pinchot Crk., White Mtns.,  
s-c CA/NV border  
-- loc'n of last fresh HP;  
1 or 2 itinerant indiv's**

**Greenmonster Cnyn., Monitor Range, central NV**  
**last stronghold within site; loc'n of type specimen**



**Peterson Crk., Shoshone Range, central NV**  
**(recent local extinction; multi-scale patchiness of talus habitat)**



# Anatomy of a decline: *persistence*

- 6 local extinctions from historic to end of my 1<sup>st</sup> sampling (**once every 10.7 yrs**)

- 4 add'l local extinctions from 1<sup>st</sup> to end of my 2<sup>nd</sup> sampling (**once every 2.2 yrs**)

- Old evidences

## 3 periods of sampling

Historic 1898-1956

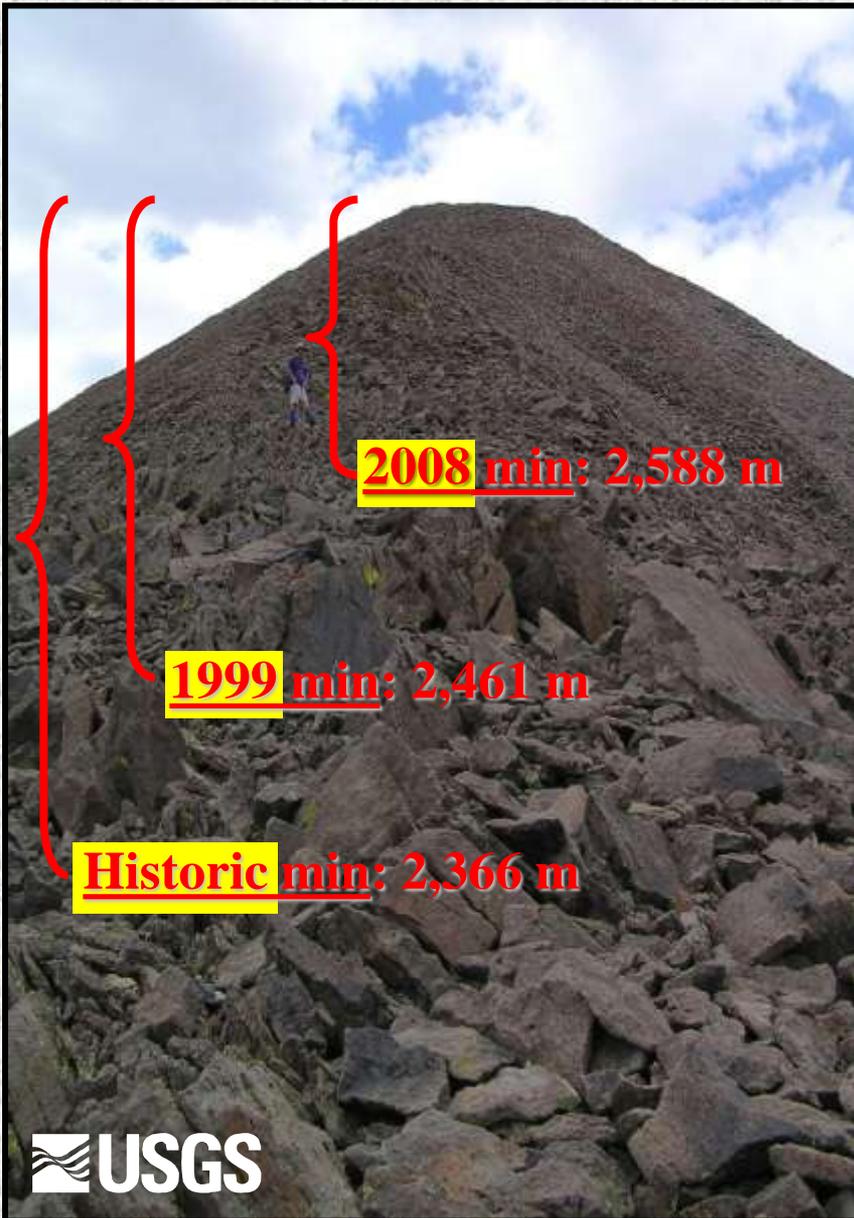
Recent\_1 1994-1999

Recent\_2 2003-2008

**$N = 25$  historical locations**



# Anatomy of a decline: *upslope migrations*



- Minimum elevation of detections, Historic to my first (1990s) sampling: **13.2 m per decade**
- Minimum elev. of detections, 1<sup>st</sup> to 2<sup>nd</sup> sampling: **145.1 m per decade**
  - Parmesan & Yohe (2003) meta-analysis: **6.1 m per decade**
- No  $\Delta$  in max, mean, or median elev, at most sites
- At lower-elevation margins, apparent: a) loss of animals on S-facing slopes, and b) reduced animal densities



# Other species are shifting, too ...

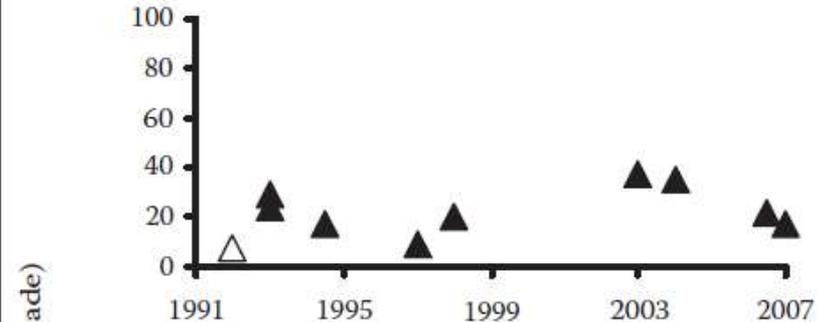
## Observed across diverse taxa

### Evidence from Single Species Studies for Upward Shifts in Elevational Distributions Linked to Climate Change

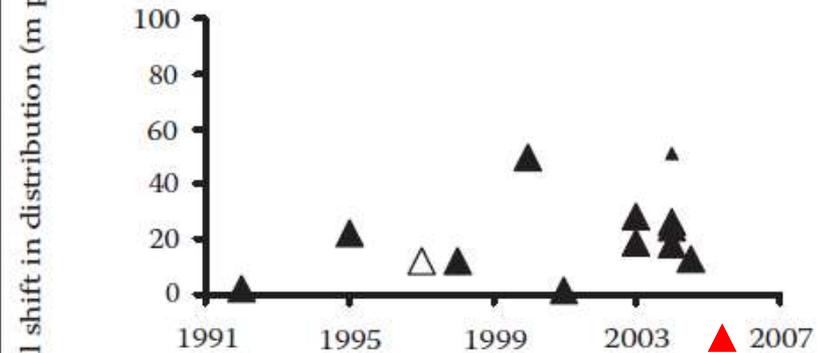
Species and Evidence for Range Shift	Location (Lat°)	References
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<i>Euphydryas editha</i> (butterfly)	Western North America (30–53N)	Parmesan 1996, 2005
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<b>Upper Elevation Expansion</b>		
<i>Thaumetopoea pityocampa</i> (moth)	Italy (46N)	Battisti et al. 2005, 2006
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Wilson & Gutiérrez 2012

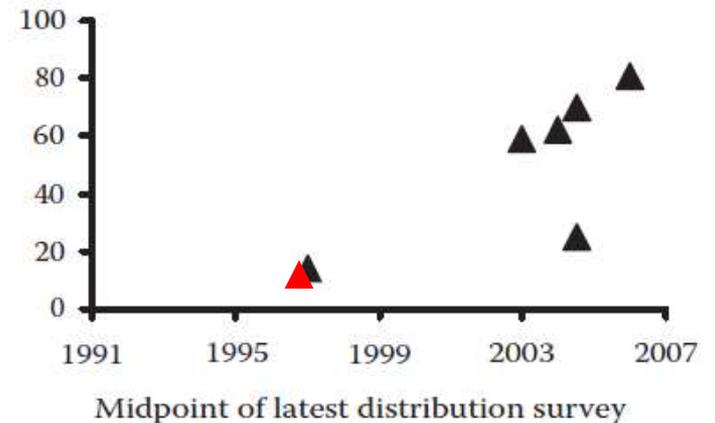
a) Mean elevation



b) Upper limit



c) Lower limit



Elevational shift in distribution (m per decade)

Midpoint of latest distribution survey

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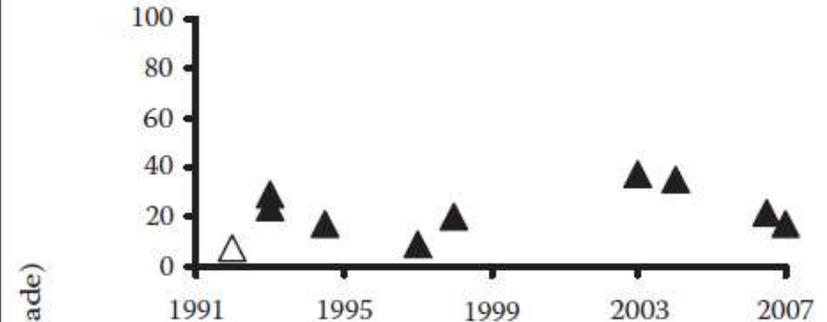
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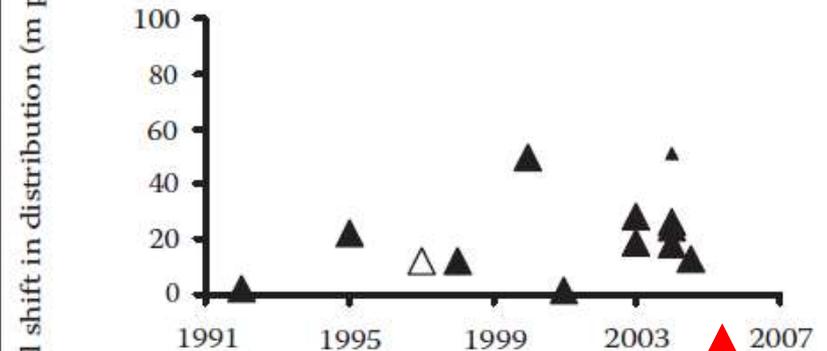
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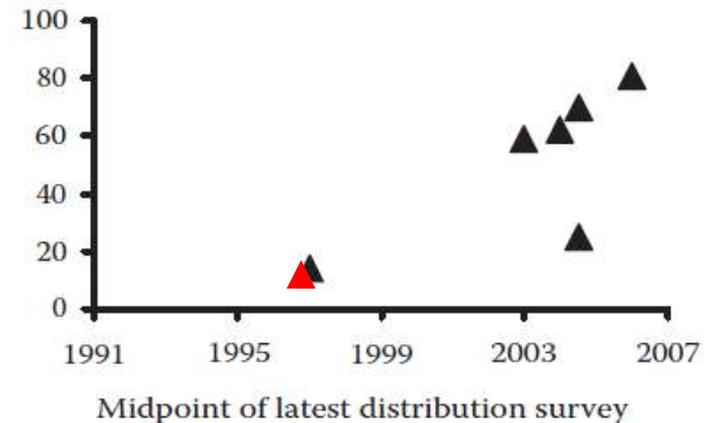
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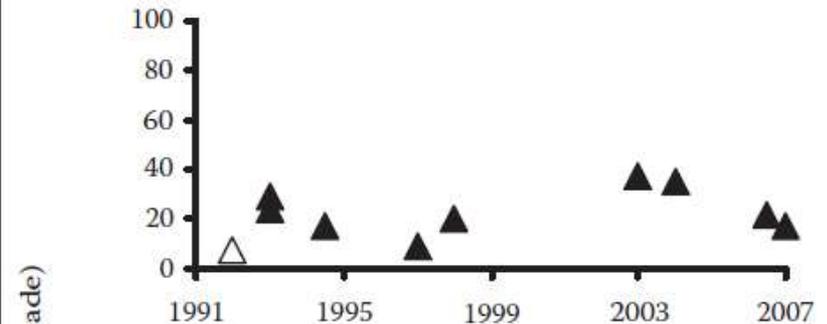
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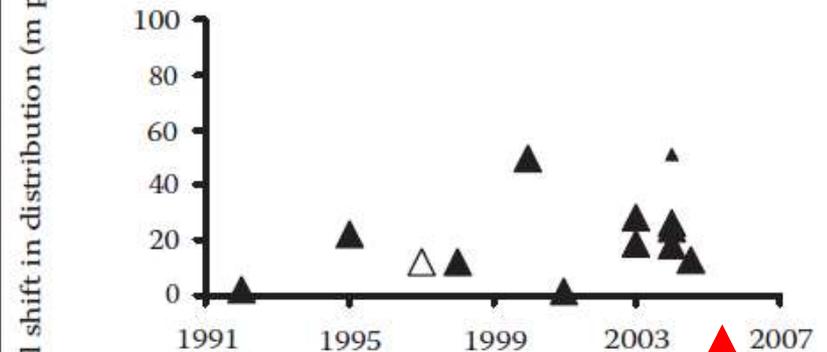
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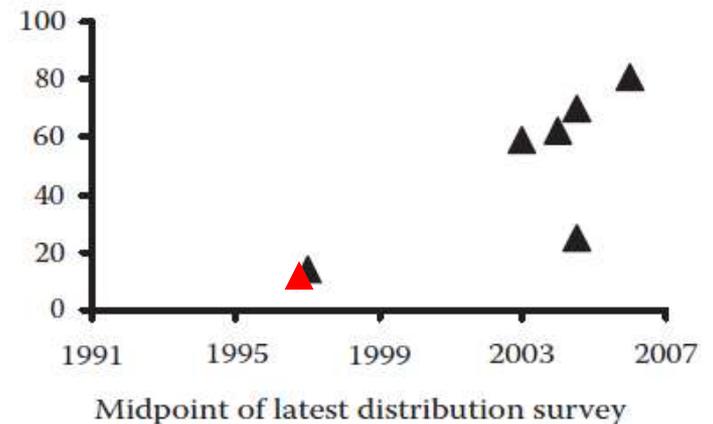
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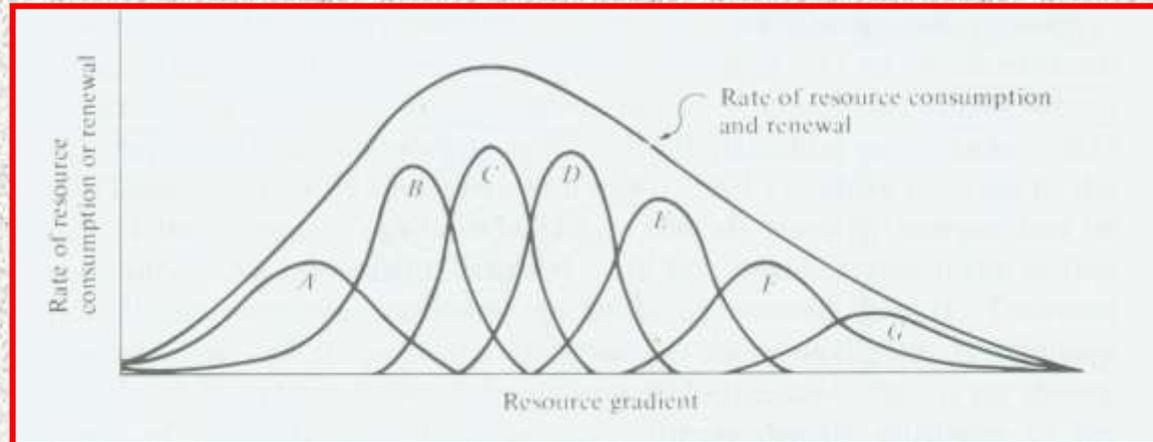


Midpoint of latest distribution survey

# Ecological-niche models

- **Forecasting future responses to climate change**
- **Relate to niche concept**

- **Most assume niches are conserved**

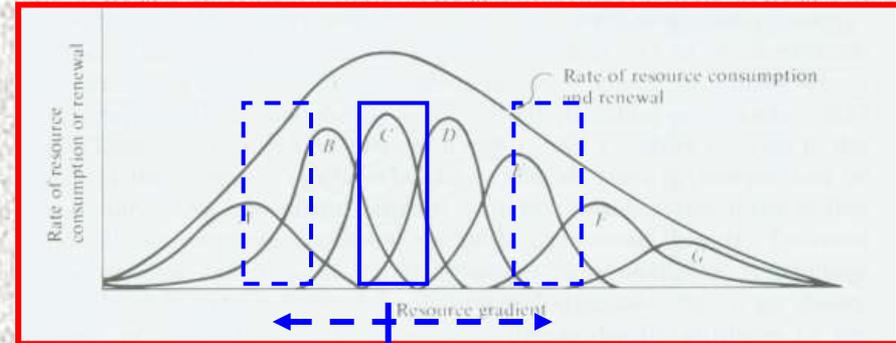


- **Many sources of uncertainty**
- **Often rely on *coarse-scale* data, analysis**
- **Frequently *presence-only* data, modeling**

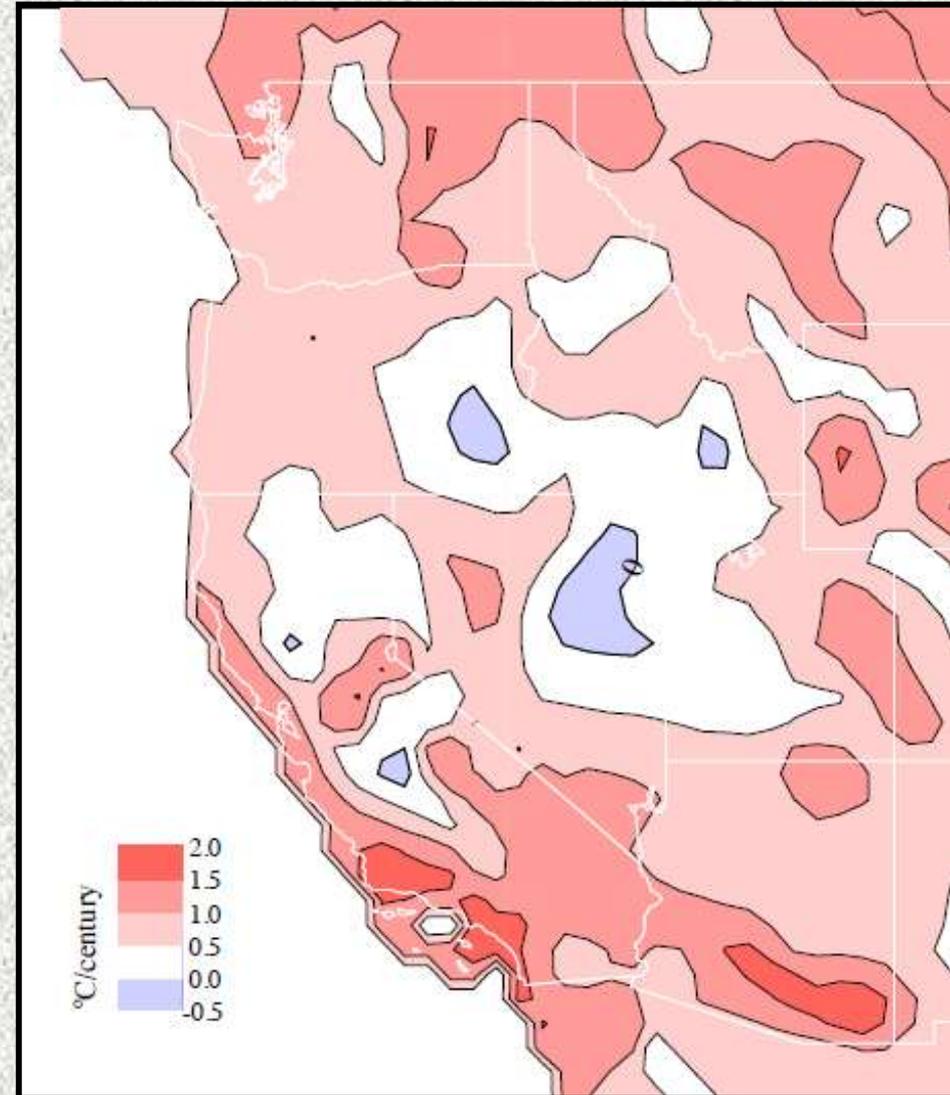
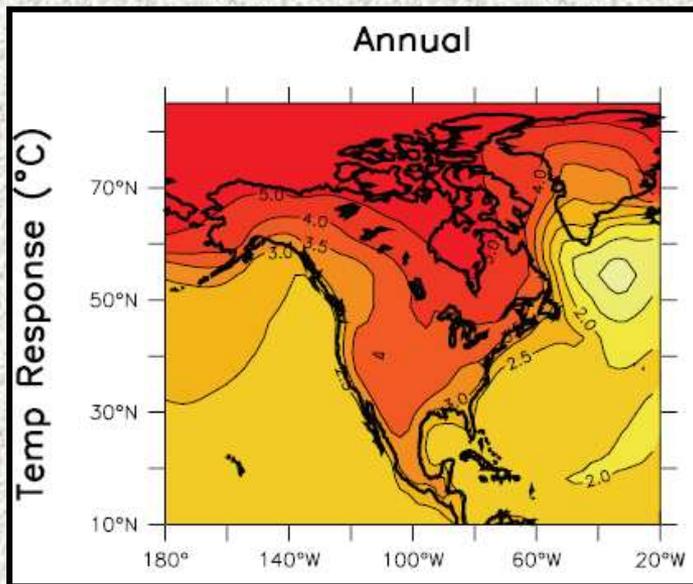
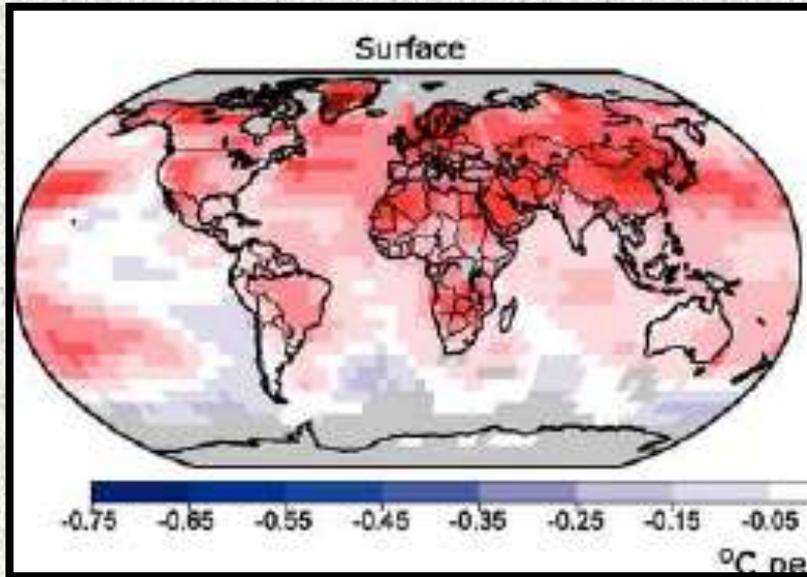
# Modeling changes in species distributions

- Observed evol'n in thermal performance
- Sufficient genetic diversity for local adapt'n
- Pika extirpations latitudinally distributed

**Local adaptation:**  
Losses determined  
by *how much*  
*climate has changed*



# Modeling changes in species distributions: *spatial heterogeneity in amount of climate changes*



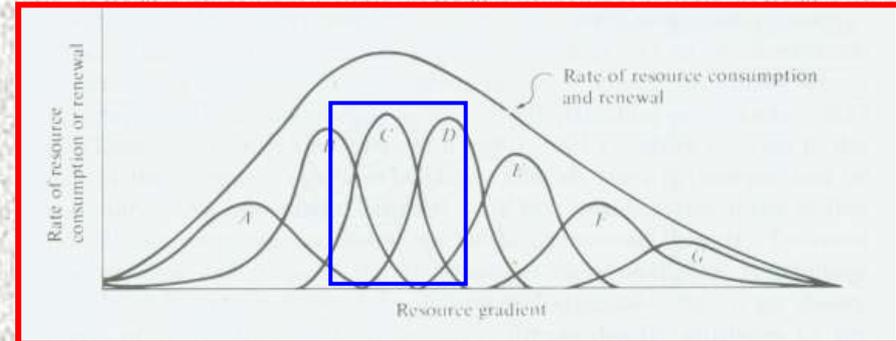
# Modeling changes in species distributions

- Observed evolution in thermal performance
- Sufficient genetic diversity for local adaptation
- Pika extirpations latitudinally distributed
- ***HOWEVER ...***

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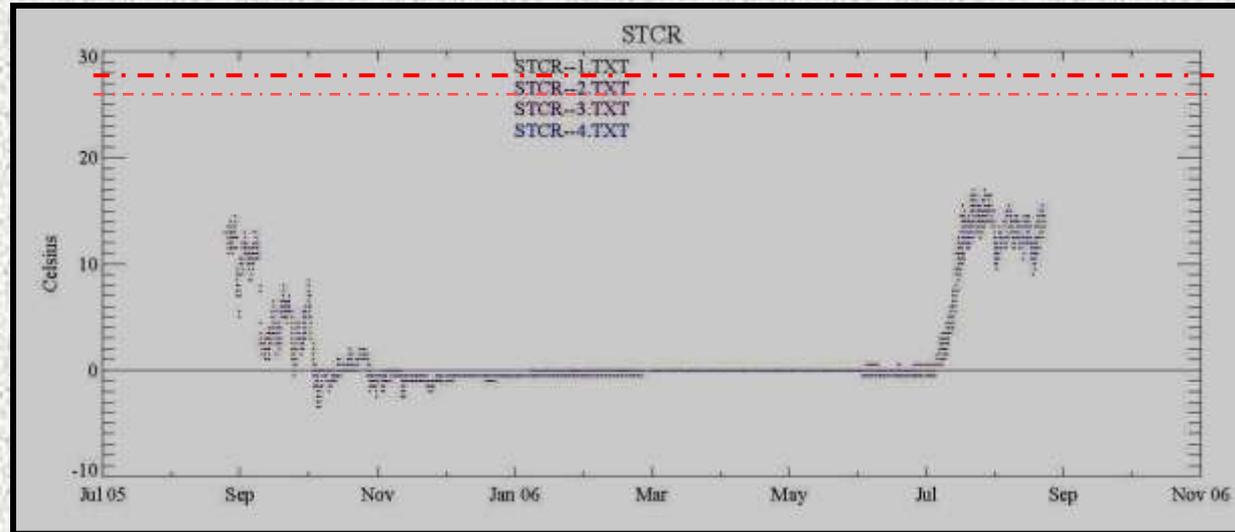


- Most species' losses at S. edge of range
- Rate of climatic change has been **RAPID**
- Energetic, physiological constraints

**Fixed-dimensions niche:**  
Losses determined by  
*proximity to edge of bio-*  
*climatic niche* (i.e.,  
relative status of climate)

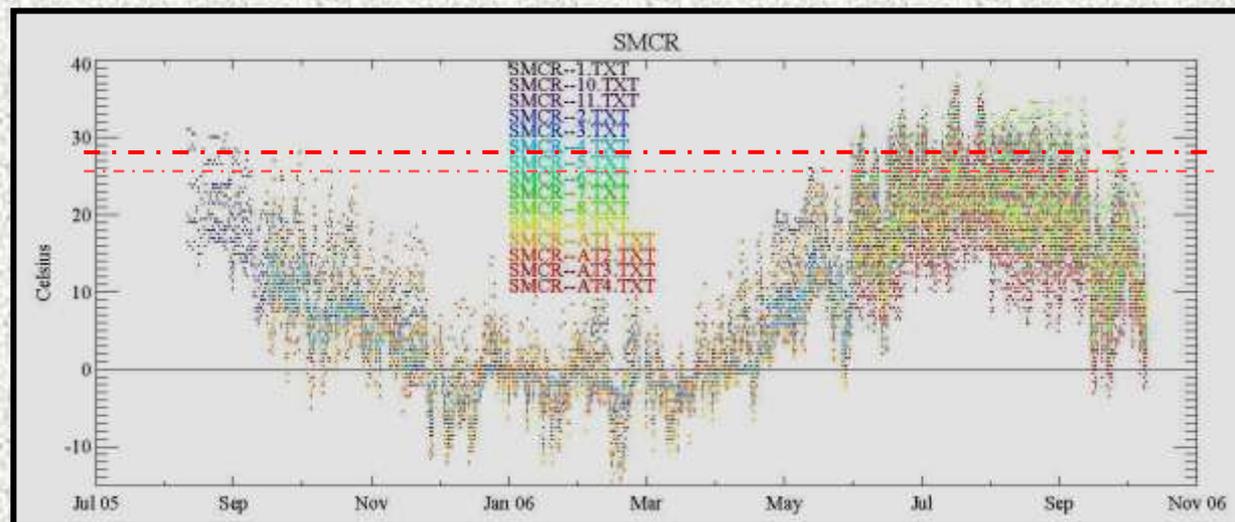
# Potential mechanisms of GCC on montane spp.: *summer heat stress*

- **Pika-occupied** sites rarely had within-talus temps above pika-lethal thresholds



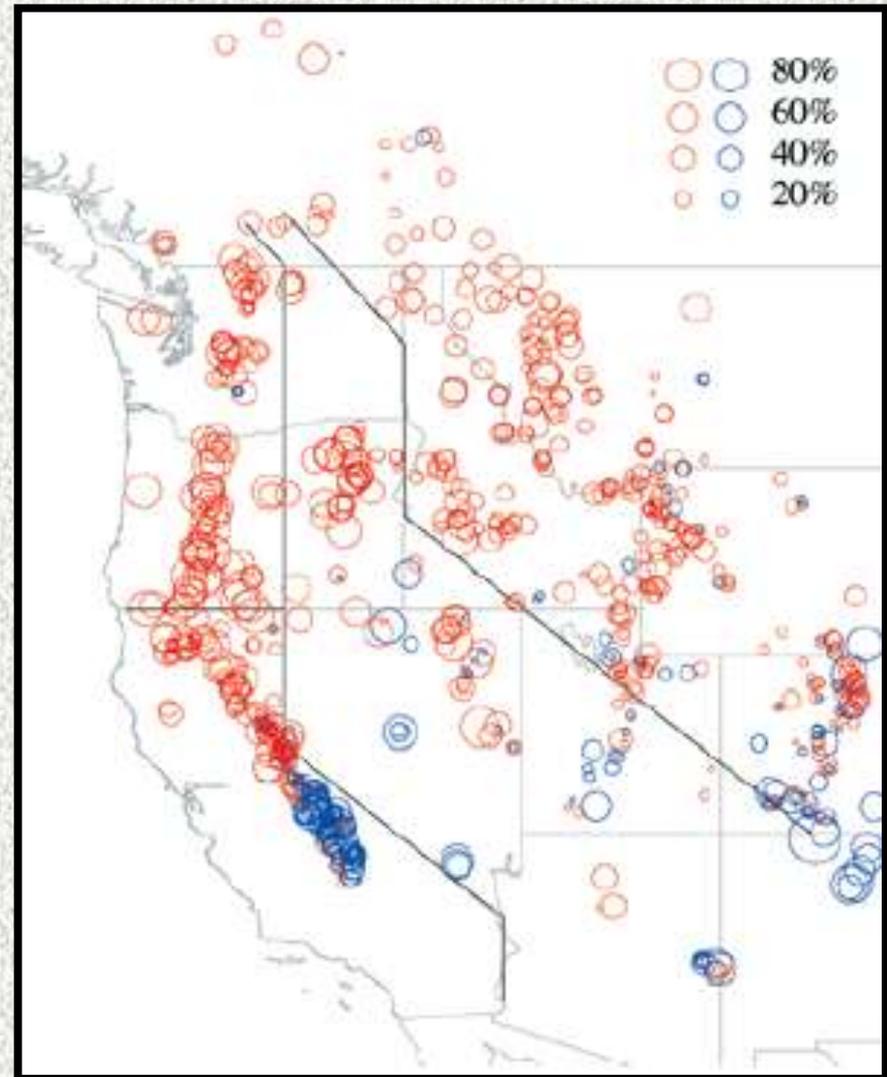
[Beever et al. 2010, \*Ecol. Appl.\*](#)

- **Locally-extinct** sites more often had within-talus temps above pika-lethal thresholds



# Potential mechanisms of GCC on montane spp.: *winter cold stress*

- **Decrease in snowpack (SWE) across 924 snow-course sites in western N. America, 1950-1997 (April 1)**

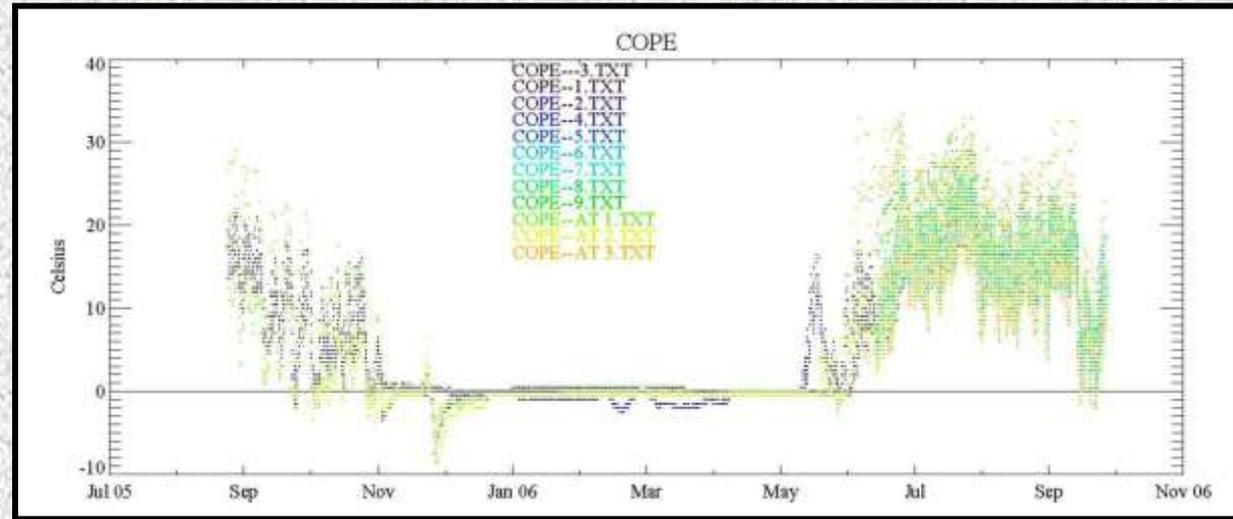


Mote et al. 2003

FWS Safeguarding WL: May 2012

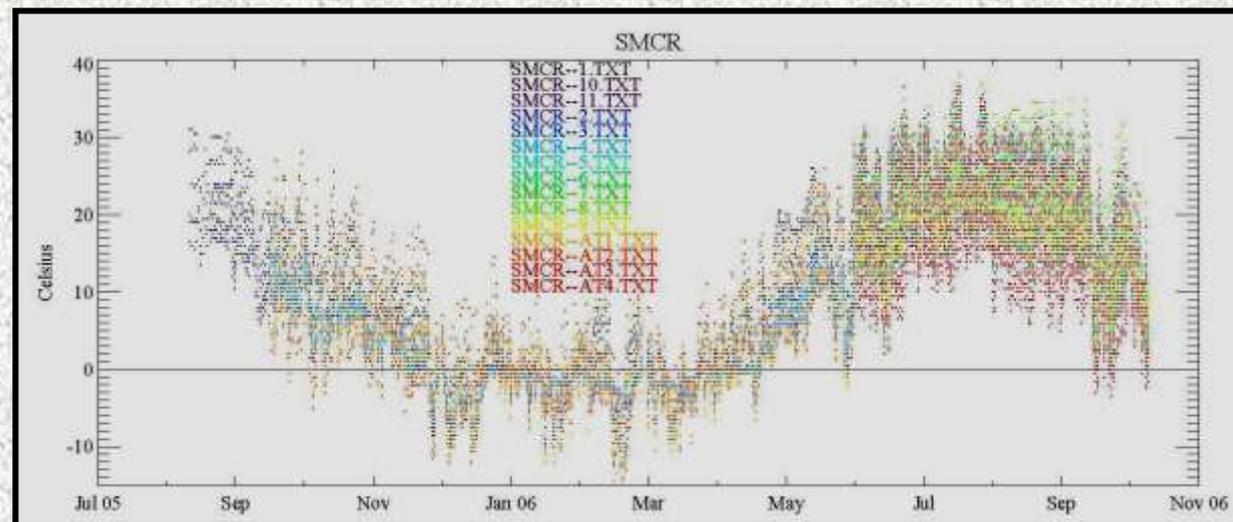
# Potential mechanisms of GCC on montane spp.: *winter cold stress (cont'd.)*

- Most **pika-occupied** sites snow-covered 0.5 – 8.2 months/yr



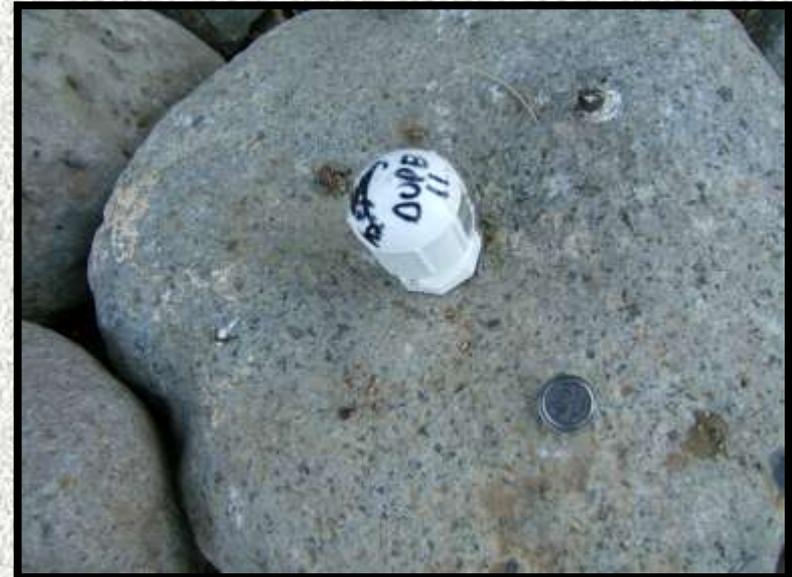
Beever et al. 2010, *Ecol. Appl.*

- 8 of 10 **locally extinct** sites never had snow cover >2 weeks



# Testing: *field and analytical methods*

- ~200 iButtons set in previously pika-occupied taluses (7/05-Present)
- Status of **site** and **talus occupancy** checked  $\geq 5$  times, 2005-2008
- Metrics for 3 mechanisms of direct thermal stress
  - Acute heat stress: # d  $> 28^{\circ}$  C
  - Chronic heat stress: average summer (JJA) temperature
  - Winter cold stress: #d  $< 0$  or  $-5^{\circ}$  C
- Central hypothesis: comparison of data fit between A) field conditions in 2005-06 (i-button data) and B) *magnitude of change* from 1945-1975 to 1976-2006; also investigated *total stress* (1945-2006 avg.)



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# Top models for patterns of persistence

Beever et al. 2010

Model	$-2 \times \log(L)$	$K$	$AIC_c$	$\Delta AIC_c$	Akaike weight	Evidence ratio <sup>-1</sup>
Mean summer temperature (2005–2006)	22.559	2	27.104	0.000	0.148	1.000
Days below $-5^\circ\text{C}$ (1945–2006) and mean summer temperature (1945–2006)	20.330	3	27.473	0.368	0.123	0.832
Days above $28^\circ\text{C}$ (2005–2006)	23.337	2	27.883	0.779	0.100	0.677
Mean summer temperature (2005–2006) and days above $28^\circ\text{C}$ (2005–2006)	20.774	3	27.916	0.812	0.099	0.666
Mean summer temperature (1945–2006)	23.954	2	28.500	1.396	0.074	0.498

- **Recent variables prominent among top models**
- **Importance of types of stress not previously well appreciated**
- **No clear-cut most-plausible single model**

# Importance of different variables for persistence

Beever et al. 2010

Predictor	Mean Akaike weight per model	Sign of effect
Recent mean summer temperature (°C)	0.0629	6-
Long-term mean summer temperature (°C)	0.0530	6-
Recent no. days above 28°C	0.0530	6-
Cumulative no. days below -5°C	0.0491	4-
Recent no. days below 0°C	0.0271	2-, 2+
Cumulative no. days above 28°C	0.0263	6-
Recent no. days below -5°C	0.0252	3-, 1+
Cumulative no. days below 0°C	0.0207	3-, 1+
Change in no. days above 28°C	0.0037	6-
Change in no. days below 0°C	0.0034	4-
Change in mean summer temperature (°C)	0.0022	1-, 5+
Change in no. days below -5°C	0.0014	3-, 1+

- “Climate-change” variables very poorly predicted patterns of loss
- Acute heat stress a relatively poor predictor (behavior)
- Concern @ historic low-temp variables

# Evidence of climatic influence on pikas

- **EXPERIMENTAL: Vulnerability to direct heat stress** (Smith 1974)
- **Hotter, drier macroclimates at extirpated vs. extant sites**
  - **PRISM-modeled data, AND iButtons in taluses across Basin**

	<i>iButton field data, 2005-2006</i>			
	# Days > 28°C	Avg summer temperature (°C)	# Days < 0°C	# Days < -5°C
Pika-extant sites ( <i>N</i> = 15 sites)	2.8 ± 1.0	12.05 ± 1.01	204.4 ± 13.2	15.0 ± 4.6
Pika-extirpated sites ( <i>N</i> = 10)	10.9 ± 4.0	17.02 ± 0.72	159.6 ± 9.7	28.7 ± 7.8

Beever et al. 2010, *Ecol. Appl.*

# Evidence of climatic influence on pikas

- **Within mixed-occupancy sites, extirpated patches 2.5° C warmer during summer than occupied patches**
- **Extant sites received 1.43x greater PPT during 1961-1990 than did 6 sites initially lost, & 1.75x greater PPT during 1971-2000 than 4 sites of recent loss (PRISM)**
- **Change in population-size index most negative at southernmost sites**
- **Loss of pikas from low-elevation sites in 4 ranges even though higher-elevation populations in the same range persisted**

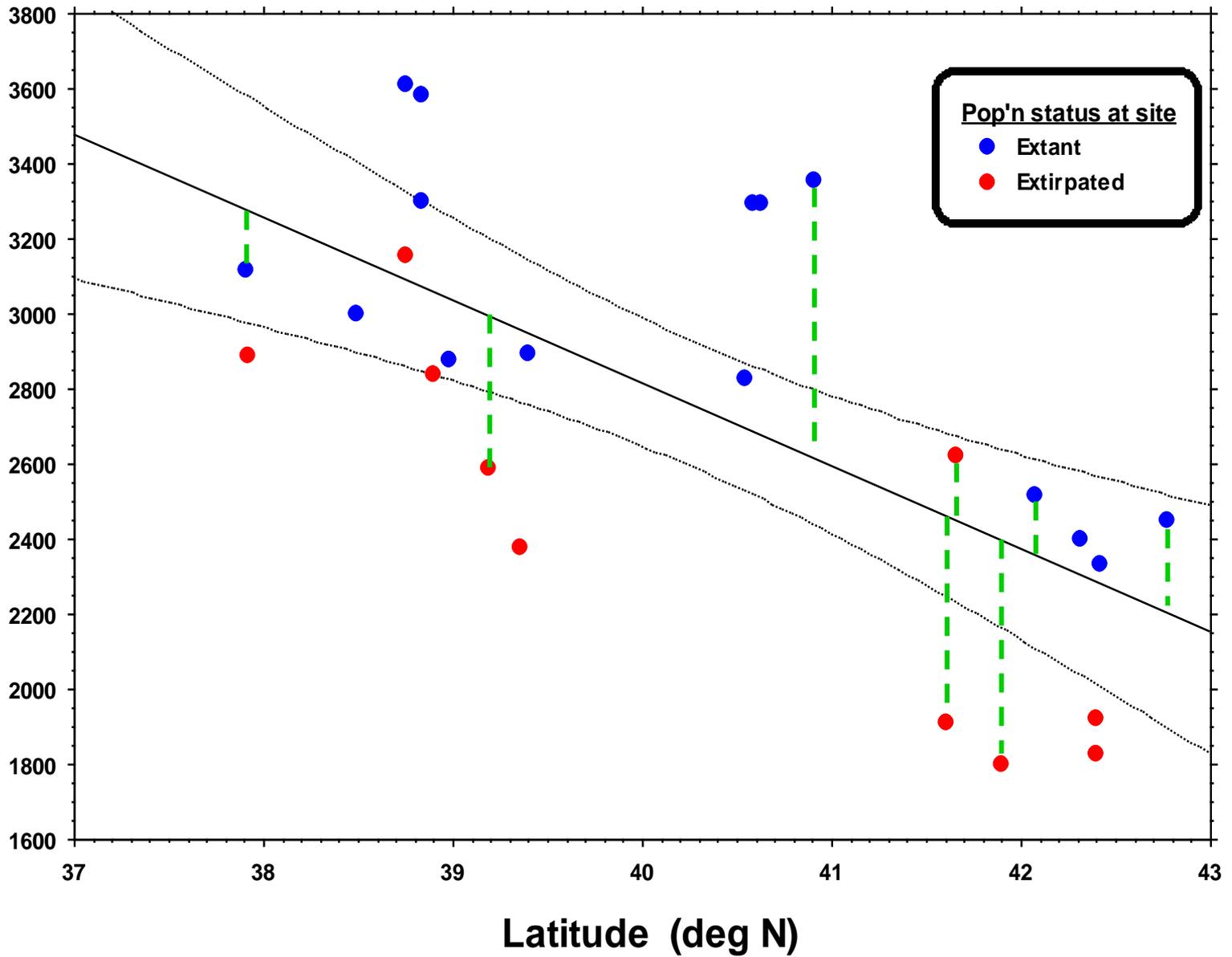
# Different drivers of extirpation? (a)

- **Compare identity and rank of top models from extirpations during 20<sup>th</sup> Century, last decade**

# Different drivers of extirpation? (a)<sub>1</sub>

- Compare identity and rank of top models from extirpations during 20<sup>th</sup> Century, last decade
- ‘Local Refuge’ (MaxElev) alone best predicted persistence from historic records through 1999
- However, for 1999-2008, **null model was plausible** for describing patterns w/Local Refuge
- Residual variable that accounts for site latitude alone strongly predicted pattern, and appeared in 26/28 of plausible models, across all 3 periods

Max. elev'n (in m) of nearby talus



# Different drivers of extirpation? (a)<sub>2</sub>

- Compare identity and rank of top models from extirpations during 20<sup>th</sup> Century, last decade
- **0/8 models plausible with using Local Refuge for 20th-Century persistence appeared in Recent top models; 8/10 of 20th-Century top models plausible with using residual appeared in Recent top models, but ranks differed**

# Different drivers of extirpation? (b)

- Ranks of variable weights shifted dramatically, last decade vs. during 20<sup>th</sup> Century

	'20th Century' (historic to 1999)	'Recent' (1999 to 2009)
<i>(a) Maximum elevation of habitat within upper bound of dispersal distance</i>		
Predictor variable (listed in order of decreasing weight per model)	MaxElev RngHab AugMaxT DistRd GrzPre99	AugMaxT RngHab DistRd GrzPost99 MaxElev
Sites correctly classified*	24/25	16/19
Average of  (weighted P[occ] - occupancy status)  †	0.123	0.276
<i>(b) Residual of Maximum elevation of local habitat on latitude</i>		
Predictor variable (listed in order of decreasing weight per model)	RngHab MaxElevR DistRd GrzPre99 AugMaxT	MaxElevR AugMaxT RngHab GrzPost99 DistRd
Sites correctly classified*	22/25	18/19
Average of  (weighted P[occ] - occupancy status)  †	0.185	0.169

Beever et al. 2011, GCBiol.

# Different drivers of extirpation? (c)

- Use model-averaging of 20<sup>th</sup>-Century results to predict probability of extirpation *after* 1990s surveys for each site;
- test whether probabilities accurately predicted extirpations

## RESULT:

Logistic regression:  $\chi^2_{1,17} = 0.03, P = 0.86$

Beever et al. 2011, *GCBiol.*

# Different drivers of extirpation? (d)

- Compare model-averaged coefficients between 20<sup>th</sup> Century and Overall periods

## RESULTS:

Beever et al. 2011, *GCBiol.*

- MaxElevR ~2x higher for Overall
- Presence of grazing >5x higher for Overall
- Amt of talus habitat in mtn range >9x lower for Overall
- Distance to nearest rd *no longer in top models*

# So WHAT ???

- **Mammals worldwide typically have been endangered primarily by habitat loss, over-exploitation, lg. body size**  
*Schipper et al. 2008*
- **Climate is the single-strongest factor effecting distributional change (increasingly so), yet interacts with other factors to affect population dynamics**
- **When microclimate data are lacking, the residual measure can provide a first estimate of site vulnerability**
- **Early-warning signs of change are valuable**
- **Knowing past dynamics may not help forecast future**

# So WHAT ???

- If talus areas are no longer habitable, relocation efforts in vain
- Species will likely respond individualistically to climatic influence; thus, extrapolation to other species may be risky



# Resiliency: assisted re-introductions(?)

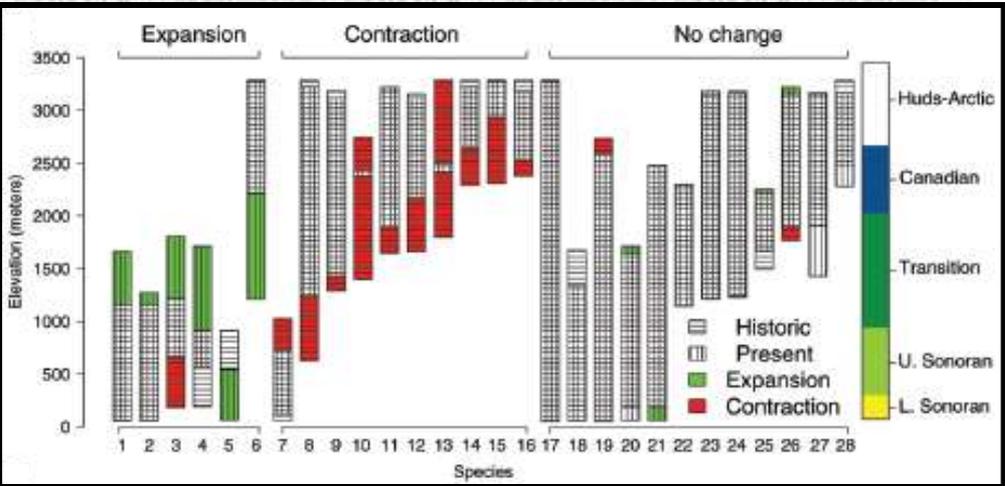
- In instances where re-introductions are considered, recipient locations should have meso- and micro-climates that are conducive to the focal species' persistence



# On extrapolation ...

## Grinnell re-survey, YOSE

No.	Species	P(G)	P(C)	Original elevation range (m)	Range limit change (m)	
<i>Range expansions</i>						
1	<i>Microtus californicus</i>	0.81	0.58	57–1160	+505 U	Elev
2	<i>Reithrodontomys megalotis</i>	0.99	0.87	57–1160	+112 U	Elev
3	<i>Peromyscus truei</i> *	0.99	0.93	183–1220	+589 U, +468 L	Era*
4	<i>Chaetodippus californicus</i>	0.28	0.19	193–914	+800 U	Era*
5	<i>Sorex ornatus</i>	0.32	0.93	549–914	-485 L	Era
6	<i>Sorex monticolus</i>	0.99	0.97	2212–3287	-1003 L	Era
<i>Range contractions</i>						
7	<i>Dipodomys heermanni</i>	0.16	0.98	57–1025	+63 L, -293 U	Era*
8	<i>Microtus longicaudus</i>	0.99	0.98	623–3287	+614 L	Era
9	<i>Zapus princeps</i>	0.98	0.90	1291–3185	+159 L, -64 U	Era
10	<i>Tamias senex</i>	0.95	0.71	1402–2743	+1007 L, -334 U	Elev
11	<i>Spermophilus lateralis</i>	0.70	0.89	1646–3200	+244 L	Era*
12	<i>Sorex palustris</i>	0.39	0.23	1658–3155	+512 L	Era
	<i>Neotoma cinerea</i> *	0.90	0.71	1798–3287	+609 L, -719 U	Era*
	<i>Spermophilus beldingi</i> *	0.98	0.98	2286–3287	+355 L	Elev
	<i>Tamias alpinus</i>	0.92	0.95	2307–3353	+629 L	Era
	<i>Ochotona princeps</i> †	NA	NA	2377–3871	+153 L	NA
<i>No change</i>						
	<i>Peromyscus maniculatus</i> *	0.99	0.99	57–3287	No change	Era*
	<i>Thomomys bottae</i> †	NA	NA	57–1676	No change	NA
	<i>Spermophilus beecheyi</i>	0.50	0.82	61–2734	-250 U	Era*
	<i>Neotoma macrotis</i>	0.90	0.91	183–1646	+67 U	Elev
	<i>Peromyscus boylii</i>	0.98	0.97	183–2469	-122 L	Elev
	<i>Sorex trowbridgii</i>	0.71	0.88	1160–2286	No change	Elev
	<i>Microtus montanus</i> *	0.81	0.98	1217–3155	No change	Elev
	<i>Tamiasciurus douglasi</i> *†	NA	NA	1229–3185	No change	NA
25	<i>Tamias quadrimaculatus</i>	0.95	0.85	1494–2210	+50 U	Era*
26	<i>Tamias speciosus</i> *	1.00	1.00	1768–3155	+128 L, +65 U	Era*
27	<i>Thomomys monticola</i> †	NA	NA	1905–3155	No change	NA
28	<i>Marmota flaviventris</i> †	NA	NA	2469–3353	No change	NA



## Responses differed across taxa

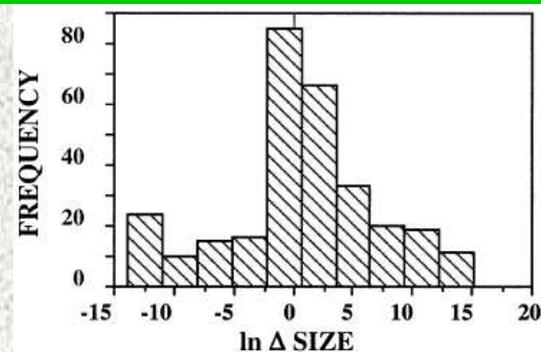
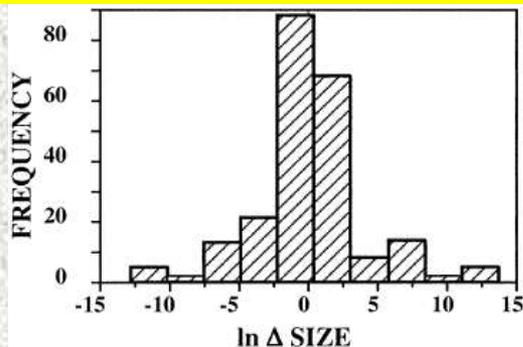


# Species have shifted differently ...

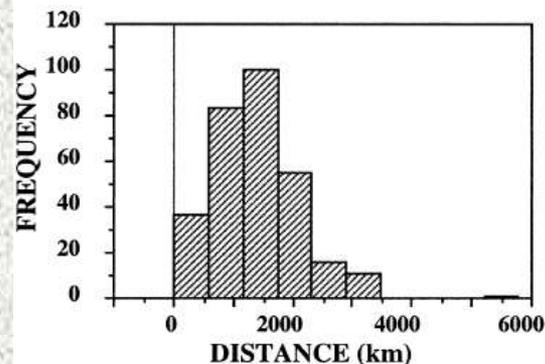
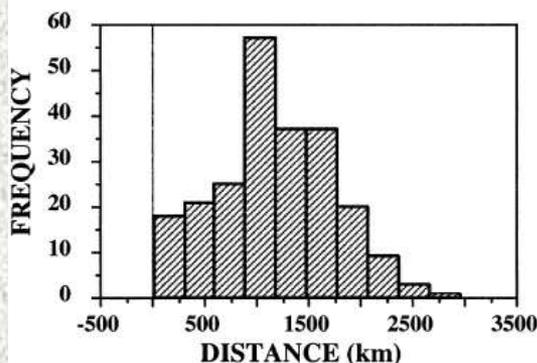
Lyons 2003, *J. Mammal.*

• During paleo times, too ... wildlife also shifted, diversely

Δ in size of geographic range



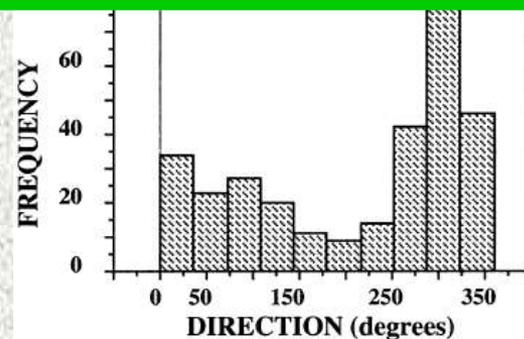
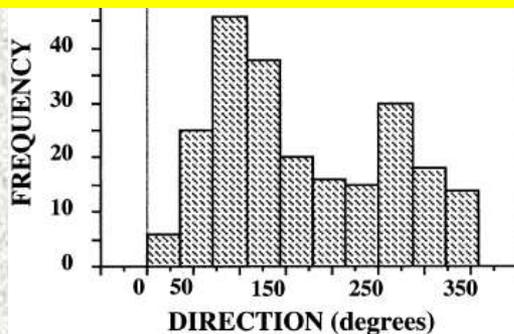
Distance of range shift of centroid



Pre-Glacial to Glacial

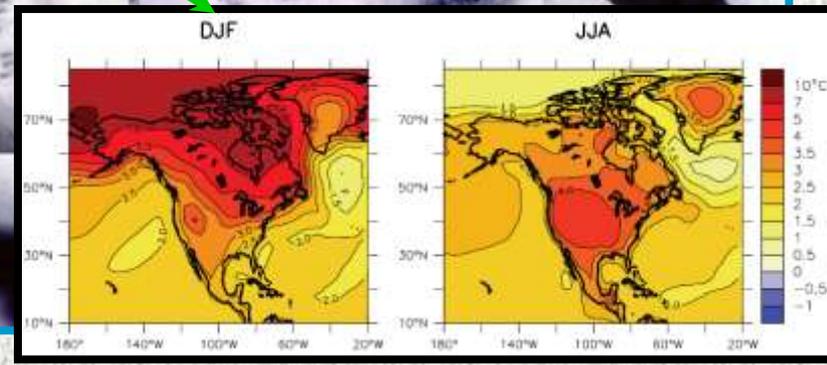
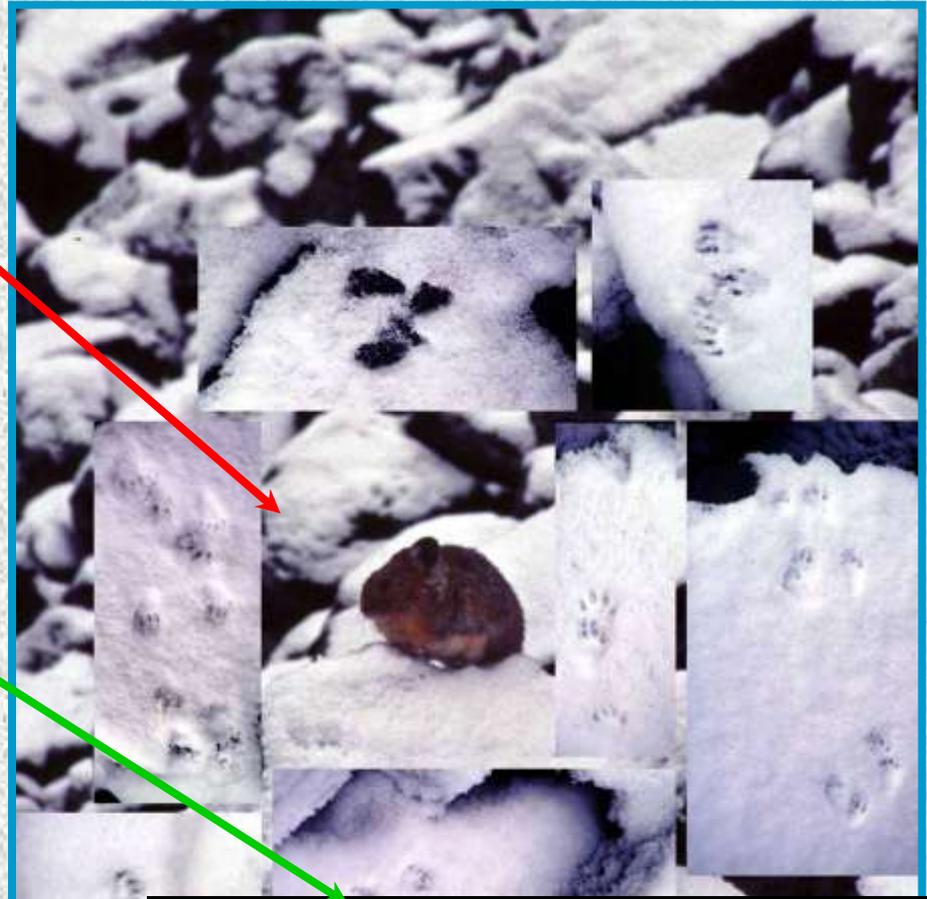
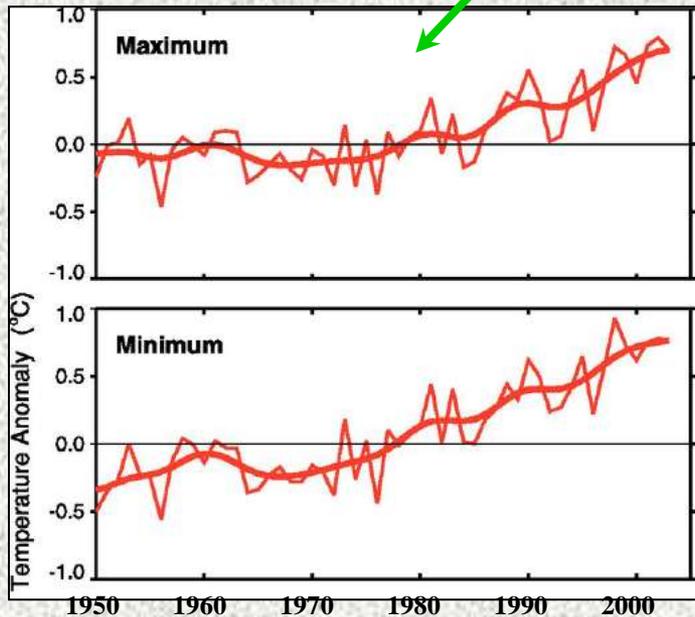
Glacial to Holocene

Azimuth of range shift of centroid



# Interpretation

- Importance of winter cold stress agrees with facts that:
  - a) pikas are active yr-round (consequent E concerns), &
  - b) T minima have risen > T maxima



# Benefits of broad-scale approaches

- Provide greater collective sample size, power
- Broader domain of inference where results apply
- Allow discrimination of broad-scale patterns from local anomalies
- Allow detection and identification of cross-scale dynamics, synergistic interactions, and ecological thresholds
- Allow for assessment of greater diversity of potential system drivers

# Methodological frontiers



- Airborne, geo-rectified IR and veg. sensors
- Derived ecohydrologic variables
- Physiological & genetic studies; facets; ENM modeling => sufficiency of areas, connectivity
- Bayesian approaches



# Final take-home messages

- **Must proceed from solid understanding of life- and natural history**



**Wolverine**



**Whitebark pine**

Behavioral plasticity to accommodate climate stress needs to be understood and incorporated

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- **Multi-pronged approaches will provide best insights into specific mechanisms by which GCC acts upon species**



# Final take-home messages



- **Distributional determinants may vary across species' ranges**
  - **context-dependencies** likely to profoundly affect wildlife responses
    - **Distributional changes in pikas have been observed in ne CA, NV, se OR, the Cascades, UT, CO, s. ID, YOSE, BC, & other locations; *variable magnitudes*.**
  - **decoupling of climate envelopes and static habitats**

# Final take-home messages

- Must proceed from solid understanding of life- and natural history
- Empirical verification of trend necessary for some wildlife spp.
- Multi-pronged approaches will provide best insights into specific mechanisms by which GCC acts upon species
- Distributional determinants can vary across species' ranges
  - hypothesize that winter cold stress more important, here
  - disjunction between climate envelopes and static habitats
- **Behavioral plasticity to accommodate climate stress needs to be understood and incorporated**

# *Thanks !*

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WWF Climate Change Program  
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