



# Index of Humboldt parameters covered

col.env.....	3	R.....	14
color.ramp.....	19	rarefy.dist.....	6
e.var.....	3	rarefy.units.....	6
env.reso.....	5	reduce.env.....	8-11
env.trim.....	7	reductype.....	8-11
env.trim.type.....	7	sp1.....	4
env1.....	3	sp2.....	4
env2.....	3	thresh.espace.z.....	15
kern.smooth.....	13	trim.buffer.sp1.....	7
nae.window.....	12	trim.buffer.sp2.....	7
non.analogous.environments...	10-11	trim.mask1.....	7
p.boxplot.....	18	trim.mask2.....	7
p.overlap.....	16		
p.scatter.....	17		

## Input Parameters:

col.env

e.var

env 1

env 2

# Input Files

**Environmental Data:** environmental variables for all sites of the study area 1 (env1) or study area 2 (env1).

Column names should be x, y, X1,X2,... Xn; with X1-Xn being any string label. If env1=env2, input the same file twice. I typical import tab delimited (.txt) or comma separated files (.csv).

Column numbers as R sees them:

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21
x																					
y																					
BIO_1																					
BIO_2																					
BIO_3																					
BIO_4																					
BIO_5																					
BIO_6																					
BIO_7																					
BIO_8																					
BIO_9																					
BIO_10																					
BIO_11																					
BIO_12																					
BIO_13																					
BIO_14																					
BIO_15																					
BIO_16																					
BIO_17																					
BIO_18																					
BIO_19																					

**col.env** Only if reductype= "STANDARD" (else ignore this parameter), this is the parameter specifies the number of columns to trim environmental space on. This can be any number of columns.

### EXAMPLE

```
col.env= c(3:10)
e.var= c(3:21)
```

**e.var** The selection of variables to include in all of the analyses. Note: this is a separate, more generalized parameter than **col.env**, but must contain all variables specified in **col.env**

## Input Parameters:

sp1

sp2

# Input Files

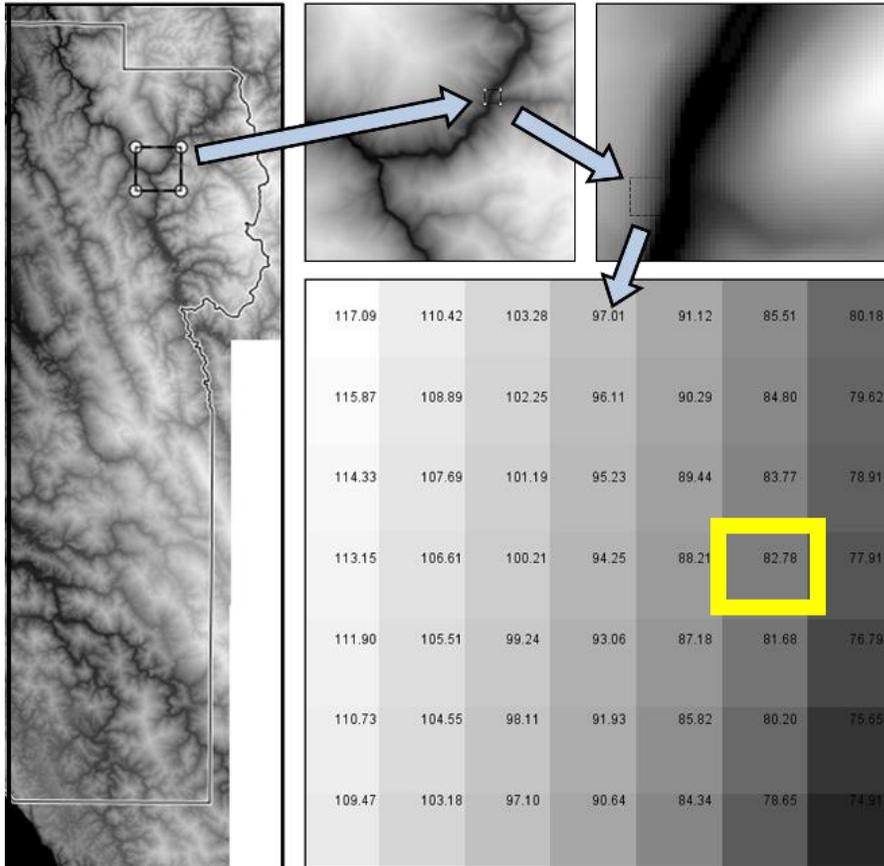
**Species or Population Data:** occurrence sites for species/population 1 (**sp1**) at study area 1 (**env1**) or species/population 2 (**sp2**) at study area 2 (**env2**). Column names should be 'sp','x','y'. I typical import tab delimited (.txt) or comma separated files (.csv).

sp	x	y
Conium_maculatum_Native	-4.82	37.84
Conium_maculatum_Native	-4.84	38.74
Conium_maculatum_Native	-3.64944	40.22694
Conium_maculatum_Native	9.15	41.38333
Conium_maculatum_Native	-2.762	42.556
Conium_maculatum_Native	-2.626	42.916
Conium_maculatum_Native	3.62621	46.7645
Conium_maculatum_Native	3.46469	46.8137
Conium_maculatum_Native	3.27334	46.8646
Conium_maculatum_Native	3.09221	47.0176
Conium_maculatum_Native	1.64331	48.0519
Conium_maculatum_Native	3.72252	48.2556
Conium_maculatum_Native	2.39086	48.4391
Conium_maculatum_Native	1.96675	48.6847
Conium_maculatum_Native	9.96	49.77
Conium_maculatum_Native	-5.13957	49.992
Conium_maculatum_Native	-5.01148	50.17534

Input  
Parameter:  
env.reso

# Input Files

**env.reso** = the resolution of the input environmental data (**env1** and **env2**) in decimal degrees. This will be obtained from the raster GIS data used to create environmental data files (**env1** and **env2**)



EXAMPLE

env.reso=0.41669

Property	Value
<b>Raster Information</b>	
Columns and Rows	4320, 1800
Cell Size (X, Y)	0.083333338, 0.083333338
Format	GRID
Source Type	Generic
Pixel Type	signed integer
Pixel Depth	16 Bit

0.083333338 decimal degrees

Note:

Here the raster data were clipped to the desired extent in ArcGIS. Then one of the clipped rasters was converted to a point dataset and latitude and longitude was added to this shapefile. Then all the raster layers were sampled using the point dataset. Lastly the final point shapefile was saved as a ".csv" or text file, cleaned up in Excel and the saved for importation into R.

Input  
Parameters:

rarefy.dist  
rarefy.units

# Preparing Occurrence Data

**Why rarefy occurrence data?** For optimal performance, this method requires input occurrence data to be spatially independent. It is common for researchers to introduce environmental biases into their analyses from spatially autocorrelated input occurrences. This causes niche estimation to be over-fit towards environmental bias resulting from sampling bias introduced from spatially clustered occurrences. The elimination of spatial clusters of localities is important for proper evaluation of a species niche. When spatial clusters of localities exist, species niches can be overly concentrated in certain E-space.

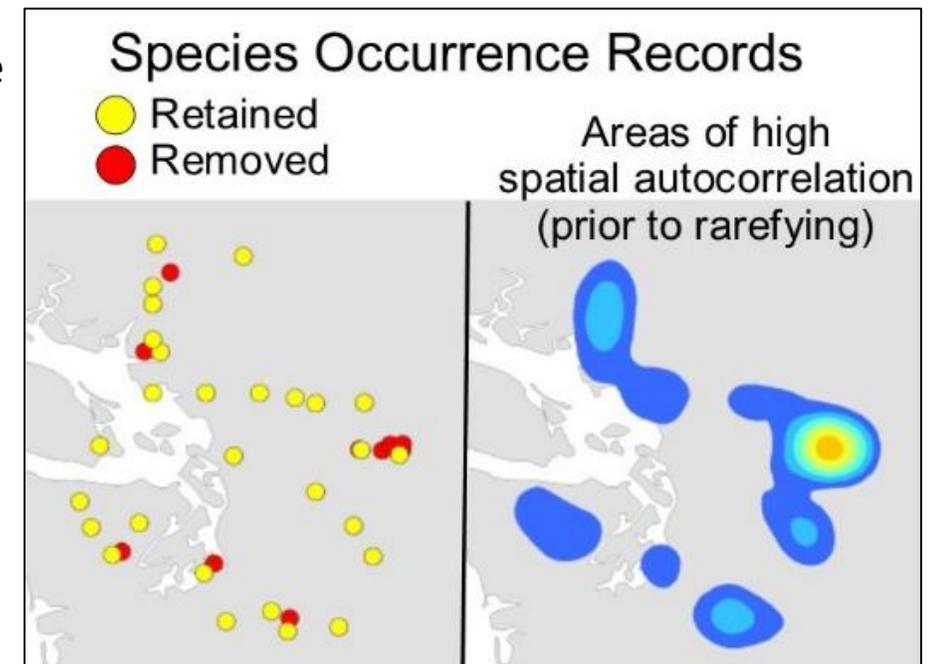
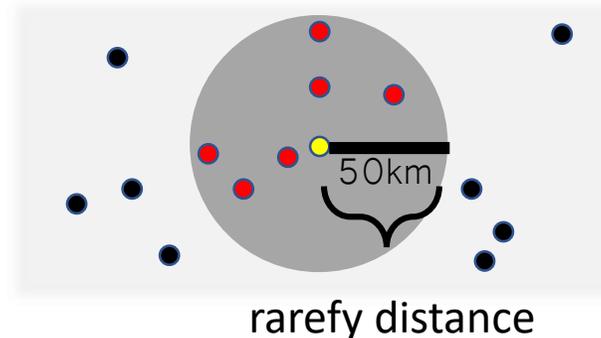
The **rarefy.dist** and **rarefy.units** parameters addresses this issue by spatially filtering locality data by a user input distance, reducing occurrence localities to a single point within the specified Euclidian distance.

**rarefy.dist** Remove occurrences closer than a minimum distance to each other (this function uses the *humboldt.occ.rarefy* function). Values need to be in km[recommended] or decimal degrees. Note: rarefy.dist=0 will remove no occurrences

**rarefy.units** The units of **rarefy.dist** parameter, either "km" for kilometers or "dd" for decimal degrees

## EXAMPLE

rarefy.dist= 50  
rarefy.units="km"



# Accessible Environments

## Input Parameters:

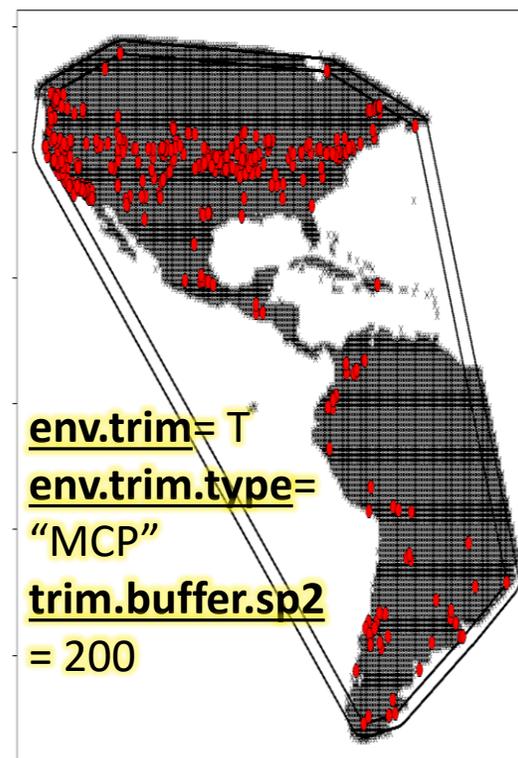
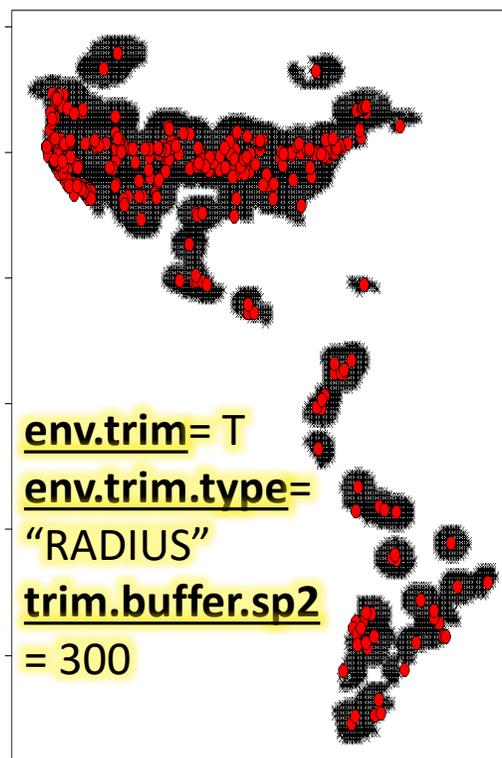
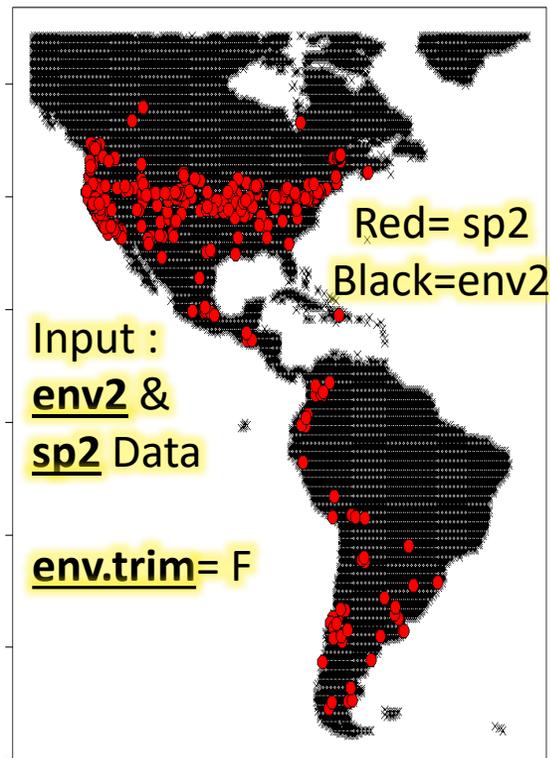
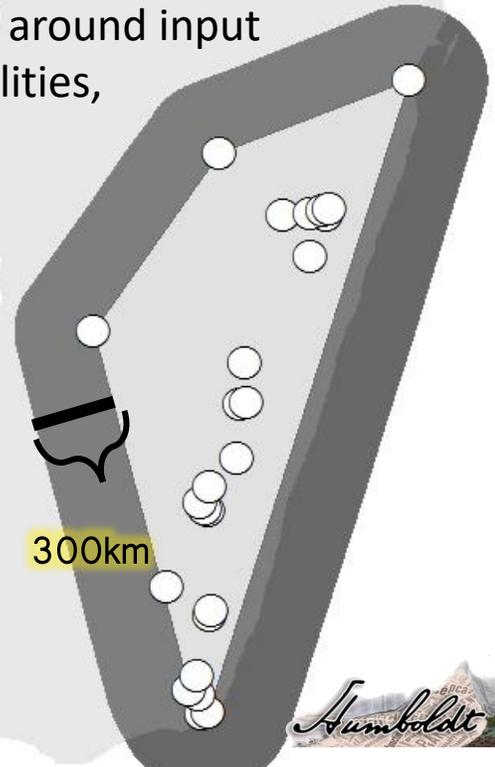
env.trim  
env.trim.type  
trim.buffer.sp1  
trim.buffer.sp2  
trim.mask1  
trim.mask2

**Why trim environments input?** The niche divergence test requires input environments to be reduced to ONLY habitats that are accessible to the species in geography. In this test it is important to clarify the habitat a species currently exists in and the availability of adjacent habitats. Humboldt present three ways to do this: via a buffered minimum convex polygon of input occurrence localities, buffering input occurrence localities, & via a user input mask (an advanced method using an input polygon shapefile WGS1894, input as trim.mask1 or trim.mask2 = "R environment name").

env.trim.type = "MCP"  
trim.buffer.sp1 = 300

These parameters creates a minimum-convex-polygon around input occurrence localities, which is then buffered by the parameter: trim.buffer.sp1

Here the buffer is 300km.



# Analogous and Non-analogous E-space

## Input Parameters:

```
reduce.env
reductype
```

**Why do analogous and non-analogous environments matter?** The niche divergence test requires input environments to be only analogous environments. Thus, environmental space that is **only shared** among habitats is used in the quantitative tests of niche similarity. Thereby differences between species are not solely due to different access to environments that can lead to the false appearance of different niches.

## Step 1: Trim E-space Limits

**reduce.env** The format to trim environments input to analogous space. If `reduce.env=1`, the second input environment (`env2`) will be trim the match the first input (`env1`). If `reduce.env=2`, both input environments trimmed so that extents of both are identical (the lower maximum value observed in `env1` and `env2` and the higher minimum value observed in `env1` and `env2` will be used to trim environmental space for each PC/environmental variable) If `reduce.env=0`, you will skip trimming environmental space

**reducetype** Only used if **reduce.env**= 1 or 2. The **reducetype** parameter specifies the format for how to reduce environmental space ("PCA" or "STANDARD"). If **reducetype**="PCA", the environmental space will be trimmed based on two principal components. If **reducetype**="STANDARD", the environmental space will be trimmed by each included variable specified in **col.env**.

### EXAMPLE

```
reduce.env= 2
reducetype="PCA"
```

## Input Parameters:

reduce.env

reductype

# Step 1: Trim E-space Limits

Below are examples of how the **reduce.env** and **reductype** parameters trim the range of environments. Note this only limits the max and min values of environments to similar ranges. Unique combinations of E-space not shared between environments (non-analogous) likely still exist.

	Post <b>env.trim</b> Environments		<b>reduce.env=0</b> <b>reductype=NA</b>		<b>reduce.env=1</b> <b>reductype=</b> "PCA"		<b>reduce.env=1</b> <b>reductype=</b> "STANDARD"		<b>reduce.env=2</b> <b>reductype=</b> "PCA"		<b>reduce.env=2</b> <b>reductype=</b> "STANDARD"	
	Env1 Input	Env2 Input	Env1 Output	Env2 Output	Env1 Output	Env2 Output	Env1 Output	Env2 Output	Env1 Output	Env2 Output	Env1 Output	Env2 Output
Bio1: max value	291	314	291	314	-	-	291	291	-	-	291	291
Bio1: min value	-226	-177	-226	-177	-	-	-226	-177	-	-	-177	-177
Bio12: max value	2729	8463	2729	8463	-	-	2729	2729	-	-	2729	2729
Bio13: min value	5	0	5	0	-	-	5	0	-	-	5	5
PC1: max value	5.12	3.41	5.12	3.41	5.12	3.41	-	-	3.41	3.41	-	-
PC1: min value	-3.35	-4.17	-3.35	-4.17	-3.35	-3.35	-	-	-3.35	-3.35	-	-
PC2: max value	7.54	2.17	7.54	2.17	7.54	2.17	-	-	2.17	2.17	-	-
PC2: min value	-1.24	-12.1	-1.24	-12.1	-1.24	-1.24	-	-	-1.24	-1.24	-	-

EXAMPLE reduce.env= 2, reductype="STANDARD", col.env=c(3,14), e.var= c(3:21)

# Step 2: Remove Non-Analogous E-space

```

Input Parameters:
reduce.env
reductype
non.analogous.environments

```

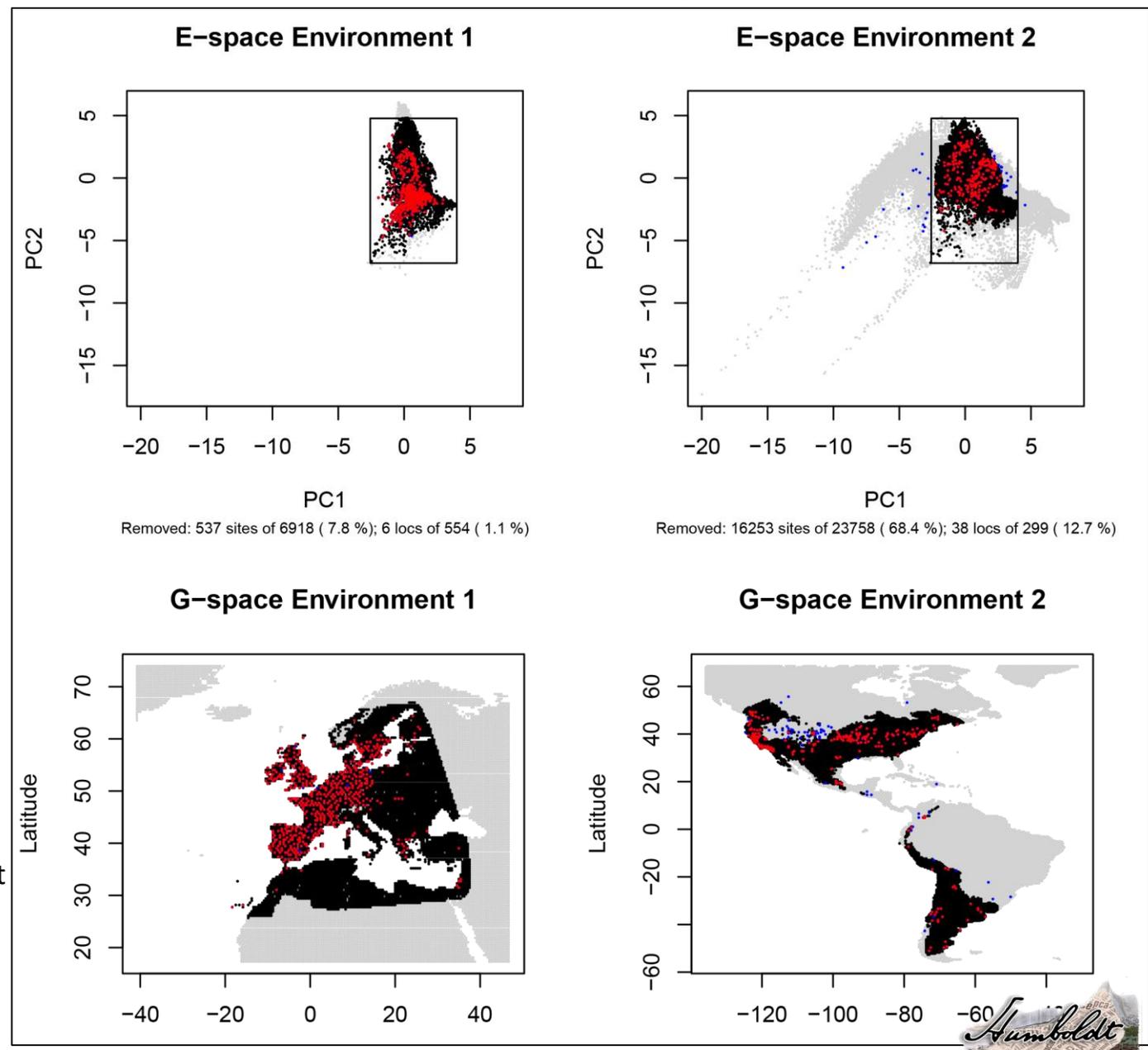
## Removing non-analogous environments

### Step 1: Trim E-space Limits

### Step 2: Remove non-analogous E-space

The second step is to remove E-space that is not identical among both habitats (option only available for **reductype="PCA"**). The **reduce.env** parameter trims habitats to shared max and mins, however often, combinations of PC1 & PC2 values are unique to one habitat and cannot be occupied by species in the other habitat.

- Occurrence localities **retained** after E-space is reduced to analogous environments
- Occurrence localities **removed** after E-space is reduced to analogous environments
- E-space **retained** after after input environments are reduced to analogous environments
- E-space **removed** after input environments are reduced to analogous environments\*\*



# Analogous and Non-analogous E-space

## Input Parameters:

reduce.env

reductype

non.analogous.environments

## Removing non-analogous environments

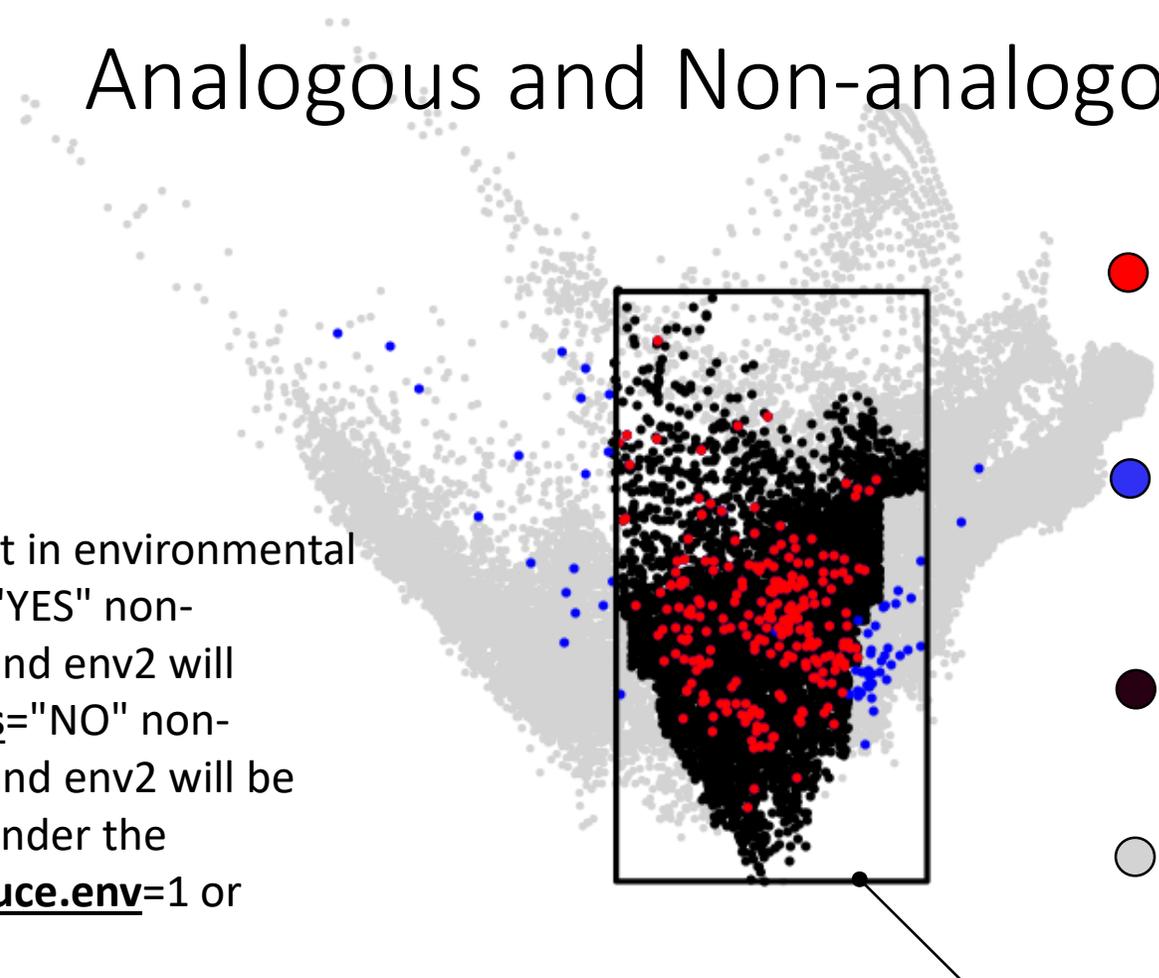
Do you allow non-analogous environment in environmental space? If non.analogous.environments="YES" non-analogous environments between env1 and env2 will be retained. If non.analogous.environments="NO" non-analogous environments between env1 and env2 will be removed. This parameter is only usable under the combinations of reductype="PCA" & reduce.env=1 or reductype="PCA" & reduce.env=2

### EXAMPLE

reduce.env= 2

reducetype="PCA"

non.analogous.environments="YES"



- Occurrence localities **retained** after E-space is reduced to analogous environments
- Occurrence localities **removed** after E-space is reduced to analogous environments
- E-space **retained** after E-space is reduced to analogous environments
- E-space **removed** after E-space is reduced to analogous environments\*\*

Black box depicts shared E-space (based only on maximum and minimum values of each PCs). If non.analogous.environments= "YES", all values within this box will be retained\*\* If non.analogous.environments= "NO", only areas shared between both environments are retained within this box.

\*\*points can also be removed via the env.trim parameter, which trims input environments in geographic space to a buffered minimum convex polygon of the focal species occurrence localities. This reduces the input environmental space to accessible habitats to the focal species.



## Input Parameters:

`nae.window`

## Analogous and Non-analogous E-space

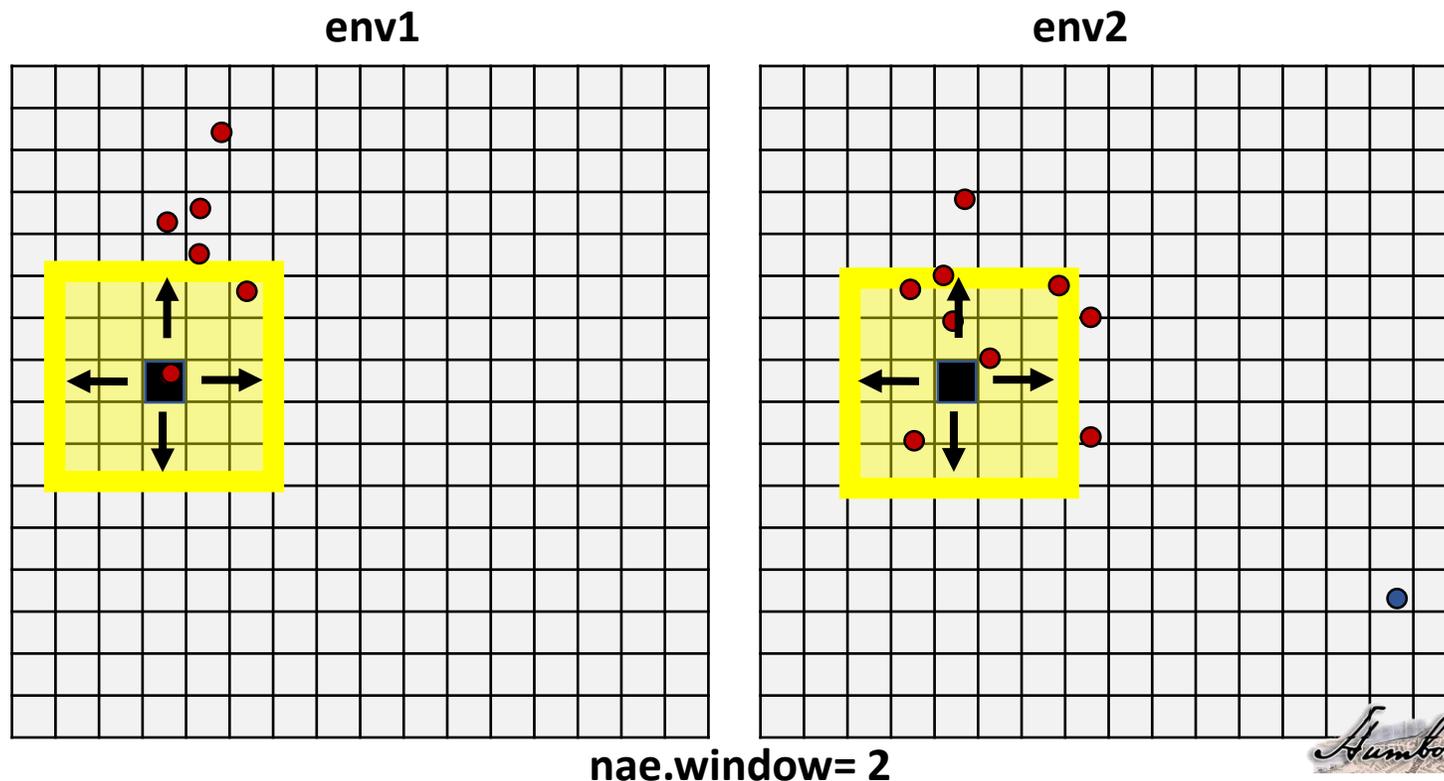
## Fine tuning the removing non-analogous environments

The trimming of E-space is a bit of an art. Do points need to be perfect matches for them to be analogous or do you allow for a little variation? The `nae.window` parameter depicts of how tight or generalized the overlap of data need to be to be considered analogous. The `nae.window` parameter depicts the spatial window from which non-analogous environments will be quantified. The non-analogous environments are characterized by gridding the E-space of `env1` and `env2` into a `R` x `R` grid (e.g. 100 x 100). If `nae.window` parameter= 0, points absent from a cell in one environment will be removed from the other. If `nae.window`>0, values absent from a window (or neighborhood of cells) in one environment will be trimmed from the other. The `nae.window` value characterizes the number of cells to search from the focal cell of environmental space values in the other environment. The larger the `nae.window` value, the fewer non-analogous environments removed. This parameter allows imperfect overlap of environments. If areas of environmental space are a little patchy between environments—but generally present—a larger `nae.window` value will retain more of the patch environments. The default value is a `nae.window`=5

## EXAMPLE

`nae.window= 2`

- Retained
- Removed



Input Parameters:

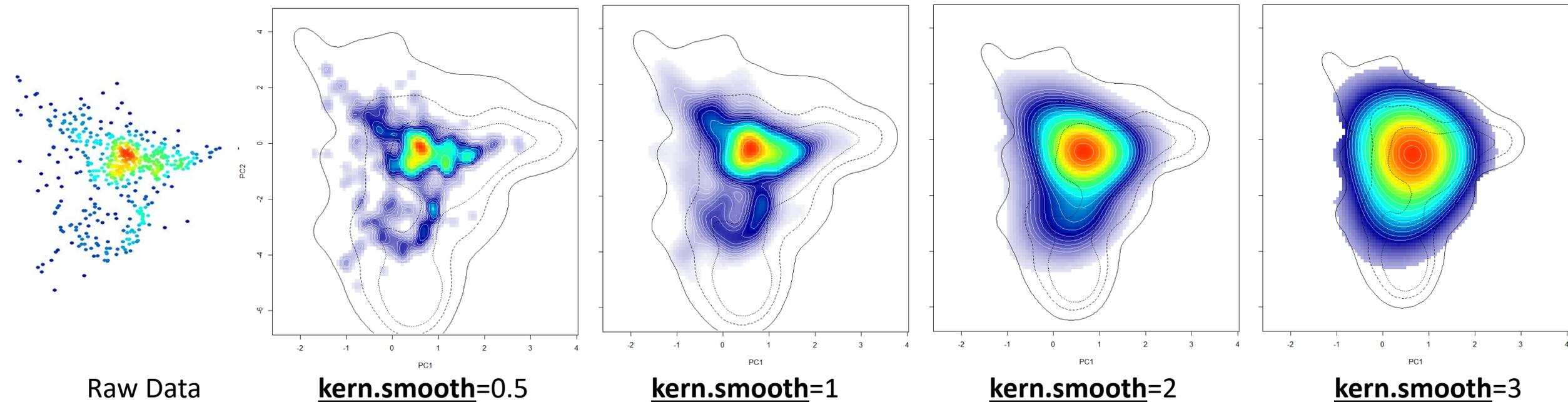
`kern.smooth`

# Quantifying Niches

The point data need to be converted to a continuous surface to quantify the species' niches.

There are three parameters that control: the shape of the surface (**kern.smooth**), the spatial resolution of the E-space (**R**), and the threshold at which low suitability habitats are considered part of the niche (**thresh.space.z**).

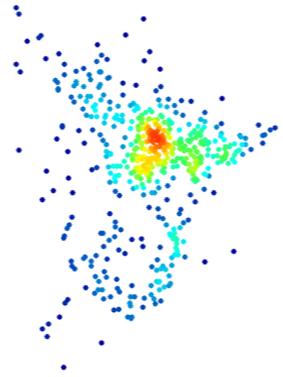
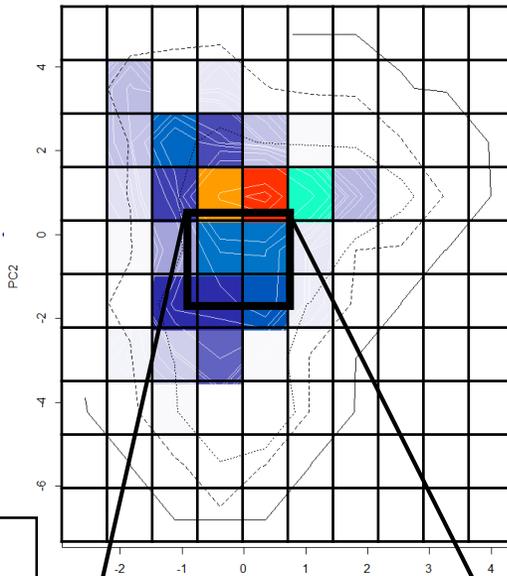
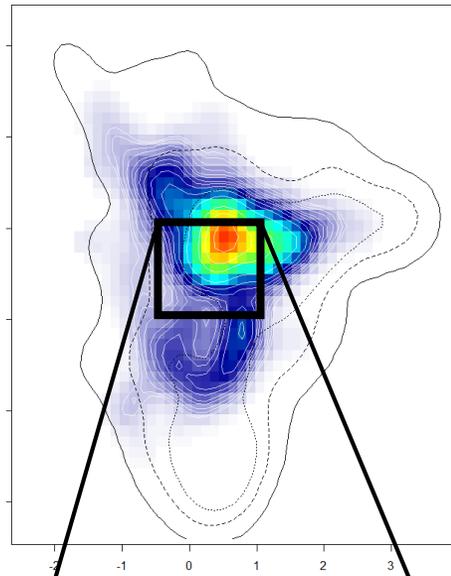
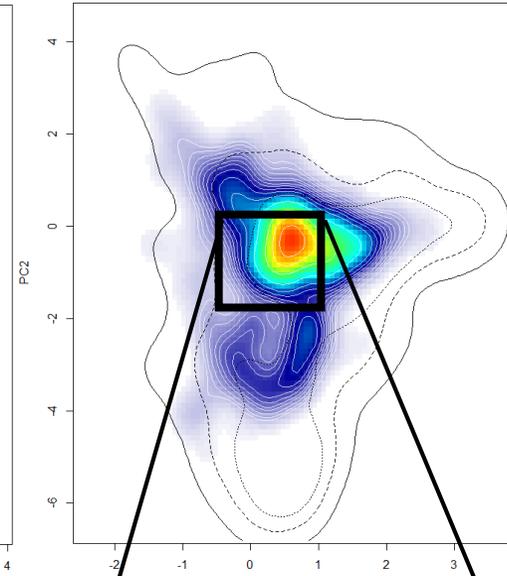
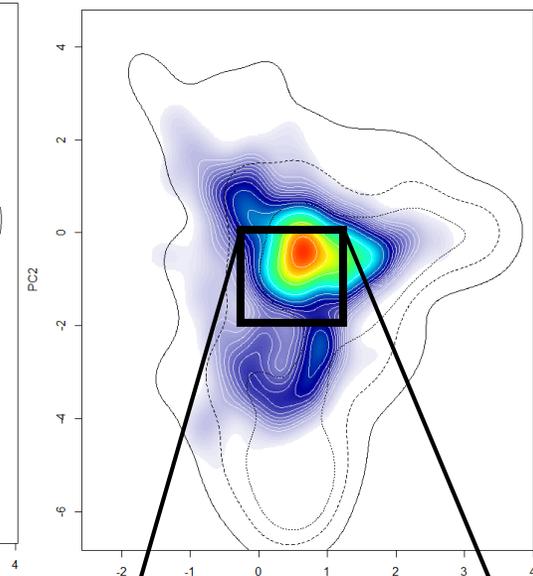
The **kern.smooth** parameter dictates the scale at which kernel smoothing occurs on environmental data. Larger values (i.e. 2) increase scale (making E-space transitions smoother and typically larger) and smaller values (i.e. 0.5) decrease scale (making occupied E-space clusters more dense and irregular). The default value is 1.



# Quantifying Niches

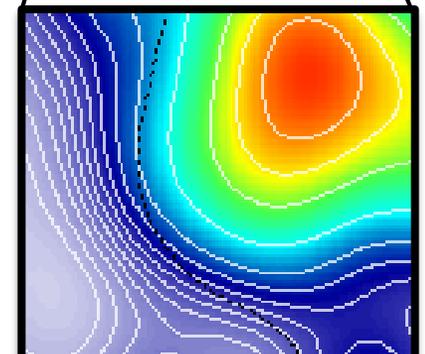
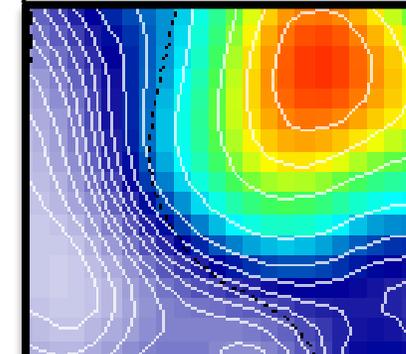
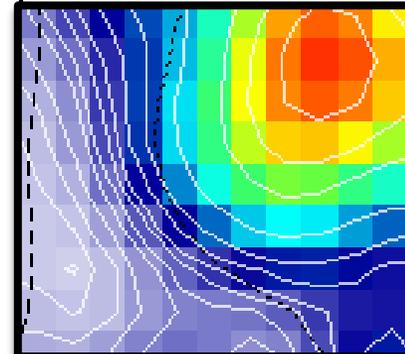
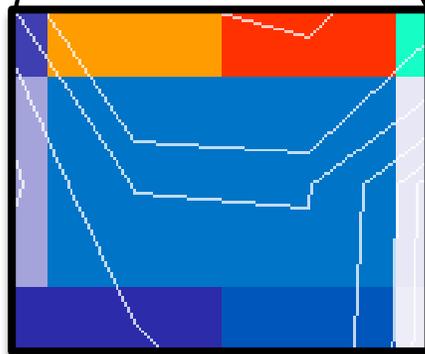
**R** is the resolution of grid in environmental space (RxR). The default value is 100. The larger this value is, the slower the analyses will be. Thus, you want to hit the sweet spot of having high enough, but not too high.

Raw Data

**R=10****R=50****R=100****R=500**

6

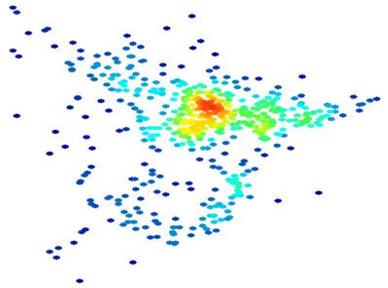
6

**R=6**

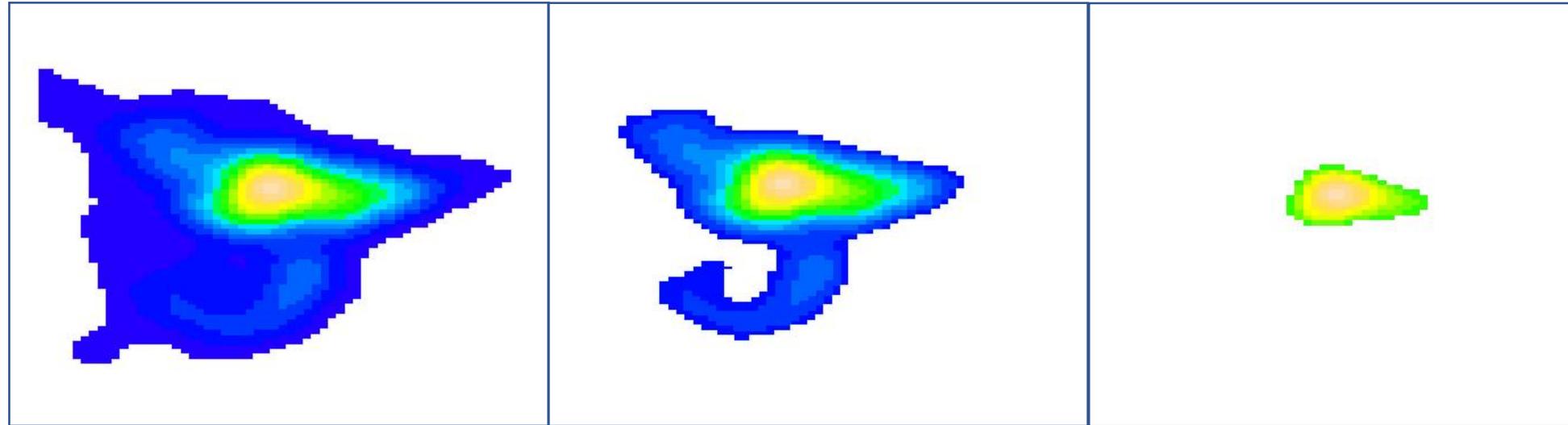
# Quantifying Niches

Input Parameters:  
`thresh.espace.z`

The **thresh.espace.z** parameter is an experimental parameter and controls the level at which values below the kernel density  $z$  values are removed for comparison and quantification of niches. Higher values will increase the value from which the low-density areas are removed from the environmental space. Basically values above this are retained and values below are removed. Input values should range from 0.0001-0. Default=0.001



Raw Data



**thresh.espace.z**=0.001

**thresh.espace.z**=0.01

**thresh.espace.z**=0.1



Input Parameters:

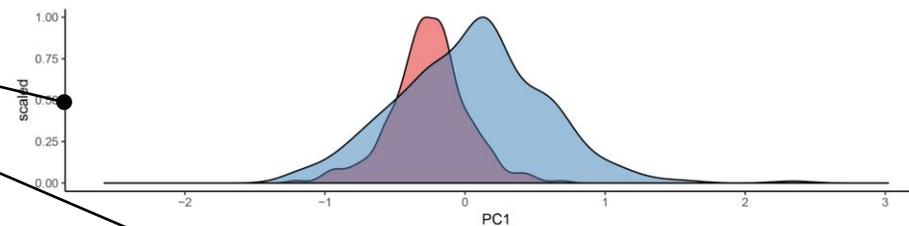
p.overlap

# Plotting Both Niches in a Single Plot

When **p.overlap=T**, both species niches are plotted in a single plot.

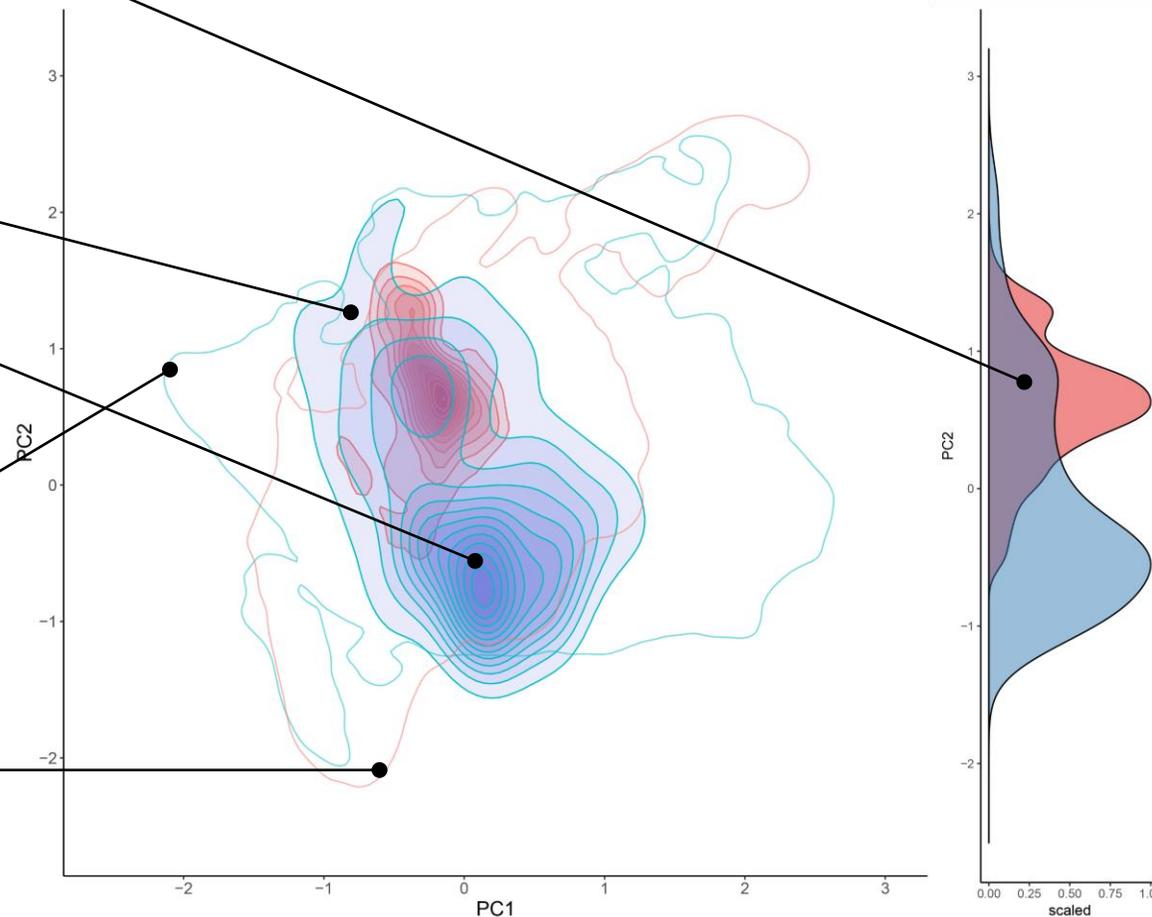
## Histogram Density Plots

For each PC, the density of each species E-space is displayed



## Filled Kernel Density Isoleths (red= sp1 and blue= sp2\*)

Lines representing the kernel density isopleths from 1-100% kernel densities. The number of bins is set at default and adjusts so that the number contour bins best display the data. Here 11 were selected, thus, as follows are the plotted kernel isopleths: 0.01,0.1,0.2,0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, 1



## Empty Kernel Density Isoleths (red= env1 and blue= env2\*)

Lines representing the 1% kernel density isopleths of the environments (not species) depicting outside boundaries of E-space. Sometimes species values erroneously exceed these due to the smooth parameter of kernel densities estimates.

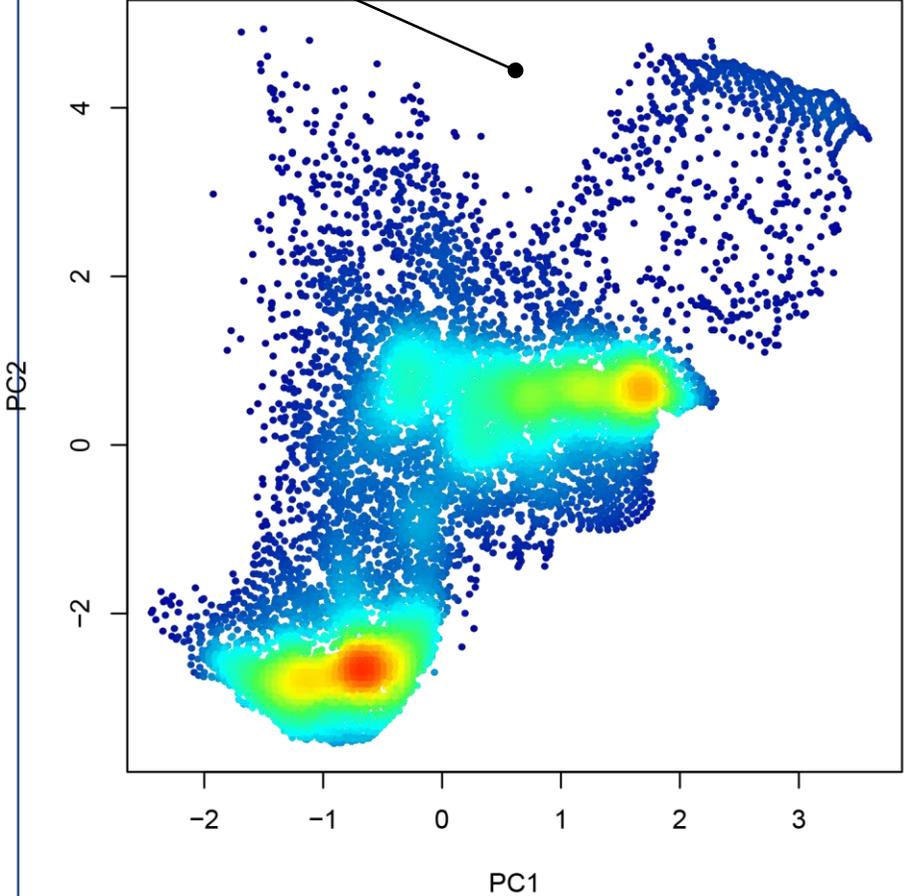
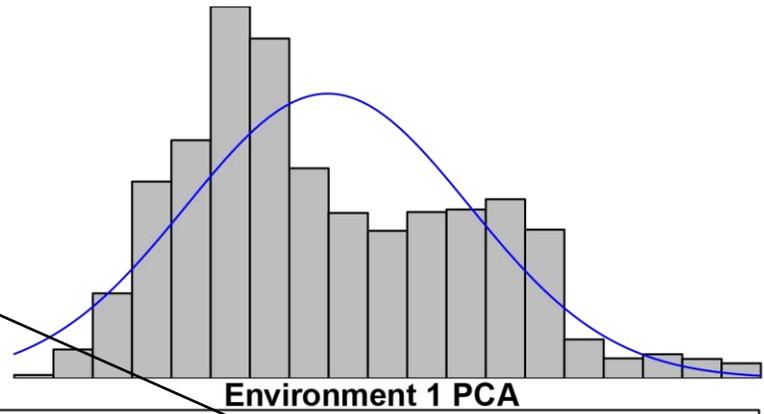
**\*if swap=T, colors will be switched in plots**

Input Parameters:  
p.scatter

# Plotting Raw Data

When **p.scatter**=T, raw analysis data is plotted in a series of plots

**Density Plots of Raw E-space Data**  
A plot of raw data of species' and environment E-space. Cooler colors represent lower and warmer higher densities of E-space.



Input Parameters:

p.boxplot

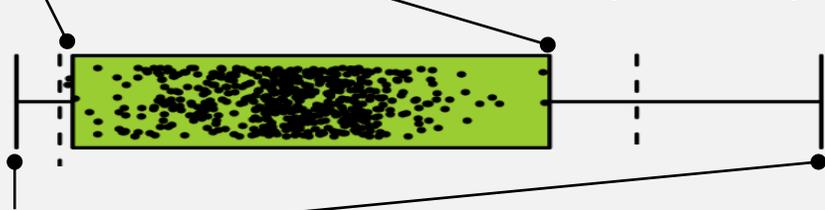
# Plotting E-space Boxplots

When **p.boxplot**=T, both species E-space are plotted

## E-space Box Plots

For each PC, the one dimensional E-space is Displayed.

**Lower and Upper Box:** Minimum and Maximum PC values for species' E-space



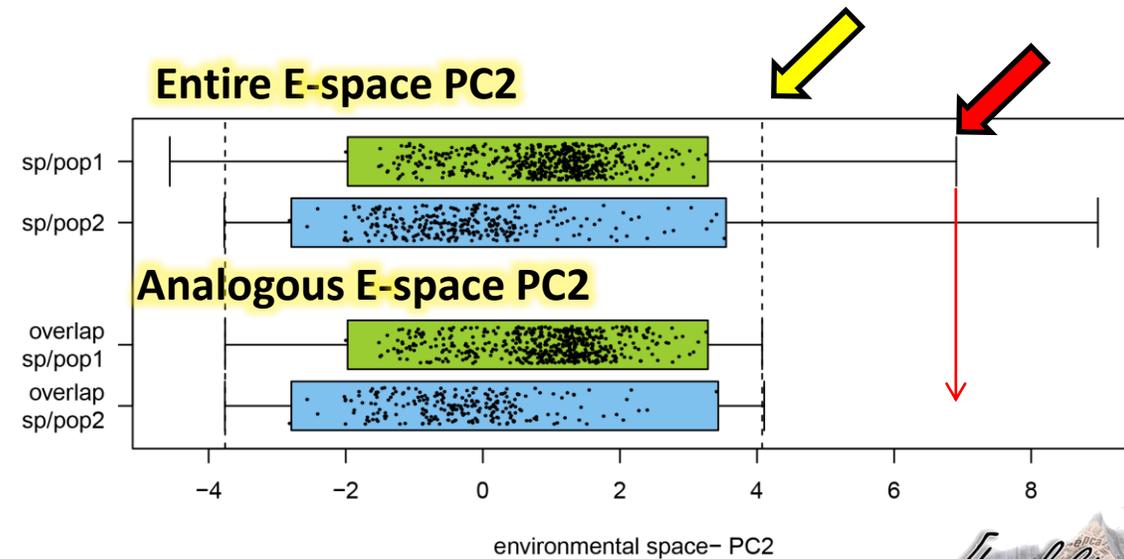
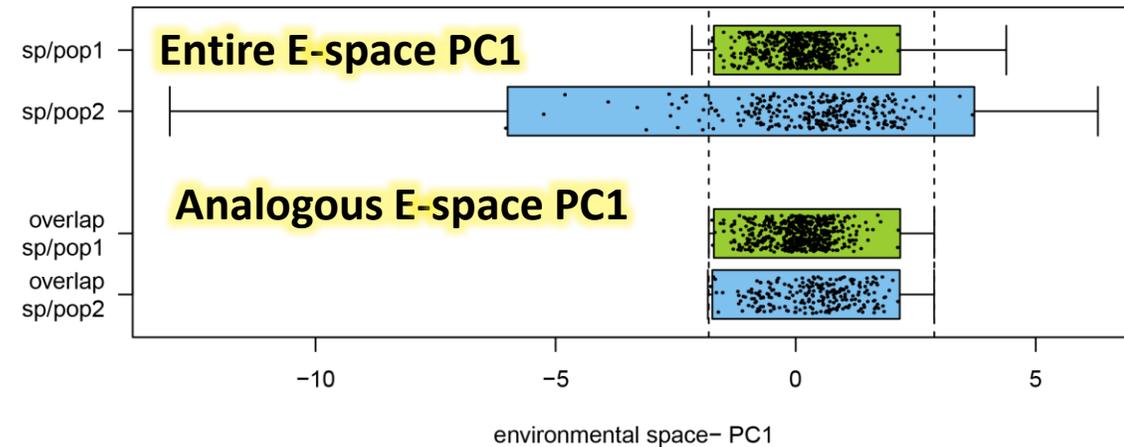
**Lower and Upper Whiskers:** Minimum and Maximum PC values for each environment' E-space

**Dotted lines:** Maximum and Minimum value of shared (analogous) E-space of both environments

**Dots:** PC values for each occurrence record

## E-space truncation in 1 dimension can affect other

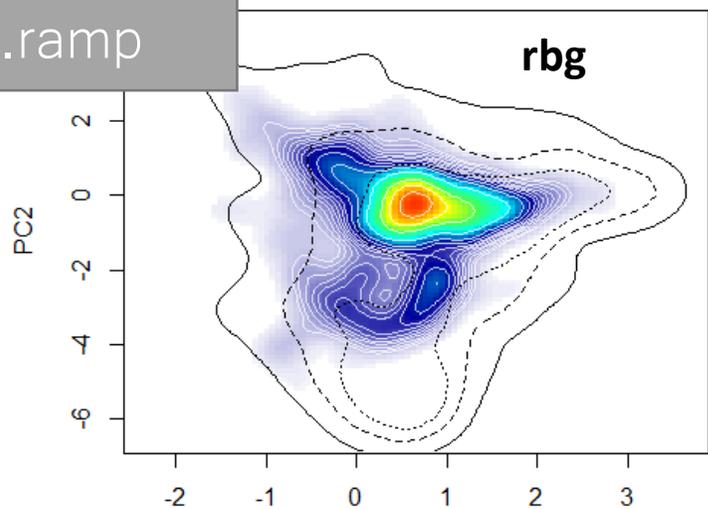
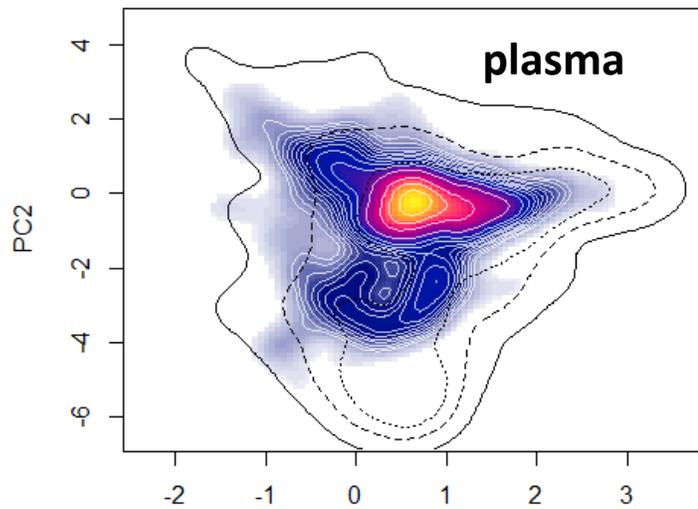
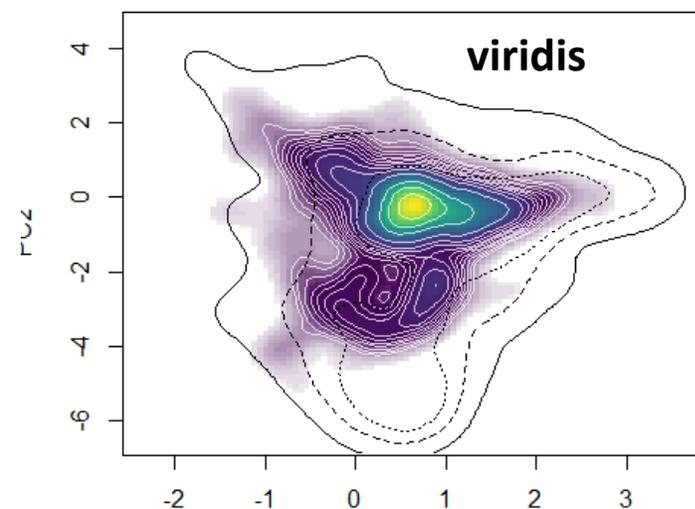
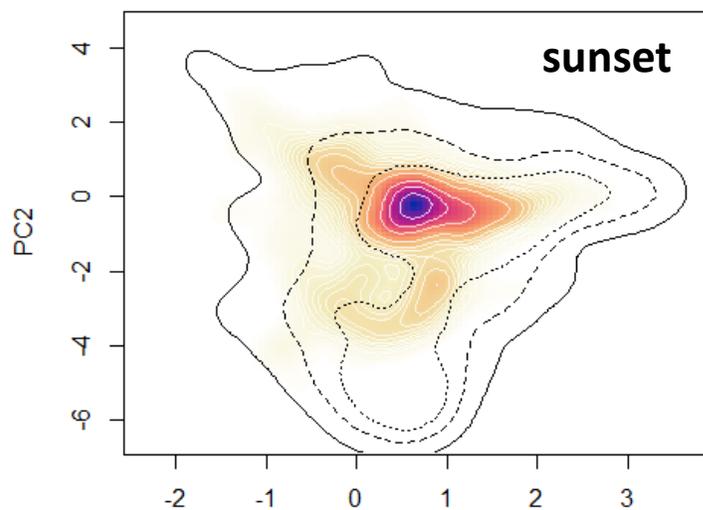
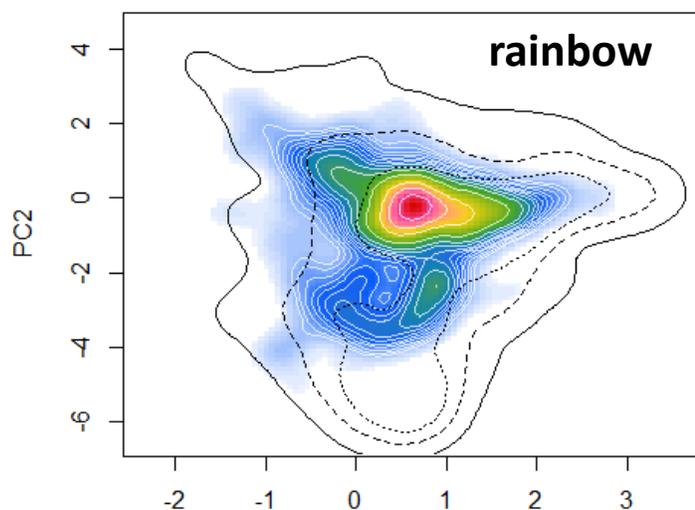
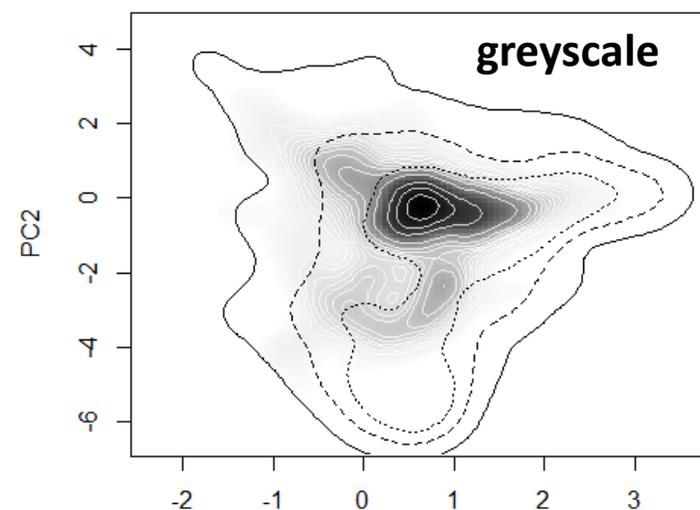
Sometimes the reduction of E-space in PC1, results in truncation of E-space in PC2. In absence of trimming E-space of PC1, the upper shared limit of E-space for PC2 should be ~7 (red arrow). Those values were lost in trimming PC1 and within the remaining analogous E-space, ~4 (yellow arrow) becomes the shared upper limit for PC2.



## Color Schemes

Input Parameters:

color.ramp

color.ramp= 1 (default)color.ramp= 2color.ramp= 3color.ramp= 4color.ramp= 5color.ramp= 6