

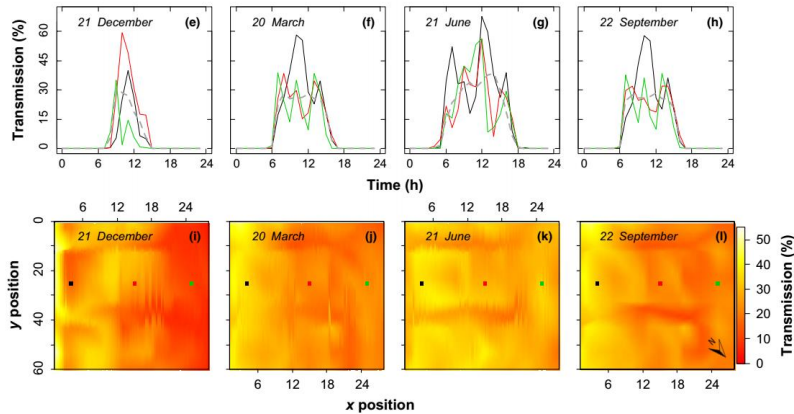
# Exchange event between the European phenomic community and industry

## Progress on Design and analysis of phenotyping experiments across multiple platforms

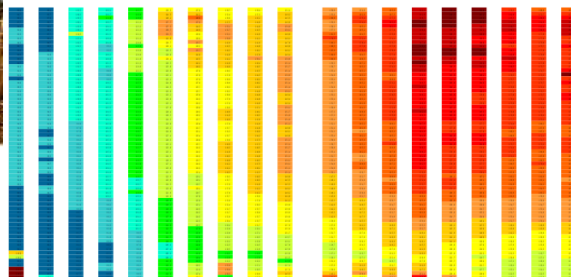
Emilie Millet & Fred van Eeuwijk (WUR)

Design the experiments and analyse the phenomics data with appropriate tools and methods

Platform = variability space and time



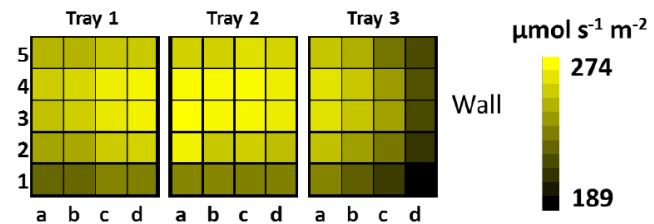
*Cabrera-Bosquet et al. 2016*



*Christian Jeudy, Christophe Salon*

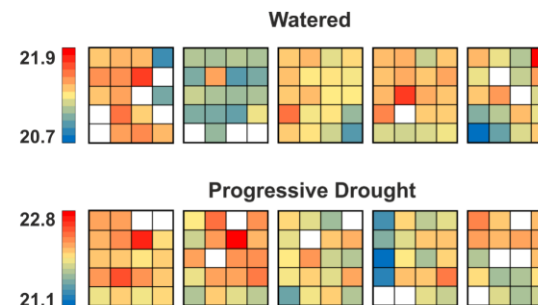


Wall



Front

*Mirko Pavicic, Kristiina Himanen*



Design the experiments and analyse the phenomics data with appropriate tools and methods

Platform = variability space and time

Platform = repeated measurement on thousands of units (plants, organs, etc.)

→ need to consider tools for **designing and analysing**

1. Include and correct for variability = increase precision of phenotype
2. Keep track of decisions = traceability (FAIR) + data re-use
3. Automation of the process = save time
4. New software/tools publicly available = adoption by the community

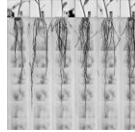
# The analysis of phenomic data : new strategies

## Design experiment



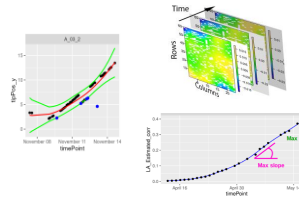
Expe  
map

## Perform experiment & pictures analysis



Raw  
data

## Raw data analysis

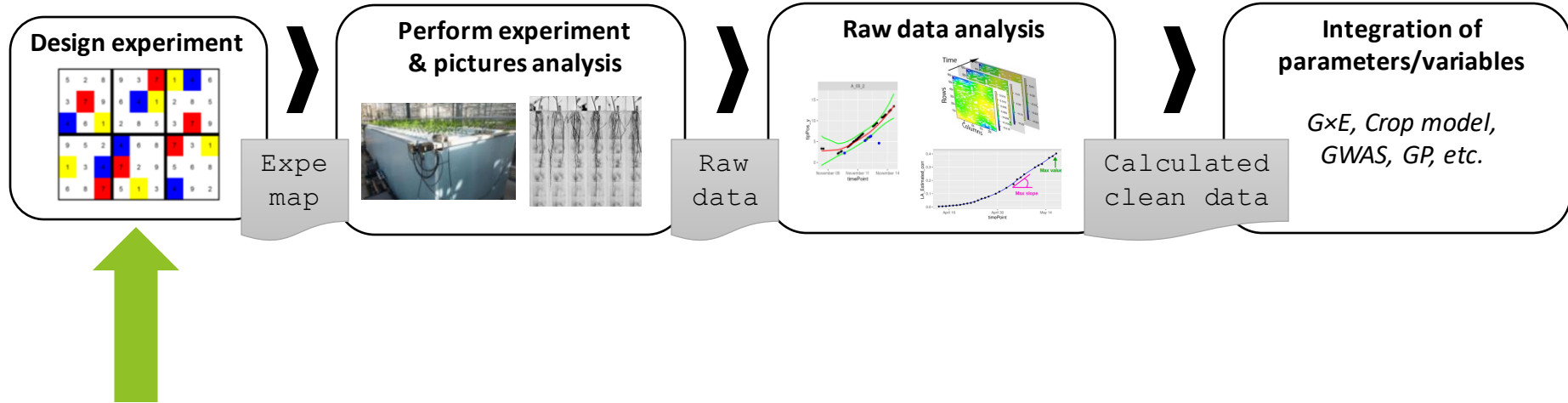


Calculated  
clean data

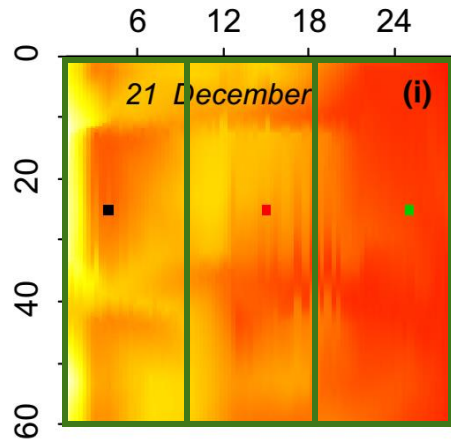
## Integration of parameters/variables

*G×E, Crop model,  
GWAS, GP, etc.*

# The analysis of phenomic data : new strategies



The main goal of the experimental design: random allocation of treatment but not only...

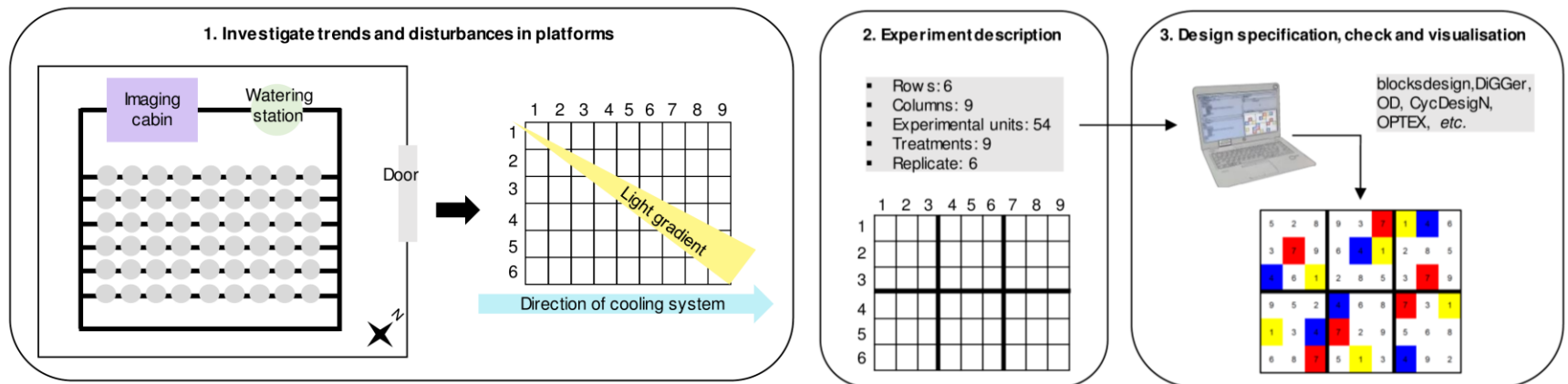


Expected environmental variability in 1 or 2 directions (rows and columns):

→ define **set of experimental units** that are considered to be **internally homogeneous: block**.

→ **restrict the randomization of a two-way layout** to assign the treatments to experimental units in a balanced way: **latinization**

Simple procedure to help choosing the design:



A user-friendly interface of a design generator has been developed (web app)

Available designs:

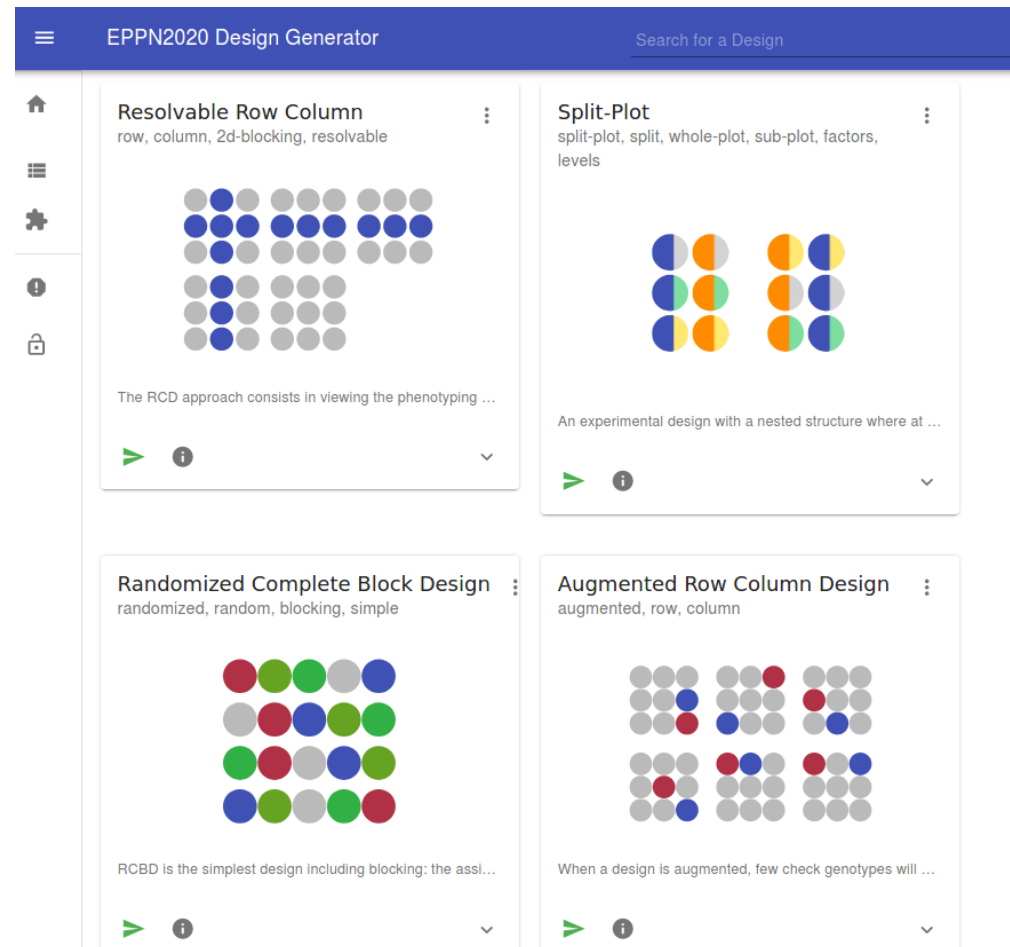
- Resolvable Row-Column (augmented)
- Randomized Complete Block
- Augmented Row-Col
- Split-Plot
- Partially replicated (p-rep)

Create a «facility» based on the layout of his own platform.

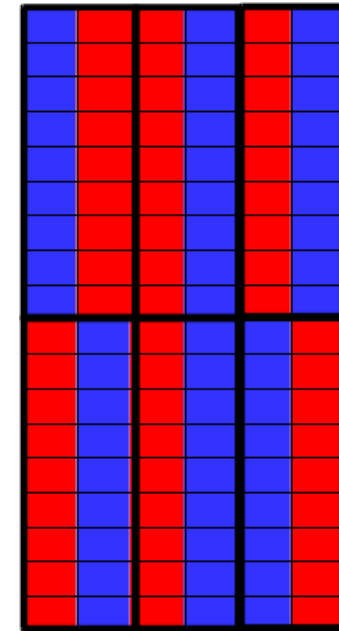
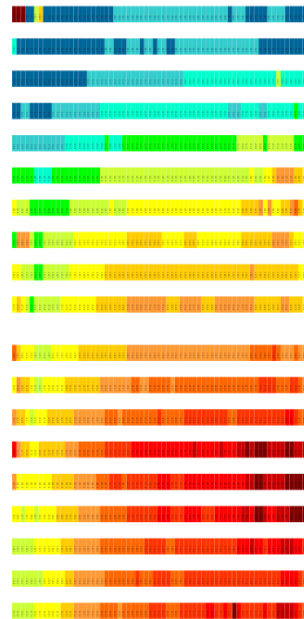
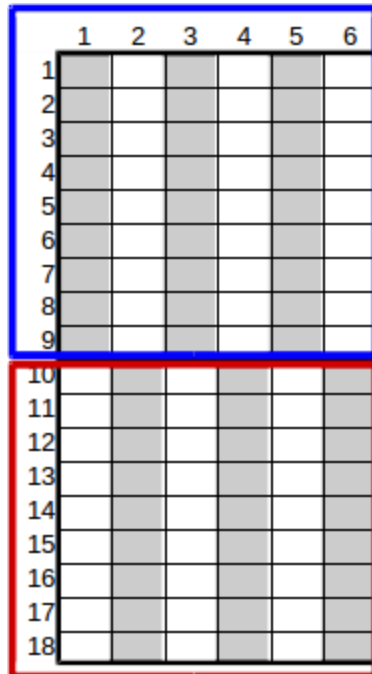
⇒ the user specify the design details/layout (number of genotypes, the number of replicates, etc.)

⇒ gives warnings when mis-specified

⇒ allows visualisation of the 2D map of the experiment



9 genotypes x 2 water level x 6 repetitions  
→ **How to design this?**



Split the treatment in two part of the platform and  
randomize the genotypes  
-> What if there is environmental variability?  
-> Cannot assess the water effect

Split into main block (water) and subplot  
(genotypes) and latinized the genotypes  
-> What software?



## The design generator web app

**VSNI**

Robert Horne,  
Darren Murray

1

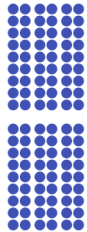
Facility  
Please pick a facility.

Blocks (Replicates)

Design options  
This step is optional.

Results  
Only available once Design is generated.

"SPPU" (rows: 18; columns: 6).



Number of Replicates \*  
6

Rows per Replicate \*  
9

Columns per Replicate \*  
2


GRID HORIZONTAL VERTICAL

BACK NEXT

2

Block Dimensions: (rows: 9; columns: 6). Note: Other blocks not shown.

SUBSTRATE COLOR GENOTYPE



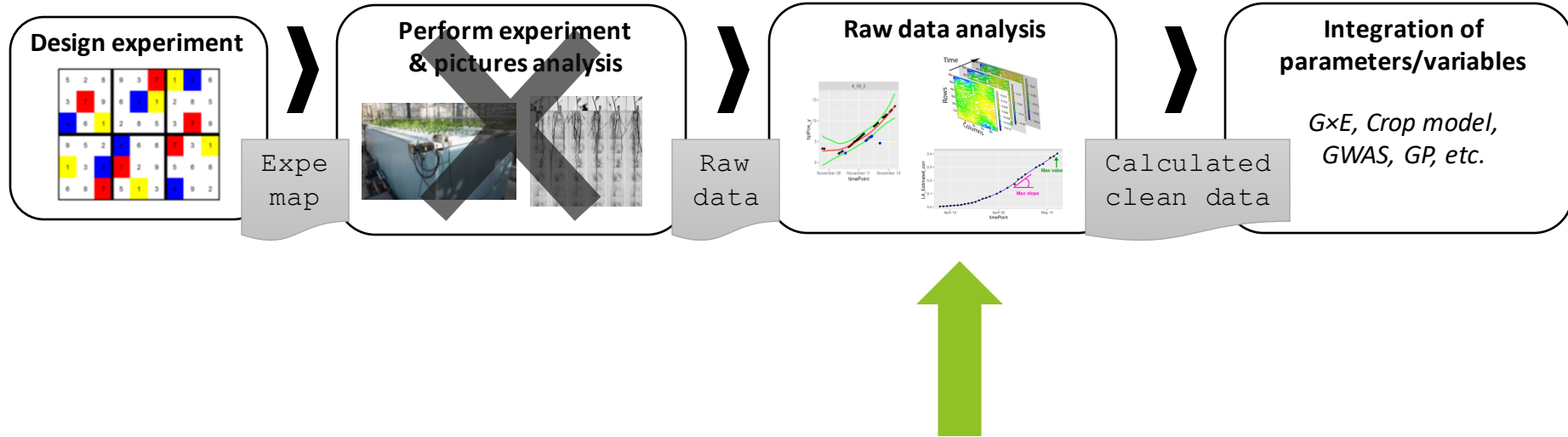
Treatment Factors:

Factor	Levels	Applied where?	Layout	Actions
Substrate	2	Whole Plot	Grid	
Genotype	27	Sub Plot	Grid	

3



# The analysis of phenomic data : new strategies



In classical field trial = multiple environments + one time measurement (yield)

1. correction for spatial variability per environment

2. combining environments

= stage-wise analysis

Or 1 + 2 at once

=> Two methods can be equivalent if weights (error) are carried on from stage 1 to stage 2

In phenomics = multiple experimental series + multiple time measurement (+ multiple scales)

→ **How to model spatial and temporal trends in high-dimensional data?**

- Time + Space in 3D models?
- Longitudinal modelling → spatial correction?
- Spatial correction → time modelling?
- How to detect outliers?

Advantages of stage-wise:

- making diagnostic of individual experiments
- clean individual experiment data
- sometimes required to obtained starting values of one-stage analysis

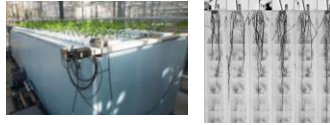
# The analysis of phenomic data : new strategies

## Design experiment



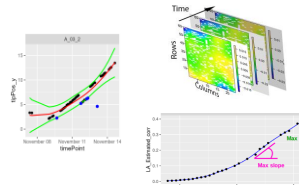
Expe  
map

## Perform experiment & pictures analysis



Raw  
data

## Raw data analysis

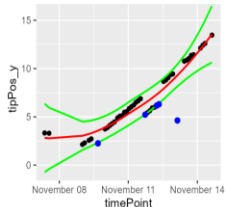


Calculated  
clean data

## Integration of parameters/variables

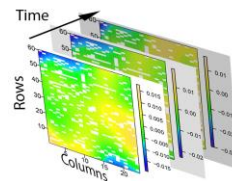
GxE, Crop model,  
GWAS, GP, etc.

## Detect outlying single observation



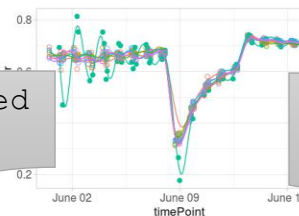
Raw  
clean  
data

## Correct for spatial trend



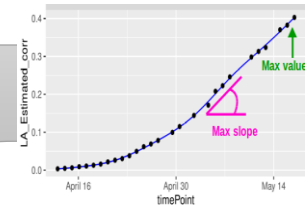
Corrected  
data

## Detect outlying time series



Corrected  
clean data

## Extract parameters from time course



## R package: statgenHTP

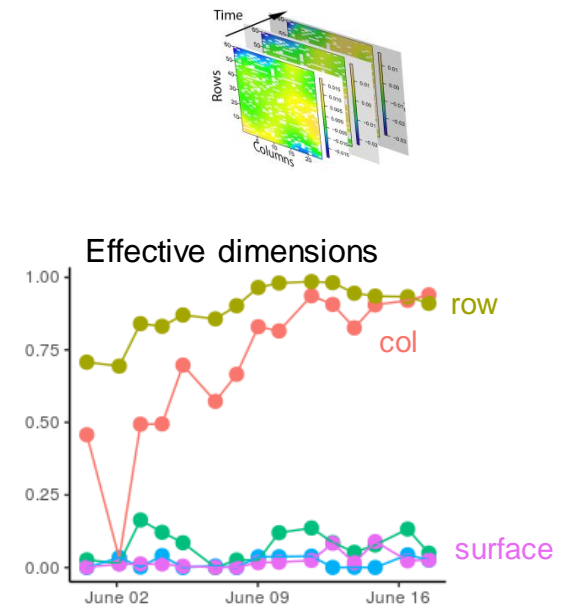
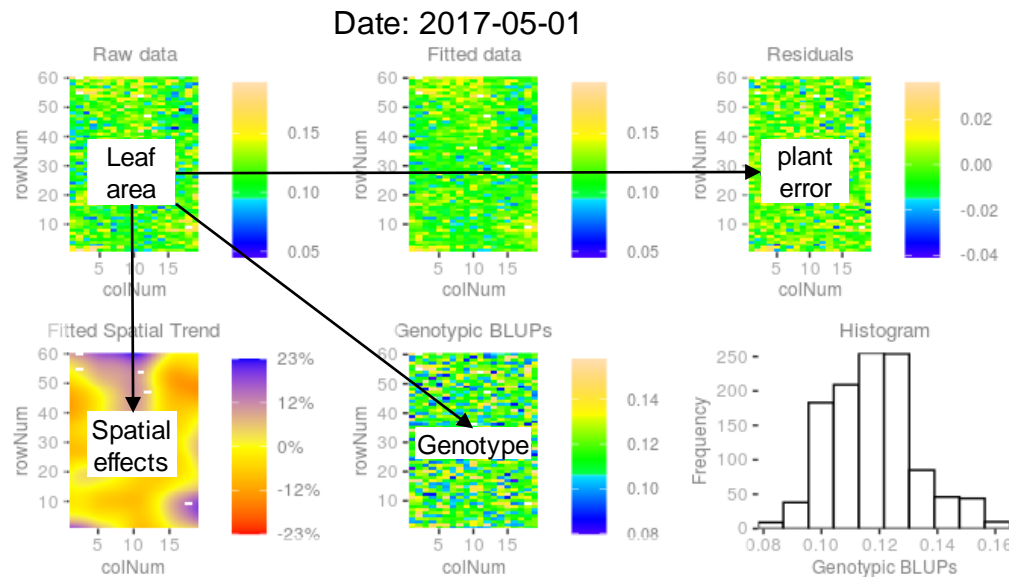


statgenHTP: High Throughput Phenotyping (HTP) Data Analysis

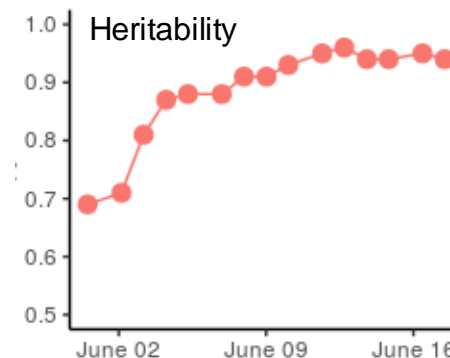
Phenotypic analysis of data coming from high throughput phenotyping (HTP) platforms, including different types of outlier detection, spatial analysis, and parameter estimation. The package is being developed within the EPPN2020 project (<<https://eppn2020.plant-phenotyping.eu/>>). Some functions have been created to be used in conjunction with the R package 'asreml' for the 'ASReml' software, which can be obtained upon purchase from 'VSN' international (<<https://www.vsnl.co.uk/software/asreml/>>).

Version: 1.0.1

Accurately separate the genetic (treatment) effects from the spatial effects at each time point using flexible 2-dimensional P-spline surfaces (SpATS in R):

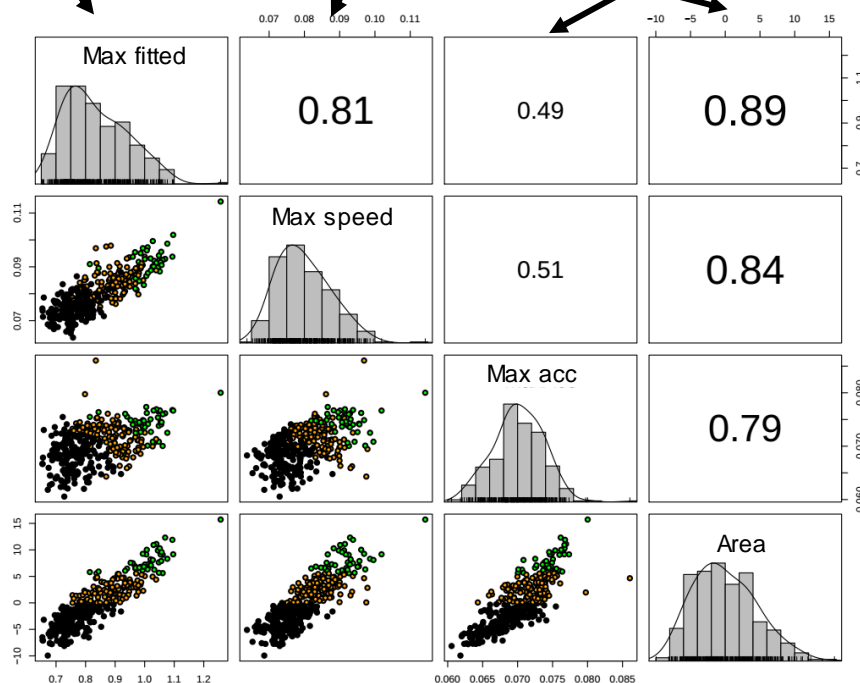
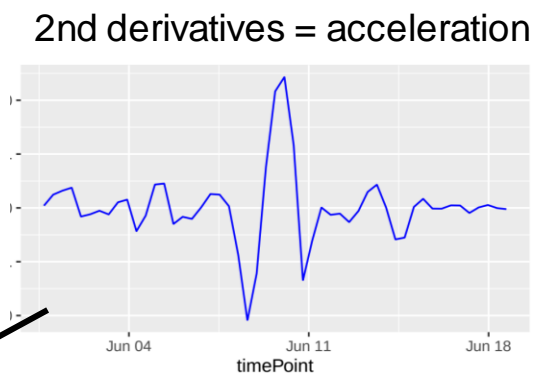
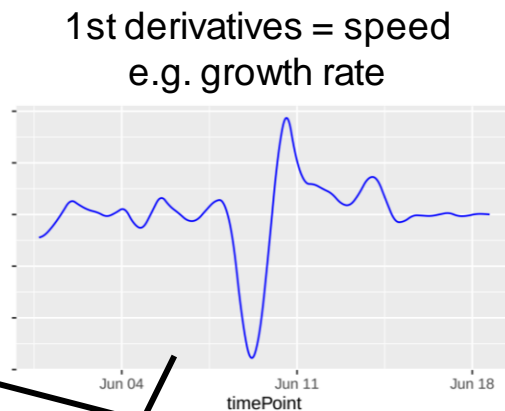
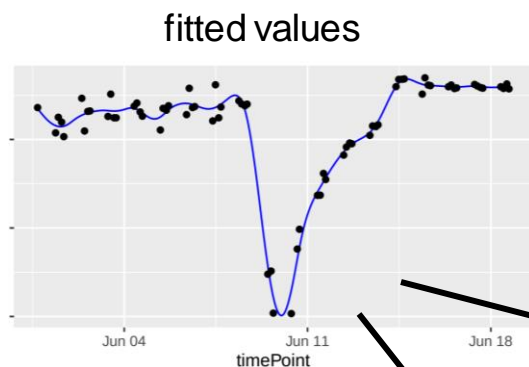


- (1) Correction following the partition by Welham et al. 2004  
→ data at the **experimental/observational unit level**  
or
- (2) Prediction  
→ data at the **treatment (genotype) level**



# Estimation of time-independent variables

Fit splines on corrected data:



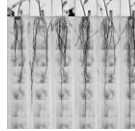
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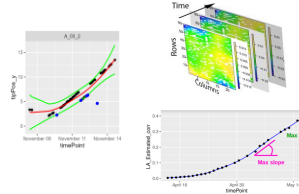
Expe  
map

## Perform experiment & pictures analysis



Raw  
data

## Raw data analysis



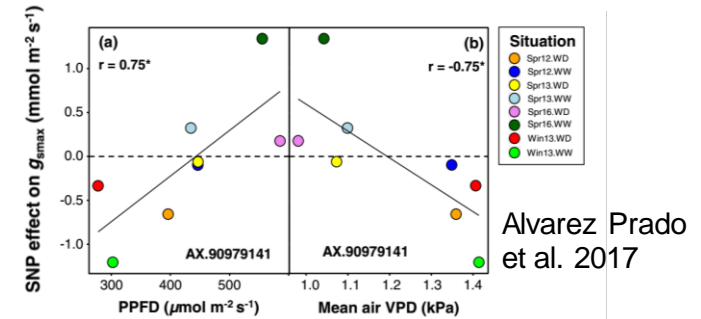
Calculated  
clean data

## Integration of parameters/variables

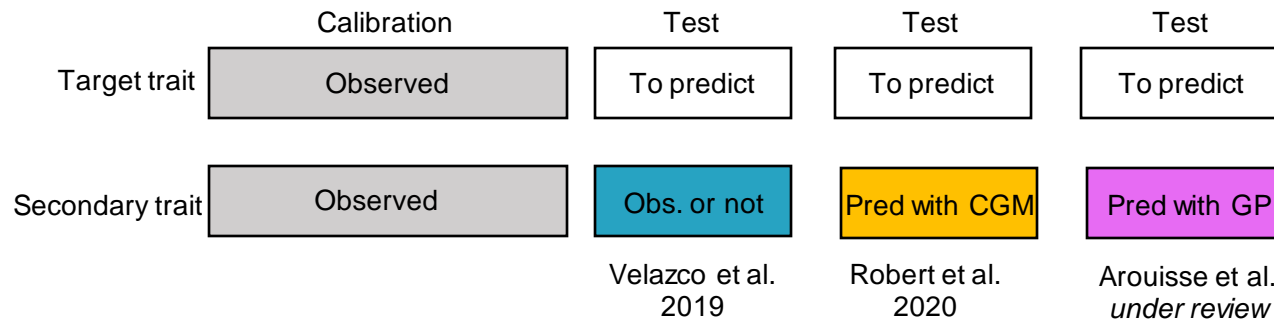
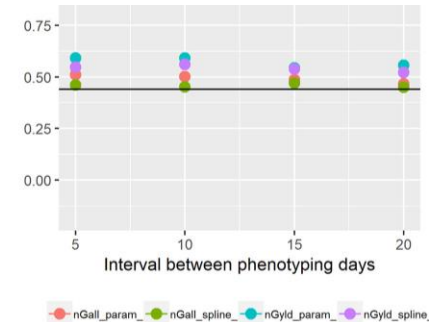
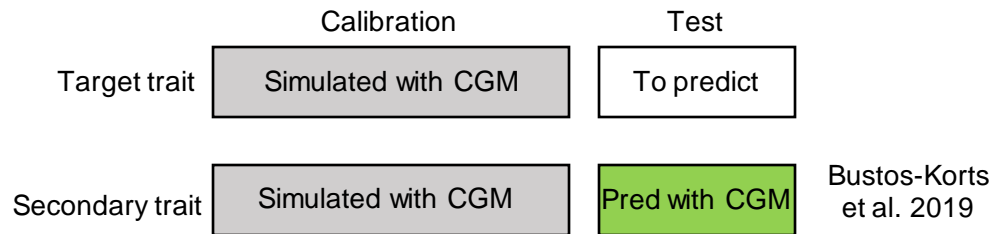
*G×E, Crop model,  
GWAS, GP, etc.*



Multiple experiments =  $G \times E$  and  $QTL \times E$  in platform too  
 → allelic effect vs. environmental cov  
 → multi-trait QTL



Using phenomic traits helps better predict the target trait (yield)



redrawn from Robert et al. 2020



Development of new technology = big data in phenomics

→ Need to adapt existing methods and to streamline the analysis/cleaning process

- Experimental design also in platform → Design generator + procedures
- Strategy for data analysis in multiple stages → statgenHTP R package
- Part of a pipeline of analysis with multiple modules

Collaborative work, multi-disciplines

- Dialog with platform managers/users
- Platform user decision/expertise
- Link with IS and data re-use
- EMPHASIS standards





F van Eeuwijk  
W Kruijer  
M Boer  
B van Rossum  
M Aarts  
R Boesten



R Horne  
D Murray



F Fiorani  
O Muller  
H Porter



C Rodriguez  
D Perez



I Sanchez  
N Hilgert



J Doonan  
G Garzon Martinez



K Olšovská  
M Kovár



F Tardieu  
C Welcker  
L Cabrera-Bosquet



X Draye  
S Passot



T Pridmore  
D Wells  
C Sturrock



R Rincint  
V Allard  
J Legouis



S Dhondt  
V Storme



K Himanen  
M Pavicic