

One-way alpha design

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One-way ANOVA & pairwise comparison post hoc tests in an alpha design.

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```
# (install &) load packages
pacman::p_load(
  agridat,
  conflicted,
  desplot,
  emmeans,
  ggtext,
  lme4,
  lmerTest,
  multcomp,
  multcompView,
  tidyverse)

# handle function conflicts
conflicts_prefer(dplyr::filter)
conflicts_prefer(dplyr::select)
```

```
conflicts_prefer(lmerTest::lmer)
```

1 Data

This example is taken from Chapter “3.8 Analysis of an α -design” of the course material “Mixed models for metric data (3402-451)” by [Prof. Dr. Hans-Peter Piepho](#). It considers data published in John and Williams (1995) from a yield (t/ha) trial laid out as an alpha design. The trial had 24 genotypes (**gen**), 3 complete replicates (**rep**) and 6 incomplete blocks (**block**) within each replicate. The block size was 4.

1.1 Import

The data is available as part of the {agridat} package and needs no further formatting:

```
dat <- as_tibble(agridat::john.alpha)
dat

# A tibble: 72 x 7
   plot rep  block gen  yield  row  col
   <int> <fct> <fct> <fct> <dbl> <int> <int>
1     1  R1    B1    G11  4.12     1     1
2     2  R1    B1    G04  4.45     2     1
3     3  R1    B1    G05  5.88     3     1
4     4  R1    B1    G22  4.58     4     1
5     5  R1    B2    G21  4.65     5     1
6     6  R1    B2    G10  4.17     6     1
7     7  R1    B2    G20  4.01     7     1
8     8  R1    B2    G02  4.34     8     1
9     9  R1    B3    G23  4.23     9     1
10    10  R1    B3    G14  4.76    10     1
# i 62 more rows
```

1.2 Explore

We make use of `dlookr::describe()` to conveniently obtain descriptive summary tables. Here, we get can summarize per block and per cultivar.

```

dat %>%
  group_by(gen) %>%
  dlookr::describe(yield) %>%
  select(2:n, mean, sd) %>%
  arrange(desc(n), desc(mean)) %>%
  print(n = Inf)

```

```

# A tibble: 24 x 4
   gen      n mean    sd
  <fct> <int> <dbl> <dbl>
1 G01      3  5.16 0.534
2 G05      3  5.06 0.841
3 G12      3  4.91 0.641
4 G15      3  4.89 0.207
5 G19      3  4.87 0.398
6 G13      3  4.83 0.619
7 G21      3  4.82 0.503
8 G17      3  4.73 0.379
9 G16      3  4.73 0.502
10 G06      3  4.71 0.464
11 G22      3  4.64 0.432
12 G14      3  4.56 0.186
13 G02      3  4.51 0.574
14 G18      3  4.44 0.587
15 G04      3  4.40 0.0433
16 G10      3  4.39 0.450
17 G11      3  4.38 0.641
18 G08      3  4.32 0.584
19 G24      3  4.14 0.726
20 G23      3  4.14 0.232
21 G07      3  4.13 0.510
22 G20      3  3.78 0.209
23 G09      3  3.61 0.606
24 G03      3  3.34 0.456

```

```

dat %>%
  group_by(rep, block) %>%
  dlookr::describe(yield) %>%
  select(2:n, mean, sd) %>%
  arrange(desc(mean)) %>%
  print(n = Inf)

```

```
# A tibble: 18 x 5
  rep   block     n mean    sd
  <fct> <fct> <int> <dbl> <dbl>
1 R2    B3      4  5.22 0.149
2 R2    B5      4  5.21 0.185
3 R2    B6      4  5.11 0.323
4 R2    B4      4  5.01 0.587
5 R1    B5      4  4.79 0.450
6 R1    B1      4  4.75 0.772
7 R1    B6      4  4.58 0.819
8 R3    B1      4  4.38 0.324
9 R1    B3      4  4.36 0.337
10 R1   B4      4  4.33 0.727
11 R3    B3      4  4.30 0.0710
12 R1    B2      4  4.29 0.273
13 R2    B2      4  4.23 0.504
14 R3    B4      4  4.22 0.375
15 R3    B5      4  4.15 0.398
16 R2    B1      4  4.12 0.411
17 R3    B2      4  3.96 0.631
18 R3    B6      4  3.61 0.542
```

Additionally, we can decide to plot our data:

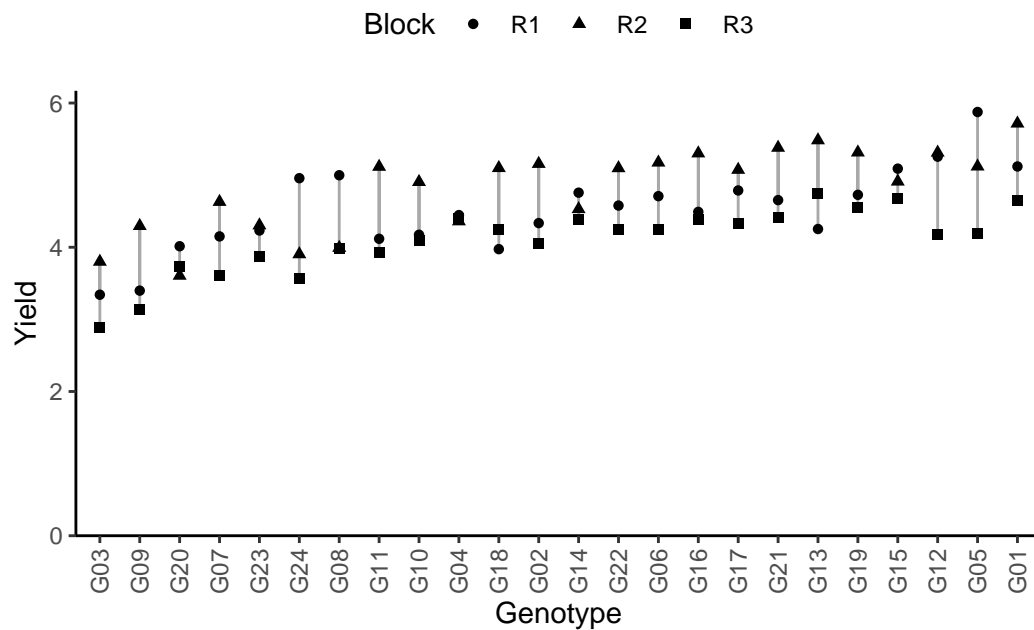
```
# sort genotypes by mean yield
gen_order <- dat %>%
  group_by(gen) %>%
  summarise(mean = mean(yield)) %>%
  arrange(mean) %>%
  pull(gen) %>%
  as.character()

ggplot(data = dat) +
  aes(
    y = yield,
    x = gen,
    shape = rep
  ) +
  geom_line(
    aes(group = gen),
    color = "darkgrey"
  ) +
```

```

geom_point() +
scale_x_discrete(
  name = "Genotype",
  limits = gen_order
) +
scale_y_continuous(
  name = "Yield",
  limits = c(0, NA),
  expand = expansion(mult = c(0, 0.05))
) +
scale_shape_discrete(
  name = "Block"
) +
guides(shape = guide_legend(nrow = 1)) +
theme_classic() +
theme(
  legend.position = "top",
  axis.text.x = element_text(angle = 90, vjust = 0.5)
)

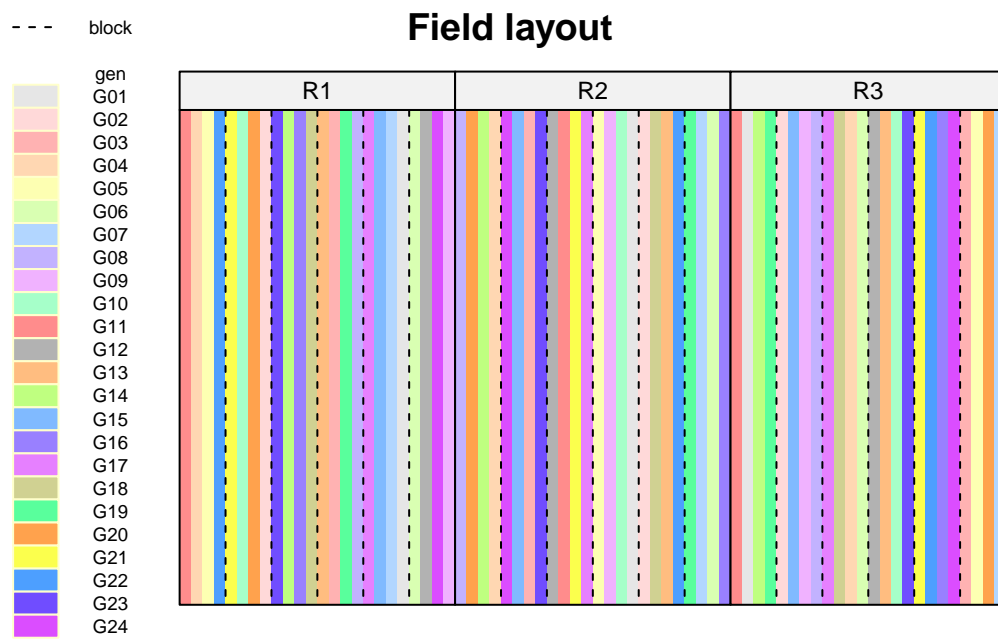
```



Finally, since this is an experiment that was laid with a certain experimental design (= a non-resolvable augmented design) - it makes sense to also get a field plan. This can be done

via `desplot()` from `{desplot}`.

```
desplot(  
  data = dat,  
  flip = TRUE, # row 1 on top, not on bottom  
  form = gen ~ row + col | rep, # fill color per genotype, headers per replicate  
  out1 = block, # lines between incomplete blocks  
  out1.gpar = list(col = "black", lwd = 1, lty = "dashed"), # line type  
  main = "Field layout", # title  
  key.cex = 0.6,  
  layout = c(3, 1) # force all reps drawn in one row  
)
```



2 Modelling

Finally, we can decide to fit a linear model with `yield` as the response variable and (fixed) `gen` and `block` effects. There also needs to be term for the 18 incomplete blocks (*i.e.* `rep:block`) in the model, but it can be taken either as a fixed or a random effect. Since our goal is to compare genotypes, we will determine which of the two models we prefer by comparing the average standard error of a difference (s.e.d.) for the comparisons between adjusted genotype means - the lower the s.e.d. the better.

```
# blocks as fixed (linear model)
mod.fb <- lm(yield ~ gen + rep +
            rep:block,
            data = dat)

mod.fb %>%
  emmeans(pairwise ~ "gen",
          adjust = "tukey") %>%
  pluck("contrasts") %>% # extract diffs
  as_tibble() %>% # format to table
  pull("SE") %>% # extract s.e.d. column
  mean() # get arithmetic mean
```

```
[1] 0.2766288
```

```
# blocks as random (linear mixed model)
mod.rb <- lmer(yield ~ gen + rep +
              (1 | rep:block),
              data = dat)

mod.rb %>%
  emmeans(pairwise ~ "gen",
          adjust = "tukey",
          lmer.df = "kenward-roger") %>%
  pluck("contrasts") %>% # extract diffs
  as_tibble() %>% # format to table
  pull("SE") %>% # extract s.e.d. column
  mean() # get arithmetic mean
```

```
[1] 0.2700388
```

As a result, we find that the model with random block effects has the smaller s.e.d. and is therefore more precise in terms of comparing genotypes.

 Model assumptions met? (click to show)

It would be at this moment (i.e. after fitting the model and before running the ANOVA), that you should check whether the model assumptions are met. Find out more in the [summary article “Model Diagnostics”](#)

3 ANOVA

Based on our model, we can then conduct an ANOVA:

```
ANOVA <- anova(mod.rb, ddf = "Kenward-Roger")
ANOVA

Type III Analysis of Variance Table with Kenward-Roger's method
      Sum Sq Mean Sq NumDF  DenDF F value    Pr(>F)
gen 10.5070  0.45683     23  35.498   5.3628 4.496e-06 ***
rep   1.5703  0.78513      2  11.519   9.2124  0.004078 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Accordingly, the ANOVA's F-test found the cultivar effects to be statistically significant ($p < .001^{***}$).

4 Mean comparison

Besides an ANOVA, one may also want to compare adjusted yield means between cultivars via post hoc tests (t-test, Tukey test etc.).

```
mean_comp <- mod.rb %>%
  emmeans(specs = ~ gen) %>% # adj. mean per genotype
  cld(adjust = "Tukey", Letters = letters) # compact letter display (CLD)

mean_comp
```

gen	emmean	SE	df	lower.CL	upper.CL	.group
G03	3.50	0.199	44.3	2.85	4.15	ab
G09	3.50	0.199	44.3	2.85	4.15	a c
G20	4.04	0.199	44.3	3.39	4.69	abcd
G07	4.11	0.199	44.3	3.46	4.76	abcd
G24	4.15	0.199	44.3	3.50	4.80	abcd
G23	4.25	0.199	44.3	3.60	4.90	abcd
G11	4.28	0.199	44.3	3.63	4.93	abcd
G18	4.36	0.199	44.3	3.71	5.01	abcd
G10	4.37	0.199	44.3	3.72	5.02	abcd
G02	4.48	0.199	44.3	3.83	5.13	abcd
G04	4.49	0.199	44.3	3.84	5.14	abcd

G22	4.53	0.199	44.3	3.88	5.18	abcd
G08	4.53	0.199	44.3	3.88	5.18	cd
G06	4.54	0.199	44.3	3.89	5.19	b d
G17	4.60	0.199	44.3	3.95	5.25	d
G16	4.73	0.199	44.3	4.08	5.38	d
G12	4.76	0.199	44.3	4.11	5.40	d
G13	4.76	0.199	44.3	4.11	5.41	d
G14	4.78	0.199	44.3	4.13	5.42	d
G21	4.80	0.199	44.3	4.15	5.44	d
G19	4.84	0.199	44.3	4.19	5.49	d
G15	4.97	0.199	44.3	4.32	5.62	d
G05	5.04	0.199	44.3	4.39	5.69	d
G01	5.11	0.199	44.3	4.46	5.76	d

Results are averaged over the levels of: rep
 Degrees-of-freedom method: kenward-roger
 Confidence level used: 0.95
 Conf-level adjustment: sidak method for 24 estimates
 P value adjustment: tukey method for comparing a family of 24 estimates
 significance level used: alpha = 0.05
 NOTE: If two or more means share the same grouping symbol,
 then we cannot show them to be different.
 But we also did not show them to be the same.

Note that if you would like to see the underlying individual contrasts/differences between adjusted means, simply add `details = TRUE` to the `cld()` statement. Furthermore, check out the [Summary Article “Compact Letter Display”](#).

Finally, we can create a plot that displays both the raw data and the results, *i.e.* the comparisons of the adjusted means that are based on the linear model.

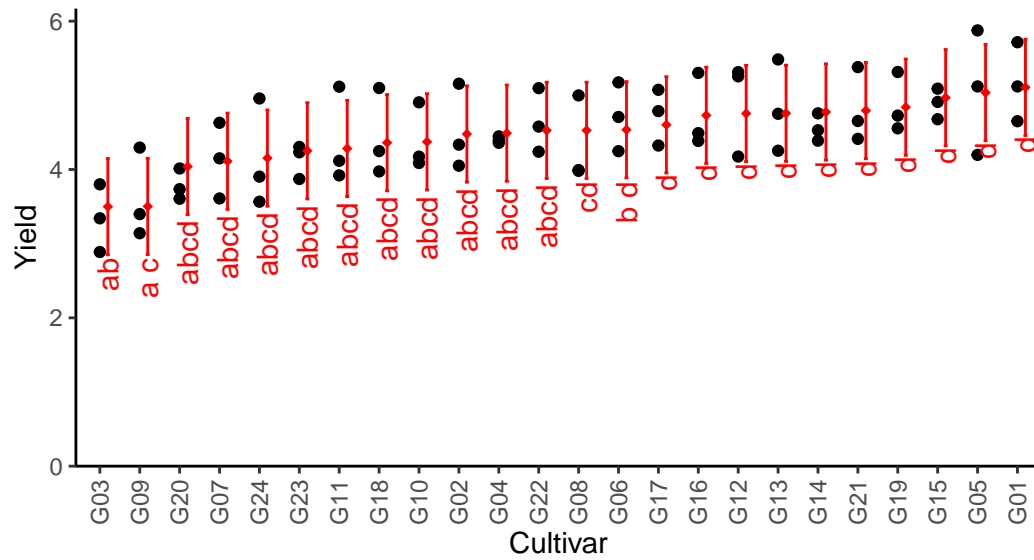
```
# reorder genotype factor levels according to adjusted mean
my_caption <- "Black dots represent raw data. Red diamonds and error bars represent adjusted means"

ggplot() +
  # green/red dots representing the raw data
  geom_point(
    data = dat,
    aes(y = yield, x = gen)
  ) +
  # red diamonds representing the adjusted means
  geom_point(
```

```

data = mean_comp,
aes(y = emmean, x = gen),
shape = 18,
color = "red",
position = position_nudge(x = 0.2)
) +
# red error bars representing the confidence limits of the adjusted means
geom_errorbar(
  data = mean_comp,
  aes(ymin = lower.CL, ymax = upper.CL, x = gen),
  color = "red",
  width = 0.1,
  position = position_nudge(x = 0.2)
) +
# red letters
geom_text(
  data = mean_comp,
  aes(y = lower.CL, x = gen, label = str_trim(.group)),
  color = "red",
  angle = 90,
  hjust = 1.1,
  position = position_nudge(x = 0.2)
) +
scale_x_discrete(
  name = "Cultivar",
  limits = as.character(mean_comp$gen)
) +
scale_y_continuous(
  name = "Yield",
  limits = c(0, NA),
  expand = expansion(mult = c(0, 0.05))
) +
labs(caption = my_caption) +
theme_classic() +
theme(plot.caption = element_textbox_simple(margin = margin(t = 5)),
      plot.caption.position = "plot",
      axis.text.x = element_text(angle = 90, vjust = 0.5))

```



Black dots represent raw data. Red diamonds and error bars represent adjusted means with 95% confidence limits per cultivar. Means followed by a common letter are not significantly different according to the Tukey-test.

John, J. A., and E. R. Williams. 1995. "Cyclic and Computer Generated Designs." *Biometrical Journal* 38 (7): 778–78. <https://doi.org/10.1002/bimj.4710380703>.