

# The role of wetting vs. drying on SOM destabilization

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# Objectives

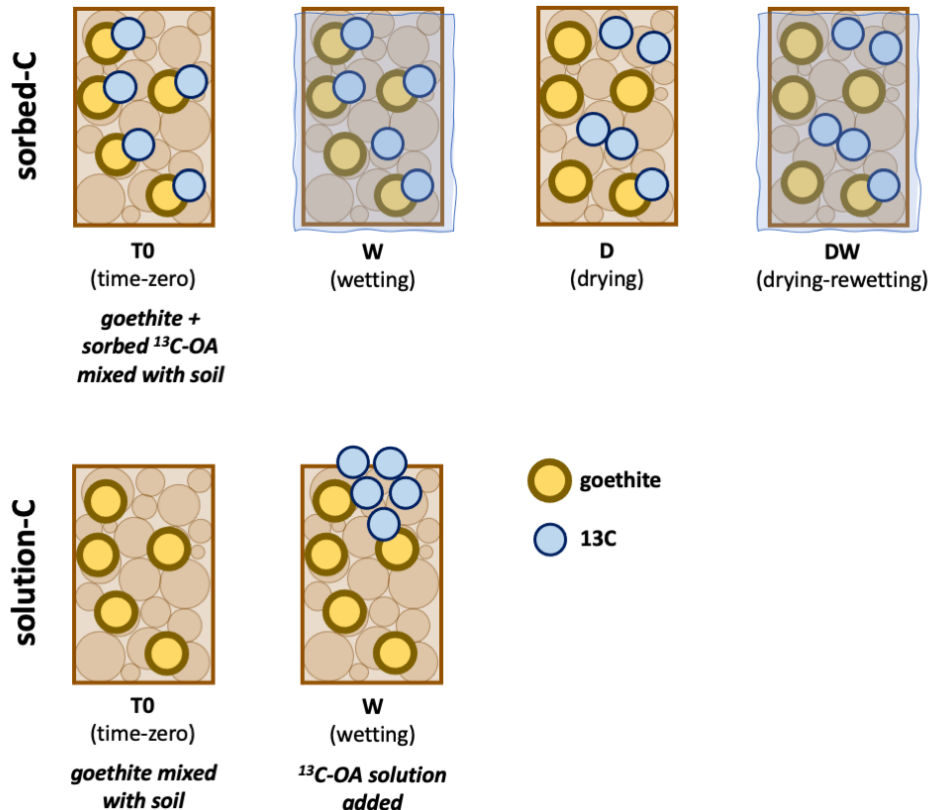
To partition biochemical and physicochemical protection/destabilization mechanisms for soil C by their sensitivities to wetting and drying.

## Hypotheses

1. drying and wetting processes destabilize C via different mechanisms: desorption and cell lysis (drying), vs. increased hydrologic connectivity and improved spatial access to substrate (wetting).
  - under conditions of drying, desorption of C from clay-sorbed substrate will result in [lag and release] of desorption/respiration. vs. cell lysis means we will have a pulse of more  $^{12}\text{C}$  vs. the newly added  $^{13}\text{C}$
2. drying will have a stronger effect on soil C destabilization, compared to wetting
  - drying will have a higher ratio of  $^{13}/^{12}\text{C}$  because the respiration will be more destabilization vs. wetting/access to native soil C.

# Experiment

60 g field-moist Palouse soil (~58 g oven-dry equivalent) in pint size Mason jars  
labelled substrate ( $^{13}\text{C}$ -oxalic acid, 18.66 % enriched) was added to the soil, in adsorbed or solution form.

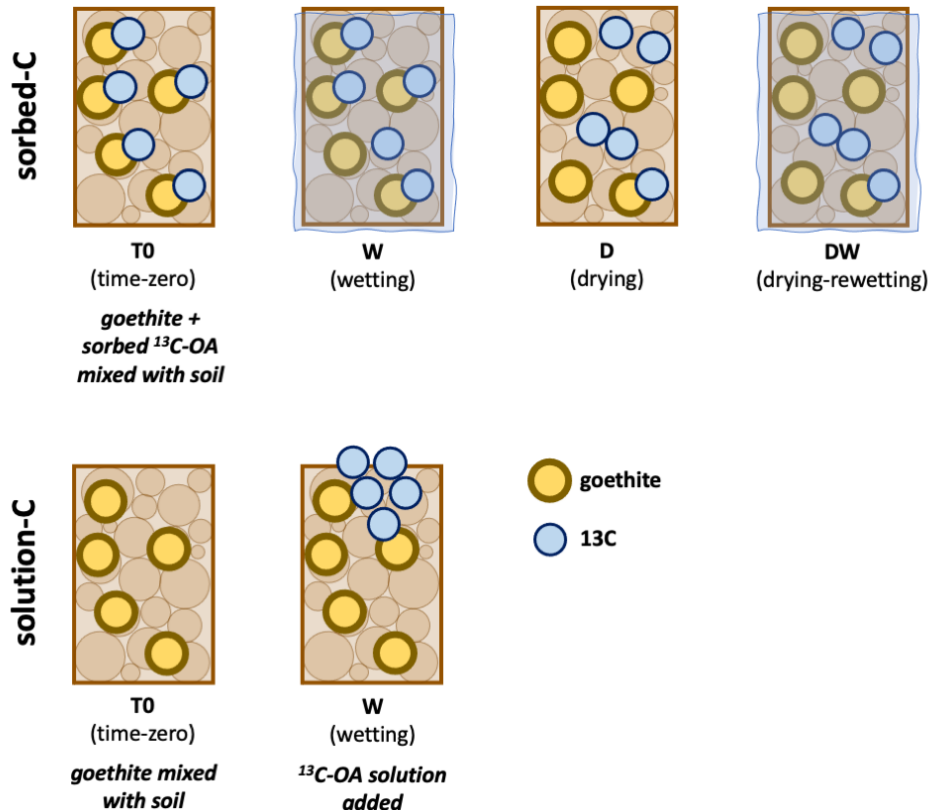


## adsorbed-C:

- 20 mL of labelled OA solution mixed with 5 g goethite; rinsed 3x; labelled goethite resuspended in 5 mL water; added as paste to soil units
- The soil was subjected to one of the following treatments, after which the jars were sealed:
  1. time-zero: jars sealed as soon as goethite was added
  2. wetting: 20 mL water added
  3. drying: soil allowed to air-dry until constant weight
  4. drying-rewetting: soil allowed to dry, then rewet with 20 mL water
- The jars were sealed for 48 hours.

# Experiment

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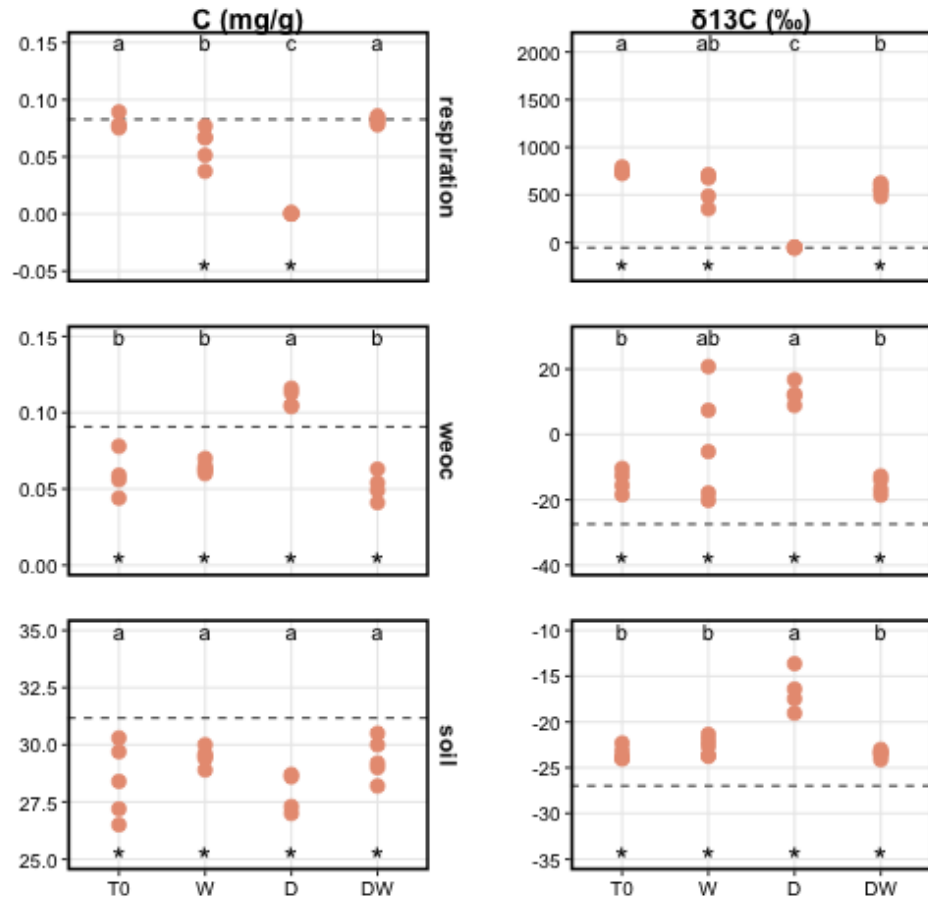
## solution-C:

- a parallel experiment was performed as a control for priming effects of the added OA
- 5 g goethite mixed with soil; 16 mL of labelled OA solution added to the soil-goethite mixture
- The soil was subjected to one of the following treatments, after which the jars were sealed:
  1. time-zero: goethite mixed with soil
  2. wetting: 16 mL  $^{13}\text{C}$ -OA solution added
- The jars were sealed for 48 hours.

# Analyses

1. **respiration:** headspace samples were collected after the 48-hr incubation. analyzed for CO<sub>2</sub> and <sup>13</sup>C-CO<sub>2</sub>
  - LI-7810 (LI-COR), LGR
2. **WEOC** (water extractable organic C): organic carbon extracted and analyzed for C content and <sup>13</sup>C
  - extracts were dried to powder and analyzed as solid samples
  - VariolIsotope Cube/Isoprime precisION IRMS (Elementar)
3. **total soil C:** soil was dried and analyzed for C and <sup>13</sup>C content
  - VariolIsotope Cube/Isoprime precisION IRMS (Elementar)

# sorbed-C



dashed line = avg of control samples

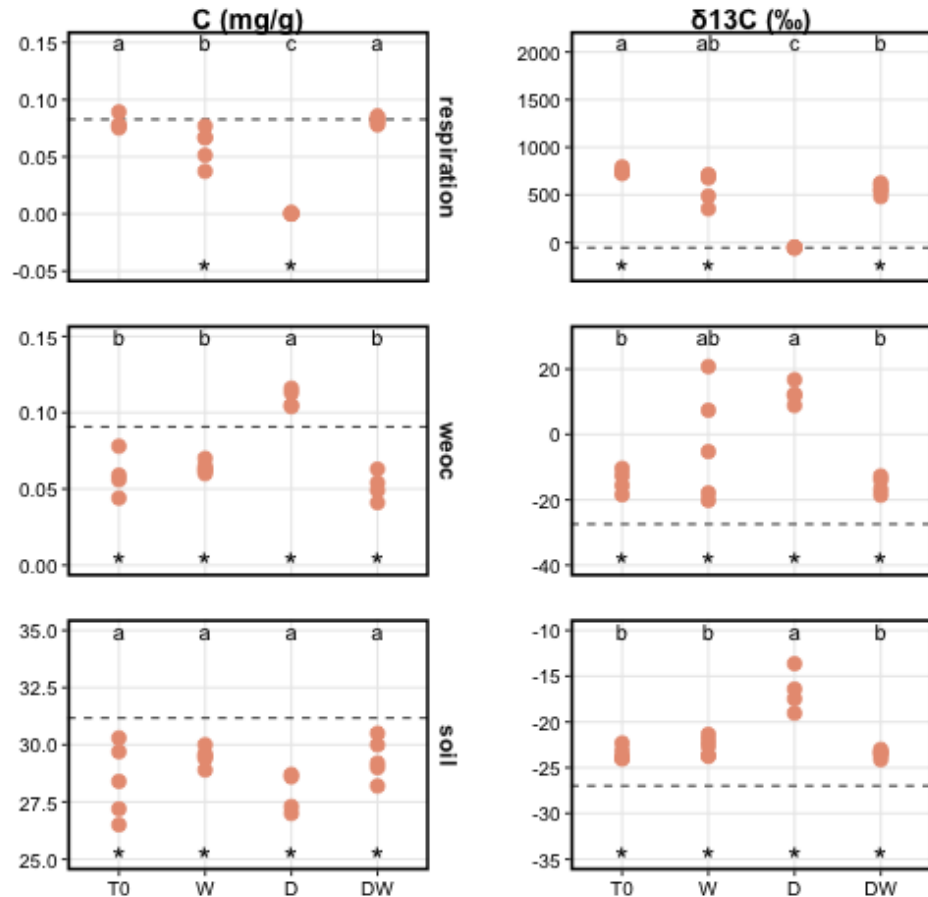
## Figure Caption

C concentrations and  $\delta^{13}\text{C}$  values (relative to Vienna Pee Dee Belemnite) for T0 (time-zero), W (wetting), D (drying), and DW (drying-rewetting) treatments.

Different letters denote significant differences across treatments.

Asterisks denote significant differences from the unlabelled control samples (dashed line).

# sorbed-C

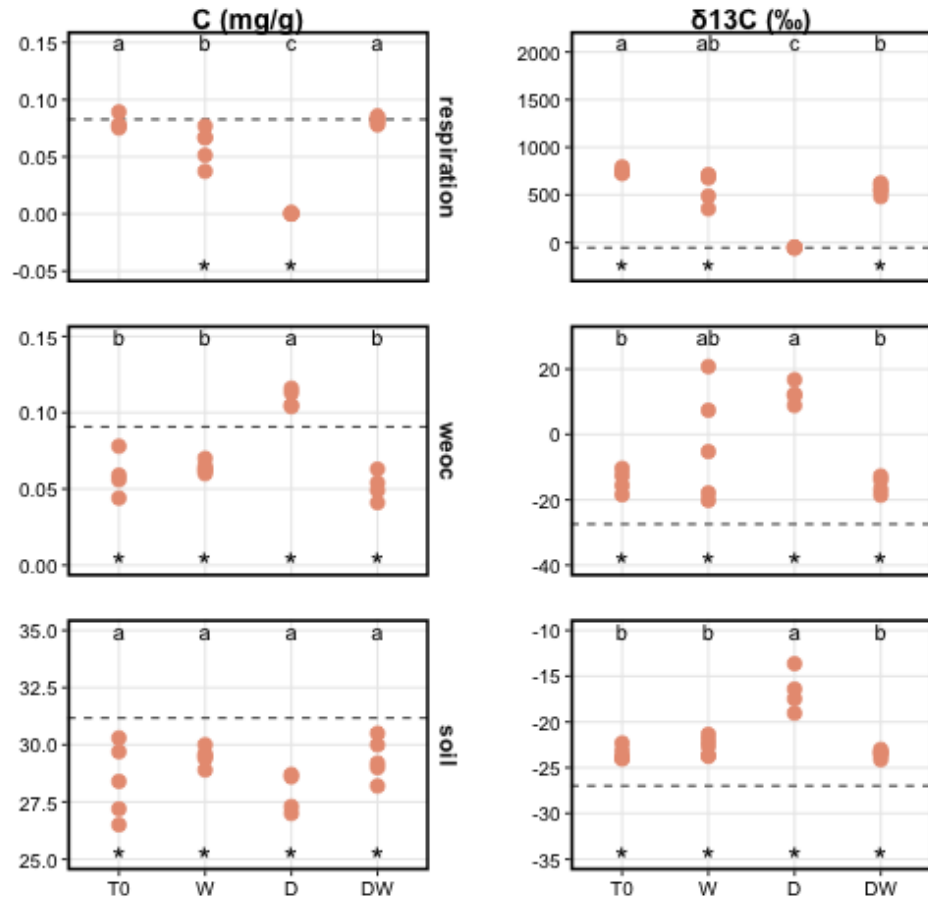


dashed line = avg of control samples

## respiration

- C amendment did not increase respiration
- however, the  $\text{CO}_2$  evolved was significantly enriched throughout (except for D, which had almost no respiration)
  - thus, the added OA did contribute to the  $\text{CO}_2$  evolved, but did not increase overall  $\text{CO}_2$  evolved
  - respiration in these soils is not C limited?
  - no priming effect?
  - perhaps low microbial load, so low respiration potential
  - *even adding oxalic acid in solution form did not increase  $\text{CO}_2$  significantly, see solution-C slides*

# sorbed-C



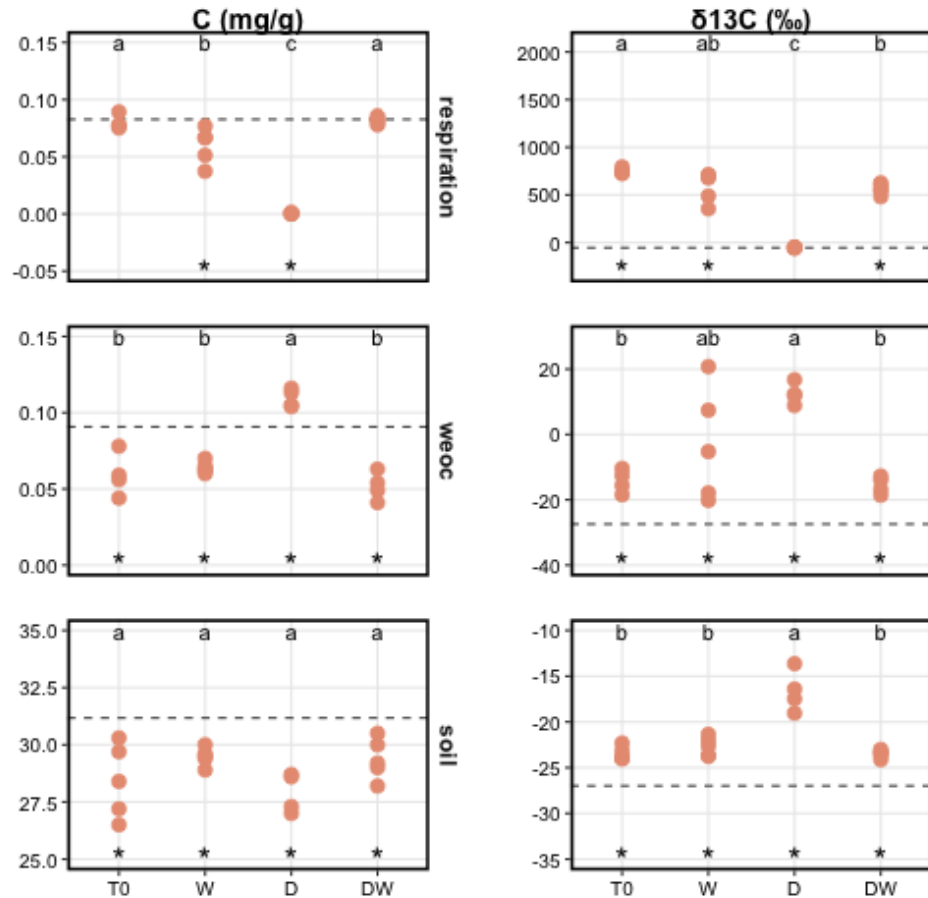
dashed line = avg of control samples

## respiration

- simply mixing the labelled clay with the soil did not change the amount of  $\text{CO}_2$  evolved (T0).
- wetting decreased the evolved  $\text{CO}_2$  by 0.05 mg C/g.
  - negative priming as the oxalic acid was released?
  - high moisture suppressed respiration?
  - lag effect?



# sorbed-C

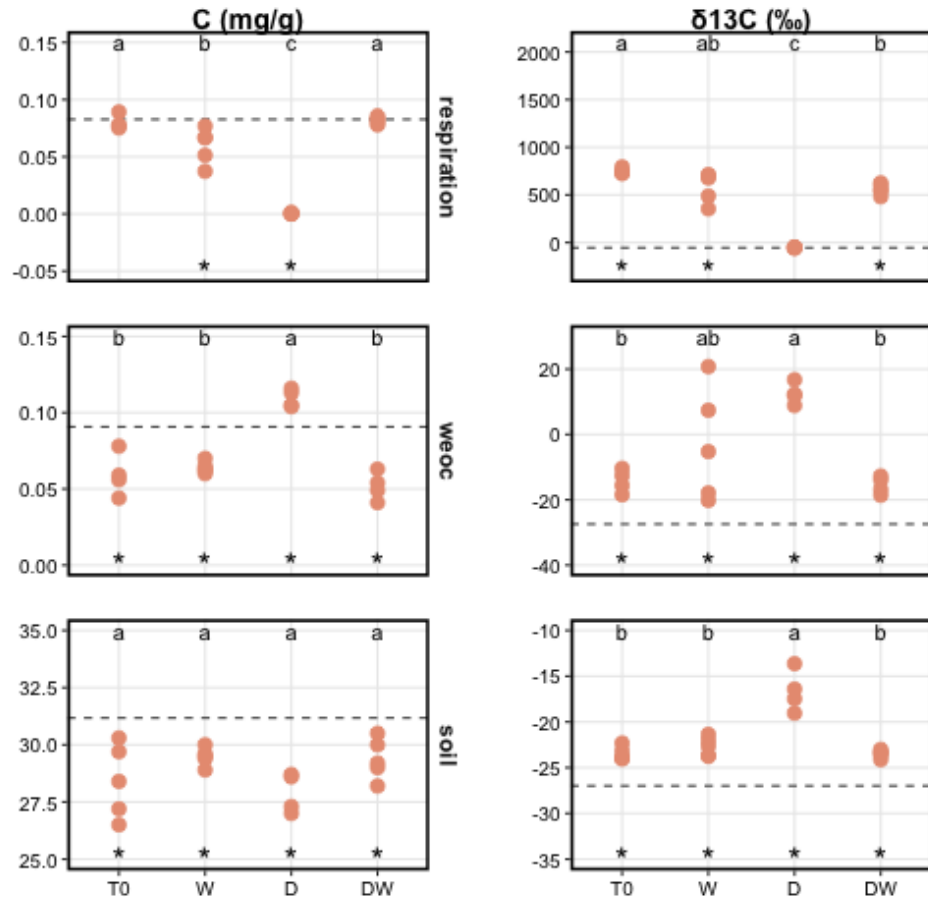


dashed line = avg of control samples

## respiration

- CO<sub>2</sub> enrichment did not differ between W and DW samples.
  - *we expected DW > W*
  - W and DW mobilized C equally?
- but more CO<sub>2</sub> was evolved after DW, compared to W
  - drought effects: destabilization of C, microbial necromass/osmolytes

# sorbed-C

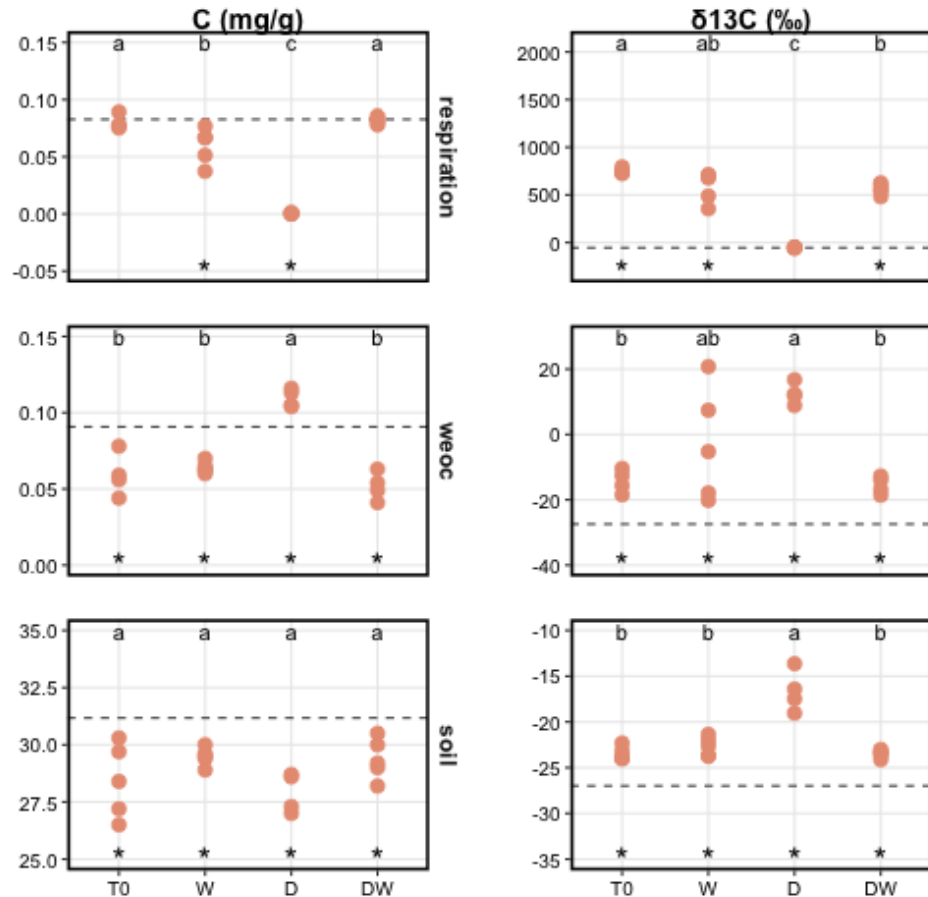


dashed line = avg of control samples

## WEOC

- WEOC concentrations were lower than control for all except D
  - because the C was consumed for respiration? (*inconsistent with resp data*)
- D had the greatest WEOC concentrations
  - (a) desorption of C, (b) microbial necromass/osmolytes, and/or (c) C was not respired and therefore accumulated

# sorbed-C

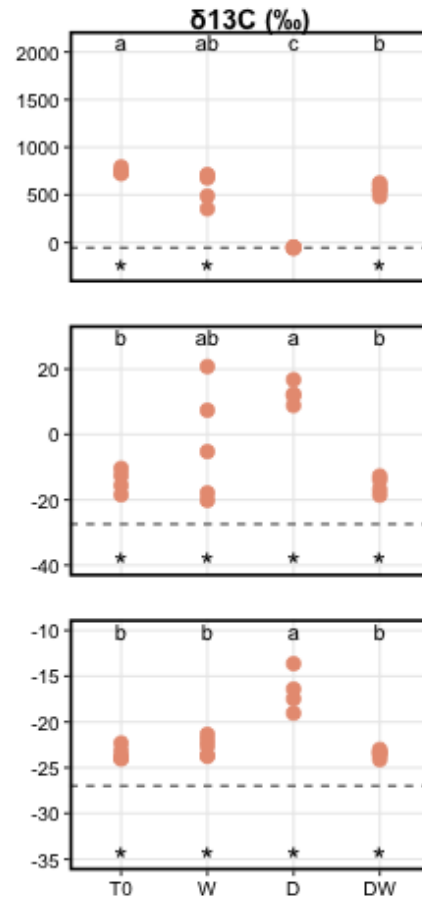
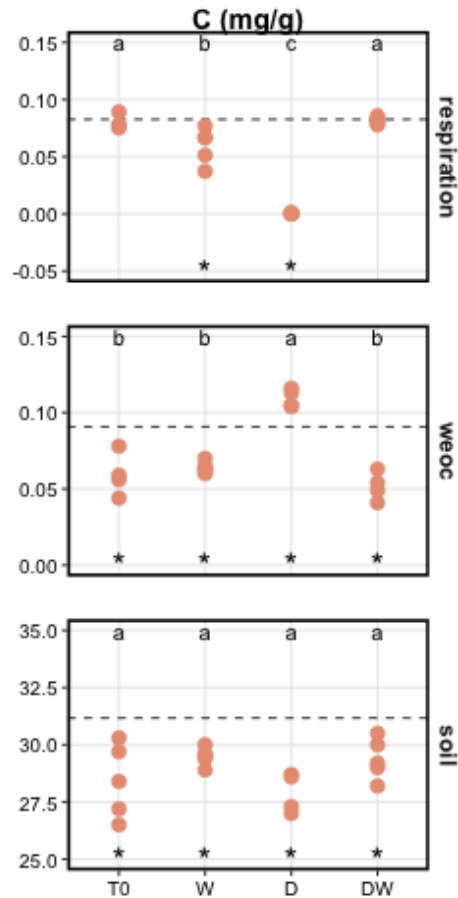


dashed line = avg of control samples

## WEOC

- WEOC was enriched throughout
- W and D were the most enriched
  - because both wetting and drying processes destabilized C
- note: W and D had similar enrichment, but W had lower WEOC concentrations. So less  $^{13}\text{C}$  was destabilized by wetting?
  - also: mass balance slide shows T0 and W had similar  $^{13}\text{C}$ , and D had increased  $^{13}\text{C}$ . So drying destabilized more C than W?

# sorbed-C

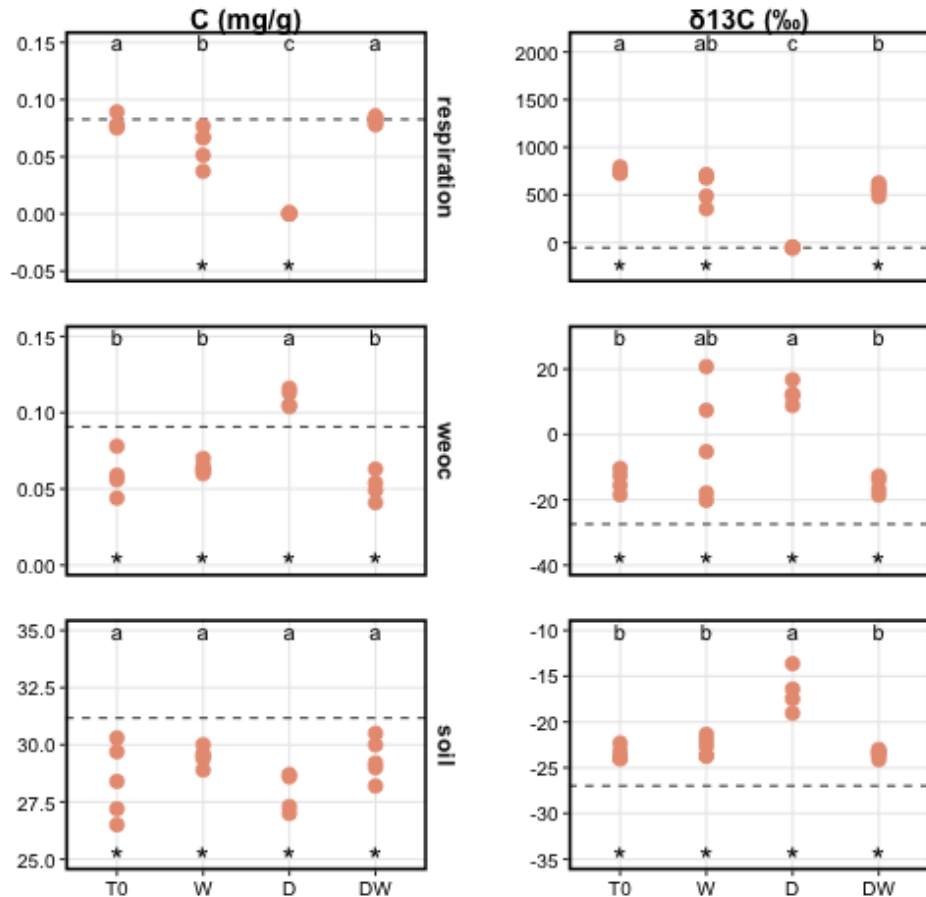


dashed line = avg of control samples

## WEOC

- W had greater variability than D (STATS)
  - preferential wetting patterns?
  - "weaker/non-uniform destabilization" than drying?

# sorbed-C

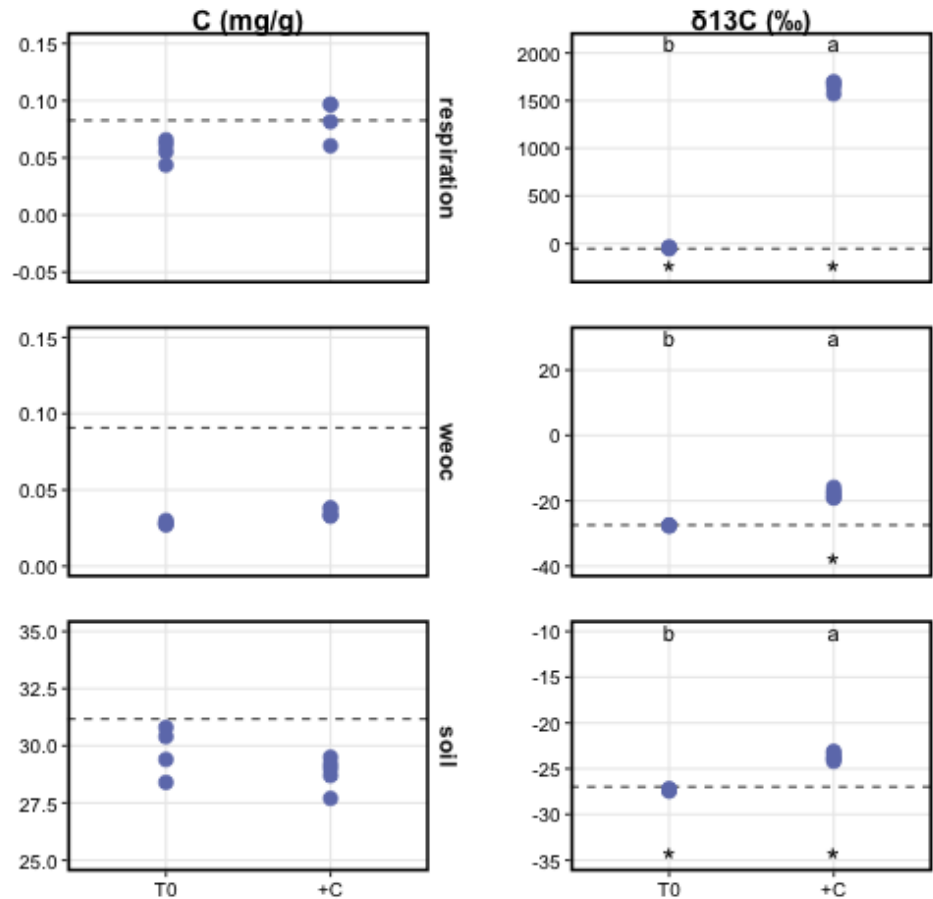


dashed line = avg of control samples

## soil C

- soil C concentrations were lower than control for all
  - because the C was consumed for respiration? (*inconsistent with resp data*)
- soil C was enriched for all,
- with greatest enrichment in D
  - ???
  - accumulation of unrespired C?
  - but D also had the lowest C concentrations (non-sig), and  $^{13}\text{C}$  ug/g was similar to T0 (see mass balance slide)

# solution-C



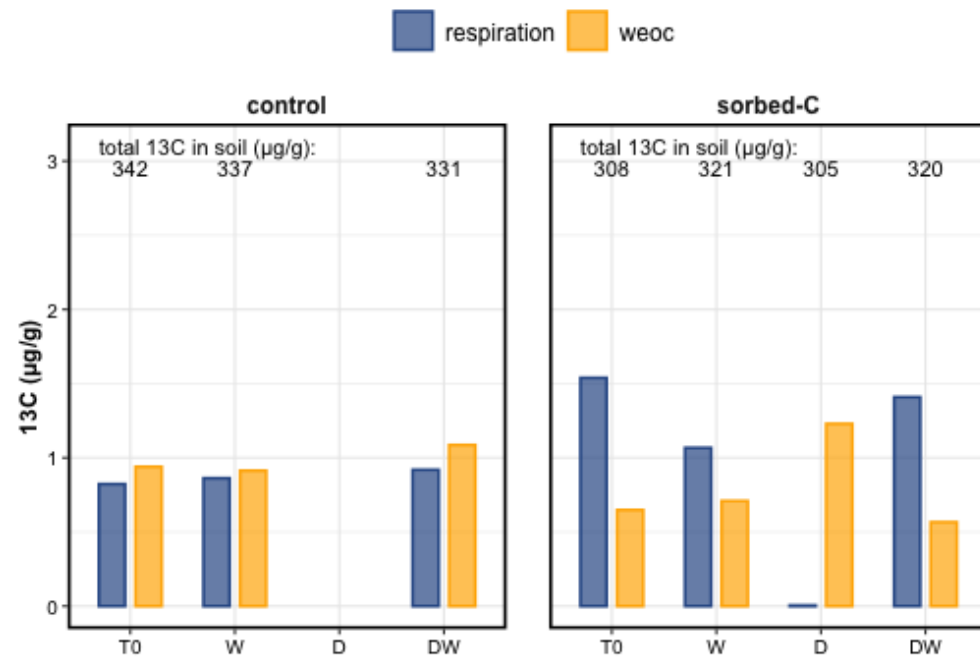
dashed line = avg of control samples

T0 = goethite mixed with soil

+C =  $^{13}\text{C}$ -OA added in solution form

- respiration, WEOC, and soil were all enriched after addition of soluble  $^{13}\text{C}$ -oxalic acid
- respiration: more enriched from solution-C (1648 ‰) compared to sorbed-C (586 ‰)
  - OA was more available in solution form than in adsorbed form
- but despite increased OA contribution, overall  $\text{CO}_2$  evolved did not increase.

# Mass Balance: sorbed-C



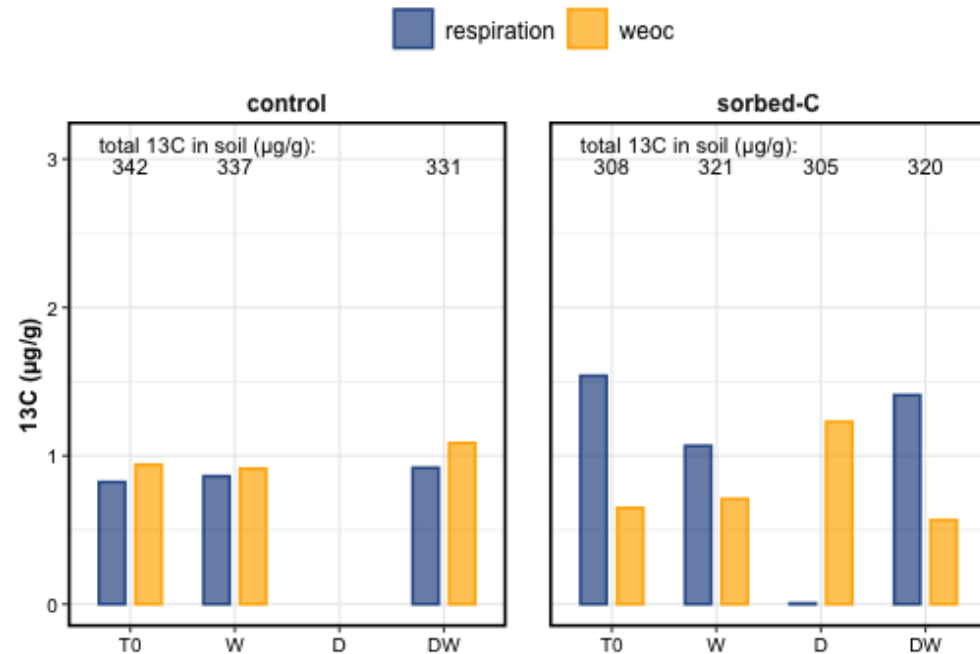
493 µg (7.72 µg/g) <sup>13</sup>C was added in sorbed form.

<sup>13</sup>C µg/g in sorbed-C soils compared to control soils:

	resp	soil	weoc
T0	+0.716	-34.1	-0.291
W	+0.205	-15.6	-0.204
D	NA †	NA †	NA †
DW	+0.490	-11.3	-0.520

† no control samples

# Mass Balance: sorbed-C



## isotope retention

493 µg (7.72 µg/g)  $^{13}\text{C}$  was added in sorbed form.

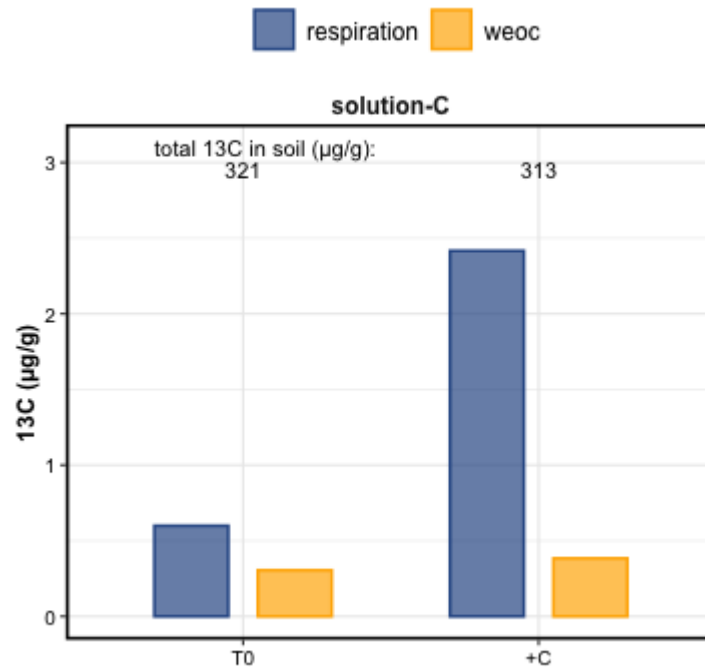
But the treated soils showed decreased  $^{13}\text{C}$  values.  
(???)

comparing added  $^{13}\text{C}$  with (resp + soil) difference,  
there is a missing piece of  $^{13}\text{C}$

- methane? no, because  $\text{CH}_4$  production was very, very low, and did not change with treatment/OA addition.



# Mass Balance: solution-C



417  $\mu\text{g}$  (6.54  $\mu\text{g/g}$ )  $^{13}\text{C}$  was added as solution

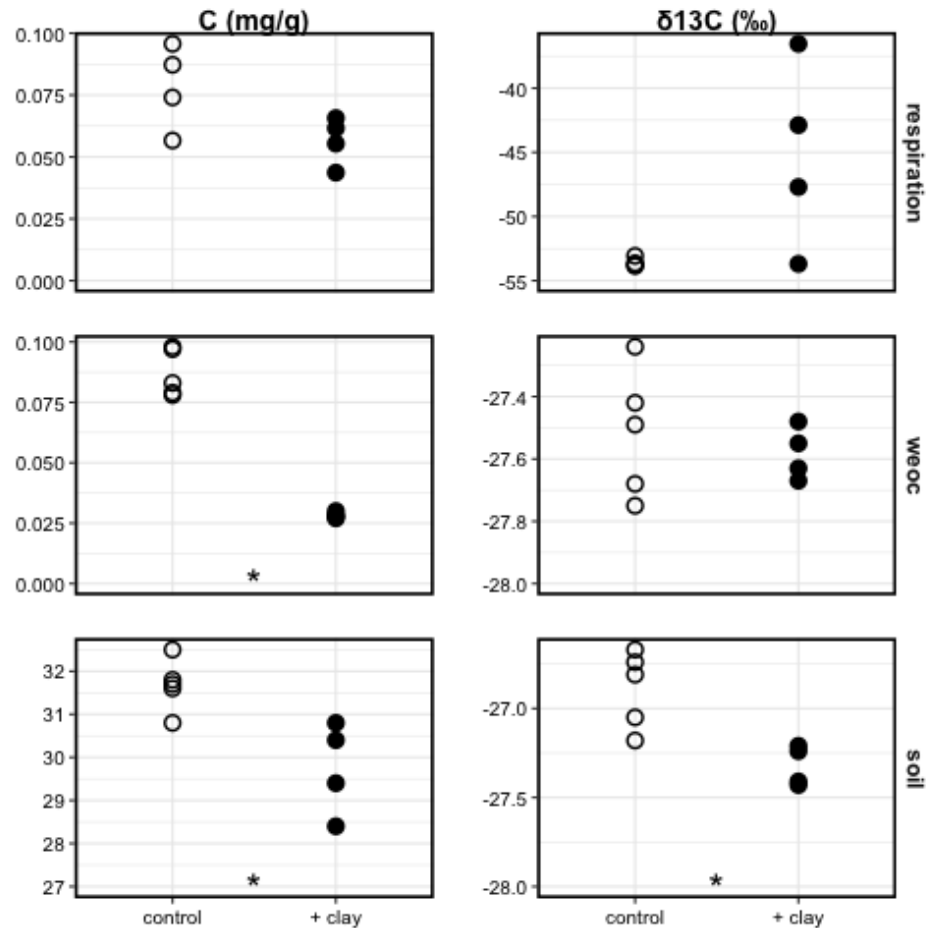
- very little change in WEOC- $^{13}\text{C}$
- some increase in resp- $^{13}\text{C}$
- decrease in soil- $^{13}\text{C}$

**SO WHERE DID THE  $^{13}\text{C}$  GO???**

$^{13}\text{C}$   $\mu\text{g/g}$  in +C compared to T0 (time zero):

	resp	soil	weoc
+C	+1.82	-8	+0.08

## extra: How did clay addition influence C?



We compare time-zero of control (soil only) with time-zero of solution-C treatment (soil + goethite)

- respiration did not change significantly, but enrichment was more variable
  - ???
- weoc decreased, but enrichment did not change
  - previously available sorbed onto goethite
- soil C decreased, as did enrichment
  - ???

# Conclusions

- wetting did not destabilize much sorbed-C
- drying released more sorbed-C than wetting

# MAJOR QUESTIONS

1. do we use  $\delta^{13}\text{C}$  or  $\text{C}^{13}$  mass (mass balance) to determine extent of destabilization?
2. missing  $^{13}\text{C}$ ?

# Session Info

Date run: 2021-01-29

Slides prepared using **xaringan**

```
## R version 4.0.2 (2020-06-22)
## Platform: x86_64-apple-darwin17.0 (64-bit)
## Running under: macOS Catalina 10.15.7
##
## Matrix products: default
## BLAS:   /System/Library/Frameworks/Accelerate.framework/Versions/A/Frameworks/vecLib.framework/Versions/A/libBLAS.dylib
## LAPACK: /Library/Frameworks/R.framework/Versions/4.0/Resources/lib/libRlapack.dylib
##
## locale:
## [1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8
##
## attached base packages:
## [1] stats      graphics  grDevices  utils      datasets  methods   base
##
## other attached packages:
## [1] agricolae_1.3-3 patchwork_1.1.1 outliers_0.14  PNWColors_0.1.0
## [5] drake_7.13.0    forcats_0.5.0  stringr_1.4.0  dplyr_1.0.2
## [9] purrr_0.3.4     readr_1.4.0    tidyr_1.1.2    tibble_3.0.4
## [13] ggplot2_3.3.3   tidyverse_1.3.0
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