

this time: SD; normal curve;
 next time: experimental design;
 probability

Wed: JD ch
 (017)
 LN pp. 95-118

AMS7
 27 Jan
 2016

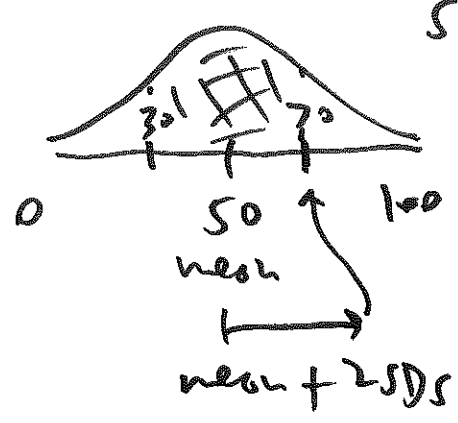
①

hwk 1 due this Wed

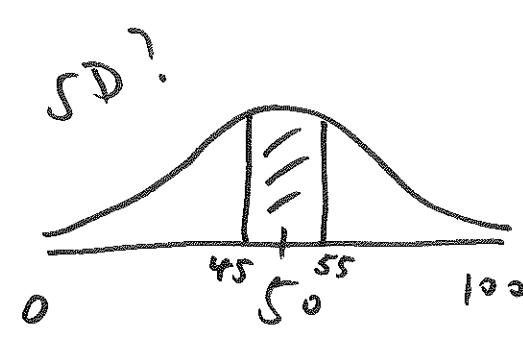
JD

new office hrs: MWF 11.45am - 12.45pm

in BE 357C



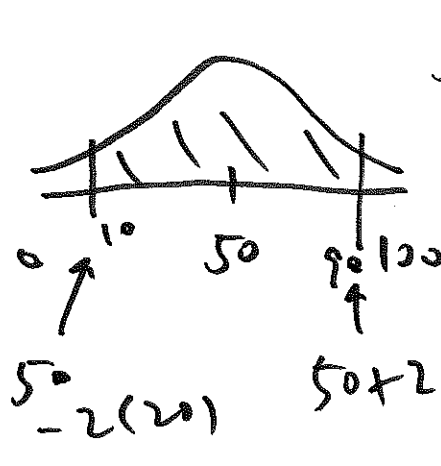
SD 10?
 no still too small



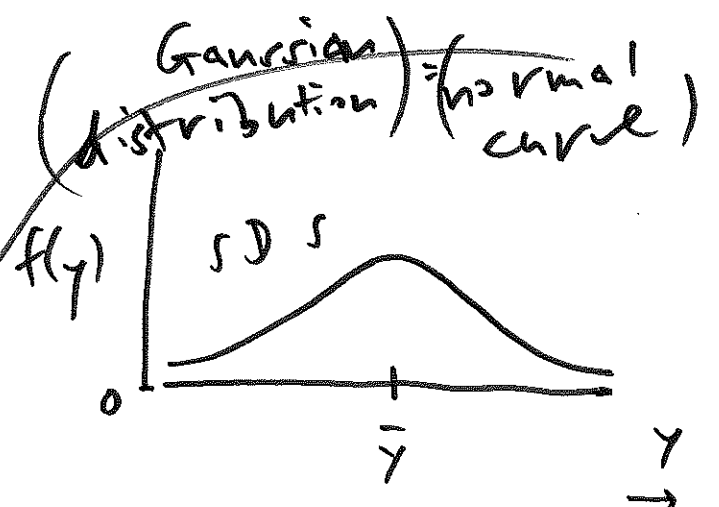
SD 5?
 no too small



SD 50?
 no too big



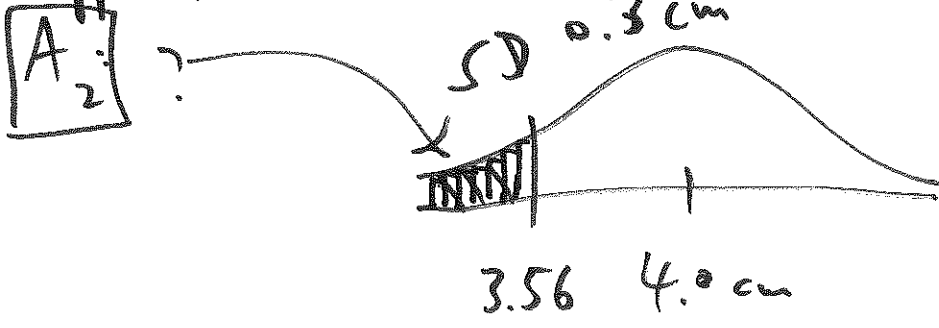
SD 20?
 yes about right



Q: $\% \leq 3.56 \text{ cm}$?

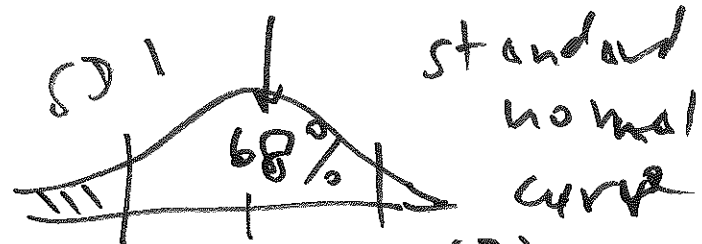
exact
 A₁: $\frac{2}{24} =$ ②
 8.3%

approximate



butterfly
 why length

emp. rule (part 2)

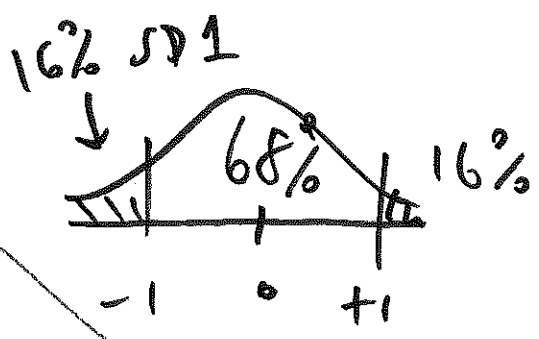
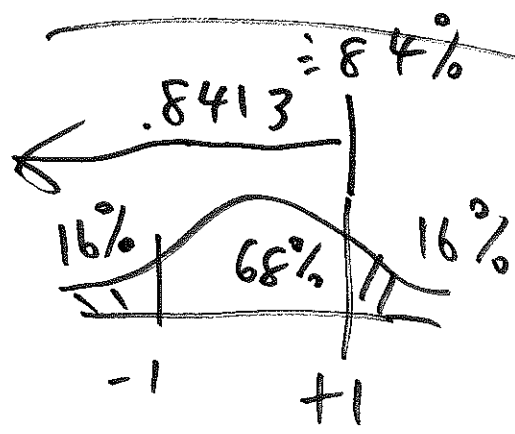


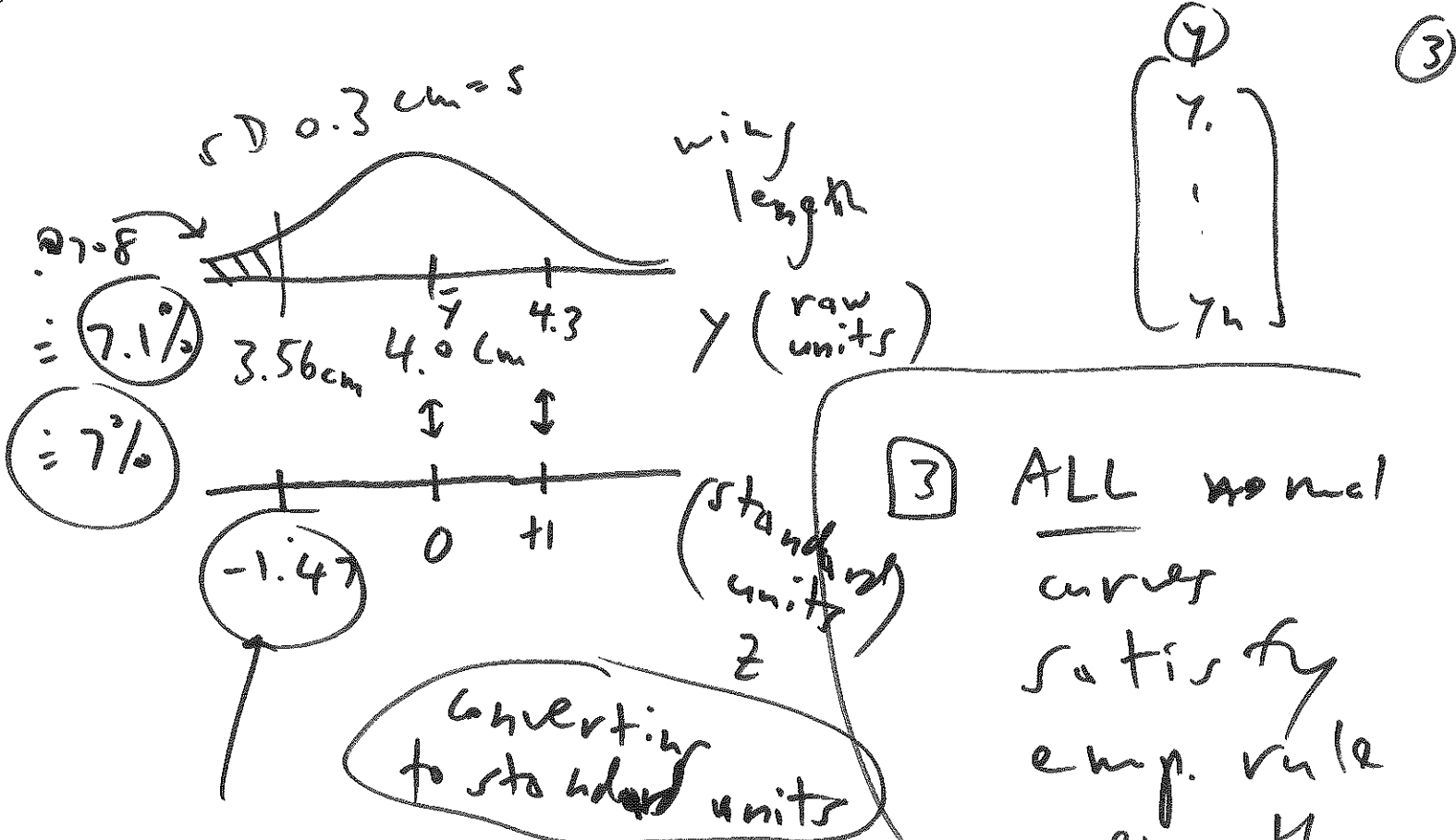
1. normal curves are symmetric

2. total area under each normal curve is 100% = 1.0

-1 0 +1 (z)
 -1.00
 table:
 .1587 ← decimal
 "

$15.87\% \approx 16\%$





3 ALL normal curves satisfy emp. rule exactly

converting to standard units

$$\frac{3.56 \text{ cm} - 4.0 \text{ cm}}{0.3 \text{ cm}} = \frac{x - \text{mean}}{SD} = \frac{y - \bar{y}}{s}$$

$$= \frac{-0.44}{0.3} = -1.47$$

controlled (11.55)

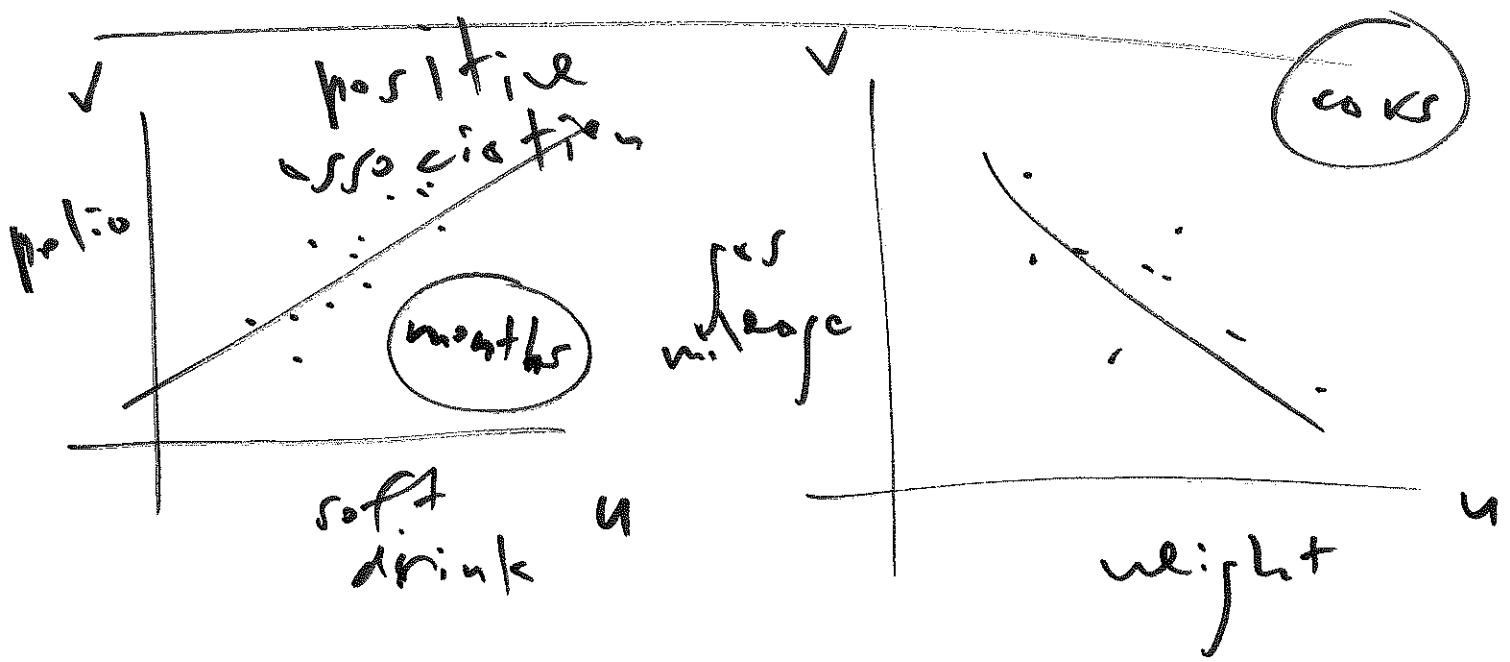
Experiment: there is a treatment (T) group & a control (C) group, & experimenters control who goes into which group

Y (outcome) : cortex weight ⁽⁴⁾

X (treatment) : T vs. C

Z (potential confounding factor) : genetic background (cortex weight)

(PCF)



Y (outcome)

control wt

X (treatment)

T enriched

C deprived

Z (genetics)

as Z ↑, Y ↑ ✓

as X goes from C to T,

Z ↑ ↓ ✓

PCFs are the enemy in experimental design because they cause bias in conclusions

how defeat PCFs?

hold them constant in T/C comparison