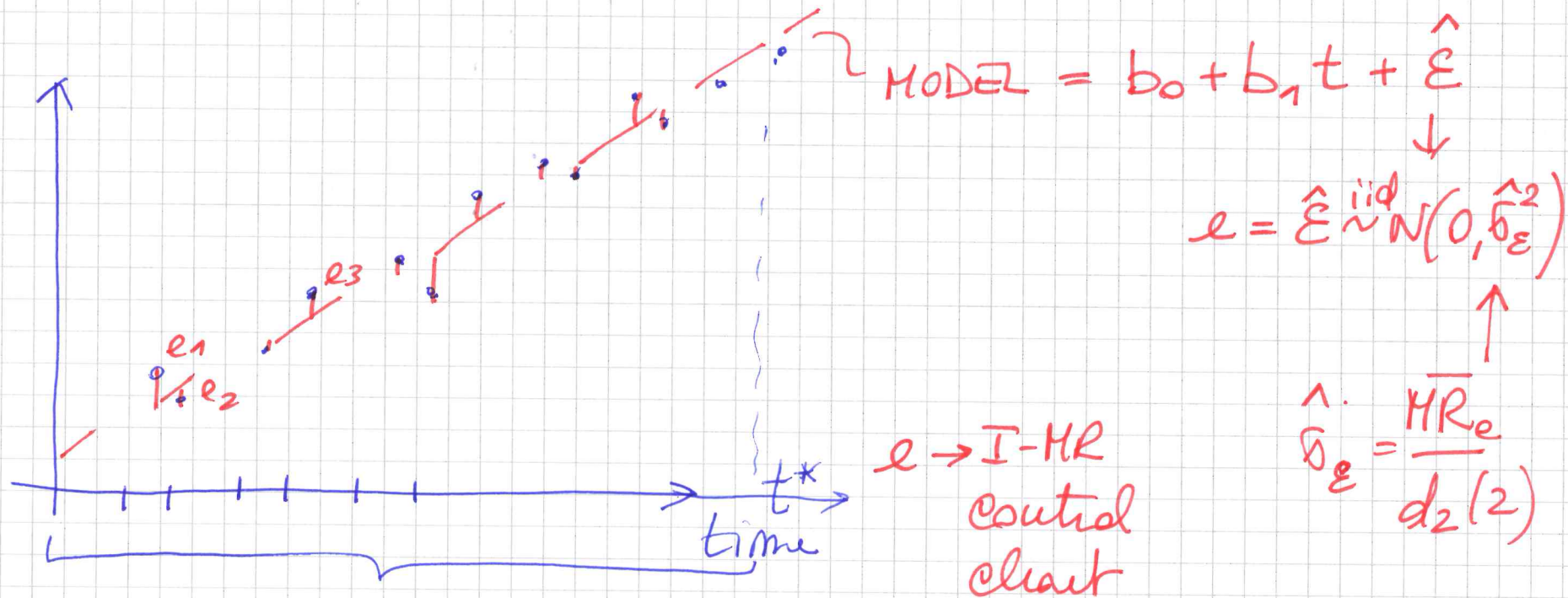


QDA 2023.05.17

①

SPC or SQM with NON iid data

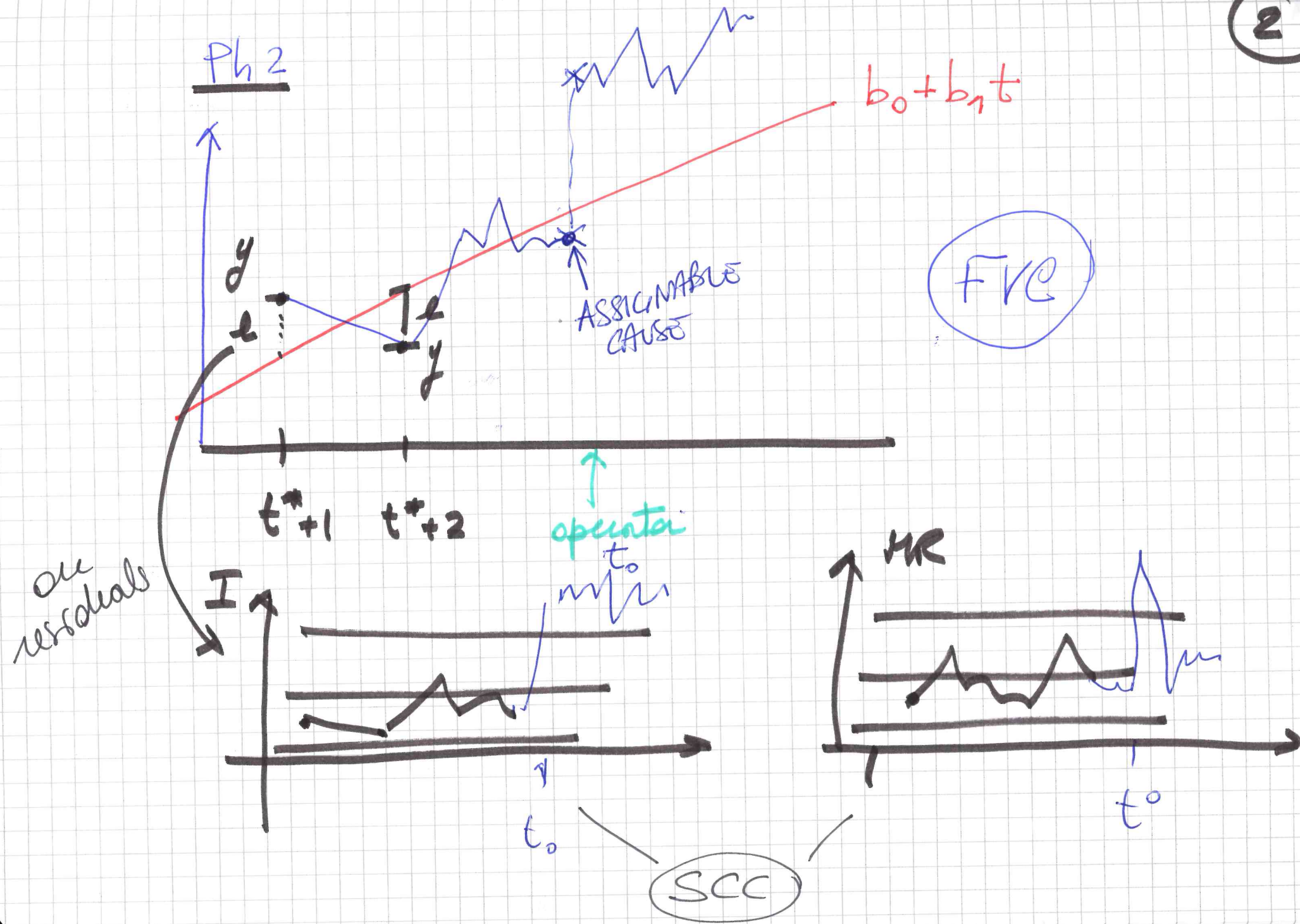


Ph 1



MODEL (complete, with all the unknown parameters estimated) for the IN CONTROL DATA

2



SAMPLING STRATEGY

$m > 1$

Ishikawa example

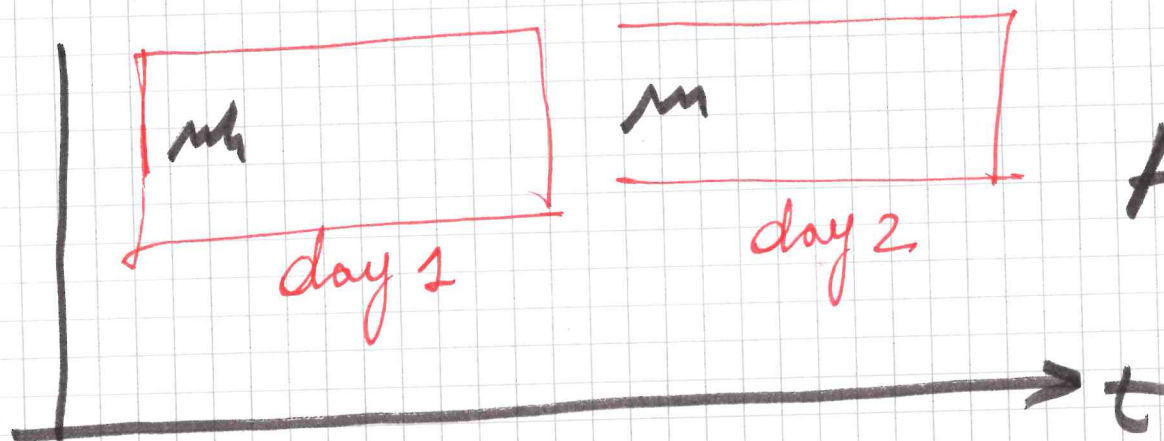


ex AR(1) (3)

mm

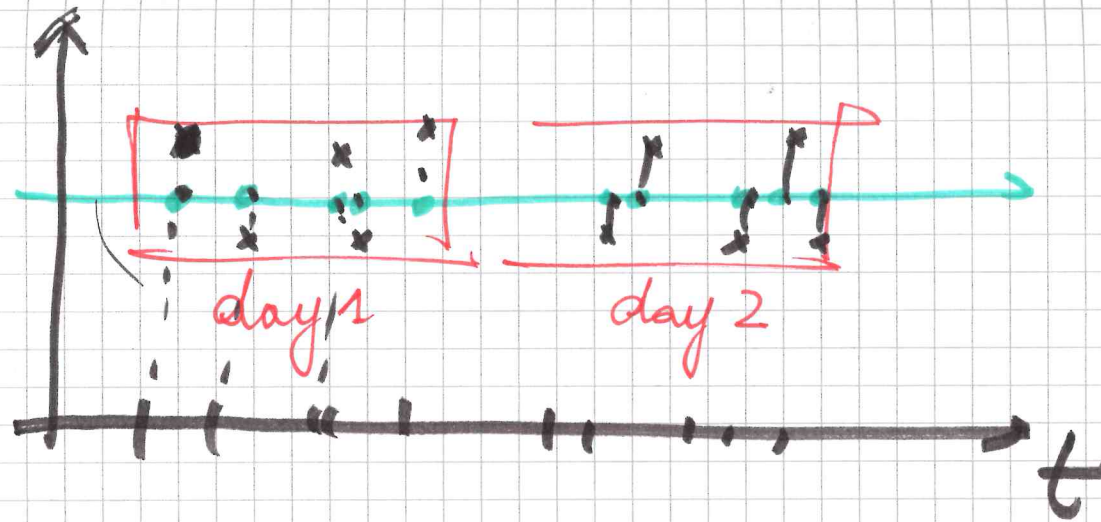
h constant

$$X_t = \beta_0 + \beta_1 X_{t-1} + \epsilon_t$$



AR?

④



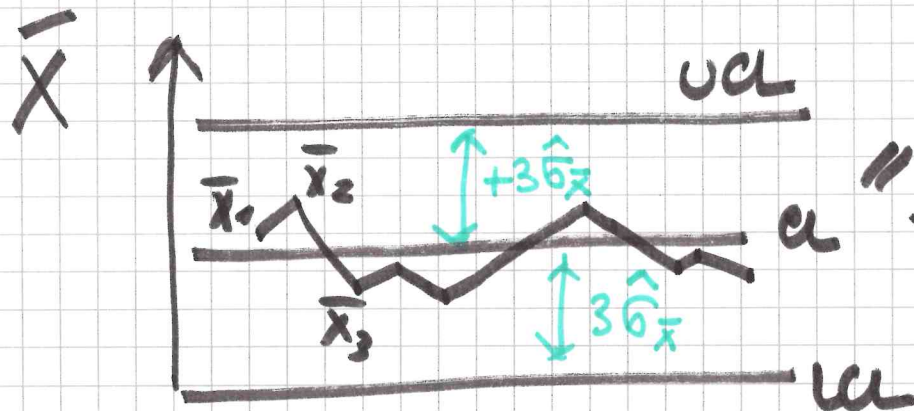
RND
SAMPUNG

~~$X \sim N(\mu, \sigma^2)$~~

$m > 1$

VIOLATION
of the iid
assumption

↓ ↓
batch $m=5$ batch $m=5$



a "HUGGING" or
STRATIFICATION

it seems that the control
chart on the \bar{X} is
overestimating
the variability
of \bar{X}

(5)

Remind that

$$X \stackrel{\text{iid}}{\sim} N(\mu, \sigma^2)$$

 \Downarrow

$$\bar{X} = \frac{1}{n} \sum_i X_i \rightarrow E(\bar{X}) = E\left(\frac{1}{n} \sum X_i\right) =$$

$$= \frac{1}{n} \sum \underbrace{E(X_i)}_{\mu} = \frac{1}{n} n \mu = \mu$$

ASSUMPTIONS
in
 \bar{X} ee

$$\sigma_{\bar{X}}^2$$

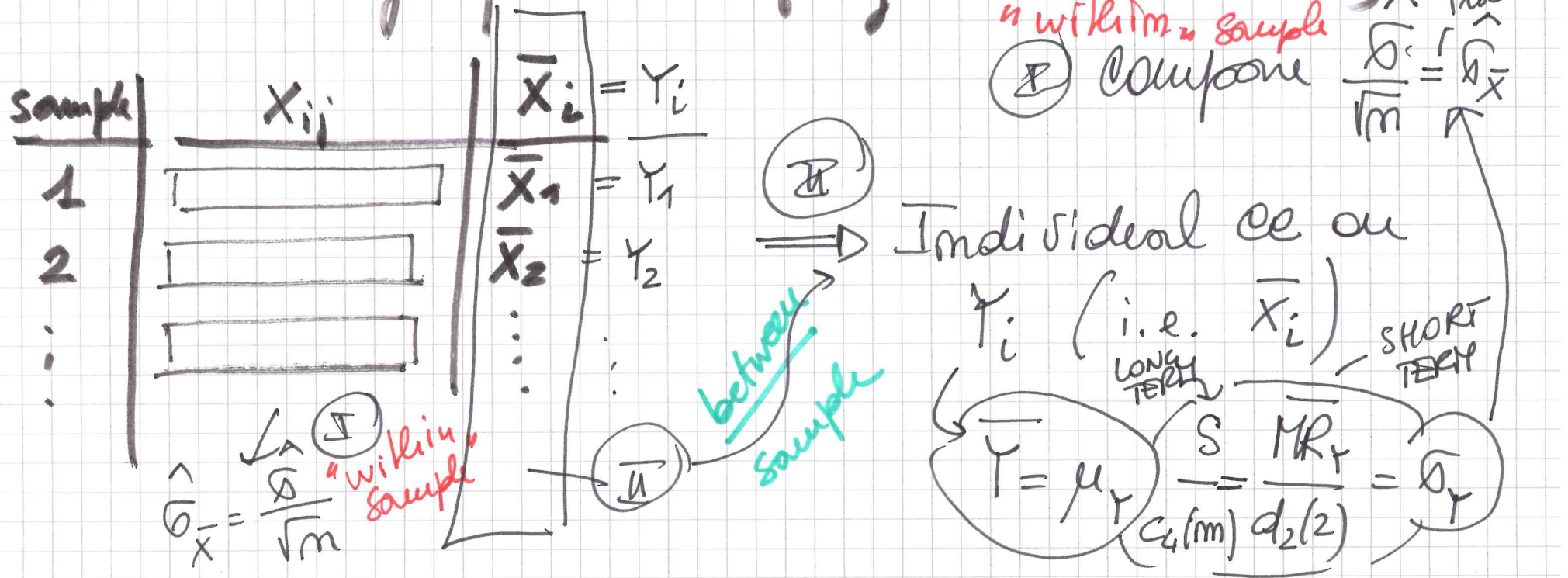
$$V(\bar{X}) = V\left(\frac{1}{n} \sum X_i\right) = \frac{1}{n^2} V(\sum X_i)$$

ASSUMING
NO COV
UNCORRELATED

$$= \frac{1}{n^2} \sum V(X_i) = \frac{n \sigma^2}{n^2} = \frac{\sigma^2}{n}$$

If X_i are non iid $\Rightarrow \sigma_{\bar{X}}^2 = \frac{\sigma^2}{m}$ in the 6
 \bar{X} ee can be over- or under-estimating
 the real $V(\bar{X})$

If I suspect HUGGING or STRATIFICATION (or any other strange pattern of \bar{X}_i in the X bar ee)



when the sampling strategy is structured

X_i

day
1 6.00

1 10.

1

1

1

2

2

2

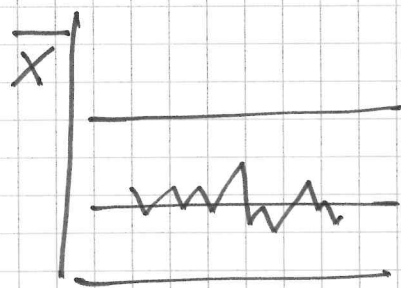
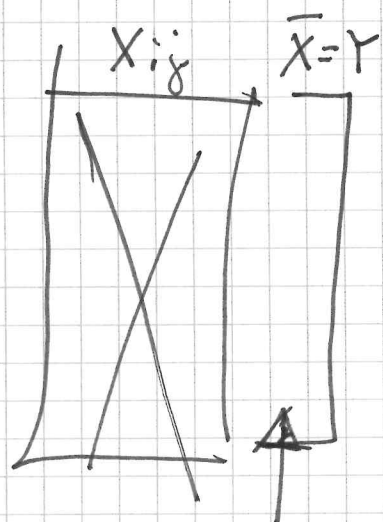
2

2

data originally in the first day, first sample
(first row)
→ ACF
PACF
autocor?

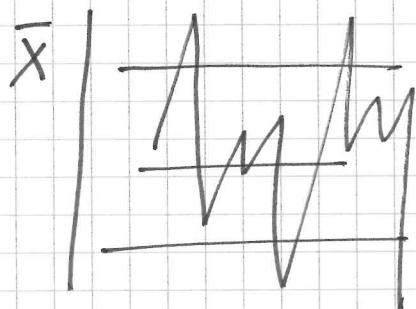
data in the second day (second row)

$m > 1$ \rightarrow plots in \bar{X} ce



HUGGING

\rightarrow suspect iid assumption are violated



OVERDISPERSION

What can we do?

between
sample

try
to
model
 X_{ij} 's

just take all the \bar{X} 's as a new sequence of quality indicator (i.e., a new random variable Y) and proceed with a \bar{I} ce on this new Y 's
pros: it is working?
cons: losing information

batching / gapping *

best model $X_{ij} = f(\overbrace{i, j}^{\text{ARIMA}}, \text{dummy}, \text{trend}) + \varepsilon$

pros: better

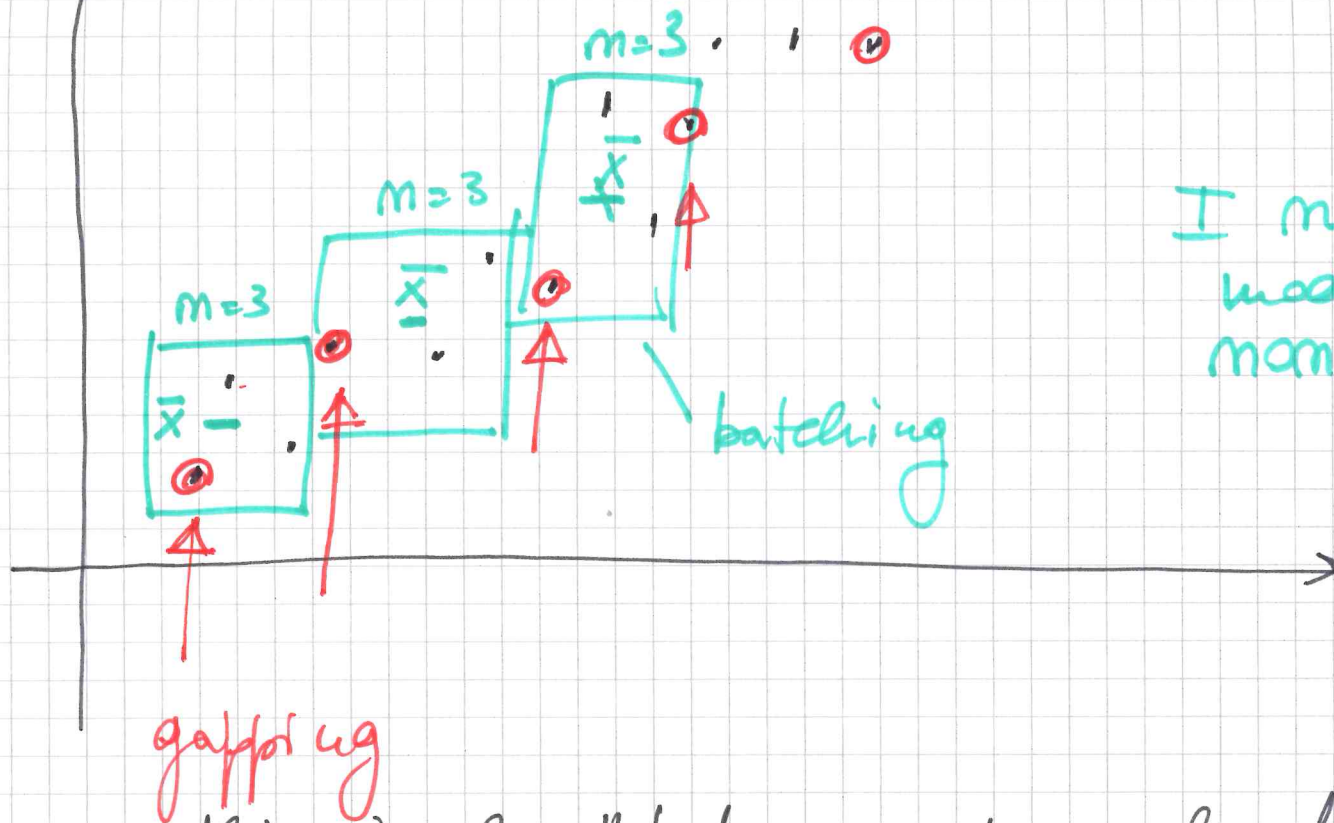
model for the data, without
losing info

\downarrow
FVC
+
SCC

cons: finding the right
model can be a
complex task

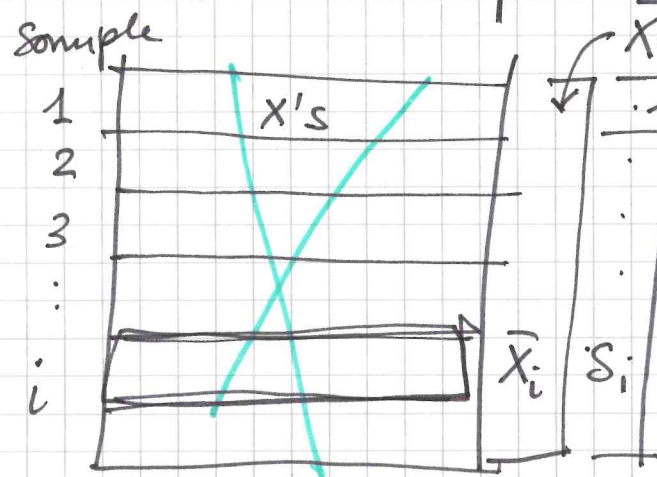
* a note: gapping or batching can work only for
stationary processes

ex: trend = non stationary process



I need to model the non stationarity!

You can use this similar "between" approach for any statistic taken at each sample



Individual cell on \bar{Y}
 Individual cell on \bar{W}
 (Box Cox?)