



Karolinska  
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SciLifeLab

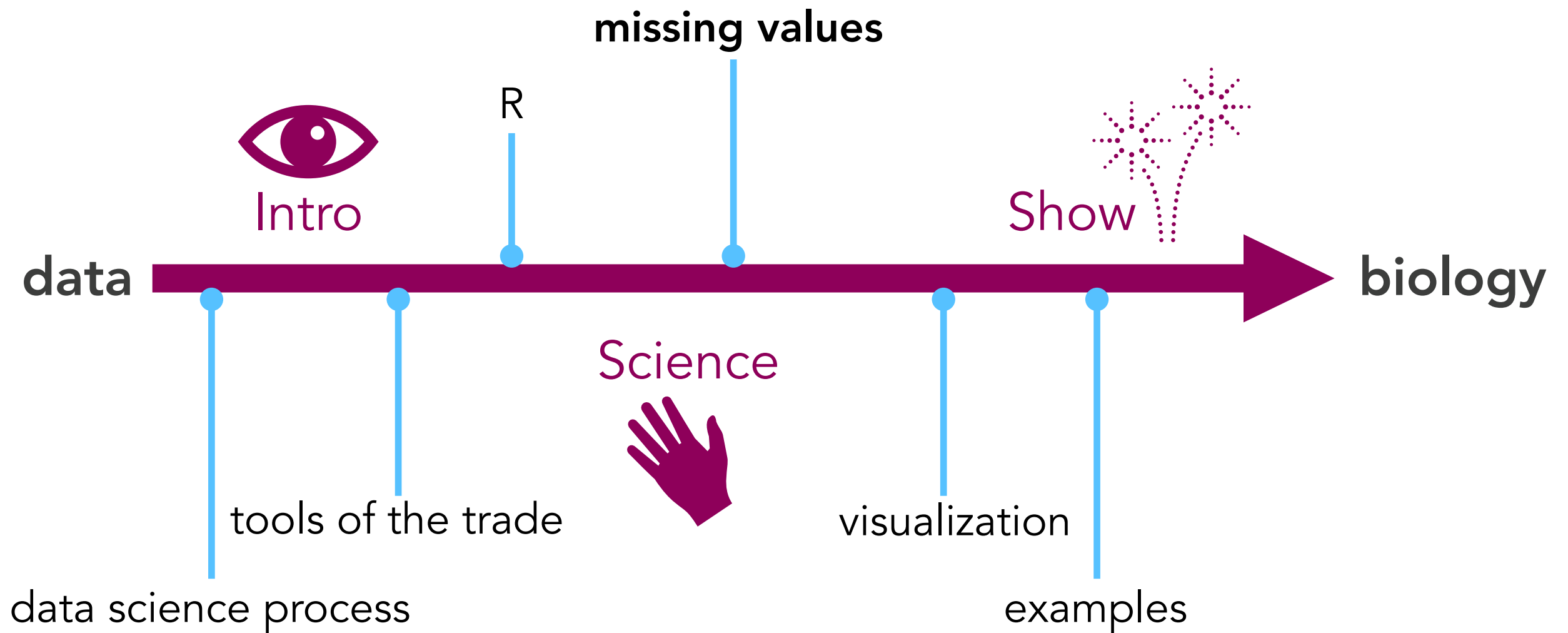


# Missing value imputation

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# Three types of missing values

A large purple circle containing the text "MCAR".

MCAR

missing completely at random

no cause to missingness

A large purple circle containing the text "MAR".

MAR

missing at random

missing values can be  
explained by the available data

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missing data can be ignored

A large purple circle containing the text "NMAR".

NMAR

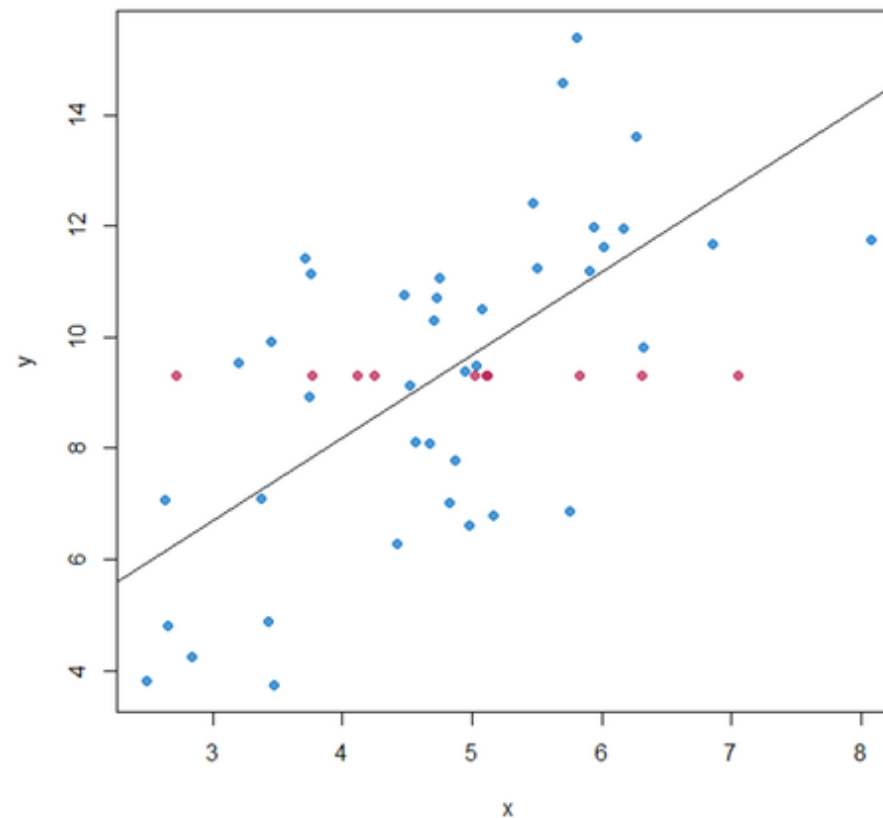
not missing at random

missing data **cannot** be ignored

There are *numerous* methods to fill missing values.

# Keep the sample statistics

impute with **mean** | **median** | **mode** of the data

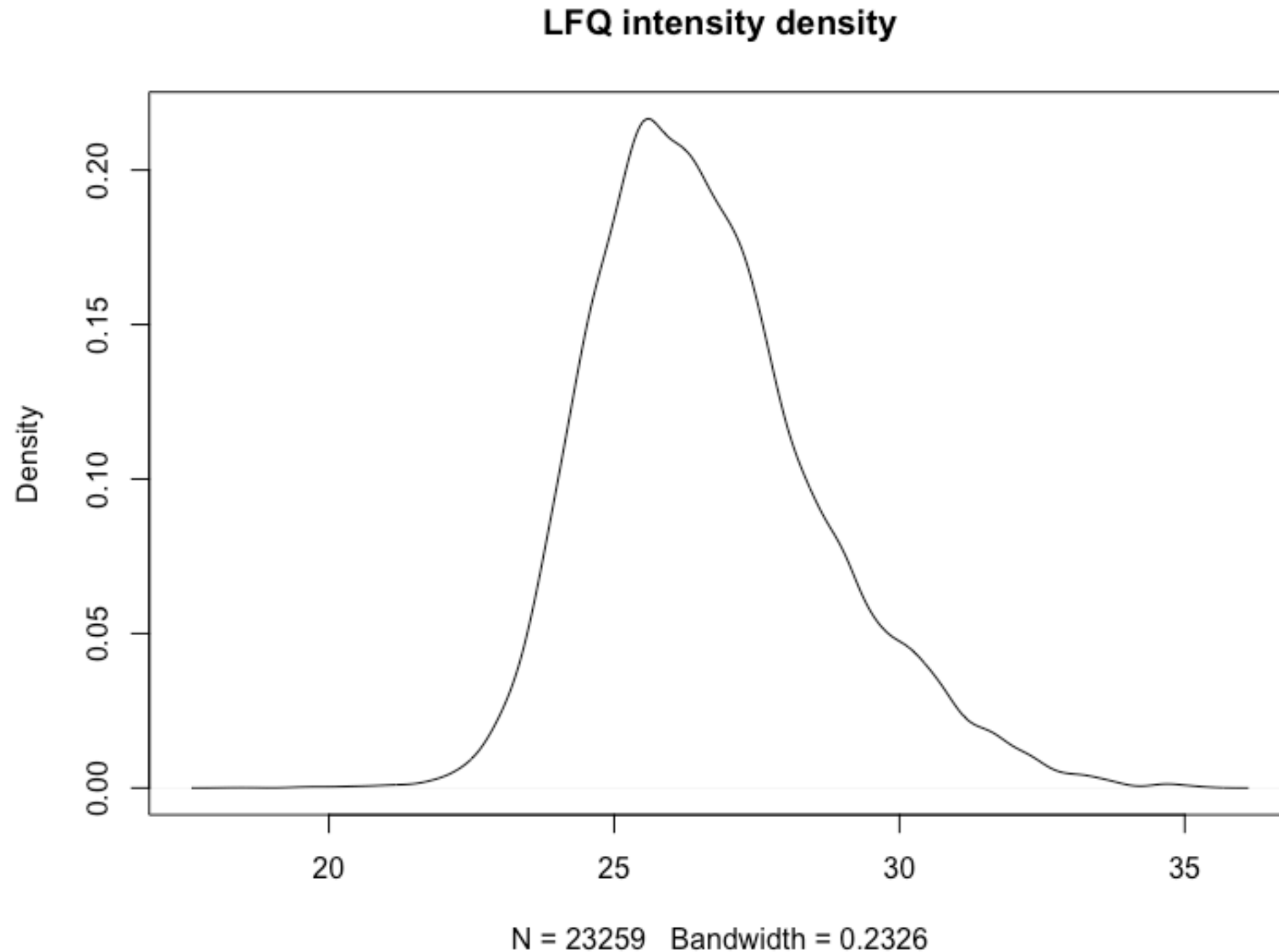


**not optimal!**

changes variance, introduces bias, ...

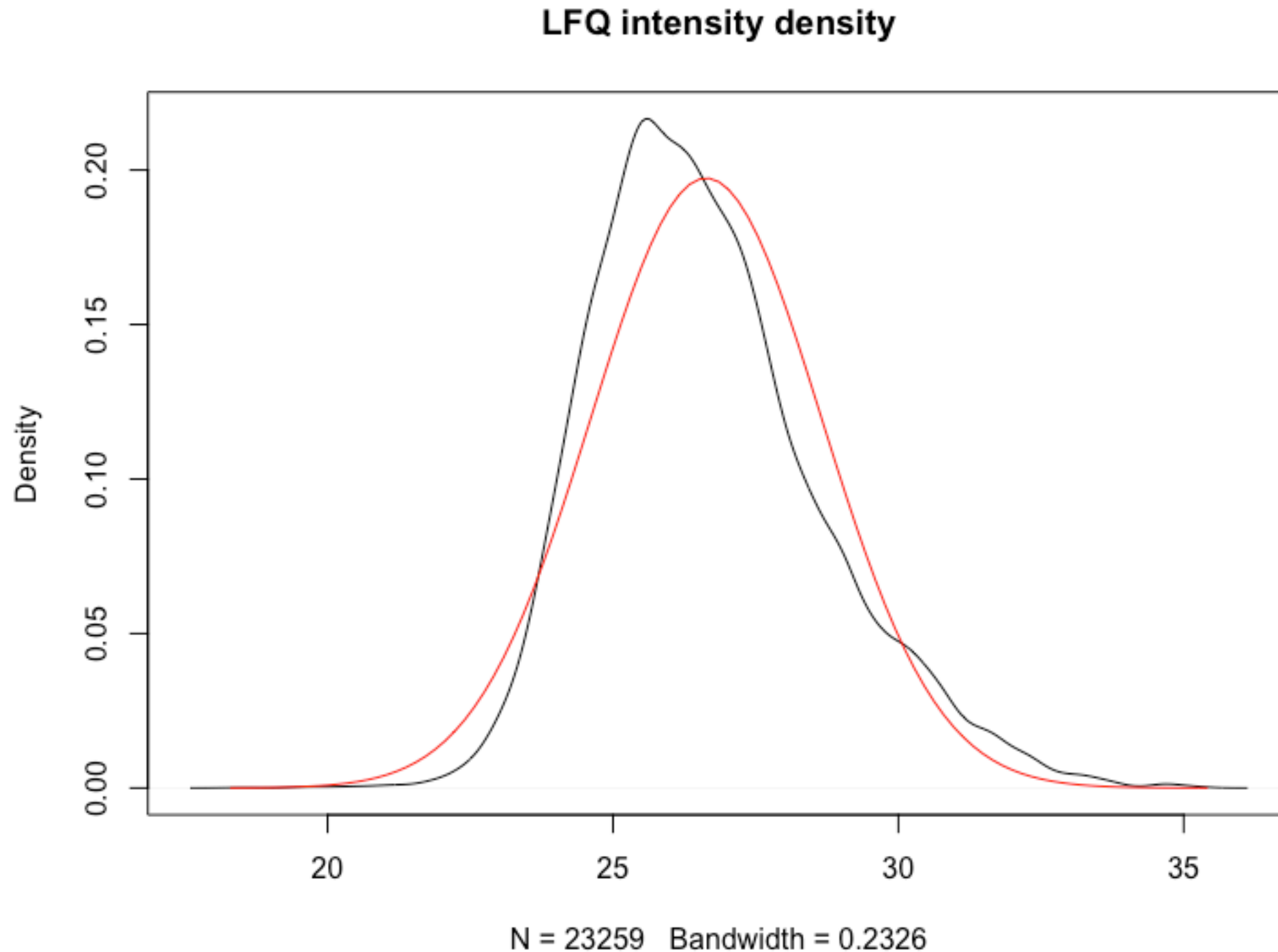
# A usual dataset

Caro's dataset (HepG2 + **NC1** or DMSO)

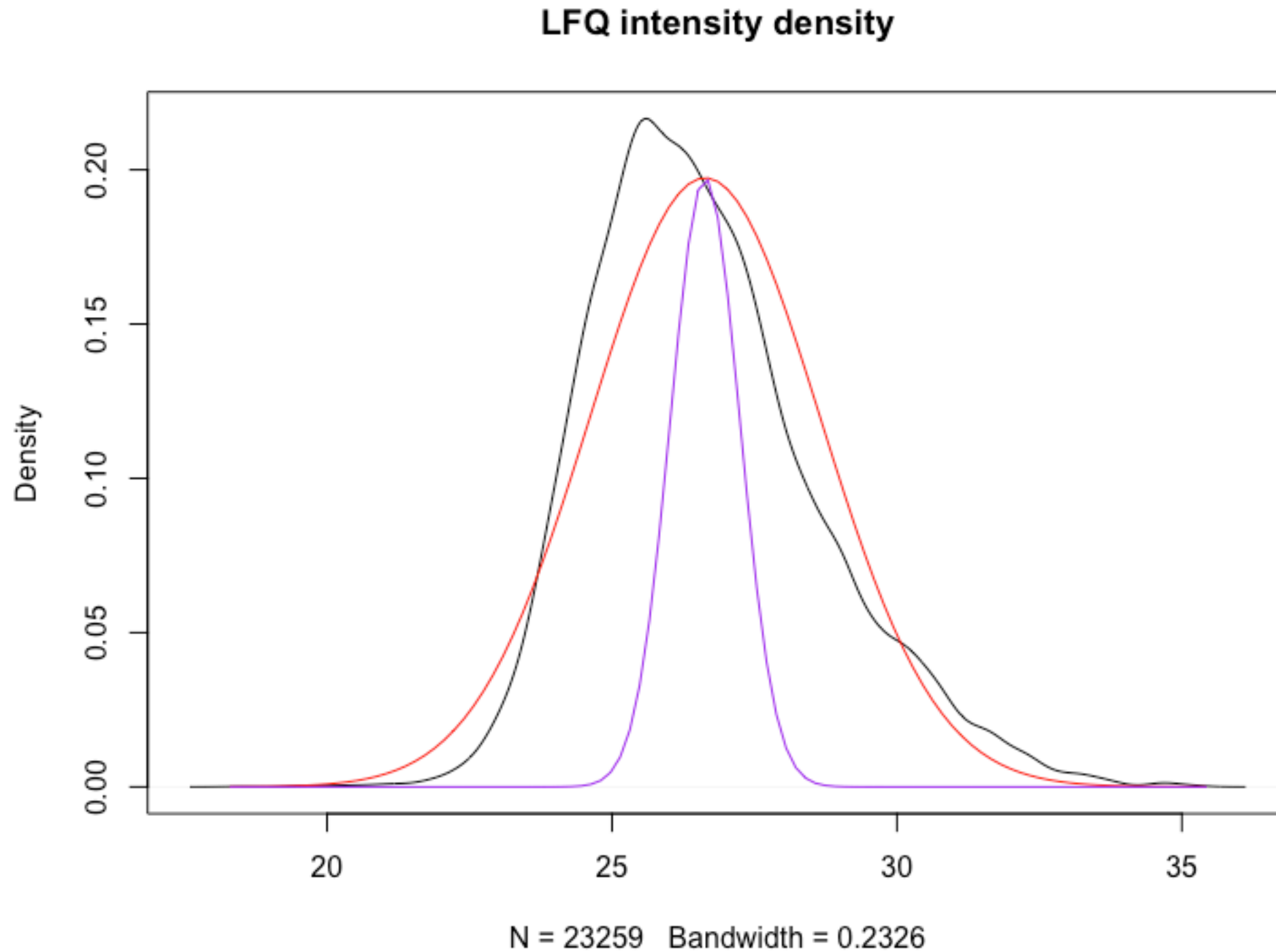


# Infer a normal distribution

Disclaimer: This is how I understood Perseus does it. ;)

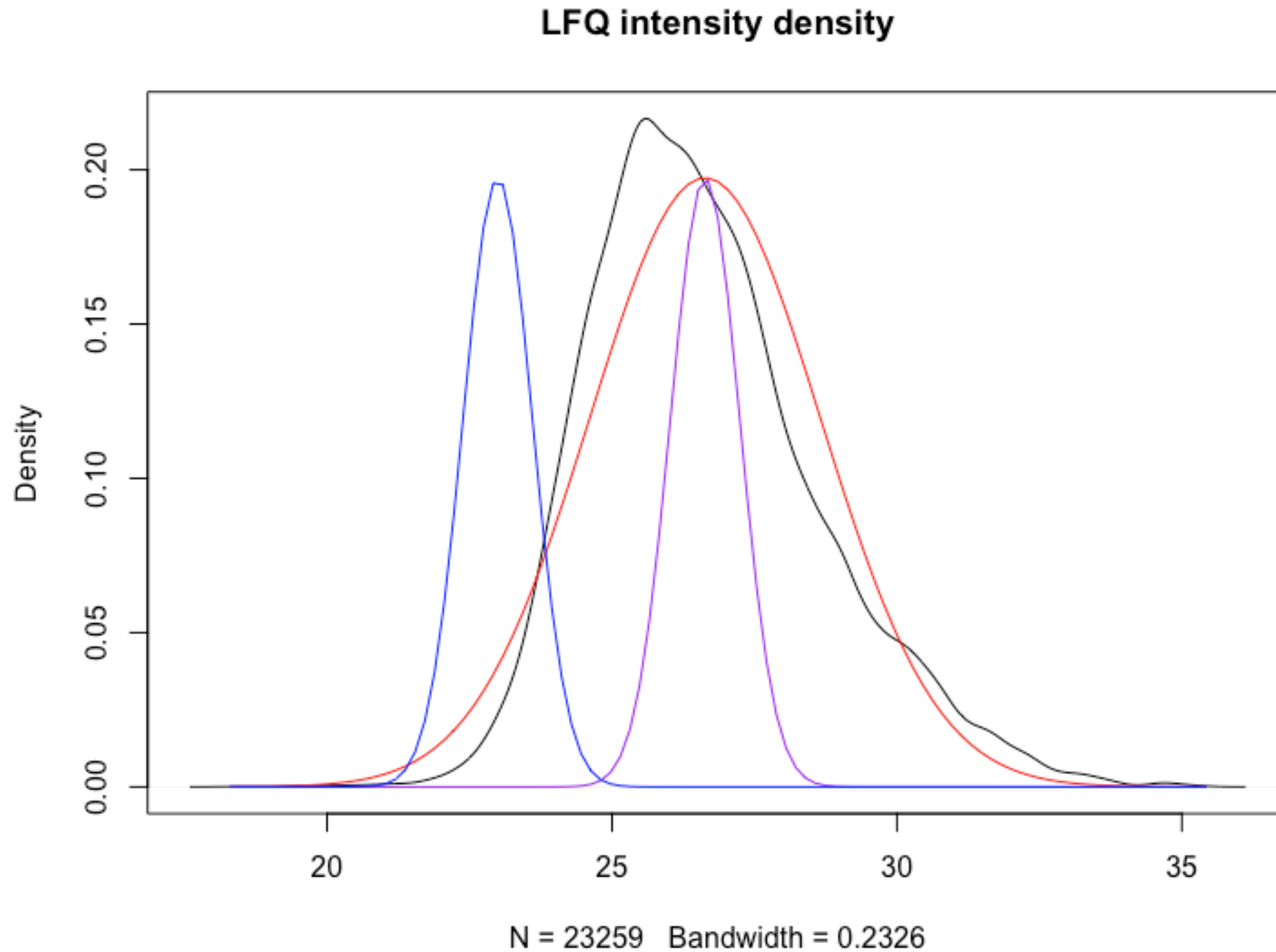


# Make the distribution narrow



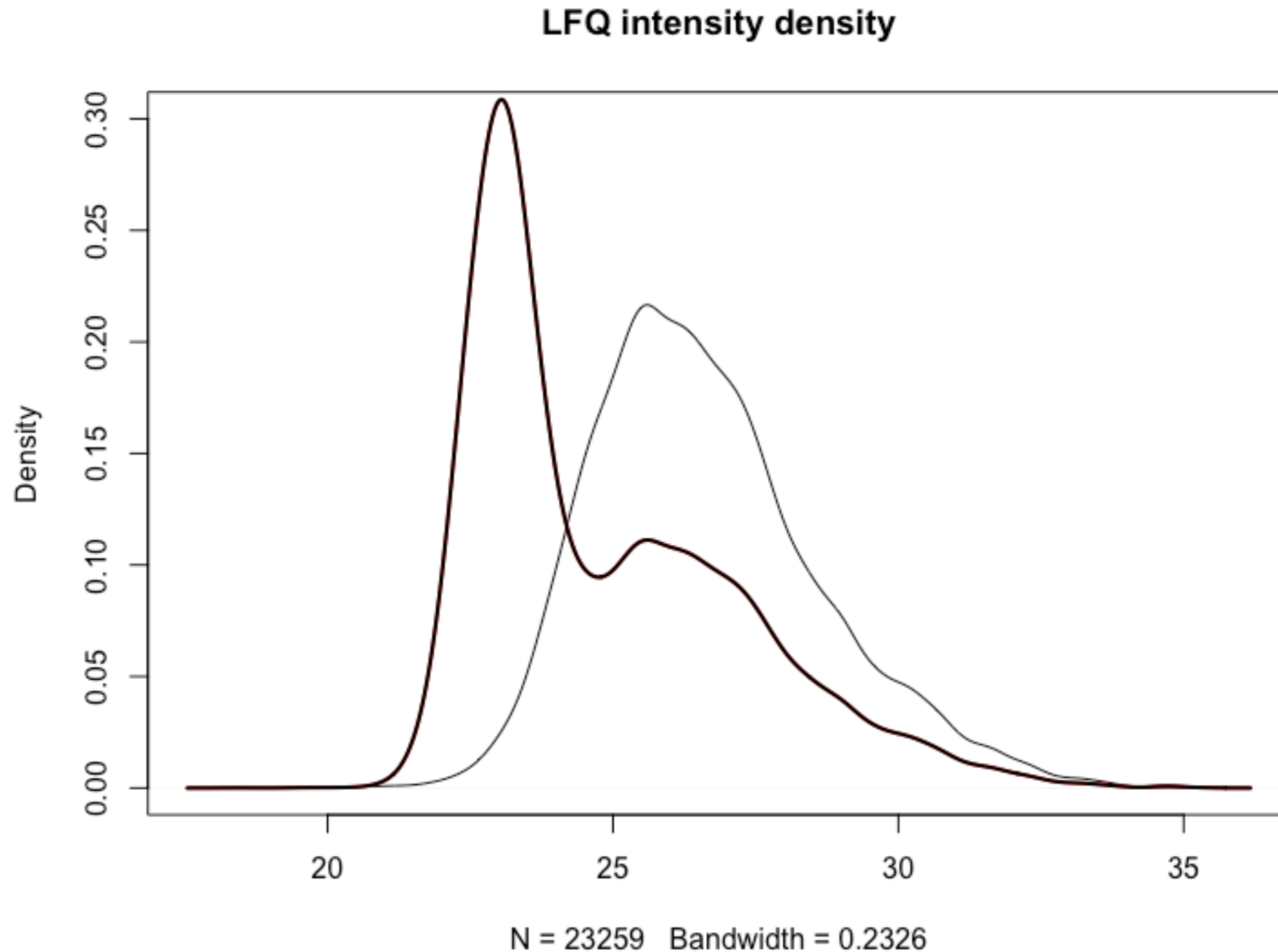


# And shift it **downwards**



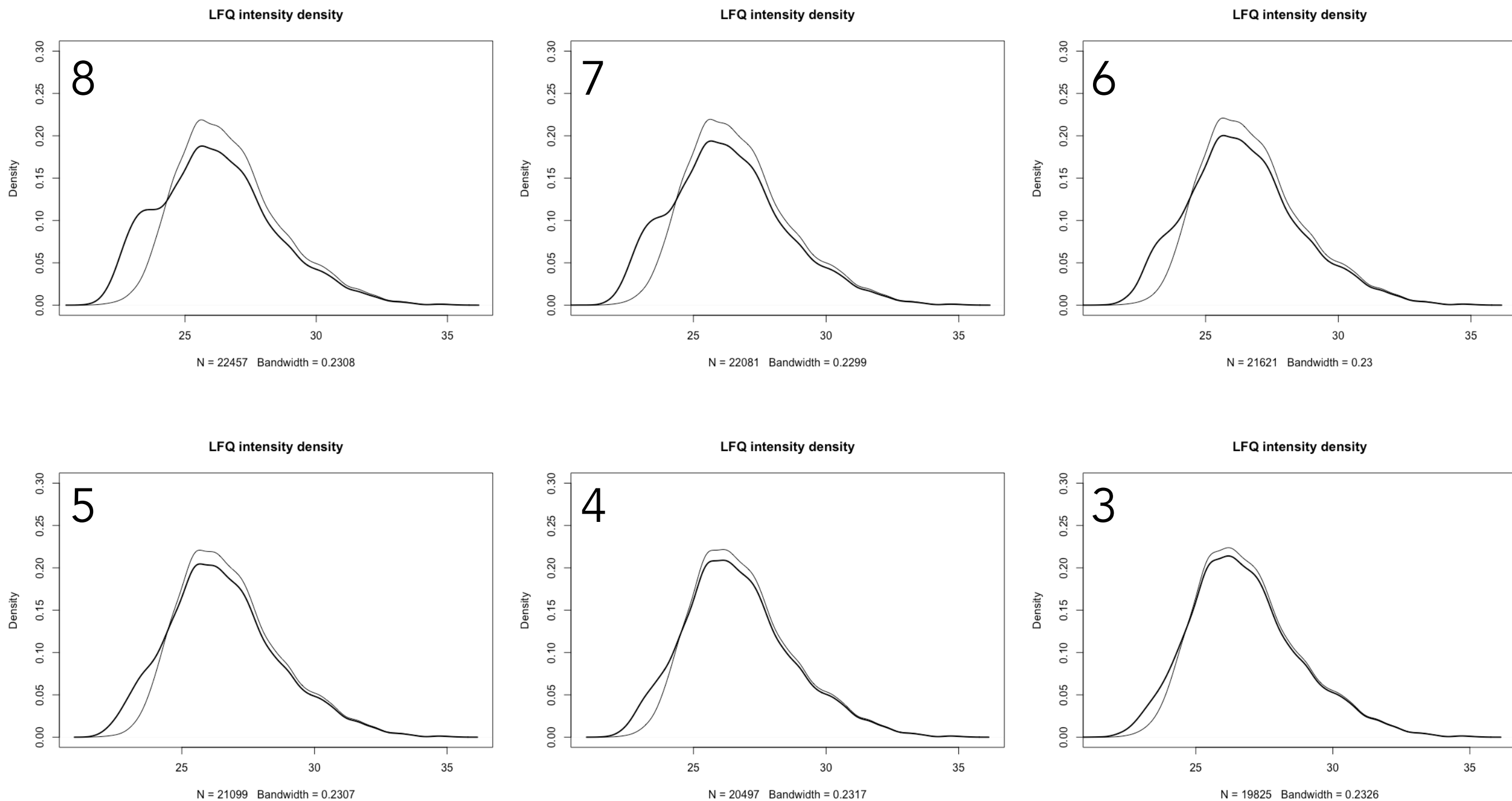
# The new distribution models low intensity values

# Take values from the new distribution



# Prefilter your data!

# Max x missing values allowed



**Stay in touch with  
your raw data!**

# Imputation guidelines

...according to my experiences

- 1 Try different parameters (filters, fittings).
- 2 Always keep the raw data in mind.  
Especially when interpreting your data.
- 3 Plot, plot, plot.

**Now let's do it in R!**



# Your tasks

Split in groups of two

Walk through the code for missing value imputation (`impute.R`)

Try to understand the lines

**modify the filtering for number NAs allowed**

**modify the width of the new distribution**

**<https://github.com/higsch/bioinfo-workshop>**

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