

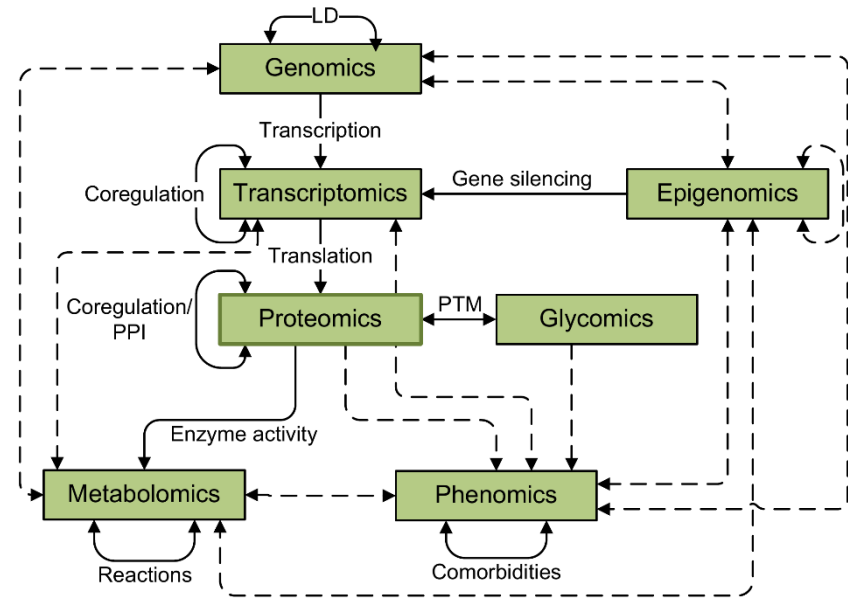
Part 1: overview

- Principal Component Analysis (PCA)
- Partial Least Squares (PLS)
 - Maximal covariance principle
- Two-way Orthogonal PLS (O2PLS)
- Post-hoc analyses using external databases



Background

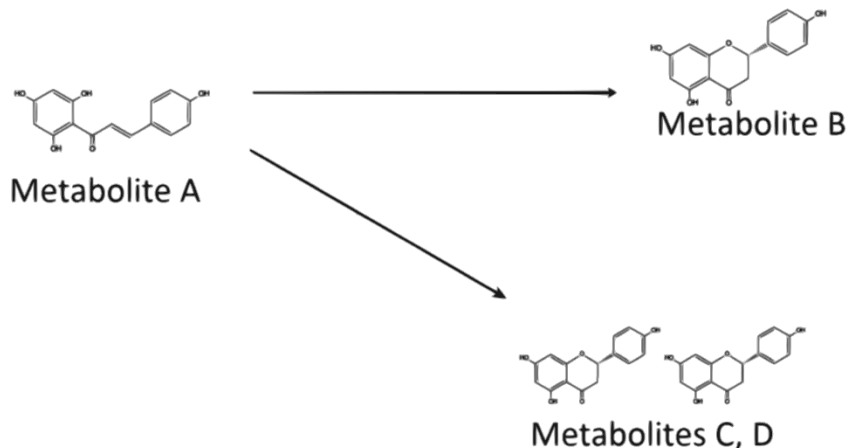
- Recent advances in technology provided many types of biological datasets (multi-omics data)
- Different levels of biological variation measured
- Need for integrative approaches: combine data and extract information



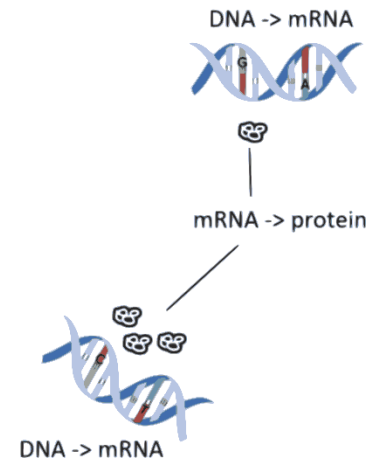
Integrative approach: aims

- How does variation between omics datasets relate?
- Which types of features induce this variation?
- Can we benefit from a joint/integrative analysis?

Metabolomics

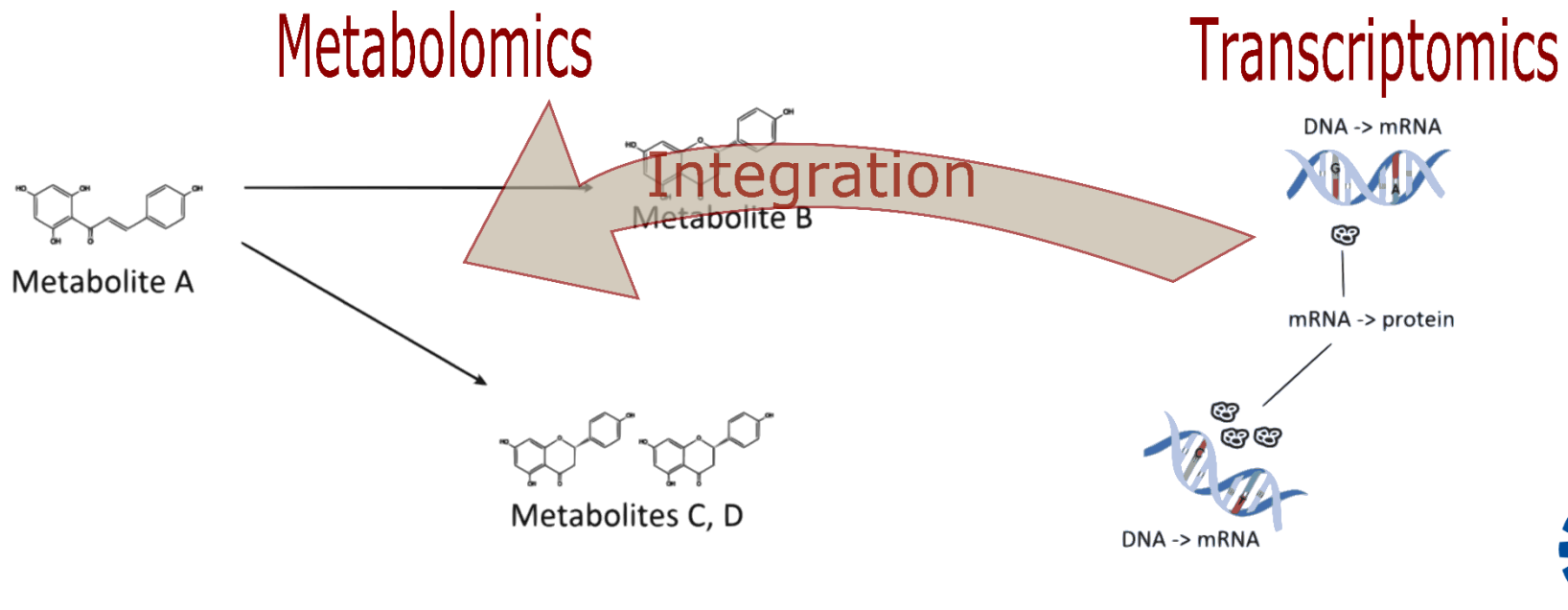


Transcriptomics



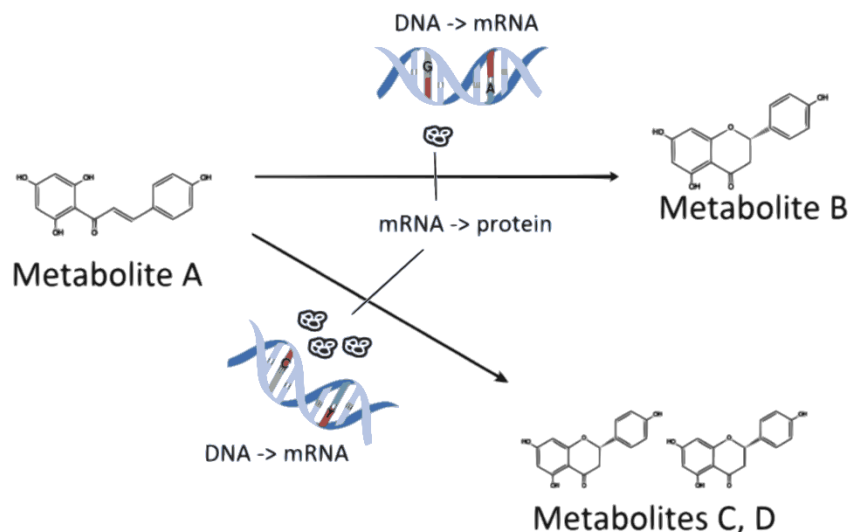
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Integrative approaches: means

There is typically

- High correlation *among* features
 - Genes are correlated
 - Metabolites are correlated
- Relation *between* features from two datasets
- Dimension reduction approach: few components drive association



Example: bivariate data

- Suppose we have two genes, and two metabolites x_1, x_2, y_1, y_2
- The variance of x_1 is larger than of x_2
- The variance of y_2 is larger than of y_1
- Only x_2 and y_1 are correlated
- Which variables will get high weight with PCA? Why?
- Which variables should get high weight when you look at the relation between x and y ?



Partial Least Squares (PLS)

- Let X and Y be two data matrices
 - Size: N times p and q , respectively
 - p and q can be very large
- Recall: in PCA, variance is maximized
- We are interested in the **c**ovariation between X and Y



Partial Least Squares (PLS)

- Objective: maximize covariance between both linear combinations
- The combinations are again relative
- Can be calculated using
 - A singular value decomposition (*svd* in R)
- Similar interpretation as PCA, except that we focus on covariance



Example: PLS

```
library(OmicsPLS)
```

```
gene1 <- rnorm(100)
```

```
gene2 <- rnorm(100,sd=0.75)
```

```
metab1 <- rnorm(100)
```

```
metab2 <- gene2
```

```
X <- cbind(gene1, gene2)
```

```
Y <- cbind(metab1, metab2)
```

```
svd(X,0,1)$v
```

```
o2m(X, Y, 1, 0, 0)$W.
```

These are the weights for PCA

```
      [,1]  
[1,] -0.9882865  
[2,] -0.1526100
```

These are the weights for PLS

Data is not centered, proceeding...

```
      [,1]  
gene1 0.02574932  
gene2 0.99966843
```



Partial Least Squares: summary

- For given datasets X and Y , we want to inspect their relation
- We consider directions of maximal covariance
- This direction is represented by weights for each feature, calculated by a singular value decomposition
- The projections of the data onto these weights are called the scores: $t = Xw$ and $u = Yc$
- One can interpret or plot the weights and scores to understand which features/samples are most important



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