GBIO0015

# Homework 1

## A tour in genetic epidemiology

### **Important dates:**

• Presentation date: 10 April 2017

#### Mark:

• 25 points

#### **Instruction:**

- Read Bessonov et al. (2015) paper and take Evans et al. (2013) as supporting document to address the following questions in your slides presentation.
- You can work in groups. One presentation per group is allowed (but everyone should present a part) and a score will be a group score.
- Presentation duration: 20min.
- Reference papers could be found from:

# http://www.montefiore.ulg.ac.be/~chaichoompu/CK/?Courses

### Questions

- 1) What is the study design:
  - a. Samples: unrelated individuals? Cohort? Case-control? Family? Would that make a difference for the subsequent analyses?
  - b. Markers: which markers are used? How many?
- 2) How were dependencies (LD) between markers treated? This relates to "pruning". Why is this important?
- 3) How were markers coded in analytic models? Does this matter in terms of final epistasis results?
- 4) Which analytic framework / model was used? Regression? Parametric or non-parametric approach?
- 5) How was multiple testing dealt with?
- 6) What can be used as prior biological information? Is this information also relevant post analysis?
- 7) Which measures were taken to strengthen the findings?
  - a. Does agreement between different analytic tools imply validity of findings?
  - b. Replication?
  - c. Validation?

Address all of the questions and work out in more details, using the aforementioned papers or information you can retrieve from:

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