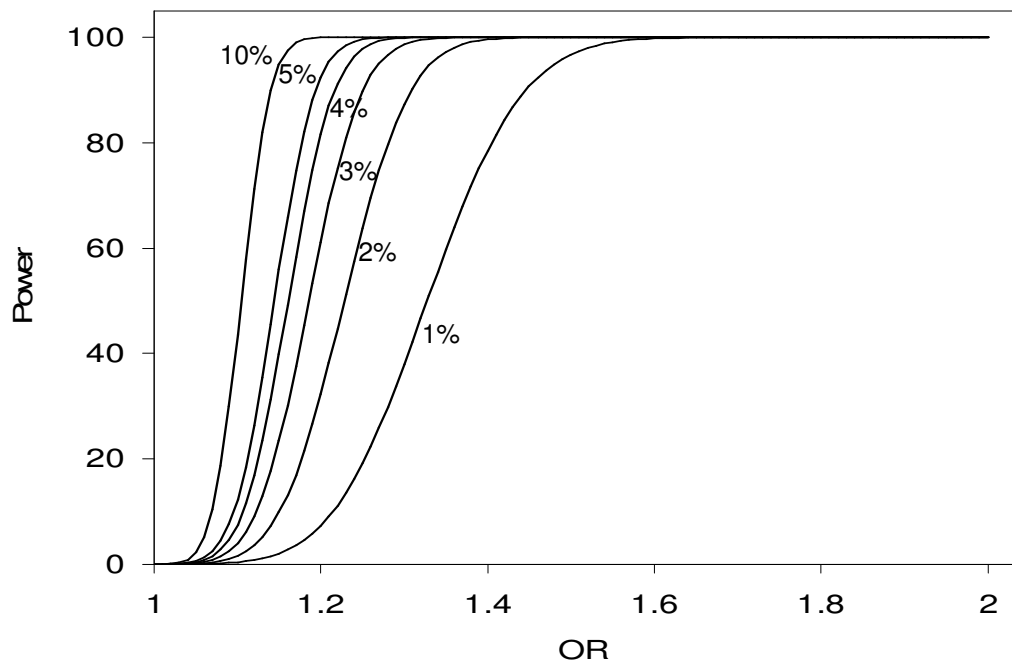


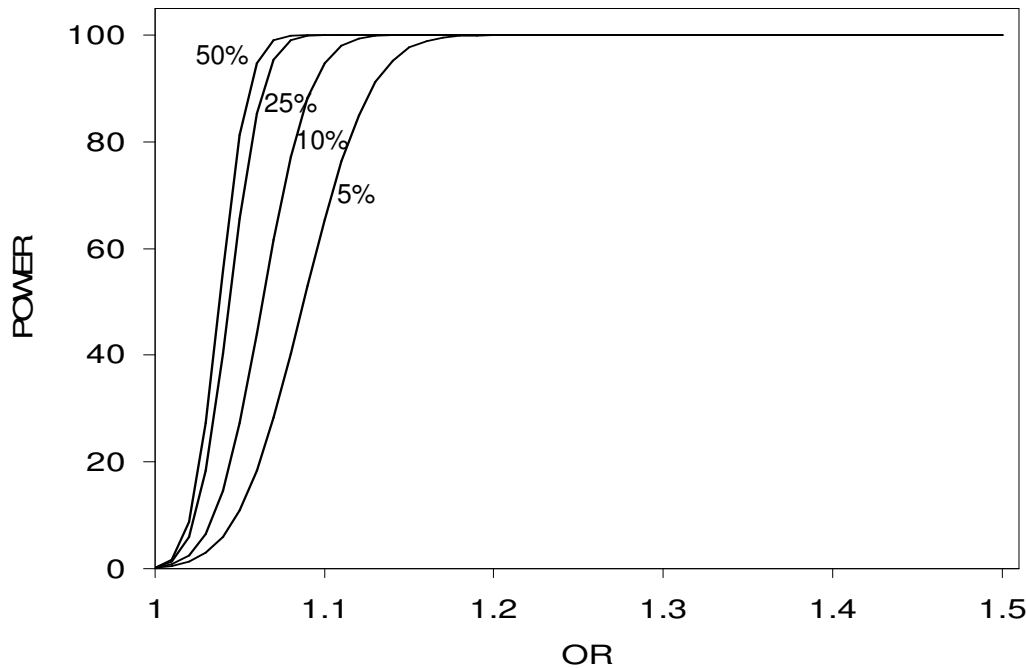
Figure S1. Power to detect associated variants in discovery and replication stages.

a) Discovery stage meta-analysis.



Power to detect an association with $\alpha = 10^{-4}$ (two-sided) assuming a per-allele effect and a discovery stage study size of 11,202 coronary disease cases and 30,733 controls (equivalent to the European studies in the discovery stage) across a range of minor allele frequencies (1%,2%,3%,4%,5%,10%). These power calculations assume that there is no between-study heterogeneity.

b) Replication stage meta-analysis.



Power to detect an association with $\alpha = 1.9 \times 10^{-3}$ (one-sided) assuming a per-allele effect and a replication stage study size of 17,121 coronary disease cases and 40,473 controls (equivalent to the whole replication stage) range of minor allele frequencies (5%,10%,25%,50%). These power calculations assume that there is no between-study heterogeneity.