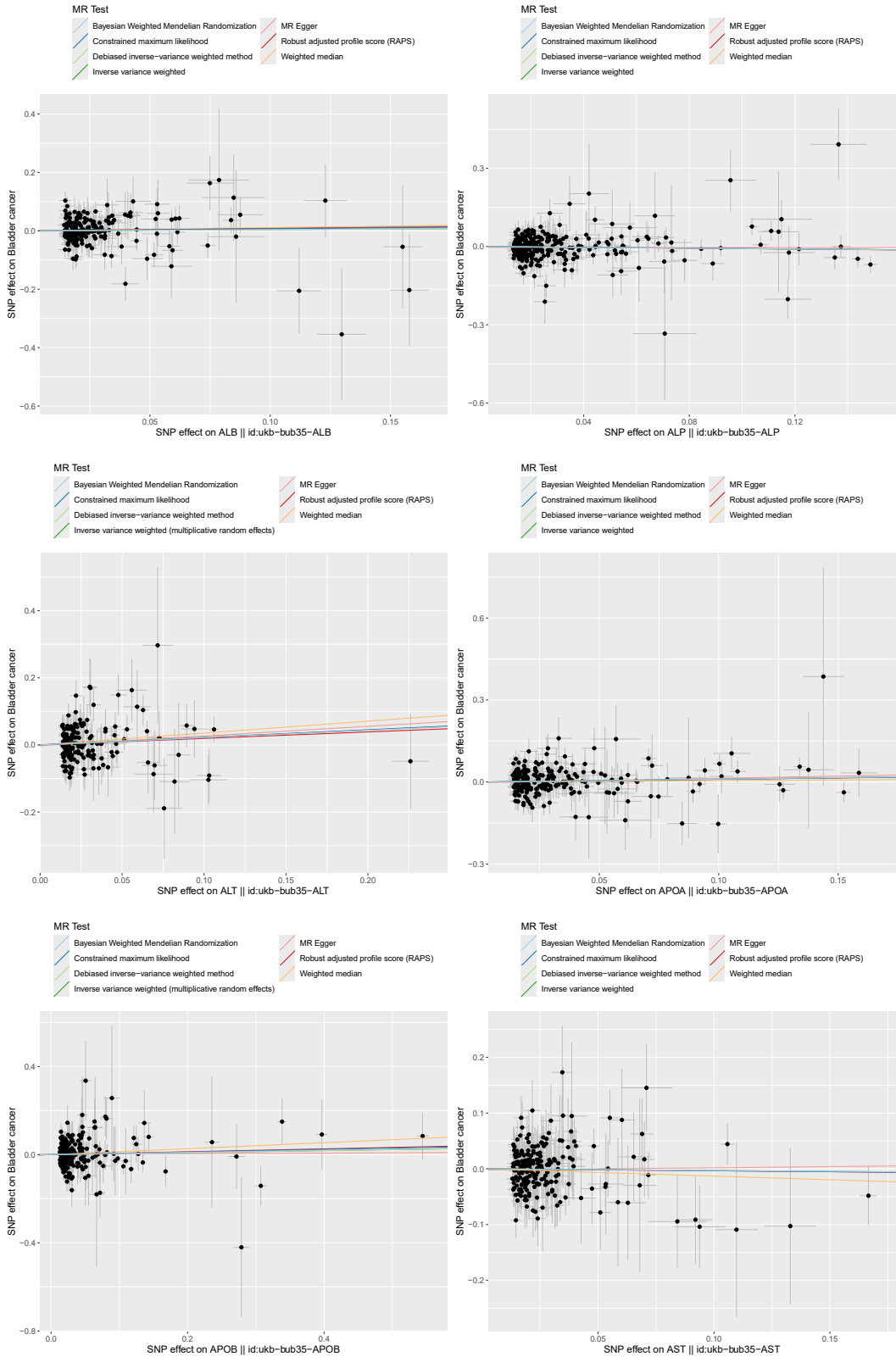
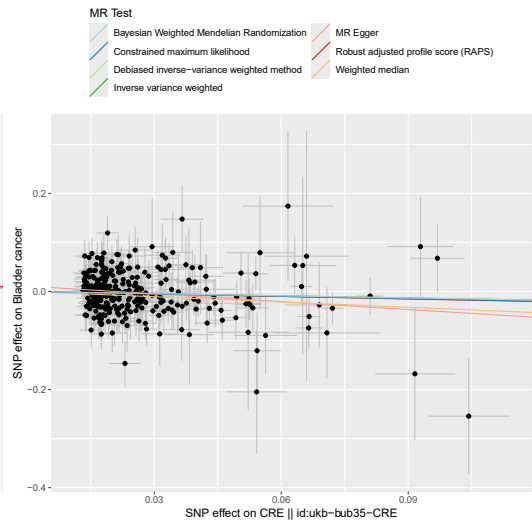
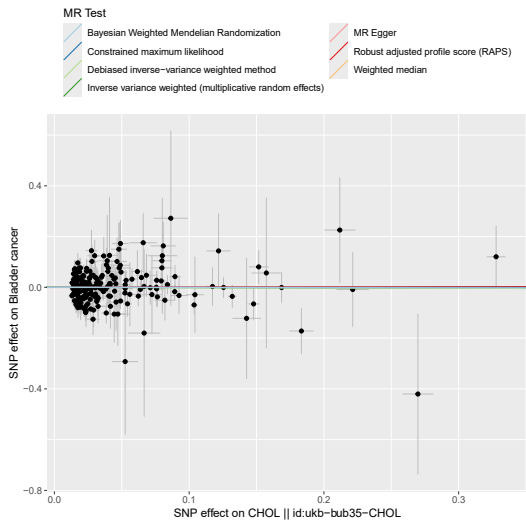
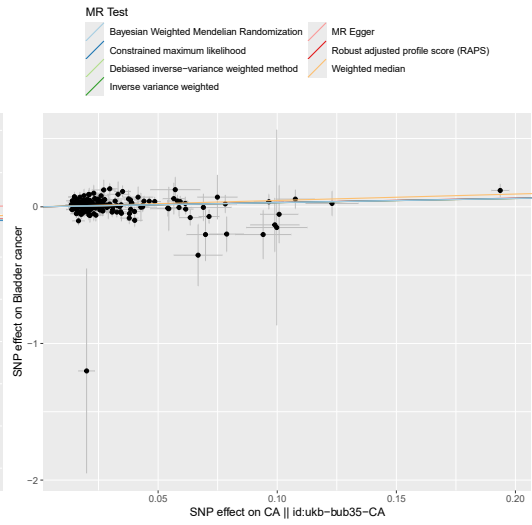
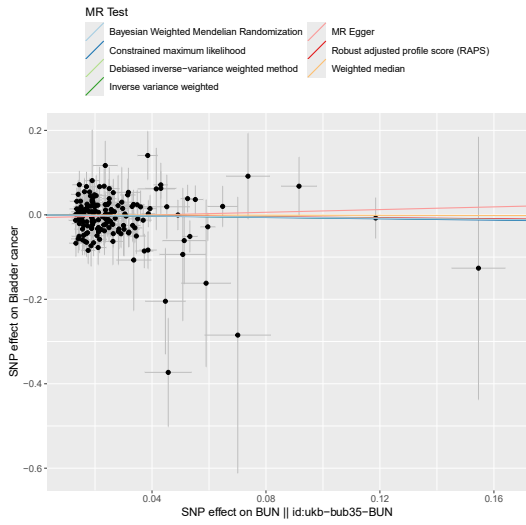
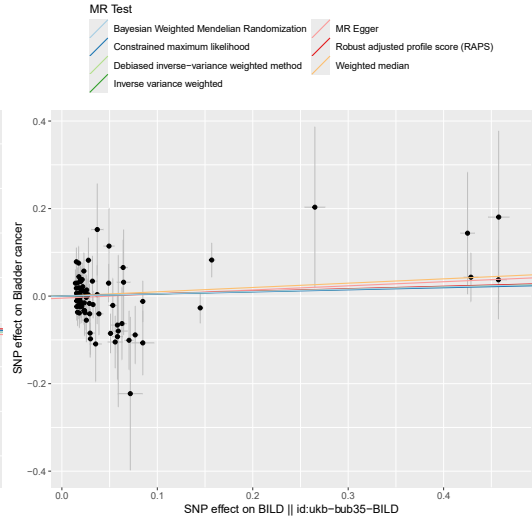
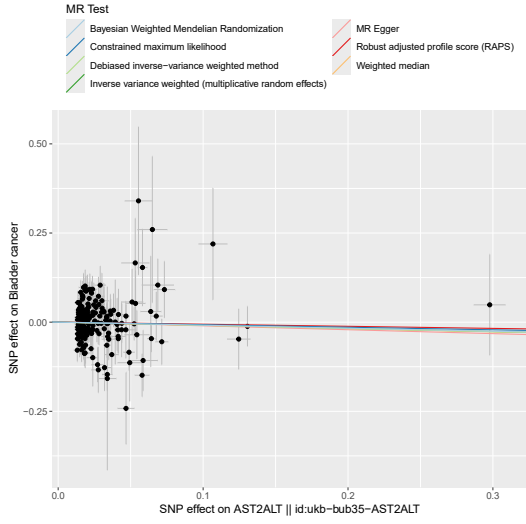
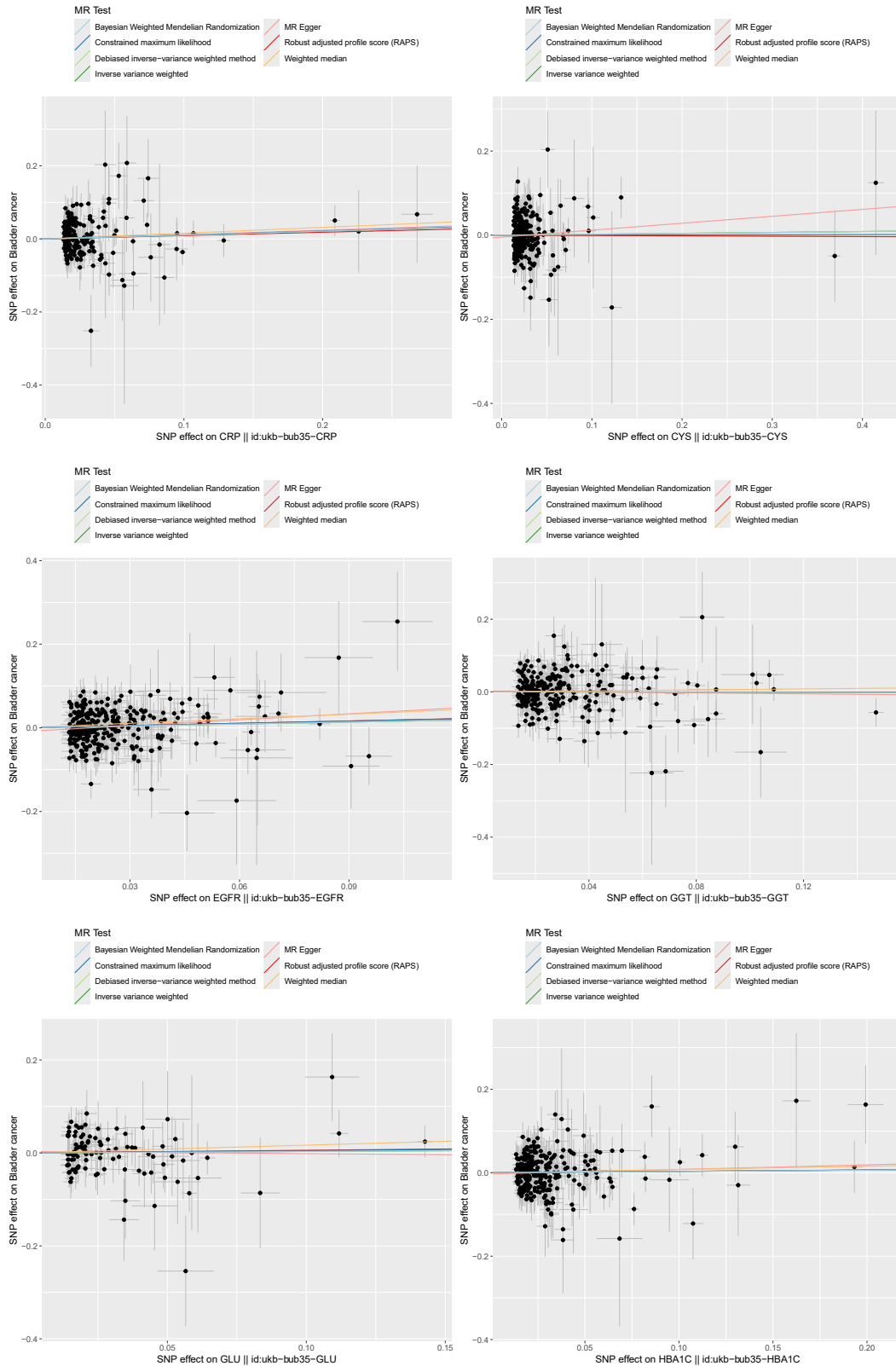
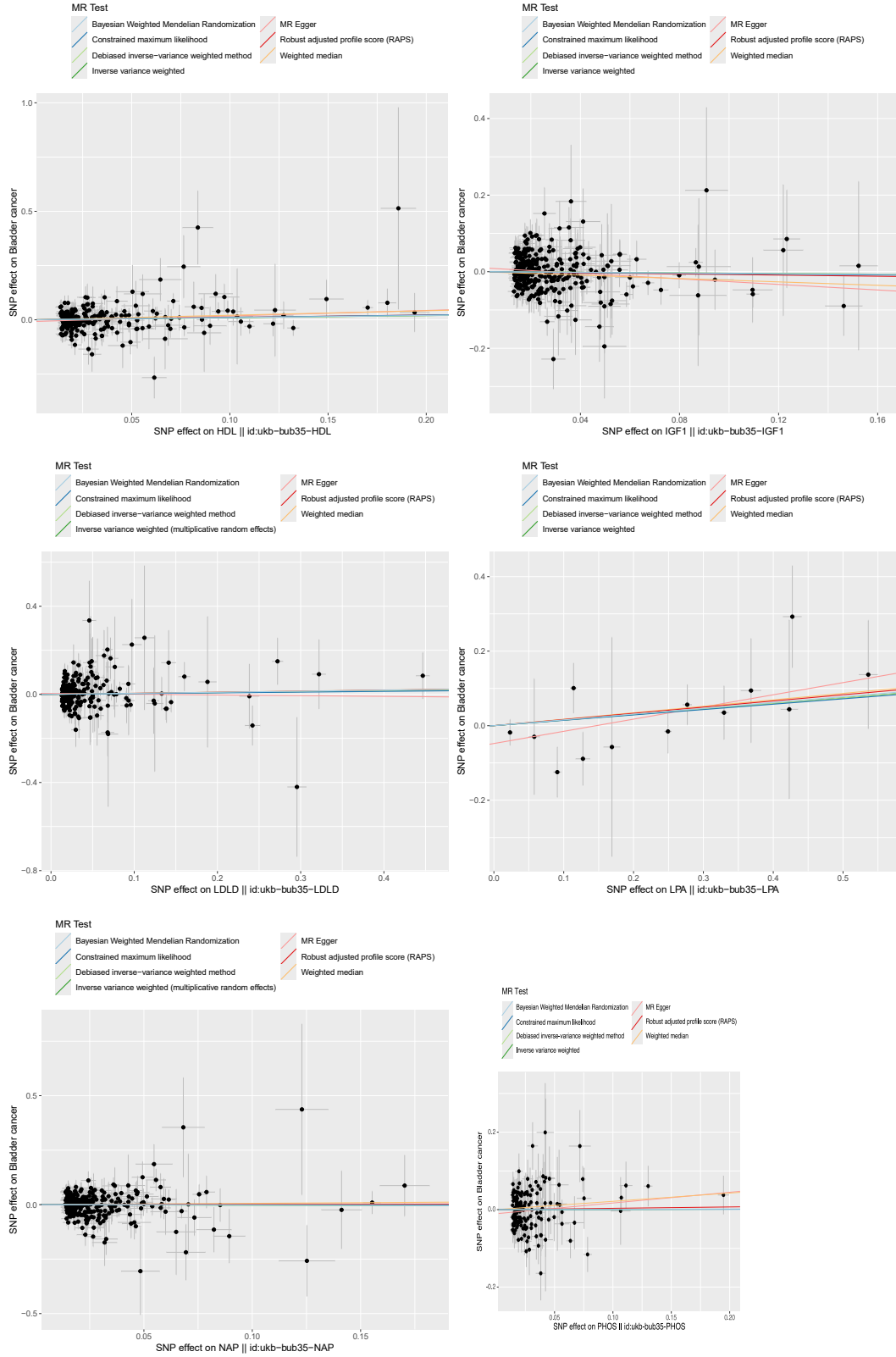


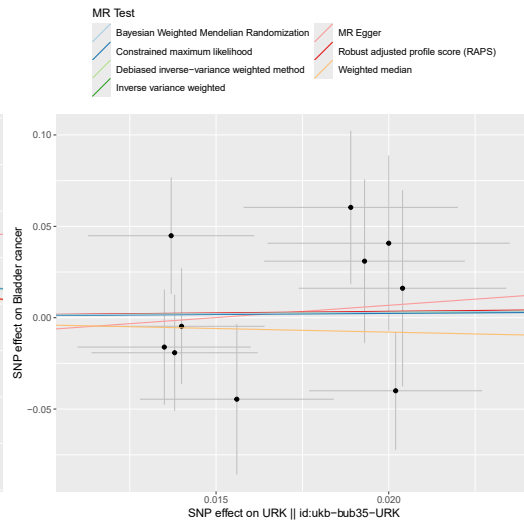
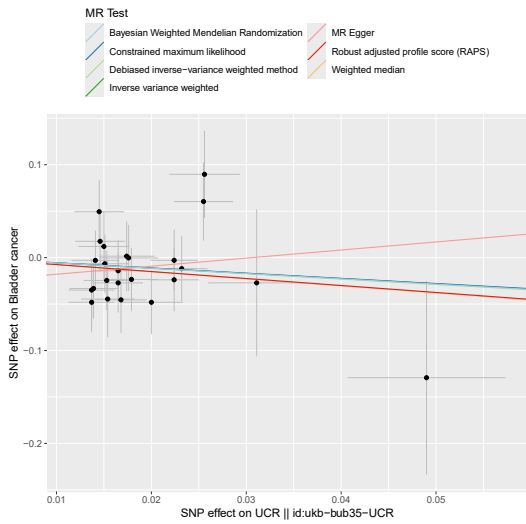
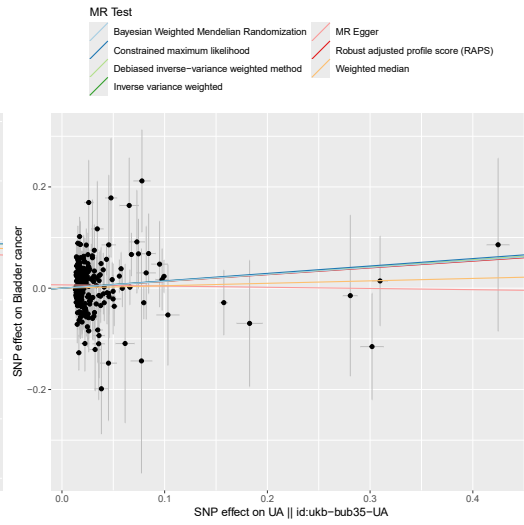
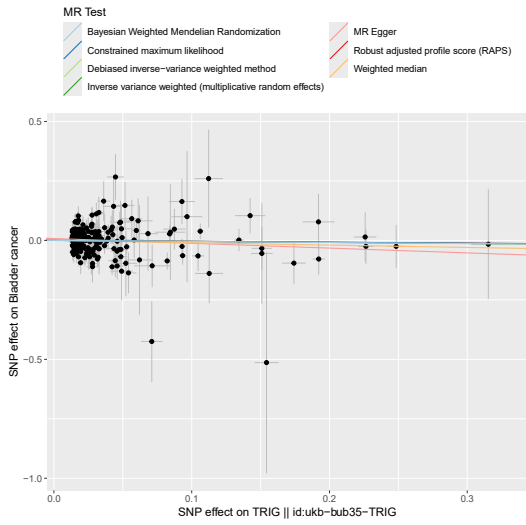
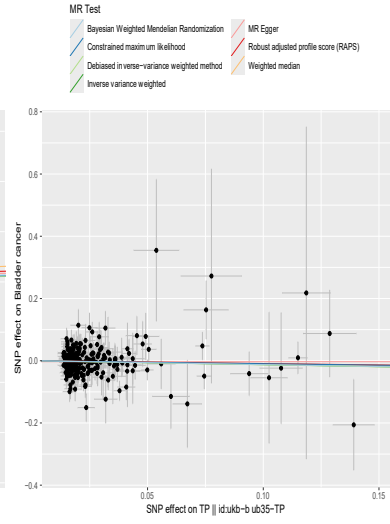
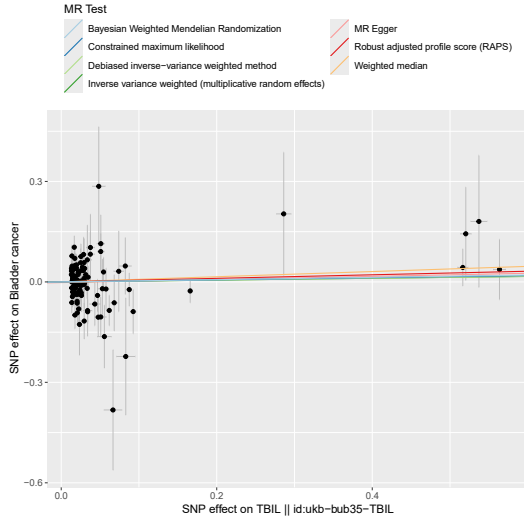
Scatter plot of Bca

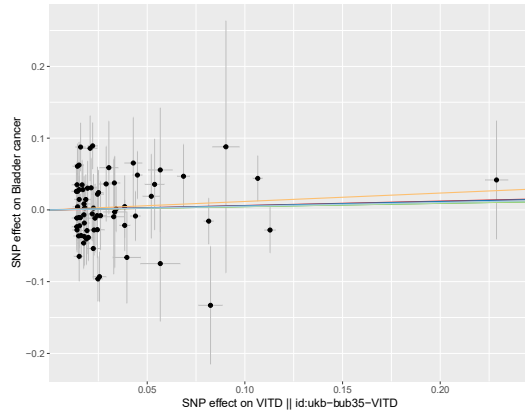
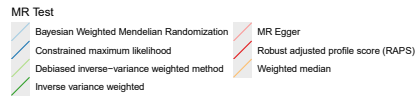
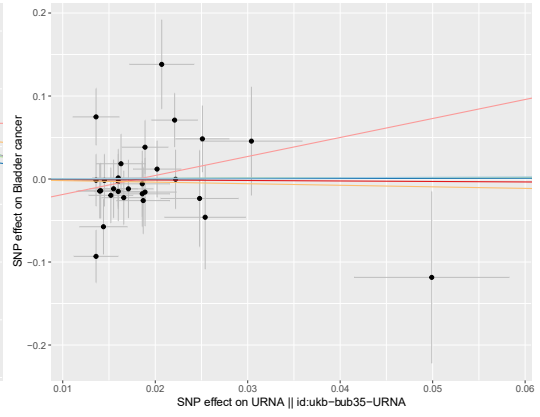
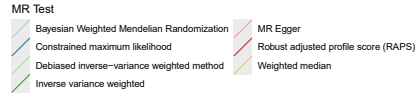
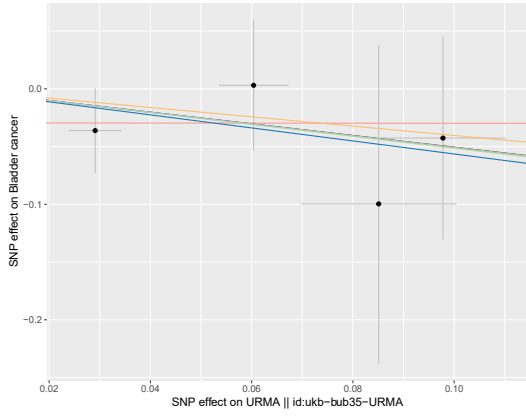
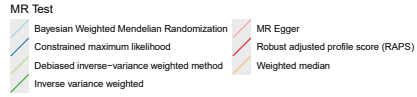




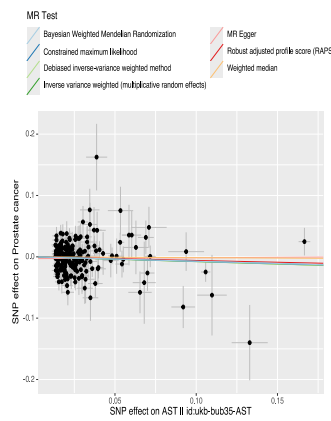
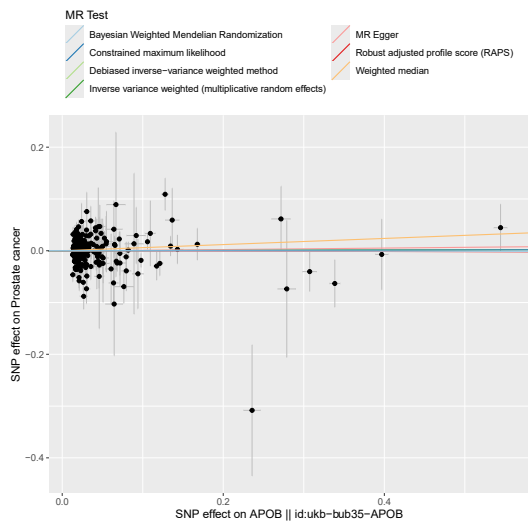
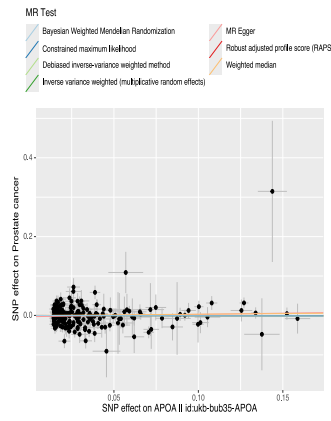
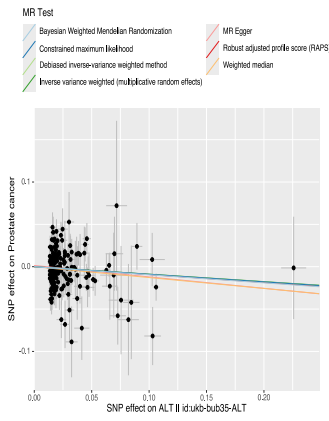
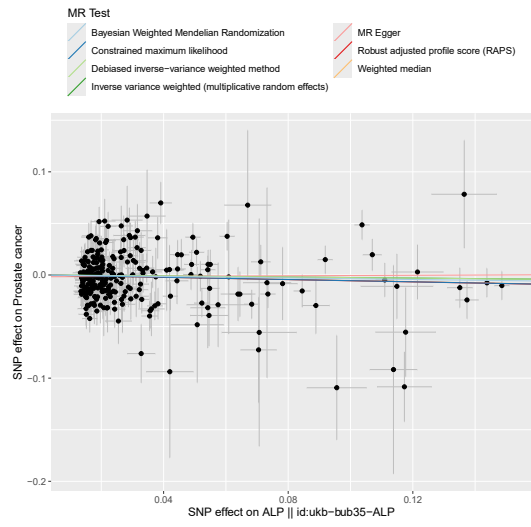
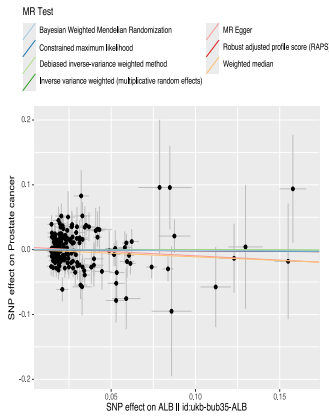


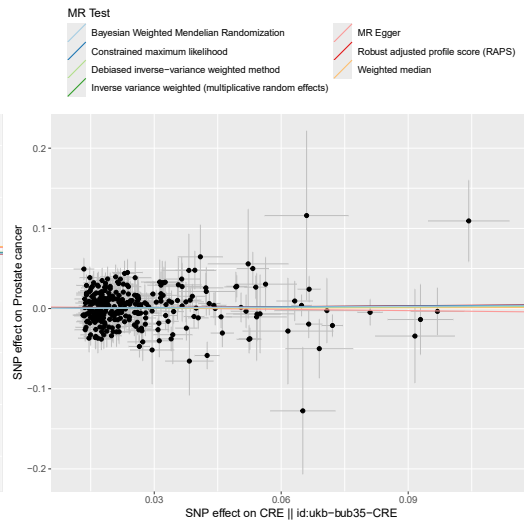
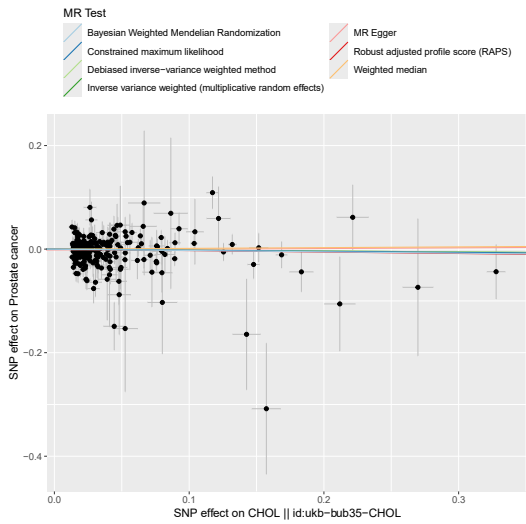
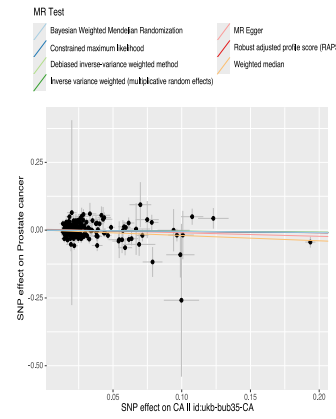
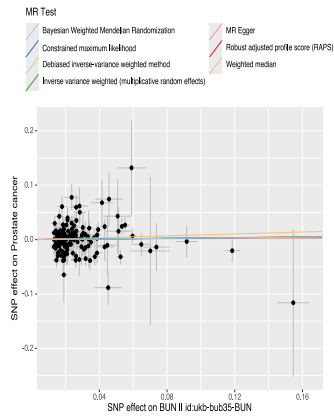
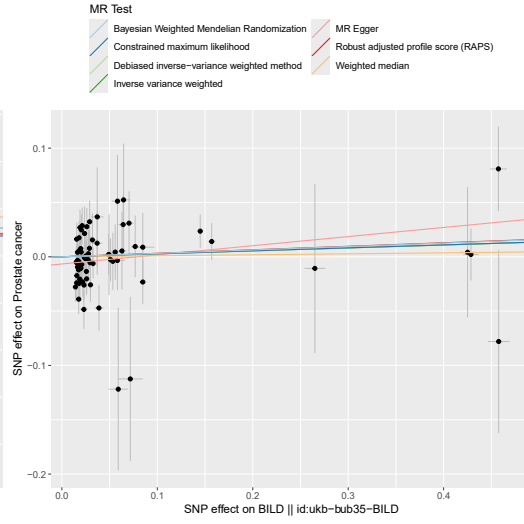
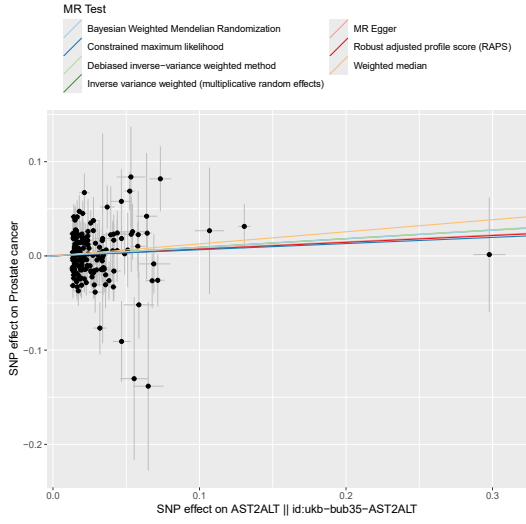


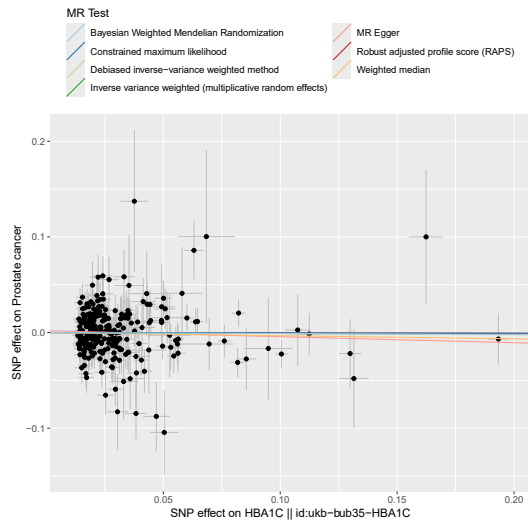
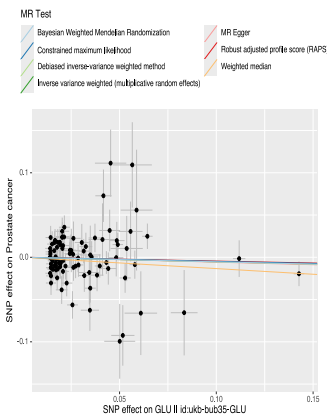
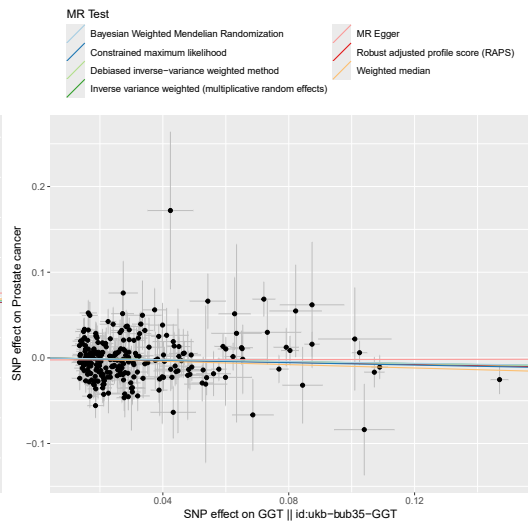
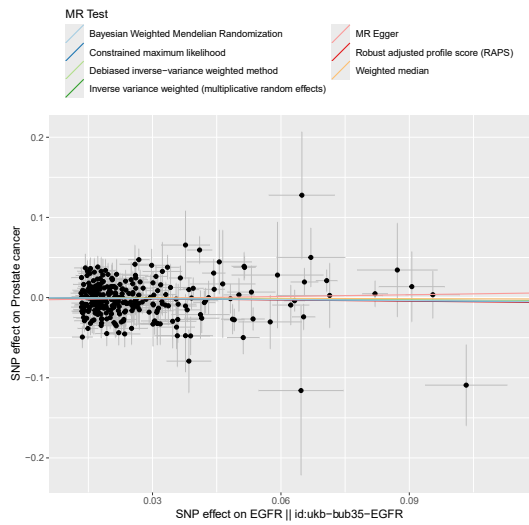
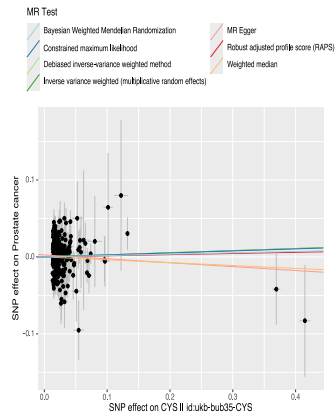
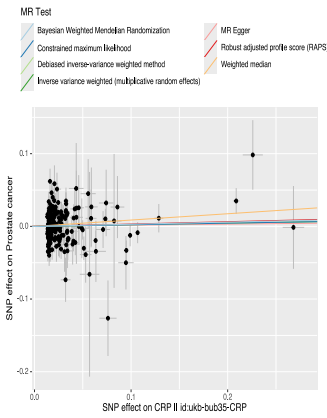


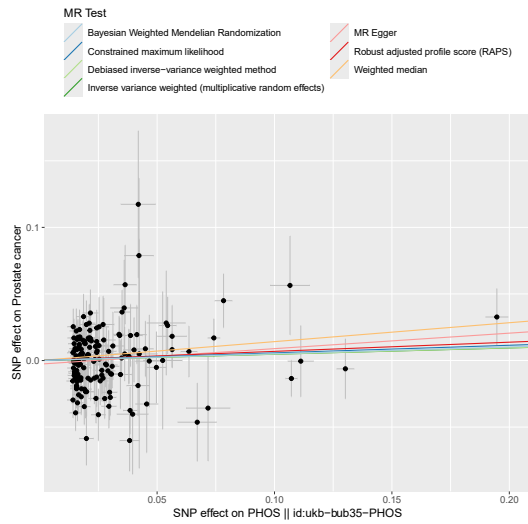
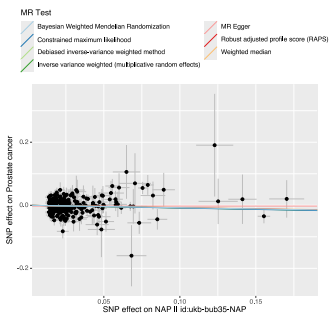
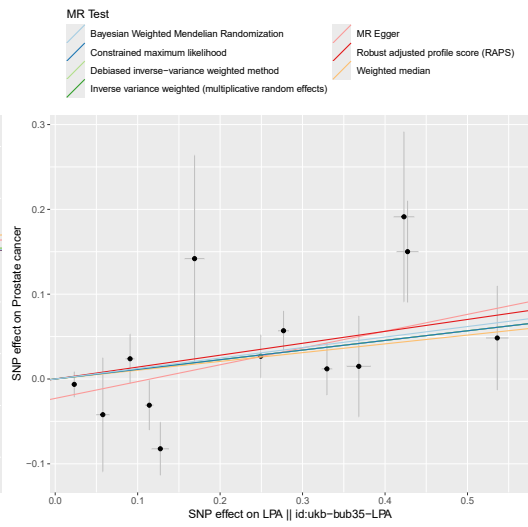
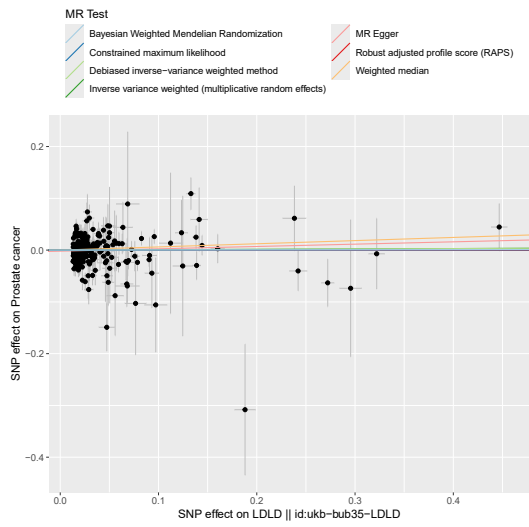
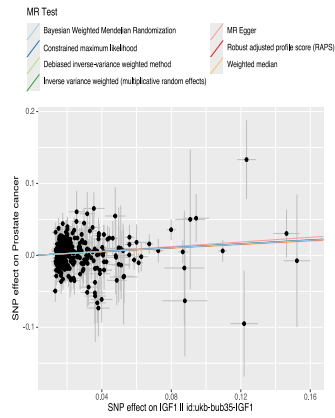
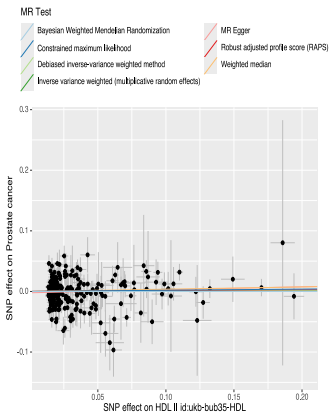


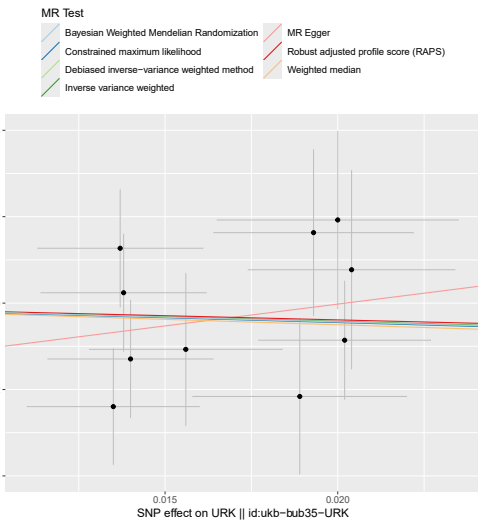
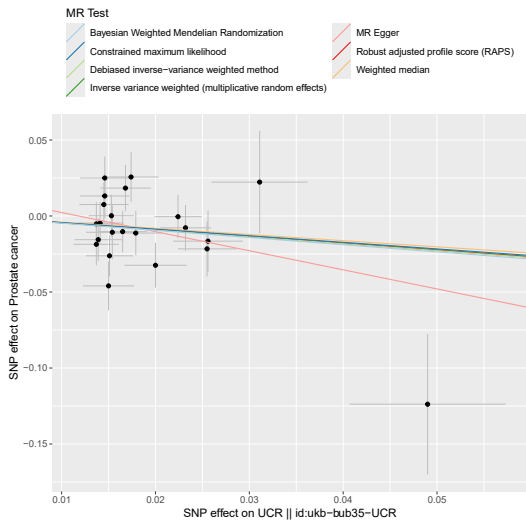
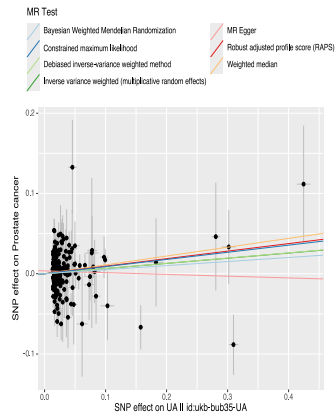
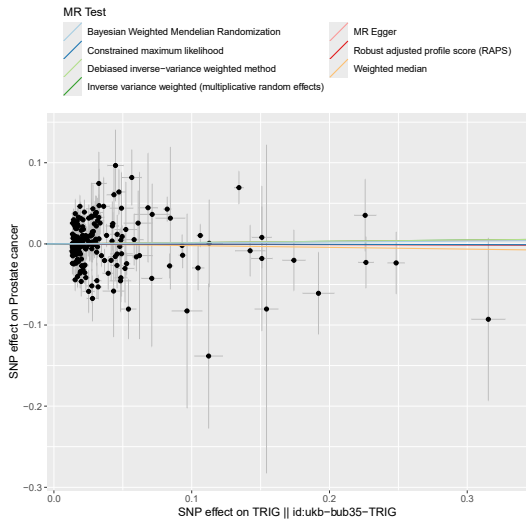
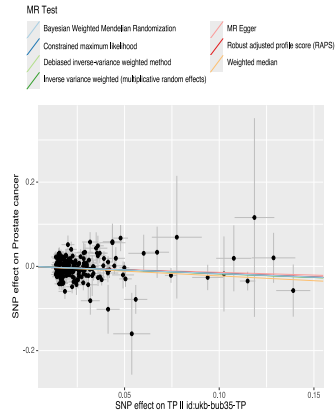
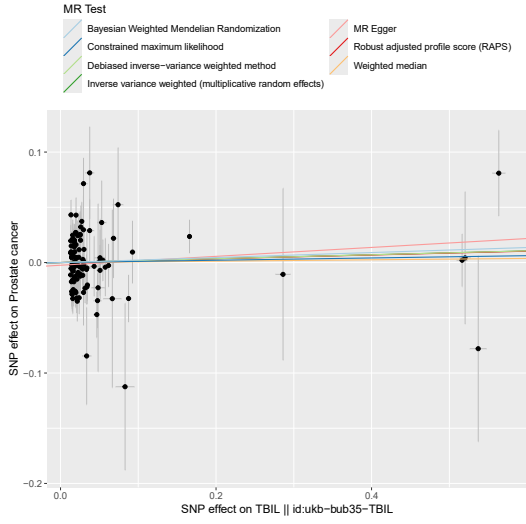
Scatter plot of PCa

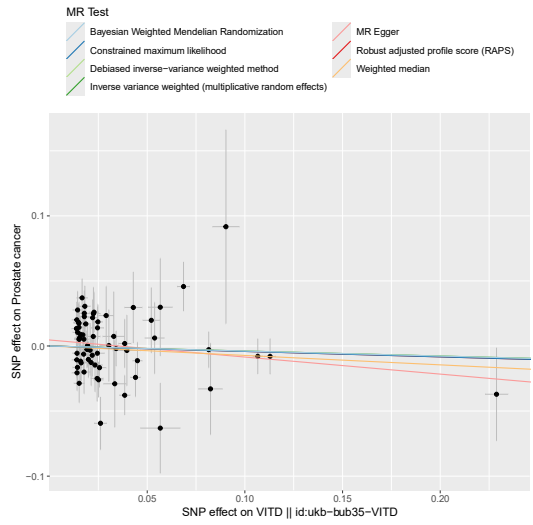
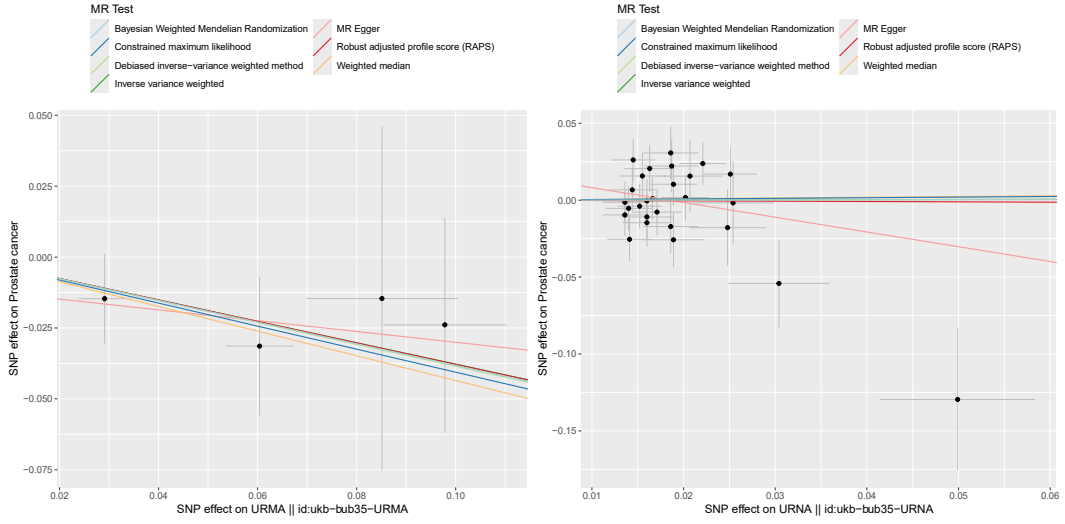




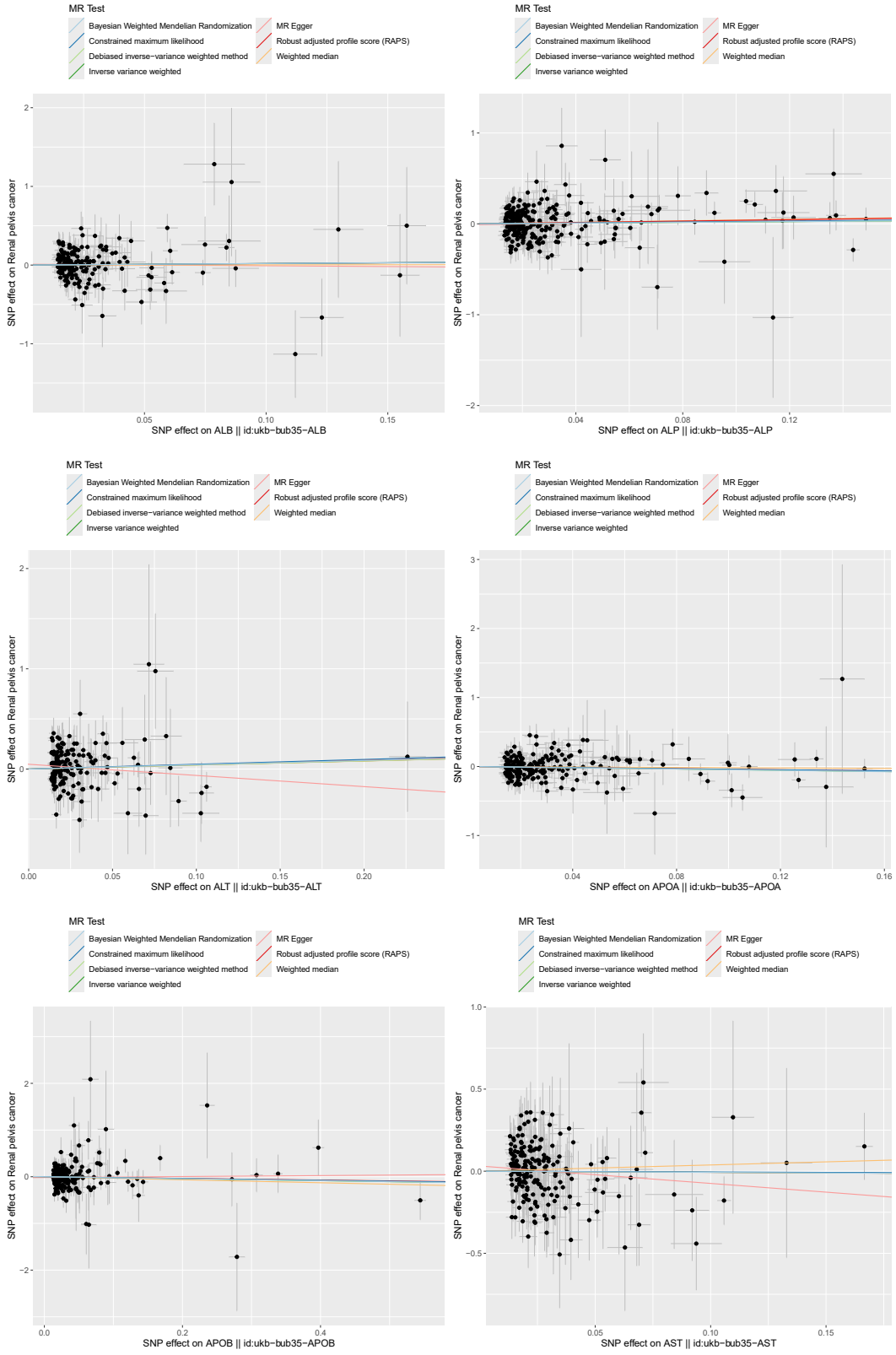


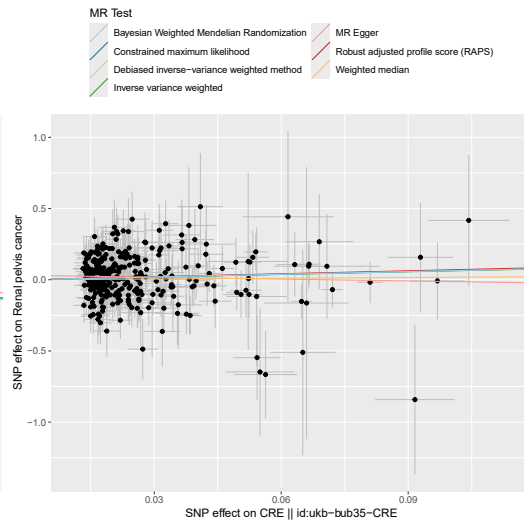
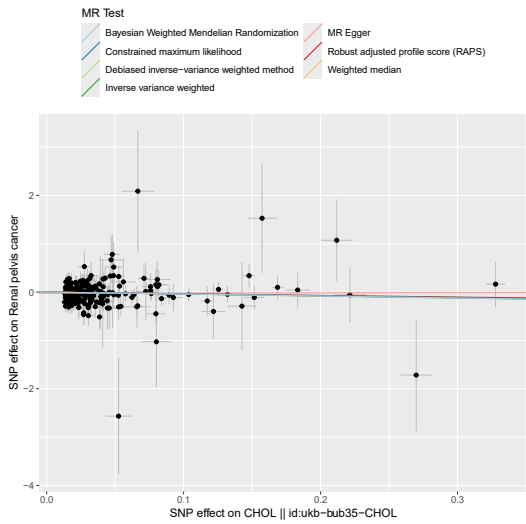
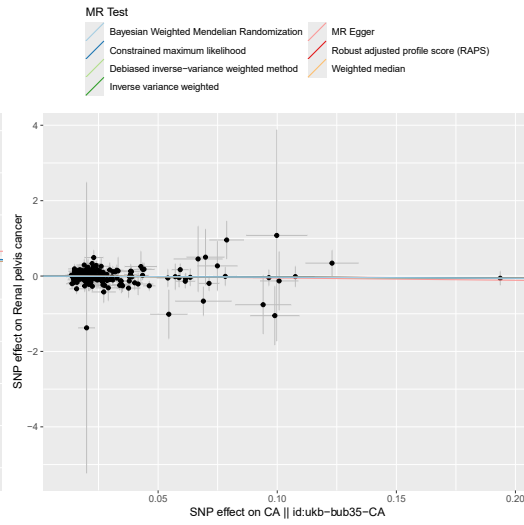
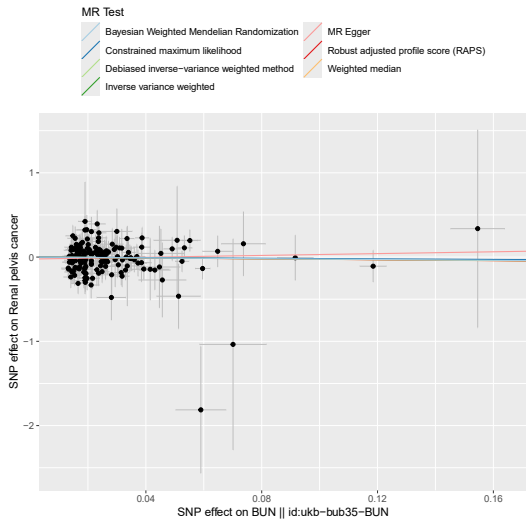
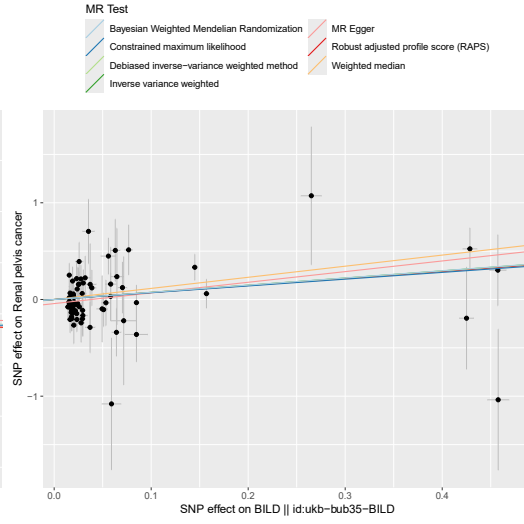
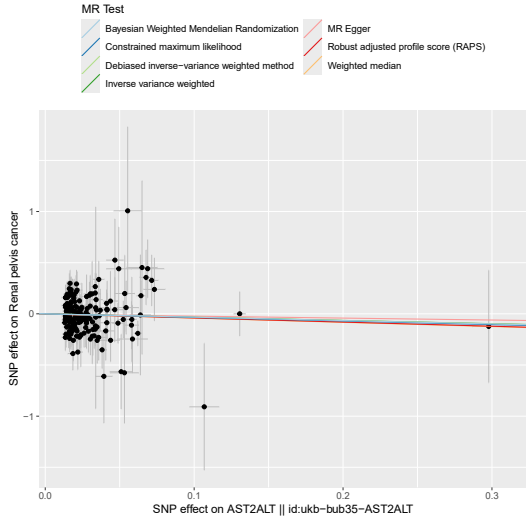


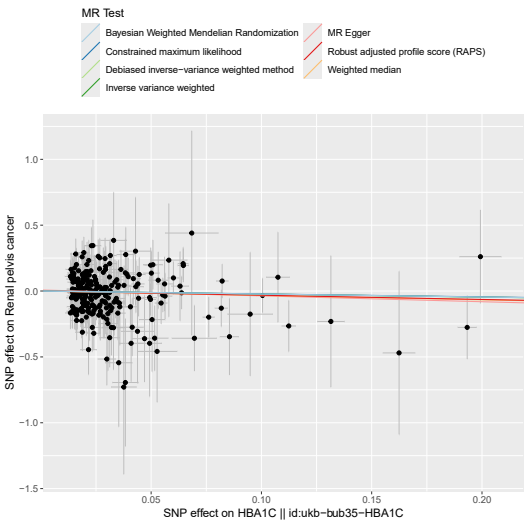
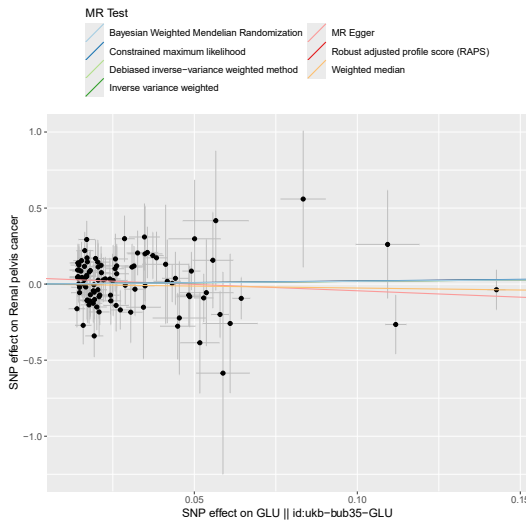
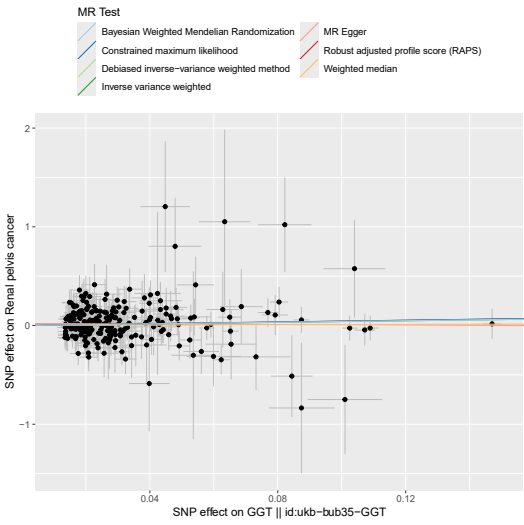
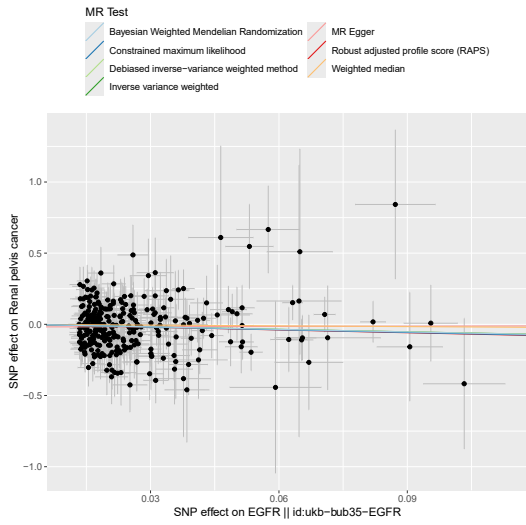
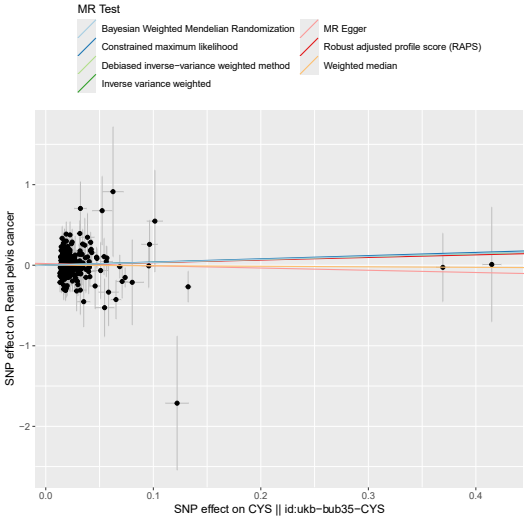
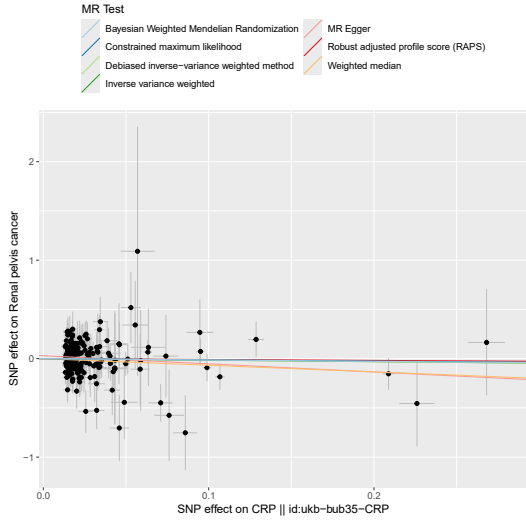


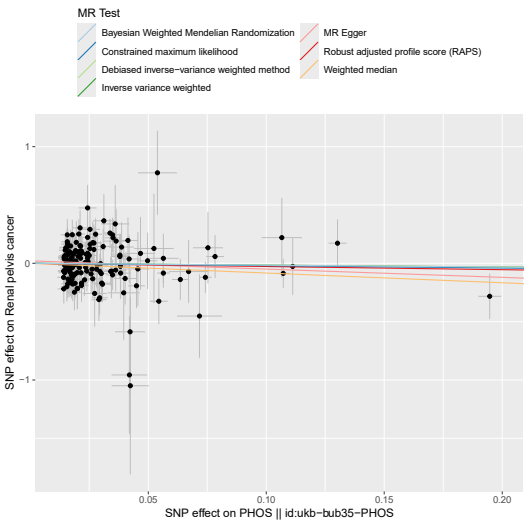
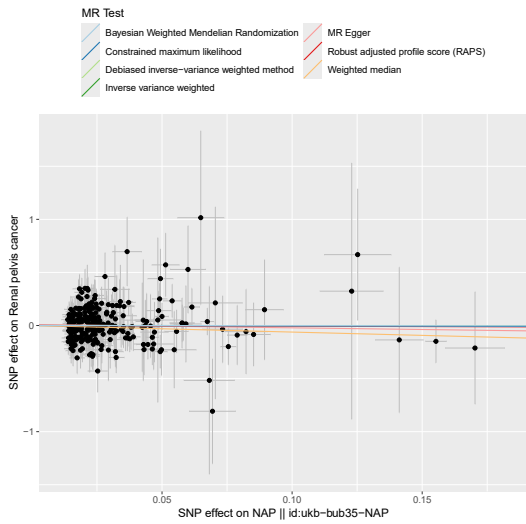
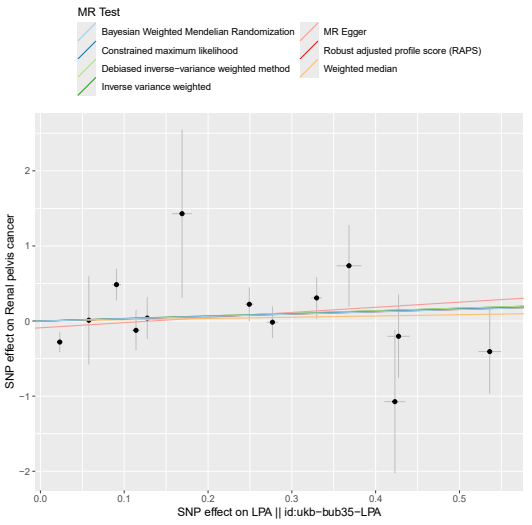
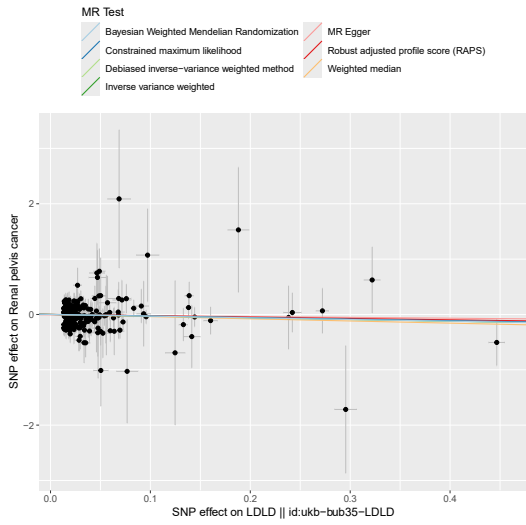
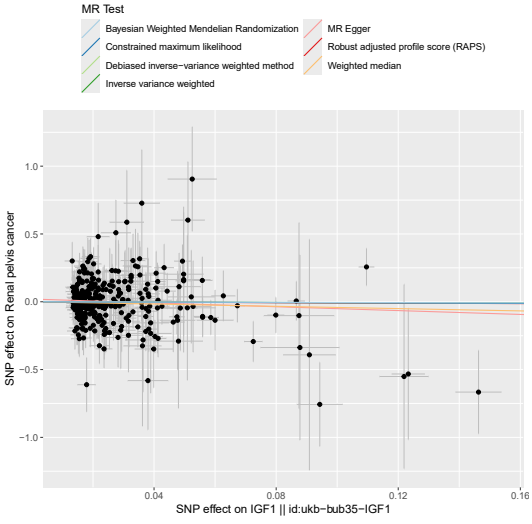
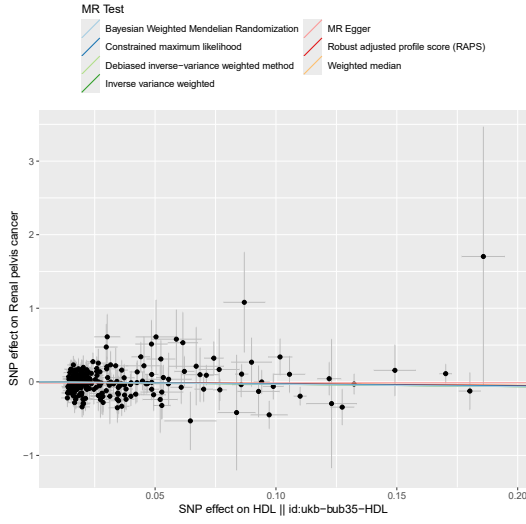


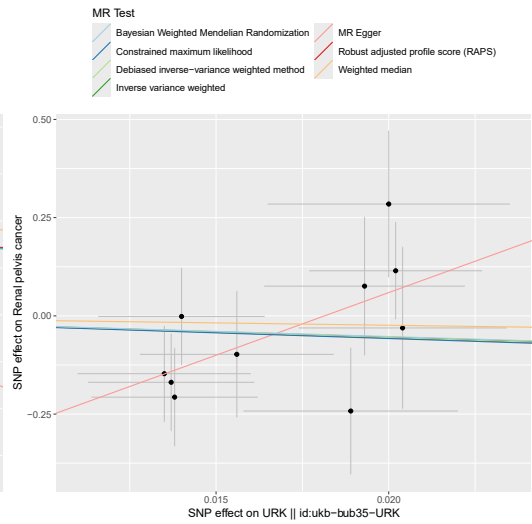
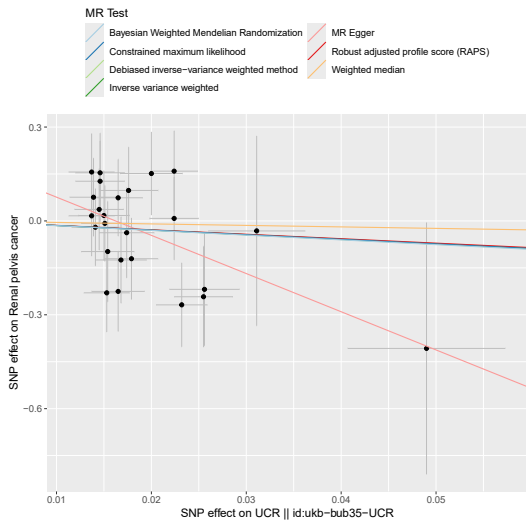
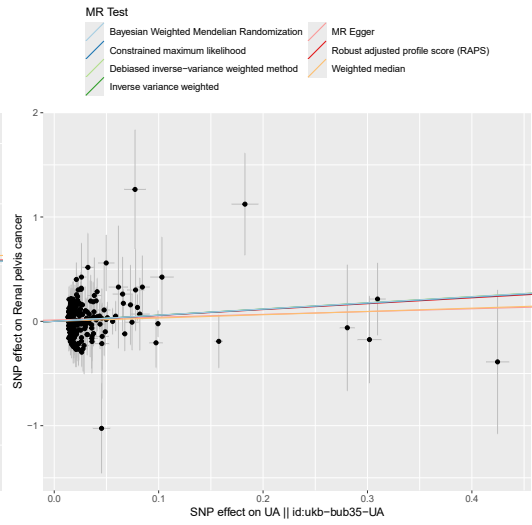
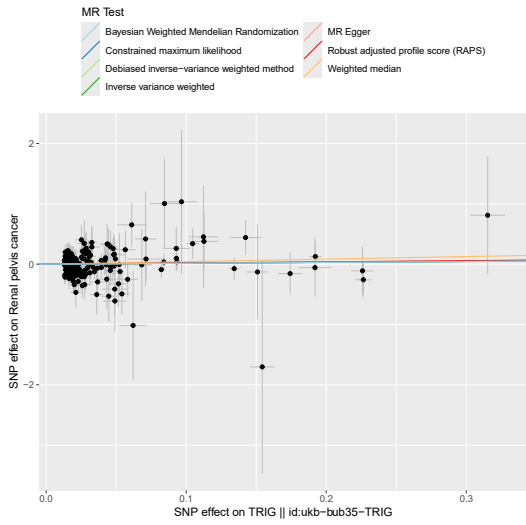
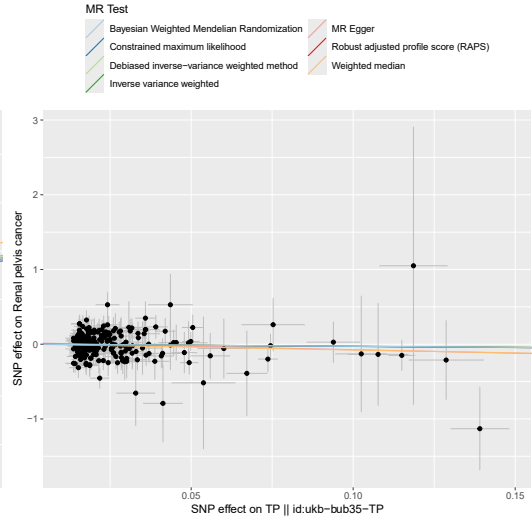
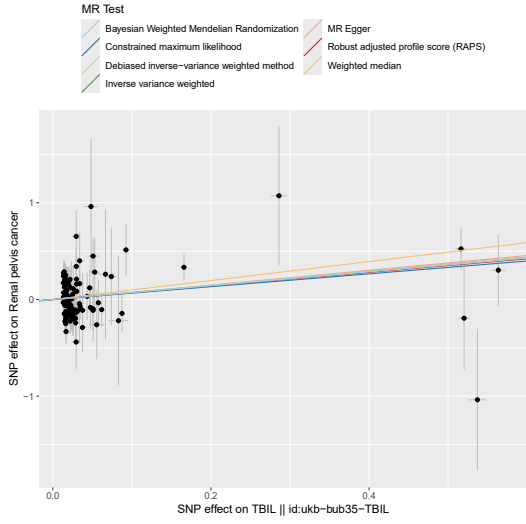
Scatter plot of RPCa

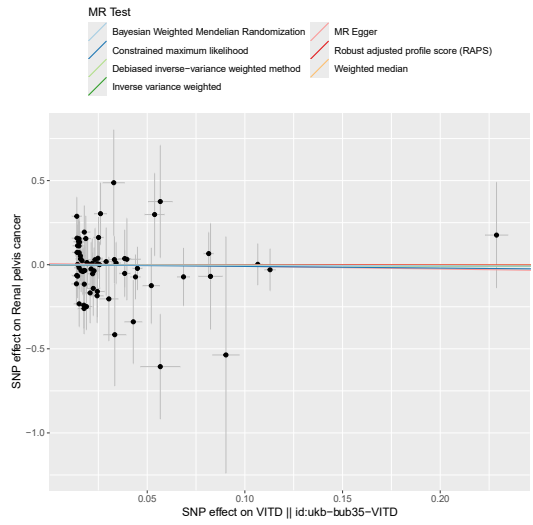
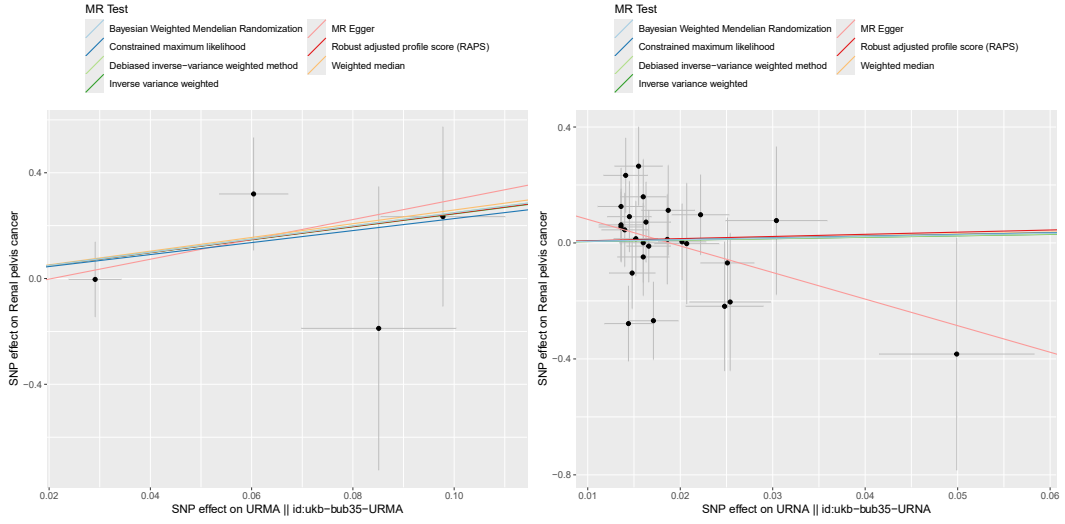




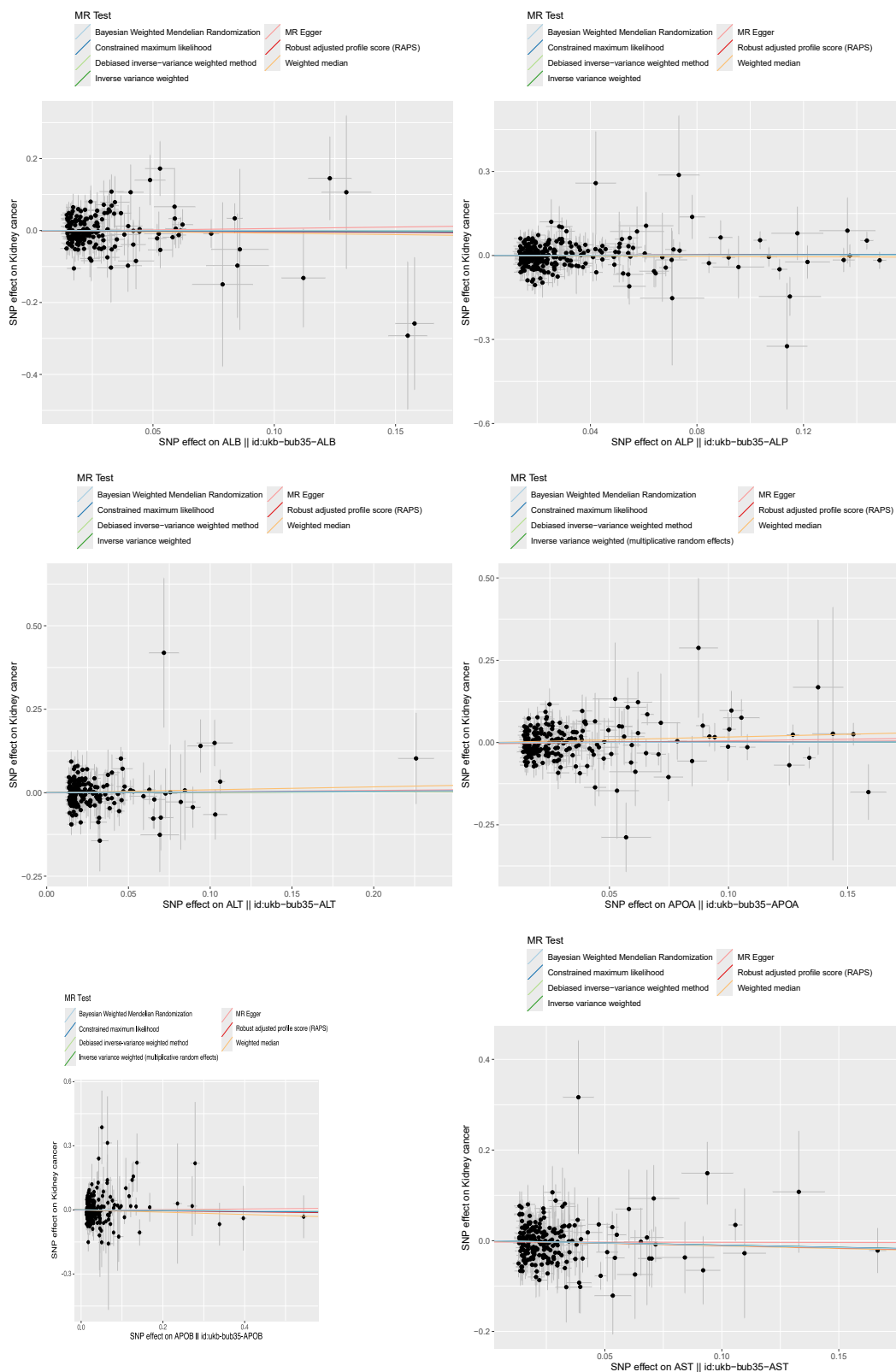


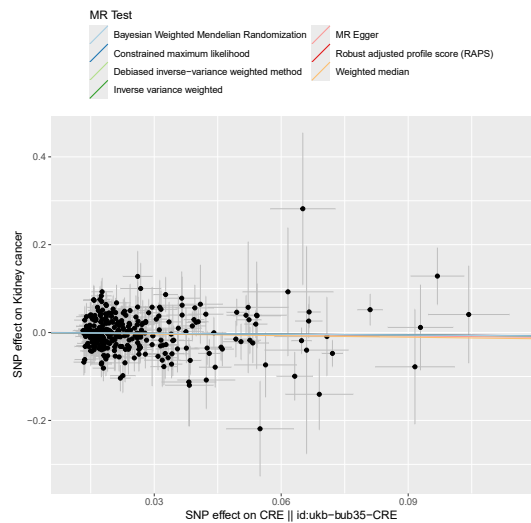
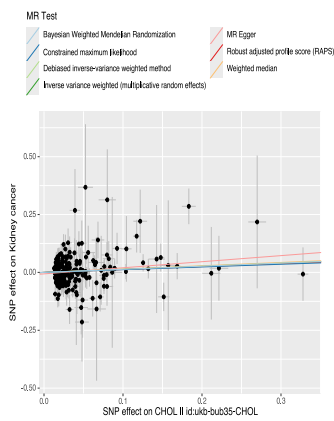
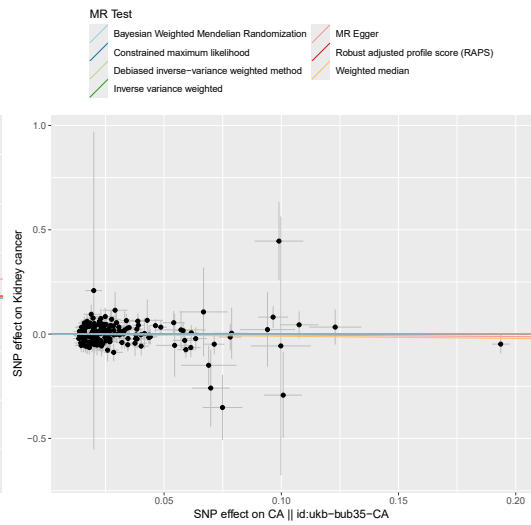
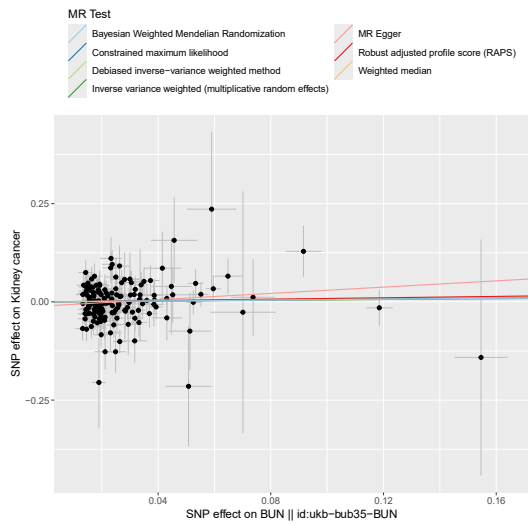
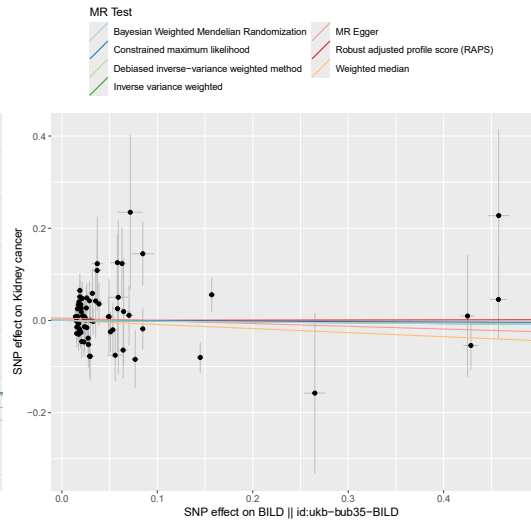
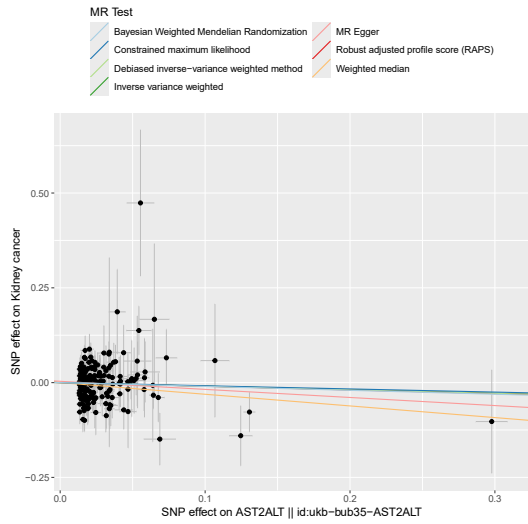


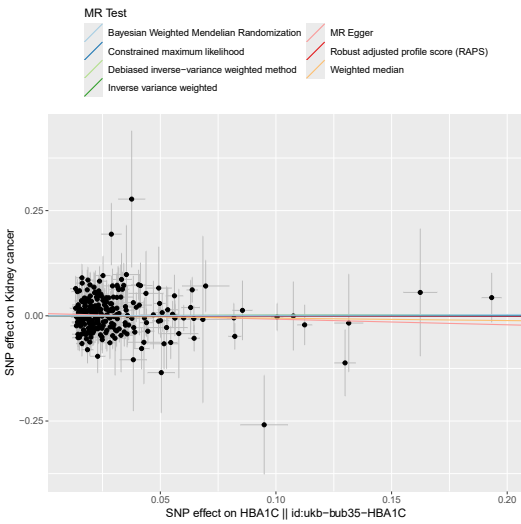
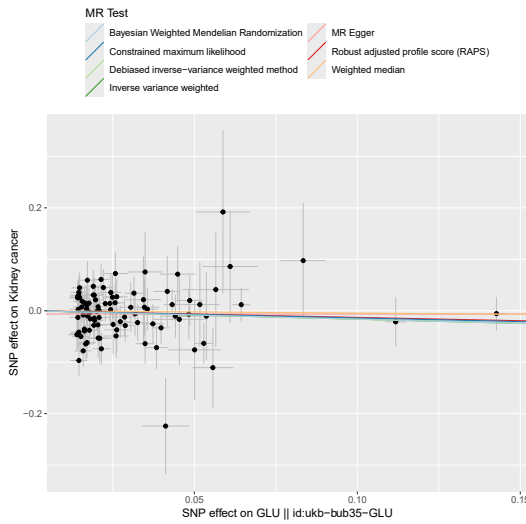
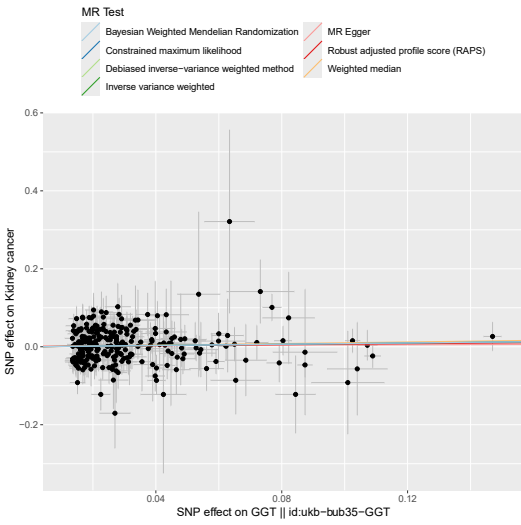
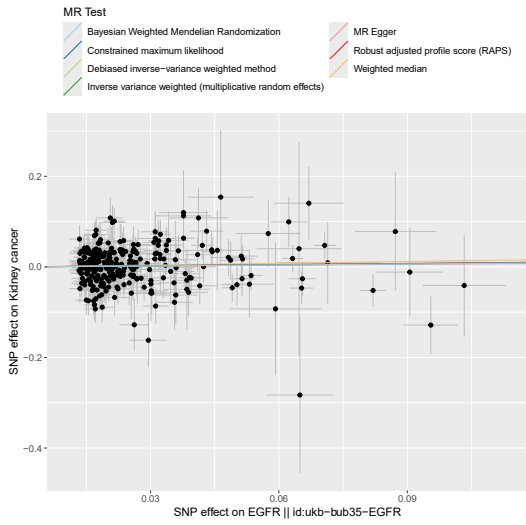
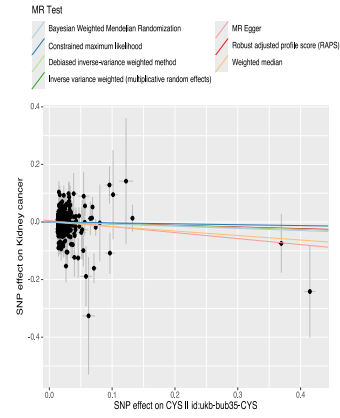
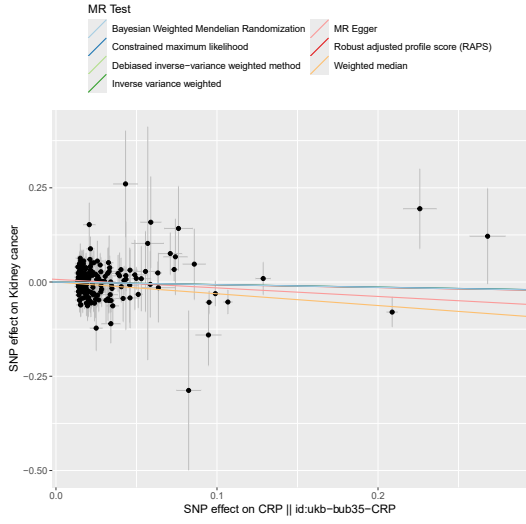


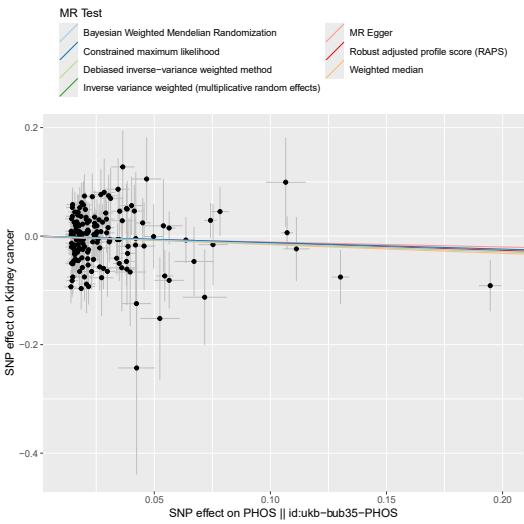
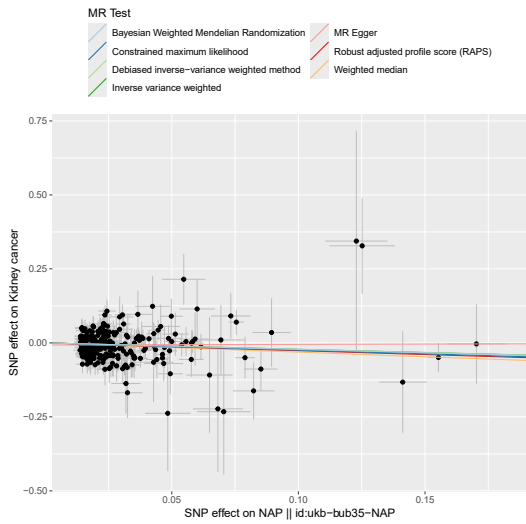
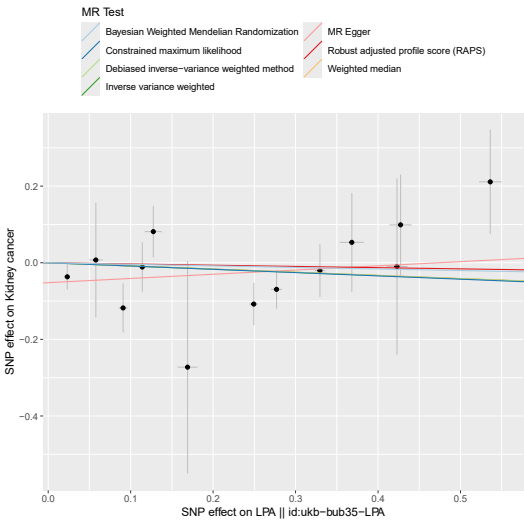
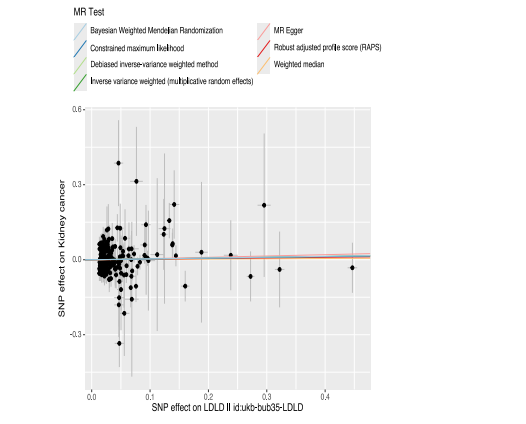
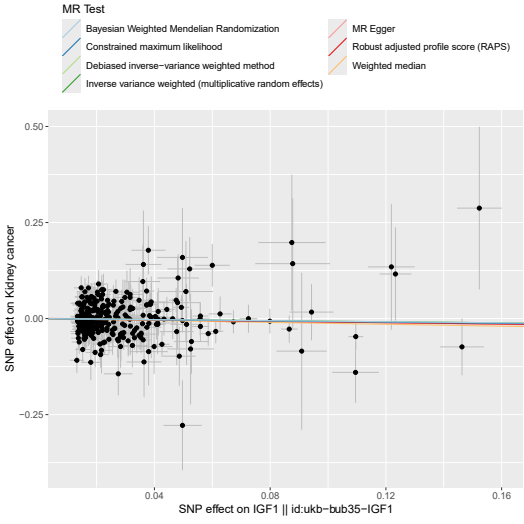
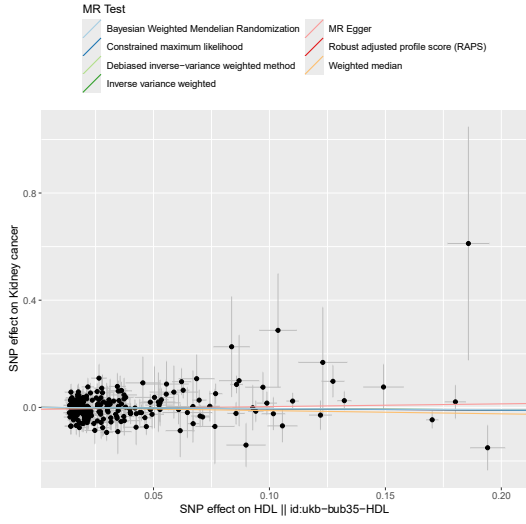


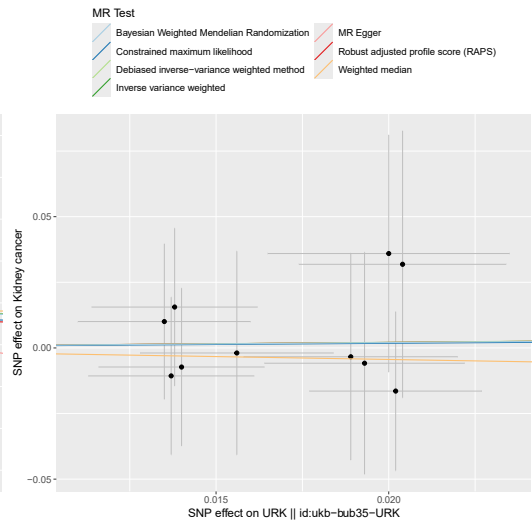
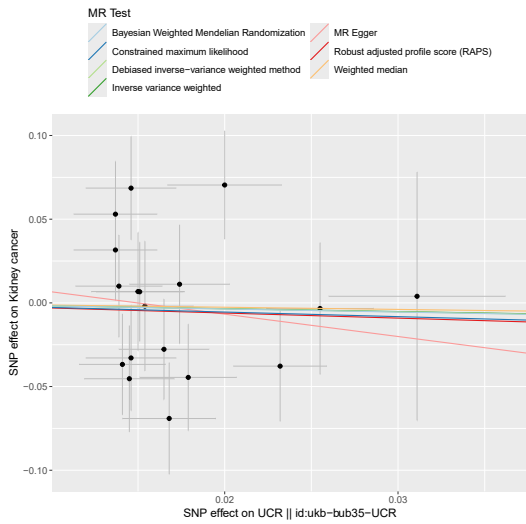
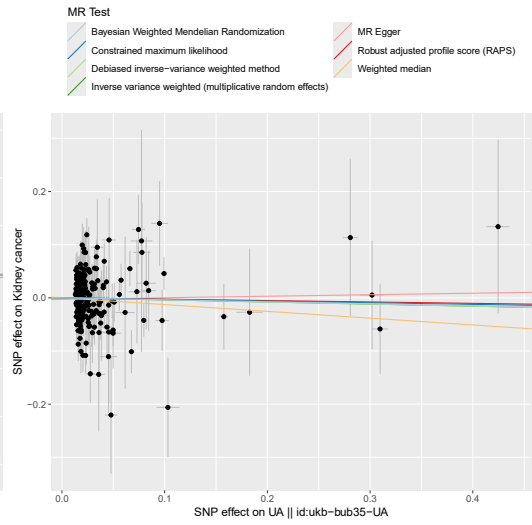
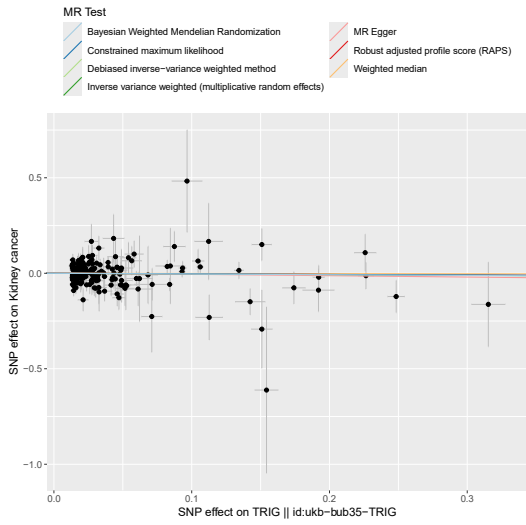
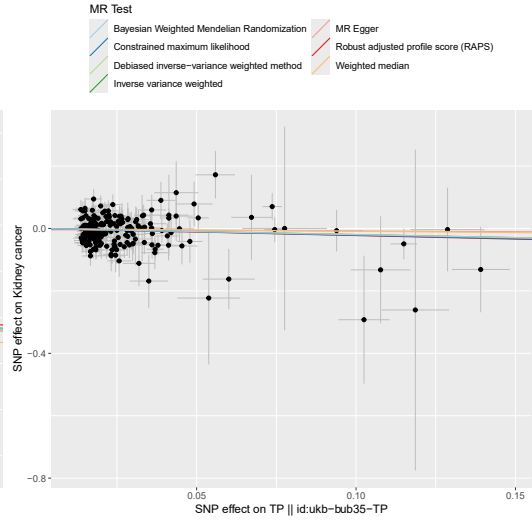
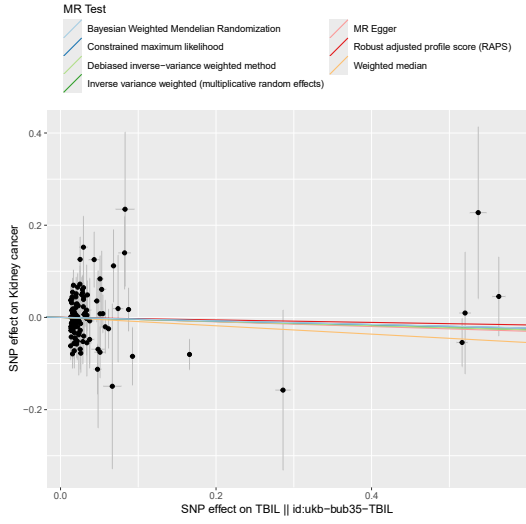
Scatter plot of KCa

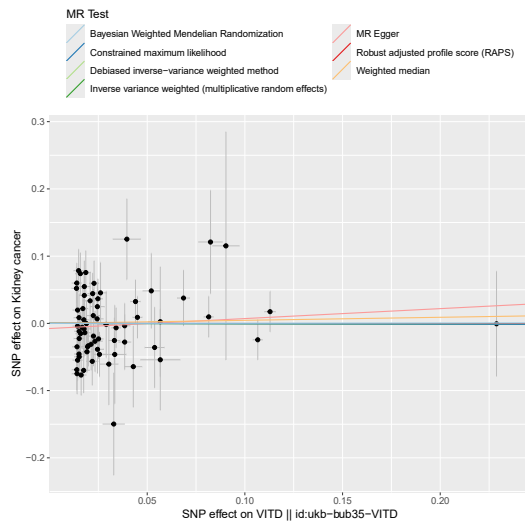
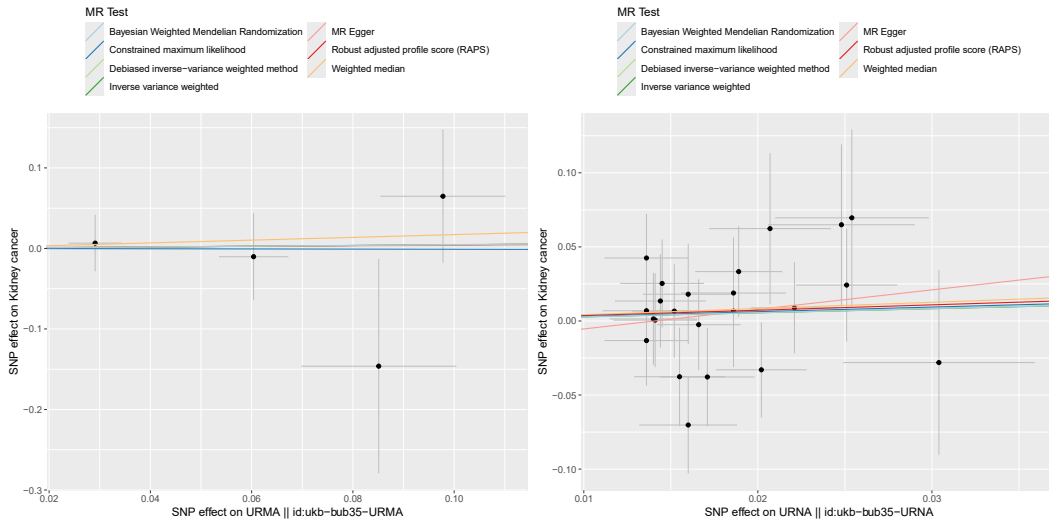




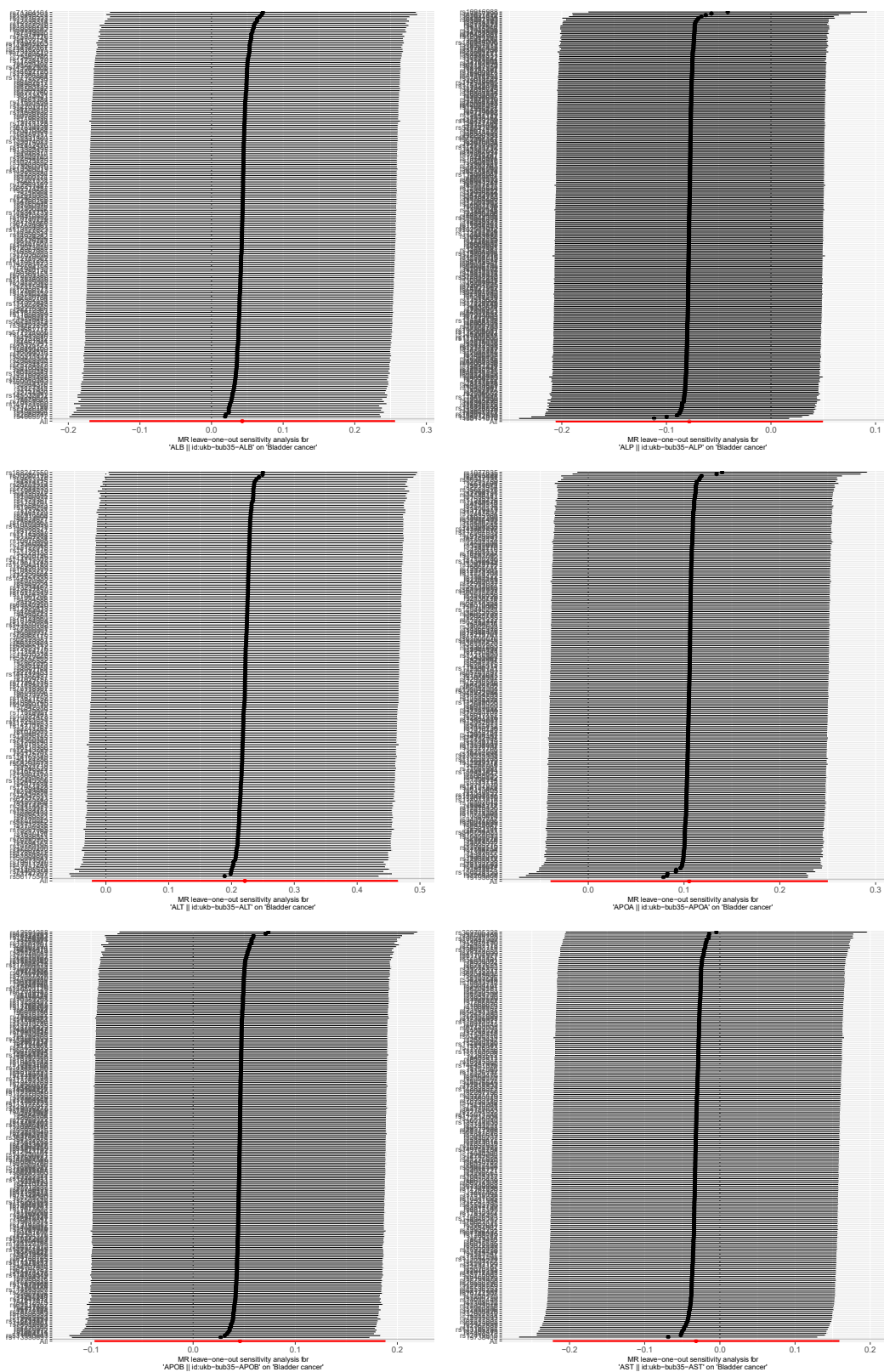


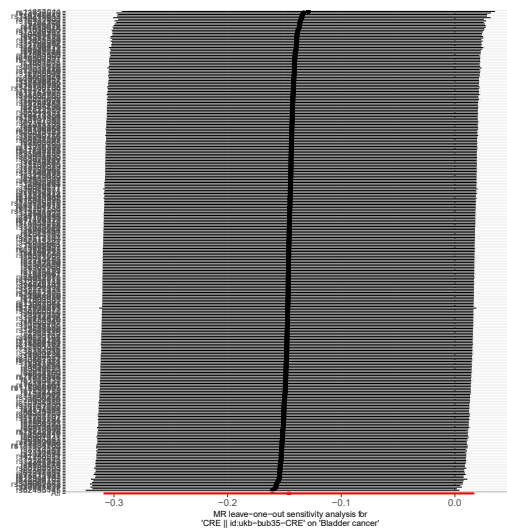
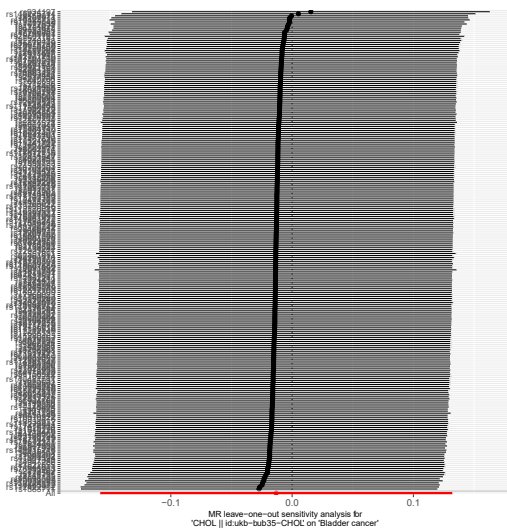
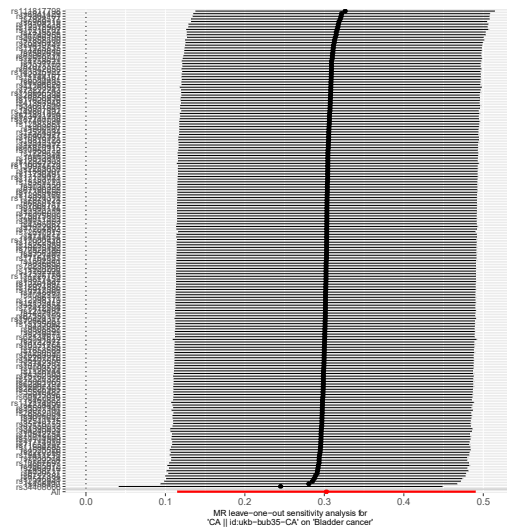
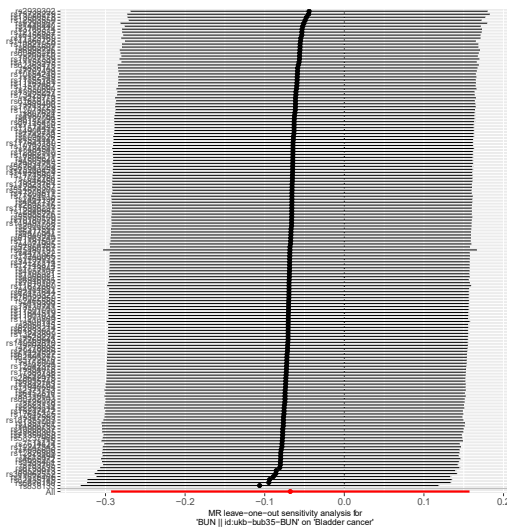
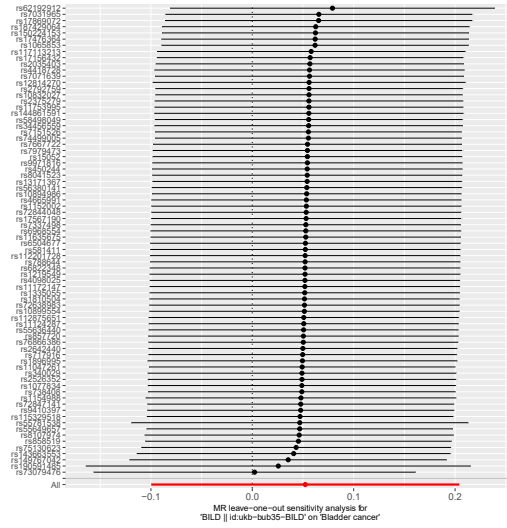
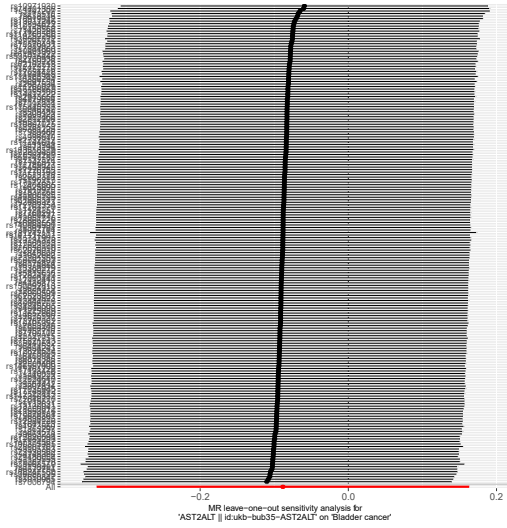


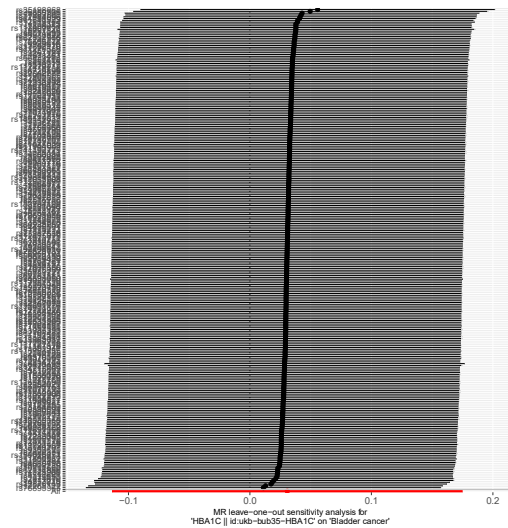
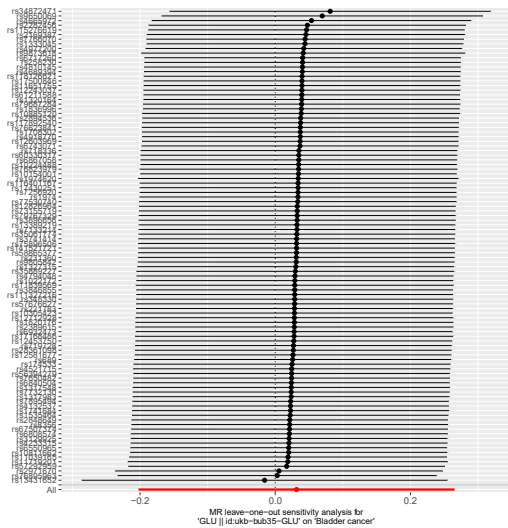
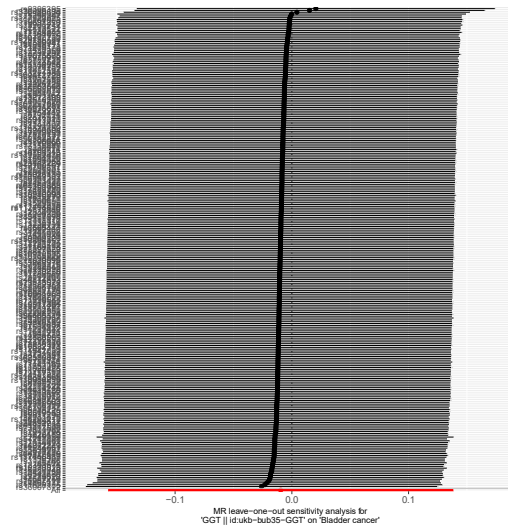
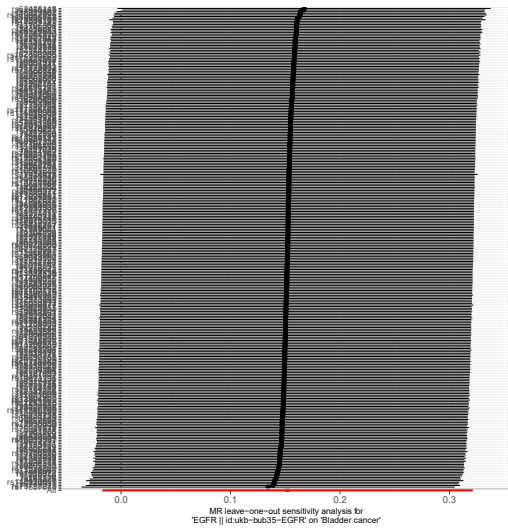
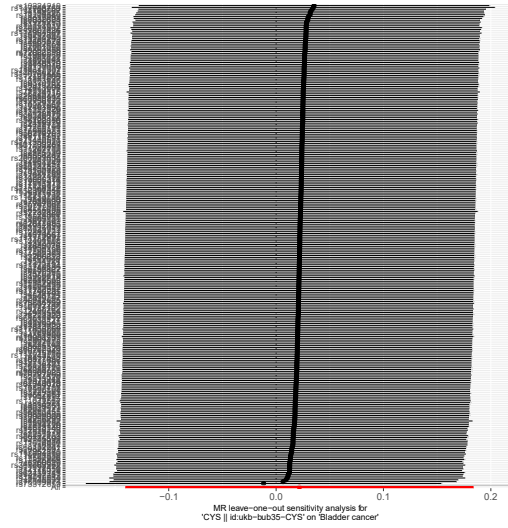
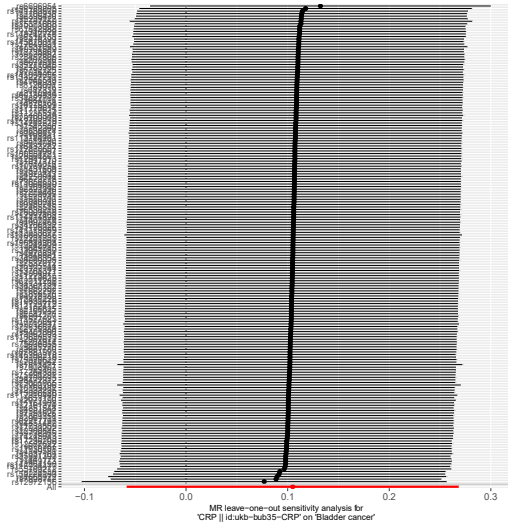


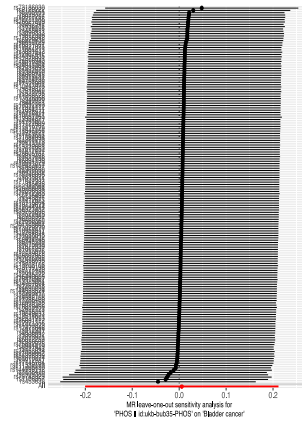
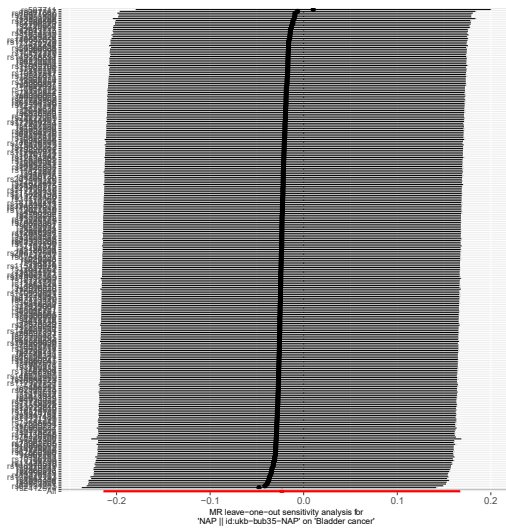
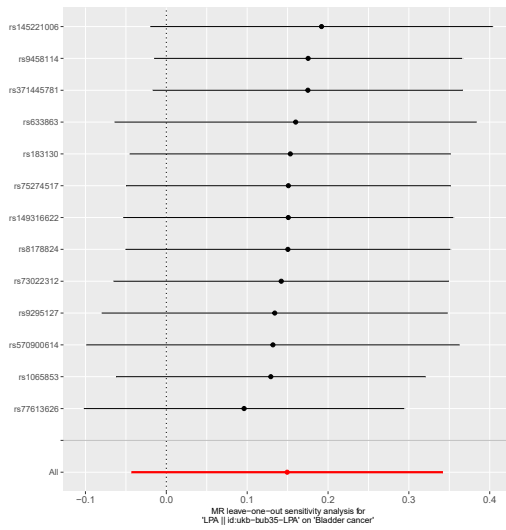
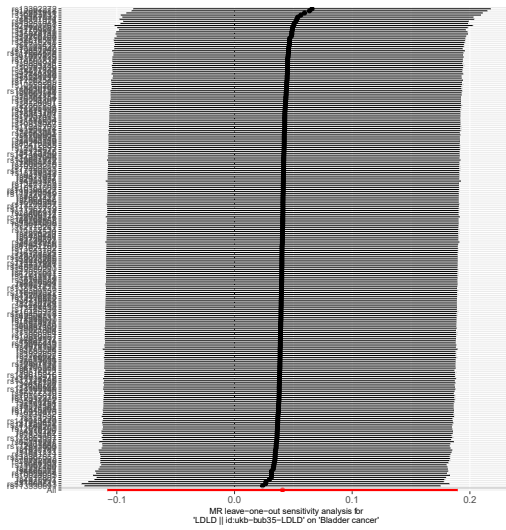
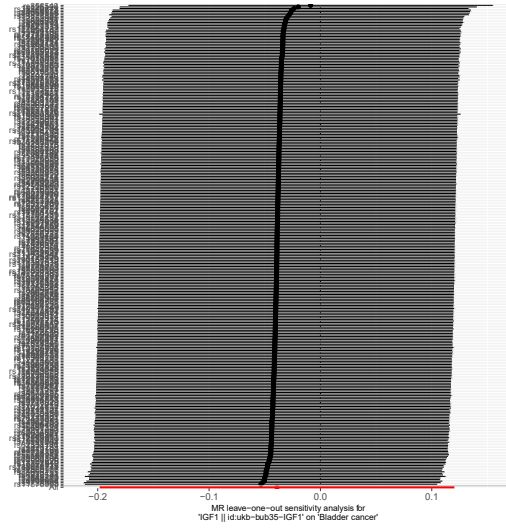
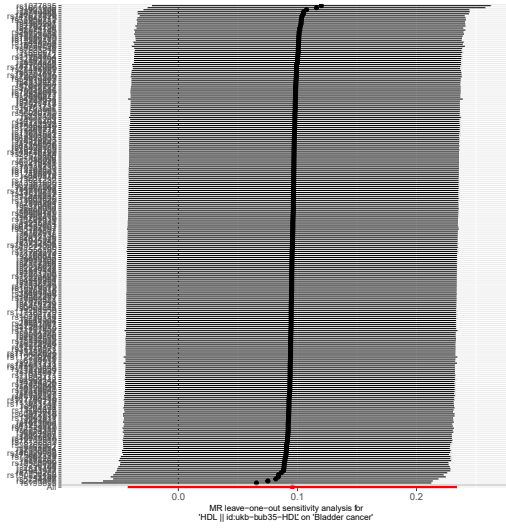


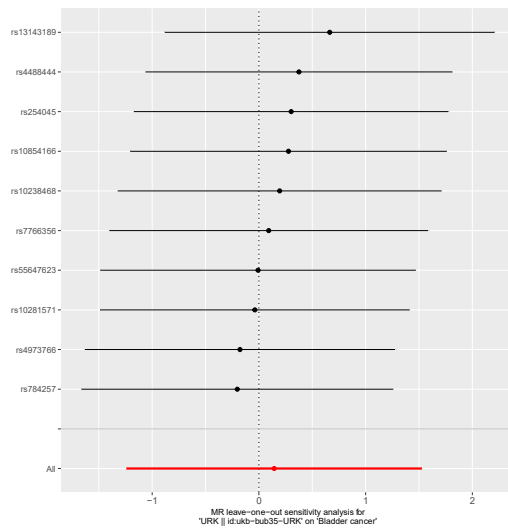
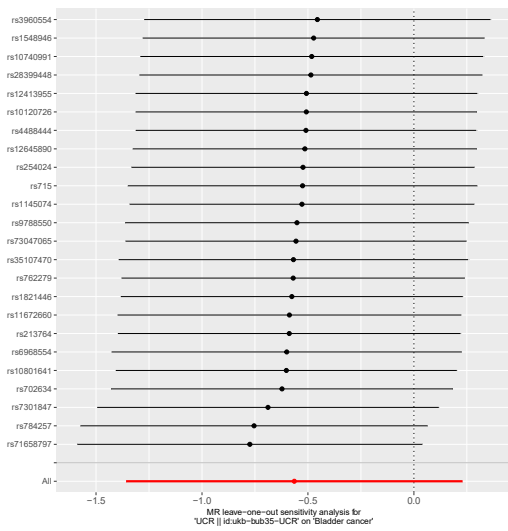
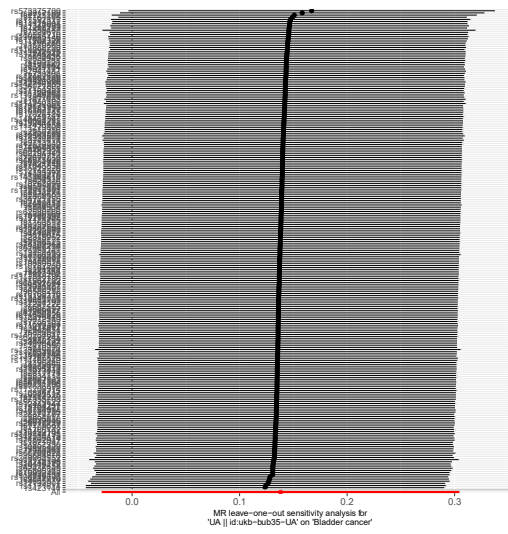
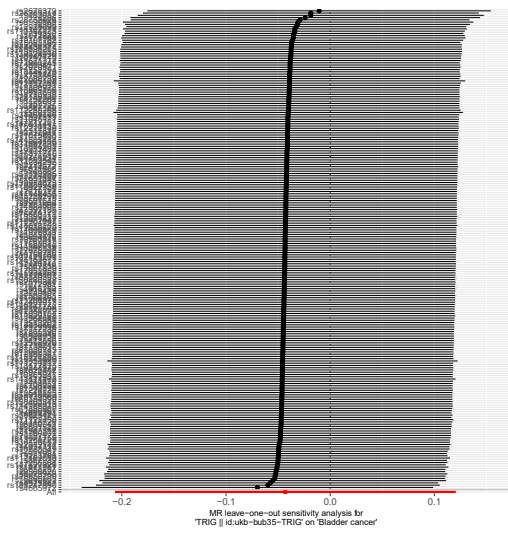
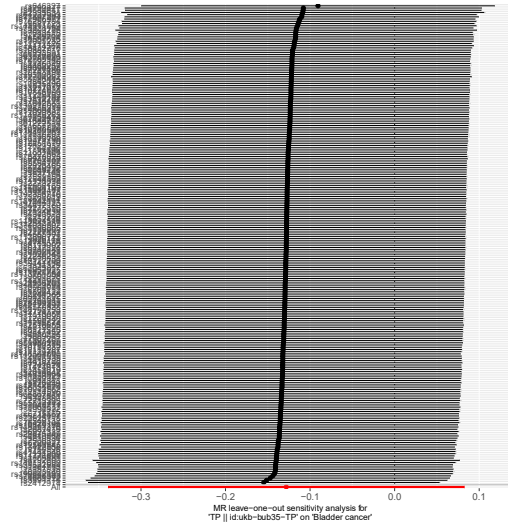
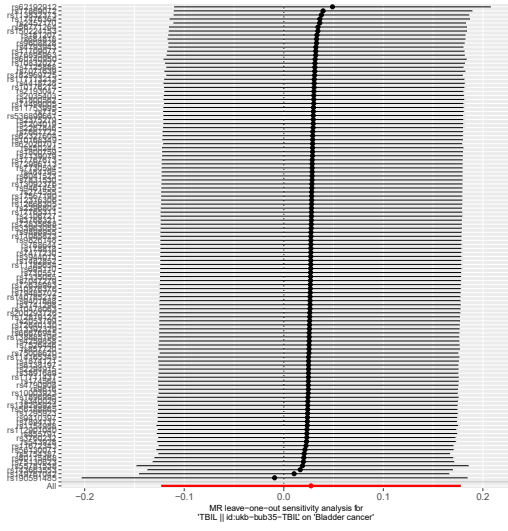
BCa's leave-one-out method

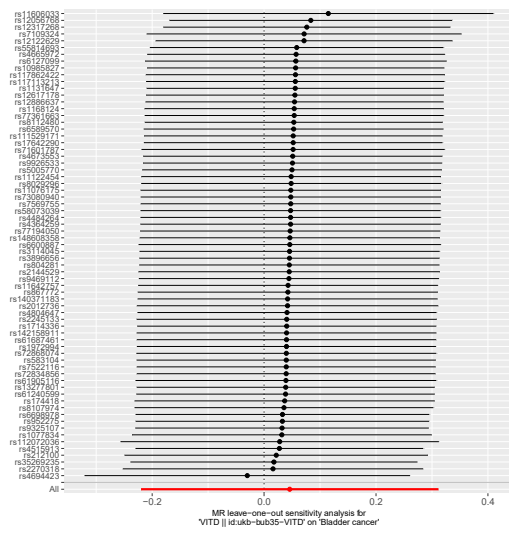
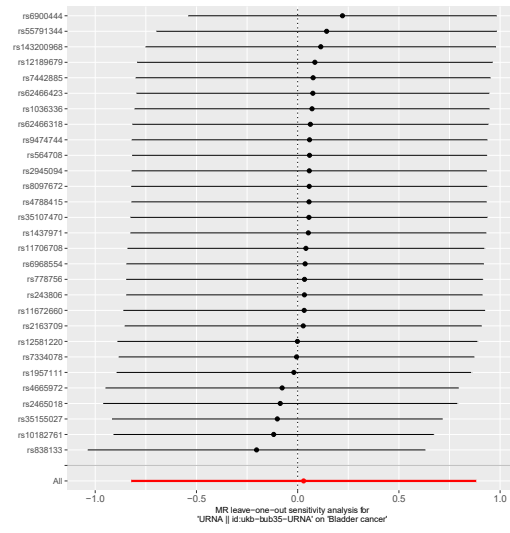
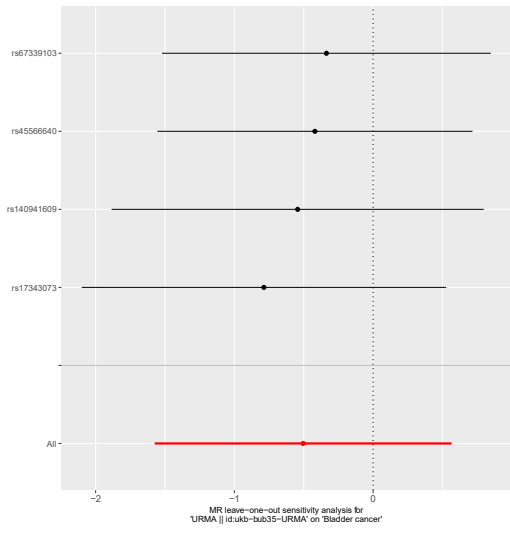




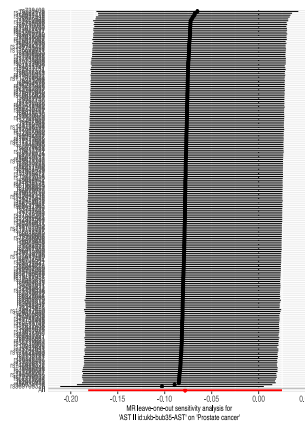
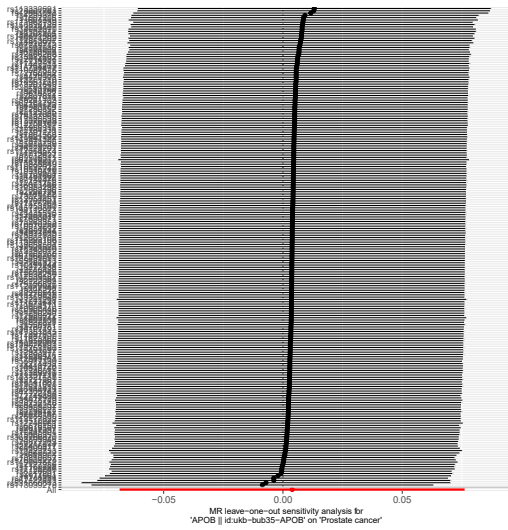
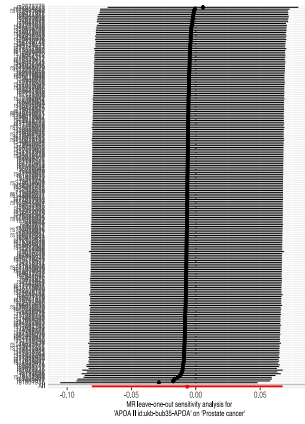
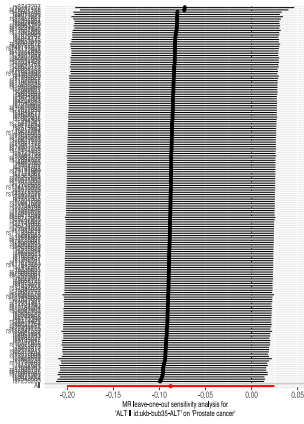
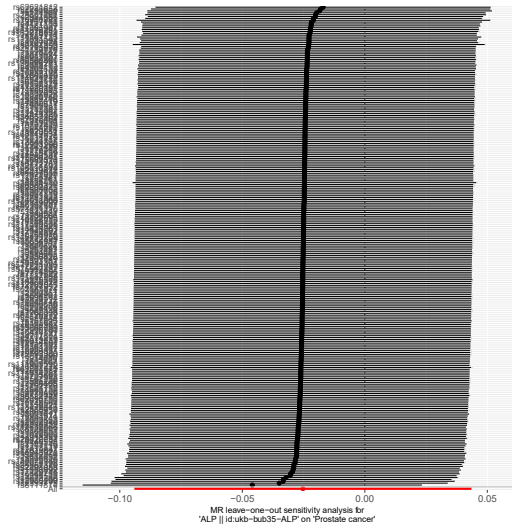
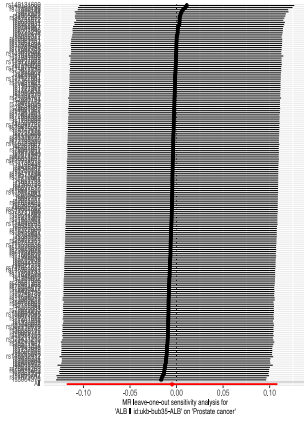


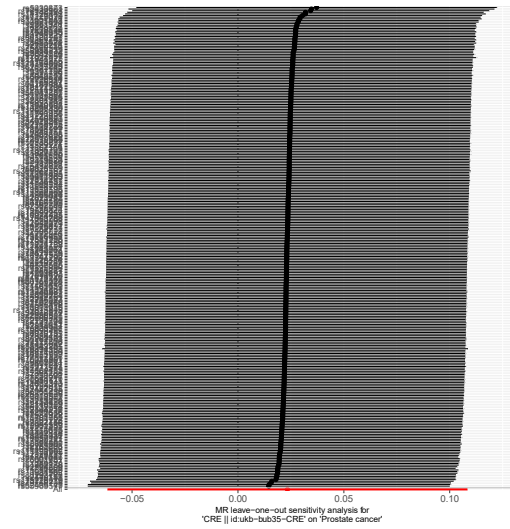
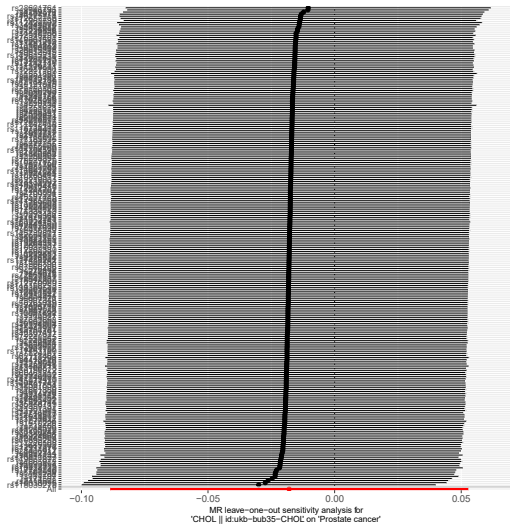
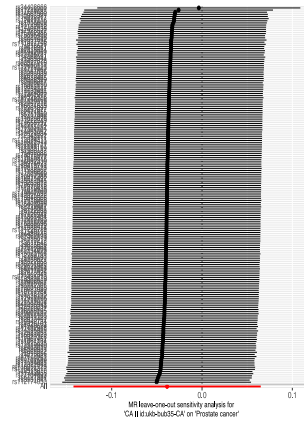
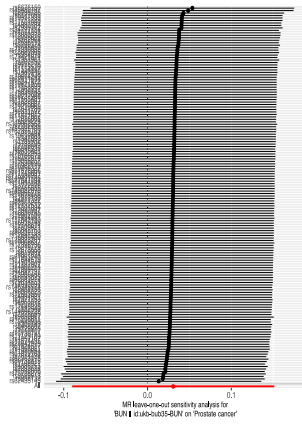
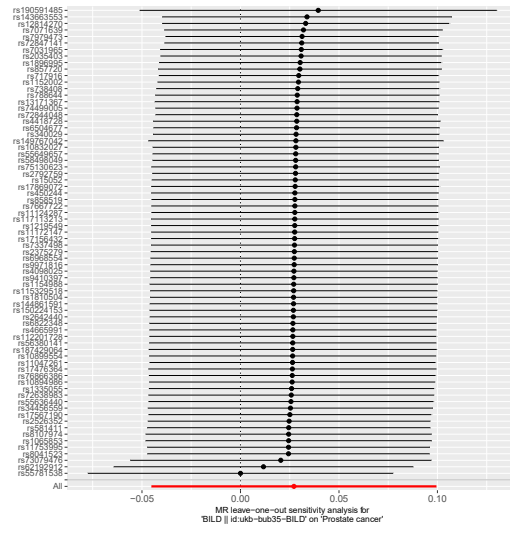
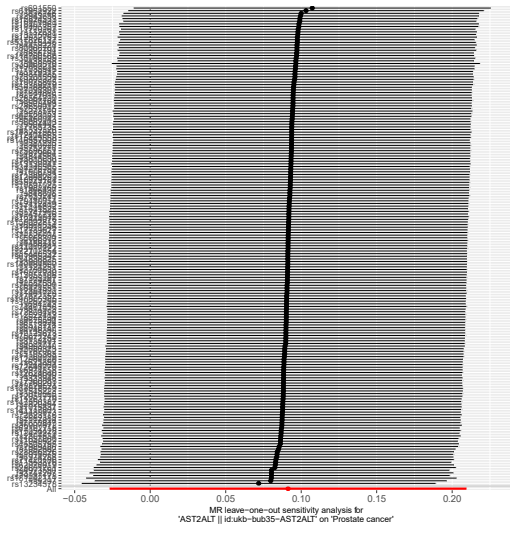


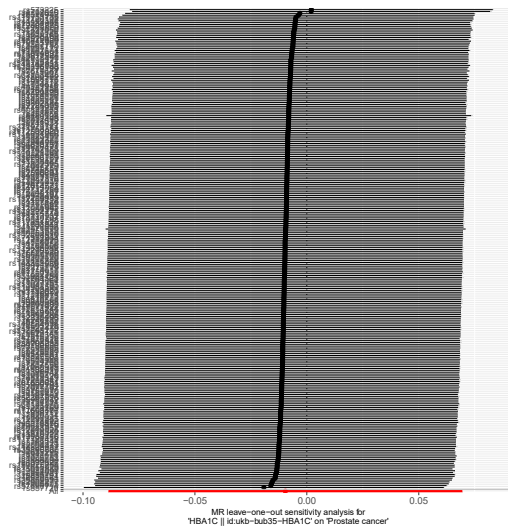
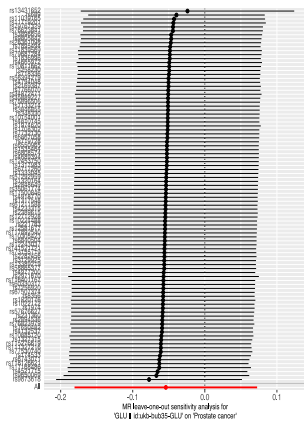
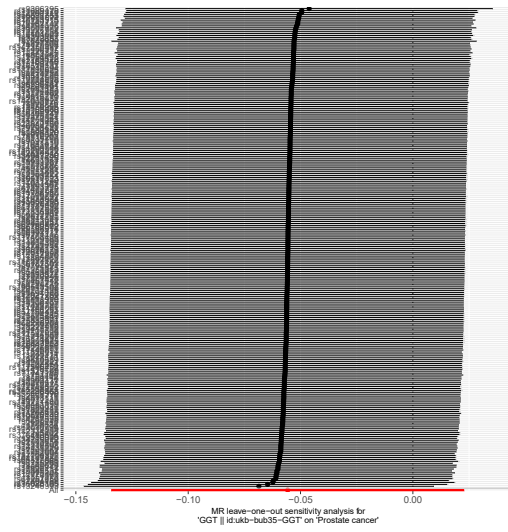
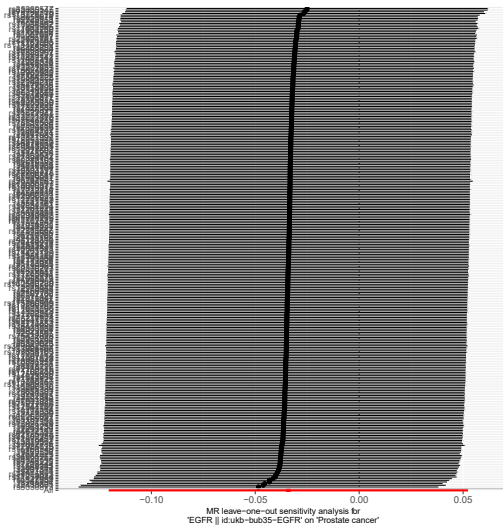
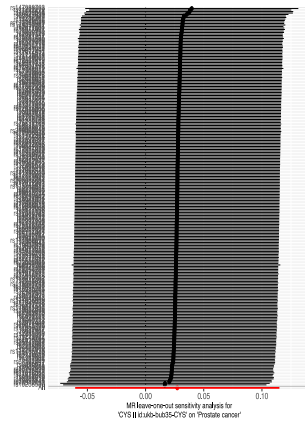
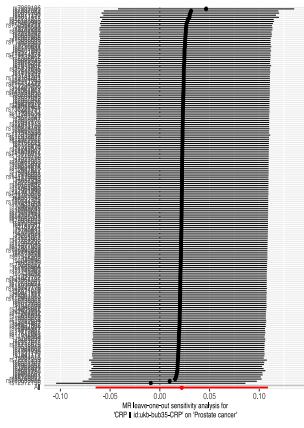


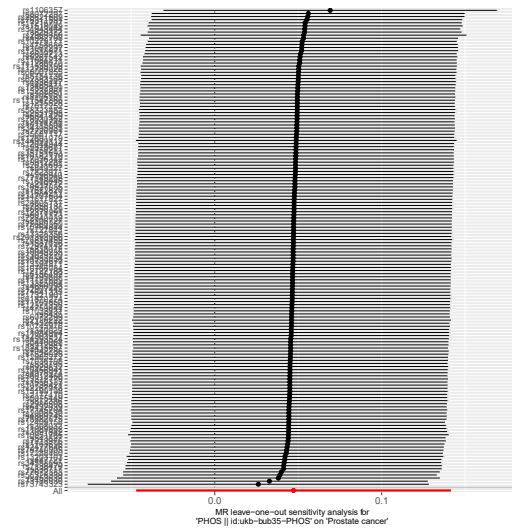
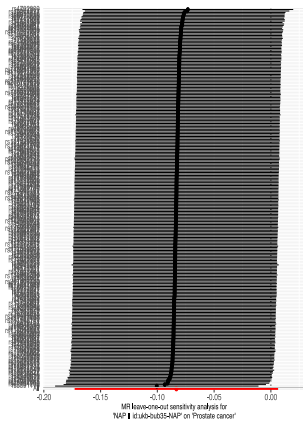
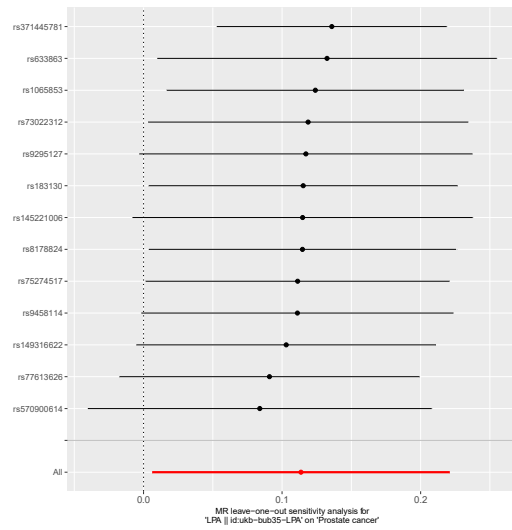
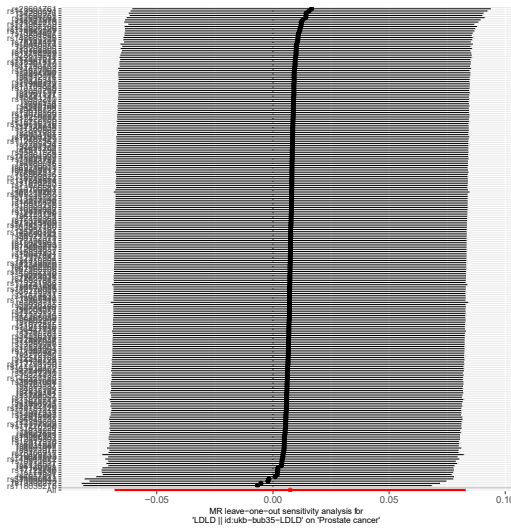
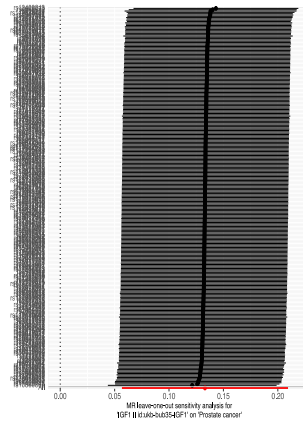
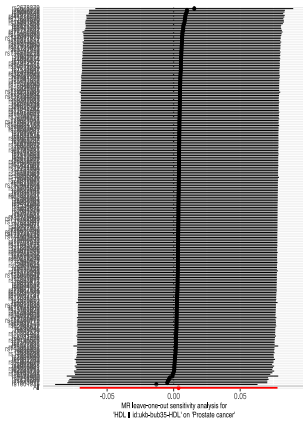


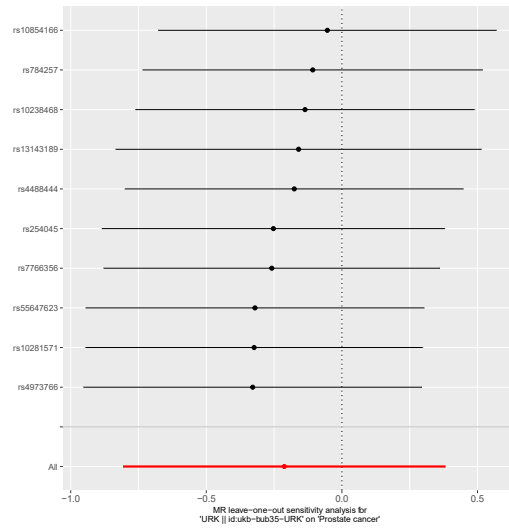
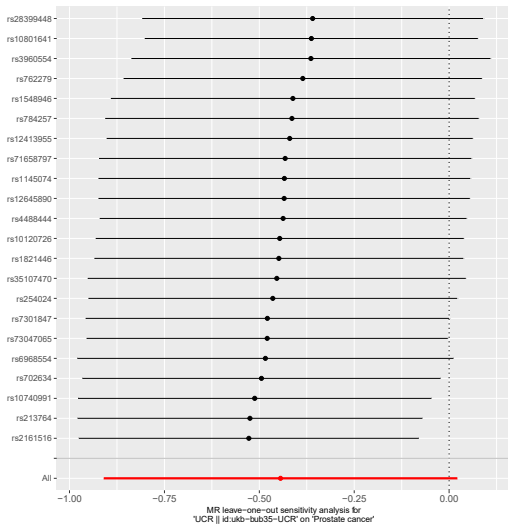
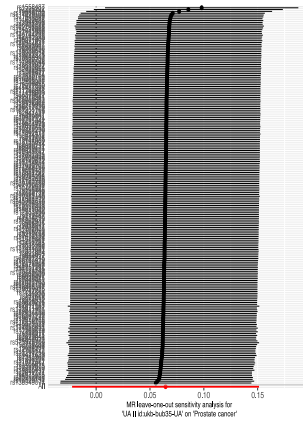
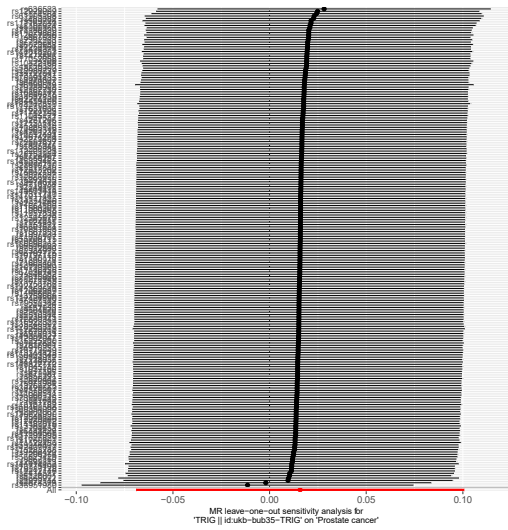
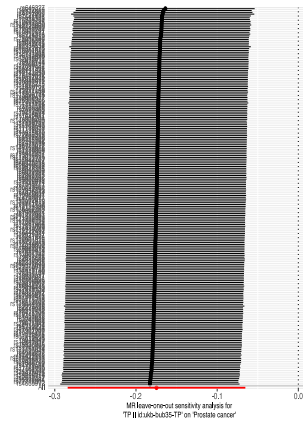
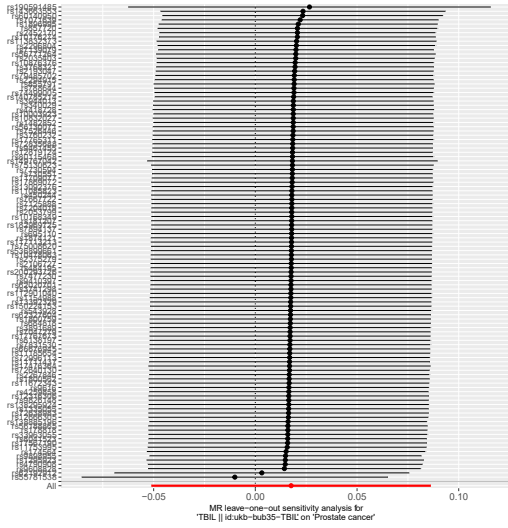
PCa's leave-one-out method

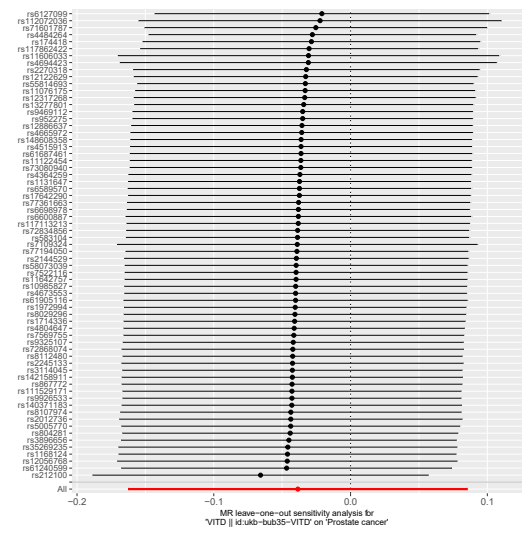
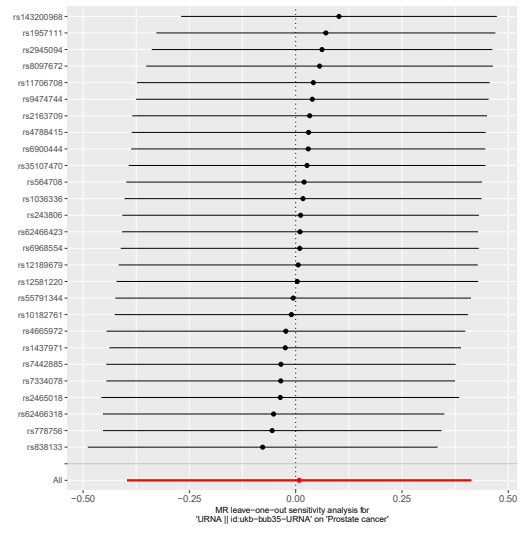
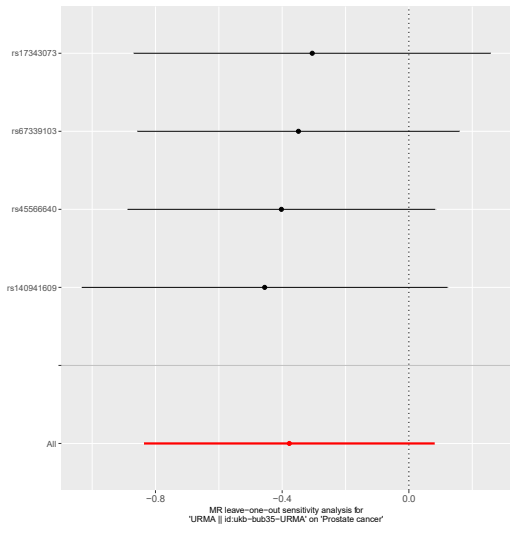




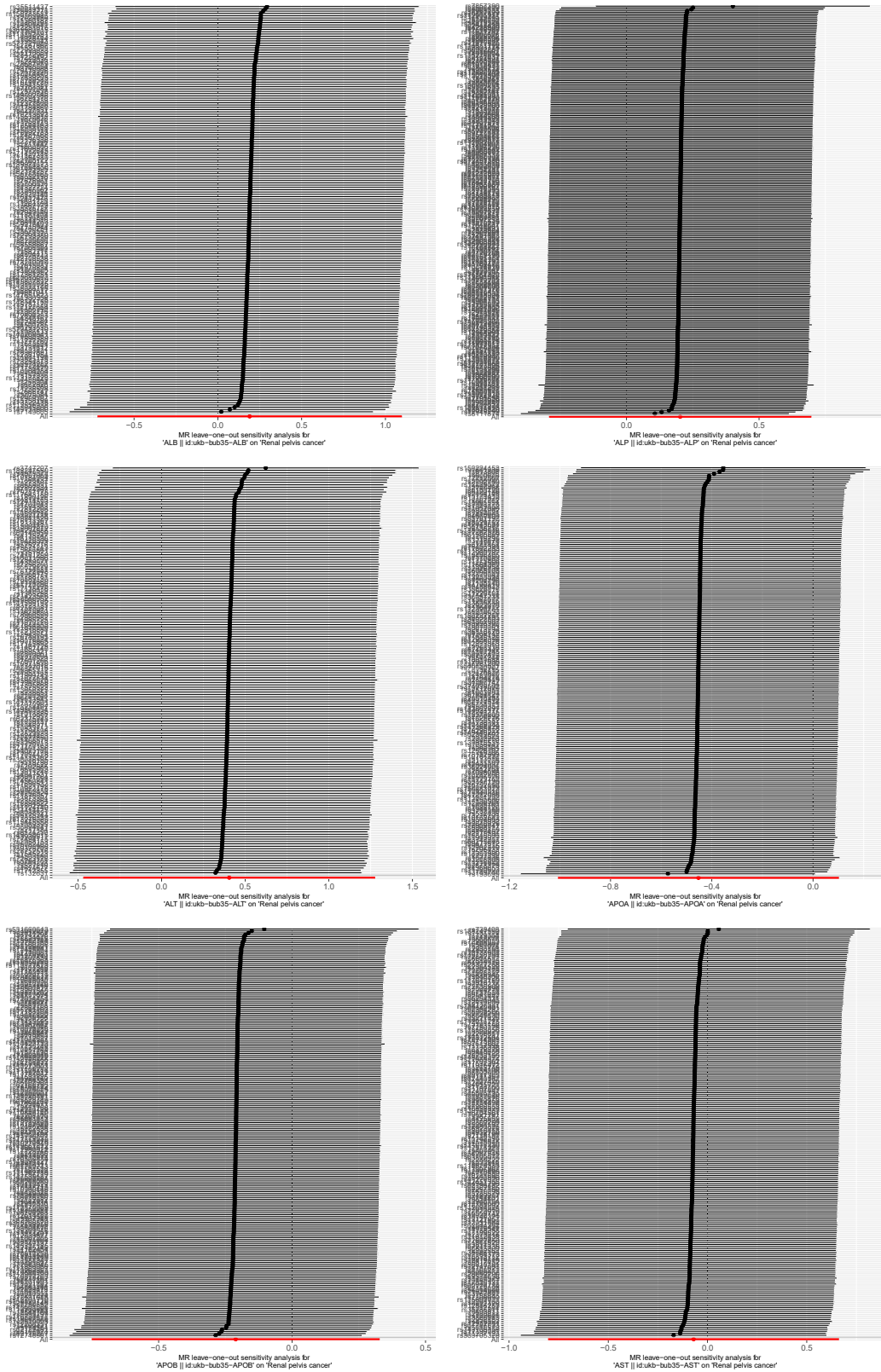


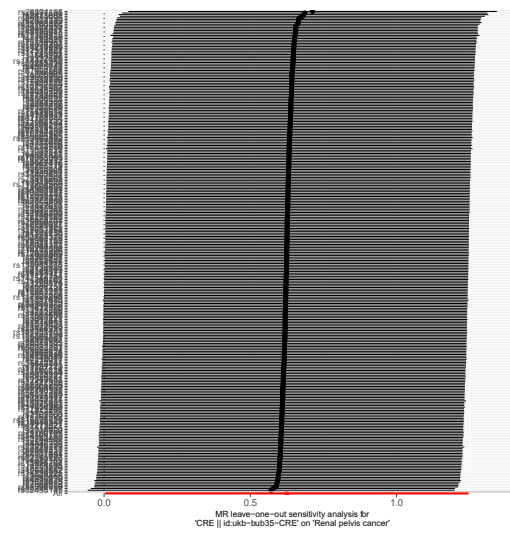
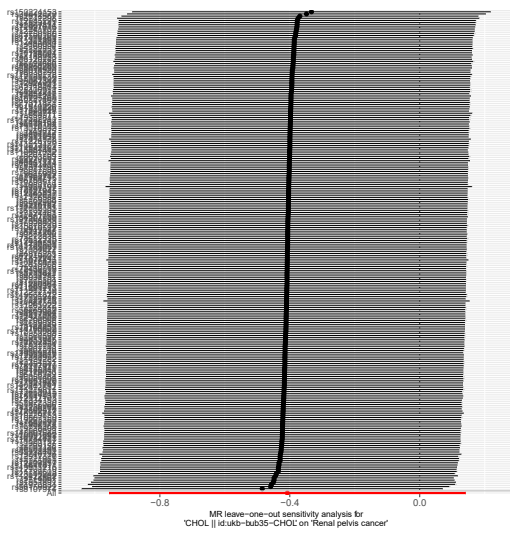
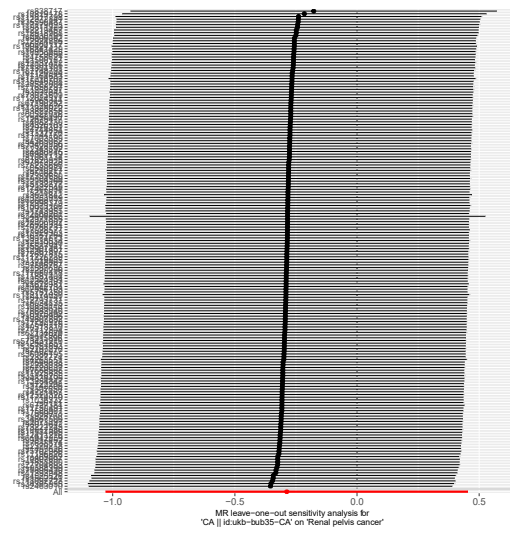
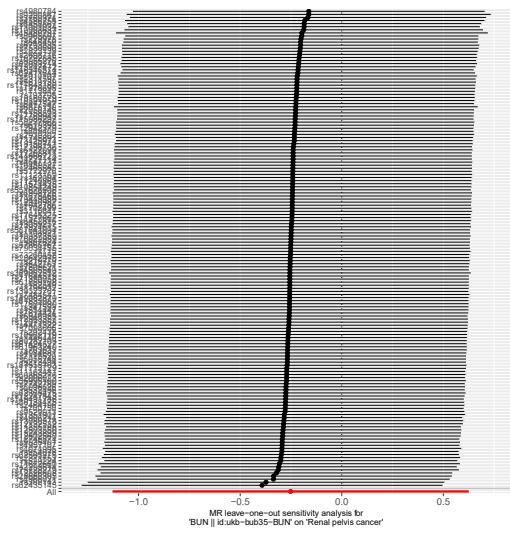
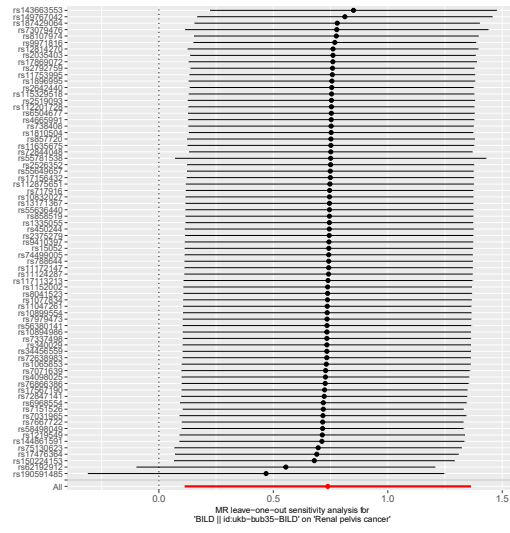
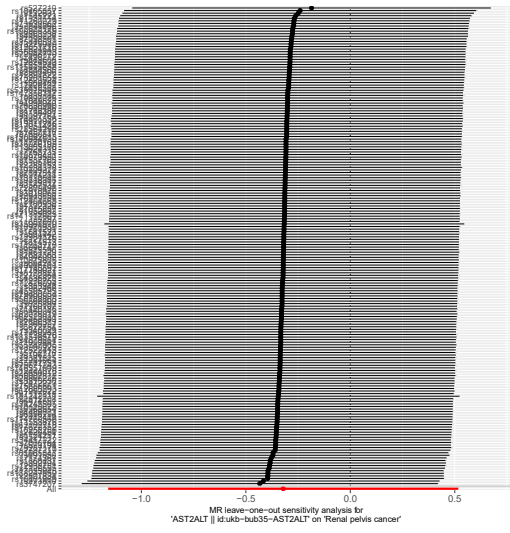


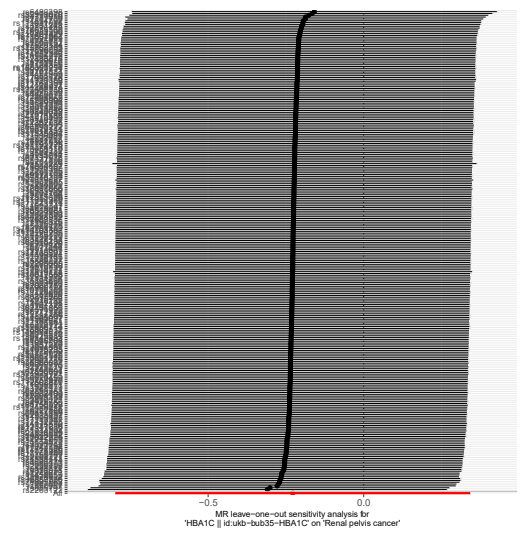
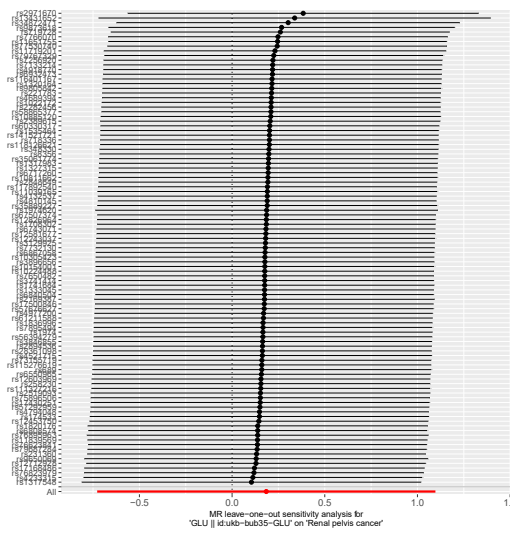
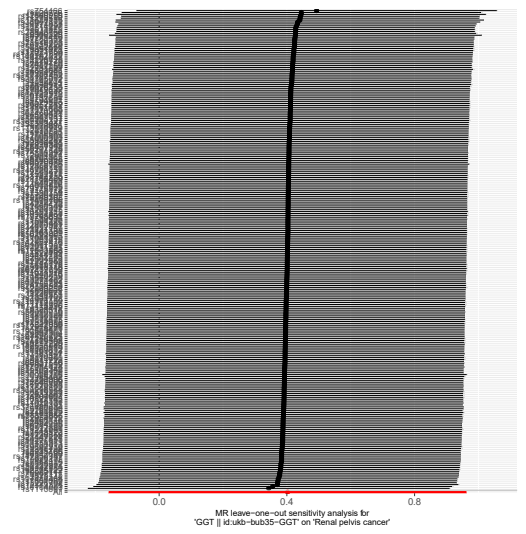
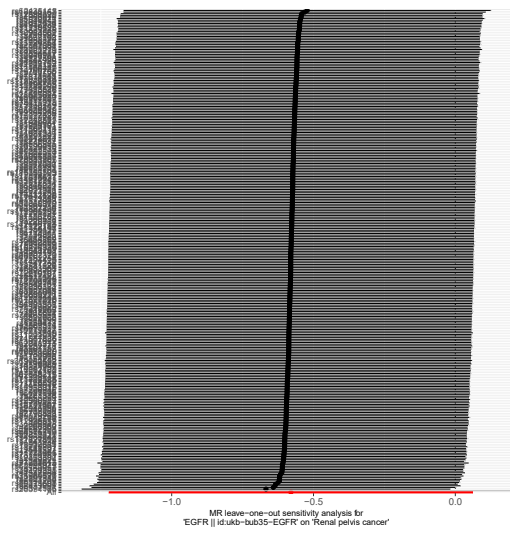
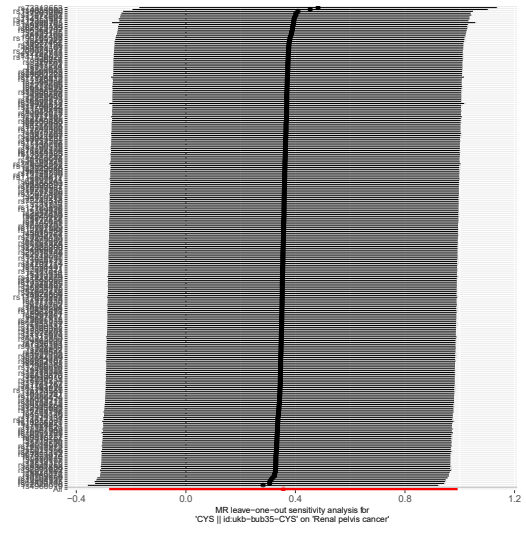
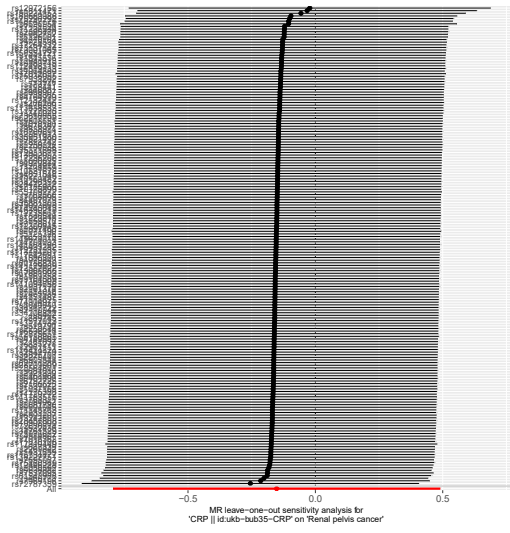


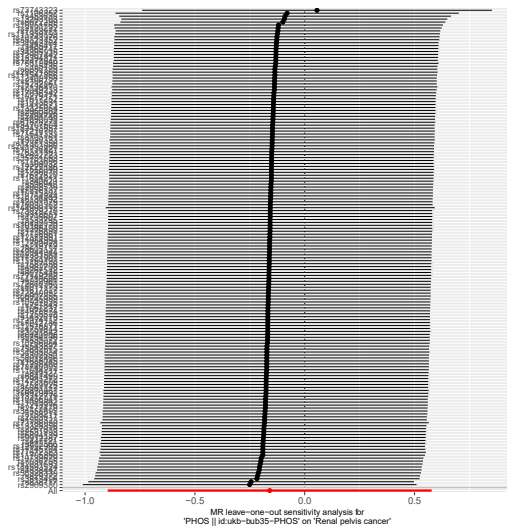
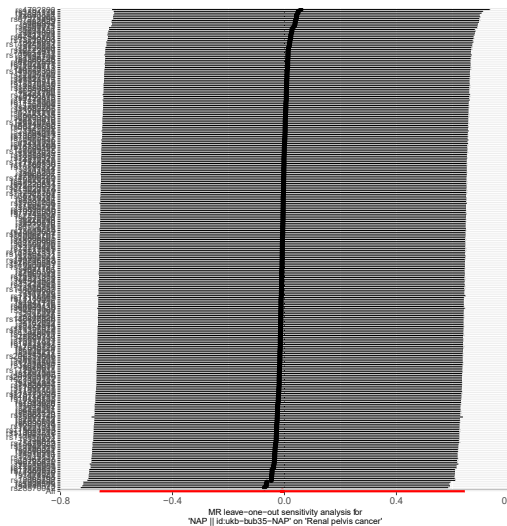
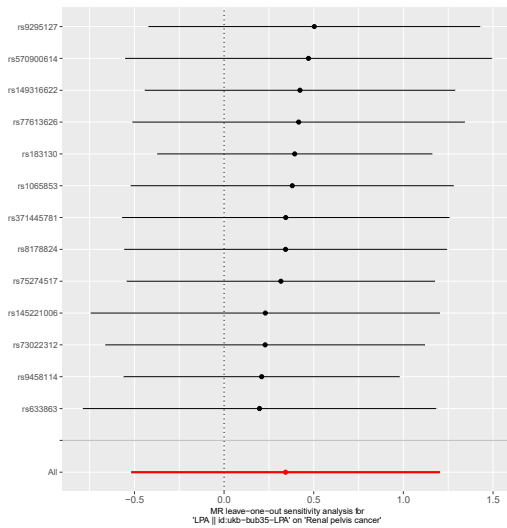
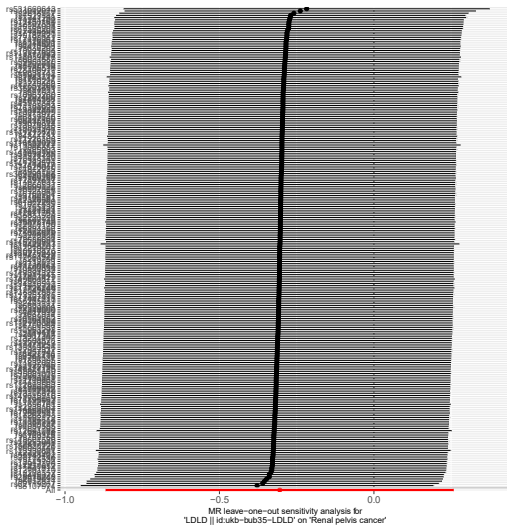
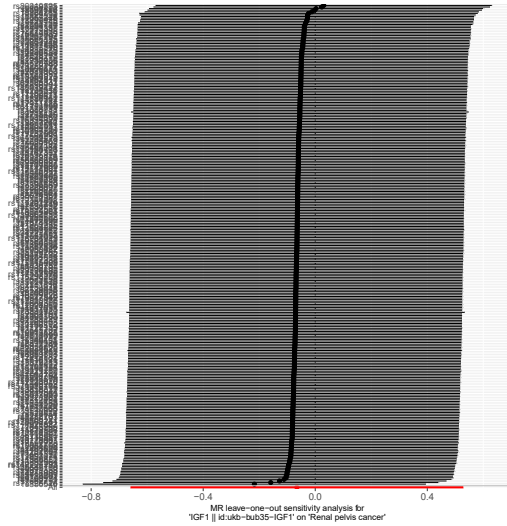
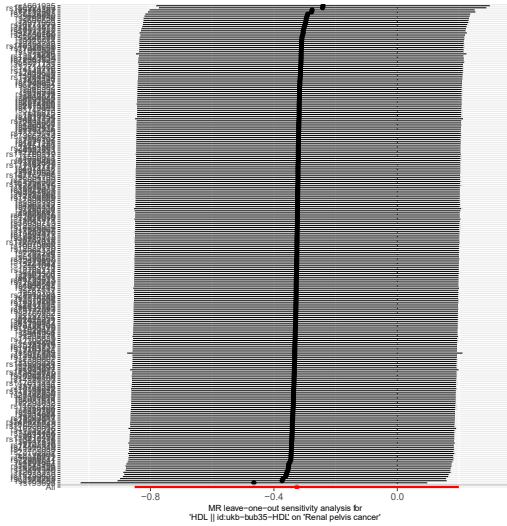


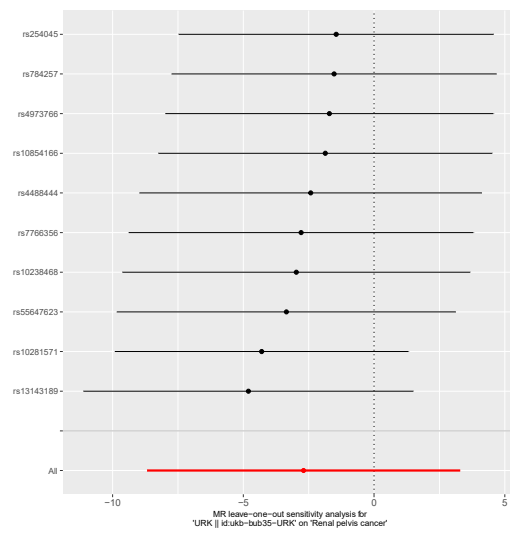
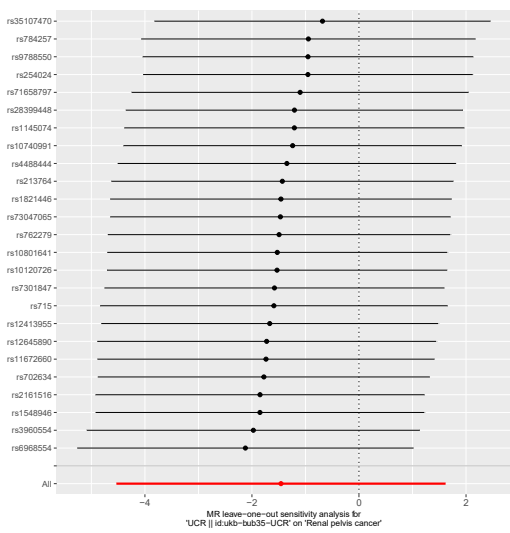
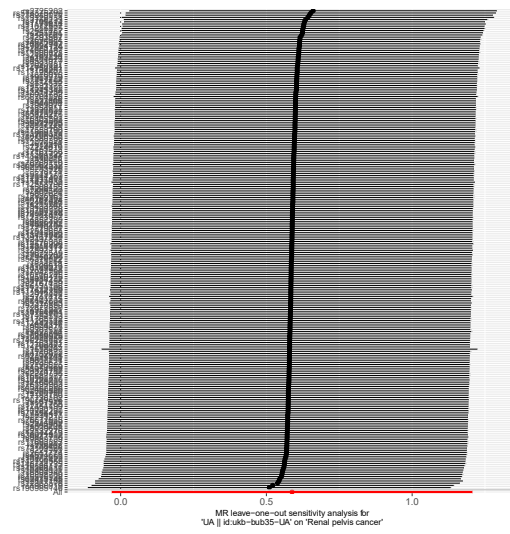
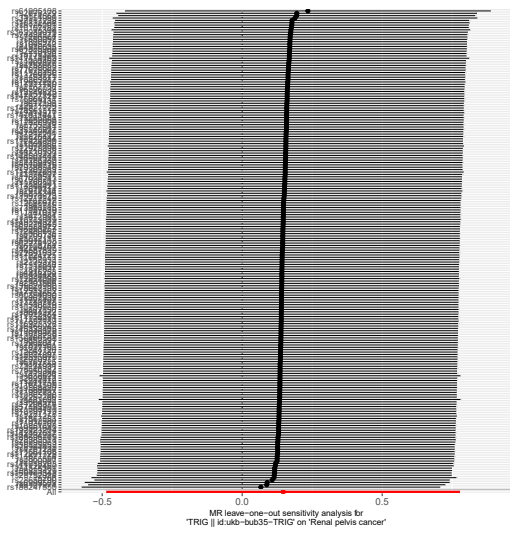
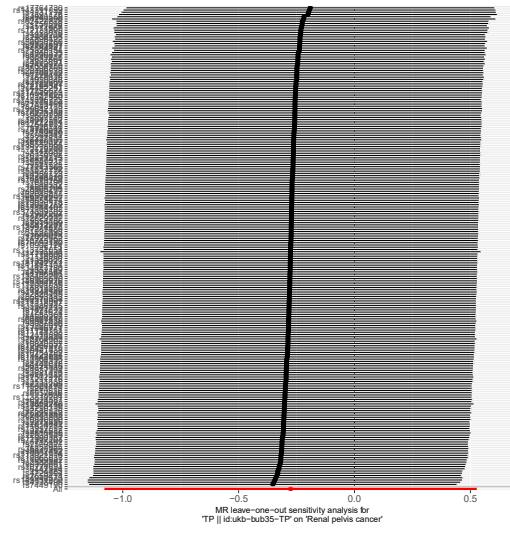
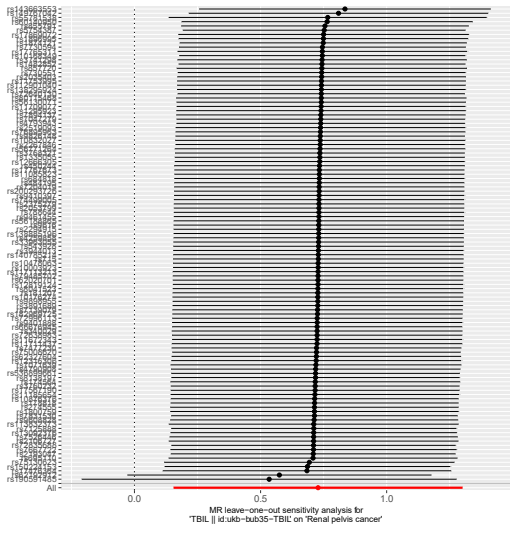
RPCa's leave-one-out method

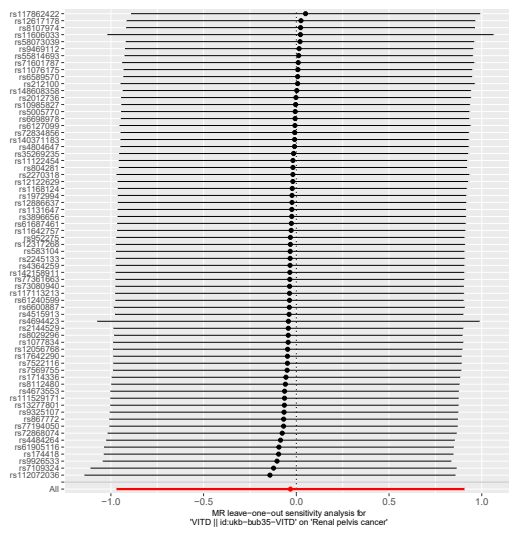
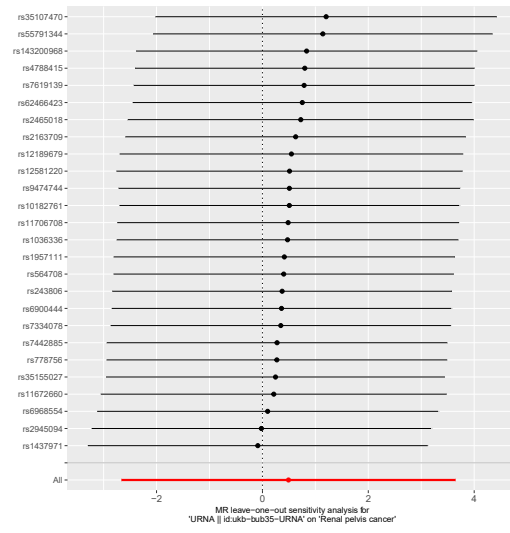
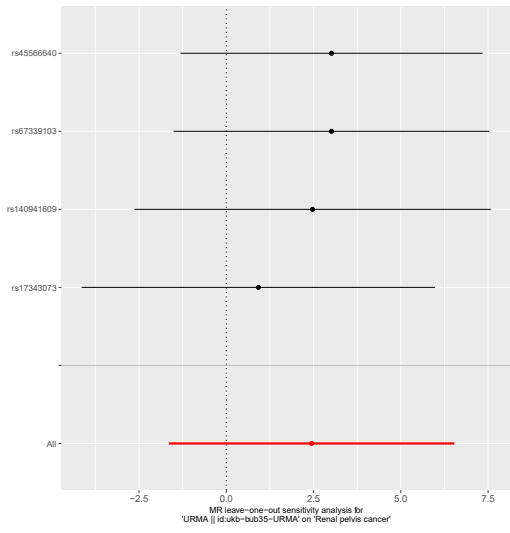




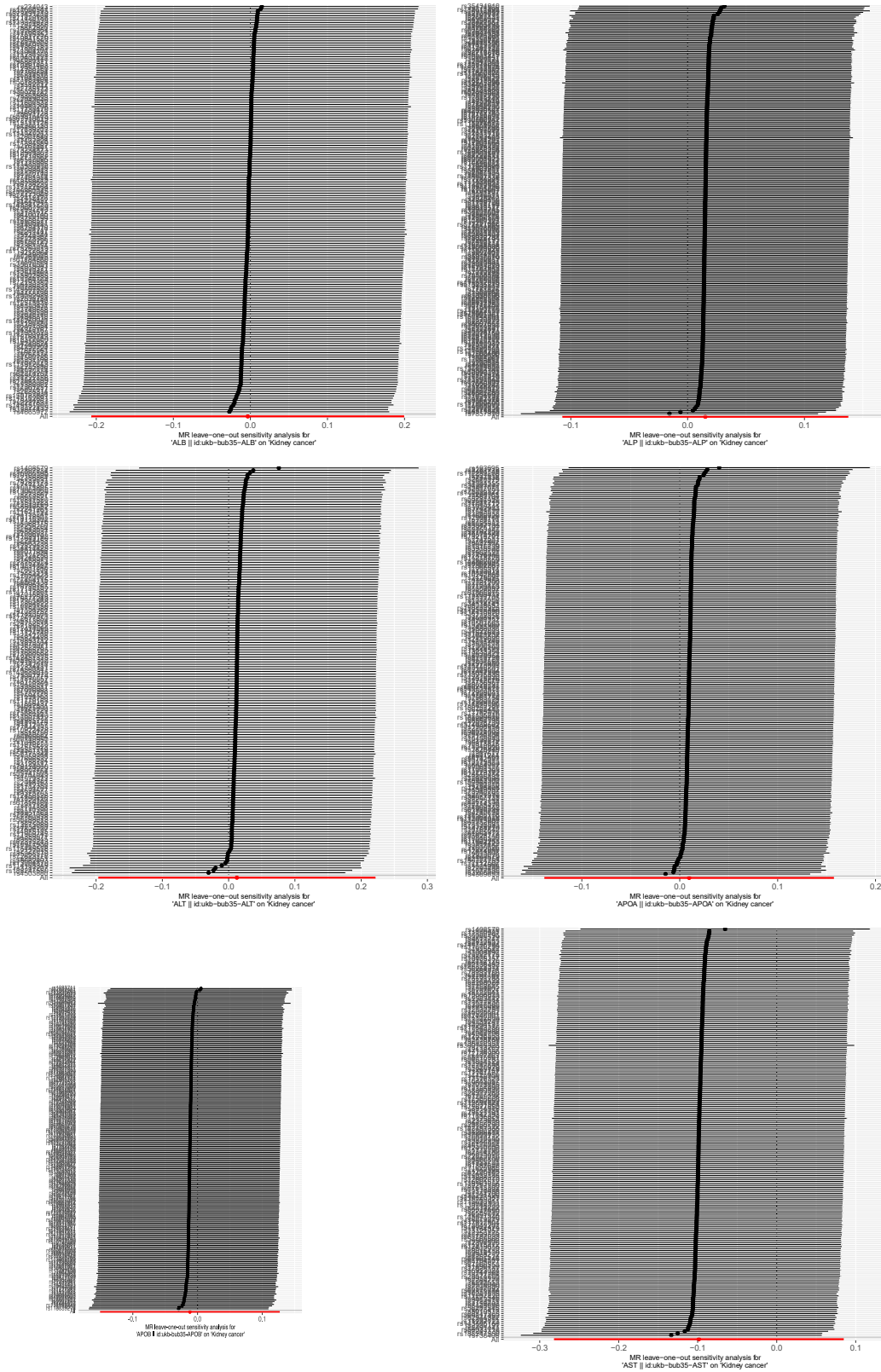


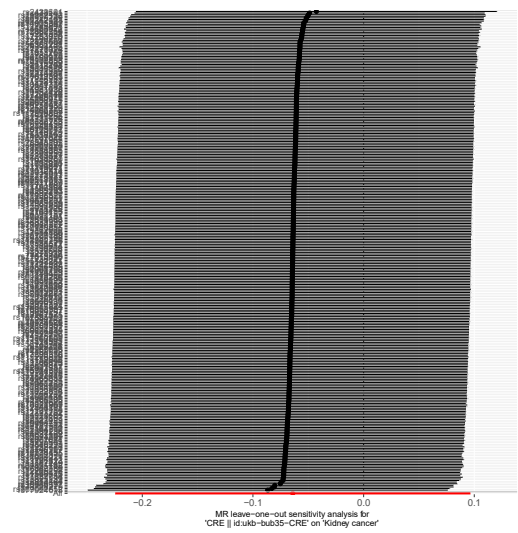
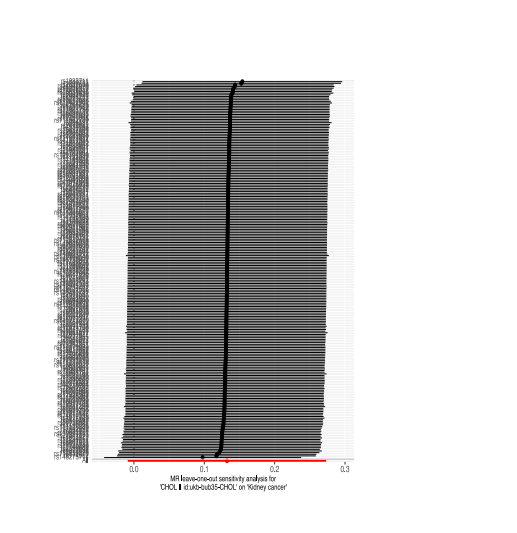
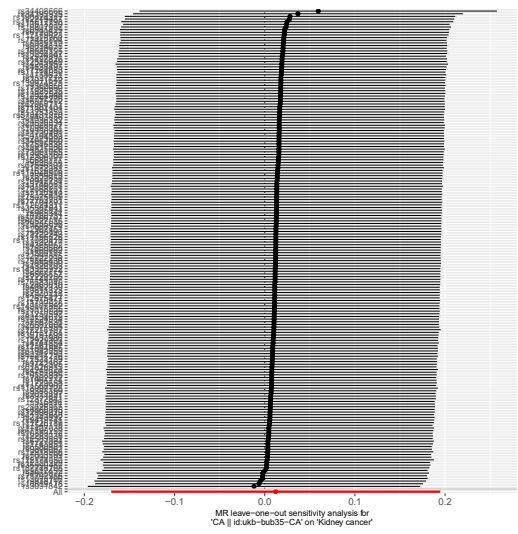
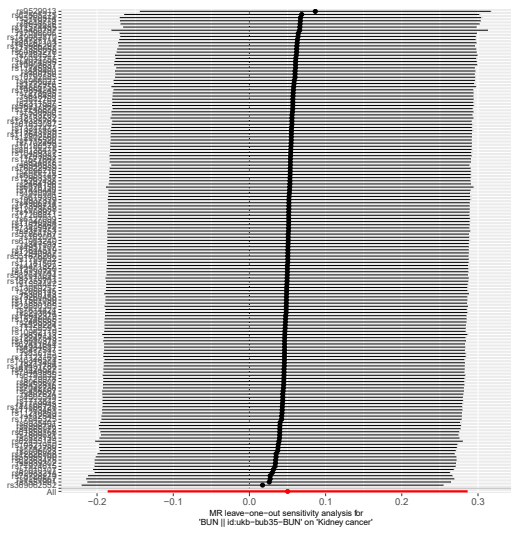
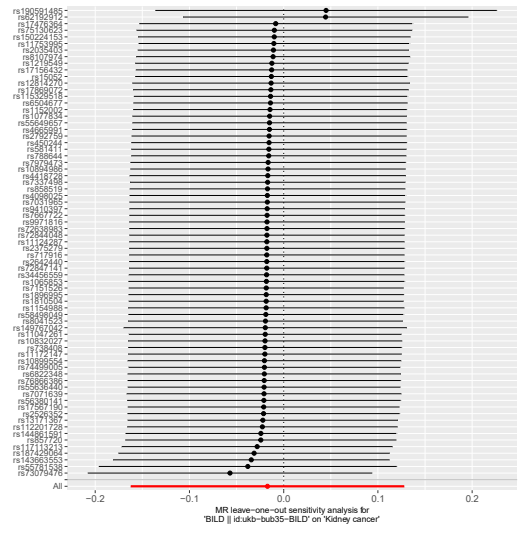
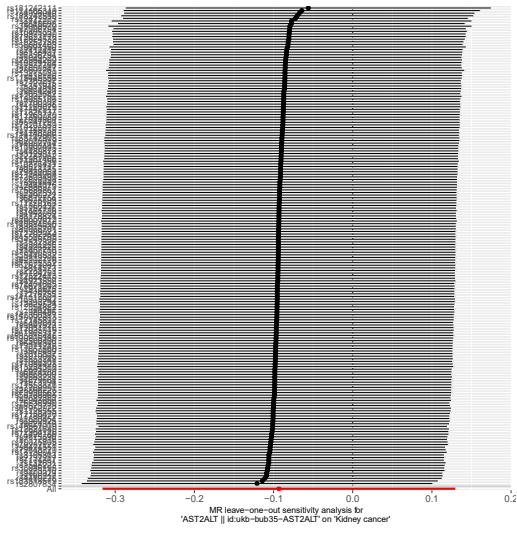


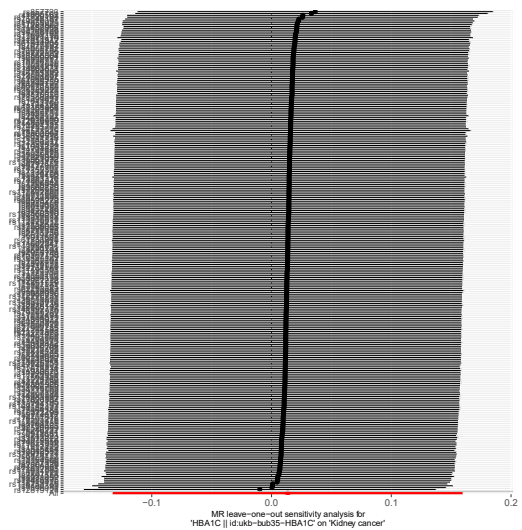
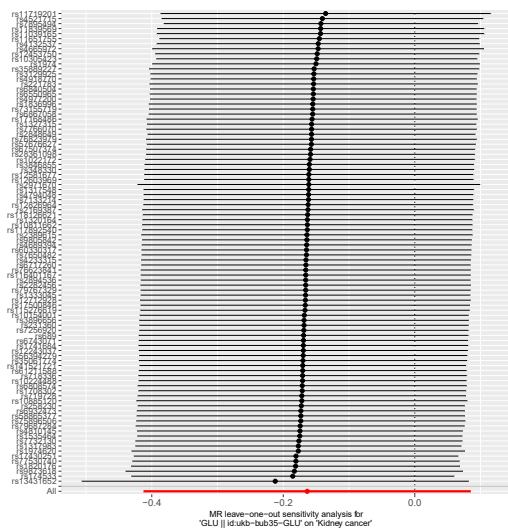
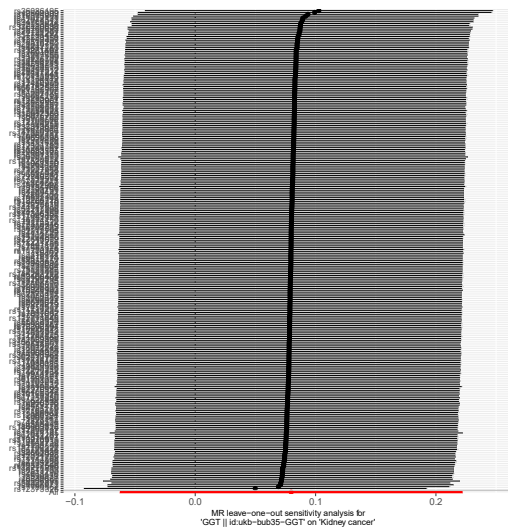
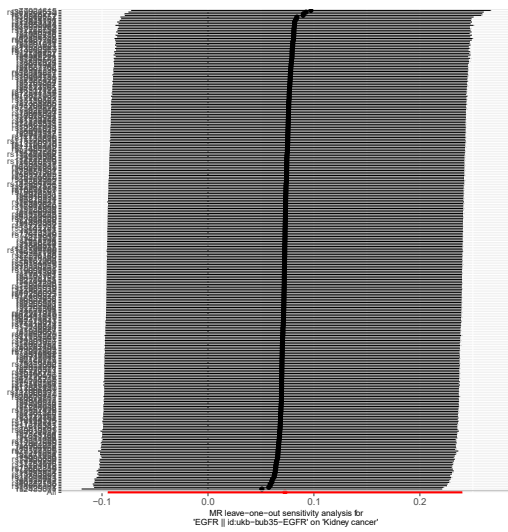
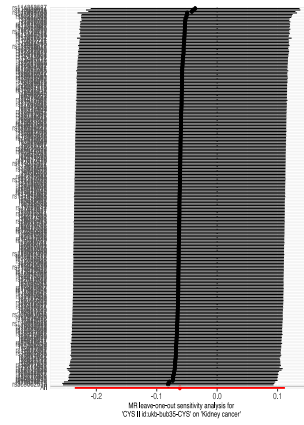
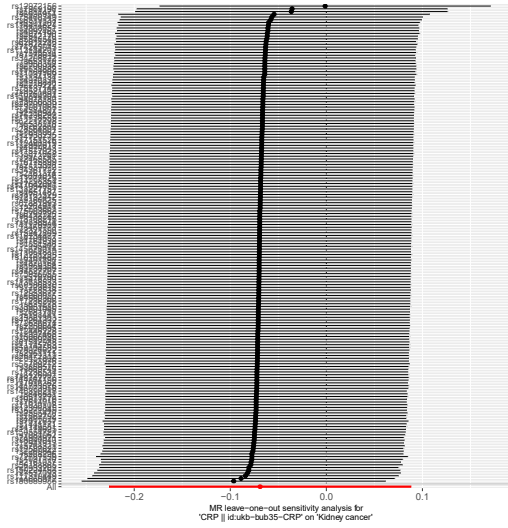


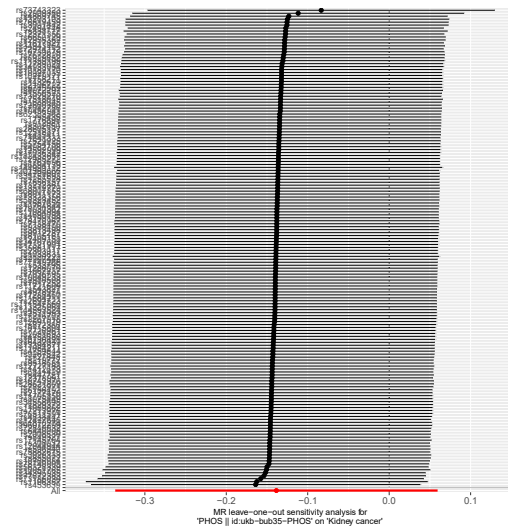
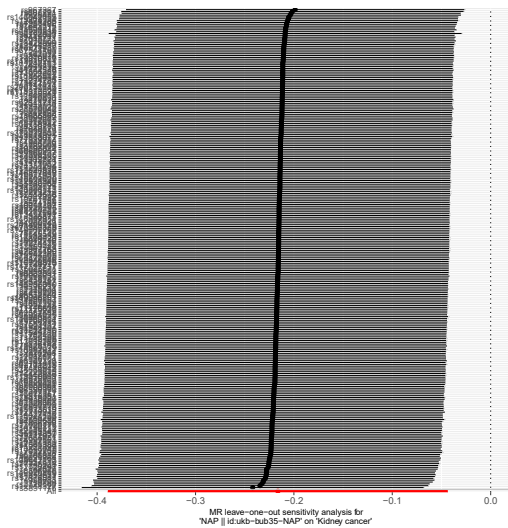
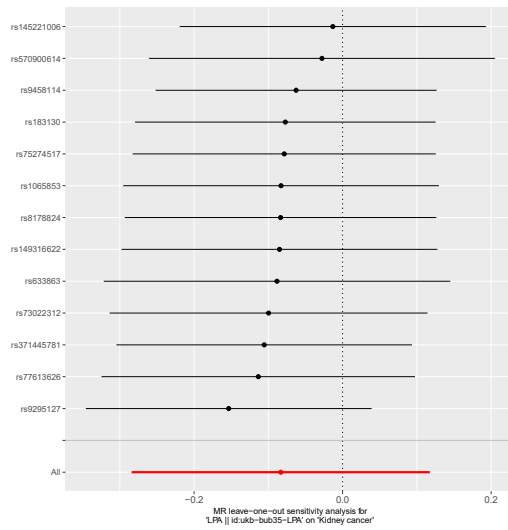
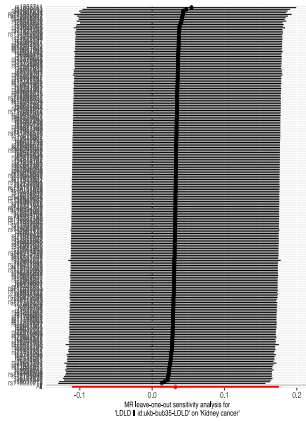
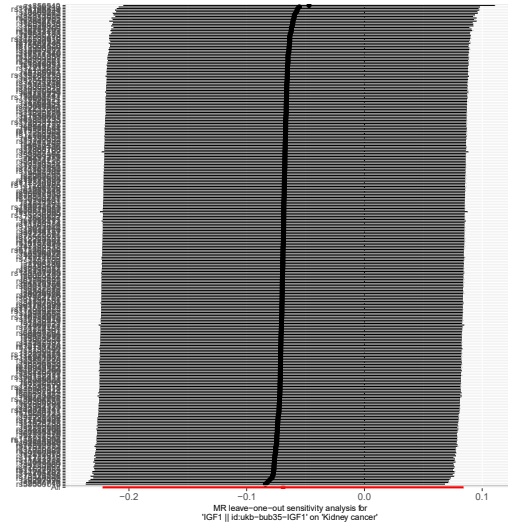
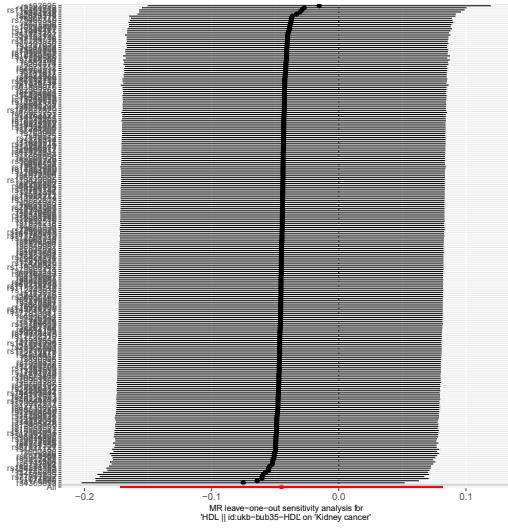


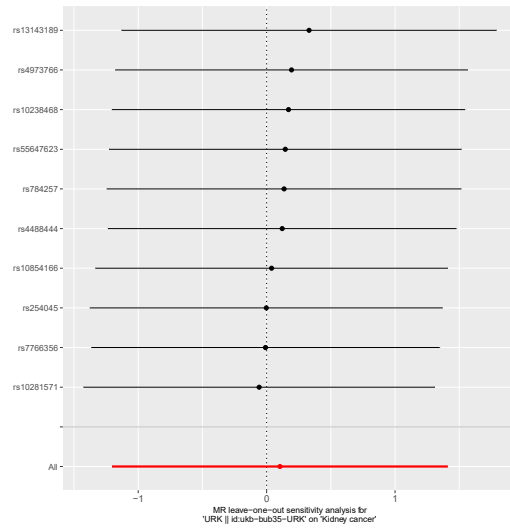
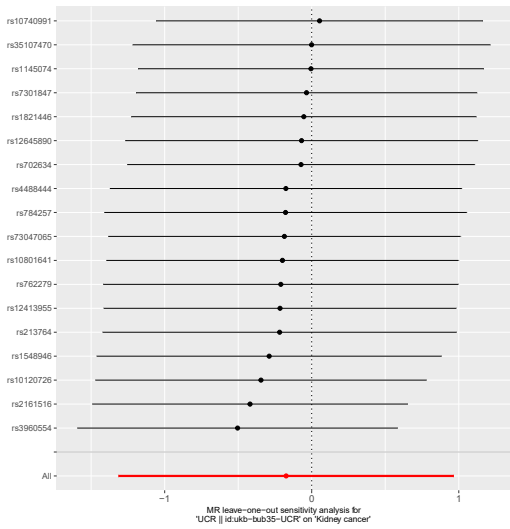
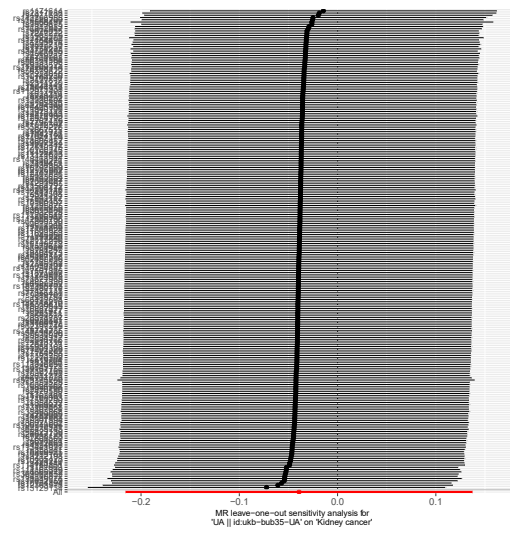
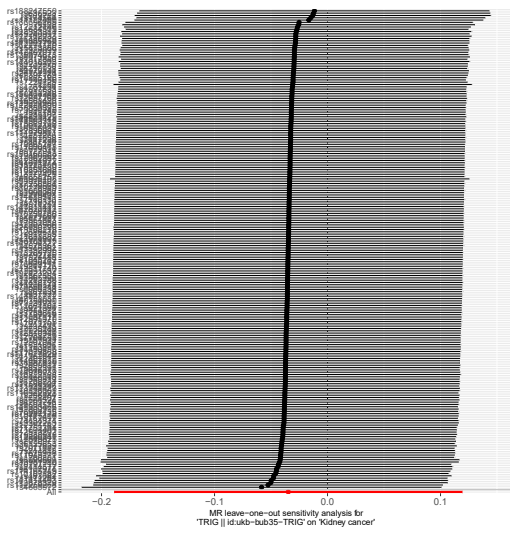
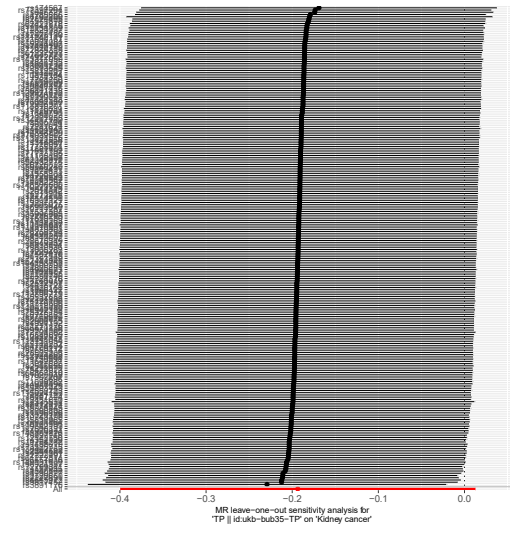
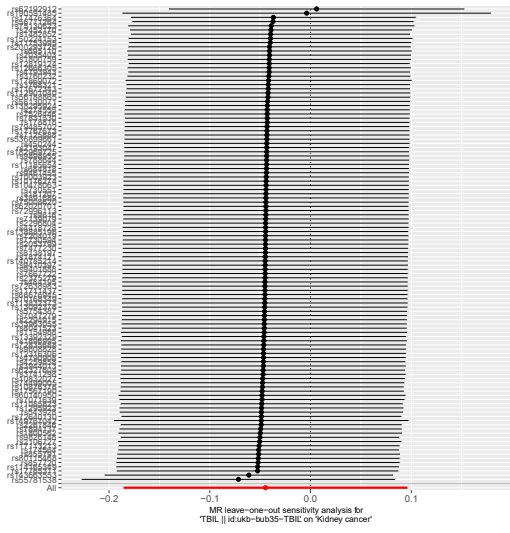
KCa's leave-one-out method

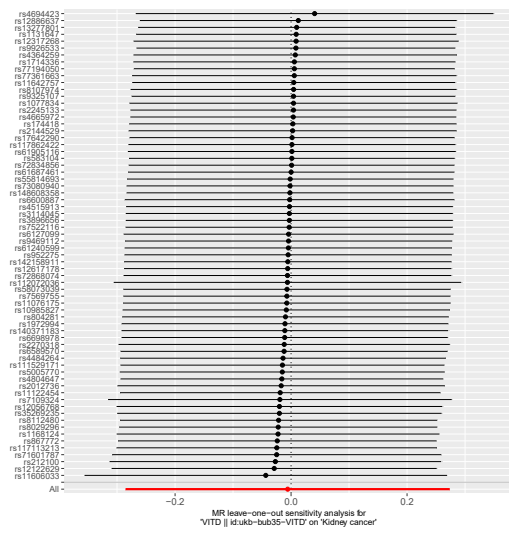
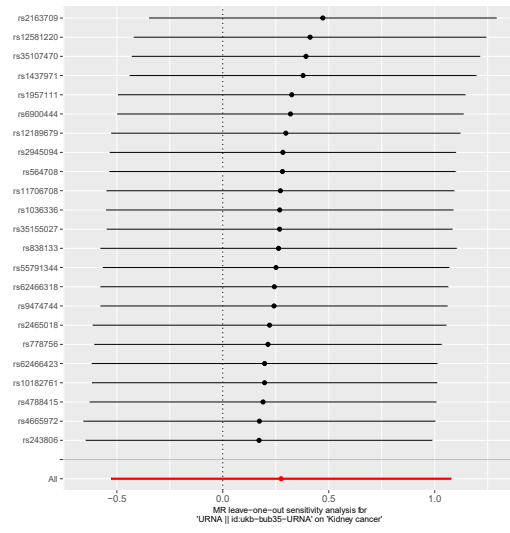
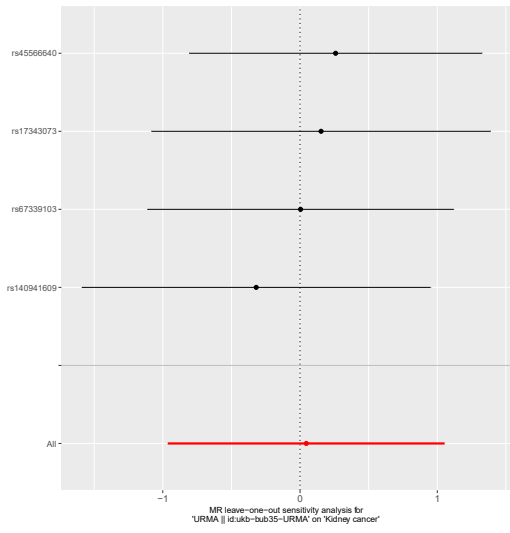




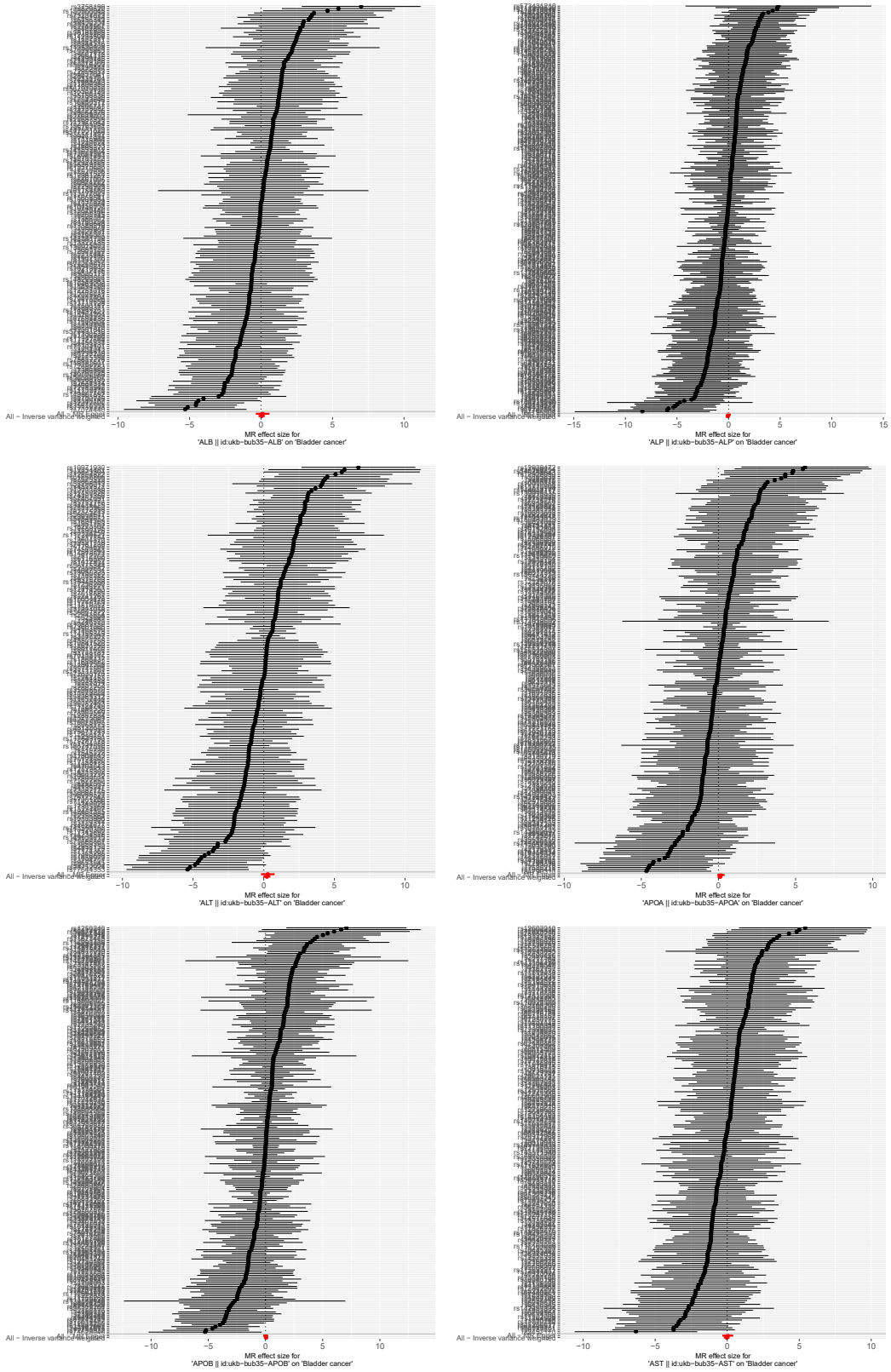


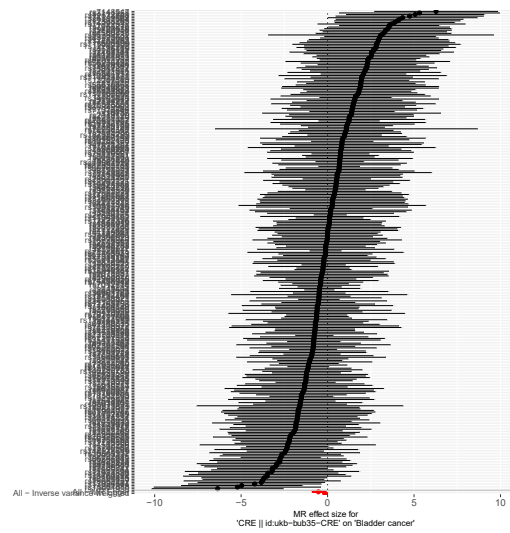
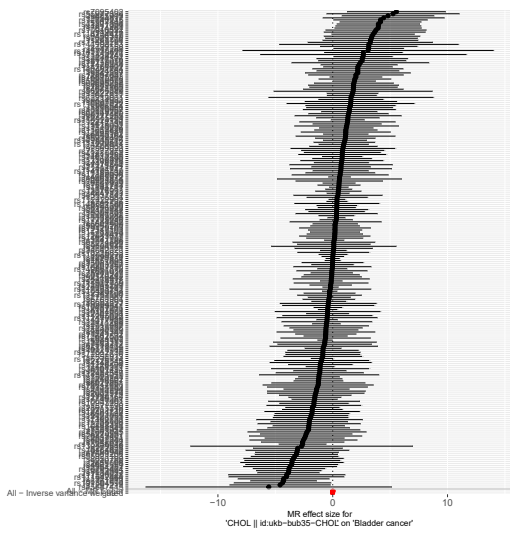
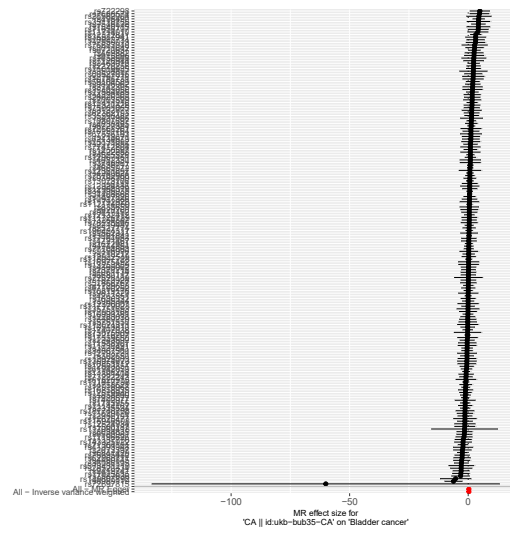
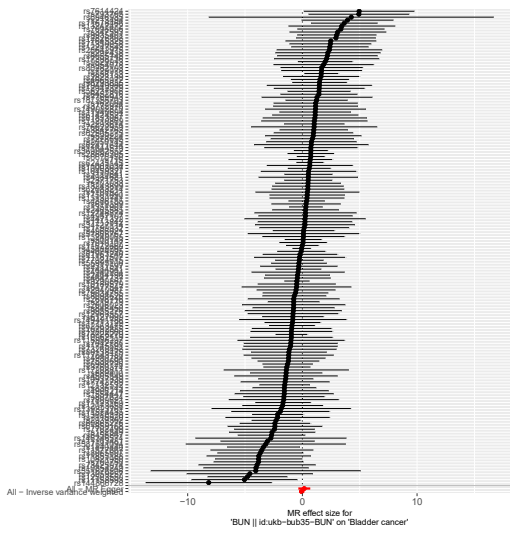
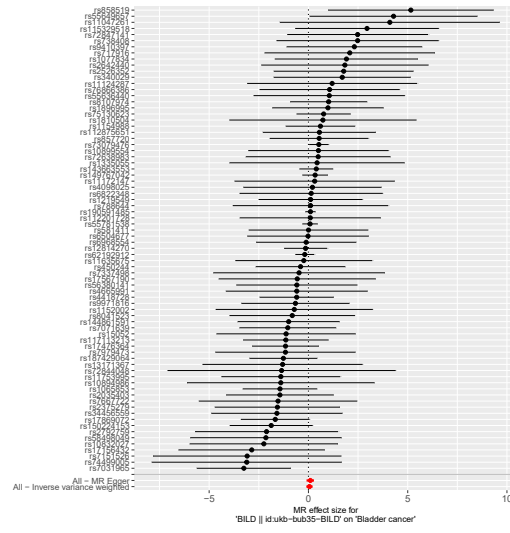
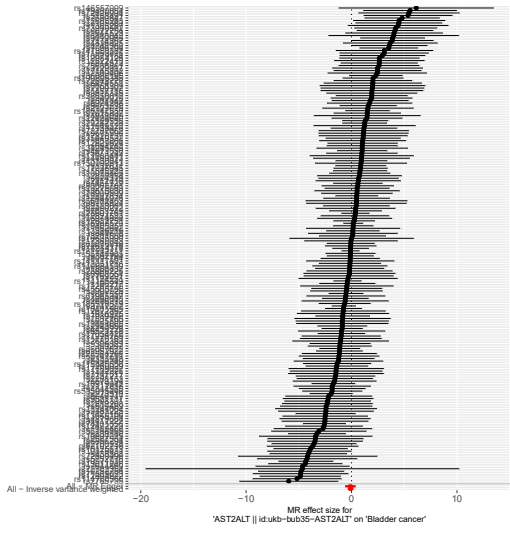


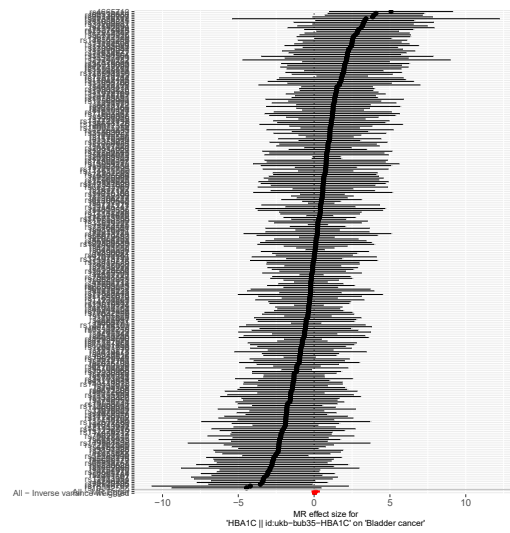
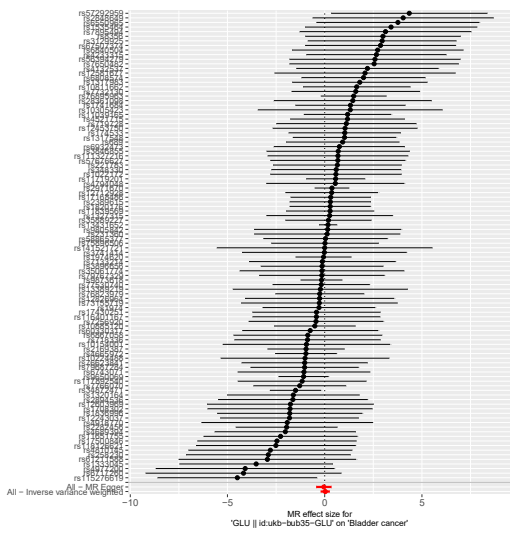
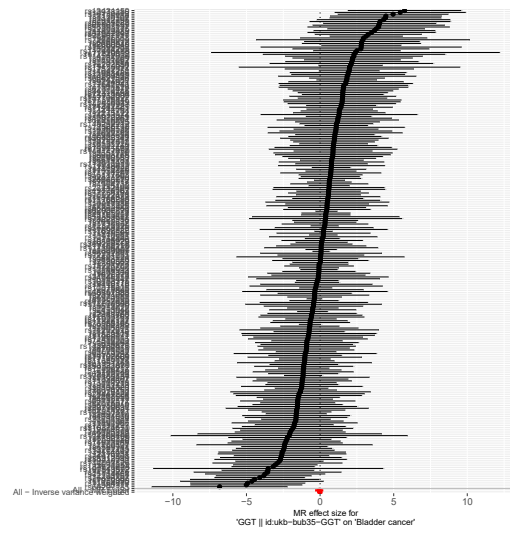
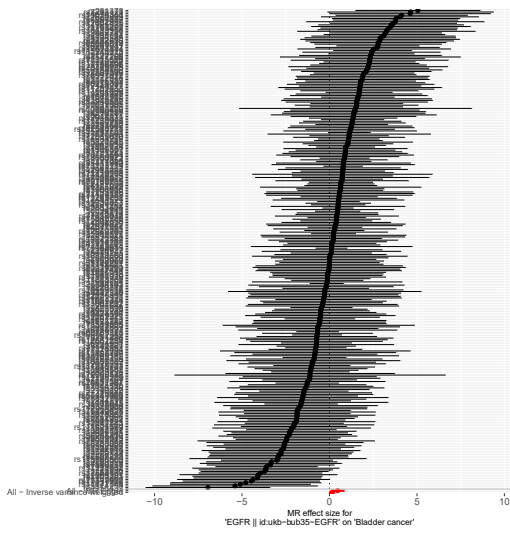
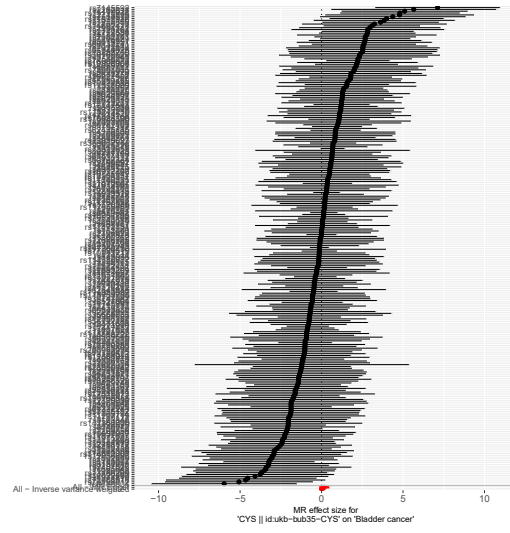
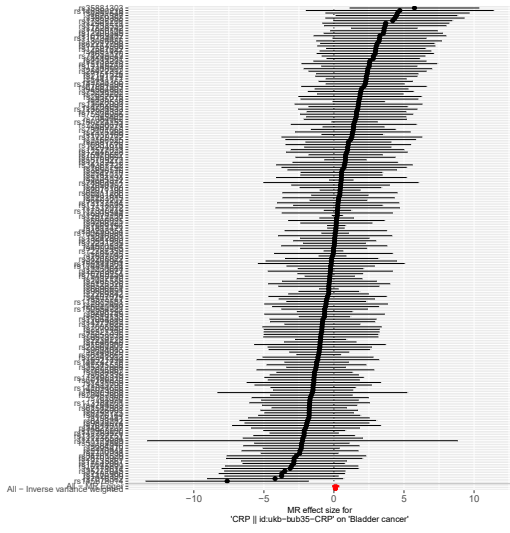


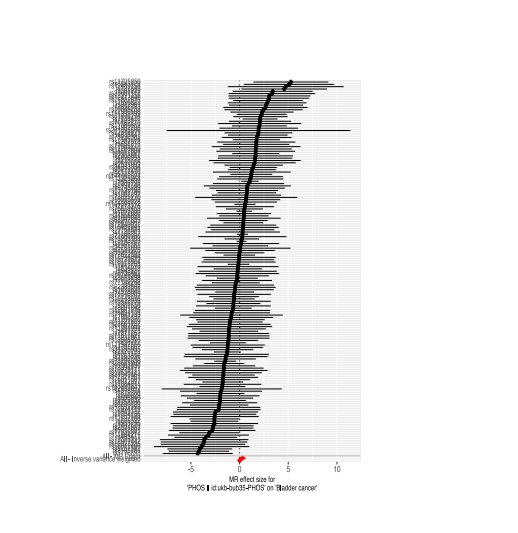
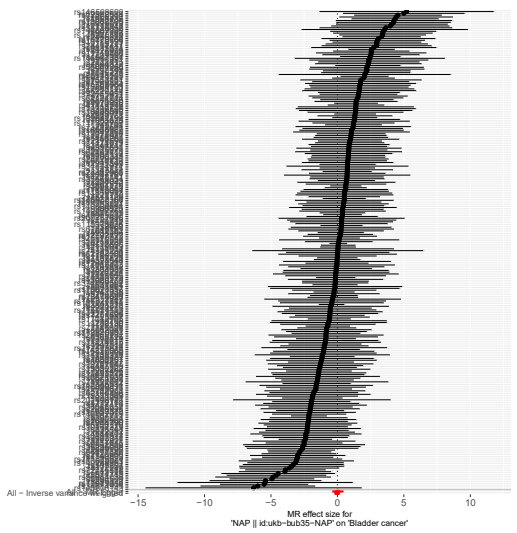
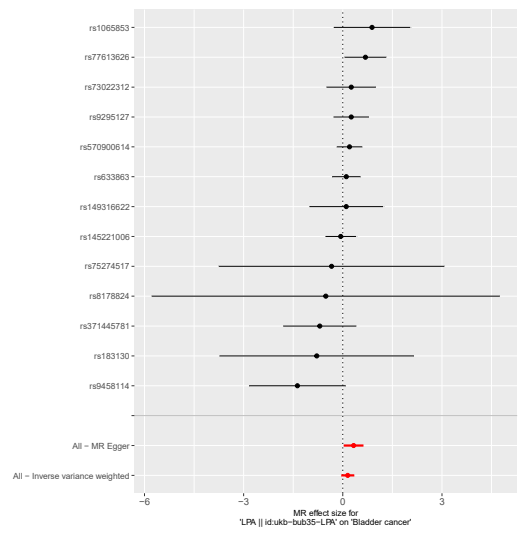
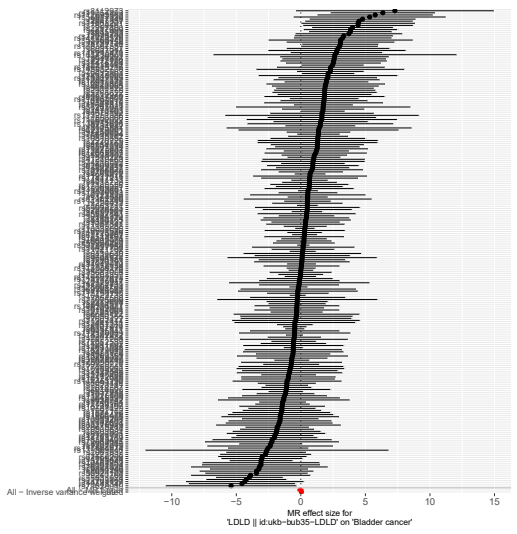
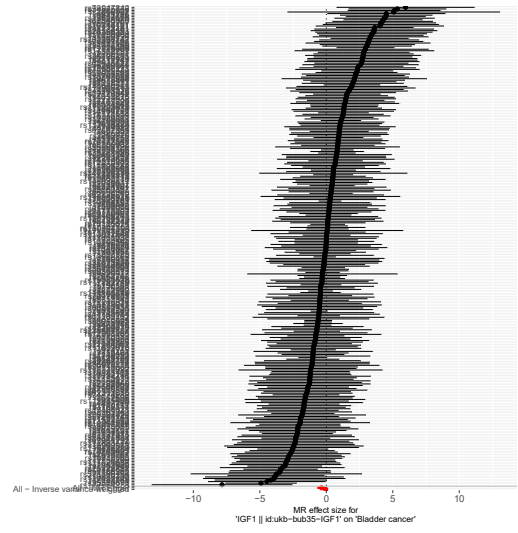
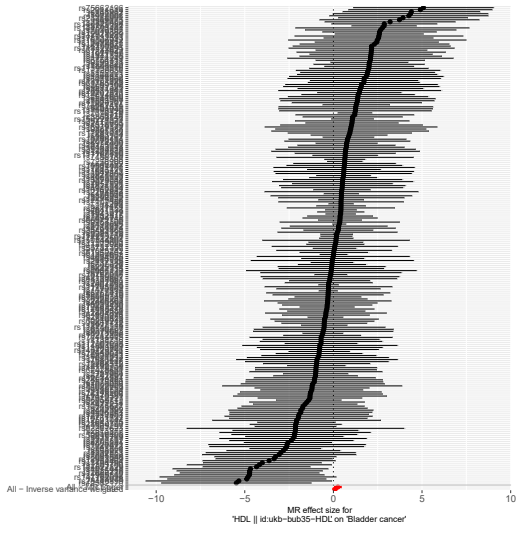


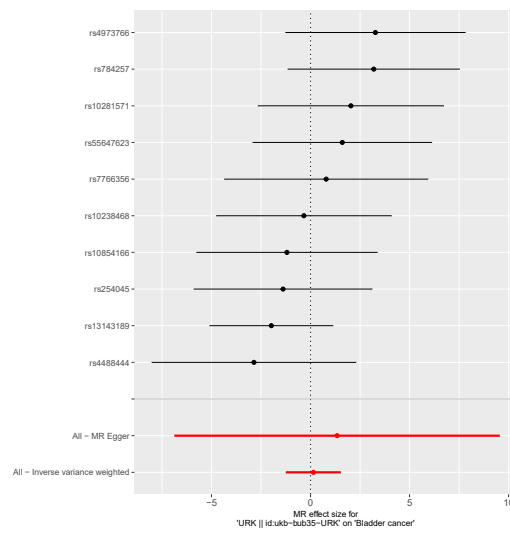
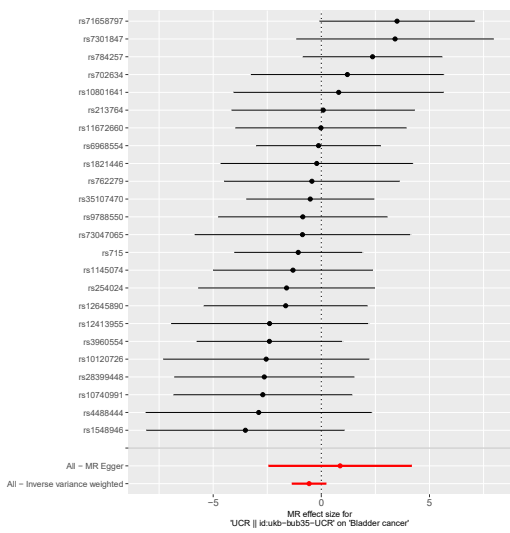
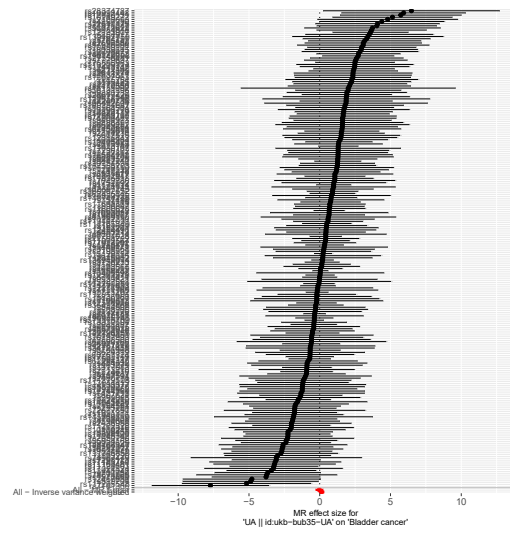
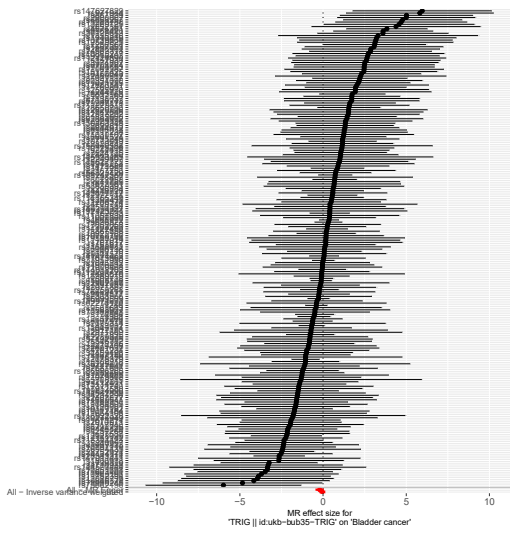
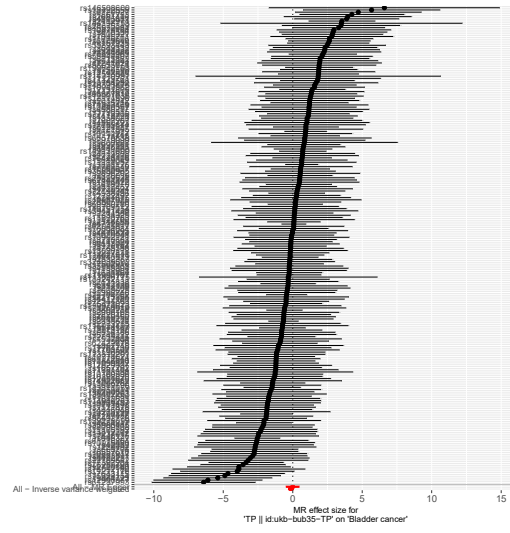
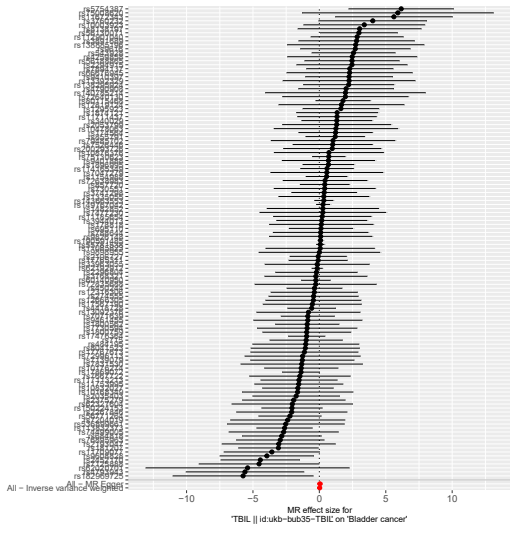
Forest plot of BCa

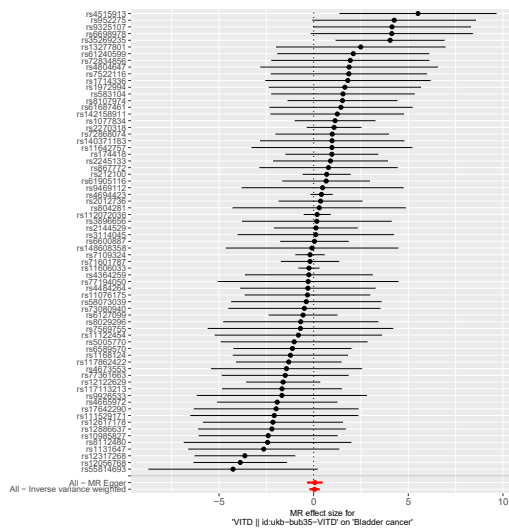
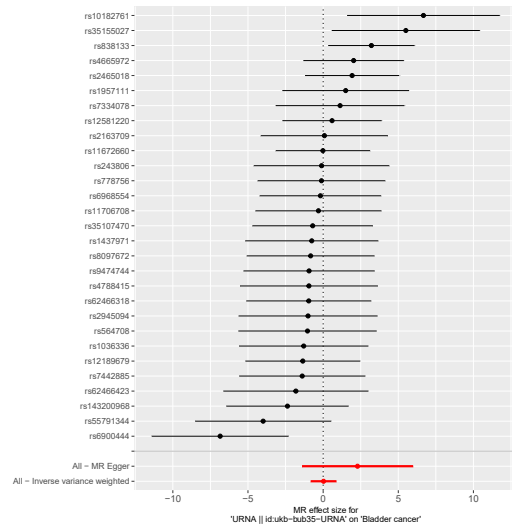
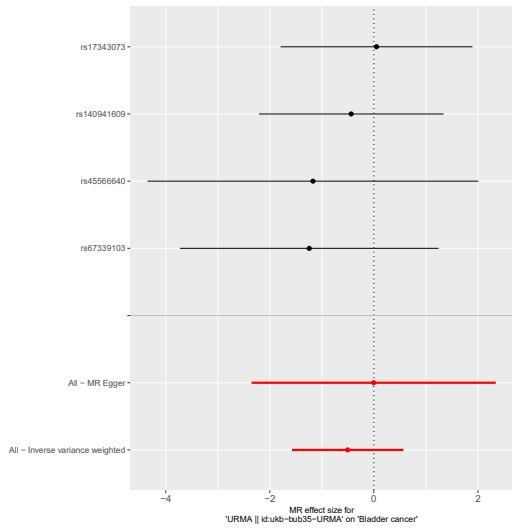




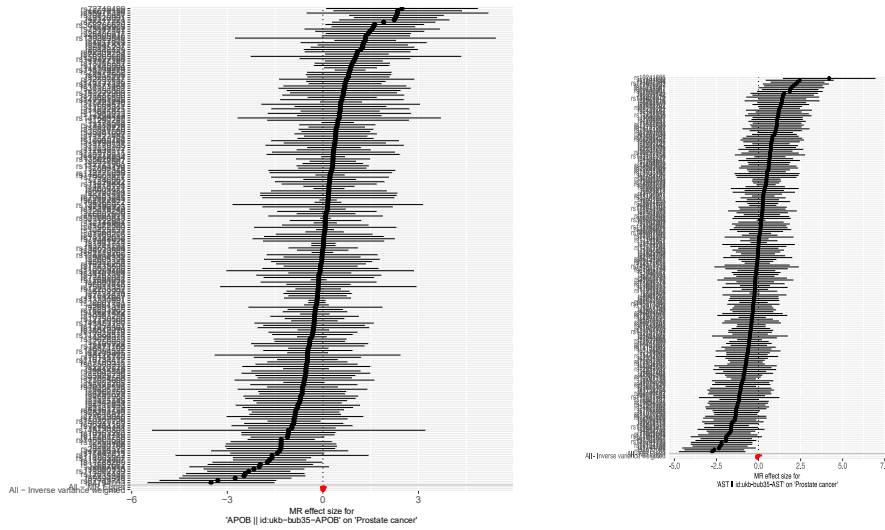
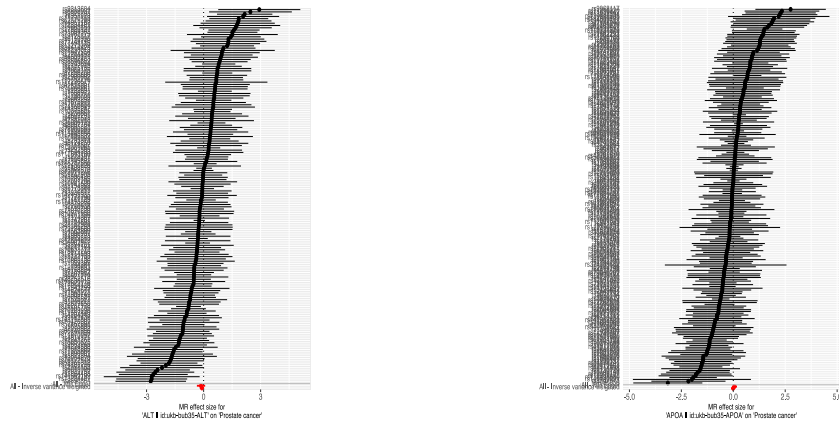
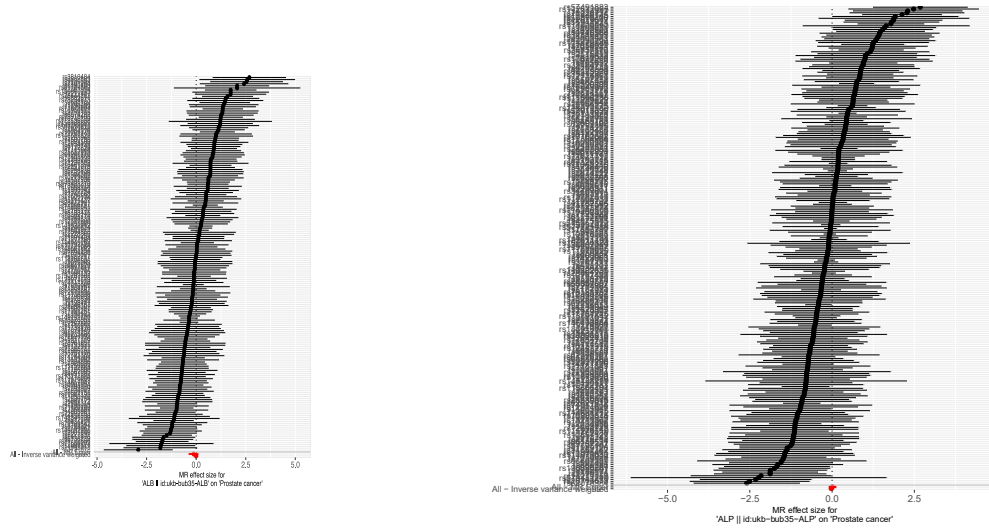


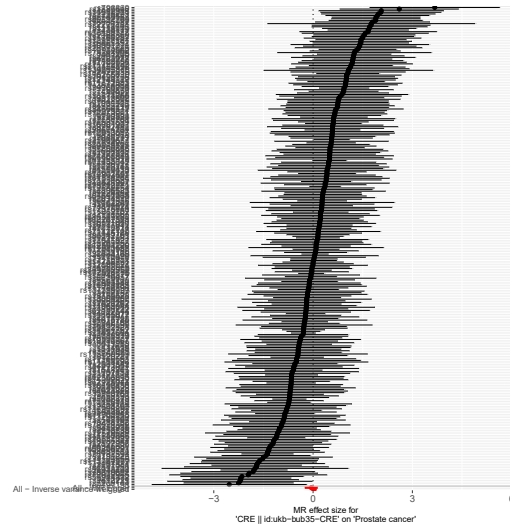
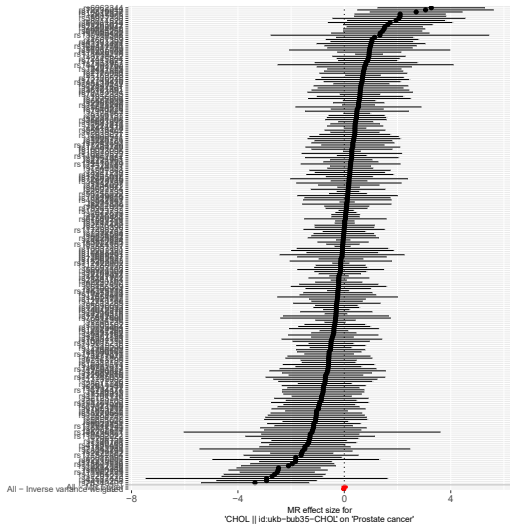
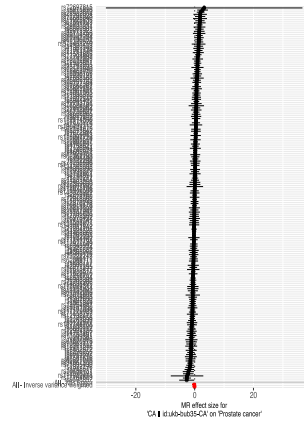
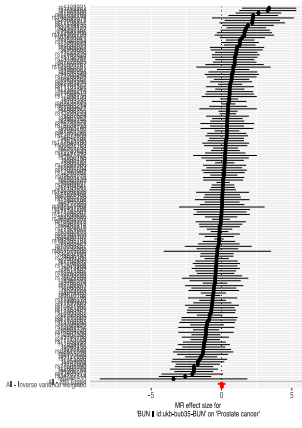
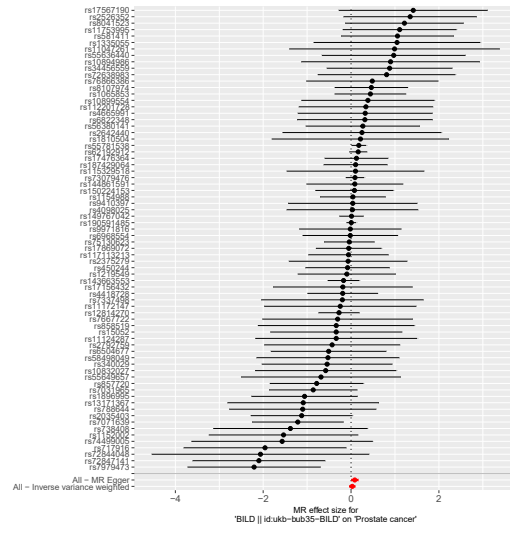
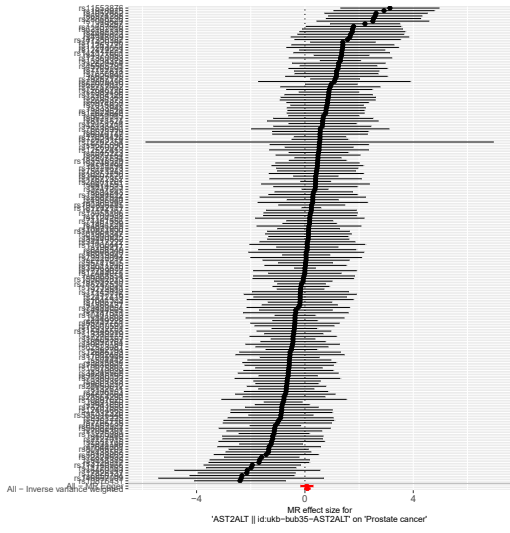


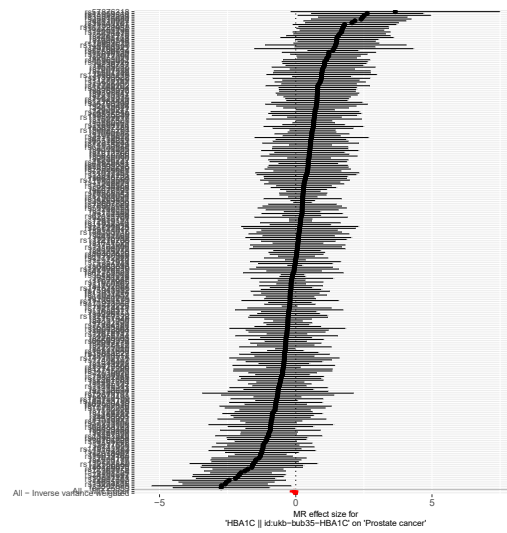
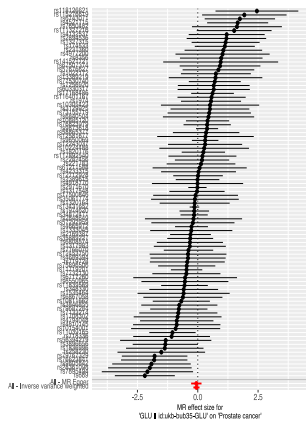
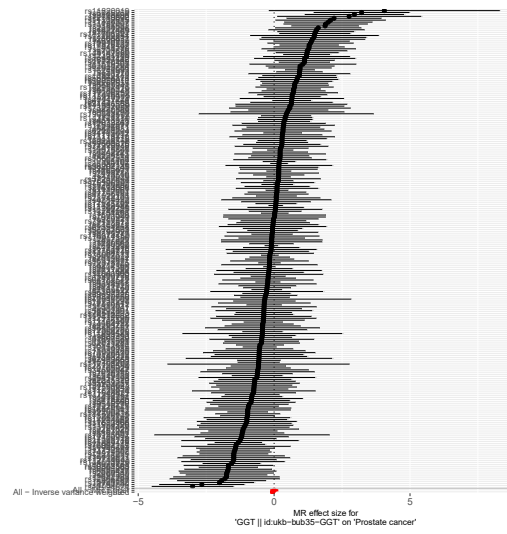
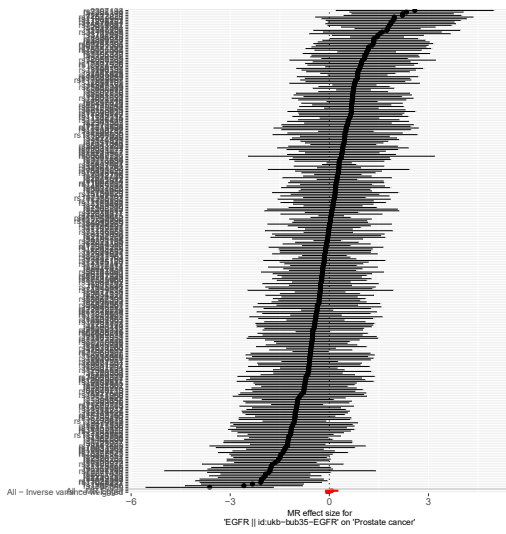
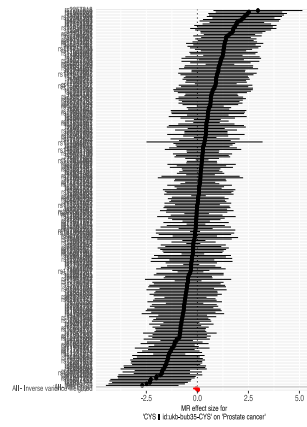
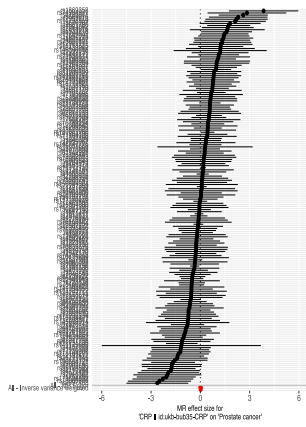


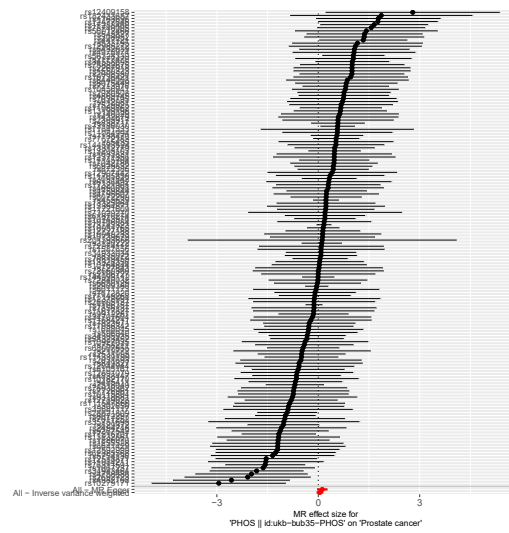
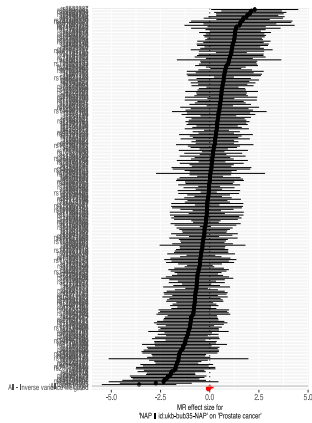
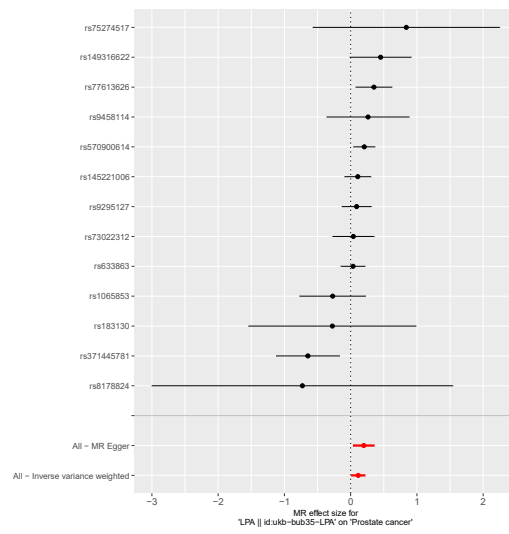
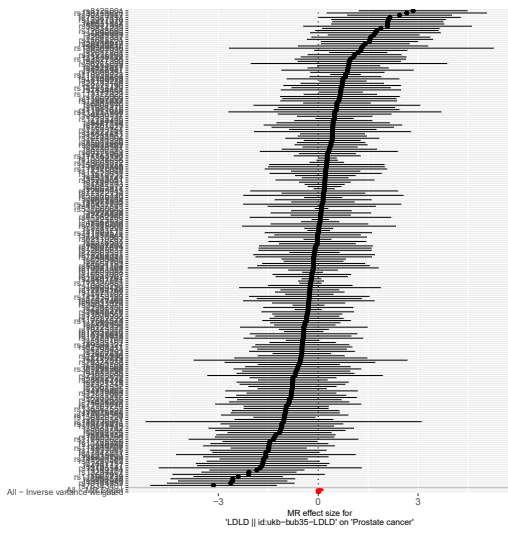
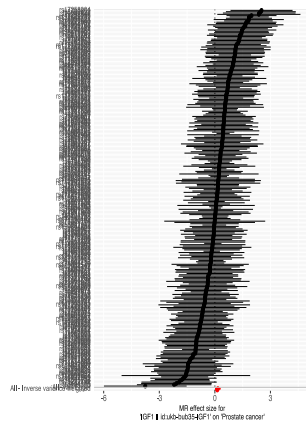
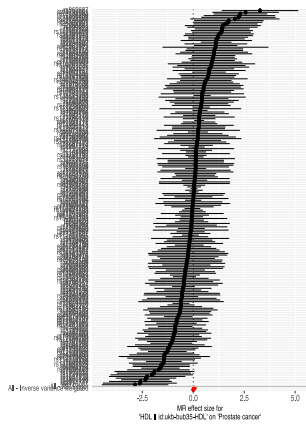


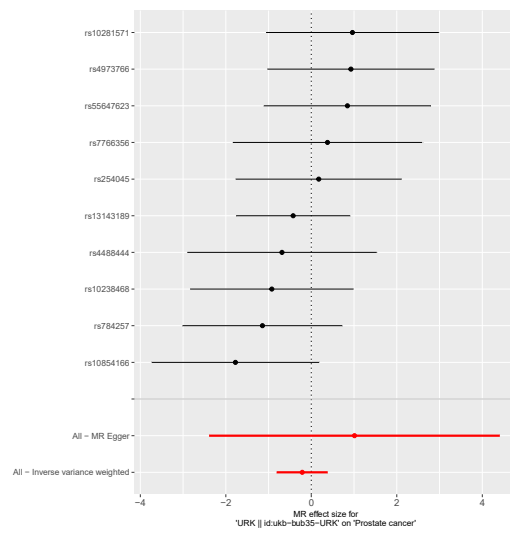
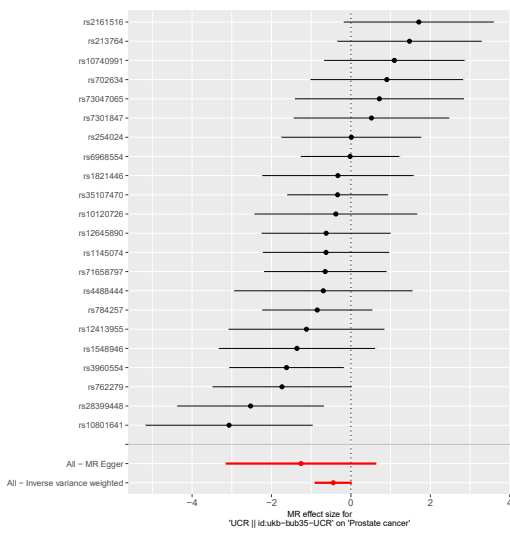
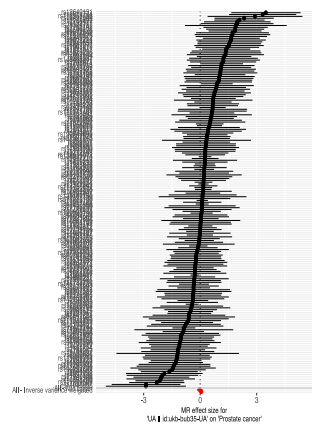
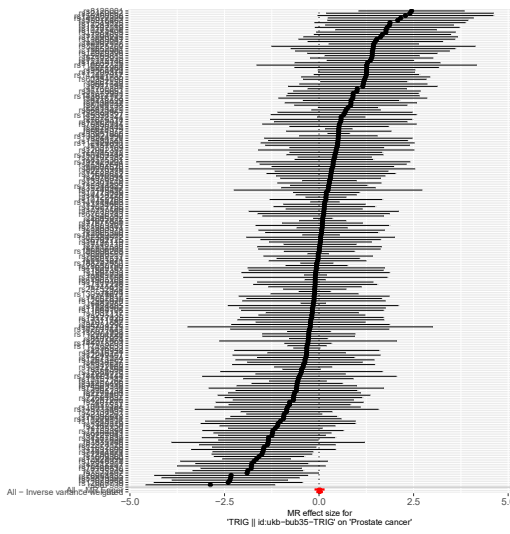
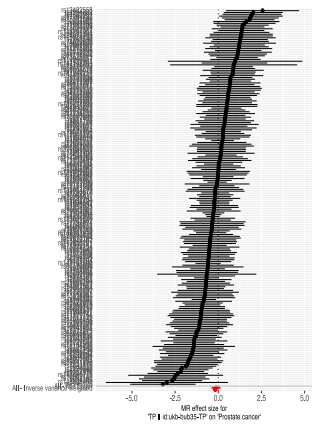
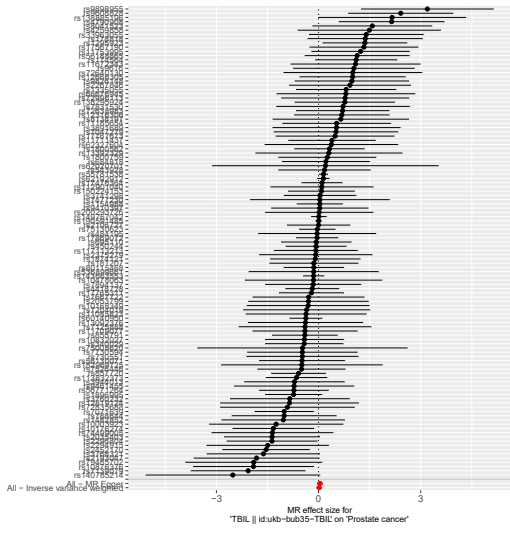
Forest plot of PCa

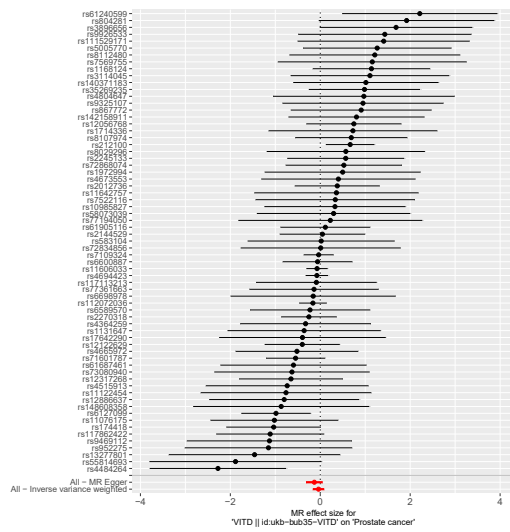
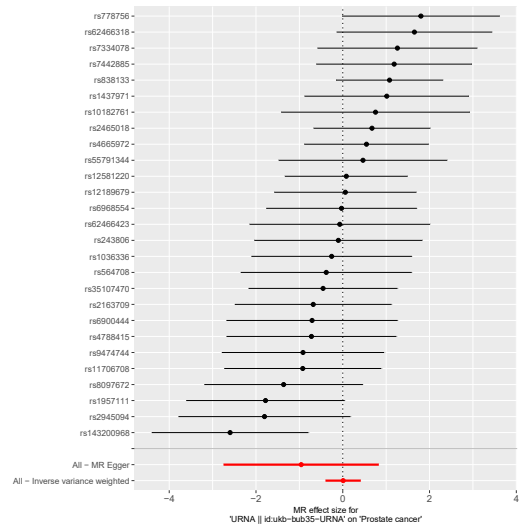
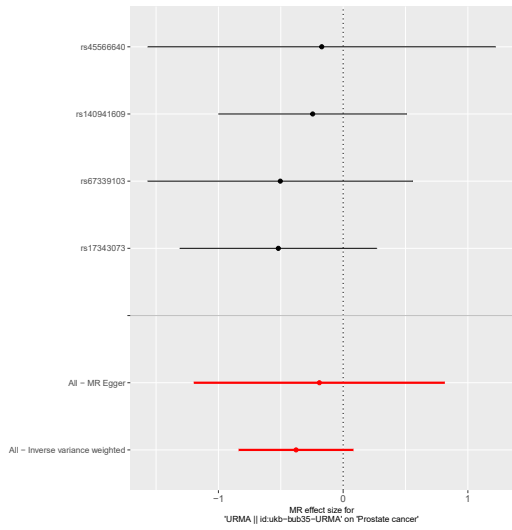




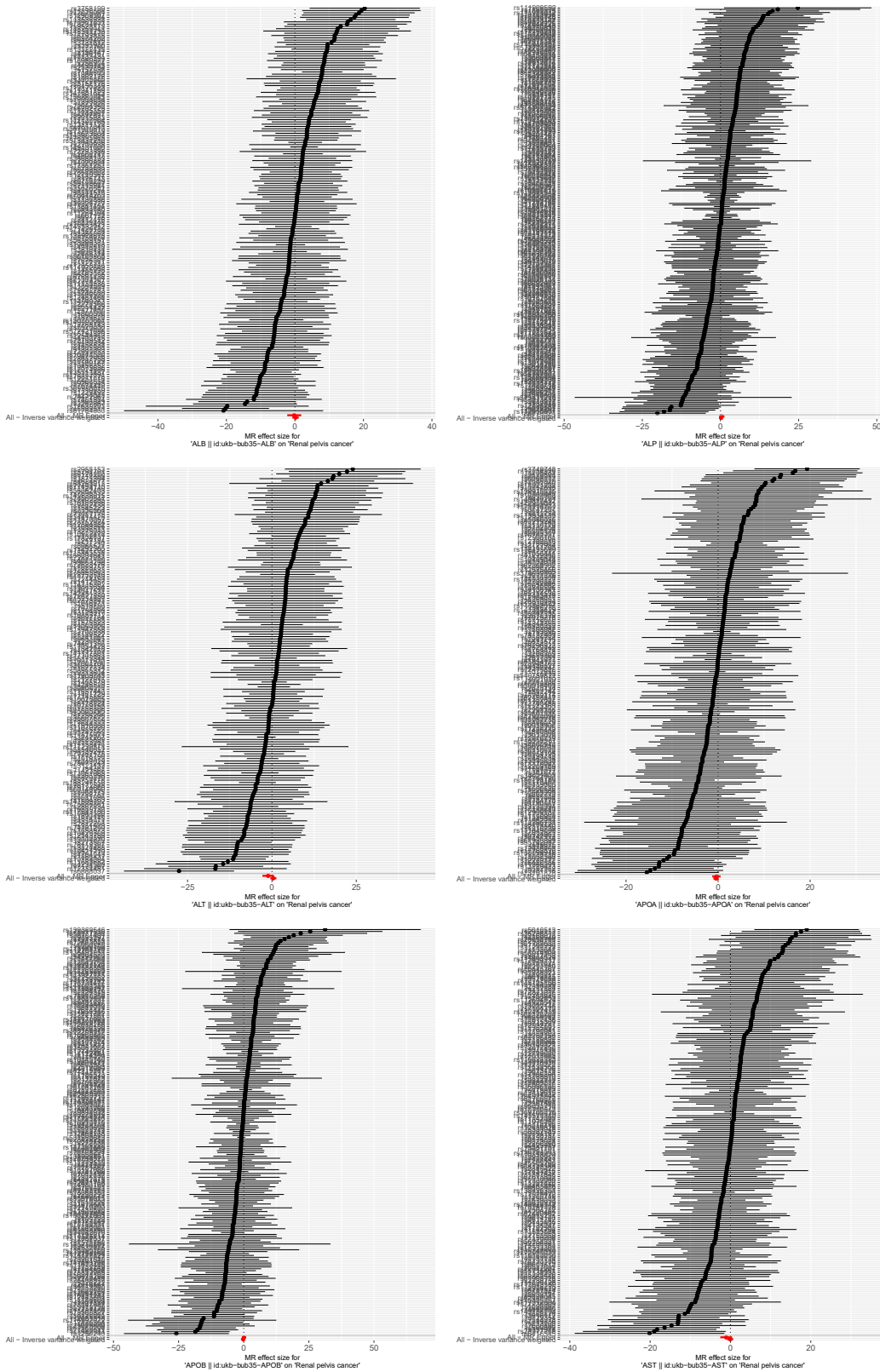


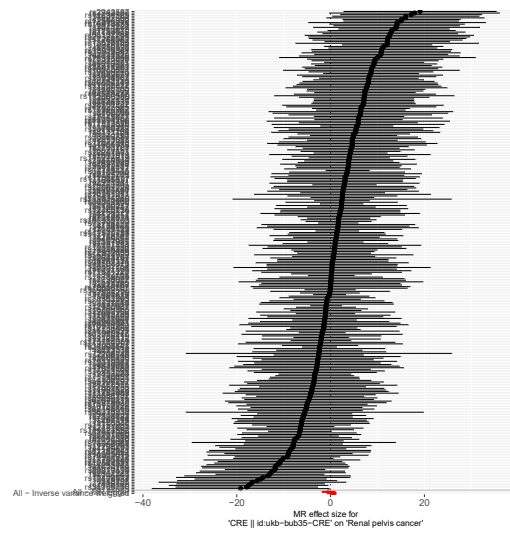
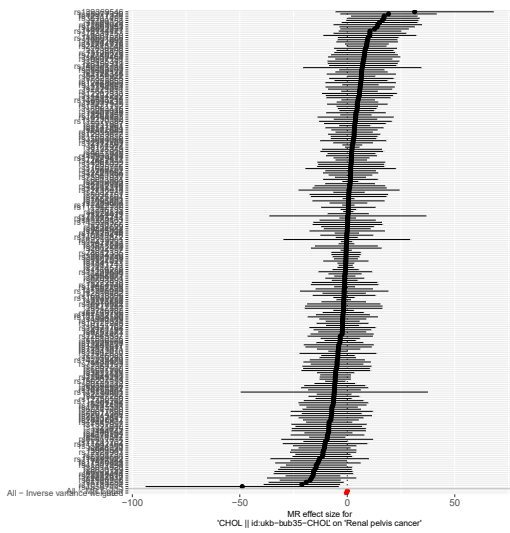
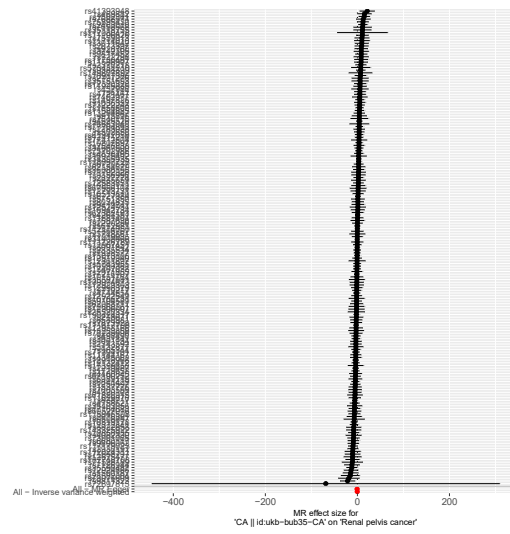
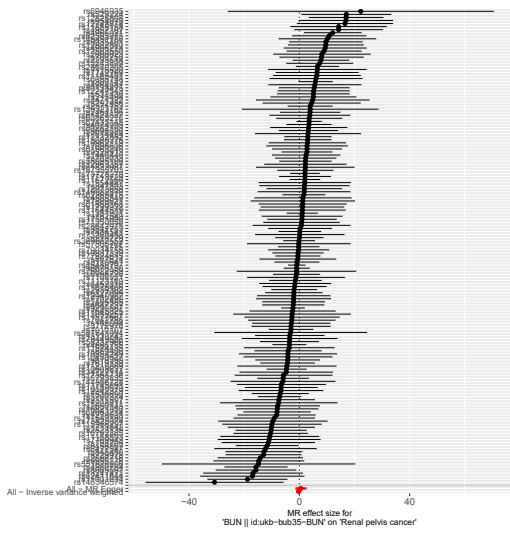
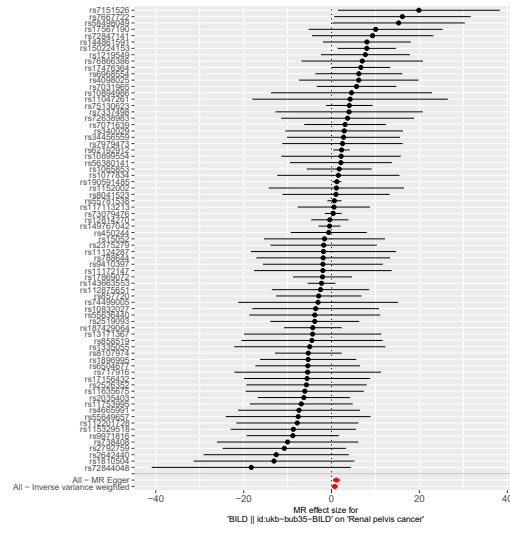
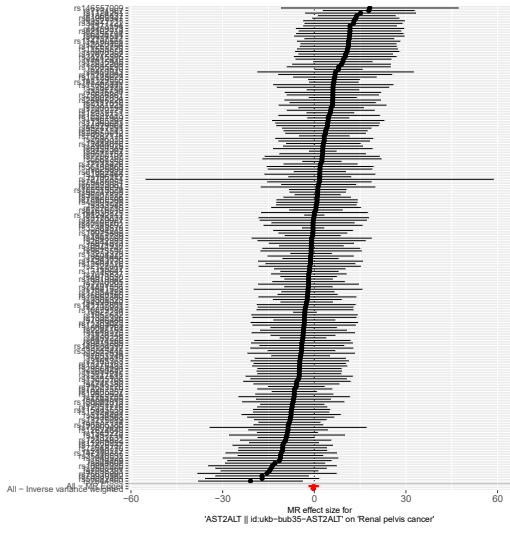


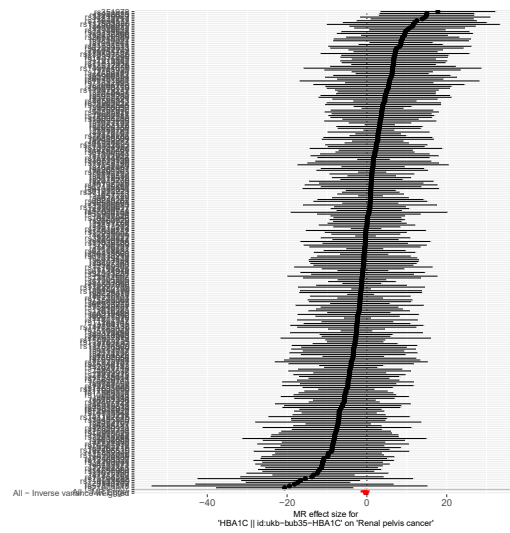
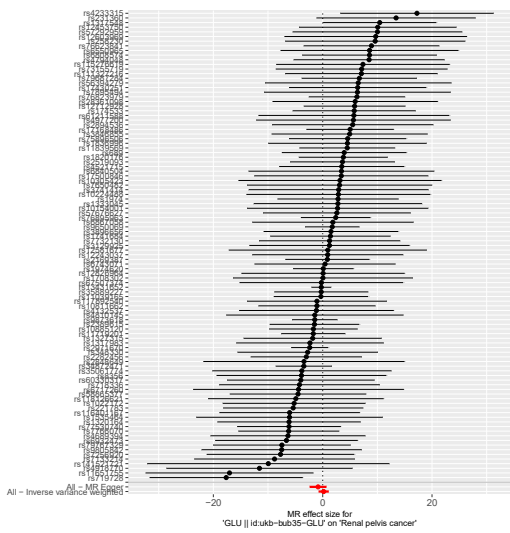
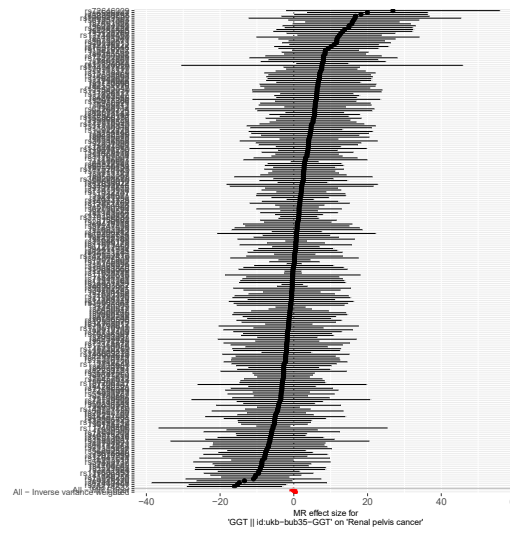
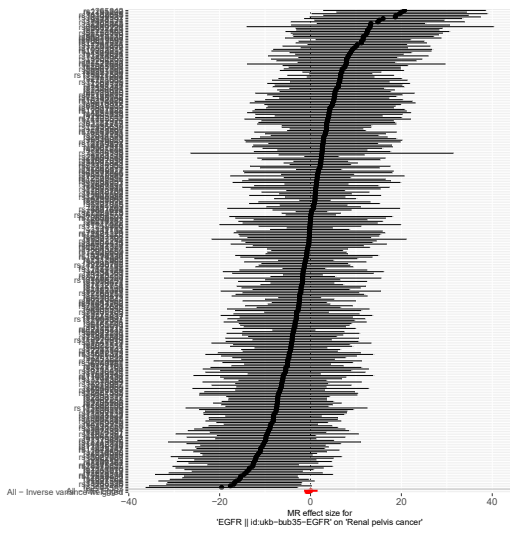
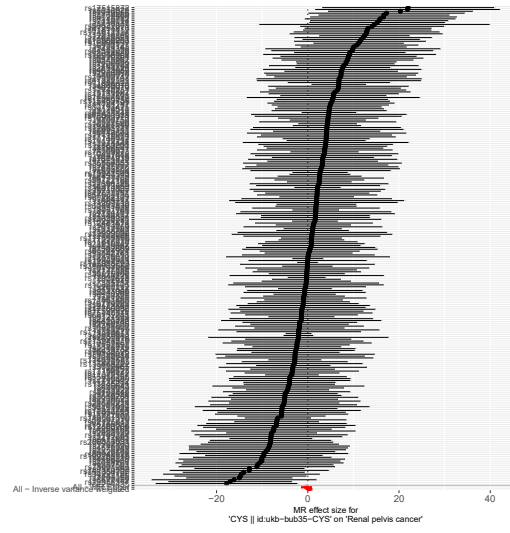
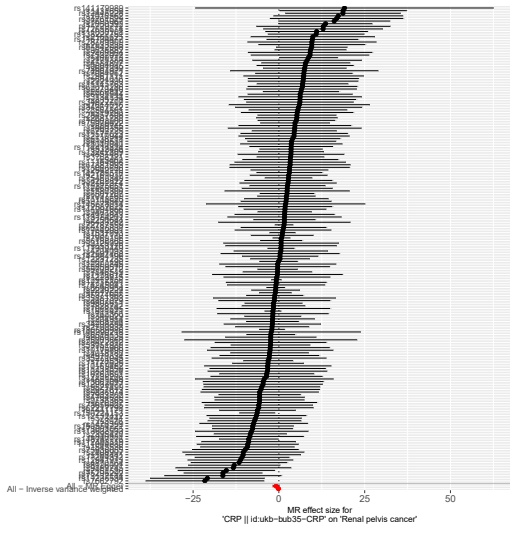


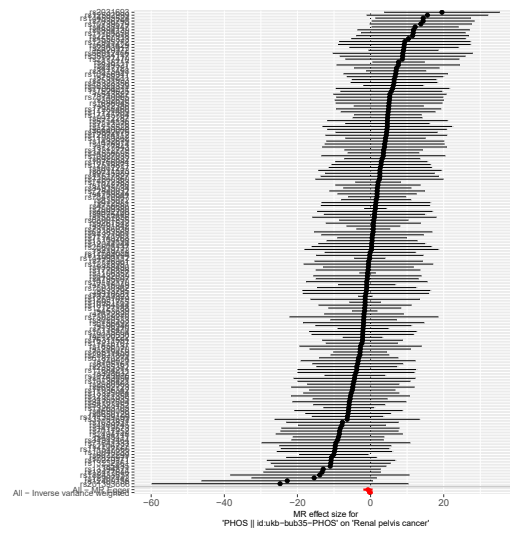
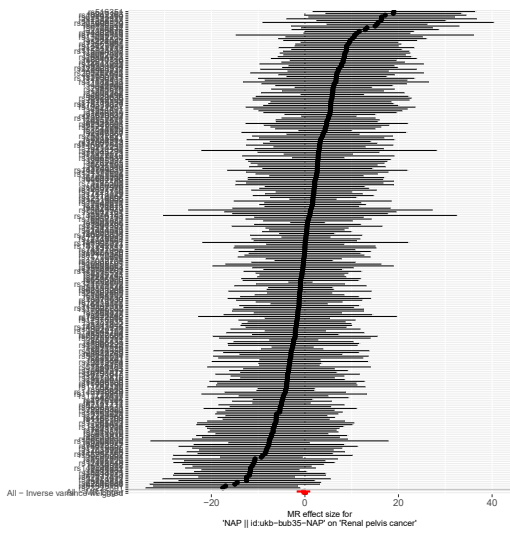
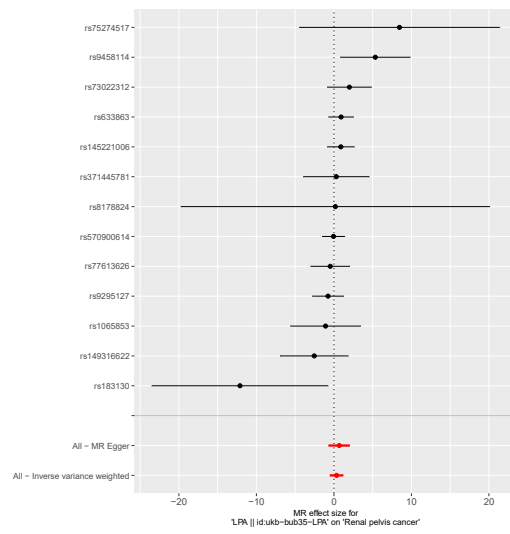
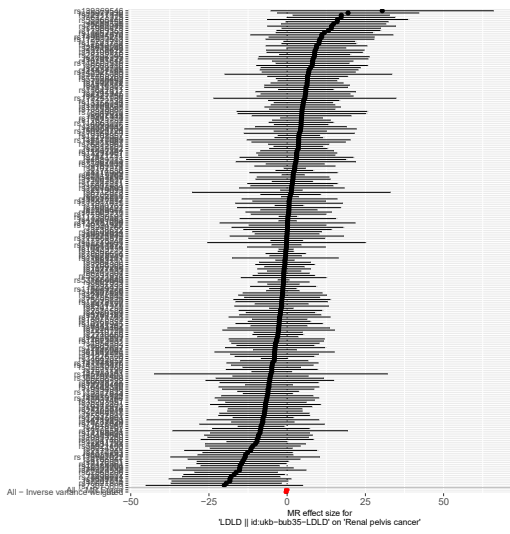
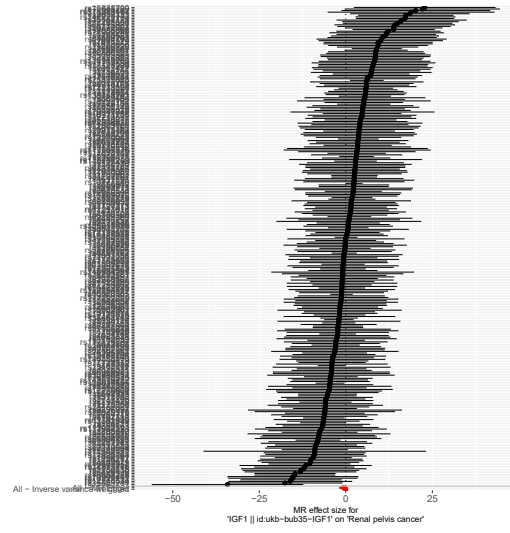
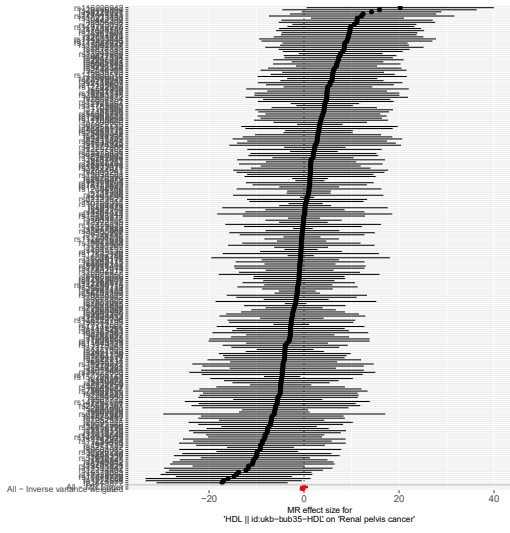


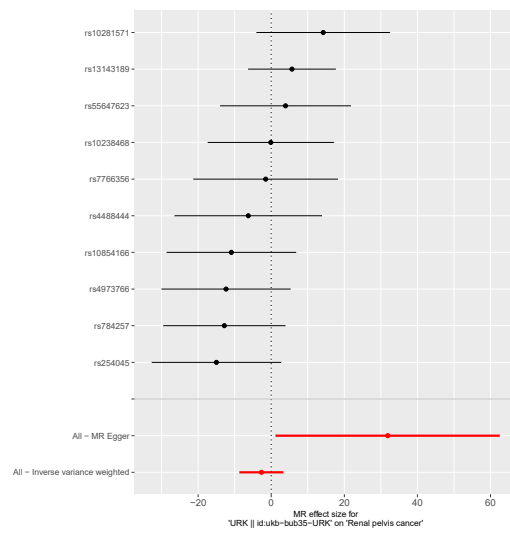
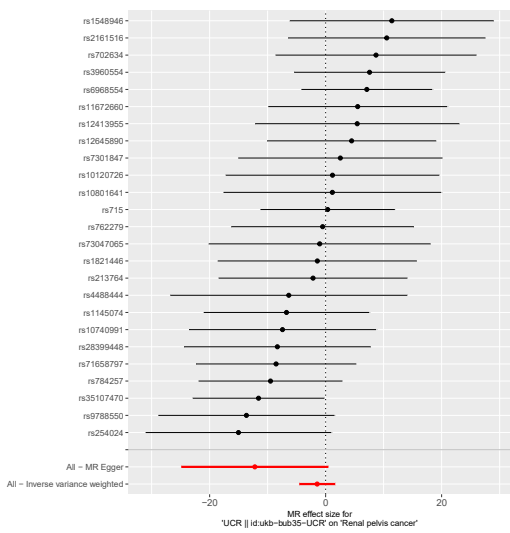
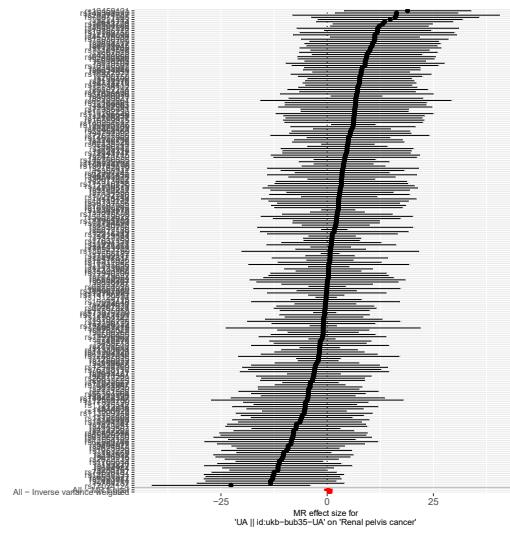
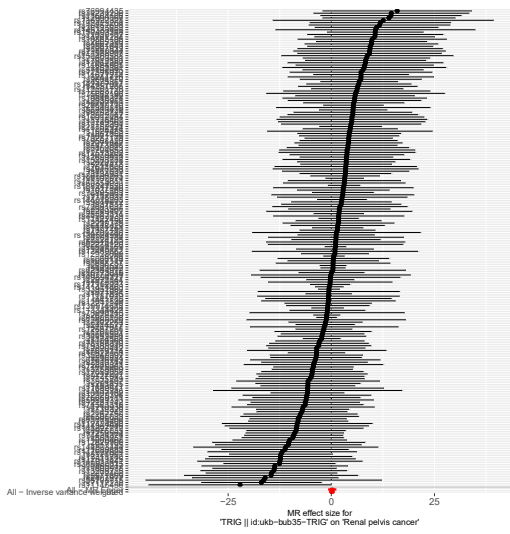
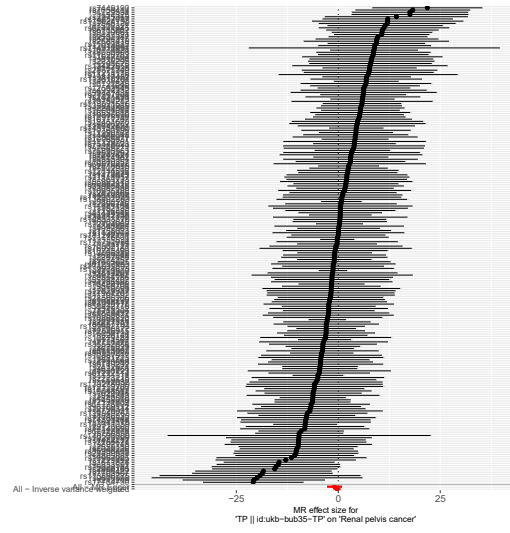
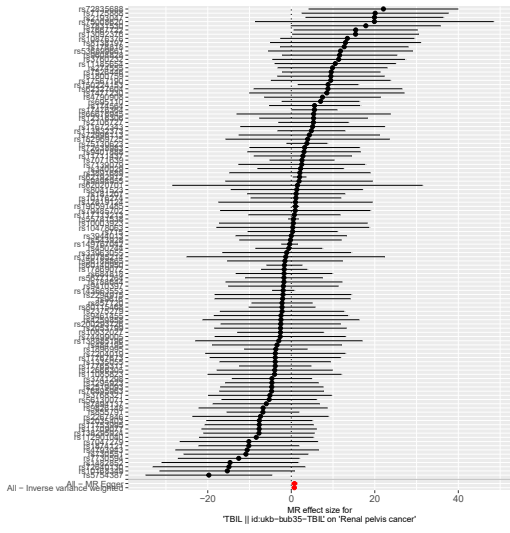
Forest plot of RPCa

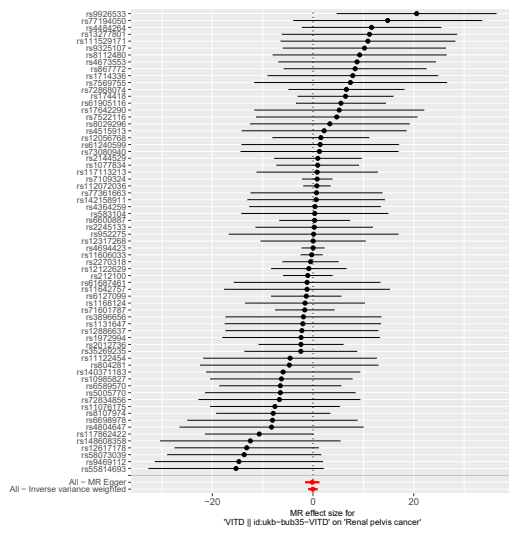
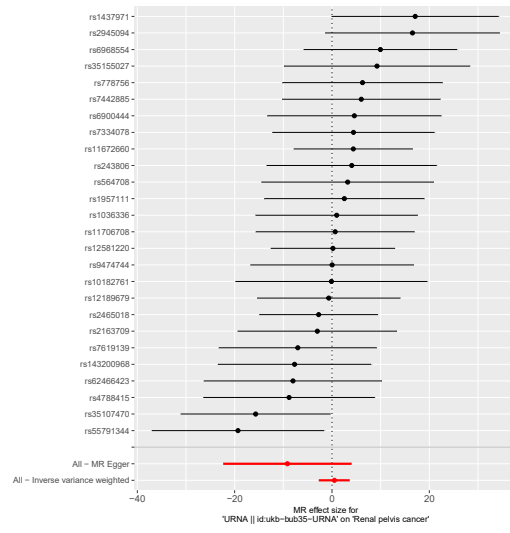
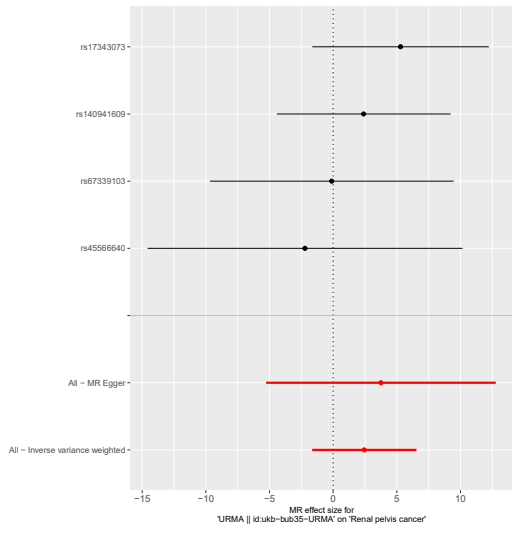




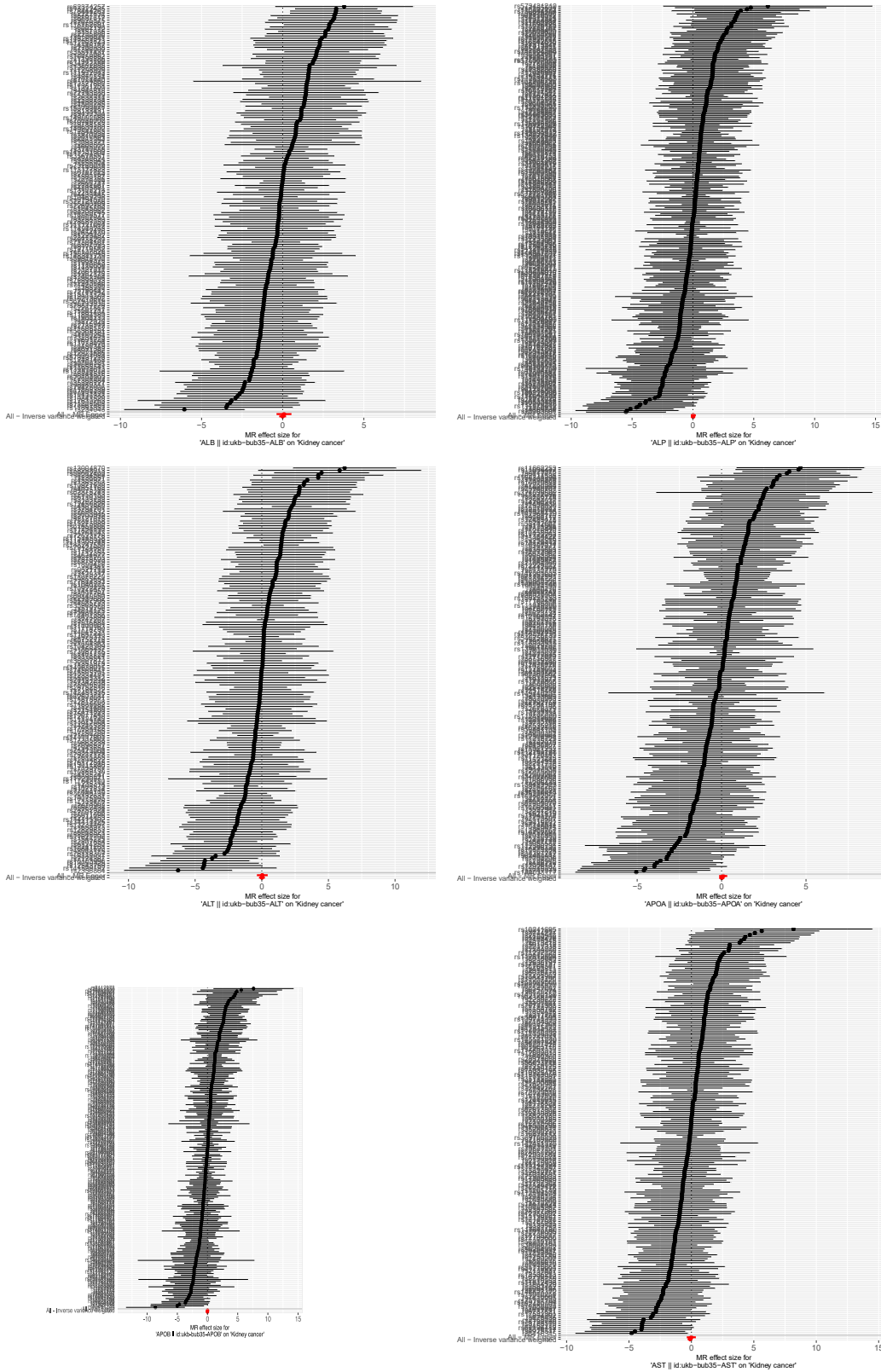


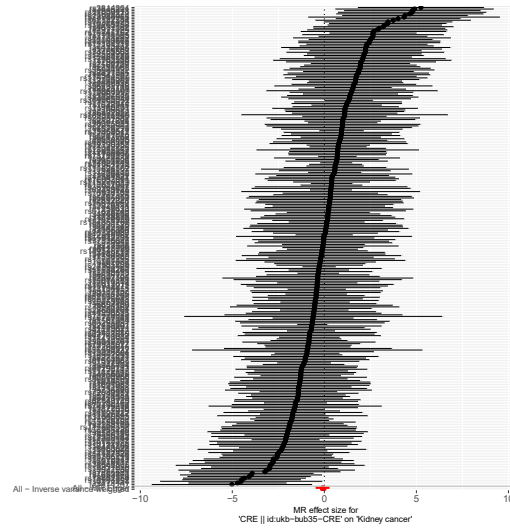
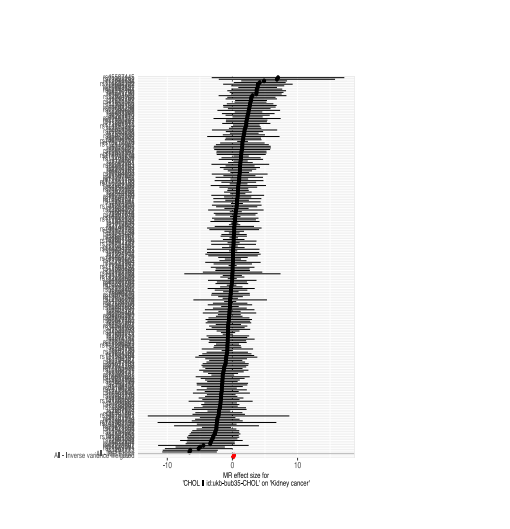
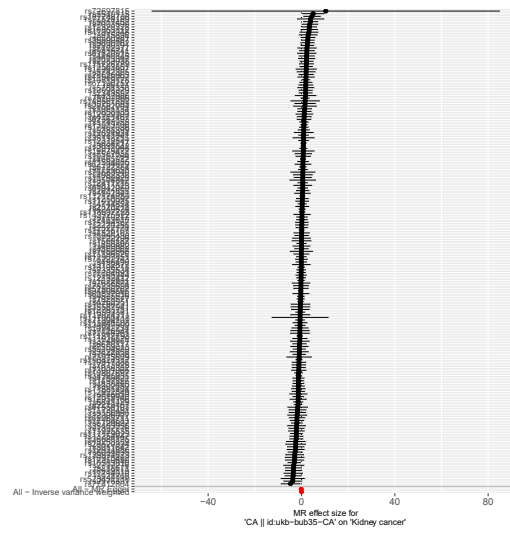
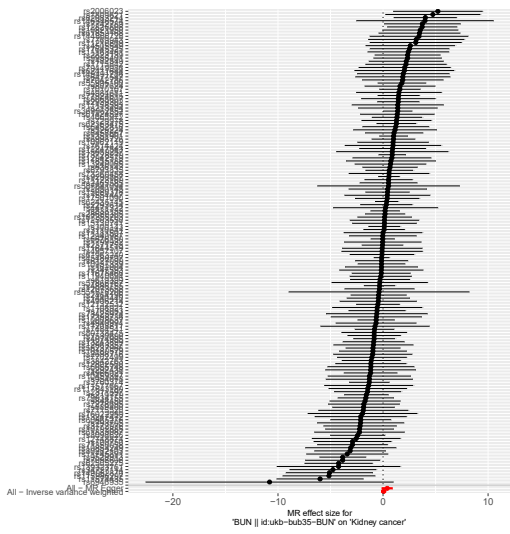
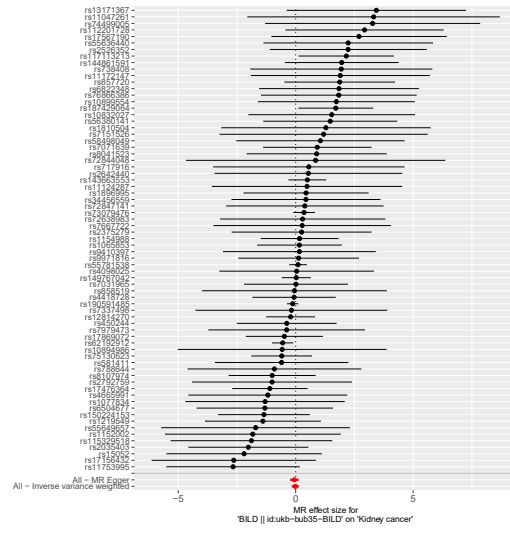
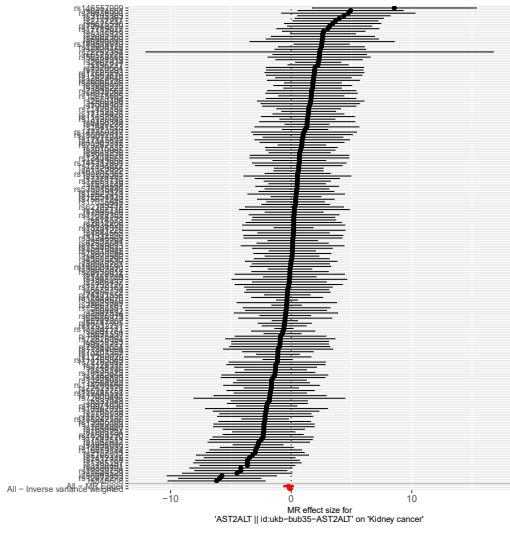


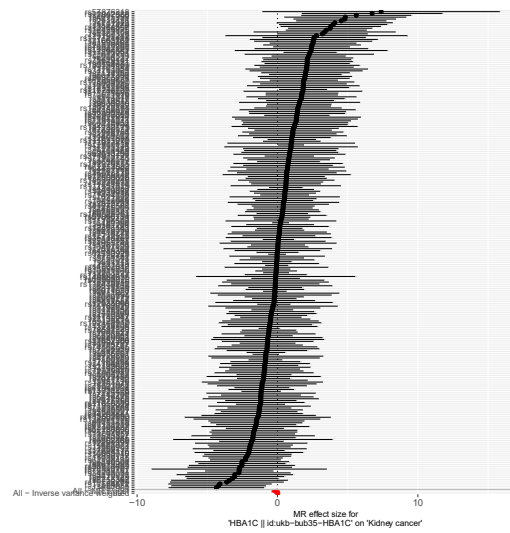
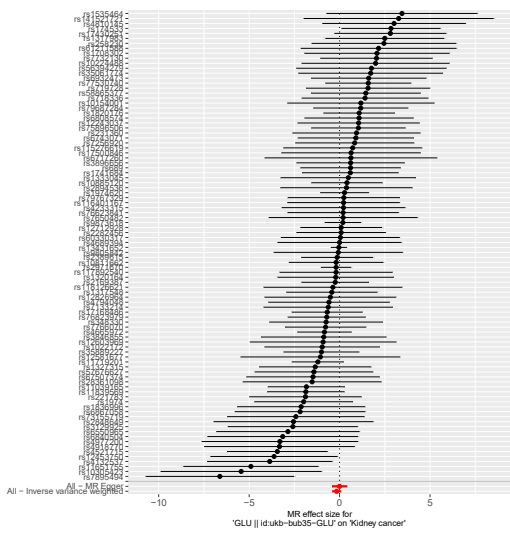
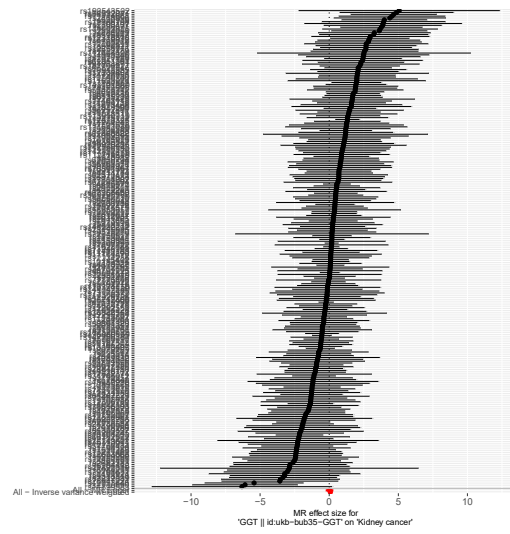
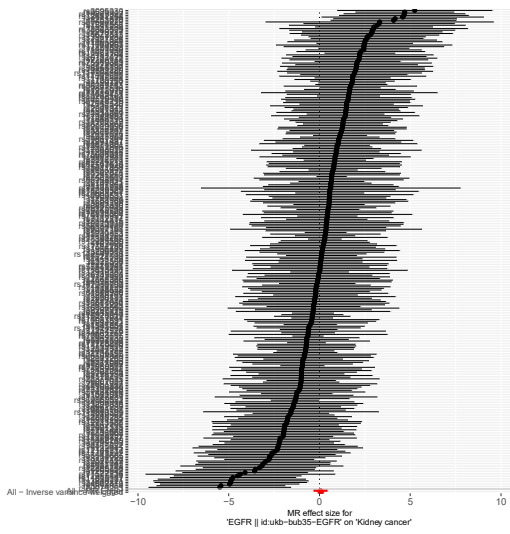
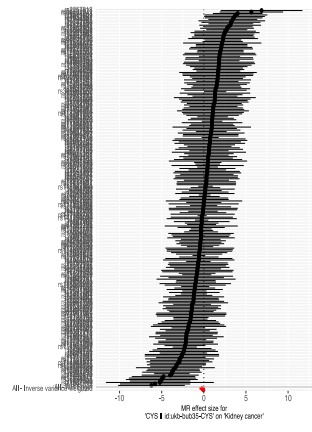
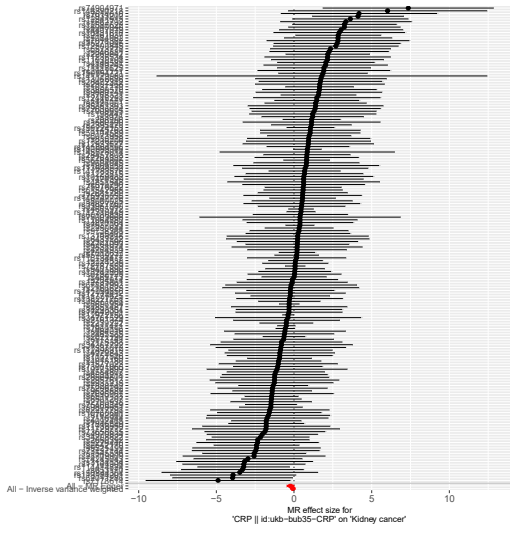


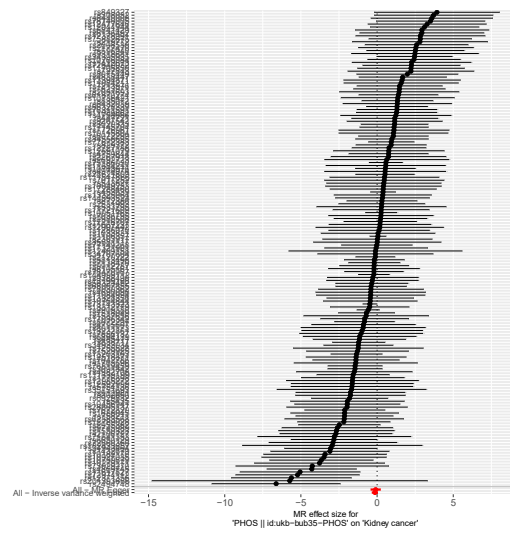
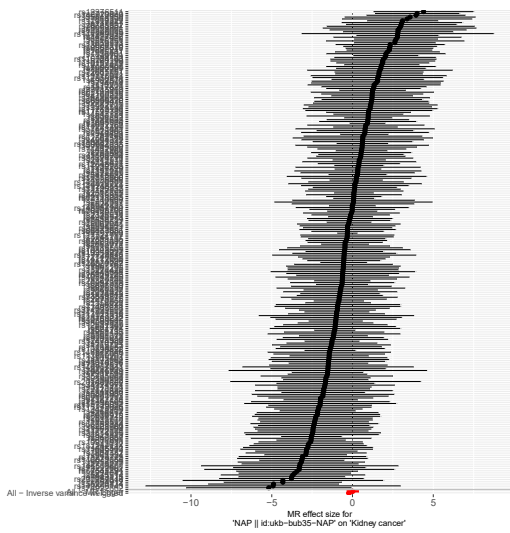
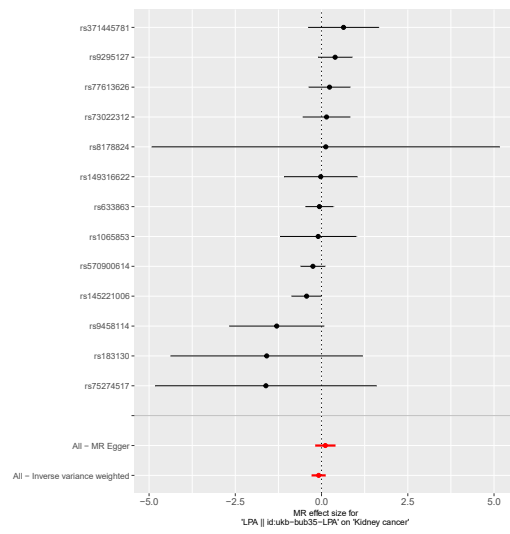
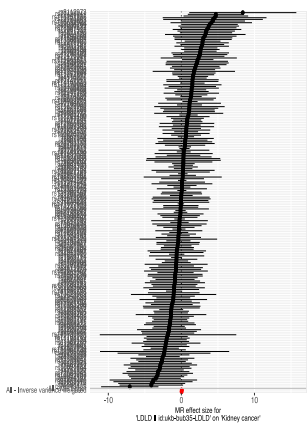
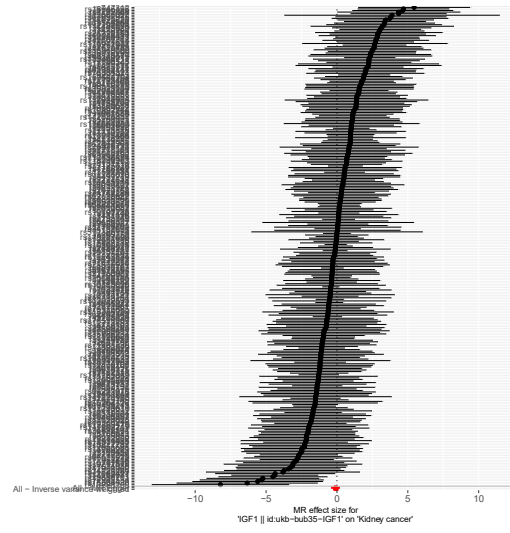
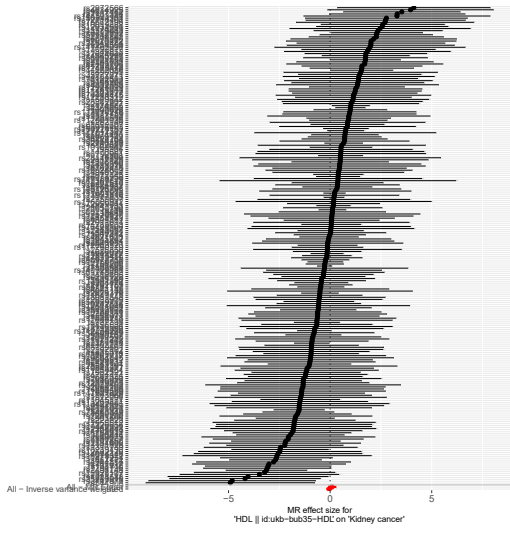


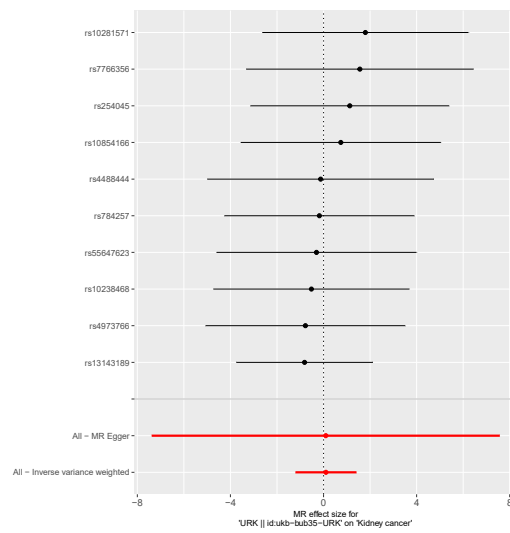
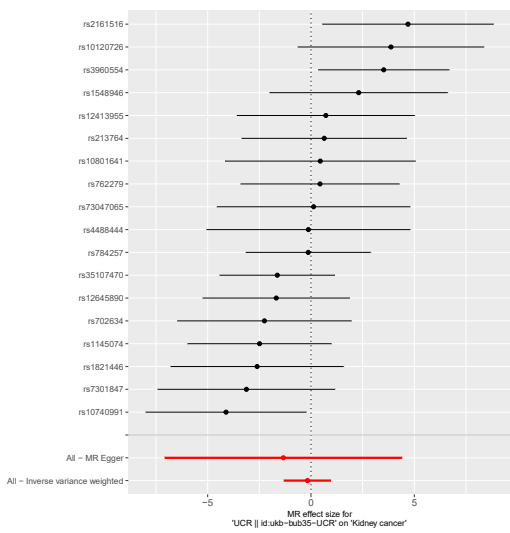
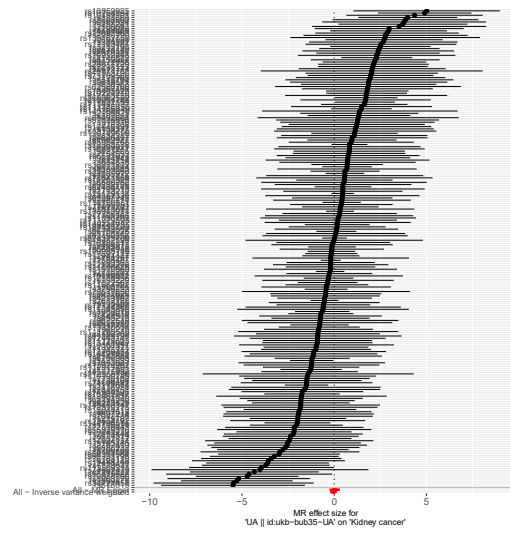
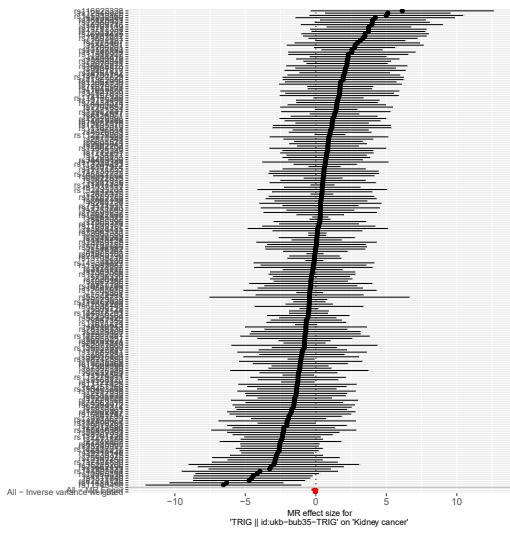
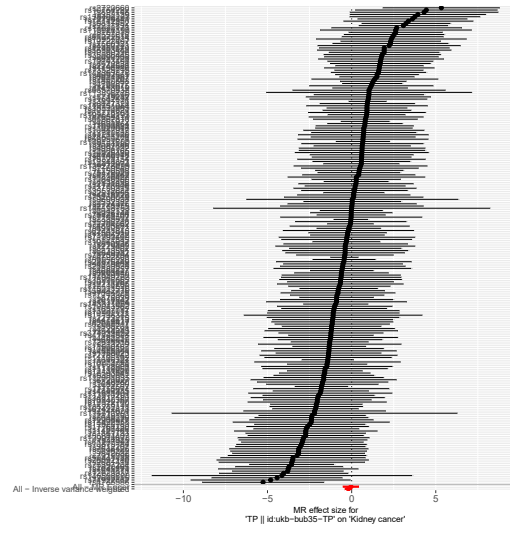
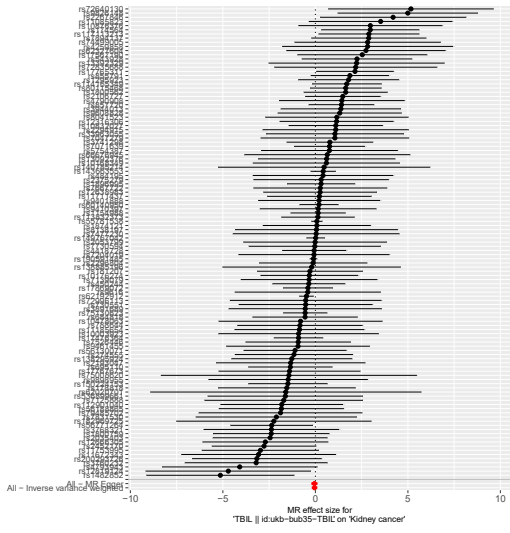
Forest plot of KCa

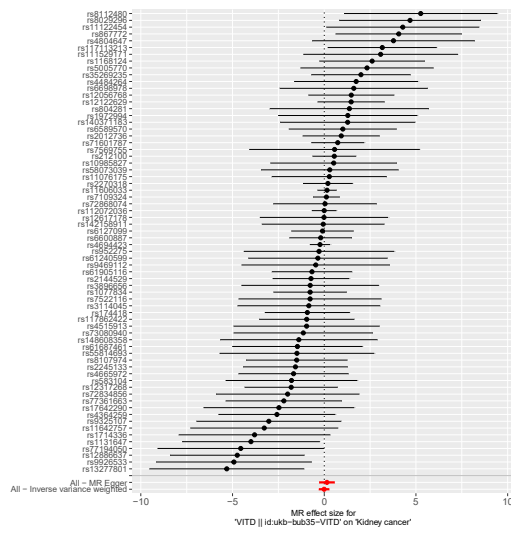
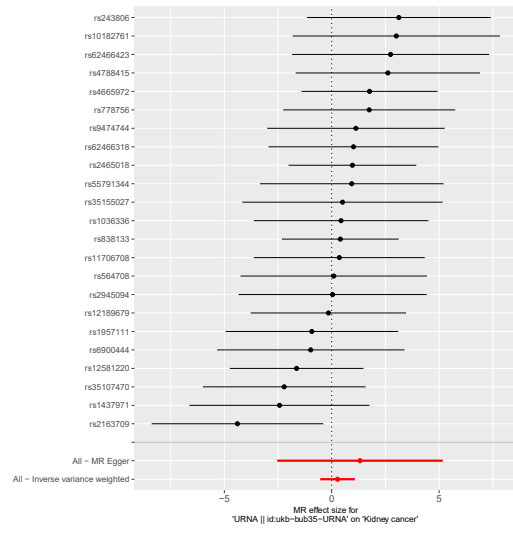
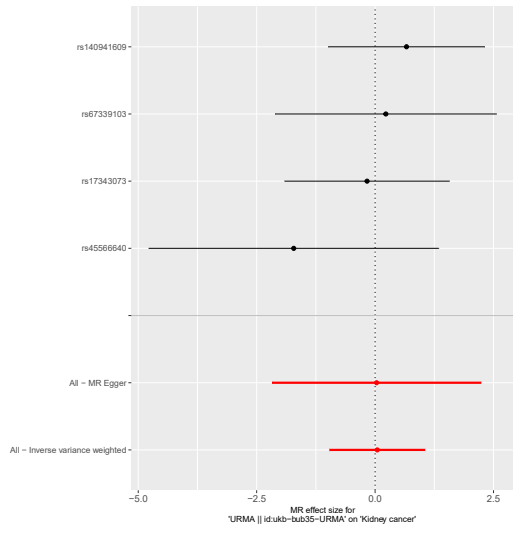




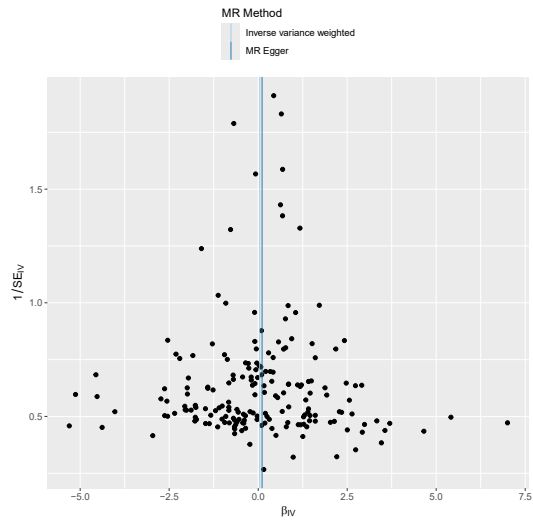




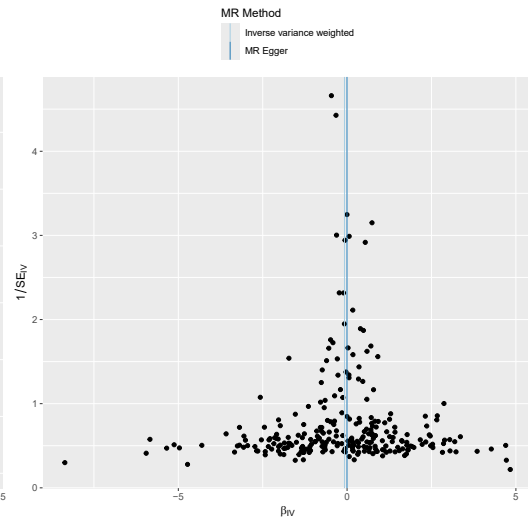




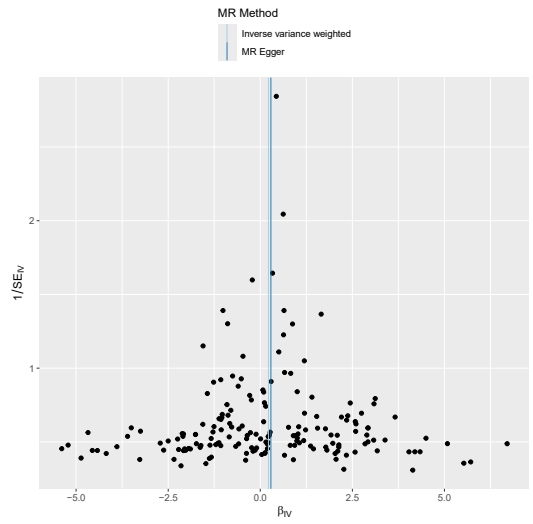
funnel plot of BCa



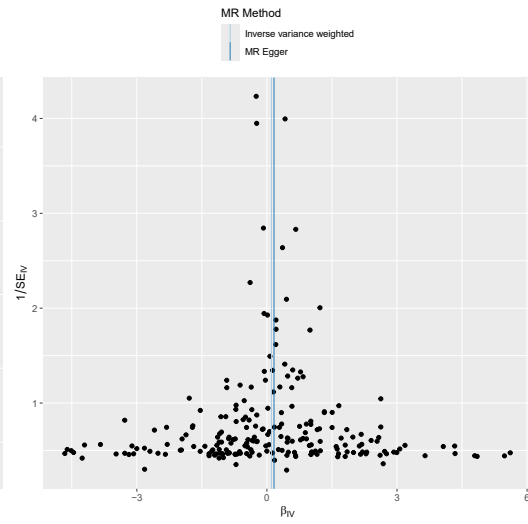
ALB



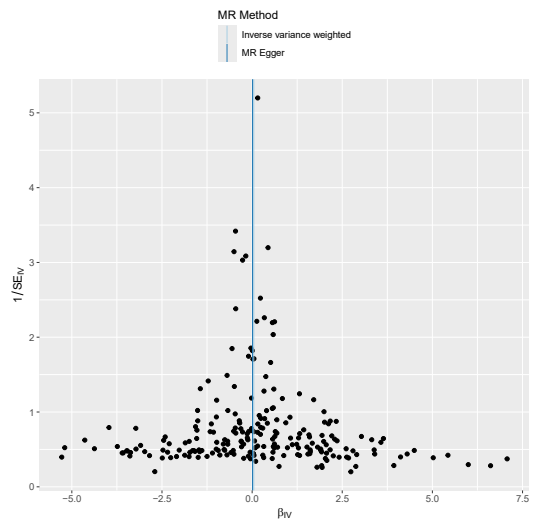
ALP



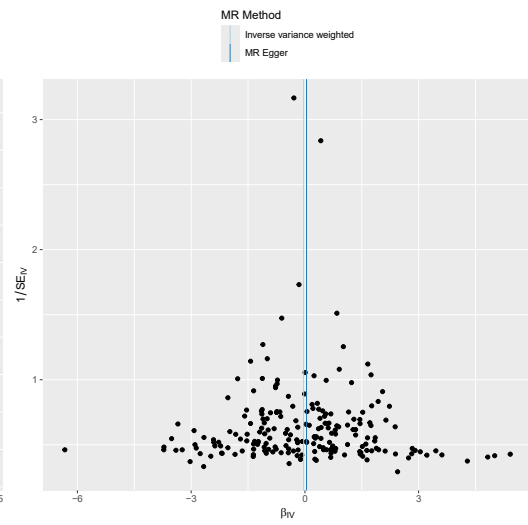
ALT



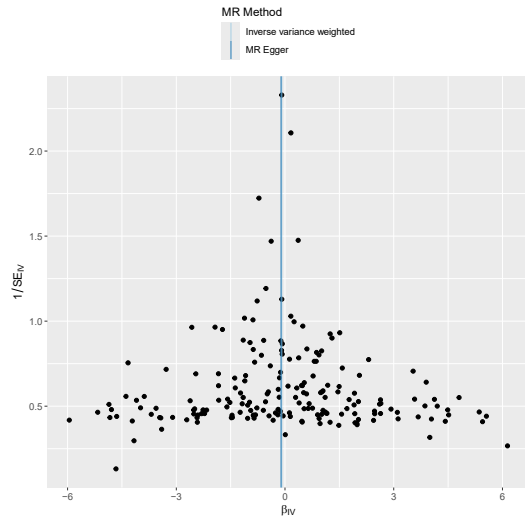
APOA



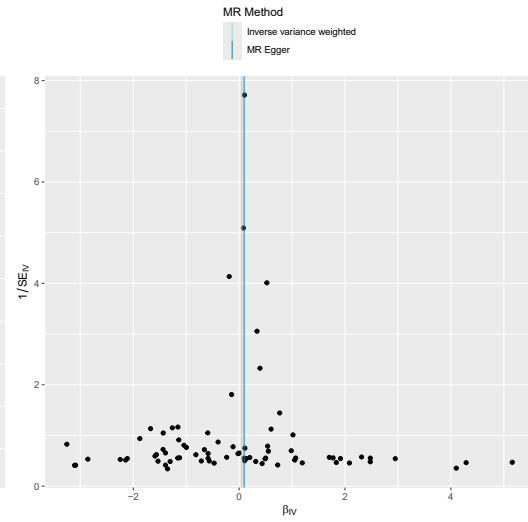
APOB



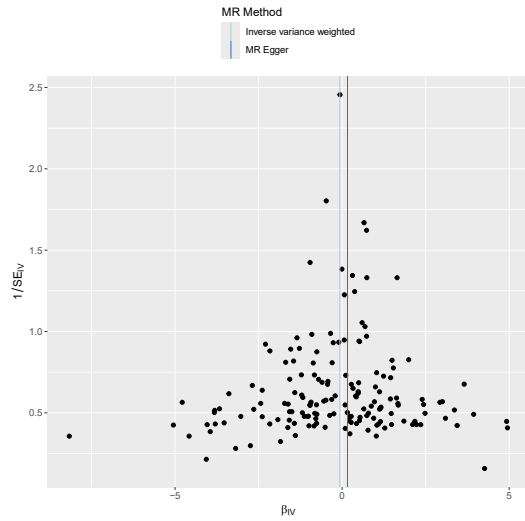
AST



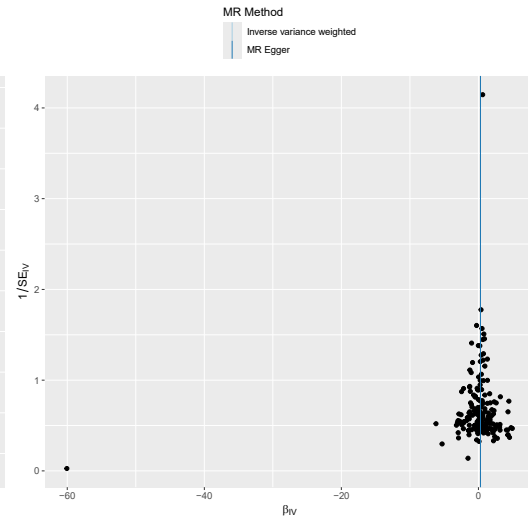
AST/ALT



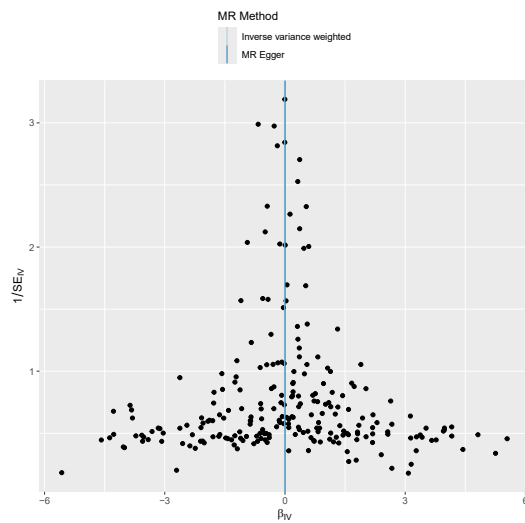
BILD



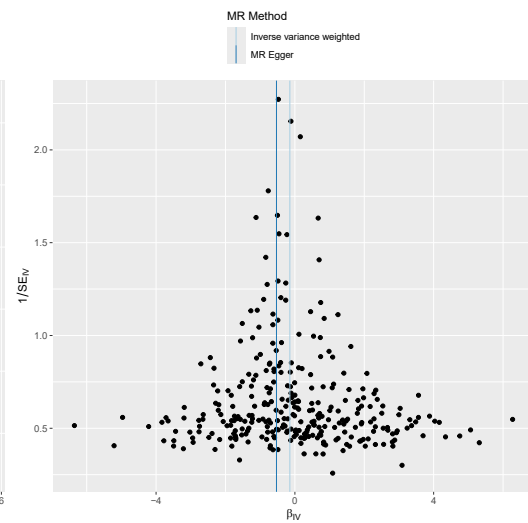
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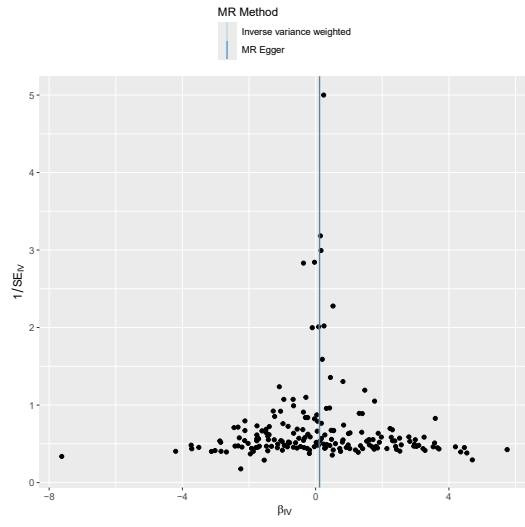
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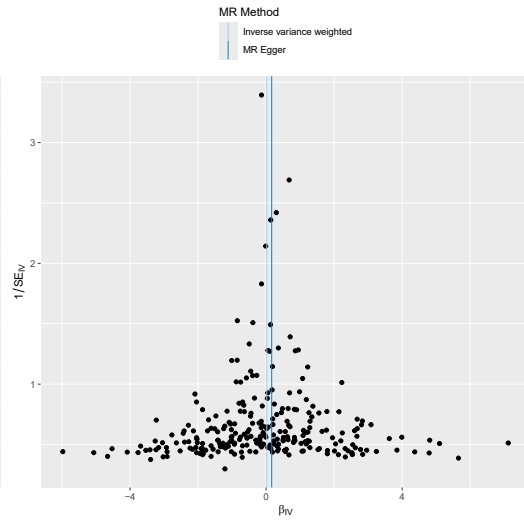
CHOL



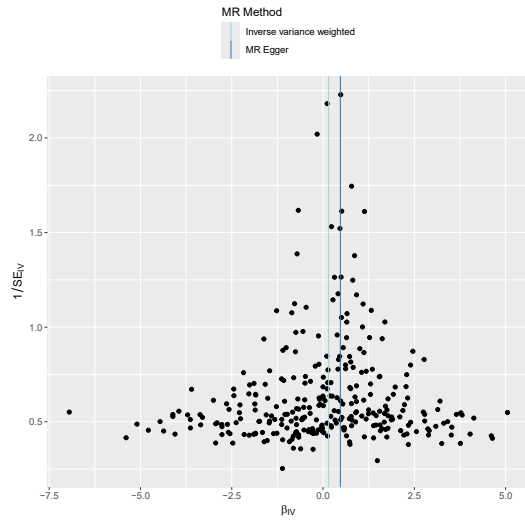
CRE



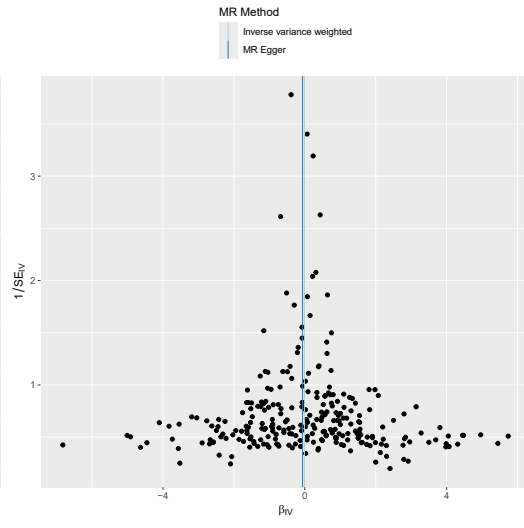
CRP



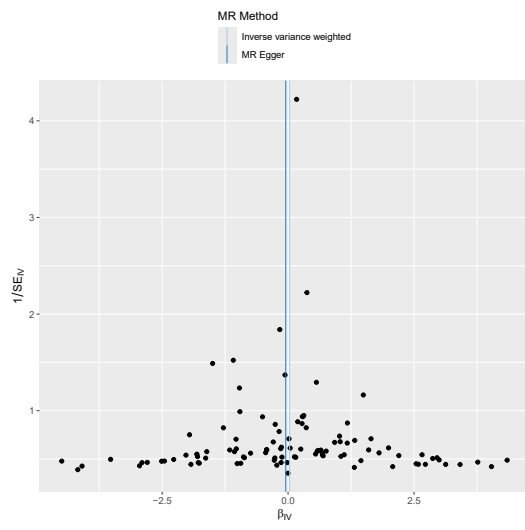
CYS



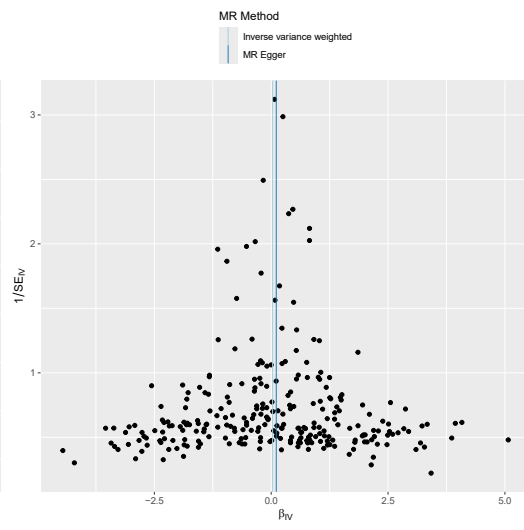
EGFR



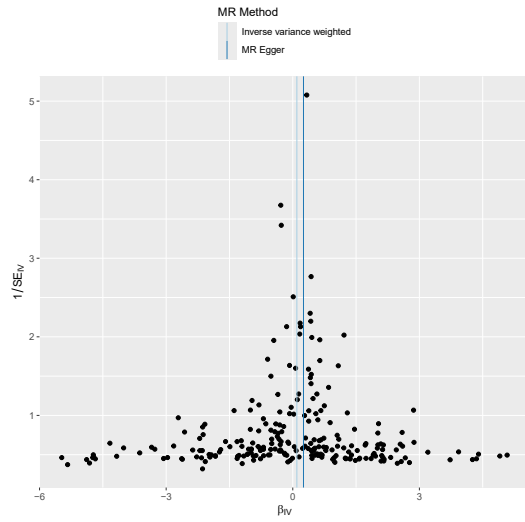
GGT



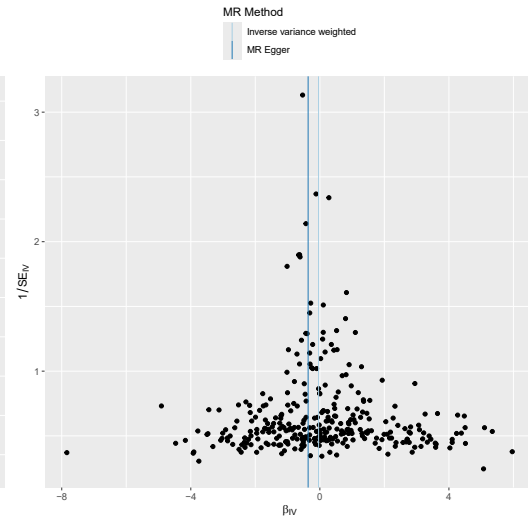
GLU



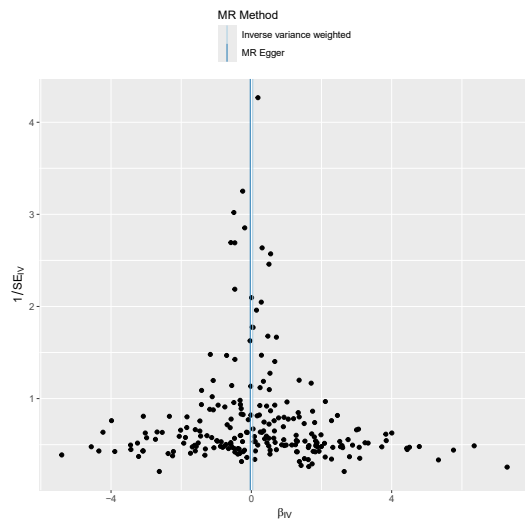
HBA1C



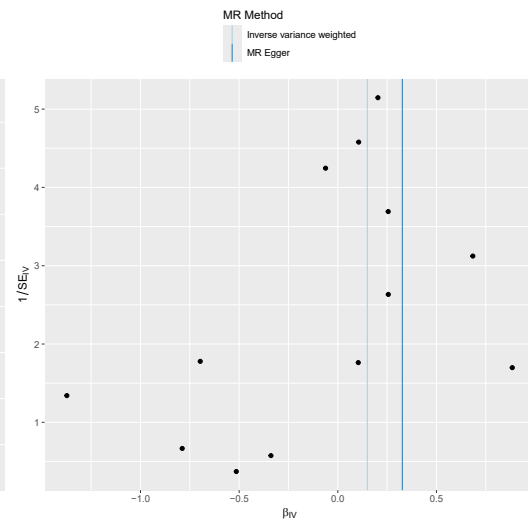
HDL



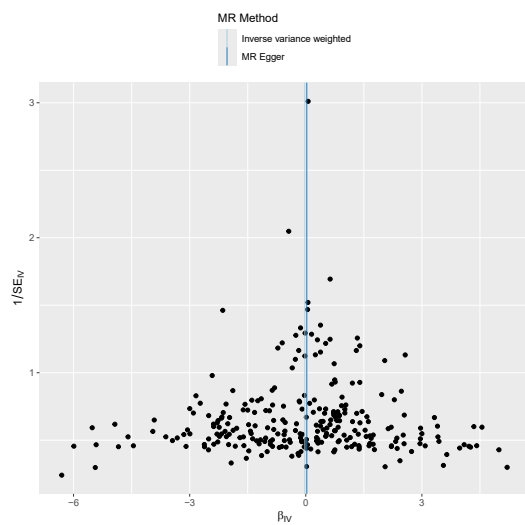
IGF-1



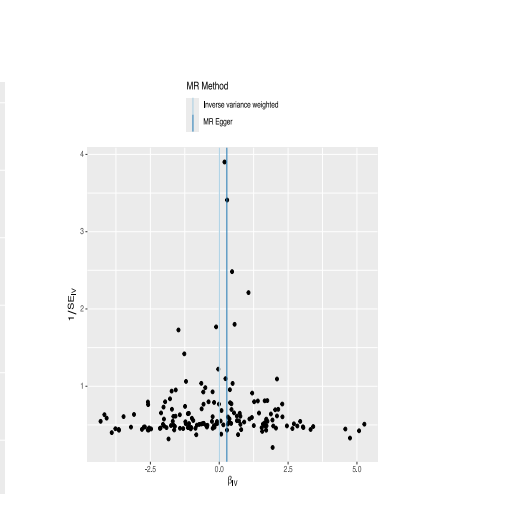
LDL



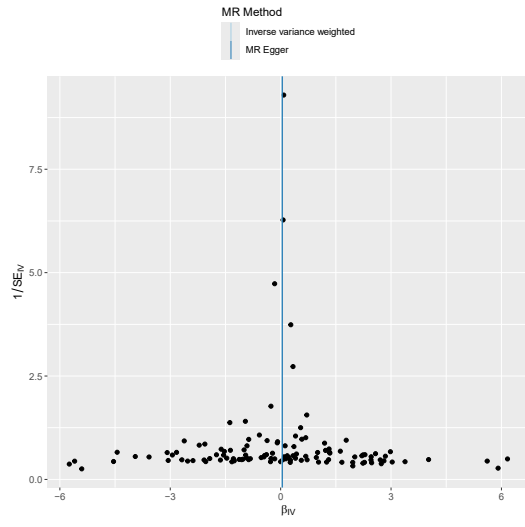
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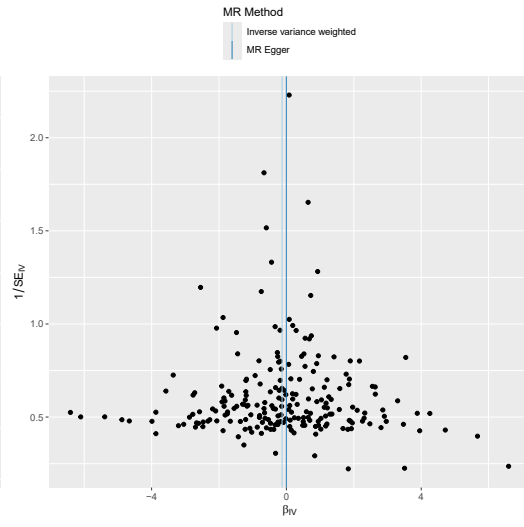
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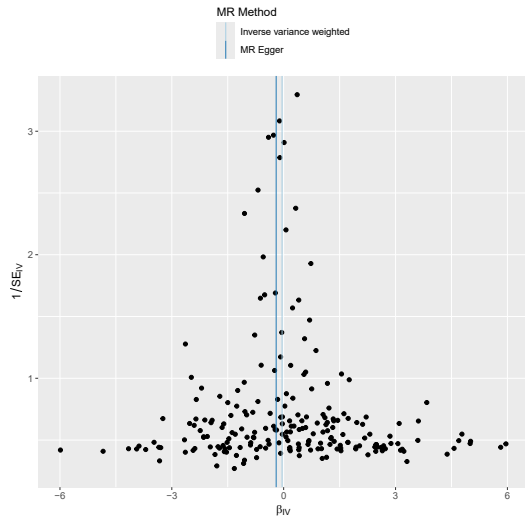
PHOS



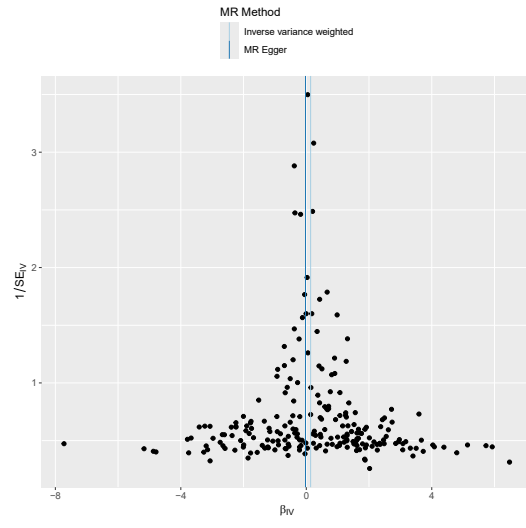
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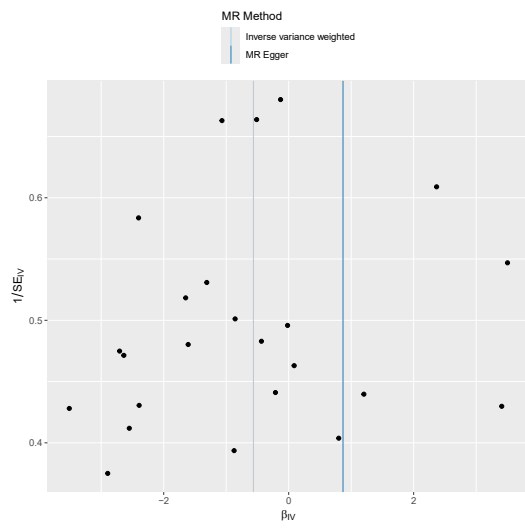
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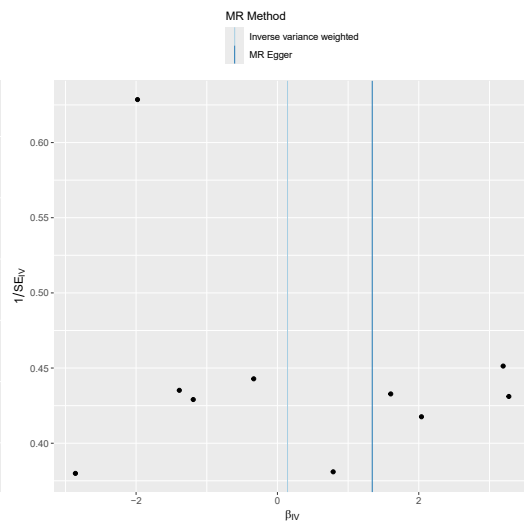
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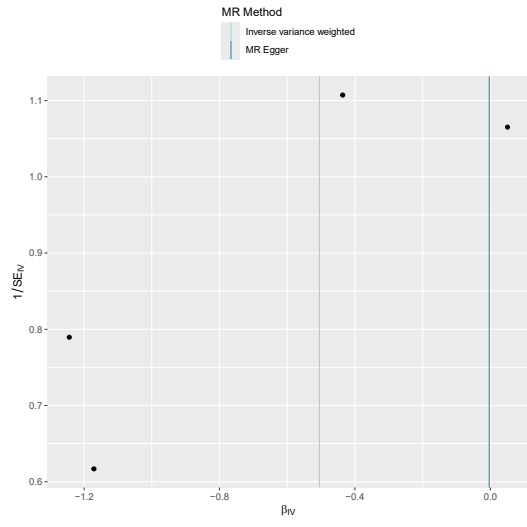
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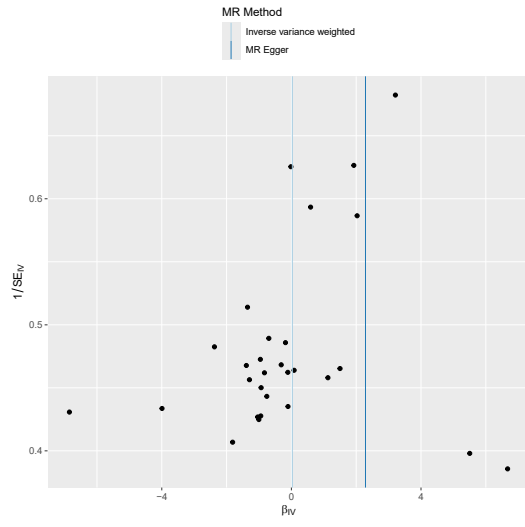
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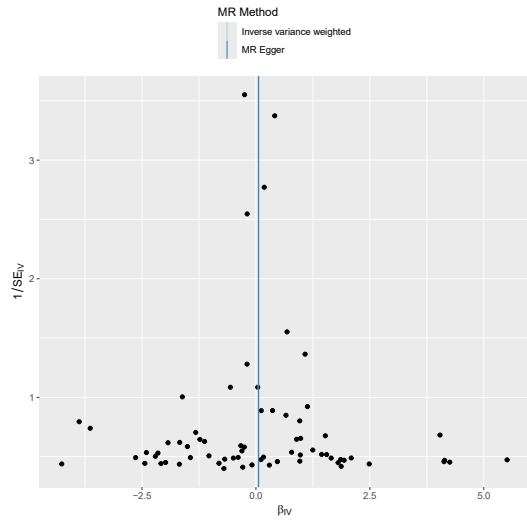
URK



URMA

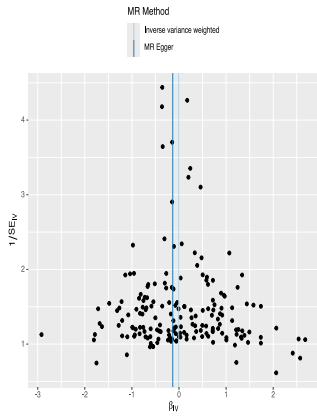


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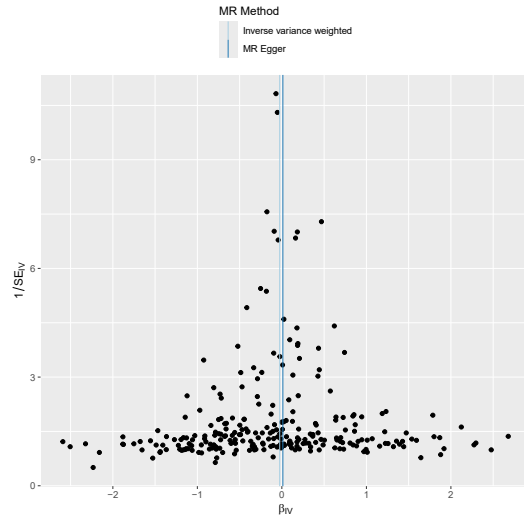


VITD

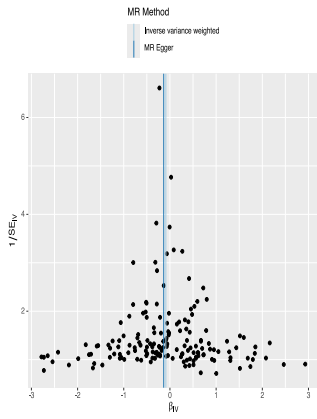
funnel plot of PCA



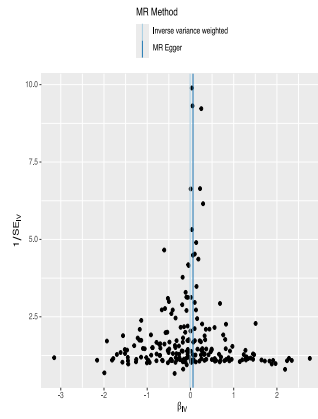
ALB



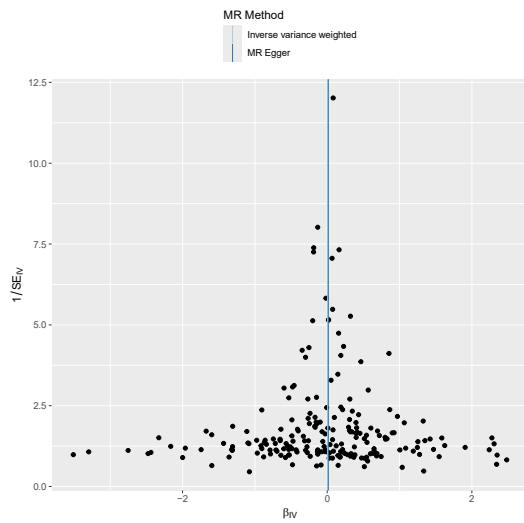
ALP



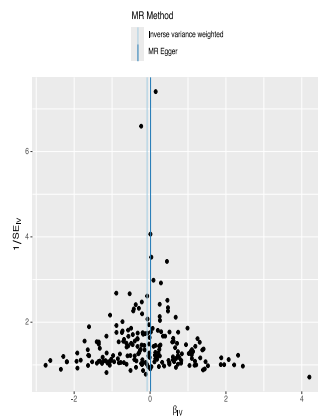
ALT



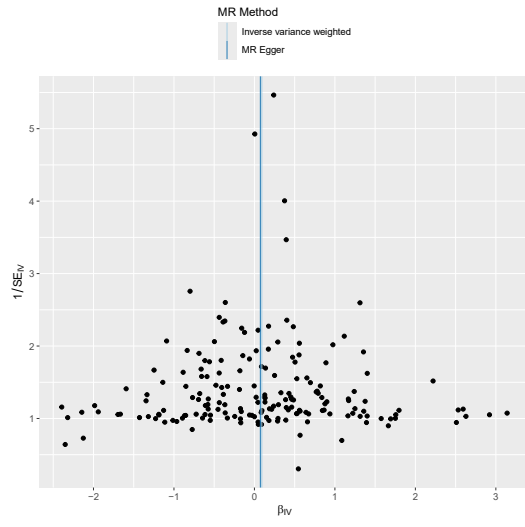
APOA



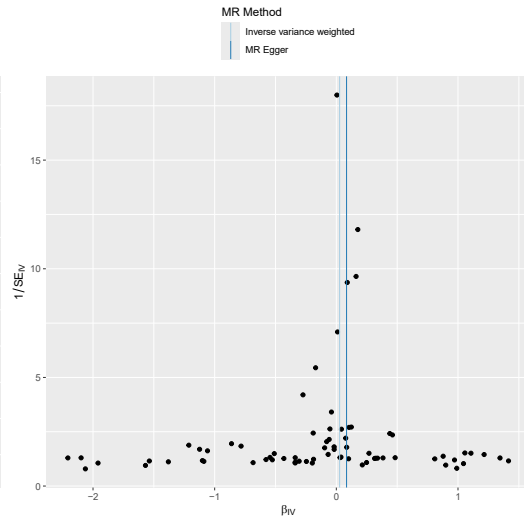
APOB



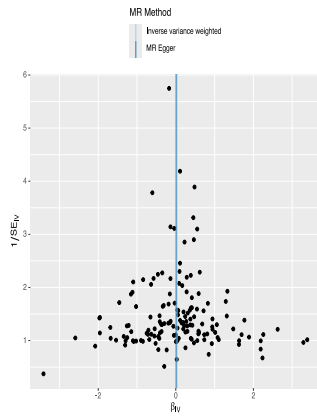
AST



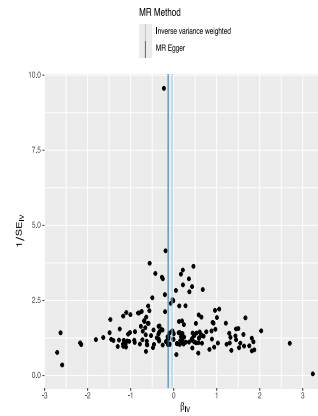
AST/ALT



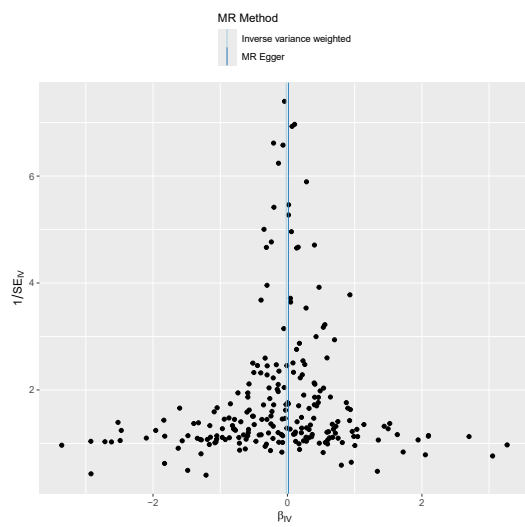
BILD



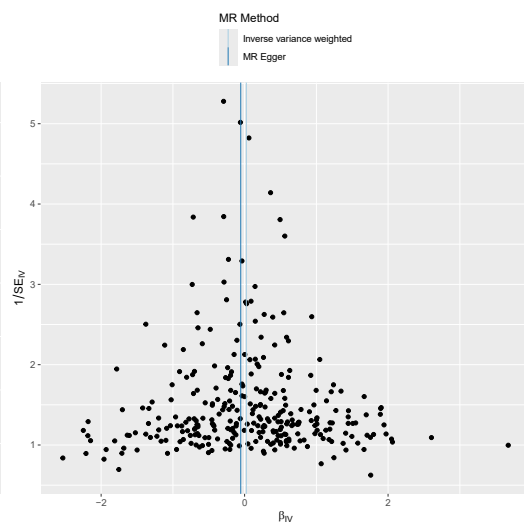
BUN



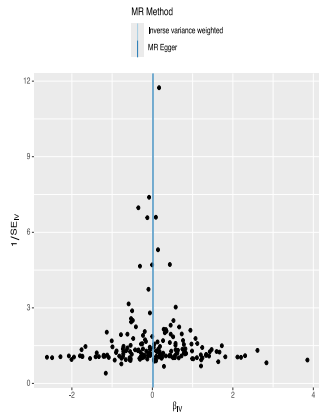
CA



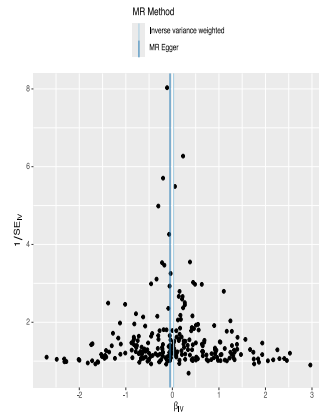
CHOL



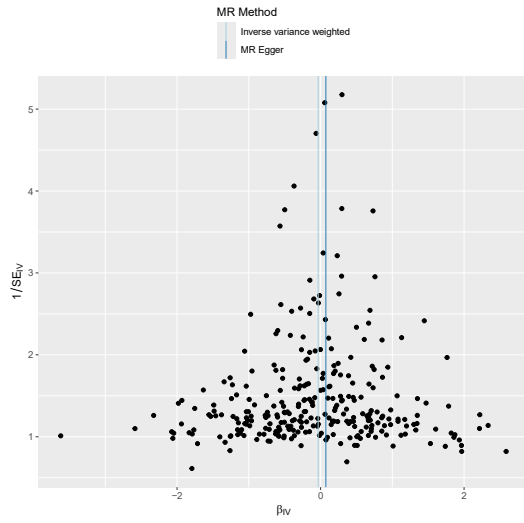
CRE



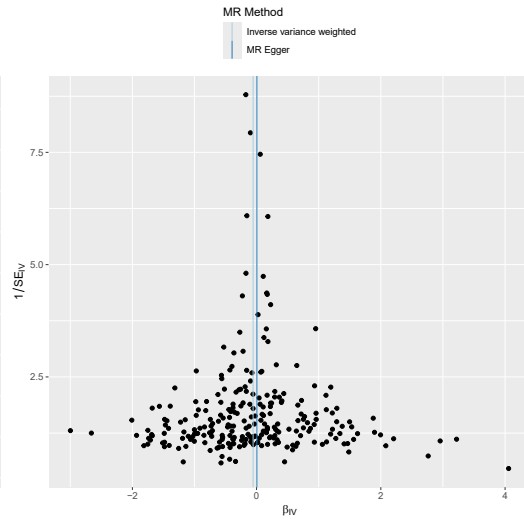
CRP



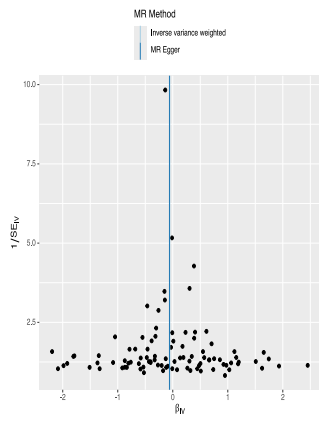
CYS



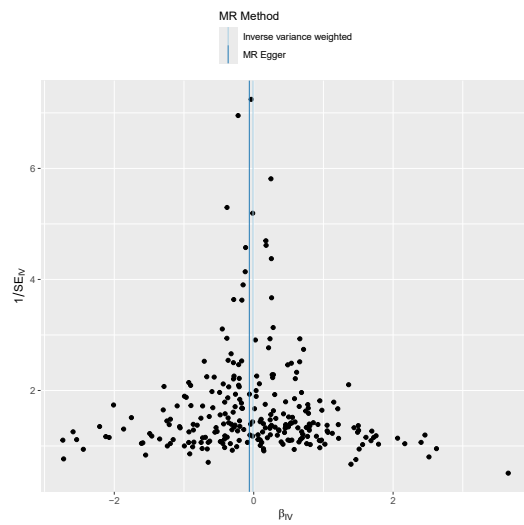
EGFR



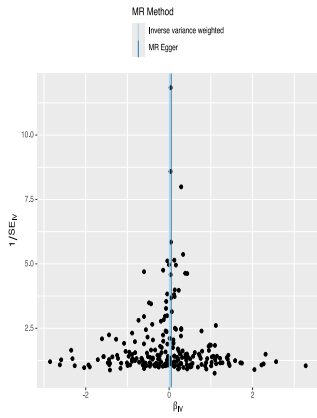
GGT



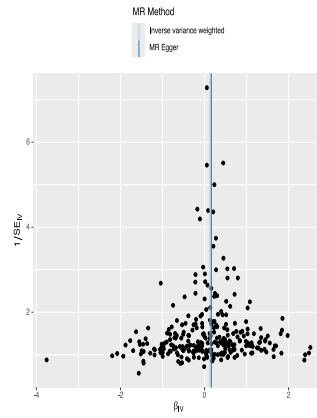
GLU



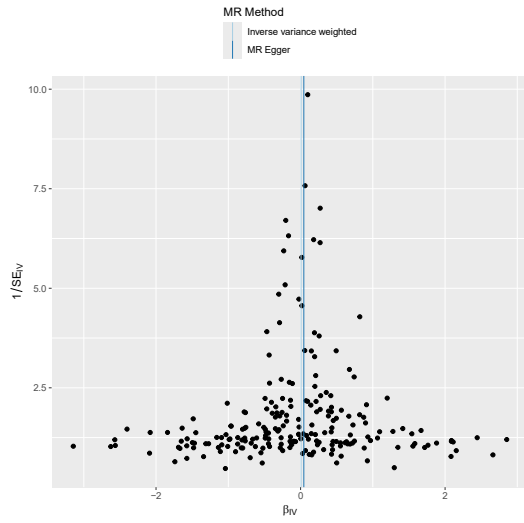
HBA1C



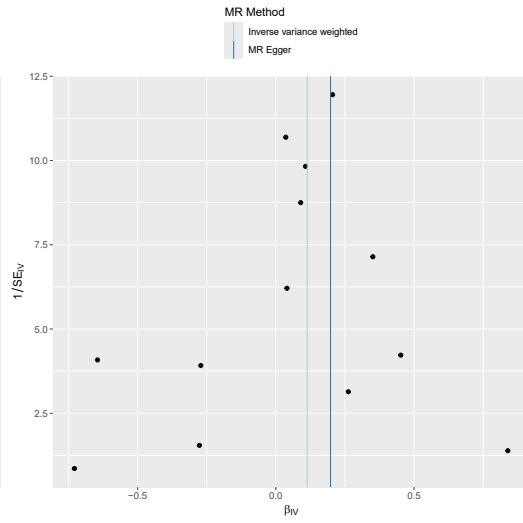
HDL



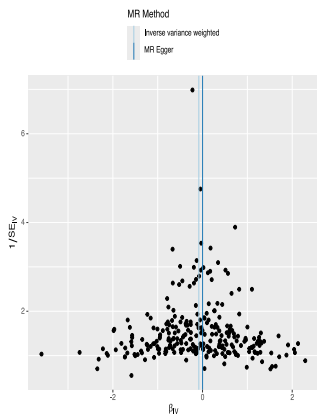
IGF-1



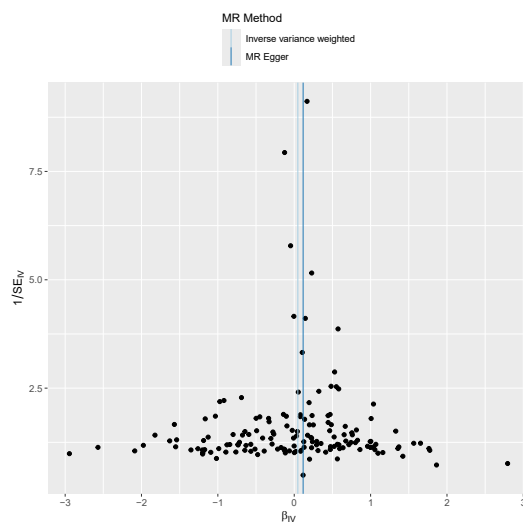
LDLD



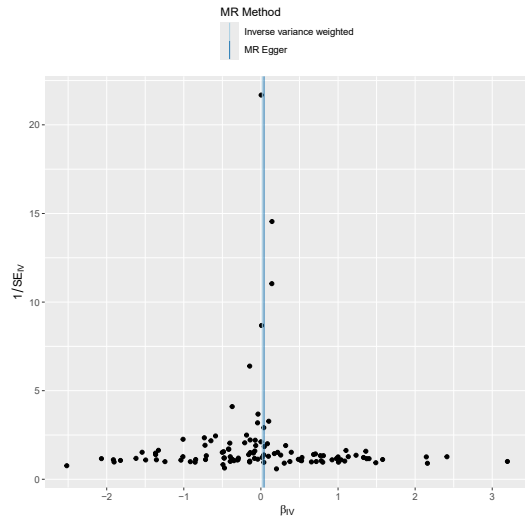
LPA



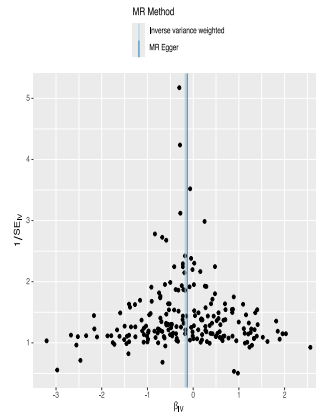
NAP



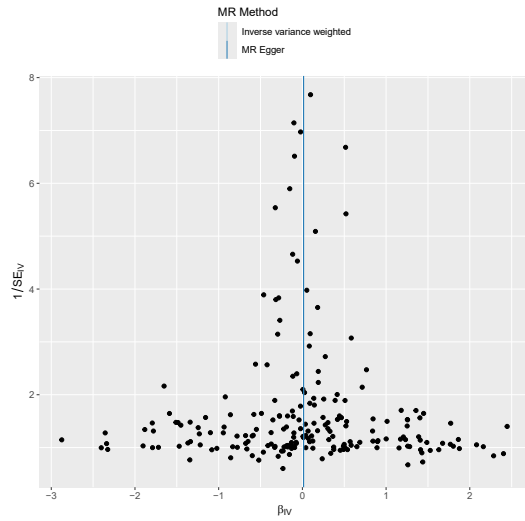
PHOS



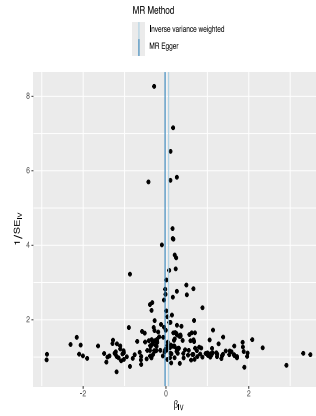
TBIL



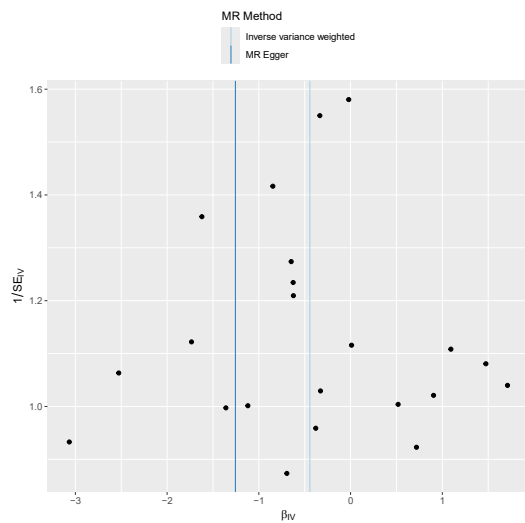
TP



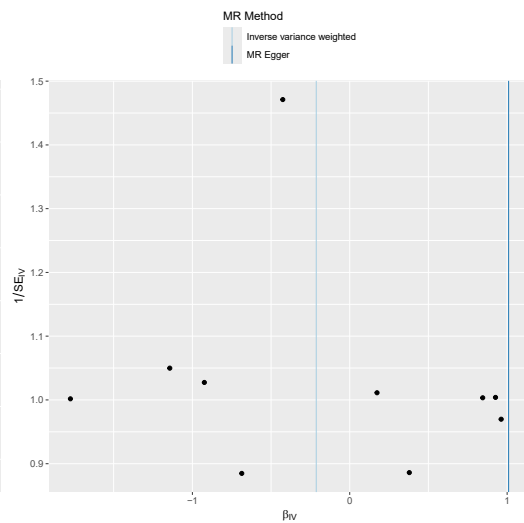
TRIG



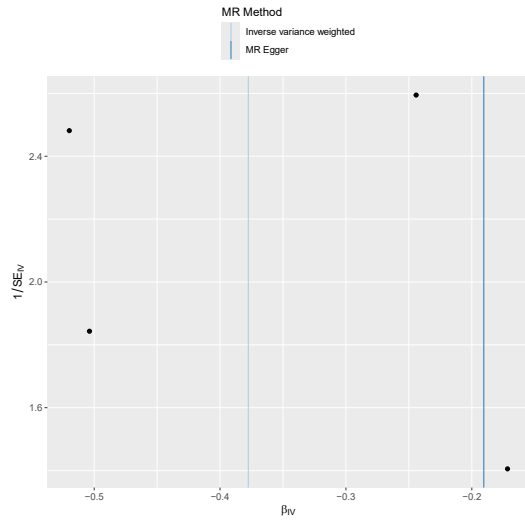
UA



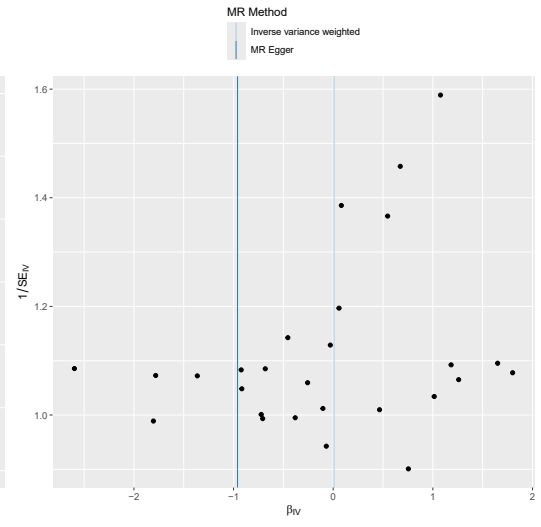
UCR



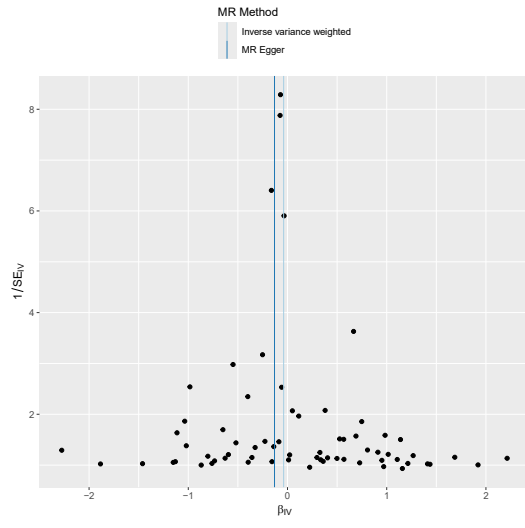
URK



URMA

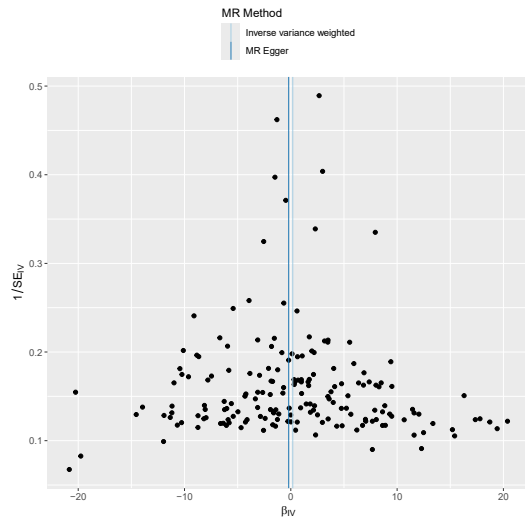


URNA

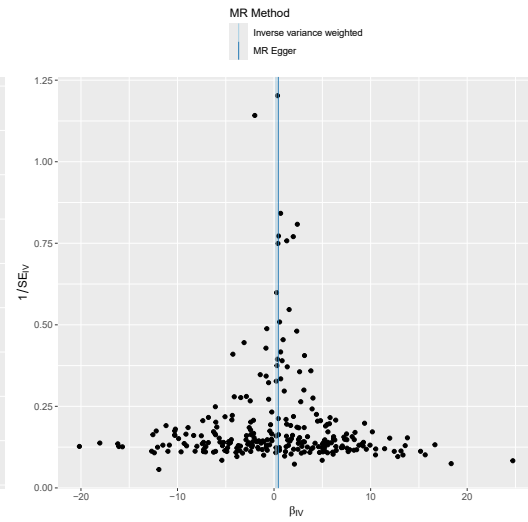


VITD

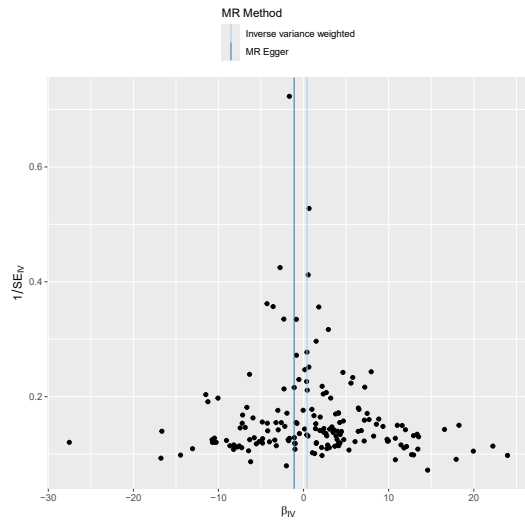
funnel plot of RPCa



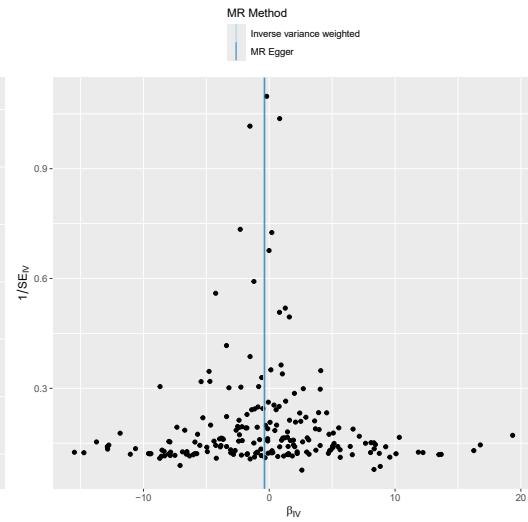
ALB



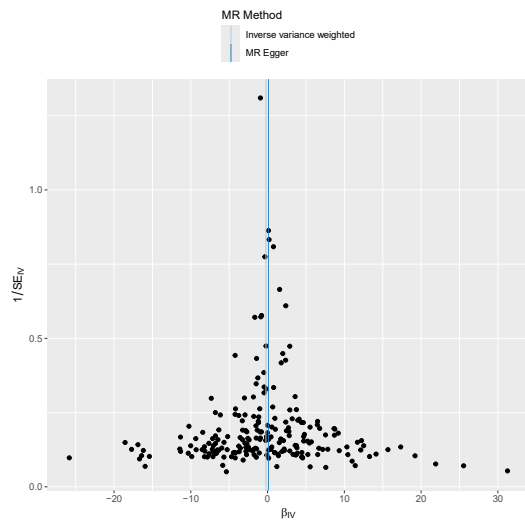
ALP



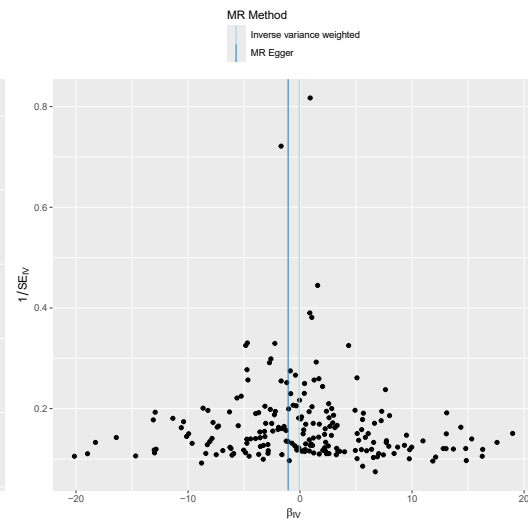
ALT



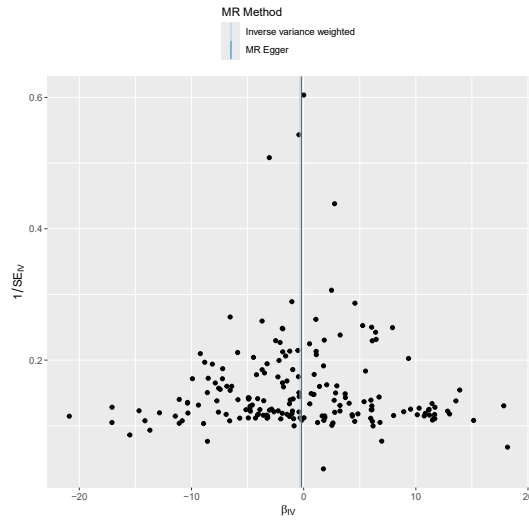
APOA



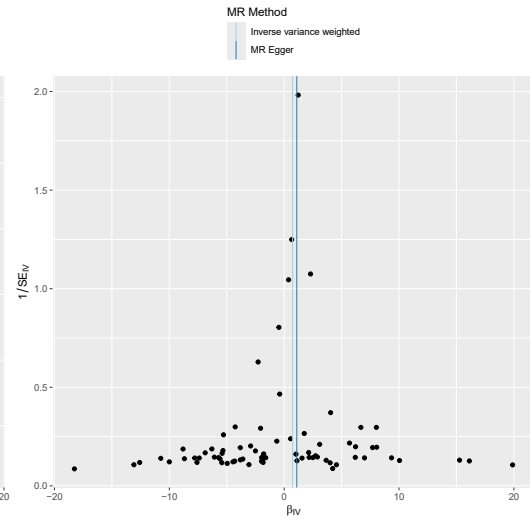
APOB



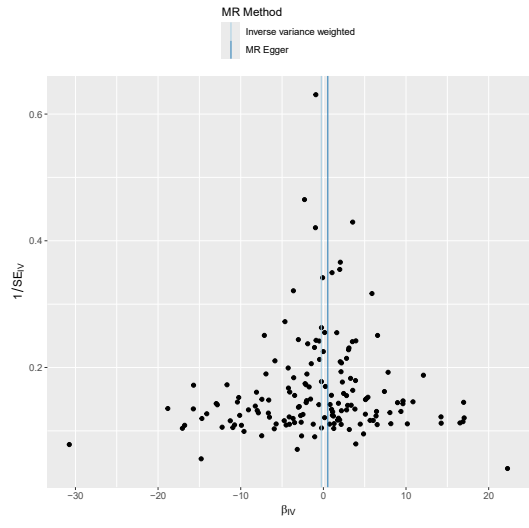
AST



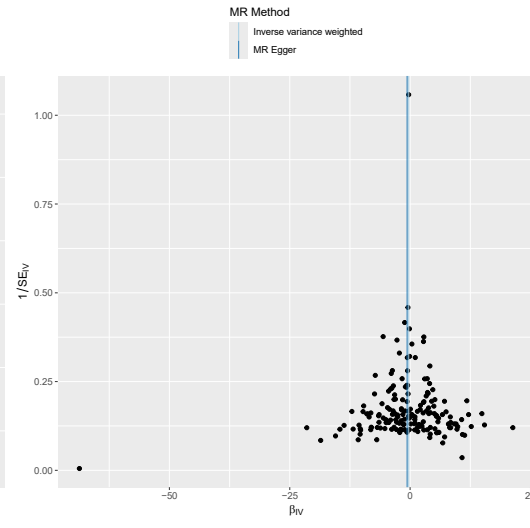
AST/ALT



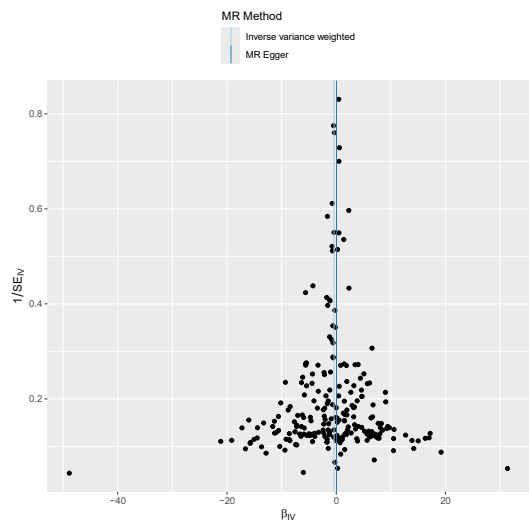
BILD



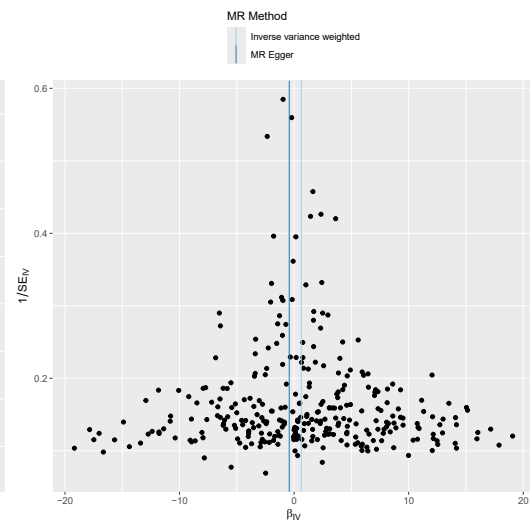
BUN



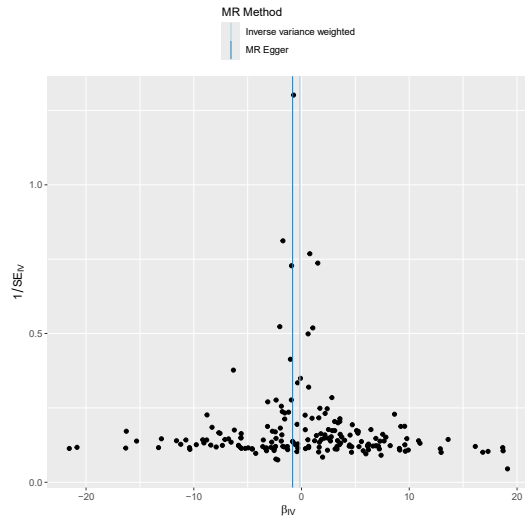
CA



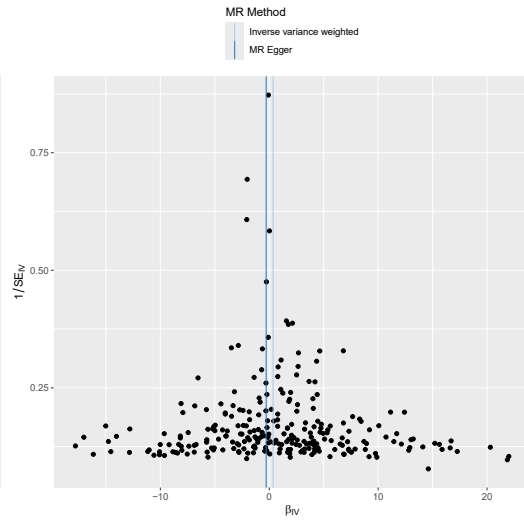
CHOL



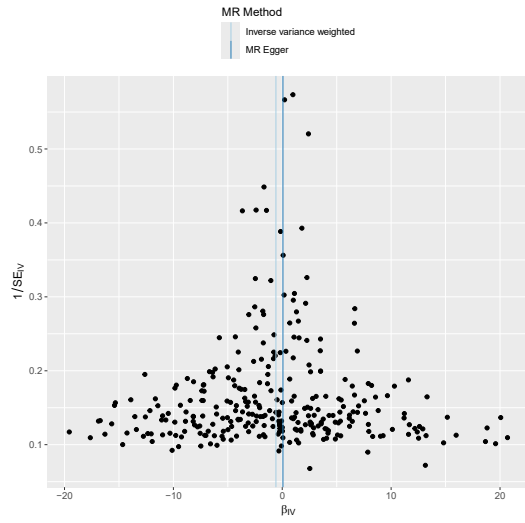
CRE



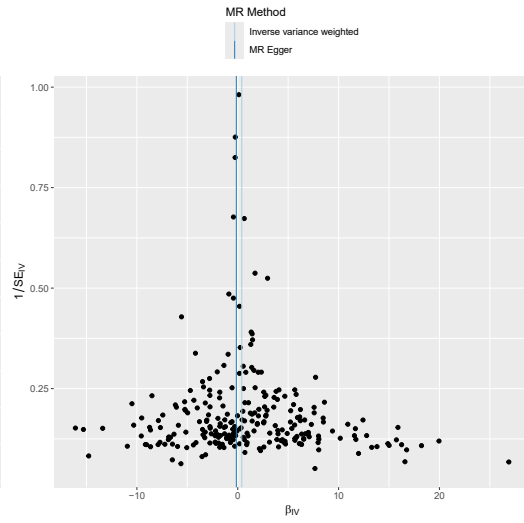
CRP



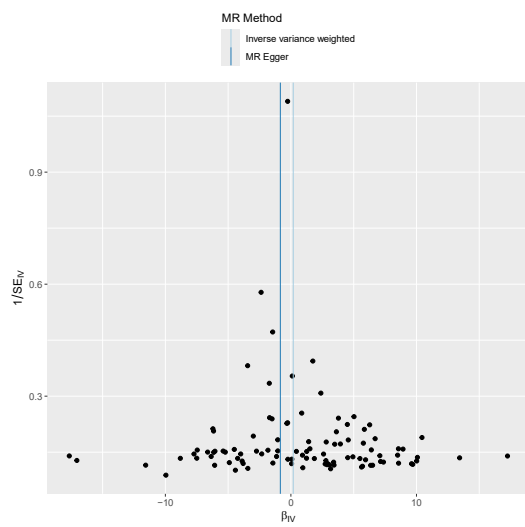
CYS



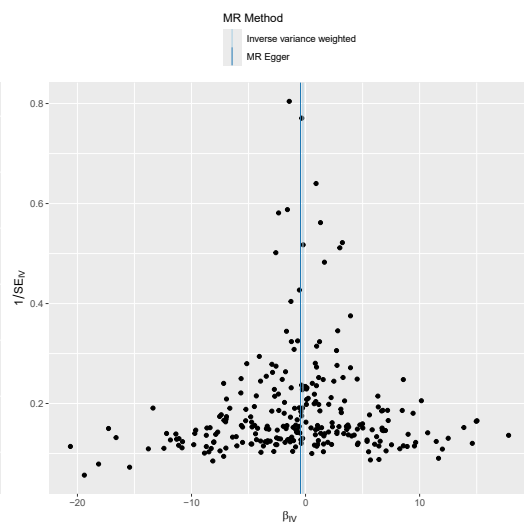
EGFR



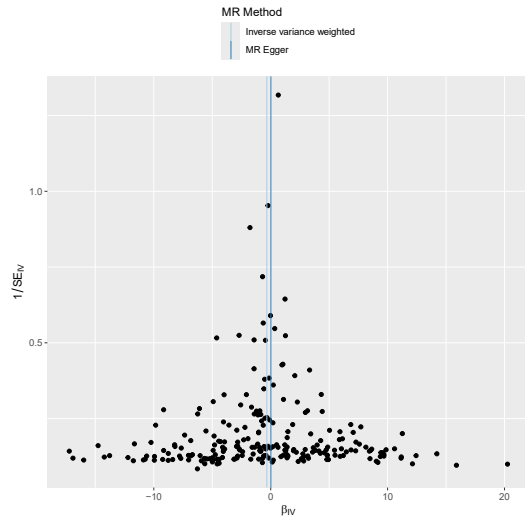
GGT



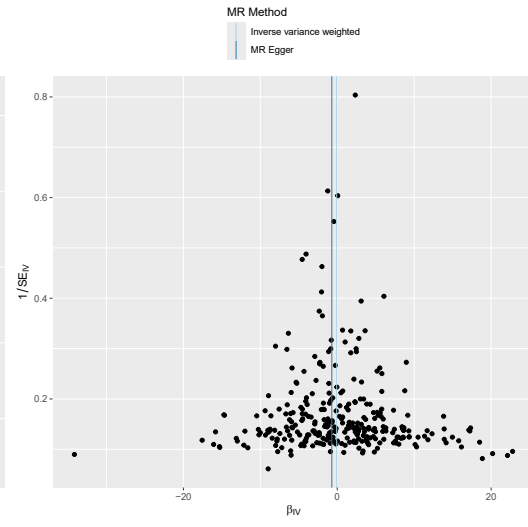
GLU



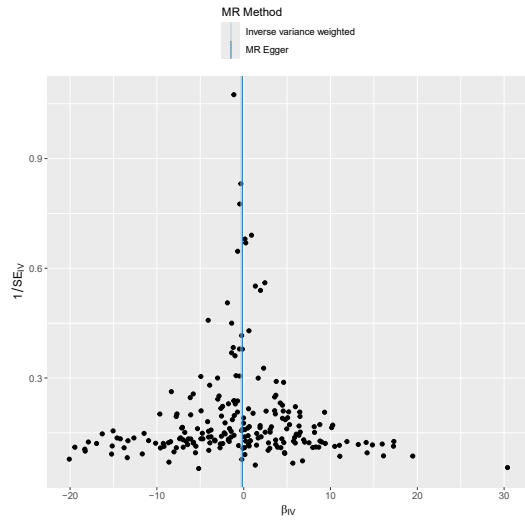
HBA1C



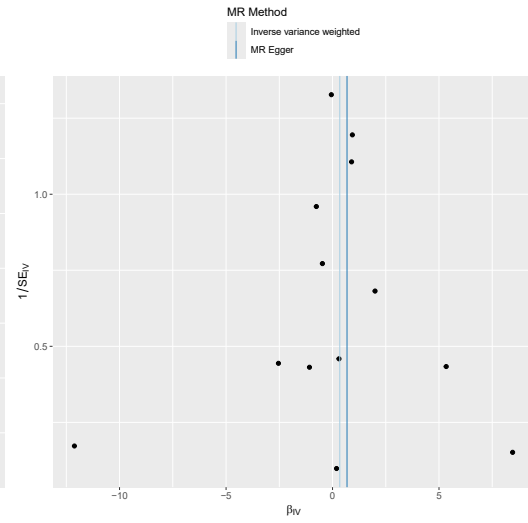
HDL



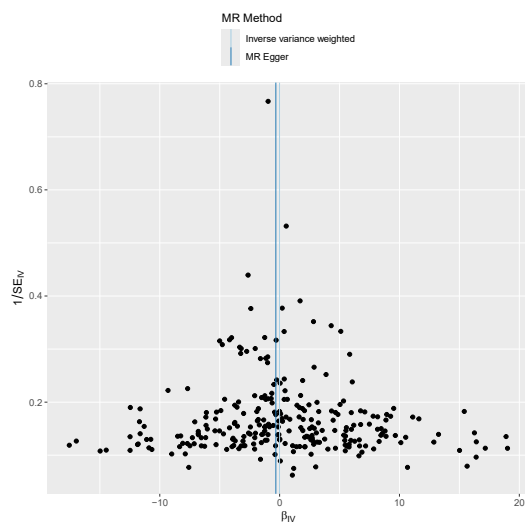
IGF-1



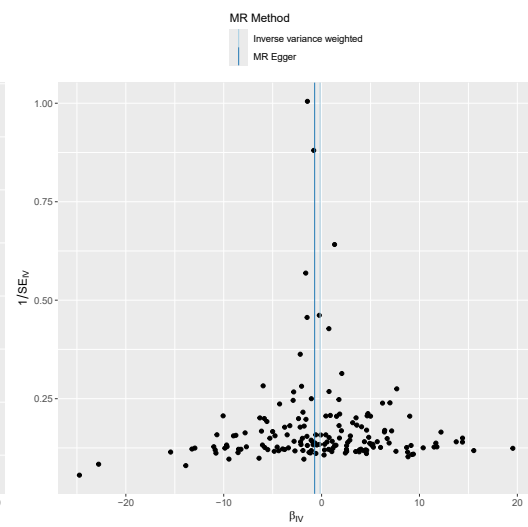
LDLD



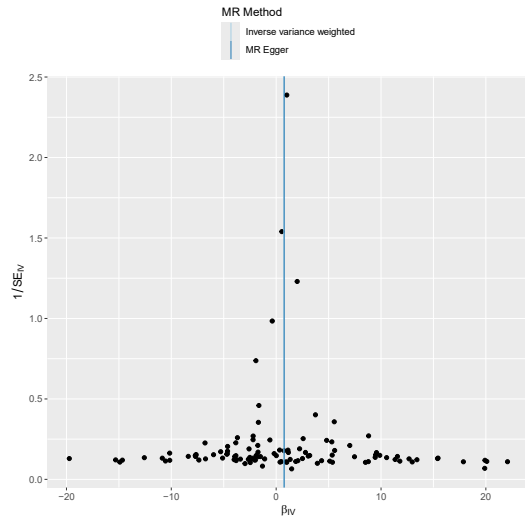
LPA



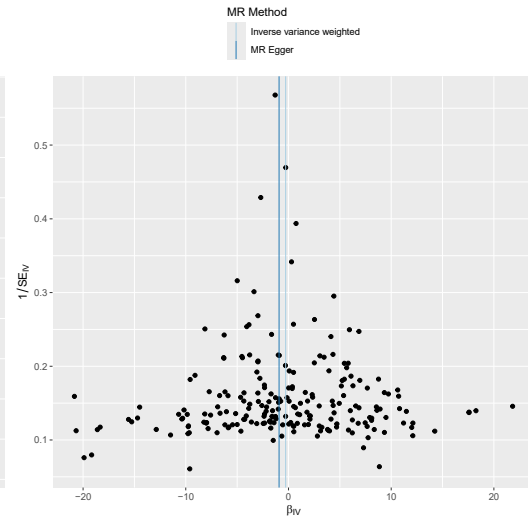
NAP



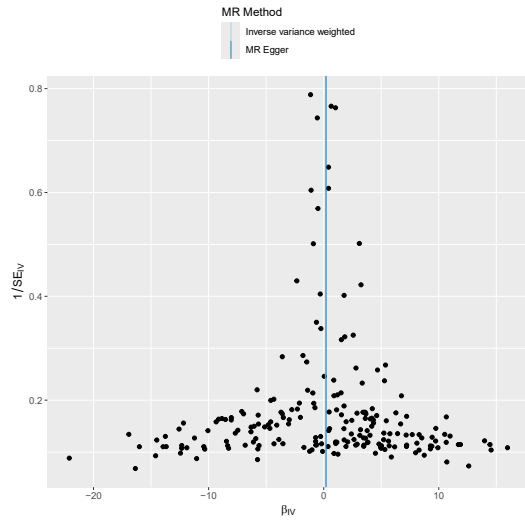
PHOS



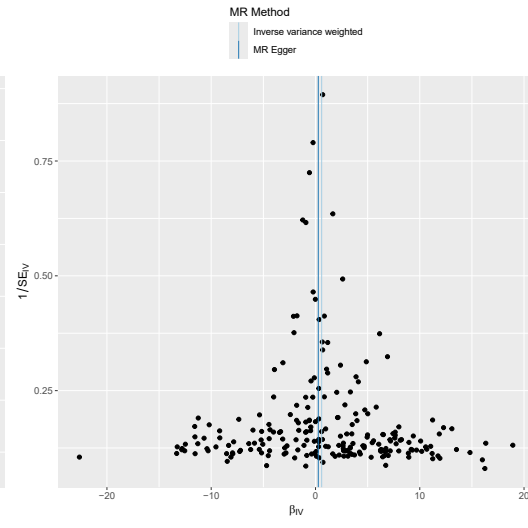
TBIL



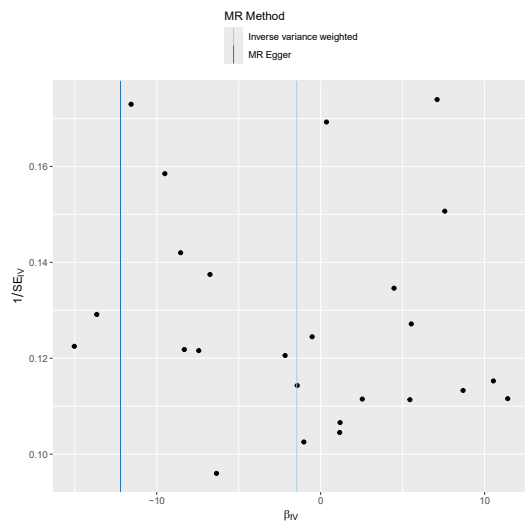
TP



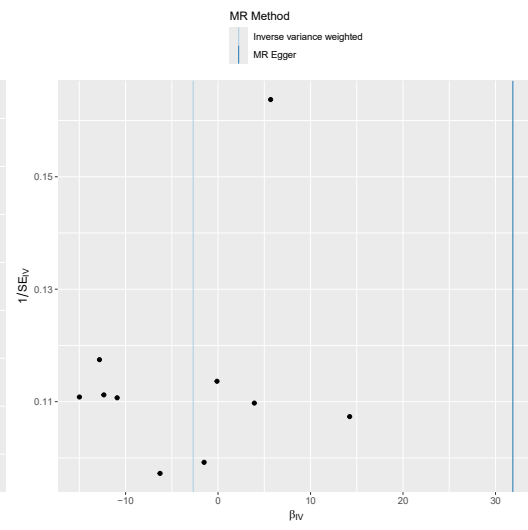
TRIG



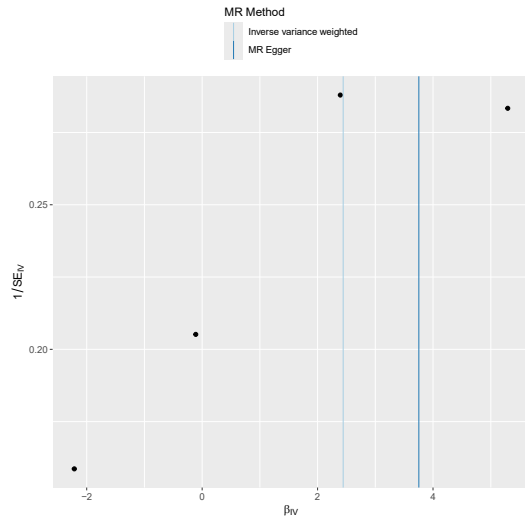
UA



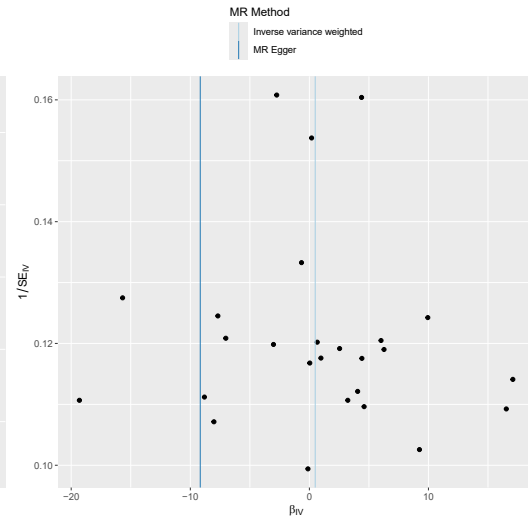
UCR



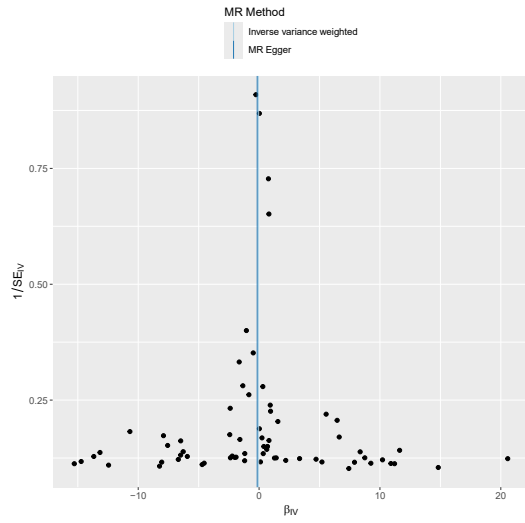
URK



URMA

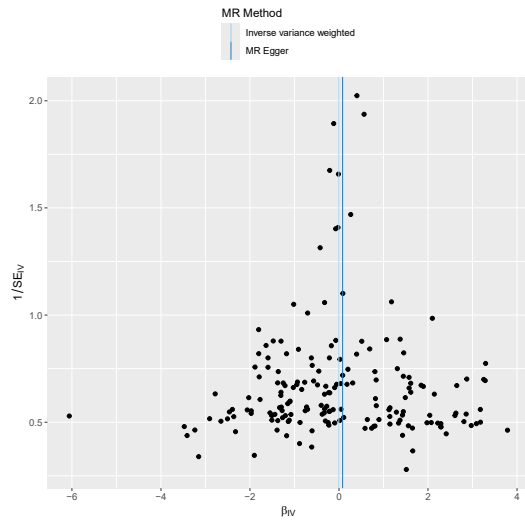


URNA

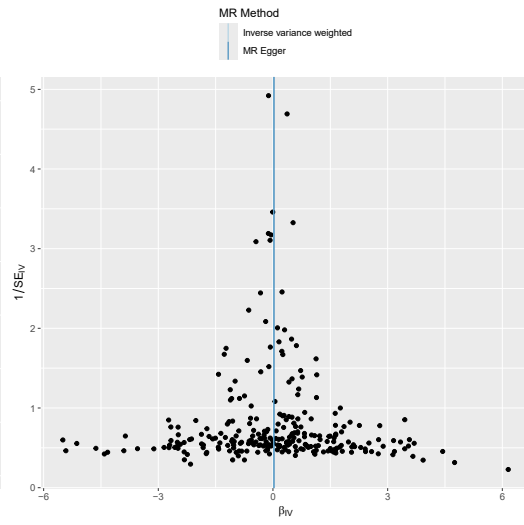


VITD

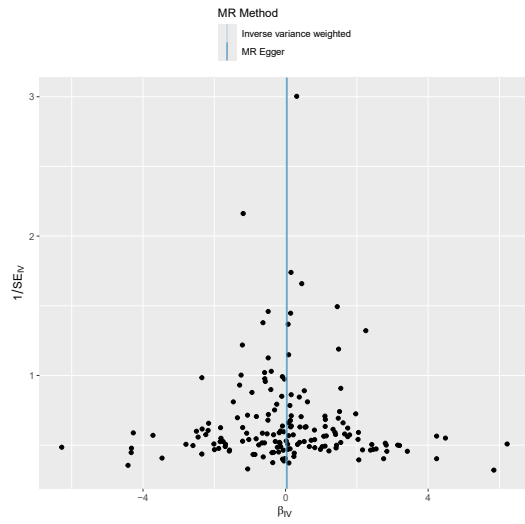
funnel plot of KCa



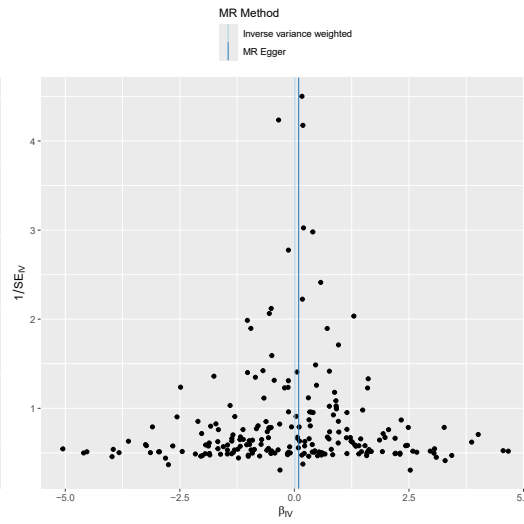
ALB



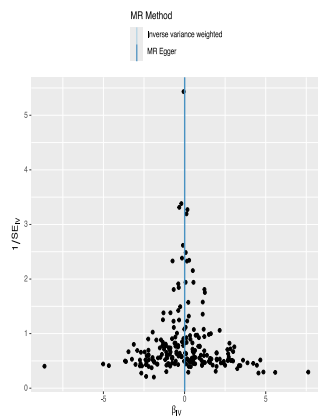
ALP



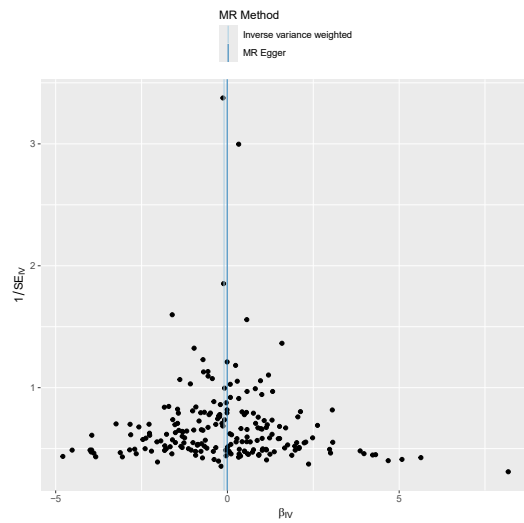
ALT



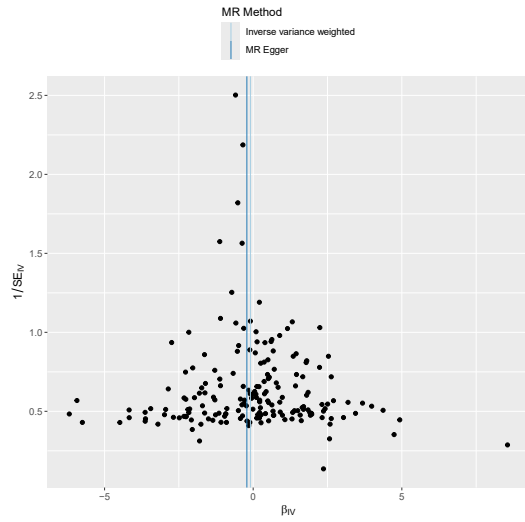
APOA



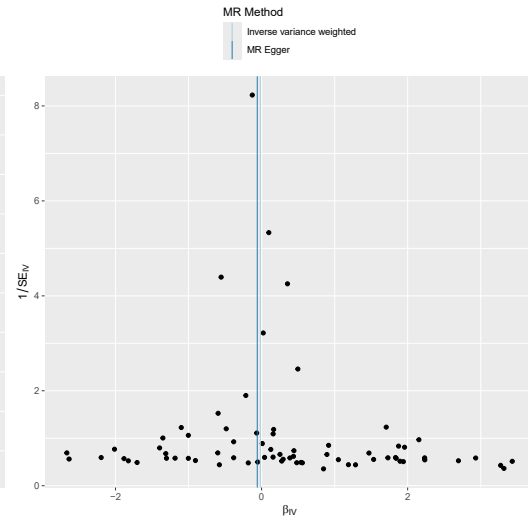
APOB



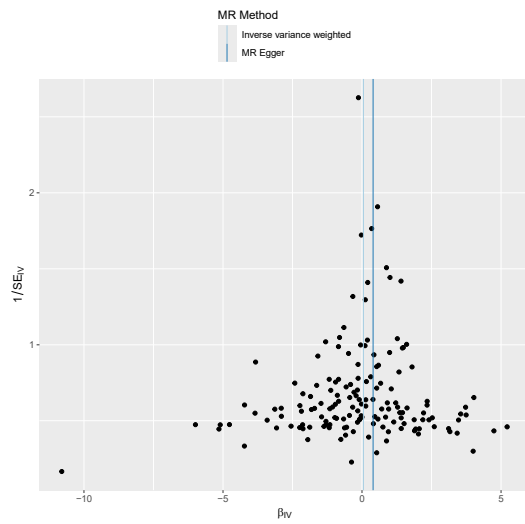
AST



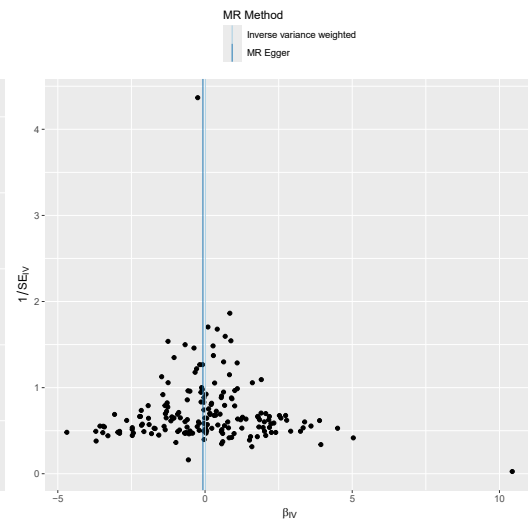
AST/ALT



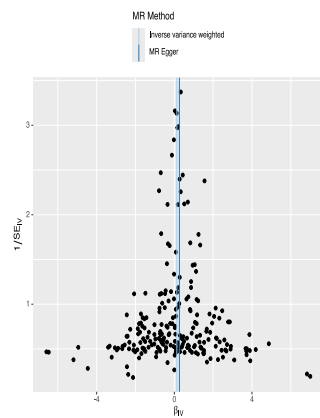
BILD



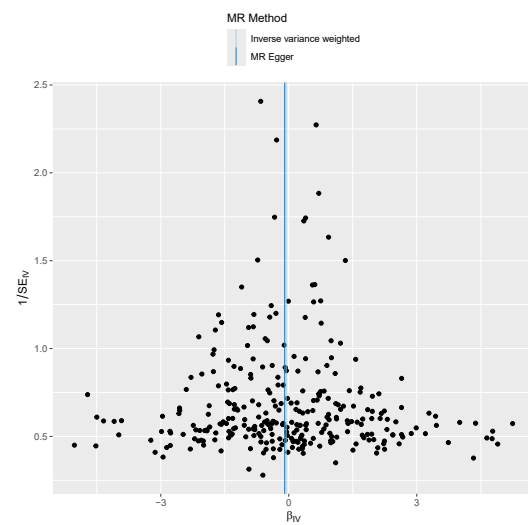
BUN



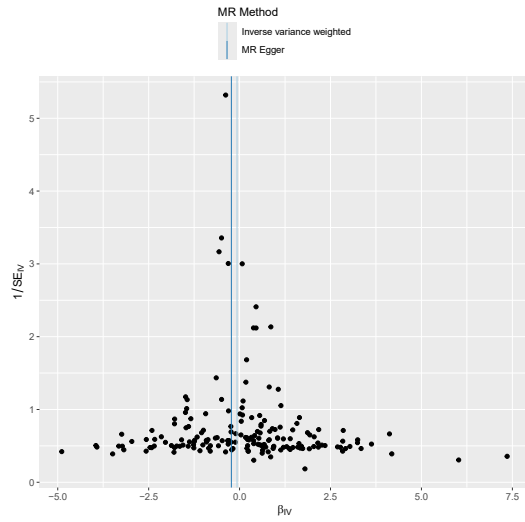
CA



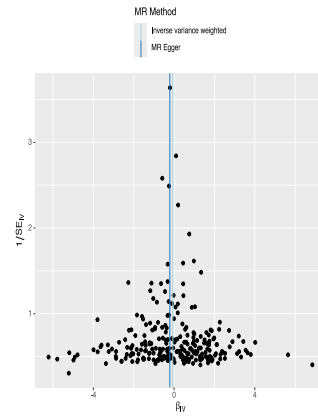
CHOL



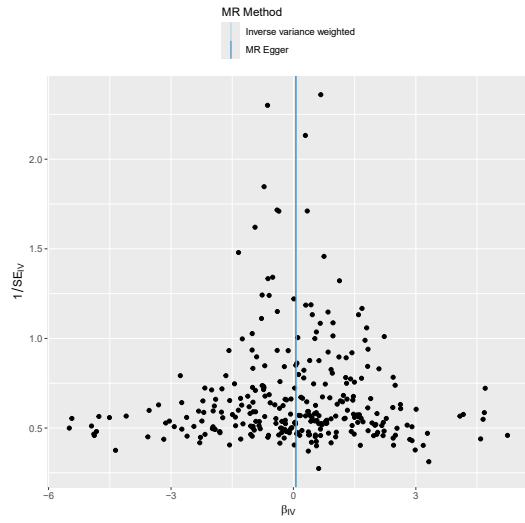
CRE



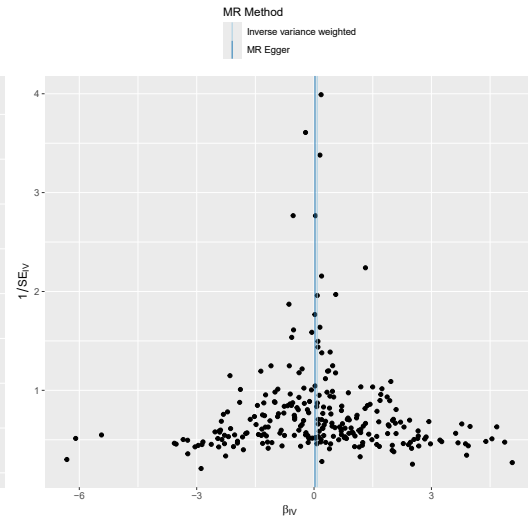
CRP



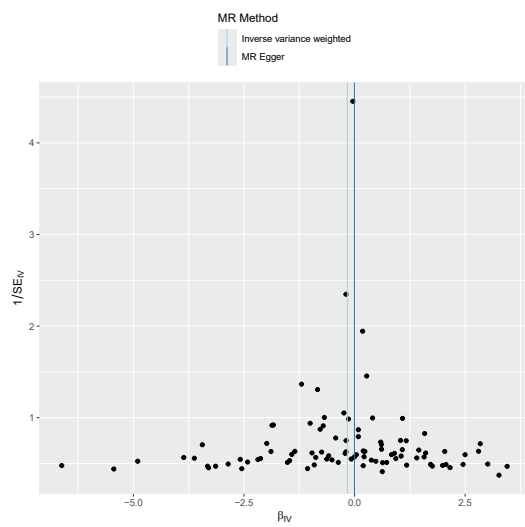
CYS



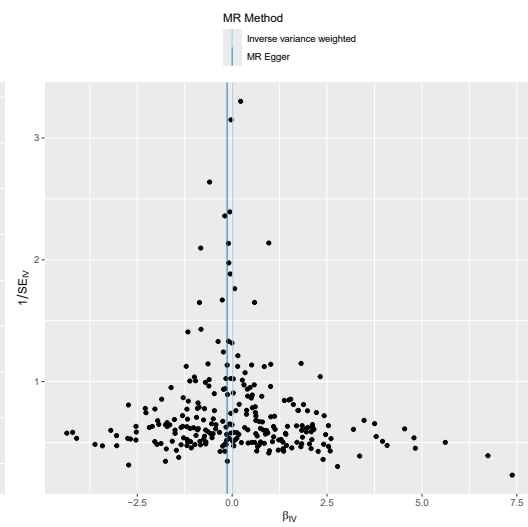
EGFR



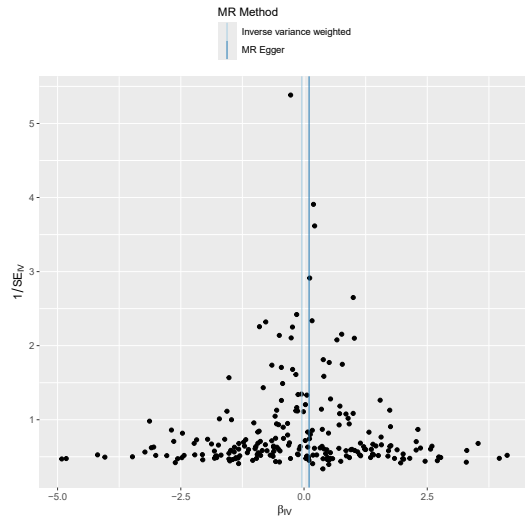
GGT



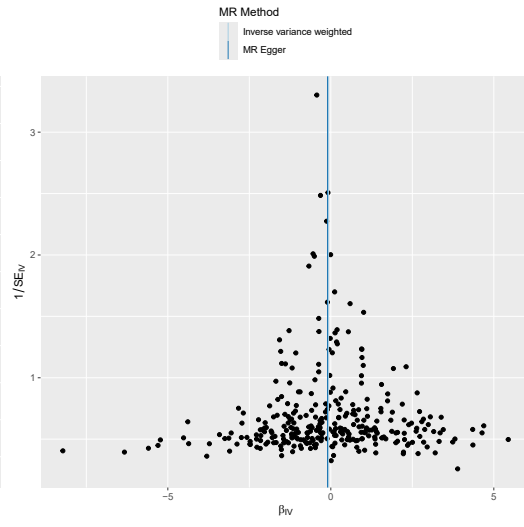
GLU



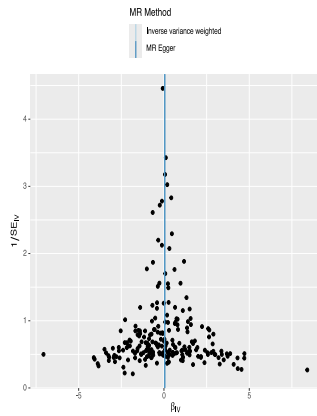
HBA1C



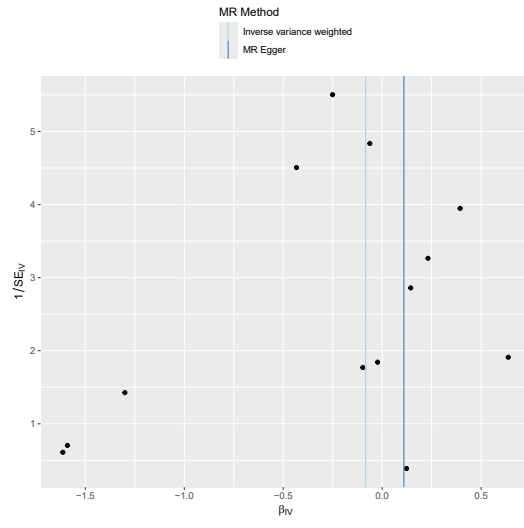
HDL



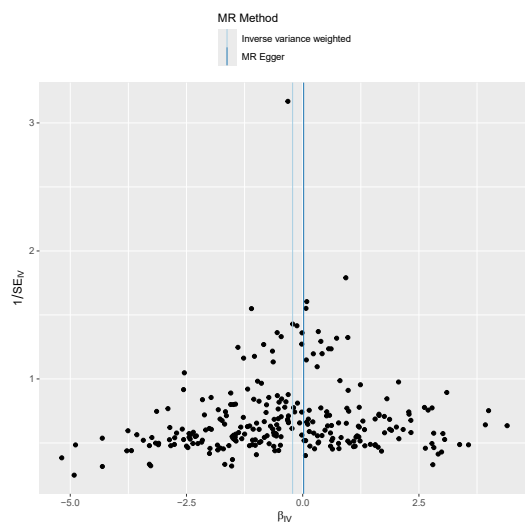
IGF-1



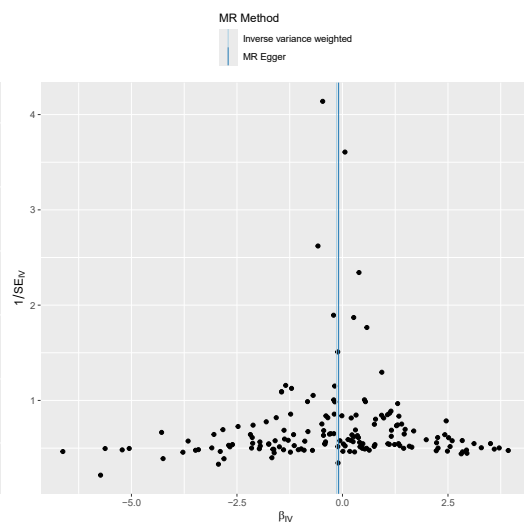
LDLD



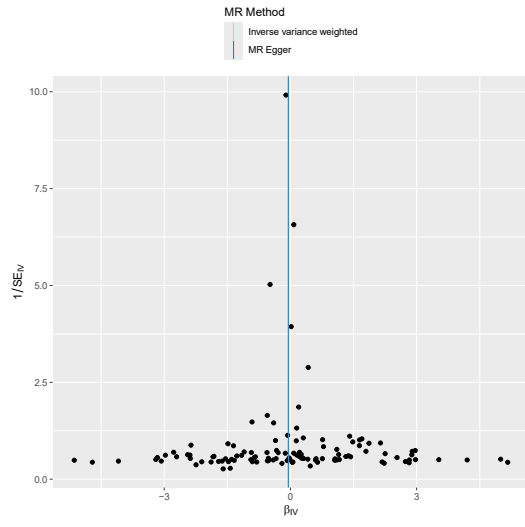
LPA



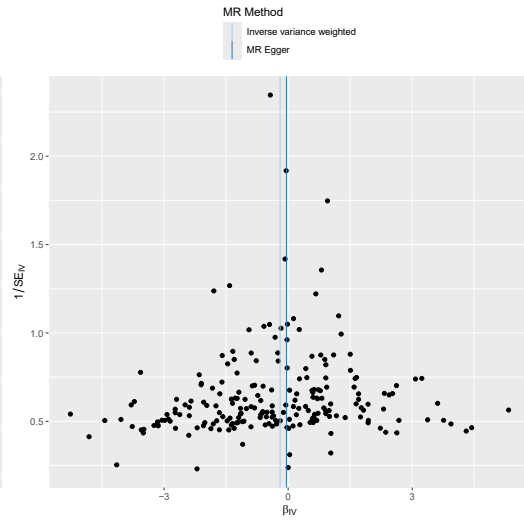
NAP



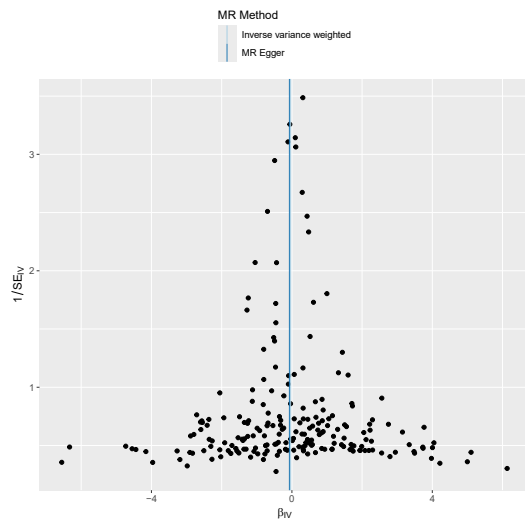
PHOS



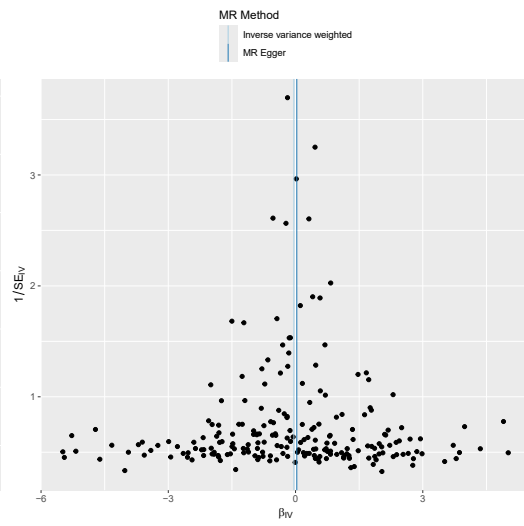
TBIL



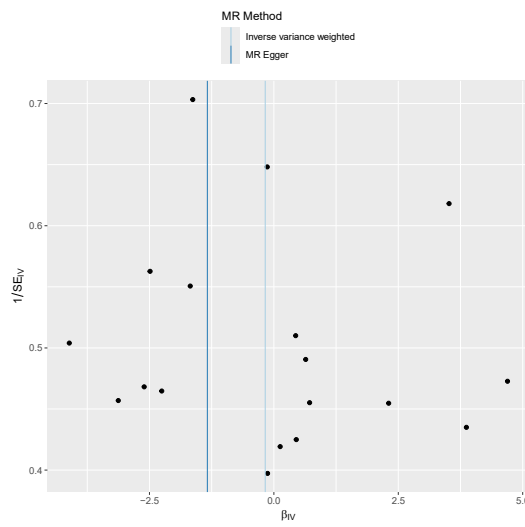
TP



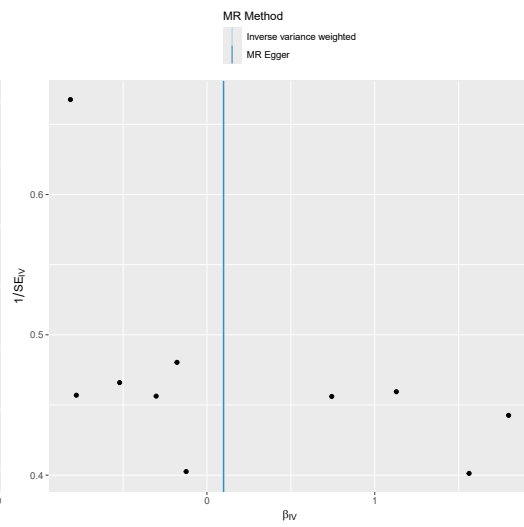
TRIG



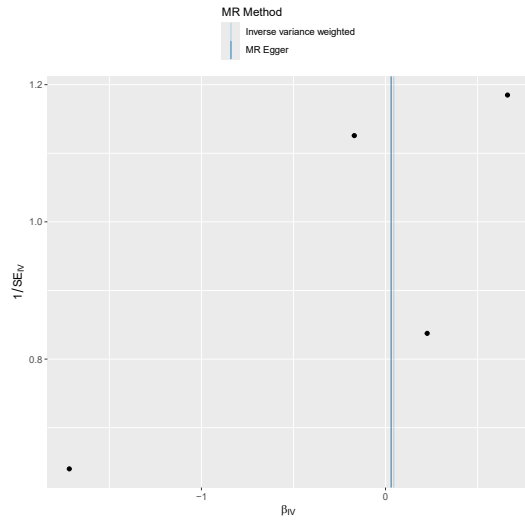
UA



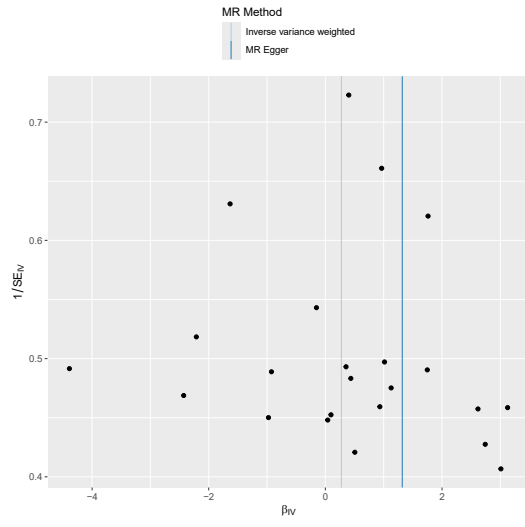
UCR



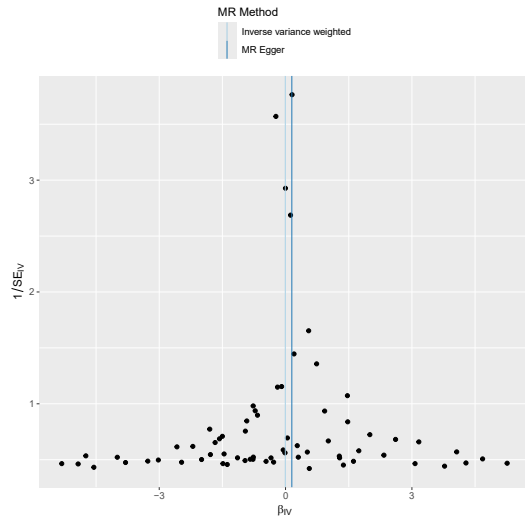
URK



URMA



URNA



VITD

STROBE-MR checklist of recommended items to address in reports of Mendelian randomization studies^{1 2}

Item No.	Section	Checklist item
1	TITLE and ABSTRACT	Indicate Mendelian randomization (MR) as the study's design in the title and/or the abstract if that is a main purpose of the study
INTRODUCTION		
2	Background	Explain the scientific background and rationale for the reported study. What is the exposure? Is a potential causal relationship between exposure and outcome plausible? Justify why MR is a helpful method to address the study question
3	Objectives	State specific objectives clearly, including pre-specified causal hypotheses (if any). State that MR is a method that, under specific assumptions, intends to estimate causal effects
METHODS		
4	Study design and data sources	Present key elements of the study design early in the article. Consider including a table listing sources of data for all phases of the study. For each data source contributing to the analysis, describe the following: <ul style="list-style-type: none"> a) Setting: Describe the study design and the underlying population, if possible. Describe the setting, locations, and relevant dates, including periods of recruitment, exposure, follow-up, and data collection, when available. b) Participants: Give the eligibility criteria, and the sources and methods of selection of participants. Report the sample size, and whether any power or sample size calculations were carried out prior to the main analysis c) Describe measurement, quality control and selection of genetic variants d) For each exposure, outcome, and other relevant variables, describe methods of assessment and diagnostic criteria for diseases e) Provide details of ethics committee approval and participant informed consent, if relevant
5	Assumptions	Explicitly state the three core IV assumptions for the main analysis (relevance, independence and exclusion restriction) as well assumptions for any additional or sensitivity analysis
6	Statistical methods: main analysis	Describe statistical methods and statistics used <ul style="list-style-type: none"> a) Describe how quantitative variables were handled in the analyses (i.e., scale, units, model) b) Describe how genetic variants were handled in the analyses and, if applicable, how their weights were selected c) Describe the MR estimator (e.g. two-stage least squares, Wald ratio) and related statistics. Detail the included covariates and, in case of two-sample MR, whether the same covariate set was used for adjustment in the two samples d) Explain how missing data were addressed e) If applicable, indicate how multiple testing was addressed

7	Assessment of assumptions	Describe any methods or prior knowledge used to assess the assumptions or justify their validity
8	Sensitivity analyses and additional analyses	Describe any sensitivity analyses or additional analyses performed (e.g. comparison of effect estimates from different approaches, independent replication, bias analytic techniques, validation of instruments, simulations)
9	Software and pre-registration	<ul style="list-style-type: none"> a) Name statistical software and package(s), including version and settings used b) State whether the study protocol and details were pre-registered (as well as when and where)
RESULTS		
10	Descriptive data	<ul style="list-style-type: none"> a) Report the numbers of individuals at each stage of included studies and reasons for exclusion. Consider use of a flow diagram b) Report summary statistics for phenotypic exposure(s), outcome(s), and other relevant variables (e.g. means, SDs, proportions) c) If the data sources include meta-analyses of previous studies, provide the assessments of heterogeneity across these studies d) For two-sample MR: <ul style="list-style-type: none"> i. Provide justification of the similarity of the genetic variant-exposure associations between the exposure and outcome samples ii. Provide information on the number of individuals who overlap between the exposure and outcome studies
11	Main results	<ul style="list-style-type: none"> a) Report the associations between genetic variant and exposure, and between genetic variant and outcome, preferably on an interpretable scale b) Report MR estimates of the relationship between exposure and outcome, and the measures of uncertainty from the MR analysis, on an interpretable scale, such as odds ratio or relative risk per SD difference c) If relevant, consider translating estimates of relative risk into absolute risk for a meaningful time period d) Consider plots to visualize results (e.g. forest plot, scatterplot of associations between genetic variants and outcome versus between genetic variants and exposure)
12	Assessment of assumptions	<ul style="list-style-type: none"> a) Report the assessment of the validity of the assumptions b) Report any additional statistics (e.g., assessments of heterogeneity across genetic variants, such as I^2, Q statistic or E-value)
13	Sensitivity analyses and additional analyses	

		a) Report any sensitivity analyses to assess the robustness of the main results to violations of the assumptions
		b) Report results from other sensitivity analyses or additional analyses
		c) Report any assessment of direction of causal relationship (e.g., bidirectional MR)
		d) When relevant, report and compare with estimates from non-MR analyses
		e) Consider additional plots to visualize results (e.g., leave-one-out analyses)
DISCUSSION		
14	Key results	Summarize key results with reference to study objectives
15	Limitations	Discuss limitations of the study, taking into account the validity of the IV assumptions, other sources of potential bias, and imprecision. Discuss both direction and magnitude of any potential bias and any efforts to address them
16	Interpretation	<p>a) Meaning: Give a cautious overall interpretation of results in the context of their limitations and in comparison with other studies</p> <p>b) Mechanism: Discuss underlying biological mechanisms that could drive a potential causal relationship between the investigated exposure and the outcome, and whether the gene-environment equivalence assumption is reasonable. Use causal language carefully, clarifying that IV estimates may provide causal effects only under certain assumptions</p> <p>c) Clinical relevance: Discuss whether the results have clinical or public policy relevance, and to what extent they inform effect sizes of possible interventions</p>
17	Generalizability	Discuss the generalizability of the study results (a) to other populations, (b) across other exposure periods/timings, and (c) across other levels of exposure
OTHER INFORMATION		
18	Funding	Describe sources of funding and the role of funders in the present study and, if applicable, sources of funding for the databases and original study or studies on which the present study is based
19	Data and data sharing	Provide the data used to perform all analyses or report where and how the data can be accessed, and reference these sources in the article. Provide the statistical code needed to reproduce the results in the article, or report whether the code is publicly accessible and if so, where
20	Conflicts of Interest	All authors should declare all potential conflicts of interest

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1. Skrivankova VW, Richmond RC, Woolf BAR, Yarmolinsky J, Davies NM, Swanson SA, et al. Strengthening the Reporting of Observational Studies in Epidemiology using Mendelian Randomization (STROBE-MR) Statement. JAMA. 2021;under review.
2. Skrivankova VW, Richmond RC, Woolf BAR, Davies NM, Swanson SA, VanderWeele TJ, et al. Strengthening the Reporting of Observational Studies in Epidemiology using Mendelian Randomisation (STROBE-MR): Explanation and Elaboration. BMJ. 2021;375:n2233.