

Contents

| | |
|---|------------|
| Glossary | xi |
| Acronyms | xii |
| 1 Introduction and Literature Review | 1 |
| 1.1 Cancer Research in the Post-Genomic Era | 1 |
| 1.1.1 Cancer is a Global Health Issue | 2 |
| 1.1.1.1 The Genetics and Molecular Biology of Cancers | 3 |
| 1.1.2 The Genomics Revolution in Cancer Research | 3 |
| 1.1.2.1 High-Throughput Technologies | 4 |
| 1.1.2.2 Bioinformatics and Genomic Data | 5 |
| 1.1.3 Genomics Projects | 5 |
| 1.1.3.1 The Cancer Genome Project | 6 |
| 1.1.3.2 The Cancer Genome Atlas Project | 6 |
| 1.1.4 Genomic Cancer Medicine | 8 |
| 1.1.4.1 Cancer Genes and Driver Mutations | 8 |
| 1.1.4.2 Precision Cancer Medicine | 9 |
| 1.1.4.3 Molecular Diagnostics and Pan-Cancer Medicine | 9 |
| 1.1.4.4 Targeted Therapeutics and Pharmacogenomics | 10 |
| 1.1.5 Systems and Network Biology | 11 |
| 1.2 Synthetic Lethal Cancer Medicine | 12 |
| 1.2.1 Synthetic Lethal Genetic Interactions | 13 |
| 1.2.2 Synthetic Lethal Concepts in Genetics | 13 |
| 1.2.3 Synthetic Lethality in Model Systems | 15 |
| 1.2.3.1 Synthetic Lethal Pathways and Networks | 15 |
| 1.2.3.2 Evolution of Synthetic Lethality | 16 |
| 1.2.4 Synthetic Lethality in Cancer | 17 |
| 1.2.5 Clinical Impact of Synthetic Lethality in Cancer | 18 |
| 1.2.6 High-throughput Screening for Synthetic Lethality | 20 |
| 1.2.6.1 Synthetic Lethal Screens | 21 |
| 1.2.7 Computational Prediction of Synthetic Lethality | 22 |
| 1.2.7.1 Bioinformatics Approaches to Genetic Interactions . . | 22 |
| 1.2.7.2 Comparative Genomics | 23 |
| 1.2.7.3 Analysis and Modelling of Protein Data | 26 |
| 1.2.7.4 Differential Gene Expression | 28 |
| 1.2.7.5 Data Mining and Machine Learning | 29 |

| | | |
|----------|--|-----------|
| 1.2.7.6 | Mutual Exclusivity and Bimodality | 32 |
| 1.2.7.7 | Rationale for Further Development | 33 |
| 1.3 | E-cadherin as a Synthetic Lethal Target | 33 |
| 1.3.1 | The <i>CDH1</i> gene and its Biological Functions | 34 |
| 1.3.1.1 | Cytoskeleton | 34 |
| 1.3.1.2 | Extracellular and Tumour Micro-environment | 34 |
| 1.3.1.3 | Cell-Cell Adhesion and Signalling | 35 |
| 1.3.2 | <i>CDH1</i> as a Tumour (and Invasion) Suppressor | 35 |
| 1.3.2.1 | Breast Cancers and Invasion | 35 |
| 1.3.3 | Hereditary Diffuse Gastric (and Lobular Breast) Cancer | 36 |
| 1.3.4 | Cell Line Models of <i>CDH1</i> Null Mutations | 37 |
| 1.4 | Summary and Research Direction of Thesis | 38 |
| 1.4.1 | Thesis Aims | 39 |
| 2 | Methods and Resources | 41 |
| 2.1 | Bioinformatics Resources for Genomics Research | 41 |
| 2.1.1 | Public Data and Software Packages | 41 |
| 2.1.1.1 | Cancer Genome Atlas Data | 42 |
| 2.1.1.2 | Reactome and Annotation Data | 43 |
| 2.2 | Data Handling | 43 |
| 2.2.1 | Normalisation | 43 |
| 2.2.2 | Sample Triage | 43 |
| 2.2.3 | Metagenes and the Singular Value Decomposition | 44 |
| 2.2.4 | Candidate Triage and Integration with Screen Data | 46 |
| 2.3 | Techniques | 46 |
| 2.3.1 | Statistical Procedures and Tests | 47 |
| 2.3.2 | Gene Set Over-representation Analysis | 48 |
| 2.3.3 | Clustering | 48 |
| 2.3.4 | Heatmap | 48 |
| 2.3.5 | Modelling and Simulations | 49 |
| 2.3.5.1 | Receiver Operating Characteristic Curves | 50 |
| 2.3.6 | Resampling Analysis | 50 |
| 2.4 | Pathway Structure Methods | 51 |
| 2.4.1 | Network and Graph Analysis | 51 |
| 2.4.2 | Sourcing Graph Structure Data | 52 |
| 2.4.3 | Constructing Pathway Subgraphs | 52 |
| 2.4.4 | Network Analysis Metrics | 53 |
| 2.5 | Implementation | 54 |
| 2.5.1 | Computational Resources and Linux Utilities | 54 |
| 2.5.2 | R Language and Packages | 55 |
| 2.5.3 | High Performance and Parallel Computing | 58 |
| 3 | Methods Developed During Thesis | 60 |
| 3.1 | A Synthetic Lethal Detection Methodology | 60 |
| 3.2 | Synthetic Lethal Simulation and Modelling | 62 |
| 3.2.1 | A Model of Synthetic Lethality in Expression Data | 63 |

| | | |
|----------|---|-----------|
| 3.2.2 | Simulation Procedure | 67 |
| 3.3 | Detecting Simulated Synthetic Lethal Partners | 70 |
| 3.3.1 | Binomial Simulation of Synthetic Lethality | 70 |
| 3.3.2 | Multivariate Normal Simulation of Synthetic Lethality | 72 |
| 3.3.2.1 | Multivariate Normal Simulation with Correlated Genes | 74 |
| 3.3.2.2 | Specificity with Query-Correlated Pathways | 82 |
| 3.4 | Graph Structure Methods | 84 |
| 3.4.1 | Upstream and Downstream Gene Detection | 84 |
| 3.4.1.1 | Permutation Analysis for Statistical Significance | 85 |
| 3.4.2 | Simulating Gene Expression from Graph Structures | 86 |
| 3.5 | Customised Functions and Packages Developed | 90 |
| 3.5.1 | Synthetic Lethal Interaction Prediction Tool | 90 |
| 3.5.2 | Data Visualisation | 91 |
| 3.5.3 | Extensions to the iGraph Package | 92 |
| 3.5.3.1 | Sampling Simulated Data from Graph Structures | 92 |
| 3.5.3.2 | Plotting Directed Graph Structures | 92 |
| 3.5.3.3 | Computing Information Centrality | 94 |
| 3.5.3.4 | Testing Pathway Structure with Permutation Testing | 94 |
| 3.5.3.5 | Metapackage to Install iGraph Functions | 95 |
| 4 | Synthetic Lethal Analysis of Gene Expression Data | 96 |
| 4.1 | Synthetic Lethal Genes in Breast Cancer | 97 |
| 4.1.1 | Synthetic Lethal Pathways in Breast Cancer | 98 |
| 4.1.2 | Expression Profiles of Synthetic Lethal Partners | 100 |
| 4.1.2.1 | Subgroup Pathway Analysis | 103 |
| 4.2 | Comparing Synthetic Lethal Gene Candidates | 105 |
| 4.2.1 | Primary siRNA Screen Candidates | 105 |
| 4.2.2 | Comparison with Correlation | 105 |
| 4.2.3 | Comparison with Primary Screen Viability | 108 |
| 4.2.4 | Comparison with Secondary siRNA Screen Validation | 110 |
| 4.2.5 | Comparison to Primary Screen at Pathway Level | 111 |
| 4.2.5.1 | Resampling Genes for Pathway Enrichment | 113 |
| 4.2.6 | Integrating Synthetic Lethal Pathways and Screens | 118 |
| 4.3 | Synthetic Lethal Pathway Metagenes | 119 |
| 4.4 | Replication in Stomach Cancer | 121 |
| 4.5 | Discussion | 122 |
| 4.5.1 | Strengths of the SLIPT Methodology | 122 |
| 4.5.2 | Synthetic Lethal Pathways for E-cadherin | 123 |
| 4.5.3 | Replication and Validation | 125 |
| 4.5.3.1 | Integration with siRNA Screening | 125 |
| 4.5.3.2 | Replication across Tissues | 126 |
| 4.6 | Summary | 126 |

| | | |
|----------|--|------------|
| 5 | Synthetic Lethal Pathway Structure | 128 |
| 5.1 | Synthetic Lethal Genes in Reactome Pathways | 128 |
| 5.1.1 | The PI3K/AKT Pathway | 129 |
| 5.1.2 | The Extracellular Matrix | 131 |
| 5.1.3 | G Protein Coupled Receptors | 134 |
| 5.1.4 | Gene Regulation and Translation | 134 |
| 5.2 | Network Analysis of Synthetic Lethal Genes | 136 |
| 5.2.1 | Gene Connectivity and Vertex Degree | 137 |
| 5.2.2 | Gene Importance and Centrality | 138 |
| 5.2.2.1 | Information Centrality | 138 |
| 5.2.2.2 | PageRank Centrality | 140 |
| 5.3 | Relationships between Synthetic Lethal Genes | 141 |
| 5.3.1 | Detecting Upstream or Downstream Synthetic Lethality | 142 |
| 5.3.2 | Resampling for Synthetic Lethal Pathway Structure | 144 |
| 5.4 | Discussion | 146 |
| 5.5 | Summary | 148 |
| 6 | Simulation and Modelling of Synthetic Lethal Pathways | 149 |
| 6.1 | Synthetic Lethal Detection Methods | 150 |
| 6.1.1 | Performance of SLIPT and χ^2 across Quantiles | 150 |
| 6.1.1.1 | Correlated Query Genes affects Specificity | 154 |
| 6.1.2 | Alternative Synthetic Lethal Detection Strategies | 156 |
| 6.1.2.1 | Correlation for Synthetic Lethal Detection | 156 |
| 6.1.2.2 | Testing for Bimodality with BiSEp | 158 |
| 6.2 | Simulations with Graph Structures | 159 |
| 6.2.1 | Performance over Graph Structures | 160 |
| 6.2.1.1 | Simple Graph Structures | 160 |
| 6.2.1.2 | Constructed Graph Structures | 162 |
| 6.2.2 | Performance with Inhibitions | 165 |
| 6.2.3 | Synthetic Lethality across Graph Structures | 170 |
| 6.2.4 | Performance within a Large Simulated Datasets | 174 |
| 6.3 | Simulations in More Complex Graph Structures | 178 |
| 6.3.1 | Simulations over Pathway-based Graphs | 179 |
| 6.3.2 | Pathway Structures in a Large Simulated Datasets | 182 |
| 6.4 | Discussion | 185 |
| 6.4.1 | Simulation Procedure | 185 |
| 6.4.2 | Comparing Methods with Simulated Data | 186 |
| 6.4.3 | Design and Performance of SLIPT | 187 |
| 6.4.4 | Simulations from Graph Structures | 189 |
| 6.5 | Summary | 190 |
| 7 | Discussion | 192 |
| 7.1 | Synthetic Lethality and <i>CDH1</i> Biology | 192 |
| 7.1.1 | Established Functions of <i>CDH1</i> | 193 |
| 7.1.2 | The Molecular Role of <i>CDH1</i> in Cancer | 193 |
| 7.2 | Significance | 194 |

| | | |
|----------|---|------------|
| 7.2.1 | Synthetic Lethality in the Genomic Era | 194 |
| 7.2.2 | Clinical Interventions based on Synthetic Lethality | 196 |
| 7.3 | Future Directions | 197 |
| 7.4 | Conclusions | 199 |
| | Bibliography | 201 |
| A | Sample Quality | 225 |
| A.1 | Sample Correlation | 225 |
| A.2 | Replicate Samples in TCGA Breast Cancer Data | 228 |
| B | Software Used for Thesis | 232 |
| C | Mutation Analysis in Breast Cancer | 241 |
| C.1 | Synthetic Lethal Genes and Pathways | 241 |
| C.2 | Synthetic Lethal Expression Profiles | 242 |
| C.3 | Comparison to Primary Screen | 245 |
| C.3.1 | Resampling Analysis | 247 |
| C.4 | Compare SLIPT genes | 249 |
| D | Metagene Analysis | 251 |
| D.1 | Pathway Signature Expression | 251 |
| D.2 | Synthetic Lethal Reactome Metagenes | 255 |
| E | Intrinsic Subtyping | 256 |
| F | Stomach Expression Analysis | 258 |
| F.1 | Synthetic Lethal Genes and Pathways | 258 |
| F.2 | Comparison to Primary Screen | 262 |
| F.2.1 | Resampling Analysis | 264 |
| F.3 | Metagene Analysis | 266 |
| G | Synthetic Lethal Genes in Pathways | 267 |
| H | Network Analysis for Mutation SLIPT | 274 |
| I | Pathway Structure for Mutation SLIPT | 277 |
| J | Performance of SLIPT and χ^2 | 279 |
| J.1 | Correlated Query Genes affects Specificity | 285 |
| K | Simulations on Graph Structures | 291 |
| K.0.1 | Simulations from Inhibiting Graph Structures | 292 |
| K.1 | Simulation across Graph Structures | 295 |
| K.2 | Simulations from Complex Graph Structures | 299 |
| K.2.1 | Simulations from Complex Inhibiting Graphs | 302 |
| K.3 | Simulations from Pathway Graph Structures | 308 |

List of Figures

| | | |
|------|--|-----|
| 1.1 | Synthetic genetic interactions | 14 |
| 1.2 | Synthetic lethality in cancer | 17 |
| 2.1 | Read count density | 45 |
| 2.2 | Read count sample mean | 45 |
| 3.1 | Framework for synthetic lethal prediction | 61 |
| 3.2 | Synthetic lethal prediction adapted for mutation | 62 |
| 3.3 | A model of synthetic lethal gene expression | 64 |
| 3.4 | Modelling synthetic lethal gene expression | 65 |
| 3.5 | Synthetic lethality with multiple genes | 66 |
| 3.6 | Simulating gene function | 68 |
| 3.7 | Simulating synthetic lethal gene function | 68 |
| 3.8 | Simulating synthetic lethal gene expression | 69 |
| 3.9 | Performance of binomial simulations | 71 |
| 3.10 | Comparison of statistical performance | 71 |
| 3.11 | Performance of multivariate normal simulations | 73 |
| 3.12 | Simulating expression with correlated gene blocks | 75 |
| 3.13 | Simulating expression with correlated gene blocks | 76 |
| 3.14 | Synthetic lethal prediction across simulations | 78 |
| 3.15 | Performance with correlations | 79 |
| 3.16 | Comparison of statistical performance with correlation structure | 80 |
| 3.17 | Performance with query correlations | 81 |
| 3.18 | Statistical evaluation of directional criteria | 82 |
| 3.19 | Performance of directional criteria | 83 |
| 3.20 | Simulated graph structures | 87 |
| 3.21 | Simulating expression from a graph structure | 88 |
| 3.22 | Simulating expression from graph structure with inhibitions | 89 |
| 3.23 | Demonstration of violin plots with custom features | 93 |
| 3.24 | Demonstration of annotated heatmap | 93 |
| 3.25 | Simulating graph structures | 94 |
| 4.1 | Synthetic lethal expression profiles of analysed samples | 101 |
| 4.2 | Comparison of SLIPT with siRNA | 106 |
| 4.3 | Comparison of SLIPT and siRNA genes with correlation | 106 |
| 4.4 | Comparison of SLIPT and siRNA genes with correlation | 108 |
| 4.5 | Comparison of SLIPT and siRNA genes with screen viability | 109 |

| | | |
|------|---|-----|
| 4.6 | Comparison of SLIPT genes with siRNA screen viability | 109 |
| 4.7 | Resampled intersection of SLIPT and siRNA candidate genes | 114 |
| 5.1 | Synthetic lethality in the PI3K cascade | 130 |
| 5.2 | Synthetic lethality in Elastic Fibre Formation | 132 |
| 5.3 | Synthetic lethality in Fibrin Clot Formation | 133 |
| 5.4 | Synthetic lethality in the GPCRs | 135 |
| 5.5 | Synthetic lethality and vertex degree | 137 |
| 5.6 | Synthetic lethality and centrality | 139 |
| 5.7 | Synthetic lethality and PageRank | 141 |
| 5.8 | Structure of synthetic lethality resampling | 143 |
| 6.1 | Performance of χ^2 and SLIPT across quantiles | 152 |
| 6.2 | Performance of χ^2 and SLIPT across quantiles with more genes | 153 |
| 6.3 | Performance of χ^2 and SLIPT across quantiles with query correlation | 154 |
| 6.4 | Performance of χ^2 and SLIPT across quantiles with query correlation and more genes | 155 |
| 6.5 | Performance of negative correlation and SLIPT | 157 |
| 6.6 | Simple graph structures | 160 |
| 6.7 | Performance of simulations on a simple graph | 161 |
| 6.8 | Performance of simulations is similar in simple graphs | 163 |
| 6.9 | Performance of simulations on a pathway | 164 |
| 6.10 | Performance of simulations on a simple graph with inhibition | 166 |
| 6.11 | Performance is higher on a simple inhibiting graph | 167 |
| 6.12 | Performance of simulations on a constructed graph with inhibition | 168 |
| 6.13 | Performance is affected by inhibition in graphs | 170 |
| 6.14 | Detection of synthetic lethality within a graph structure | 172 |
| 6.15 | Performance of simulations including a simple graph | 175 |
| 6.16 | Performance on a simple graph improves with more genes | 176 |
| 6.17 | Performance on an inhibiting graph improves with more genes | 178 |
| 6.18 | Performance of simulations on the PI3K cascade | 181 |
| 6.19 | Performance of simulations including the PI3K cascade | 183 |
| 6.20 | Performance on pathways improves with more genes | 184 |
| A.1 | Correlation profiles of removed samples | 226 |
| A.2 | Correlation analysis and sample removal | 227 |
| A.3 | Replicate excluded samples | 228 |
| A.4 | Replicate samples with all remaining | 229 |
| A.5 | Replicate samples with some excluded | 230 |
| C.1 | Synthetic lethal expression profiles of analysed samples | 243 |
| C.2 | Comparison of mtSLIPT to short interfering RNA (siRNA) | 245 |
| C.3 | Compare mtSLIPT and siRNA genes with correlation | 249 |
| C.4 | Compare mtSLIPT and siRNA genes with correlation | 249 |
| C.5 | Compare mtSLIPT and siRNA genes with siRNA viability | 250 |
| D.1 | Pathway metagene expression profiles | 253 |

| | | |
|------|---|-----|
| D.2 | Expression profiles for estrogen receptor related genes | 254 |
| F.1 | Synthetic lethal expression profiles of stomach samples | 260 |
| F.2 | Comparison of SLIPT in stomach to siRNA | 262 |
| G.1 | Synthetic lethality in the PI3K/AKT pathway | 267 |
| G.2 | Synthetic lethality in the PI3K/AKT pathway in cancer | 268 |
| G.3 | Synthetic lethality in the Extracellular Matrix | 269 |
| G.4 | Synthetic lethality in the GPCR Downstream | 270 |
| G.5 | Synthetic lethality in the Translation Elongation | 271 |
| G.6 | Synthetic lethality in the Nonsense-mediated Decay | 272 |
| G.7 | Synthetic lethality in the 3' UTR | 273 |
| H.1 | Synthetic lethality and vertex degree | 274 |
| H.2 | Synthetic lethality and centrality | 275 |
| H.3 | Synthetic lethality and PageRank | 275 |
| I.1 | Structure of synthetic lethality resampling | 277 |
| J.1 | Performance of χ^2 and SLIPT across quantiles | 279 |
| J.2 | Performance of χ^2 and SLIPT across quantiles | 281 |
| J.3 | Performance of χ^2 and SLIPT across quantiles with more genes | 283 |
| J.4 | Performance of χ^2 and SLIPT across quantiles with query correlation | 285 |
| J.5 | Performance of χ^2 and SLIPT across quantiles with query correlation | 287 |
| J.6 | Performance of χ^2 and SLIPT across quantiles with query correlation and more genes | 289 |
| K.1 | Performance of simulations on a simple graph | 291 |
| K.2 | Performance of simulations on an inhibiting graph | 292 |
| K.3 | Performance of simulations on a constructed graph with inhibition | 293 |
| K.4 | Performance of simulations on a constructed graph with inhibition | 294 |
| K.5 | Detection of synthetic lethality within a graph structure | 295 |
| K.6 | Detection of synthetic lethality within an inhibiting graph | 297 |
| K.7 | Detection of synthetic lethality within an inhibiting graph | 298 |
| K.8 | Performance of simulations on a branching graph | 299 |
| K.9 | Performance of simulations on a complex graph | 300 |
| K.10 | Performance of simulations on a large graph | 301 |
| K.11 | Performance of simulations on a branching graph with inhibition | 302 |
| K.12 | Performance of simulations on a branching graph with inhibition | 303 |
| K.13 | Performance of simulations on a complex graph with inhibition | 304 |
| K.14 | Performance of simulations on a complex graph with inhibition | 305 |
| K.15 | Performance of simulations on a large constructed graph with inhibition | 306 |
| K.16 | Performance of simulations on a large constructed graph with inhibition | 307 |
| K.17 | Performance of simulations on the $G_{\alpha i}$ signalling pathway | 308 |
| K.18 | Performance of simulations including the $G_{\alpha i}$ signalling pathway | 309 |

List of Tables

| | | |
|-----|--|-----|
| 1.1 | Methods for predicting genetic interactions | 23 |
| 1.2 | Methods for predicting synthetic lethality in cancer | 24 |
| 1.3 | Methods used by Wu <i>et al.</i> (2014) | 25 |
| 2.1 | Excluded samples by batch and clinical characteristics. | 44 |
| 2.2 | Computers used during thesis | 54 |
| 2.3 | Linux utilities and applications used during thesis | 55 |
| 2.4 | R installations used during thesis | 56 |
| 2.5 | R Packages used during thesis | 56 |
| 2.6 | R packages developed during thesis | 58 |
| 4.1 | Candidate synthetic lethal gene partners of <i>CDH1</i> from SLIPT | 98 |
| 4.2 | Pathways for <i>CDH1</i> partners from SLIPT | 99 |
| 4.3 | Pathways for clusters of <i>CDH1</i> partners from SLIPT | 104 |
| 4.4 | ANOVA for synthetic lethality and correlation with <i>CDH1</i> | 107 |
| 4.5 | Comparison of Synthetic Lethal Interaction Prediction Tool (SLIPT) genes against secondary siRNA screen | 111 |
| 4.6 | Pathways for <i>CDH1</i> partners from SLIPT and siRNA | 112 |
| 4.7 | Pathways for <i>CDH1</i> partners from SLIPT | 115 |
| 4.8 | Pathways for <i>CDH1</i> partners from SLIPT and siRNA primary screen . | 116 |
| 4.9 | Examples of candidate metagenes synthetic lethal for <i>CDH1</i> from SLIPT | 120 |
| 5.1 | ANOVA for synthetic lethality and vertex degree | 138 |
| 5.2 | ANOVA for synthetic lethality and information centrality | 139 |
| 5.3 | ANOVA for synthetic lethality and PageRank centrality | 140 |
| 5.4 | Resampling for pathway structure of synthetic lethal detection methods | 145 |
| B.1 | Complete list of R packages used during this thesis | 232 |
| C.1 | Candidate synthetic lethal gene partners of <i>CDH1</i> from mtSLIPT . . . | 241 |
| C.2 | Pathways for <i>CDH1</i> partners from mtSLIPT | 242 |
| C.3 | Pathways for clusters of <i>CDH1</i> partners from mtSLIPT | 244 |
| C.4 | Pathways for <i>CDH1</i> partners from mtSLIPT and siRNA | 246 |
| C.5 | Pathways for <i>CDH1</i> partners from mtSLIPT | 247 |
| C.6 | Pathways for <i>CDH1</i> partners from mtSLIPT and siRNA primary screen | 248 |
| D.1 | Candidate synthetic lethal metagenes against <i>CDH1</i> from mtSLIPT . . | 255 |

| | | |
|-----|--|-----|
| E.1 | Comparison of intrinsic subtypes | 256 |
| F.1 | Synthetic lethal gene partners of <i>CDH1</i> from SLIPT in stomach cancer | 258 |
| F.2 | Pathways for <i>CDH1</i> partners from SLIPT in stomach cancer | 259 |
| F.3 | Pathways for clusters of <i>CDH1</i> partners in stomach SLIPT | 261 |
| F.4 | Pathways for <i>CDH1</i> partners from SLIPT and siRNA | 263 |
| F.5 | Pathways for <i>CDH1</i> partners from SLIPT in stomach cancer | 264 |
| F.6 | Pathways for <i>CDH1</i> partners from SLIPT in stomach and siRNA | 265 |
| F.7 | Synthetic lethal metagenes against <i>CDH1</i> in stomach cancer | 266 |
| H.1 | ANOVA for synthetic lethality and vertex degree | 276 |
| H.2 | ANOVA for synthetic lethality and information centrality | 276 |
| H.3 | ANOVA for synthetic lethality and PageRank centrality | 276 |
| I.1 | Resampling for pathway structure of synthetic lethal detection methods | 278 |

Glossary

| | |
|-------------------|--|
| bioinformatics | Statistical or computational approaches to biological data or research tools. |
| gene expression | A measure of the relative expression of each gene from the mRNA extracted from (pooled) cells. |
| genome | All of the DNA sequence in the genome. |
| genomic | The use of data from all genes in the genome. |
| graph or network | A mathematical structure modelling or depicting the relationships between elements. |
| metagene | A consistent signal of expression for a collection of genes such as a biological pathway, derived from singular value decomposition. |
| microarray | A high-throughput technique to measure presence or abundance of nucleic acid sequences from binding to probes. |
| mutation | A change in DNA sequence that disrupts gene function. |
| RNA-Seq | The generation of transcriptome data from sequencing RNA. |
| synthetic lethal | Genetic interactions where inactivation of multiple genes is inviable (or deleterious) which are viable if inactivated separately. |
| tumour suppressor | A gene potentially causes cancer, typically by disruption of functions which protect the cell from cancer. |

Acronyms

| | |
|---------|--|
| ANOVA | Analysis of Variance. |
| AUROC | Area Under the Receiver Operating Characteristic (curve). |
| BiSEp | Bimodal Subsetting Expression. |
| DNA | Deoxyribonucleic Acid. |
| FDR | False Discovery Rate. |
| HPC | High Performance Computing. |
| mtSLIPT | Synthetic Lethal Interaction Prediction Tool (against mutation). |
| NeSI | New Zealand eScience Infrastructure. |
| PI3K | Phosphoinositide 3-kinase. |
| ROC | Receiver Operating Characteristic (curve). |
| siRNA | Short Interfering RNA. |
| SLIPT | Synthetic Lethal Interaction Prediction Tool. |
| Slurm | Simple Linux Utility for Resource Management. |
| TCGA | The Cancer Genome Atlas (genomics project). |

Chapter 6

Simulation and Modelling of Synthetic Lethal Pathways

Simulation and modelling of [synthetic lethality](#) in [gene expression](#) was revisited in greater detail in this chapter, building upon the results which supported the use of [SLIPT](#) (in Section 3.3). In Chapter 3, a procedure for generating simulated data with underlying (known) [synthetic lethal](#) partners of a query gene, such as *CDH1*, was developed (as described in Section 3.2.2) by sampling from a multivariate normal distribution based on a statistical model of [synthetic lethality](#) in [gene expression](#) data (as described in Section 3.2.1). This simulation framework was applied to simulated data (in Section 3.3), including simple correlation structures to assess the statistical performance of the [SLIPT](#) methodology and support its use as a computational approach for detecting [synthetic lethal](#) candidates from [expression](#) data throughout this thesis (Chapters 4 and 5).

While this basic framework provided some support for the use of [SLIPT](#), further investigations with simulations were conducted to assess the strengths and limitations of the [SLIPT](#) methodology, compare it to alternative statistical approaches to [synthetic lethal](#) detection, and assess its performance under more complex correlation structures. Together these simulation investigations assess the performance of the [SLIPT](#) methodology, including on pathway [graph](#) structures (e.g., those discussed in Chapter 5). These results can indicate whether the [SLIPT](#) methodology robustly detects known [synthetic lethal](#) partners (and how it compares to other [bioinformatics](#) strategies) or is suitable for wider [genomics](#) applications.

These simulation investigations continue to utilise the multivariate normal simulation procedure (as applied in Section 3.3) with further refinements. The [SLIPT](#)

methodology (and the χ^2 test) were applied across a range of parameters (including altering the quantiles for detecting [synthetic lethal](#) direction) and compared to correlation as a predictor of [synthetic lethality](#). These simulations included thousands of non synthetic lethal genes and correlations with the query gene (as performed in Section 3.3).

A refined simulation procedure was developed specifically to extend the methodology described in Section 3.2 to utilise pathway [graph](#) structures for the correlation structures of simulated datasets (as described in Section 3.4.2). This methodology can be applied to simulated correlation structures across simple [graph](#) structures to test specific network modules or use [pathway](#) structures based on biological pathways. Thus [graph](#) structure and simulation approaches were combined to test whether a gene locus in a pathway affects detection by SLIPT and whether SLIPT performance is affected by [pathway](#) structure. The simulation procedure based on [graph](#) structures was applied in a computational pipeline across many parameter combinations using high-performance computing resources (as discussed in Section 2.5.3) and the core simulation functions have been released as a software package for wider use to test [bioinformatics](#) and statistical methods on [graph](#) structures (as described in Section 3.5.3).

6.1 Synthetic Lethal Detection Methods

The SLIPT methodology (as it has been applied throughout Chapters 4 and 5) was compared for alternative computational approaches to detecting [synthetic lethality](#) in simulated [gene expression](#) data. As discussed in Section 3.3, this procedure enabled testing the ability of SLIPT to detect known [synthetic lethal](#) partner genes by sampling from a statistical model of [synthetic lethality](#). While comprehensive benchmarking has not been performed, several approaches to [synthetic lethal](#) detection are considered (e.g., Pearson correlation, the χ^2 test, and testing for bimodality) to evaluate the strengths of the SLIPT methodology, including modifications to the parameters of SLIPT. The following comparisons of simulations of computational detection of [synthetic lethality](#) with different statistical rationales were performed to show the strengths of SLIPT, evaluate whether it is appropriate for further application in [genomics](#) research, and identify limitations which may be addressed with further developments.

6.1.1 Performance of SLIPT and χ^2 across Quantiles

Simulated datasets with [synthetic lethal](#) partner genes were generated using the multivariate normal simulation procedure (as described in Section 3.2.2) with performance

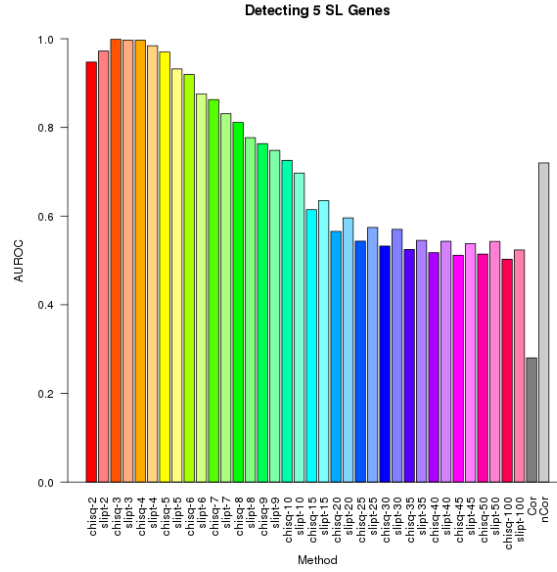
assessed using [area under receiver operating characteristic \(AUROC\)](#) analysis (as described in Section 2.3.5). [Synthetic lethal](#) detection was compared for modifications to the [SLIPT](#) methodology (as described in Section 3.1), namely that the quantiles used to define low and high [expression](#) were varied. Rather, than $1/3$ (as used throughout this thesis) the samples below the lowest $1/n$ quantile and above the highest $1/n$ quantile were used for [SLIPT](#) (and the χ^2 -test) to detect samples that exhibited low and high [expression](#) levels respectively. The quantiles tested ranged from two, splitting at the $1/2$ quantile (the median), to 100, using the lowest (1%) and highest (99%) percentiles.

This enabled testing of the threshold for low [expression](#) of genes which is most able to distinguish [synthetic lethal](#) genes, even with higher-order [synthetic lethal](#) interactions (as discussed in Section 3.2.1). Both [SLIPT](#) with the directional criteria for [synthetic lethality](#) and significance of the equivalent χ^2 test were performed for each quantile. Pearson correlation was also tested on simulated continuous [expression](#) data for [synthetic lethal](#) detection in simulated data, considering both positive and negative correlations separately as predictors of [synthetic lethality](#) for comparison with χ^2 based approaches, using discrete categories of gene function deriving from quantiles.

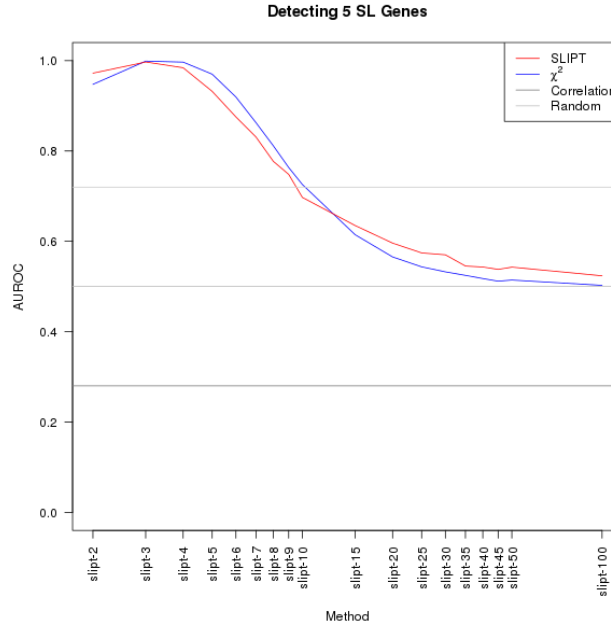
The results presented throughout this Section use the example of five [synthetic lethal](#) partners to illustrate the differences in performance between the standard [SLIPT](#) procedure (slipt-3) to n quantiles (slipt- n), the χ^2 -test on the same quantiles, and positive or negative correlation. However, similar results across different numbers of known [synthetic lethal](#) genes are shown in Appendix J. The [synthetic lethal](#) detection procedures were compared with 10,000 simulations of a small dataset of 100 genes and 1000 samples without correlation structure between genes (as performed in Section 3.3.2).

As shown in Figure 6.1, the $1/3$ -quantiles previously used have optimal performance and [SLIPT](#) has a comparable or higher performance than the χ^2 -test alone across quantiles. Pearson correlation was also tested as a predictor of [synthetic lethality](#) (i.e., whether highly positive or negative correlations with the query gene detected [synthetic lethal](#) partners). Positive correlation performed worse than random (with an [AUROC](#) lower than 0.5) as thus coexpression of genes was not predictive of [synthetic lethality](#) in simulated data. Conversely, negative correlation was predictive of [synthetic lethality](#), consistent with [synthetic lethal](#) gene activity being mutually exclusive. However, neither correlation approach performed as well as the optimal quantiles for the [SLIPT](#) procedure or the χ^2 -test.

These results are shown in both a bargraph and lineplot to show the individual results of each parameter, and to compare [SLIPT](#) with the χ^2 -test side-by-side across



(a) Barplot of χ^2 , SLIPT, and correlation.



(b) Lineplot of χ^2 , SLIPT, and correlation.

Figure 6.1: **Performance of χ^2 and SLIPT across quantiles.** Synthetic lethal detection (of 5 genes) with quantiles as on the axes. The barplot uses the same hues for each quantile (grey for correlation) and darker for χ^2 (and positive correlation). The line plot (with log-scale quantiles) is coloured according to the legend. SLIPT and χ^2 perform similarly, peaking at $1/3$ -quantiles and converging to random (0.5). Negative correlation had higher performance than positive correlation but not optimal quantiles for SLIPT or χ^2 .

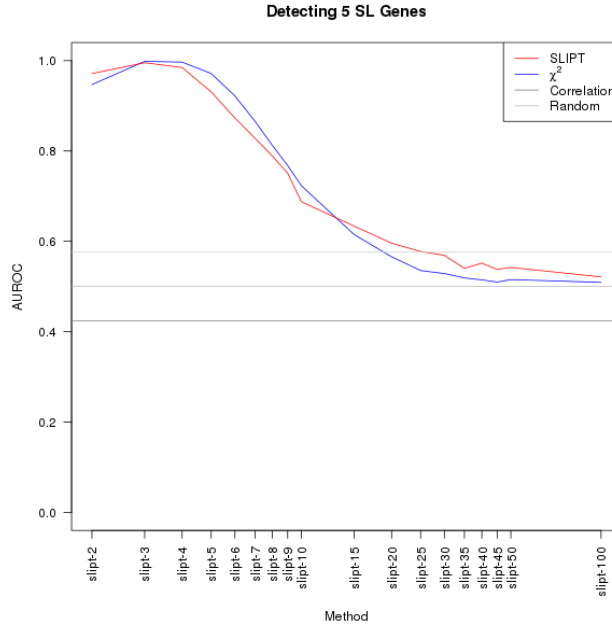


Figure 6.2: **Performance of χ^2 and SLIPT across quantiles with more genes.** **Synthetic lethal** detection (of 5 genes in 20,000) with quantiles as in axis labels. The line plot (with log-scale quantiles) is coloured according to the legend. As for simulations with fewer genes, **SLIPT** and χ^2 perform similarly, peaking at $1/3$ -quantiles and converging to random (0.5). Negative correlation had higher performance than positive correlation but not optimal quantiles for **SLIPT** or χ^2 .

quantiles. Similarly, these plots are given for detecting a range of known **synthetic lethal** partners in the simulations in Appendix Figures J.1 and J.2. These demonstrate that the findings shown for five **synthetic lethal** genes are robust across different numbers of underlying **synthetic lethal** genes.

The **synthetic lethal** detection procedures were also tested with 1000 simulations of a larger dataset of 20,000 genes and 1000 samples. While fewer simulations gives a less accurate **receiver operating characteristic (ROC)** result, this is sufficient to replicate the above findings with a feasible number of genes in a human **gene expression** dataset and assess the impact of a higher proportion of non synthetic lethal genes (potential false positives). Simulated datasets of this size were also used in Section 3.3.2 to test the specificity in a number of genes similar to that in experimental datasets for cancer **genomes**. As shown in Figure 6.2, the above findings were replicated in simulations of a larger dataset with 20,000 genes. These were also robustly replicated across varying numbers of underlying **synthetic lethal** genes (as shown in Appendix Figure J.3).

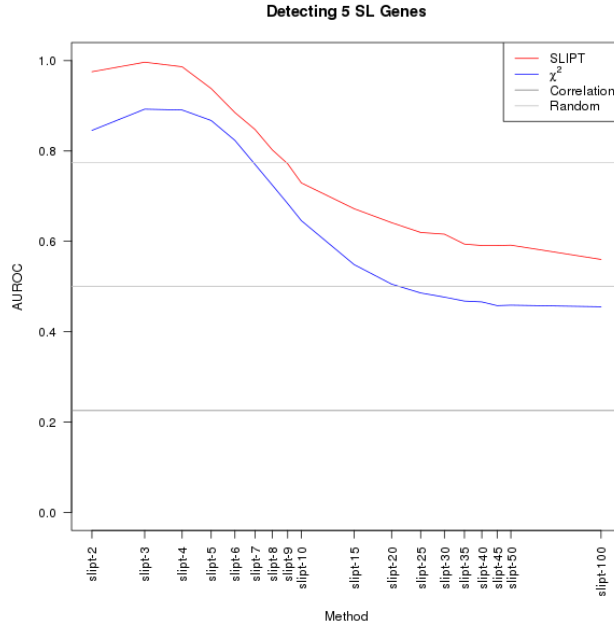


Figure 6.3: **Performance of χ^2 and SLIPT across quantiles with query correlation.** Synthetic lethal detection (of 5 genes in 100 including 5 query correlated) with quantiles as in axis labels. The line plot (with log-scale quantiles) is coloured according to the legend. SLIPT performs consistently higher than χ^2 due to higher specificity. Negative correlation performed modestly.

6.1.1.1 Correlated Query Genes affects Specificity

As discussed in Section 3.3.2.2, positively correlated genes (with the query gene) have an impact on the performance of synthetic lethal detection. SLIPT was able to distinguish these correlated genes from synthetic lethal partners and hence is likely to have a higher specificity in datasets which include positively correlated genes with the query gene (as expected in gene expression data). The synthetic lethal detection procedures were compared with 10,000 simulations of a small dataset of 100 genes (with 5 correlated with the query gene) and 1000 samples otherwise without correlation structure between genes. As shown in Figure 6.3, this specificity is reflected in the increased AUROC performance values for SLIPT (in contrast to Figure 6.1). This specificity can be attributed to the directional criteria (as described in Section 3.1) since the χ^2 -test alone performs comparatively poorly with positively correlated genes.

The synthetic lethal detection procedures were also compared with 1000 simulations of a larger dataset of 20,000 genes (with 1000 correlated with the query gene) and 1000 samples otherwise without correlation structure between genes. This simulation

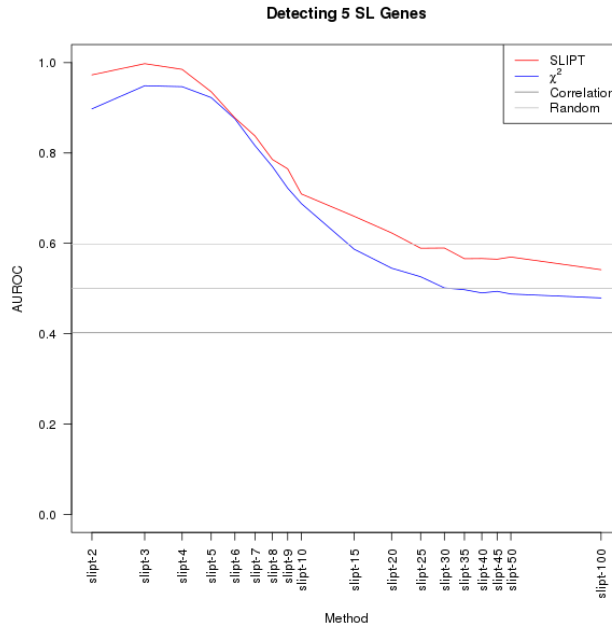


Figure 6.4: **Performance of χ^2 and SLIPT across quantiles with query correlation and more genes.** Synthetic lethal detection (of 5 genes in 20,000 including 1000 query correlated) with quantiles as in axis labels. The line plot (with log-scale quantiles) is coloured according to the legend. SLIPT performs consistently higher than χ^2 due to higher specificity. Negative correlation performed modestly.

increases the number of genes (and proportion of negative genes) to those comparable with a human [gene expression](#) dataset while maintaining a comparable 5% of positively correlated genes. As shown in Figure 6.4, SLIPT still outperforms χ^2 or negative correlation and is optimal at the $1/3$ -quantile. The difference between SLIPT and χ^2 was less pronounced in a larger dataset with many weakly correlated genes. The greater specificity of SLIPT than the χ^2 -test to distinguish positively correlated non synthetic lethal genes is not as evident with a large number of negative genes (as potential false positives). However, specificity is an important consideration in large-scale [genomics](#) analysis where there are potentially many false positives.

Nevertheless, SLIPT with $1/3$ -quantiles (as performed throughout Chapters 4 and 5), had higher performance than when other quantile thresholds were used, particularly when positive correlations were present (replicating the Section 3.3.2.2). These findings hold across different numbers of underlying [synthetic lethal](#) genes (as shown in Figures J.5 and J.6).

Together these results support the use of [SLIPT](#), particularly the use of quantiles as thresholds for gene function and specific use of $1/3$ -quantiles which perform well compared to other quantiles. A particular concern in the design of [SLIPT](#) for [expression](#) data whether the sample sizes are sufficient when the data are divided into quantiles. The [SLIPT](#) methodology further performed better for $1/3$ -quantiles (and other moderate values) than χ^2 or correlation as a predictor of [synthetic lethality](#). These results were irrespective of sample size or p-value threshold since the results replicated across sample sizes and the [AUROC](#) values were independent significance thresholds. Using a moderate number of quantiles for [SLIPT](#) ensures that there are a sufficient number of samples expected below and above them so that deviations from these are statistically detectable. These quantiles were also optimal for the χ^2 test which uses the same expected values as the [SLIPT](#) directional conditions.

6.1.2 Alternative Synthetic Lethal Detection Strategies

The [SLIPT](#) approach (and χ^2) to detect [synthetic lethality](#) from binning [expression](#) to estimate gene function also outperformed correlations, which use continuous data directly. Correlation performing poorly as a [synthetic lethal](#) detection strategy is consistent with there not necessarily being a relationship between [synthetic lethal](#) partners, which can be in distinct biological pathways, expressed at different times or in different cell types. Nevertheless, correlation is among the alternative detection methods considered in further detail.

The [BImodal Subsetting ExPression](#) (BiSEp) R package ([Wappett, 2014](#)) for using bimodality to detect [synthetic lethality](#) ([Wappett et al., 2016](#)) was also considered, along with a linear regression approach. These statistical methods span a range of computational approaches to detecting [synthetic lethality](#) and serve to compare alternatives to [SLIPT](#), supporting its design and application. However, these comparisons are able to provide supporting data from statistical modelling and simulations for the viability of the [SLIPT](#) methodology for [synthetic lethal](#) discovery in cancer (as demonstrated in Chapter 4) and further applications.

6.1.2.1 Correlation for Synthetic Lethal Detection

As expected, negative (Pearson) correlation performed better than positive correlation at detecting [synthetic lethality](#) (shown in Section 6.1.1). However, neither correlation approach performed as well as [SLIPT](#) or the χ^2 test as a predictor of [synthetic lethal](#) gene partners. It is notable that negative correlation still often performed considerably better than random chance.

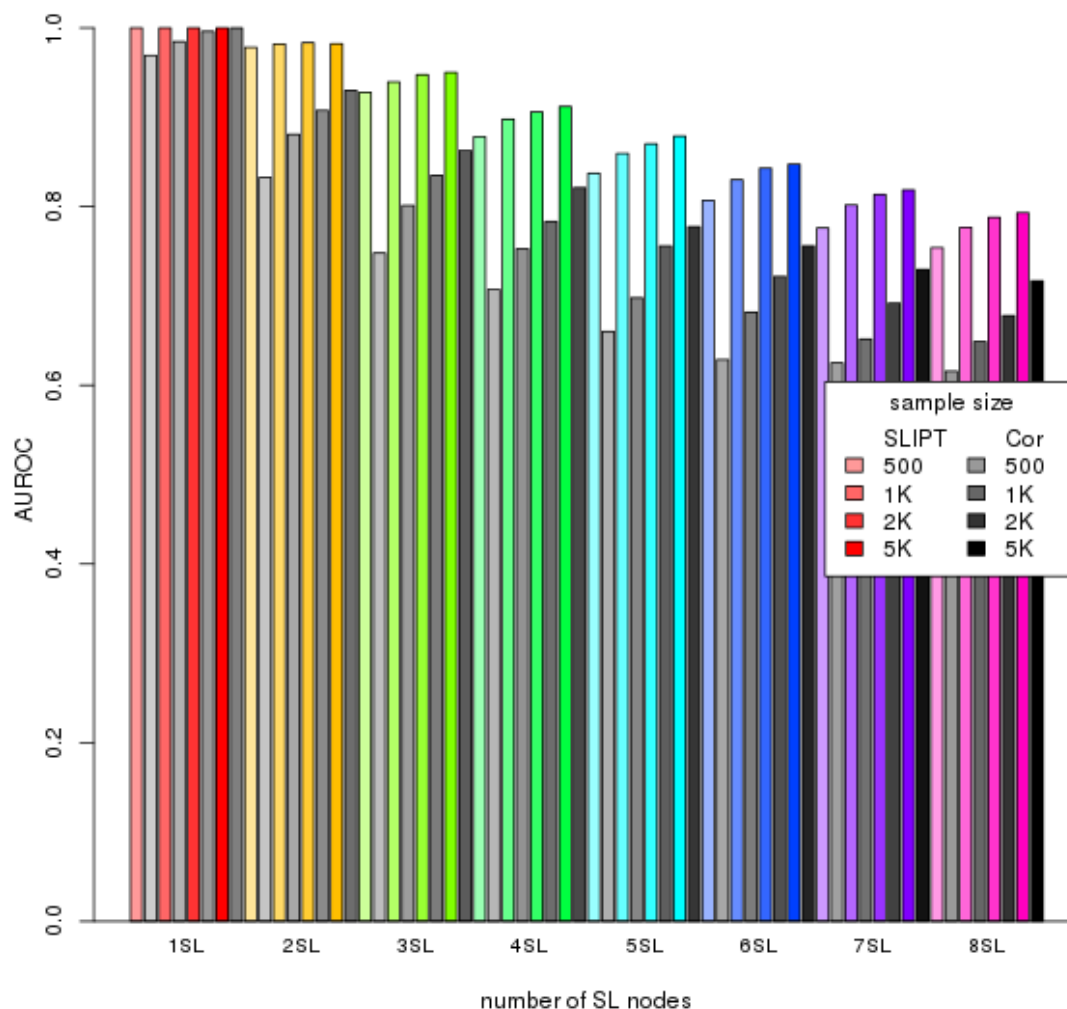


Figure 6.5: **Performance of negative correlation and SLIPT.** Synthetic lethal detection with SLIPT was compared to negative (Pearson) correlation across parameters. SLIPT consistently outperformed correlation. Both approaches had lower performance for more synthetic lethal partners and for lower sample sizes. 10,000 simulations were performed with correlation structure.

Negative correlation was compared directly to the SLIPT methodology (as described in Section 3.1) across numbers of known synthetic lethal partners and sample size (ranging from 500 to 5000). This comparison used 1000 simulations of a dataset with 20,000 genes and synthetic lethal genes from within a network (sampled as in Section 3.4.2) with a 0.8 correlation between adjacent genes. In a direct comparison of SLIPT and

negative correlation (shown in Figure 6.5), SLIPT consistently has higher performance in simulated data across parameter values and (inverse) correlation-based approaches perform modestly in comparison. Thus using thresholds to categorise expression data (as performed by SLIPT and χ^2) does not compromise the performance of these methods by losing continuous data that would be used for calculating correlations.

Both SLIPT and correlation had poorer performance with increasing numbers of the synthetic lethal genes to detect, while they had higher performance in higher sample sizes, as expected (as previously observed for SLIPT in Section 3.3). Thus the issue with detection of greater numbers of synthetic lethal genes is not specific to SLIPT but occurs across computational methods of synthetic lethal discovery in (simulated) expression data and likely stems from cryptic higher-order synthetic lethal interactions (as conservatively assumed in Section 3.2.1).

6.1.2.2 Testing for Bimodality with BiSEp

Extensive attempts were also made to compare SLIPT to the BiSEp methodology (Wappett *et al.*, 2016), a statistical approach to identify synthetic lethal gene pairs from mutually exclusive relationships using bimodal distributions. This synthetic lethal detection methodology is also designed for expression analysis in cancer and is readily available as an (open-source) R package (Wappett, 2014), a practice which facilitates adoption and testing of the methodology on the same datasets and simulations procedures as previously used for SLIPT.

The BiSEp package is designed for global testing of all potential gene pairs in the genomes for synthetic lethality rather than focusing on the search space of potential partners of the query gene. This approach was unable to detect synthetic lethal gene pairs in the The Cancer Genome Atlas (TCGA) breast cancer expression dataset (Koboldt *et al.*, 2012). However, this may be due to stringent thresholds under the multiple testing of millions of potential gene pairs.

For a direct comparison with the query-based SLIPT approach, the source code of the BiSEp R functions was modified to test solely for the partners of a specific gene. This approach was still unable to detect synthetic lethal partners of *CDH1* in TCGA breast cancer expression data (Koboldt *et al.*, 2012), even with the detection thresholds for bimodality and significance greatly relaxed from those which the package defaults to.

To circumvent multiple testing issues, BiSEp only tests gene pairs for synthetic lethality between genes with a detectable bimodal distribution. However, even with relaxed thresholds, bimodal distributions were not detectable in the normalised TCGA data

(Koboldt *et al.*, 2012). Such normalisation Ritchie *et al.* (2015) is standard practice for expression datasets generated from microarrays or RNA-Seq and therefore BiSEp may not be appropriate to apply to this data. However, it is noted that BiSEp may also use other data types such as DNA copy number or cell line data for which it may be more applicable (Wappett *et al.*, 2016).

Nevertheless, attempts were made to test BiSEp on simulated datasets with underlying synthetic lethal genes (using the procedures described in Sections 3.2.2 and 3.4.2). However, BiSEp was also unable to detect genes with bimodal distributions of genes (and thus unable to detect synthetic lethality) in a limited number of computationally intensive simulations. Therefore investigations on a wider range of parameters were not performed.

6.2 Simulations with Graph Structures

The simulations of synthetic lethality performed in Section 3.3 included correlated blocks of genes as a rudimentary representation of pathway structure and co-regulated genes. The simulation procedure was enhanced here to account for more complex pathway structures by sampling from multivariate normal distributions with correlation structure derived from graph structures (as described in Section 3.4.2). This approach enabled the simulation of synthetic lethal pathways with known correlation structure and synthetic lethal partners (of a gene not in the pathway). Using this procedure, the performance of SLIPT was evaluated under simple controlled correlation structures and complex correlations, such as those derived from biological networks (e.g., those described in Chapter 5). The SLIPT methodology was tested in artificially constructed networks to evaluate the effect of pathway structure on synthetic lethal detection. These included large biologically feasible pathways to ensure that the SLIPT methodology is robust under complex correlation structures and applicable to such complex genomics data.

These simulations combine the approach of prior simulation analyses (in Sections 3.3 and 6.1) with the graph structures for biological pathways (as used in Chapter 5). This enabled testing whether subtle or large differences in pathway structure affect synthetic lethal detection, whether inhibiting relationships (or inverse correlations) between genes affect synthetic lethal detection, and whether synthetic lethal detection varies by which gene is synthetic lethal and which genes are closely linked within the pathway structure. In addition, large numbers of synthetic lethal genes and biologically feasible numbers of genes (with many non-synthetic lethal genes) were tested to replic-

ate the findings of Sections 3.3 and 6.1 in correlated structures derived from pathway graphs, including examples of biological pathways from Reactome (Croft *et al.*, 2014).

Simple and more complex constructed graph structures were used to demonstrate the impact of pathway structure of the performance of SLIPT for synthetic lethal detection in simulations. In addition, more complex constructed graph structures were compared to the phosphoinositide 3-kinase (PI3K) and $G_{\alpha i}$ signalling pathways derived from Reactome which were used for simulation of pathway structures of biological complexity (as shown in Figure 5.1 and Appendix Figure 5.4).

6.2.1 Performance over Graph Structures

6.2.1.1 Simple Graph Structures

Simple pathway modules were used to test the effect of pathway structure on the performance of detecting synthetic lethal partners within graph structures. For an initial comparison, the graph structures (shown by Figure 6.6) were used where a gene has one upstream regulator and two downstream (Figure 6.6b) or a gene has two upstream regulators and one downstream gene (Figure 6.6b). SLIPT had a high performance in these simulations, detecting randomly selected synthetic lethal partners in both of these small simple networks (shown in Figure 6.7 and Appendix Figure K.1).

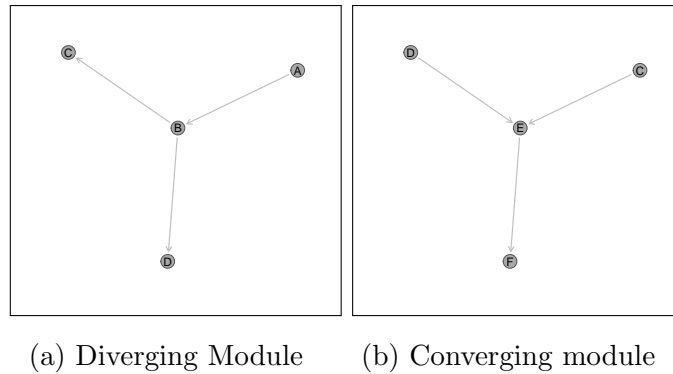
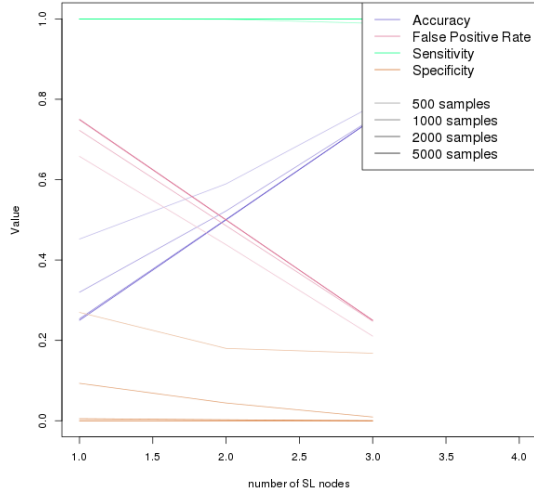
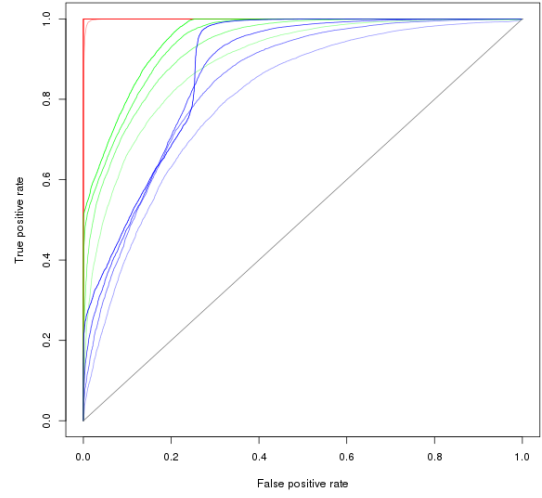


Figure 6.6: **Simple graph structures.** These simple graph structures used to demonstrate the simulation procedure. These are examples of a pathway diverging or converging respectively which enabled testing the importance of direction in pathway structures. These are used with both activating and inhibiting relationships as shown.

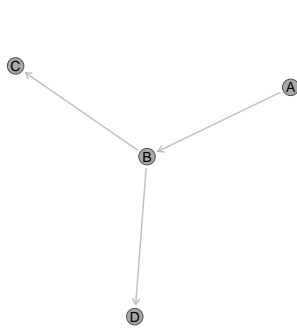
As previously observed (in Section 3.3), performance declined with higher numbers of synthetic lethal genes and lower sample sizes. However, the sensitivity of SLIPT was high with conventional p-value thresholds (adjusted by False discovery rate (FDR)).



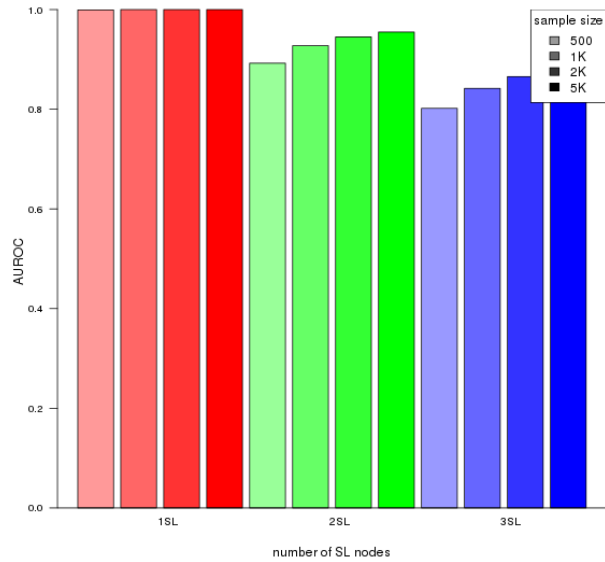
(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.7: **Performance of simulations on a simple graph.** Simulation of **synthetic lethality** was performed by sampling from a multivariate normal distribution generated from a diverging **graph** structure. Performance of **SLIPT** declines for more synthetic partners but this is mitigated by increased sample sizes (in darker colours). This manifests as a decline in specificity and the false positive rate. For each parameter value, 10,000 simulations were used. Colours of the **ROC** curves in Figure 6.7b correspond to the parameters in Figure 6.7d.

Synthetic lethal partners are often distinguishable for non synthetic lethal genes, even in simple highly correlated networks. The small number of genes and their high correlation has an impact on the **ROC** curves for higher numbers of **synthetic lethal** partners which are skewed compared to those observed previously. Specificity cannot be tested if all potential partner genes are **synthetic lethal**, which limits the number of **synthetic lethal** genes that can be tested.

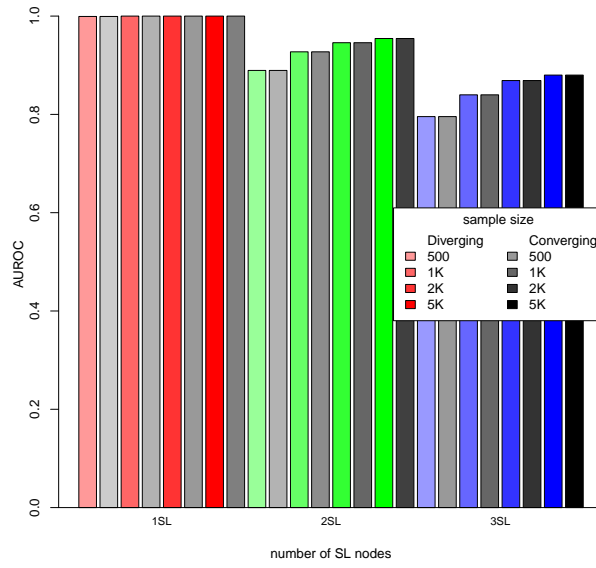
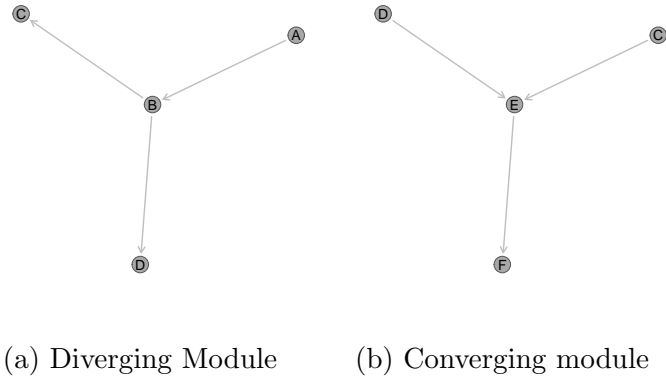
These results were consistent between the pathway modules of diverging (as shown in Figure 6.8a) and converging signals (as shown in Figure 6.8b). The **AUROC** performance and underlying curves were strikingly similar between these **graph** structures (as shown in Figure 6.7 and Appendix Figure K.1). Thus the performance of **SLIPT** was not perturbed by **pathway** structure, specifically the direction of pathway relationships since these **graph** structures also demonstrate pathways in opposite direction. In a direct comparison (shown in Figure 6.8c), the performance of simulations did not differ across parameter values in these simple **graphs** and therefore **SLIPT** is robust to pathway direction.

6.2.1.2 Constructed Graph Structures

A more complex **graph** structure was used to examine the performance of detecting **synthetic lethal** partners with **SLIPT** in simulated **expression** data with pathway correlation structures. For a simple chain of genes representing a very linear pathway (shown in Figure 6.9), the above findings were generally replicated. **SLIPT** had high performance across parameter values in small networks but was still lower for higher numbers of **synthetic lethal** genes and lower sample sizes.

When detecting **synthetic lethal** genes with **SLIPT** using adjusted (**FDR**) p-value thresholds, the performance differences were primarily due to changes in specificity, as the small numbers of **synthetic lethal** genes still had highly significant p-values. Despite lower specificity and performance in **ROC** curves, the accuracy increased and false positive rate decreased desirably with higher numbers of **synthetic lethal** genes due to the high sensitivity and the high proportion of **synthetic lethal** genes detected. Therefore the thresholds imposed by adjusted p-values appear to be appropriate for detecting **synthetic lethal** partners, even in strongly correlated pathways, at least in these small-scale test cases.

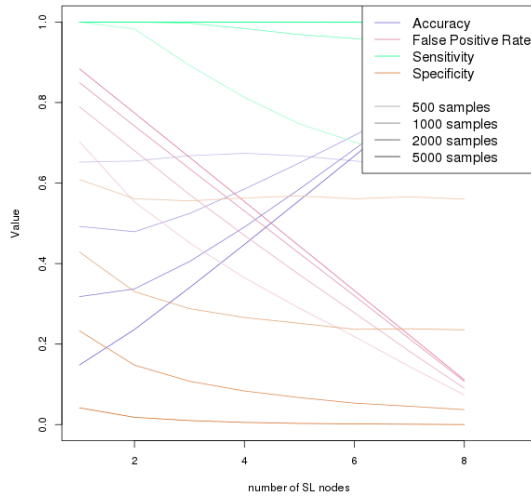
An artifact of these small test cases led to the skewed **ROC** curves (as discussed in Section 6.2.1.1), which may be the result of the low number of non-synthetic lethal genes to identify as true negatives, affecting the accuracy of specificity. This issue does not occur in larger, more complex **graph** structures, even with modest total numbers



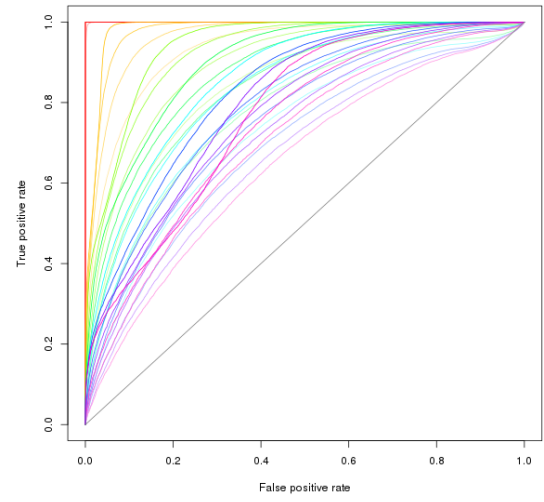
(c) Performance between Graph Structures

Figure 6.8: **Performance of simulations is similar in simple graphs.** The AUROC values for simulations of multivariate normal distributions based on each graph structure yielded indistinguishable performance across parameter values in 10,000 simulations.

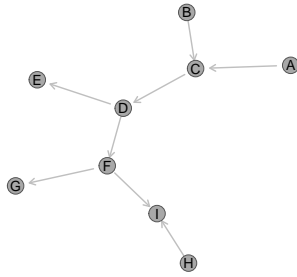
of genes and high correlations (as shown in Section 6.3). This issue is unlikely to occur in large expression datasets with many non synthetic lethal genes, as shown previously (in Section 3.3 and 6.2.1.1) with graph structures in larger datasets (in Section 6.2.4).



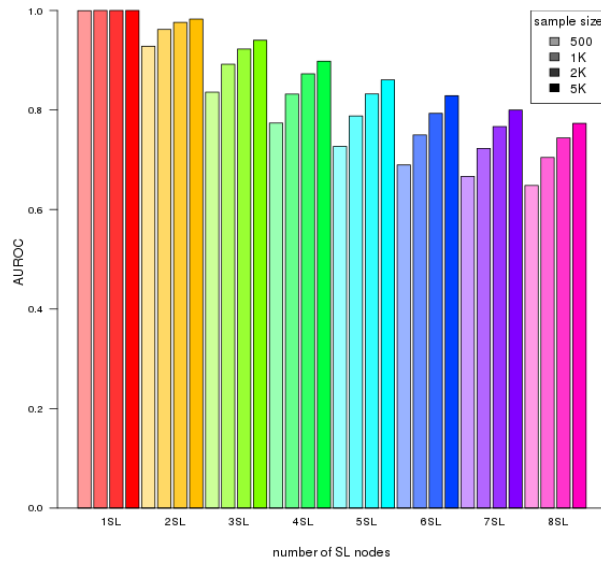
(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.9: **Performance of simulations on a pathway.** Simulation of [synthetic lethality](#) was performed by sampling from a multivariate normal distribution generated from a [pathway](#) structure. Performance of [SLIPT](#) declines for more synthetic partners and lower sample sizes (in darker colours). For each parameter value, 10,000 simulations were used. Colours of the [ROC](#) curves in Figure 6.9b correspond to the parameters in Figure 6.9d.

6.2.2 Performance with Inhibitions

Simulations of [synthetic lethality](#) in [expression](#) data were also performed with correlation structures derived from [graphs](#) containing inhibiting relationships (as are commonplace in biological pathways) which produce negative correlations. As shown in Figure 6.10, these are not an issue for detection by [SLIPT](#). Rather, the [SLIPT](#) procedure performs well on simple graph modules with highly negative correlations. With [synthetic lethal](#) detection based on p-value (adjusted by [FDR](#)), there was higher specificity, higher accuracy, and lower false positive rate in an inhibitory graph than the same graph with activating relationships (as shown by Figure 6.7).

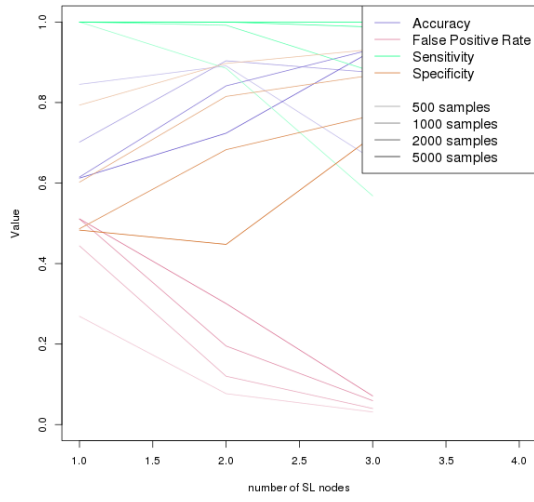
The [ROC](#) curves for an inhibiting graph also showed consistently high specificity, irrespective of detection threshold, with only the upper extreme of the curve exhibiting a skew below random performance (as shown in Figure 6.10). Nevertheless, the [AUROC](#) values show a high performance across parameter values, particularly avoiding issues with higher numbers of [synthetic lethal](#) partners (as observed in Section 6.2.1.1). However, performance was marginally lower for higher numbers of [synthetic lethal](#) genes to detect and lower sample sizes, consistent with previously observations.

Negatively correlated simulated datasets are also unperturbed by minor differences in [graph](#) structure, such as changing in the direction of the graph module. As observed for activating relationships in these graph modules, the performance was highly concordant between the graph modules (shown by similar results in Figures 6.10 and K.2).

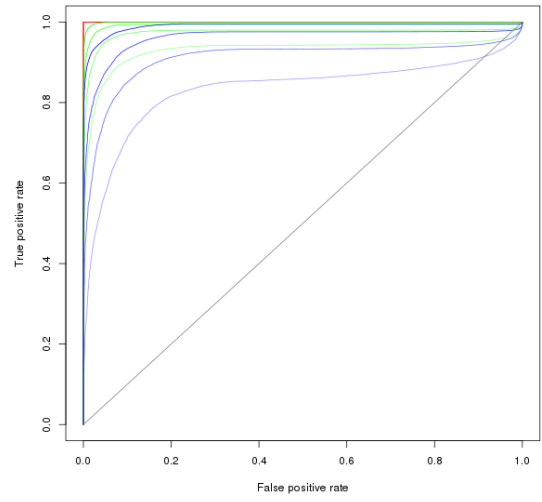
Detection of [synthetic lethality](#) by [SLIPT](#) in simulated data with inhibiting relationships outperforms simulations with activating relationships in the same [graph](#) structure (as shown in Figure 6.11). Thus [SLIPT](#) was robust in [gene expression](#) datasets with inverse correlations and performed well in them, at least in simple test cases. This is important because such relationships occur frequently in biological pathways and therefore the findings inferred from [graph](#) structures without inhibiting relationships are a conservative estimate.

The [SLIPT](#) methodology likely performs better in biological pathways (which contain negative correlations) than the [graph](#) structures discussed previously (in Section 6.2.1). This is likely since negative correlations lead to [synthetic lethal](#) partners and inversely correlated genes which are positively correlated with the query gene. As previously shown, the [SLIPT](#) methodology performs well with specificity against positively correlated query genes (in Sections 3.3.2.2 and 6.1.2.1).

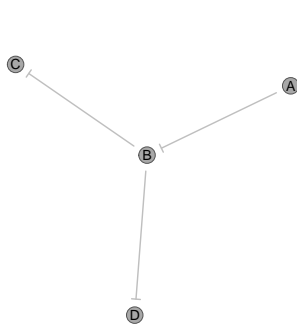
Similarly, more complex [graph](#) structures with entirely inhibiting relationships (negative correlations) also perform desirably on p-value thresholds (adjusted by [FDR](#)) and



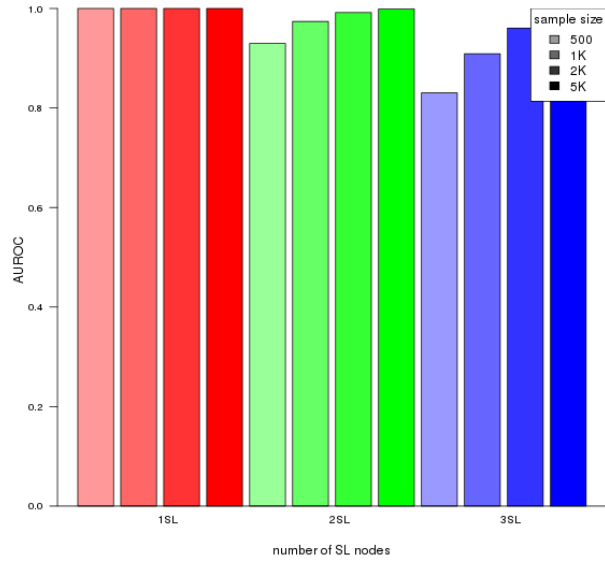
(a) Statistical evaluation



(b) Receiver operating characteristic

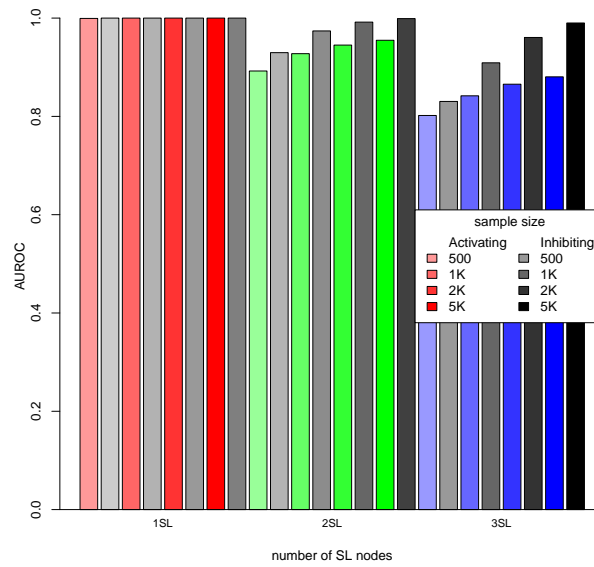
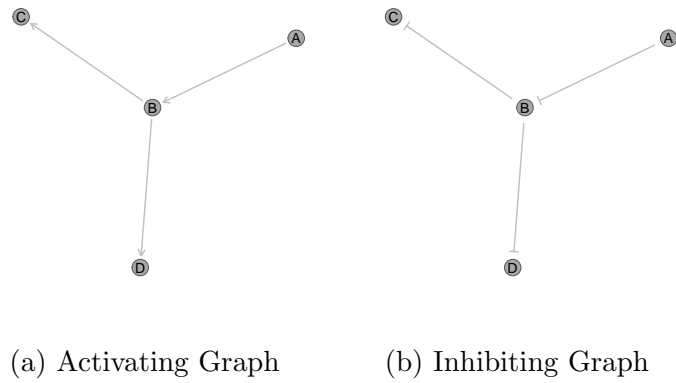


(c) Graph Structure



(d) Statistical performance

Figure 6.10: Performance of simulations on a simple graph with inhibition. Simulation of **synthetic lethality** was performed by sampling from a multivariate normal distribution generated from an inhibiting graph. Performance of **SLIPT** declined for more synthetic partners and lower sample sizes. For each parameter value, 10,000 simulations were used. Colours of the **ROC** curves in Figure 6.10b correspond to the parameters in Figure 6.10d.

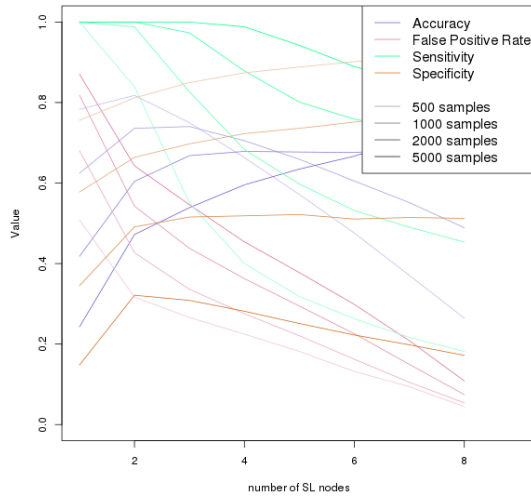


(c) Performance between Graph Structures

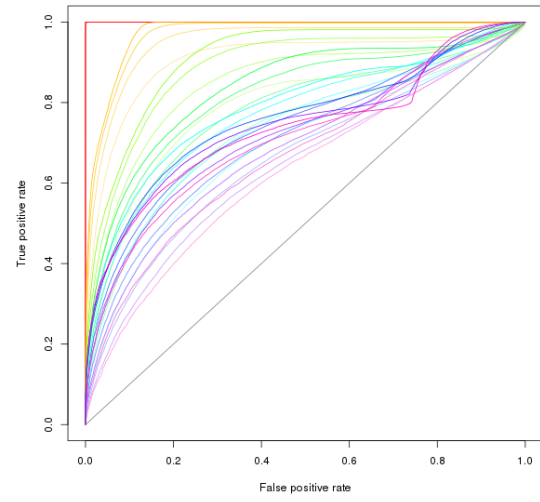
Figure 6.11: **Performance is higher on a simple inhibiting graph.** The AUROC values for simulations of multivariate normal distributions based on inhibitions in the [graph](#) structure yielded consistently higher performance across parameter values in 10,000 simulations.

have high performance across increasing numbers of [synthetic lethal](#) genes, particularly for sufficiently high sample sizes (as shown by Appendix Figure K.3). However, this is not necessarily the case for [graph](#) structures with a combination of activating and inhibiting relationships (i.e., containing positive and negative correlations). As shown by Appendix Figure K.4, such a mixed [network](#) structure does not necessarily have high performance across parameters as observed for purely inhibiting networks.

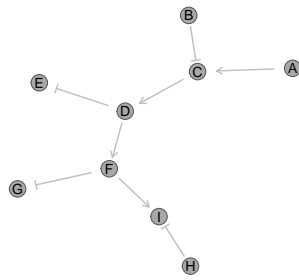
These still appear to have desirably high sensitivity, high accuracy, and low false positive rate for detecting more [synthetic lethal](#) genes, despite poor specificity. The



(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.12: **Performance of simulations on a constructed graph with inhibition.** Simulation of [synthetic lethality](#) was performed by sampling from a multivariate normal distribution generated from [pathway](#) structure with a combination of inhibitions. Performance of [SLIPT](#) declines for more synthetic partners and lower sample sizes. For each parameter value, 10,000 simulations were used.

ROC curves were particularly skewed for high proportions of the network being **synthetic lethal** and may stem from low numbers of true negative genes to detect (as discussed in Section 6.2.1.1). In a direct comparison of performance (shown in Figure 6.13), the purely inhibiting graph had consistently higher performance than the activating one, as observed for simpler **graphs** (as shown in Figure 6.11).

In contrast, the combination of activating and inhibiting relationships had slightly lower performance across parameters compared to the same **graph** structure with activating relationships. Therefore correlation structure can impact on the performance of **SLIPT** in a graph network, in either direction, specifically the addition of negative correlations. However, this may be an artifact of the simulation procedure as **synthetic lethal** genes from the correlation structure were randomly selected (without regard to their relationships), with the query gene added to ensure that conditions for **synthetic lethal** relationships were met.

This system for simulating inhibitory pathways is not ideal since it can lead to **synthetic lethal** gene combinations, by randomly selecting them, which are unlikely to occur in biological pathways. These randomly selected **synthetic lethal** genes may account for the detection results being suboptimal (i.e., difficult to detect **synthetic lethal** partners) compared to previous investigations. It is expected that inversely correlated synthetic partner genes will be highly expressed in a mutually exclusive manner such that at least one of them will be compensating for loss of the query gene in most samples, leading to a weak **synthetic lethal** signature in **expression** data in this case. Furthermore, this case may not be representative of empirical biological data with **synthetic lethal** partners of **tumour suppressor** genes which are commonly inversely correlated to the query gene (to some extent) and therefore it is unlikely that they are strongly negative correlated with each other, unless they are **synthetic lethal** partners of each other as well. It is plausible that many **synthetic lethal** partner genes will serve to separately compensate for the loss of query gene function and be positively correlated with each other. Nonetheless, these simulations demonstrate that correlation structure (particularly negative correlations) have an impact on the detection of **synthetic lethality**. However, **SLIPT** was still able to perform well across **graphs** with different activating and inhibiting relationships, and the perturbations in performance were marginal, particularly those reducing performance compared to an activating network.

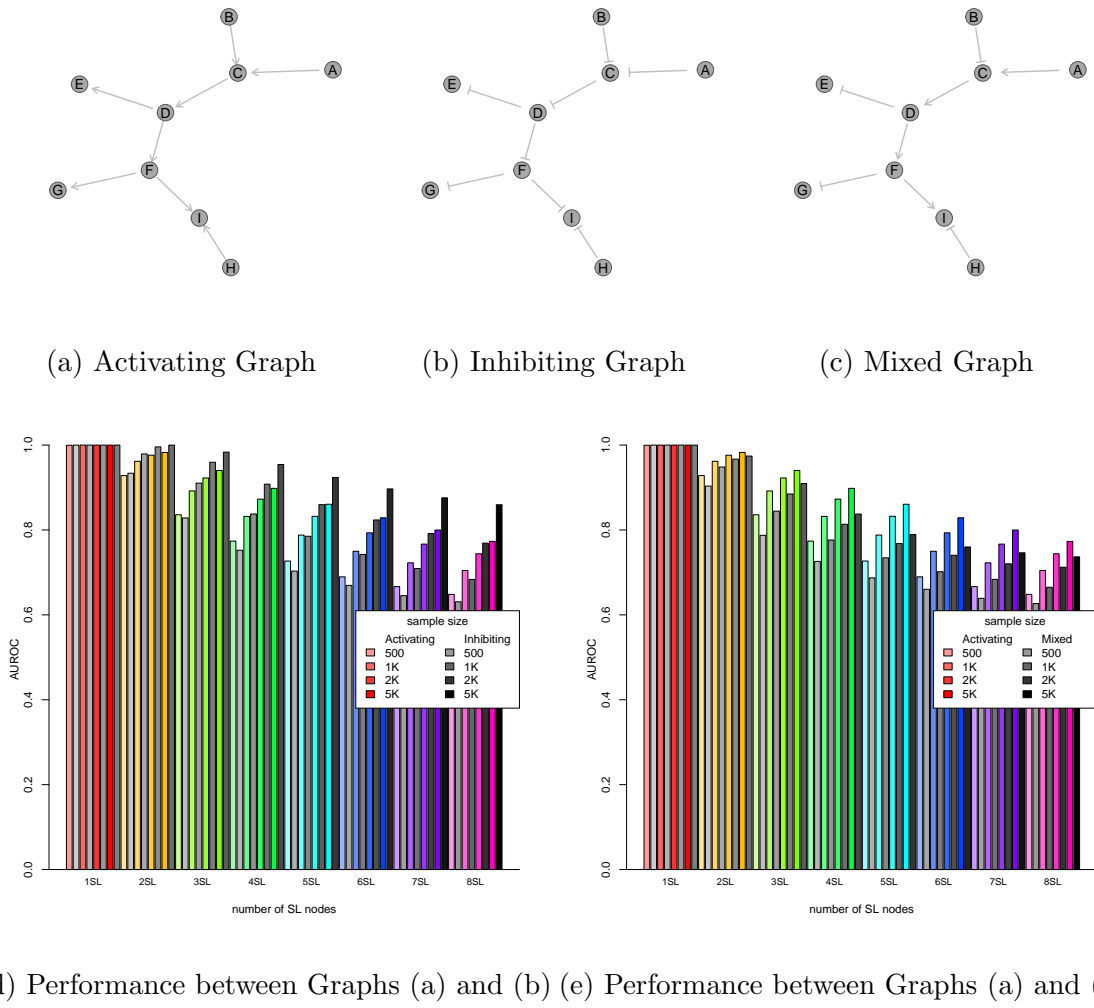


Figure 6.13: **Performance is affected by inhibition in graphs.** The AUROC values for simulations of multivariate normal distributions based on graph structure containing only inhibitions in the graph structure yielded consistently higher performance across parameter values in 10,000 simulations. A combination of activating and inhibiting relationships had lower performance but was more similar to the activating graph.

6.2.3 Synthetic Lethality across Graph Structures

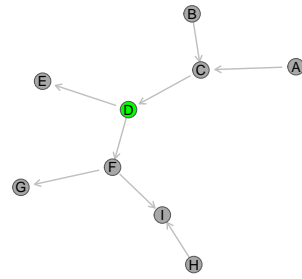
While synthetic lethal genes are distinguishable in principle from those highly positively correlated with them (as shown by ROC analysis), they are not necessarily distinguished as reflected by low specificity and high false positive rates in poorly performing simulations throughout this Section. The negative correlations are not subject to the same issue, they sometimes perturb the correlation structure between synthetic lethal partner genes making it difficult to detect many of them. Thus far, synthetic

lethal genes have been selected randomly, which is a limited approach. To examine the impact of pathway relationships in more more detail, specific genes will be selected to be synthetic lethal within a network. Replicate simulations were performed for synthetic lethal detection with a fixed synthetic lethal gene, in contrast to previous investigations (randomly selecting synthetic lethal genes). This investigation was performed to demonstrate the impact of these genes being synthetic lethal in the detection of neighbouring genes in the pathway network, under graph structure activating and inhibiting relationships.

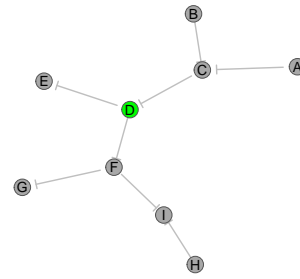
For instance, detection of a synthetic lethal gene in an activating graph structure (as shown in Figure 6.14a) is straightforward: the χ^2 values across simulations are clearly distinguishable from non synthetic lethal genes (shown in Figure 6.14c). A small number of simulations were performed for each gene being designated as synthetic lethal. In each case (of each gene being the synthetic lethal partner), the synthetic lethal gene was detectable with the highest χ^2 value, being distinguishable amongst 20,000 genes including the highly correlated graph network (as shown in Appendix Figure K.5).

This is consistent with previous observations that SLIPT performed optimally for a single synthetic lethal partner in this network (as shown in Figure 6.9). Despite optimal performance in a ROC curve, irrespective of detection threshold, many of the highly correlated genes would be detected as false positives using a conventional p-value threshold (even if adjusted by FDR) from a χ^2 test with 4 degrees of freedom as performed by SLIPT (as described in Section 3.1). In particular, the genes that were adjacent to the synthetic lethal gene “D” within the graph structure exhibited high test statistics across simulations which would often be reported as false positives (as shown in Figure 6.14c). This is not specific to example of gene “D”, with the neighbouring genes exhibiting higher χ^2 test statistics for each gene in the network when it is designated as the synthetic lethal partner (as shown in Appendix Figure K.5).

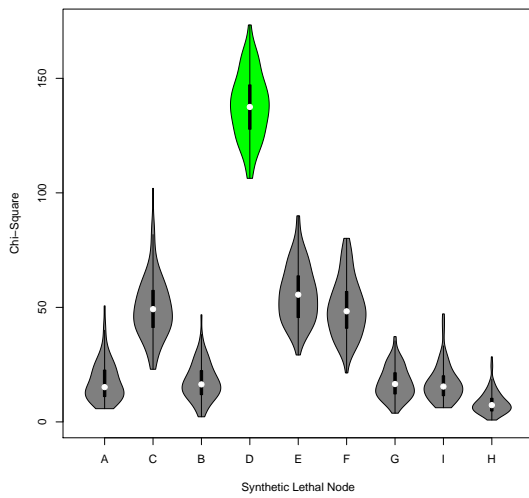
Thus the synthetic lethal signal propagated from the true synthetic lethal gene throughout the network. As such, the genes nearer to the true synthetic lethal gene (more highly correlated) had higher test statistics and were more likely to be detected by SLIPT as false positives. This tendency for adjacent genes to be detected as synthetic lethal false positives is consistent with the synthetic lethal pathways being more concordant between SLIPT in TCGA data (Koboldt *et al.*, 2012) and the siRNA screen (Telford *et al.*, 2015) than individual gene results (in Chapter 4). False positive genes are therefore still more likely to be involved in a synthetic lethal pathway by being



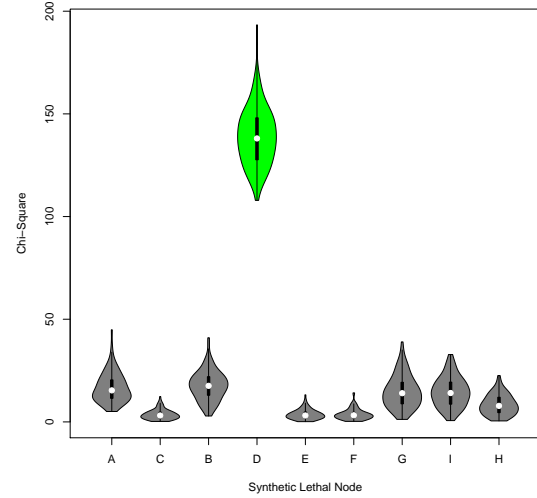
(a) Activating Graph



(b) Inhibiting Graph



(c) χ^2 distribution for Activating Graph



(d) χ^2 distribution for Inhibiting Graph

Figure 6.14: **Detection of synthetic lethality within a graph structure.** The gene “D” was designated to be **synthetic lethal** and the χ^2 value from **SLIPT** was computed for each gene across each **graph** structure. The χ^2 values were computed in 100 simulations of datasets of 20,000 genes including the **graph** structure and 1000 samples. Adjacent genes exhibited lower χ^2 values with inhibiting relationships.

correlated with a true **synthetic lethal** gene and **synthetic lethal** pathways are likely to have many genes detected by **SLIPT** giving a consensus of evidence, supporting the pathway over-representation approach in particular which may account for how it differs from pathway **metagenes**. Furthermore, **SLIPT** is still viable to detect true **synthetic lethal** partners or prioritise those most likely to be experimentally validated since those with the strongest support (i.e, higher χ^2 values and more significant p-values) are more likely to be the underlying **synthetic lethal** gene.

In contrast to an activating graph (Figure 6.14a), the immediately adjacent genes in an inhibiting graph (Figure 6.14b) had neither an elevated χ^2 test statistics indicating **synthetic lethality** nor a significant inverse effect (as shown in Figure 6.14d). Similar simulations were performed on a **graph** structure with inhibiting relationships within a dataset of 20,000 genes. The adjacent genes to the **synthetic lethal** gene “D” did not have elevated χ^2 values and therefore true **synthetic lethal** partners were highly distinguishable from non synthetic lethal genes with inhibiting relationships. This was not specific to “D” and was shown across any gene in the **graph** structure if it were designated to be the **synthetic lethal** partner of the query gene (shown in Appendix Figure K.6). This is consistent with the detection of many genes involved in kinase signalling, gene regulation, and other known cancer pathways (in Chapter 4) which frequently have inhibitory steps. These results support **SLIPT** as an appropriate approach to distinguish **synthetic lethal** partners in biological pathways, including those relevant to cancer growth and inhibition.

It should be noted, however, that the 2nd degree neighbours of the **synthetic lethal** gene still exhibited moderate χ^2 values (and are moderately correlated with the **synthetic lethal** gene). It is still possible for these to be detected as false positives as previously described for an activating **graph** structure although the presence of inhibitory relationships (and negative correlations) further increases the differences in test statistics for correlated genes and underlying **synthetic lethal** partners as shown by the extreme example (as shown in Appendix Figure K.6).

These findings are consistent with simulations in a graph containing a combination of activating and inhibiting relationships which exhibits a either of these χ^2 profiles depending on which gene is **synthetic lethal** and the relationships to adjacent genes (as shown in Appendix Figure K.7). Note that in this case, the **synthetic lethal** gene is distinguishable and inhibitory relationships within this **graph** structure make it easier to detect underlying **synthetic lethal** genes with **SLIPT** by a more highly significant χ^2 test. This contrasts with randomly selecting multiple **synthetic lethal** genes (as shown in Figure 6.13) where the performance of **SLIPT** was impeded by the inhibitory relationships between **synthetic lethal** partners in this **graph** structure. Therefore the random **synthetic lethal** genes selected previously with negative correlations between them which had poor performance are likely to have created an artifact in the simulation results as they are biologically implausible and constrain the **synthetic lethal** simulation procedure.

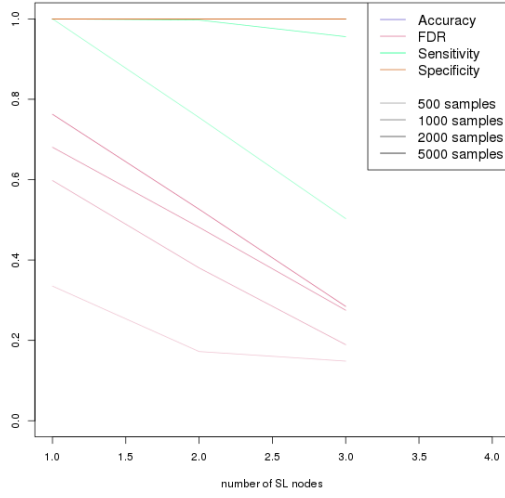
The results with one [synthetic lethal](#) partner were sufficient to infer the impact of [synthetic lethal](#) partners within pathways on neighbouring (correlated) genes. However, it is plausible that the [synthetic lethal](#) signatures in [expression](#) data would propagate through a network with multiple [synthetic lethal](#) partners as sources, provided that the correlations between [synthetic lethal](#) partners are biologically feasible. These simulations were performed on a correlated [graph](#) structure within a larger [gene expression](#) dataset of 20,000 genes (as performed in Sections 3.3 and 6.2.4), a feasible number for a full human [gene expression](#) dataset, and as such are comparable to the findings below.

6.2.4 Performance within a Large Simulated Datasets

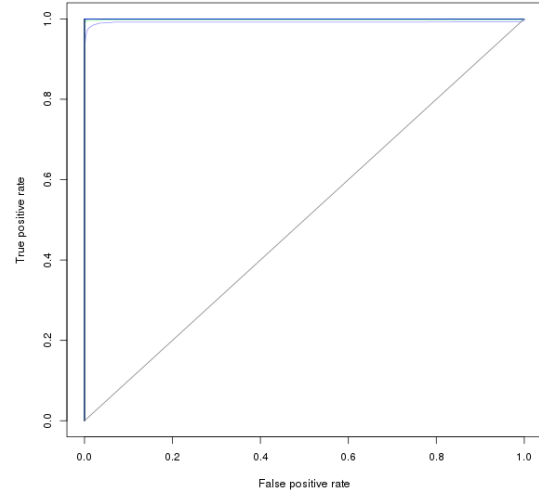
The high proportion of [synthetic lethal](#) partners in small networks (and fewer true negatives) made it difficult to accurately assess the performance of SLIPT with higher numbers of true partners to detect (as noted in Section 6.2.1.1). The addition of more non synthetic lethal genes in previous simulations increased the performance of SLIPT, particularly the specificity to reduce the number of false positives (as shown in Sections 3.3 and 6.1). The [graph](#) structures (as used in Section 6.2.1) of genes with correlations from sampling a multivariate normal distribution were included in a larger simulated dataset of 20,000 genes. This simulation procedure enabled testing of the performance of SLIPT when detecting [synthetic lethal](#) partners within correlated [graph](#) structures (of a [synthetic lethal](#) pathway) in the context of biologically feasible numbers of genes.

The simulations performed in Section 6.2.1.1 were replicated within a dataset of 20,000 genes with the remainder being composed of non synthetic lethal genes without correlation structure. The aforementioned issue with specificity in a higher number of underlying [synthetic lethal](#) genes did not occur in a simple [graph](#) structure (as shown in Figure 6.15). For such a small graph module of highly correlated genes within a [gene expression](#) dataset, detection of [synthetic lethal](#) genes within the network by SLIPT and distinguishing these from the larger dataset had high performance across parameter values. In this case, a reduction in sensitivity resulted in poorer performance, a higher number of non synthetic lethal genes were correctly identified, with a low false positive rate and high accuracy. Thus the use of stringent χ^2 p-value thresholds (adjusted by FDR) are suitable for testing for [synthetic lethality](#) in [gene expression](#) data across the number of genes in human and cancer data.

In a direct comparison with simulations in the [graph](#) structure alone (as performed in Section 6.2.1.1), detection of [synthetic lethality](#) with SLIPT performed consistently



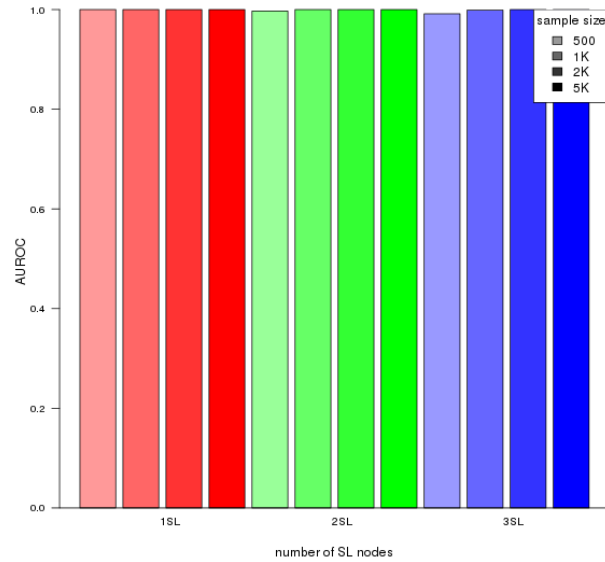
(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.15: **Performance of simulations including a simple graph.** Simulation of **synthetic lethality** was performed by sampling from a multivariate normal distribution (without correlation structure apart from the graph shown). Performance of **SLIPT** was high across parameters for detecting **synthetic lethality** in the **graph** structure within a larger dataset. The sensitivity decreased for a greater number of true positives to detect but the specificity remained high with a low false positive rate.

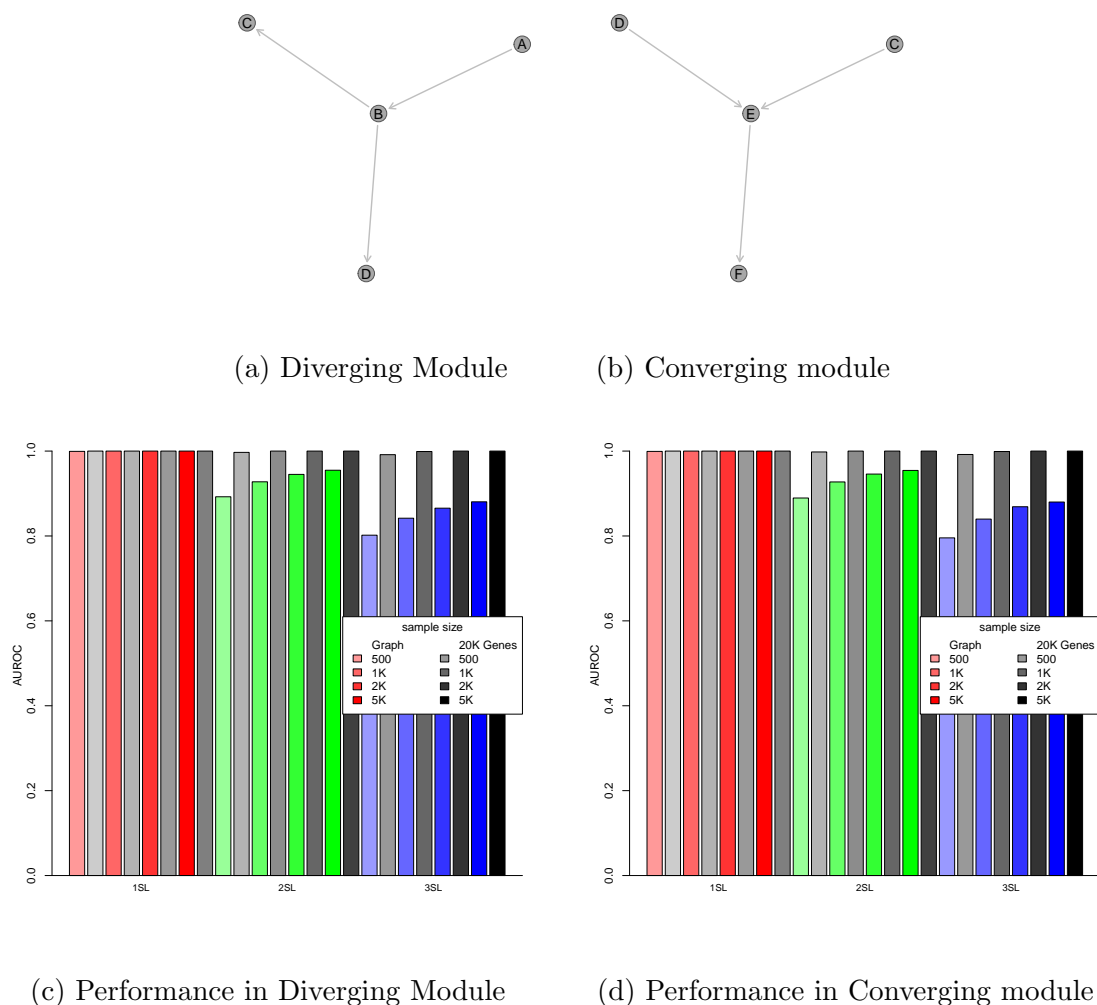


Figure 6.16: **Performance on a simple graph improves with more genes.** Simulations were performed with each of the [graph](#) structures to detect [synthetic lethal](#) partners within them. In either structure, performance of detection in a dataset containing on the [graph](#) structure (in colour) was lower than testing the [graph](#) structure within a larger dataset of non synthetic lethal genes (without correlations).

better in a larger dataset with many true negative genes to detect (as shown in Figure 6.16). The [SLIPT](#) methodology as it has a high specificity and low false positive rate, which is desirable. [SLIPT](#) is therefore applicable to large [gene expression](#) datasets where these are important considerations since the number of negative genes to correctly identify often vastly outnumbers the number of positive genes to detect.

This increase in performance with more non synthetic lethal genes to detect does not necessarily apply in an inhibiting [graph](#) structure. This an important considera-

tion since biological pathways commonly contain inhibiting relationships (and inverse correlations), although they are unlikely to occur across an entire pathway. While an increased performance for an activating graph was replicated, in this case, the performance of simulations of an entirely inhibiting graph structure did not improve within a larger dataset. Rather, the performance in the inhibiting graph structure was similar to simulations of the graph structure in isolation. It is expected that the findings based on these simulations of genes with pathway structures in smaller datasets (as described in Section 6.2.1) will be relevant to larger datasets. The simulation results in these inhibiting graph structures perform comparably or higher with more non synthetic lethal genes to distinguish from them even with inhibitory relationships within the graph structure

Performance of synthetic lethal detection of SLIPT in graph structures with inhibitions included in a larger dataset of non synthetic lethal genes did not diminish to the level of the graph structure simulated alone. In some cases (as shown in Figure 6.17), the performance of an inhibitory graph structure was consistently elevated when included within a larger dataset. However, these did not perform as well as the equivalent graph structures without inhibitory relationships within a similar dataset.

This poorer performance of inhibitory graph structures may be due to highly negatively correlated genes being false positives. These genes will be positively correlated with the query gene if they are negatively correlated with a synthetic lethal partner (i.e., within a synthetic lethal pathway). The SLIPT procedure performs well at distinguishing these positively correlated genes, as previously shown (in Sections 3.3.2.2 and 6.1.1.1). These false positives will also be a minority amongst a larger dataset of non synthetic lethal genes without correlation to the query or synthetic lethal genes.

It more likely that the poorer performance stems from negative correlations between synthetic lethal genes which makes them more difficult to individually detect (as observed in Section 6.2.2). As discussed in Section 6.2.3, this is likely an artifact of the simulation procedure selecting random synthetic lethal genes with strong inhibitory relationships between them. Therefore the poorer performance for inhibiting graphs within larger datasets is not cause for concern because the cases where SLIPT performs poorly are likely to be combinations of simulated synthetic lethal genes that are not likely to occur within biological pathways. This simulation procedure has included higher-order synthetic lethal to produce the weakest signal of synthetic lethality for individual partner genes which are still detectable by SLIPT.

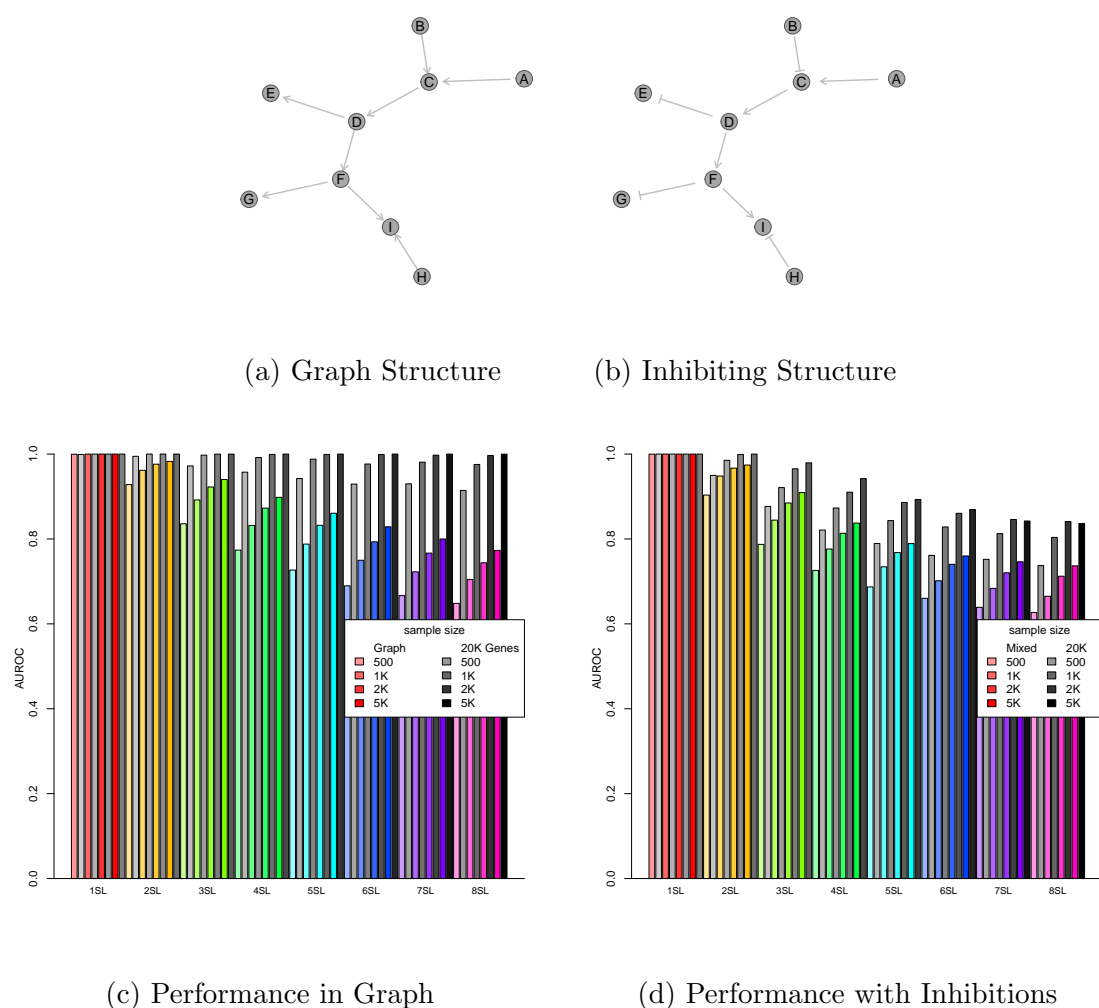


Figure 6.17: **Performance on an inhibiting graph improves with more genes.** Simulations were performed in a [graph](#) structure with activating and inhibiting relationships to detect [synthetic lethal](#) partners within them. In contrast to an activating graph, performance of detection in a dataset containing only the [graph](#) structure (in colour) was as much lower than testing the [graph](#) structure within a larger dataset of non synthetic lethal genes (without correlations) in an inhibiting [graph](#) structure with negative correlations.

6.3 Simulations in More Complex Graph Structures

Investigations with simulations based on [graph](#) structures were extended to larger [graphs](#), enabling more synthetic lethal genes within a pathway and modelling the complexity of a biological pathway. Sensitivity declines over a greater range for the number of [synthetic lethal](#) partners in a larger network with a trade-off with specificity (as shown in Appendix Figures [K.8–K.10](#)). However, the accuracy declined for greater

numbers of [synthetic lethal](#) partners and the false positive rate peaks at intermediate values. In this range, difference between simulations varied with greater sample size. The [AUROC](#) results were similar between these more complex [graph](#) structures, although the larger [graph](#) (Appendix Figure [K.10](#)) differed in sensitivity and specificity at an adjusted ([FDR](#)) p-value threshold. This difference be due to different proportions of [synthetic lethal](#) and non-synthetic lethal genes to detect, since these [graphs](#) (as shown in Appendix Figures [K.8](#) and [K.9](#)) had fewer genes.

While the [graph](#) structures (of similar size) were highly distinct, they had similar performance profiles across parameters. [SLIPT](#) is therefore robust across [pathway](#) structures and is more affected by the number or proportion of genes to detect. Findings from previous simulations in similar correlation structures (in Section [3.3](#)) should be applicable to [expression](#) data with more complex correlation structures, such as those occurring in biological pathways. Specifically, [synthetic lethal](#) partners are distinguishable from closely correlated genes in the context of a biological pathway networks, irrespective of thresholds (shown by [ROC](#)) and with the sensitivity and specificity of p-value thresholds (adjusted by [FDR](#)) as used for [SLIPT](#) (in Chapters [4](#) and [5](#)).

The findings for inhibitory [graph](#) structures were replicated with larger more complex [graph](#) structures with inhibiting relationships and more [synthetic lethal](#) genes to detect (shown in Appendix Figures [K.11–K.14](#)). In each [graph](#) structure, simulations entirely with inhibiting relationships (Appendix Figures [K.11](#), [K.13](#), and [K.15](#)) had higher performance than the equivalent graph with entirely activating relationships (Appendix Figures [K.8](#), [K.9](#), and [K.10](#)) or a combination of activating and inhibiting relationships (Appendix Figures [K.12](#), [K.14](#), and [K.16](#)). While the presence of negative correlations subtly affects the performance of [SLIPT](#), the methodology is robust across the exact structures of genes and is therefore applicable to detecting [synthetic lethal](#) genes in a range of (synthetic lethal) biological pathways with different structural relationships.

6.3.1 Simulations over Pathway-based Graphs

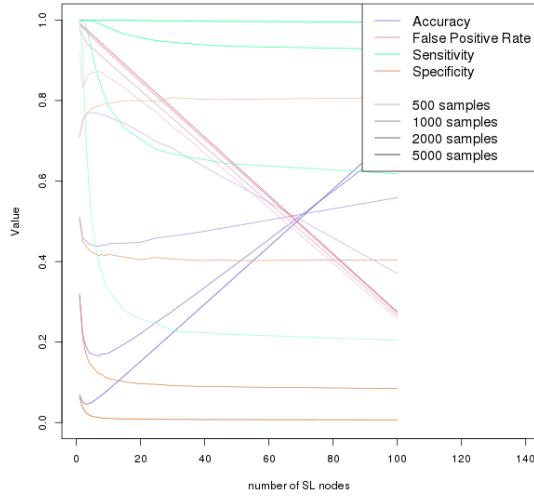
Simulations of [synthetic lethality](#) in [gene expression](#) with correlation structures thus far have used simple blocks of correlated genes (as used in Section [3.3](#)) or have been derived from constructed [graph](#) structures (as used in Section [6.2](#)). These have been used to make inferences on the impact of correlation structure but it remains to be shown whether these findings are reproducible in the complexity of the biological [network](#) structure. Specifically, [SLIPT](#) was tested on simulated data with known underlying

simulated **synthetic lethal** partners (as described in Section 3.2.2) with multivariate normal correlation structure derived from biological pathways (as described in Section 3.4.2).

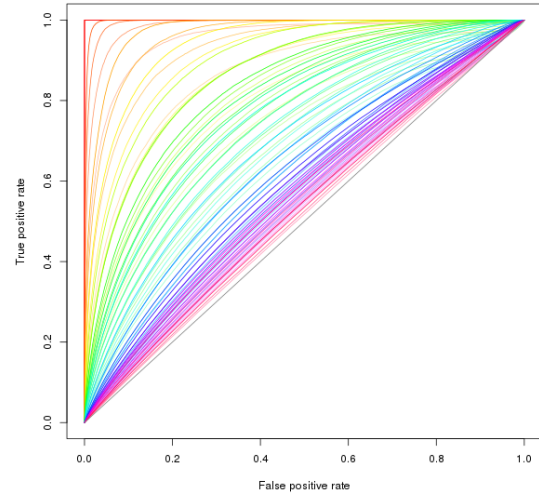
The Reactome **pathway** structure for the **PI3K** cascade (as used in Chapter 5) was used to demonstrate the simulation procedure for detecting **synthetic lethality** in the **graph** structure of a biological pathway. This pathway has clear directionality, with related signalling pathways among those identified to be **synthetic lethal** candidates (in Chapter 4). The **PI3K** pathway has a relatively moderate size (138 genes) and complexity. It is therefore suitable for comparison to previous **graph** structures of a similar scale (50–100 genes) with the complexity of a biological pathway.

The performance of **synthetic lethal** detection with **SLIPT**, in simulated **expression** data based on the Reactome **PI3K** pathway (as shown in Figure 6.18), was concordant with previous findings. **SLIPT** had high performance when detecting a low number of **synthetic lethal** genes which decreased for high numbers of **synthetic lethal** genes or lower sample sizes. In particular, the performance of simulations in the **PI3K** pathway closely resembled the simulation results for constructed **graphs** of similar scale and complexity (as shown in Appendix Figures K.8 and K.9). Using thresholds based on the χ^2 p-value (adjusted by **FDR**), simulations in the biological **PI3K** pathway had a higher sensitivity and lower specificity. While the performance decreased for more **synthetic lethal** genes to detect within the simulated **PI3K** pathway, this primarily involved a reduction in sensitivity to detect **synthetic lethal** genes rather than false positives, as the false positive rate decreased, the accuracy increased, and the specificity was relatively unperturbed (being more dependent on sample size). Thus **SLIPT** was stringent in an example of a biological **graph** structure and is appropriate for detection of **synthetic lethal** genes in complex correlation structures in **gene expression** data involving biological pathways.

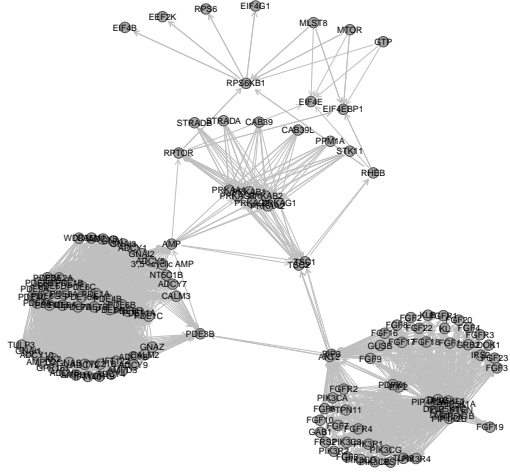
These simulations were replicated in the more complex $G_{\alpha i}$ signalling pathway (of 292 genes), which was one of the most well supported **synthetic lethal** pathways with loss of *CDH1* in cancer (in Chapters 4 and 5). This pathway showed similar relationships between sensitivity, specificity, and false positive rate with number of **synthetic lethal** partners and sample size (as shown in Appendix Figure K.17). While the overall performance was lower than for smaller networks structures, many of the findings from previous networks were replicated in a larger more complex biological network. In the $G_{\alpha i}$ signalling pathway, **SLIPT** had high performance for detecting



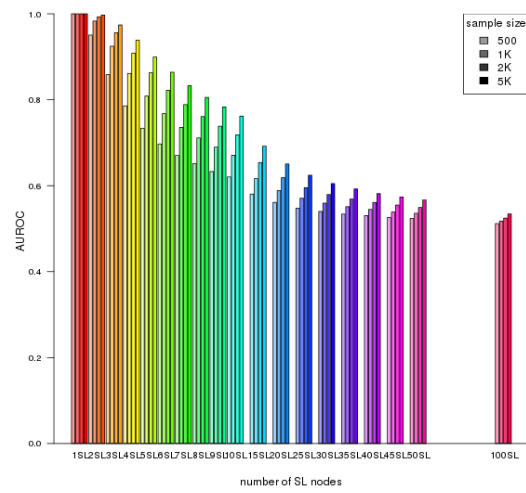
(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.18: **Performance of simulations on the PI3K cascade.** Simulation of **synthetic lethality** was performed by sampling from a multivariate normal distribution based on the Reactome PI3K cascade. Performance of **SLIPT** was high across parameters for detecting **synthetic lethality** in the **graph** structure within a larger dataset. The performance decreased for a greater number of true positives to detect but the accuracy increased with a low false positive rate.

low numbers of **synthetic lethal** genes and was highly stringent against false positives for higher numbers of **synthetic lethal** genes.

6.3.2 Pathway Structures in a Large Simulated Datasets

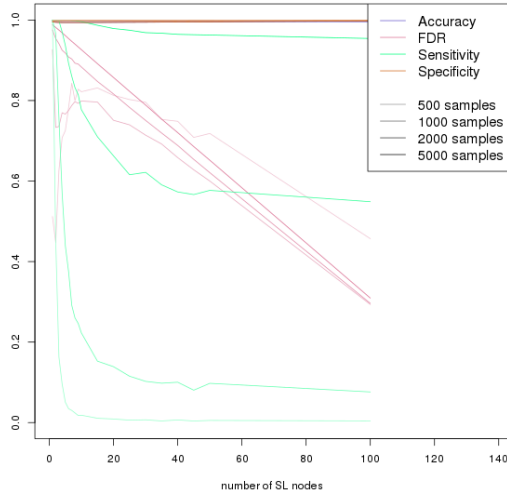
Simulations were also performed with [graph](#) structures from biological pathways included in a larger dataset to simulate [gene expression](#) data of the scale typical for human and cancer studies. These simulations (as discussed in Section 6.2.4) showed a higher specificity and therefore [SLIPT](#) had higher performance. The simulated [PI3K](#) pathway (as shown in Figure 6.19) was no exception, with high performance across parameter values, which remained high up to many genes. While the sensitivity decreased for high numbers of [synthetic lethal](#) genes to detect within the [PI3K](#) pathway, the [SLIPT](#) methodology remained accurate, with high specificity in a large simulated [gene expression](#) dataset.

Therefore the [SLIPT](#) methodology is a highly stringent approach suitable to be applied for detecting [synthetic lethal](#) genes and pathways within highly complex [expression](#) data with biological [pathway](#) structure. Even the poorly performing simulations were highly stringent, with low false positive rates, which are an important consideration in a [gene expression](#) data with many non synthetic lethal genes. The enrichment of true [synthetic lethal](#) partners among detected genes makes [SLIPT](#) valuable for triage of candidate [synthetic lethal](#) partners for further validation and for pathway analysis.

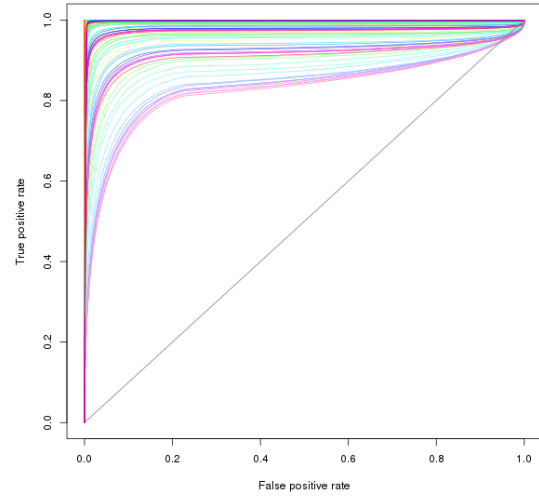
The performance of [SLIPT](#) in simulations of [synthetic lethality](#) within biological pathways was markedly higher in the context of a larger dataset of thousands of genes. As shown in a direct comparison with the [graph](#) structures alone (as shown in Figure 6.20c), performance was consistently higher across parameters in pathways of biological complexity from the Reactome database (Croft *et al.*, 2014) such as [PI3K](#) cascade and the $G_{\alpha i}$ signalling pathway (shown in Figure 6.20d and Appendix Figure K.18).

These biologically complex [graph](#) structures, based on the Reactome pathways, assumed activating relationships to test [synthetic lethal](#) detection with [SLIPT](#) in the context of complex correlation structures. Inhibiting relationships were not distinguished in the Reactome database (Croft *et al.*, 2014). However, these investigations with pathway-based [graph](#) structures indicate that the findings in constructed [graphs](#) (as used in Section 6.2) are relevant to [gene expression](#) data containing real correlated pathways. Furthermore, previous comparisons between simulations with inhibiting relationships indicated that the performance of [synthetic lethal](#) detection in an equivalent [graph](#) structure with inhibitory relationships will likely be higher.

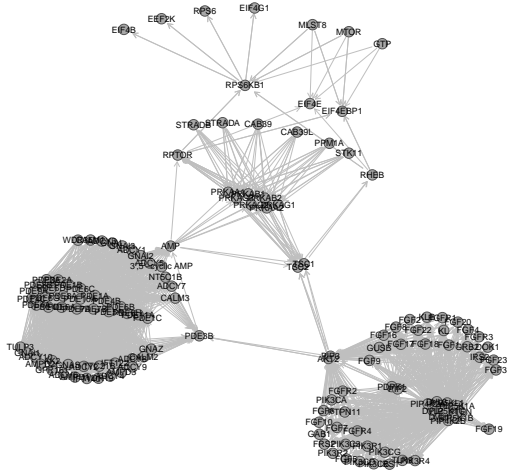
Non synthetic lethal genes, inversely correlated with the underlying [synthetic lethal](#) partners, were distinguishable by [SLIPT](#) with high specificity. [Synthetic lethal](#) genes were detectable with reasonable performance in large scale simulated [gene expression](#)



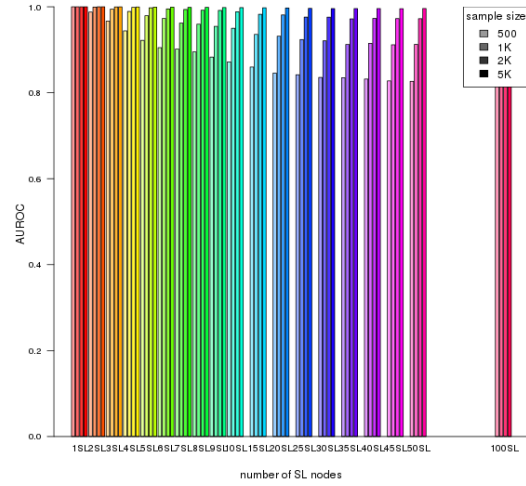
(a) Statistical evaluation



(b) Receiver operating characteristic



(c) Graph Structure



(d) Statistical performance

Figure 6.19: **Performance of simulations including the PI3K cascade.** Simulation of **synthetic lethality** was performed by sampling from a multivariate normal distribution (without correlation structure apart from the Reactome **PI3K** cascade). Performance of **SLIPT** was high across parameters for detecting **synthetic lethality** in the **graph** structure within a larger dataset. The sensitivity decreases for a greater number of true positives to detect but the specificity remains high with a low false positive rate.

data and highly (positively) correlated genes in **pathway** structures. These findings serve as a conservative estimate for the performance of **SLIPT** to detect **synthetic lethal**

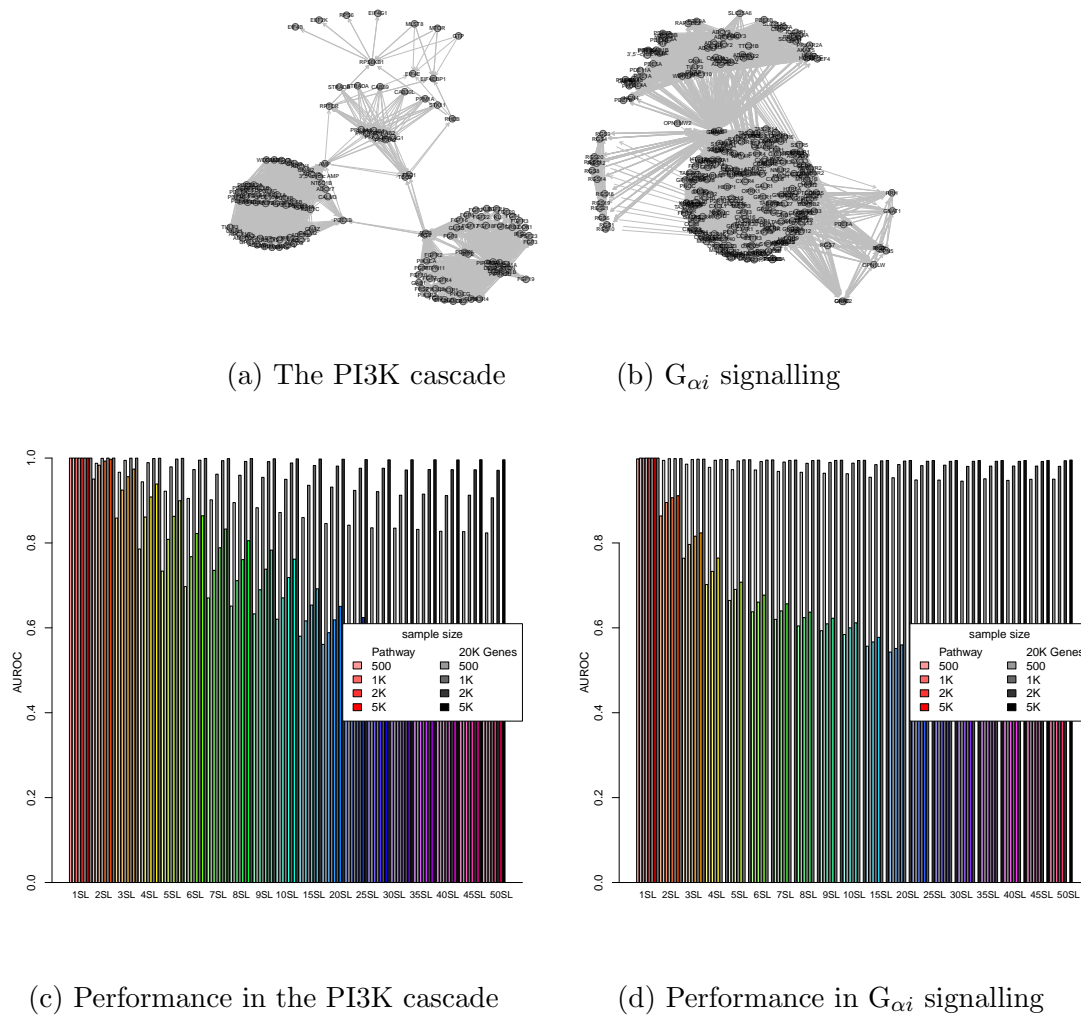


Figure 6.20: **Performance on pathways improves with more genes.** Simulations were performed in a [graph](#) structures for the [PI3K](#) cascade and $G_{\alpha i}$ signalling pathways structures to detect [synthetic lethal](#) partners within them. As for constructed graphs, performance of detection in a dataset containing only the [graph](#) structure (in colour) was as much lower than testing the [graph](#) structure within a larger dataset of non synthetic lethal genes (without correlations) for both [graphs](#) of biological complexity.

genes within a [synthetic lethal](#) biological pathway in empirical data. [Synthetic lethal](#) genes are distinguishable from correlated genes, to varying extents, in simulations. False positives are also more likely to occur within the same (synthetic lethal) pathways. Therefore [SLIPT](#) is both effective at triage of [synthetic lethal](#) candidates within a biological pathway and at identifying [synthetic lethal](#) pathways in high dimensional [gene expression](#) data.

6.4 Discussion

6.4.1 Simulation Procedure

Simulations were performed to assess the performance of the [SLIPT](#) methodology (as described in Section 3.1 and with modifications) for detecting known underlying [synthetic lethal](#) partners of a query gene. The simulation results support the findings in empirical data (in Chapters 4 and 5) by addressing whether the methodology used to generate them is accurate or has desirable statistical performance in controlled simulated conditions. These investigations include adjusting parameters such as the numbers of [synthetic lethal](#) genes which were known in empirical data to assess the performance of the [SLIPT](#) methodology across simulation parameters and characterise the datasets for which [SLIPT](#) performs well. Simulation and statistical modelling also enabled comparison of the [SLIPT](#) methodology to other statistical approaches to [synthetic lethal](#) detection in [expression](#) data.

These simulations were based on a statistical model of [synthetic lethality](#) (as described in Section 3.2.1) which was designed stringently to ensure that if [synthetic lethality](#) is detectable in the simulated datasets it would also be detectable by the same methodology in empirical [expression](#) data. The model of [synthetic lethality](#) made conservative assumptions such as the low threshold of [expression](#) for gene function or the inclusion of cryptic higher-order [synthetic lethality](#) (when testing pairwise). These assumptions decrease the likelihood that [synthetic lethal](#) signatures would be detectable in [expression](#) data. Thus it is reassuring that [synthetic lethality](#) was still detectable in under many simulation parameters as the performance of [SLIPT](#) would be expected to be higher were these assumptions to be violated in empirical data.

The simulation procedure (as described in Section 3.2.2) was designed as a computational pipeline with arguments passes to scripts. The [SLIPT](#) methodology and simulation of [expression](#) from [graph](#) structures were both used as R ([R Core Team, 2016](#)) software packages developed and released for this project (as described in Section 3.5). This design ensures that the simulations can be robustly applied across parameters with consistency between simulations apart from the differences discussed. The simulation procedure is also flexible to simulating other datasets, including [synthetic lethal](#) relationships and pathway correlation structures, should these be relevant to future investigations or [bioinformatics](#) tool development. The computational pipeline is also compatible with parallel computing and made use of [High Performance Computing \(HPC\)](#) infrastructure provided by [the New Zealand eScience Infrastructure \(NeSI\)](#) us-

ing the [Simple Linux Utility for Resource Management \(Slurm\)](#) submission system (as described in Section 2.5.3). This parallel computing pipeline enabled extensive investigations into [synthetic lethality](#) in simulated data, including approximately 2 million cpu-hours on [NeSI](#).

6.4.2 Comparing Methods with Simulated Data

Attempts were made to implement alternative [synthetic lethal](#) detection approaches such as linear regression and the [BiSEp](#) R package (discussed in Section 6.1). However, those tested were ineffective at detecting [synthetic lethality](#) in multivariate normal simulated data in comparison to [SLIPT](#). While some of the published [synthetic lethal](#) detection methods ([Jerby-Arnon *et al.*, 2014](#); [Lu *et al.*, 2015](#)) did not provide reproducible software releases for direct comparison, some of the central assumptions used in their design were tested by the statistical methods considered for [synthetic lethal](#) detection in [expression](#) data.

Another consideration is that [BiSEp](#) takes considerably more time to compute predictions than [SLIPT](#) or χ^2 which limited the number of simulations that were feasible and made it difficult to apply across parameters in the simulation pipeline (even when using supercomputing infrastructure as discussed in Section 2.5.3). The computationally intensive nature of the [BiSEp](#) procedure does not appear to be the issue for detecting [synthetic lethal](#) genes in [TCGA](#) data or simulations, although it has made more extensive simulations challenging. Rather, [BiSEp](#) is not suitable in either case since the [TCGA](#) data is normalised with [voom](#) ([Ritchie *et al.*, 2015](#)) and simulated data is generated by sampling from a multivariate normal distribution. In either case, even subtle bimodal signatures in [expression](#) data were not consistently detectable or sufficient to detect [synthetic lethality](#).

The [BiSEp](#) methodology may perform better on other data types but it cannot be directly compared with the results for [SLIPT](#) throughout this thesis which have used normalised or (multivariate) normally distributed data. Since it requires bimodal distributions, [BiSEp](#) is not suitable for stringently normalised [expression](#) data nor would it be expected to perform on (ranked) pathway [metagenes](#). Thus [SLIPT](#) represents a distinct approach more suitable for these data types whereas [BiSEp](#) may be applicable to other applications in which bimodal distributions are more frequent.

This investigation also demonstrates that implementing scientific software from other research groups is not a trivial exercise, even when released as an open-source R package. Therefore, the above results were used to evaluate [SLIPT](#) and compare it to

other statistical rationales. An comprehensive comparison to contemporary [synthetic lethal](#) detection approaches (and those released in the future) or further benchmarking is left to an impartial researcher to evaluate. The above findings show that the [SLIPT](#) approach is able to detect [synthetic lethal](#) genes in simulated data with comparable or better performance than a range of distinct statistical techniques and was appropriate for use throughout this thesis.

6.4.3 Design and Performance of SLIPT

The simulation procedure using sampling from a multivariate normal distribution was used throughout the majority of the simulation investigations in this thesis. This approach has the advantages of emulating the continuous normalised [expression](#) data used for [gene expression](#) analysis and enabled the simulation of correlation structures (as discussed in Section 3.3). These simulations scaled to datasets of comparable scale to those used in [gene expression](#) analysis with thousands of genes. The [SLIPT](#) methodology was shown to perform robustly across large numbers of genes and simple correlation structures. This includes high specificity against genes positively correlated with the query gene for which the directional [SLIPT](#) methodology was more suited to distinguishing [synthetic lethal](#) genes from than the χ^2 test without directional criteria on the number of samples observed.

These findings were expanded upon in this chapter. Specifically, different quantiles were compared for [SLIPT](#) and the χ^2 test. These approaches using threshold based discrete gene function were compared to the Pearson correlation without loss of the continuous [expression](#) data. The $1/3$ -quantiles for [SLIPT](#) (as described in Section 3.1) were optimal for both [SLIPT](#) and the χ^2 alone. In addition to being optimal for estimating the significance of [synthetic lethal](#) interactions, these quantiles were also optimal for the directional criteria of [SLIPT](#) since this method outperformed the χ^2 test and was the most different at the $1/3$ -quantile. As previously, noted this difference was more pronounced with positively correlated genes (with the query gene) for which the specificity of [SLIPT](#) improved, and was replicated in large datasets with thousands of genes as occur in human [expression](#) data. These results were not simply due to sufficient samples for significant p-values since the performance as determined by [AUROC](#) analysis is independent from significance thresholds. This indicates that the [SLIPT](#) methodology (as it has been used in Chapters 4 and 5) is optimal, with the $1/3$ -quantile having the highest performance, and as such the parameters used to design it were appropriate.

Both discrete functional approaches (SLIPT and χ^2) were able to outperform negative correlation which supports their use. In particular, this result addresses the concern that arbitrary thresholds of low and high gene function (as used by SLIPT) lose useful data by compressing the spectrum of gene expression into categorical data. However, this does not impede the performance of SLIPT if the quantiles used are optimal. The poorer performance of correlation-based detection of synthetic lethality also indicates affirms the concept of gene function for synthetic lethality being qualitative, that is expression must be sufficient for cell viability and higher expression is not necessarily higher function (as this is not the case for all genes). Furthermore, the finding that negative correlation outperforms positive correlation is also consistent with co-expression being a poor predictor of synthetic lethality compared to other approaches (Jerby-Arnon *et al.*, 2014), supporting the claims of Lu *et al.* (2015).

Compared with SLIPT, neither correlation approaches nor bimodality signatures were suitable for detecting synthetic lethality in expression data. The correlation-based approaches make assumptions about the relationship between gene expression and function which do not necessarily hold for all genes. Similarly, the bimodal approach is not appropriate for normalised data since deviations from a normal distribution have already been used for ensuring data quality, as is common practice for RNA-Seq data. A linear model or regression approach may also be used to detect synthetic lethality from relationships between expression of genes, which may be improved with conditioning on known synthetic lethal partners with multivariate regression or Bayesian priors. Similarly, synthetic lethal detection could be performed by iteratively conditioning upon the strong candidate from previous analysis. These approaches may be able to better circumvent the issues of high-order synthetic lethality and multiple testing.

Nevertheless, the above findings are sufficient to assess the performance of SLIPT and present an effective straightforward approach to synthetic lethal detection in gene expression data. Further development with linear models, Bayesian inference approaches, or comparison to existing synthetic lethal approaches (e.g., machine learning) remain as future directions. Developing and testing more sophisticated statistical approaches to synthetic lethal detection may benefit from the concepts discussed with regard to the relatively simple SLIPT methodology. Similarly, further comparisons and benchmarking of SLIPT against other computational approaches to synthetic lethal detection in gene expression data is more suitable for an independent researcher and the `slipt` R package has been released (as described in Section 3.5) for this purpose, in addition to further application in research.

6.4.4 Simulations from Graph Structures

The simple correlation structures (as used in Section 3.3) were expanded upon using the multivariate normal simulation procedure to produce correlation structures based on [graph](#) structures (as described in Section 6.2). These simulations enable further investigations into the performance of [SLIPT](#) in the context of more complex correlation structures. The simulation of [expression](#) from [network](#) structures is widely applicable to simulating pathway [expression](#) data and as such the [graphsim](#) R package has been released (as described in Section 3.5).

These investigations show that [SLIPT](#) performs robustly across datasets with different correlation structures, including those derived from [graphs](#) with the complexity of biological pathways. The [SLIPT](#) methodology was able to detect [synthetic lethal](#) genes within [synthetic lethal](#) pathways across many [graph](#) structures. This methodology performed particularly well with [synthetic lethal](#) pathways in the context of a larger dataset with a high specificity which supports [SLIPT](#) as a stringent approach to [synthetic lethal](#) detection in highly dimensional [gene expression](#) data. Together these results support the use of [SLIPT](#) in biological gene expression data since it is able to detect [synthetic lethal](#) genes in highly complex correlation structures.

Similarly, the inclusion of inhibitory relationships in [graph](#) structures was shown to increase the performance in simple networks, supporting [SLIPT](#) being applicable to biological data in which these relationships are common. While these results were not replicated in more complex inhibitory [graph](#) structures, this is likely an artifact of the simulation procedure (which randomly selects [synthetic lethal](#) genes) generating biologically implausible combinations of [synthetic lethal](#) genes which are difficult to detect. When the test statistics in simulations with a [synthetic lethal](#) gene were examined in more detail, the test statistics of the [synthetic lethal](#) gene were consistently higher and distinguishable from nearby genes in the [graph](#) structure. In contrast to previous concerns with inhibiting relationships, these differences were more pronounced with genes which had inhibitory relationships with [synthetic lethal](#) genes. While distinguishable from nearby genes in a [pathway](#) structure, the genes correlated with [synthetic lethal](#) partners still had higher test statistics than more distant genes (similar to observations with correlated genes in Section 3.3).

In addition to being able to detect [synthetic lethal](#) genes in a pathway, the proximal genes in a pathway are most likely to be false positives and therefore [SLIPT](#) is also able to detect [synthetic lethal](#) pathways. Therefore [SLIPT](#) identifies genes which are likely to be constituent of a [synthetic lethal](#) pathway and is more likely to rank underlying

[synthetic lethal](#) genes with greater significance. Together these findings support the use of [SLIPT](#) throughout this thesis, further application of [SLIPT](#), and further development of such strategies for [synthetic lethal](#) detection. Similarly, the simulation procedures developed and demonstrated for examining [synthetic lethal](#) detection in [expression](#) data using [graph](#) structures is amenable to further development and investigations into [pathway](#) structure in [expression](#) data such as predicting biological pathways from [expression](#) data or the impact of pathways on differential [expression](#) analyses.

6.5 Summary

A statistical model and simulation procedure has been developed to test the performance of the [SLIPT](#) methodology in controlled conditions, using multivariate normal distributions. This simulation procedure has been developed into a computational pipeline which was able to test the statistical performance (using stringent assumptions) of [SLIPT](#) across many parameters and compare it to alternative [synthetic lethal](#) detection strategies. The [SLIPT](#) methodology performs well at detecting small numbers of [synthetic lethal](#) genes in simple systems. It does not perform as well in more complex systems but neither do alternative strategies. The [SLIPT](#) methodology performs well compared to Pearson correlation and similar methods based on the χ^2 test. Thus [SLIPT](#) is an effective detection method for [synthetic lethal](#) relationships in [expression](#) data despite its relatively simple design.

Simulations of more complex datasets were performed, including large numbers of genes, complex correlation structure derived from [graph](#) structures, and correlations with the query gene. [SLIPT](#) performs robustly across these, including correlation structures based on complex biological pathways. The performance of [SLIPT](#) improves in larger datasets, datasets with positive correlations with the query genes, and some [graph](#) structures which include inhibiting relationships, namely those datasets more representative of [gene expression](#) in biological data. [SLIPT](#) was both capable of recurrently detecting genes within a [synthetic lethal](#) pathway, and distinguishing [synthetic lethal](#) genes from correlated with them, even in highly complex correlation structures. Therefore [SLIPT](#) is a stringent [synthetic lethal](#) detection strategy and is applicable to [gene expression](#) as previously demonstrated for the partners of *CDH1* in breast and stomach cancer in this thesis.

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