Getting started with the SPLITT library

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 \mathbf{SPLITT} is a one-header C++ library.

knitr::include_graphics("figures/UmlDiagram8.pdf")

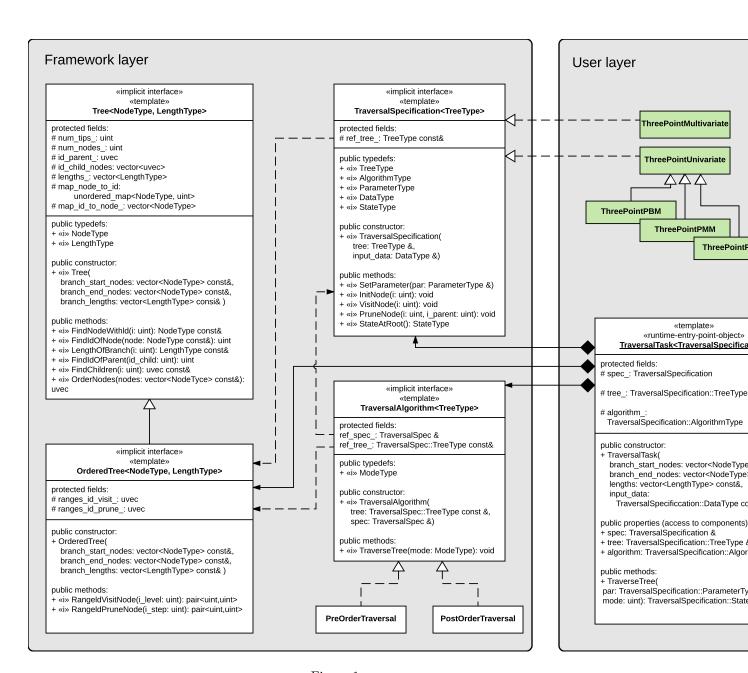
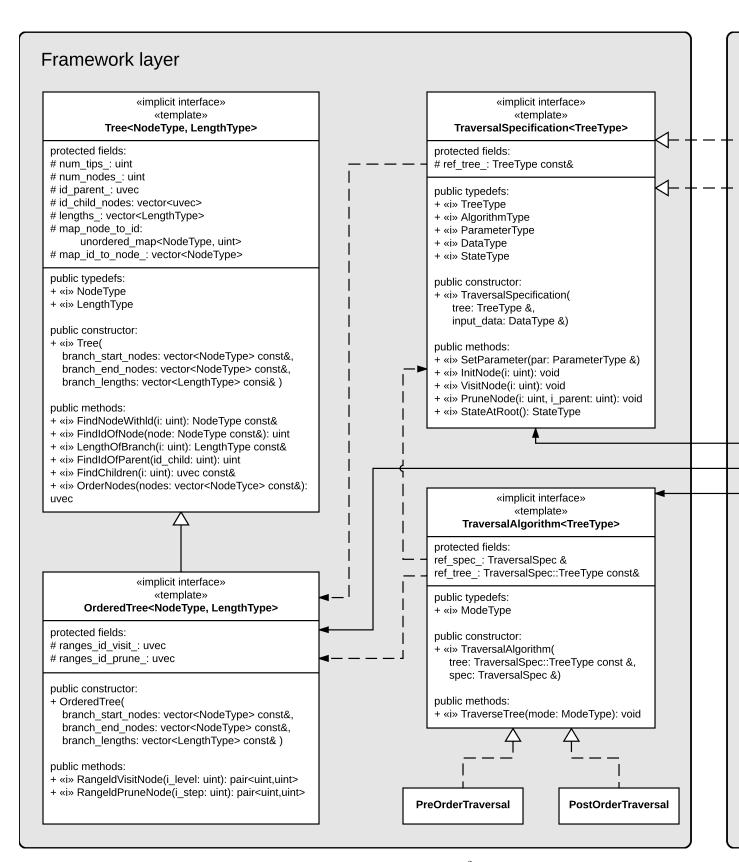


Figure 1:

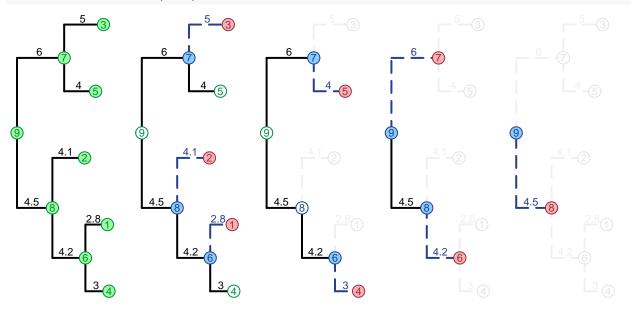


Here I show how to use **SPLITT** to estimate the broad-sense heritability, H^2 , of a numeric pathogen trait. H^2 is defined as the proportion of the trait variance attributable to genotypic variation in a population of

Packages used

```
newick <- '(((1:3,2:2.8)6:4.2,3:4.1)7:4.5,(4:4,5:5)8:6)0;'
tree <- read.tree(text=newick)</pre>
x0 < -0.1
sigma2 <- 0.25
sigmae2 <- 1
set.seed(1)
g <- rTraitCont(tree, model = "BM", root.value = x0,
                sigma = sqrt(sigma2),
                ancestor = FALSE)
x <- g + rnorm(n = length(tree$tip.label), mean = 0, sd = sqrt(sigmae2))
PMMLogLikCpp(x, tree, x0, sigma2, sigmae2, mode = 1)
## [1] -8.2
# output
# InitNode(1): x[i] t[i] a[i] b[i] c[i] : 1.9 2.8 -0.5 1.9 -2.72
# InitNode(2): x[i] t[i] a[i] b[i] c[i] : 0.96 4.1 -0.5 0.96 -1.38
# InitNode(3): x[i] t[i] a[i] b[i] c[i] : -0.74 5 -0.5 -0.74 -1.2
# InitNode(4): x[i] t[i] a[i] b[i] c[i] : 1.79 3 -0.5 1.79 -2.53
# InitNode(5): x[i] t[i] a[i] b[i] c[i] : 0.46 4 -0.5 0.46 -1.03
# InitNode(6): x[i] t[i] a[i] b[i] c[i] : nan 4.2 0 0 0
# InitNode(7): x[i] t[i] a[i] b[i] c[i] : nan 6 0 0 0
# InitNode(8): x[i] t[i] a[i] b[i] c[i] : nan 4.5 0 0 0
# InitNode(9): x[i] t[i] a[i] b[i] c[i] : nan 1.4822e-323 0 0 0
# VisitNode(1): x[i] t[i] a[i] b[i] c[i] : 1.9 2.8 -0.29 1.12 -2.25
# PruneNode(1, 6): x[i] t[i] a[j] b[j] c[j] : 1.9 2.8 -0.29 1.12 -2.25
\# \ VisitNode(2): \ x[i] \ t[i] \ a[i] \ b[i] \ c[i] : 0.96 \ 4.1 \ -0.25 \ 0.47 \ -1.5
# PruneNode(2, 8): x[i] t[i] a[j] b[j] c[j] : 0.96 4.1 -0.25 0.47 -1.5
# VisitNode(3): x[i] t[i] a[i] b[i] c[i] : -0.74 5 -0.22 -0.33 -1.45
# PruneNode(3, 7): x[i] t[i] a[j] b[j] c[j] : -0.74 5 -0.22 -0.33 -1.45
# VisitNode(4): x[i] t[i] a[i] b[i] c[i] : 1.79 3 -0.29 1.03 -2.12
# PruneNode(4, 6): x[i] t[i] a[j] b[j] c[j] : 1.79 3 -0.58 2.14 -4.36
# VisitNode(5): x[i] t[i] a[i] b[i] c[i] : 0.46 4 -0.25 0.23 -1.32
# PruneNode(5, 7): x[i] t[i] a[j] b[j] c[j] : 0.46 4 -0.47 -0.1 -2.77
# VisitNode(6): x[i] t[i] a[i] b[i] c[i] : nan 4.2 -0.26 0.97 -3.68
# PruneNode(6, 8): x[i] t[i] a[j] b[j] c[j] : nan 4.2 -0.51 1.44 -5.18
# VisitNode(7): x[i] t[i] a[i] b[i] c[i] : nan 6 -0.2 -0.04 -3.2
# PruneNode(7, 9): x[i] t[i] a[j] b[j] c[j] : nan 6 -0.2 -0.04 -3.2
# VisitNode(8): x[i] t[i] a[i] b[i] c[i] : nan 4.5 -0.24 0.67 -5.01
# PruneNode(8, 9): x[i] t[i] a[j] b[j] c[j] : nan 4.5 -0.43 0.63 -8.22
par(mfrow=c(1,5))
par(mar=c(0,0,0,0))
par(oma=c(0, 0,0, 0))
```

PlotParallelTraversal(tree)



C++ object <0x10a930ef0> of class 'SPLITT__RCPP__OrderedTreeStringNodes' <0x101f3a7f0>

```
#newick <- '(((1:4,2:4.5)11:4.3,3:4)10:1.4, (4:10.25, ((5:4,6:3.8)14:4.2,7:4.1)13:4.5, (8:6,9:6)15:6)12:2.
newick <- '(((1:4,2:4.5)11:4.3,3:4)10:3.8, (((5:3,6:2.8)14:4.2,7:4.1)13:4.5, (8:5,9:5)15:6)12:4.2)0;'
tree <- read.tree(text=newick)

par(mfrow=c(1,5))
par(mar=c(0,0,0,0))

plotParallelPruningOrder(tree, TRUE)

fig.a <- recordPlot()

newick <- "(((((1:3.2,2:3.2)10:3.2,3:3.2)11:3.2,4:3.2)12:3.2,5:3.2)13:3.2,6:3.2)14;"
tree <- read.tree(text=newick)

par(mfrow=c(1,6))
par(mar=c(0,0,0,0))
plotParallelPruningOrder(tree, TRUE)

fig.b <- recordPlot()

cowplot::plot_grid(fig.a, fig.b, nrow=2, labels = c("a", "b"))</pre>
```

Apart from base R functionality, the patherit package uses a number of 3rd party R-packages:

- For tree processing: ape v5.1 (Paradis et al. 2016);
- For reporting: data.table v1.11.2 (Dowle and Srinivasan 2016);
- For the POUMM and PP estimates: POUMM v2.1.0 (Mitov and Stadler 2017), boot v1.3.20 (Canty and Ripley 2016);
- For testing: testthat v2.0.0 (Wickham 2016).

References

Canty, Angelo, and Brian Ripley. 2016. Boot: Bootstrap R (S-Plus) Functions. R-Package. https://cran.r-project.org/web/packages/boot/index.html.

Dowle, Matt, and Arun Srinivasan. 2016. Data.table: Extension of 'Data.frame'. https://CRAN.R-project.org/package=data.table.

Mitov, Venelin, and Tanja Stadler. 2016. "The heritability of pathogen traits - definitions and estimators." *Unpublished Data* https://www.biorxiv.org/content/early/2016/06/12/058503.

———. 2017. Fast and Robust Inference of Phylogenetic Ornstein-Uhlenbeck Models Using Parallel Likelihood Calculation. BiorXiv. http://biorxiv.org/content/early/2017/05/30/115089.

———. 2018. "A Practical Guide to Estimating the Heritability of Pathogen Traits." *Molecular Biology and Evolution*, msx328. doi:10.1093/molbev/msx328.

Paradis, Emmanuel, Simon Blomberg, Ben Bolker, Julien Claude, Hoa Sien Cuong, Richard Desper, Gilles Didier, et al. 2016. Ape: Analyses of Phylogenetics and Evolution. https://CRAN.R-project.org/package=ape.

Wickham, Hadley. 2016. Testthat: Unit Testing for R. https://CRAN.R-project.org/package=testthat.