The algebra of Markov models on phylogenetic split networks

Jeremy Sumner School of Maths and Physics, University of Tasmania Joint work with Peter Jarvis and Barbara Holland

Phylomania, 29-30 Oct, 2009

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- It is well known that biological reality is more complicated.
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- There are various ways this is coped with: split networks via distances, ie. incompatible distance metrics.
- However it would be nice to generalize probability models themselves to arbitrary split networks.

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- We take e_i ≡ δ_i, where the e_i form a basis for the vector space V ≡ C^k.
- ► This allows us to speak of a *phylogenetic tensor* $P := \sum_{i_1, i_2, ..., i_n \in X} p_{i_1 i_2 ... i_n} e_{i_1} \otimes e_{i_2} \otimes ... \otimes e_{i_n} \text{ embedded in } V^{\otimes n}.$

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- It is that M₁ ⊗ M₂ occurs as a tensor product which gives stochastic independence across branches.
- Relaxing the condition is central to generalizing to incompatible split systems.

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▶ ie. For
$$k = 2$$
, forget t and take $M = e^Q$ with $Q = \begin{pmatrix} -\alpha & \beta \\ \alpha & -\beta \end{pmatrix}$.

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The general Markov model and CTMCs

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- ► These matrices form a "Lie algebra": $[L_{\alpha}, L_{\beta}] = L_{\beta} L_{\alpha}$.
- This condition is exactly what is needed to ensure that a continuous-time model is *closed* (see PJ's talk).

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- Here we have a "a ha!" moment and replace T with an arbitrary split system S.

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$$P = \exp [\mathcal{R}_1] \cdot \exp [\mathcal{R}_2] \cdot \ldots \cdot \exp [\mathcal{R}_{n-1}] \cdot \delta^n \cdot \pi,$$

where $\mathcal{R}_i = \sum_{A, |A|=i} \tau_A \mathcal{L}^{(A)}.$

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- Clearly, on an arbitrary split system the two approaches are different...
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- The L-rep mis-behaves (badly!) under marginalizations and is chronically rooted (for split systems?=contradication).

At this point it seems wise to think more carefully about what we would like to *do* with these models.

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- Special thanks to David Bryant, John Rhodes and Elizabeth Allman for additional discussions and comments.

Closing remarks

- At this point it seems wise to think more carefully about what we would like to *do* with these models.
- Hopefully the actual biological applications will be a useful guide here.
- Special thanks to David Bryant, John Rhodes and Elizabeth Allman for additional discussions and comments.

Thanks for listening!