The total external length in the evolving Kingman coalescent

Götz Kersting and Iulia Stanciu

Goethe Universität, Frankfurt am Main

CIRM, Luminy June 11–15 2012



The evolving Kingman N-coalescent (N = 5):



Moran's model with time $-\infty < t < \infty$: Links between pairs of lines appear at rate 1, independent between the different pairs. The evolving Kingman *N*-coalescent:



Kingman's coalescent at time t_1

The evolving Kingman *N*-coalescent:



The coalescent tree evolves in time.

The evolving Kingman *N*-coalescent:



Evolving time to MRCA tree topology total length total external length These results are rather different in nature, none covering any other.

Interesting aspects:

- limiting processes with
 - a.s. continuous paths versus
 - a.s. (compensated) pure jump paths
- different time scalings

A BASKET OF RESULTS

Theorem: (Pfaffelhuber/Wakolbinger, Donnelly/Kurtz, 2006)

Let $A_N(t)$ be the time to the MRCA of the evolving Kingman *N*-coalescent at time $t \in \mathbb{R}$. Then, as $N \to \infty$,

$$(A_N(t))_{t\in\mathbb{R}} \stackrel{d}{\to} A$$
,

where the limiting process $A = (A_t)_{t \in \mathbb{R}}$ is stationary, a.s. pure jump, non-Markovian.

For a related result on the two oldest families in the genealogy see Delmas, Dhersin, and Siri-Jegousse (2010).

Theorem: (Greven, Pfaffelhuber, Winter, 2009, 2010)

Let $T_N(t)$ be the tree, induced by the evolving Kingman Ncoalescent at time $t \in \mathbb{R}$ in the space of real trees furnished with the Gromov-weak topology. Then, as $N \to \infty$,

$$(T_N(t))_{t\in\mathbb{R}} \stackrel{d}{\to} T$$
,

where the limiting tree-valued process $T = (T_t)_{t \in \mathbb{R}}$ is stationary, a.s. continuous, and unique solution of a martingale problem.

Theorem: (Pfaffelhuber, Wakolbinger, Weisshaupt, 2011)

Let $L'_N(t)$ be the total length of the evolving Kingman N-coalescent at time $t \in \mathbb{R}$. Then, as $N \to \infty$,

$$(L'_N(t) - 2 \log N)_{t \in \mathbb{R}} \stackrel{d}{\rightarrow} L',$$

where the limiting process $L' = (L'_t)_{t \in \mathbb{R}}$ is stationary, a.s. pure jump, non-Markovian.

L' has *infinite quadratic variation* (Knobloch, Stanciu, Wakolbinger, 2011), thus fails to be a semimartingale. *Theorem:* (Schweinsberg, 2011)

Let $L_N''(t)$ be the total length of the evolving *Bolthausen-Sznitman N*-coalescent at time $t \in \mathbb{R}$. Then, as $N \to \infty$,

$$\left(\frac{(\log N)^2}{N}L_N''(\frac{t}{\log N}) - \log N - \log \log N\right)_{t \in \mathbb{R}} \stackrel{d}{\to} L'',$$

where the stationary limiting process $L'' = (L''_t)_{t \in \mathbb{R}}$ solves the SDE

$$dL'' = -L'' dt + dY$$

for a certain Lévy-process Y of index 1.

Theorem: (K., Stanciu, 2012, ongoing work)

Let $L_N(t)$ be the total *external length* of the evolving Kingman N-coalescent. Then

$$\left(\sqrt{\frac{N}{4\log N}} \left(L_N(\frac{t}{N}) - 2\right)\right)_{t \in \mathbb{R}} \quad \stackrel{d}{\to} \quad L \;,$$

where L is a stationary, Gaussian, a.s. continuous, with covariance function

$$Cov(L_s, L_t) = \left(\frac{1}{1+|t-s|}\right)^2$$
.

Note the different scaling of time (*real* instead of *evolutionary* periodes).

The dynamics of the external lengths:



So let for the *static* Kingman *N*-coalescent (at time t = 0)

 L_N^i = total internal branch length of level i,

in particular for i = 1

 L_N^1 = total external branch length .

How do we gain access to these quantities?

Branch numbers V_N, \ldots, V_2 and W_N, \ldots, W_2 .



$$L_N^1 = \sum_{i=2}^N V_i (T_{i-1} - T_i) , \quad L_N^2 = \sum_{i=2}^N W_i (T_{i-1} - T_i)$$

15

$$L_{N}^{1} = \sum_{i=2}^{N} V_{i}(T_{i-1} - T_{i})$$
$$= \sum_{k=1}^{N-1} T_{k}(V_{k+1} - V_{k}) + T_{1}V_{2}$$
$$\approx \sum_{k=1}^{N-1} \frac{2}{k} \Delta V_{k}$$

 ΔV_k is easy to handle for k close to N.

External branch numbers $0 = V_1, V_2, \dots, V_N$, and (total) internal branch numbers $U_1 = 1 - V_1, U_2 = 2 - V_2, \dots, U_N = 1 - V_N$



N = 1000



Let us look at the randomness within

$$V_N - N \quad (= 0)$$
$$V_{N-1} - N + 2$$
$$\vdots$$
$$V_{N-i} - N + 2i$$

:

Randomness within (V_i) close to N: $V_N - N + 1, V_{N-1} - N + 3, ...$



Randomness enters (almost) independently.

However, since we consider

$$L_N^1 \approx \sum_{k=1}^{N-1} \frac{2}{k} \Delta V_k$$

we have to understand randomness at the beginning of V_2, \ldots, V_N . So let us compare

$$V_N - N, V_{N-1} - N + 2, \dots$$
 to V_1, V_2, \dots

Randomness within (V_i) close to N and 1:

 $V_N - N + 1, V_{N-1} - N + 3, \dots$ versus $V_1 + 1, V_2 + 1, \dots$



For the (total) internal numbers $U_1 = 1 - V_1, \dots, U_N = N - V_N$ we have reversibility:

Theorem: (Janson, K. 2011)

$$(U_1, \ldots, U_{N-1}) \stackrel{d}{=} (U_{N-1}, \ldots, U_1)$$
.

For another proof see Knobloch, Stanciu, Wakolbinger (2011).

Theorem: (Janson, K. 2011)

$$\sqrt{rac{N}{4\log N}} \left(L_N^1 - 2
ight) \stackrel{d}{
ightarrow} N(0,1) \; .$$

The representation of the (total) internal numbers U_1, \ldots, U_N as diminishing urn:

- Take urn with *blue* balls, altogether N balls.
- Remove them stepwise:
 Successively remove a random pair of balls and replace it by one *orange* ball.
- If i balls are left, let U_i the number of orange balls among them and V_i the number of blue balls.

Note:

$$V_{N-i} - N + 2i = i - U_{N-i}$$
, $V_i = i - U_i$.

Now recall

Note:

 V_N,\ldots,V_2 external branch numbers W_N,\ldots,W_2 internal branch numbers level 2

 $V_N, V_{N-1}, \ldots, V_2$ is a *Markov chain* (inhomogeneous in time).

 $(V_N, W_N), (V_{N-1}, W_{N-1}), \cdots, (V_2, W_2)$ is a Markov chain, or

 W_N, \ldots, W_2 is a Markov chain, given the random environment V_N, \ldots, V_2 .

The transition probabilities:

$$P_{v,w}^{k}(v',w') = \mathbf{P}(V_{k-1} = v', W_{k-1} = w' \mid V_{k} = v, W_{k} = w)$$

$$P_{v,w}^{k}(v-2,w+1) = \frac{\binom{v}{2}}{\binom{k}{2}} \qquad P_{v,w}^{k}(v,w-2) = \frac{\binom{w}{2}}{\binom{k}{2}}$$

$$P_{v,w}^{k}(v,w) = \frac{\binom{k-v-w}{2}}{\binom{k}{2}} \qquad P_{v,w}^{k}(v-1,w-1) = \frac{vw}{\binom{k}{2}}$$

$$P_{v,w}^k(v-1,w) = \frac{v(k-v-w)}{\binom{k}{2}}$$

$$P_{v,w}^k(v,w-1) = \frac{w(k-v-w)}{\binom{k}{2}}$$

External branch numbers V_1, \ldots, V_N , and (total) internal branch numbers U_1, \ldots, U_N and internal branch numbers level 2 W_1, \ldots, W_N .



Randomness within (V_i) and (W_i) : $V_N - N + 1, V_{N-1} - N + 3, \dots$ and $V_1 + 1, V_2 + 1, \dots$ and $W_1 + 1, W_2 + 1, \dots$



Theorem: (K., Stanciu 2012)

For every $k \in \mathbb{N}$

$$\sqrt{\frac{N}{4\log N}} \left(L_N^1 - \mu_1, \dots, L_N^k - \mu_k \right) \stackrel{d}{\to} N(0, I_k) .$$

with the $k \times k$ identity matrix I_k and

$$\mu_i = \frac{2}{i} \; .$$

Idea of proof:

Reversing time seems no longer feasible.

We couple the Markov chain

 $(V_N, W_N), \ldots, (V_2, W_2)$

with two independent urns, i.e. with

 $(V_N, \tilde{V}_N), \ldots, (V_2, \tilde{V}_2)$,

where $(\tilde{V}_N, \ldots, \tilde{V}_2)$ is an independent copy of (V_N, \ldots, V_2) .

Now the urns can be reversed.

Back to the evolving coalescent:



$\operatorname{Cov}(L_s, L_t) =$

probability that a critical binary branching process consists of exactly 1 individual at time |t - s|.

