## CORRECTION

## GROWTH PROFILE AND INVARIANT MEASURES FOR THE WEAKLY SUPERCRITICAL CONTACT PROCESS ON A HOMOGENEOUS TREE

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The proof of Theorem 3 in [1] is incorrect, as it relies on a faulty use of the strong Markov property. The theorem asserts that  $\beta = \beta(\lambda)$  is strictly increasing in  $\lambda$  for  $\lambda < \lambda_2$ , where  $\lambda$  is the infection rate parameter for the contact process,  $\lambda_2$  is the upper critical value (at the transition from weak to strong survival), and  $\beta = \lim_{n \to \infty} u_n^{1/n}$  where  $u_n$  = probability that a vertex  $x_n$  at distance n from the root is ever infected, given that only the root is infected at time t = 0. In this note we shall prove the following slightly weaker result.

THEOREM 3'. If 
$$\beta(\lambda) < 1/\sqrt{d}$$
 and  $\lambda_* < \lambda$  then  $\beta(\lambda_*) < \beta(\lambda)$ .

This leaves open the possibility that  $\beta(\lambda) = 1/\sqrt{d}$  on an interval  $[\lambda_3, \lambda_2]$  of positive length. The proof of Theorem 3' below relies on the following estimate proved by Schonmann (Theorem 2) in [4]: If  $\beta(\lambda) < 1/\sqrt{d}$  then for some constant  $0 < C < \infty$  and every integer  $n \ge 1$ ,

$$(0.1) \frac{\beta(\lambda)^n}{C^n} \le u_n.$$

Schonmann's argument makes essential use of the fact, proved in [1], that  $\beta < 1/\sqrt{d}$  implies  $\eta < 1$ , where  $\eta = \lim_{t \to \infty} P\{\text{root} \in A_t\}^{1/t}$ .

The proof of Theorem 3' also uses the following elementary results.

LEMMA 1. Let  $X, X_1, X_2, \ldots$  be independent, identically distributed, positive integer-valued random variables, and let N be a geometrically distributed random variable independent of the random variables  $X_1, X_2, \ldots$  Suppose that the probability generating function  $\varphi(z) := Ez^X$  is finite for  $1 \le z < R$  and infinite at z = R. Then

(0.2) 
$$\limsup_{n \to \infty} P \left\{ \sum_{i=1}^{N} X_i > n \right\}^{1/n} > 1/R.$$

PROOF. Let  $S_N = \sum_{i=1}^N X_i$ . If N has geometric distribution with parameter  $0 , that is, if <math>P\{N = k\} = pq^k$  for all k = 0, 1, 2, ..., where q = 1 - p, then the probability generating function of  $S_N$  is  $Ez^{S_N} = p/(1 - q\varphi(z))$ . Since  $\varphi(z) \to \infty$  as  $z \to R-$ , there exists  $1 < \rho < R$  such that  $\varphi(\rho) = 1/q$ . Thus, the smallest positive singularity of the probability generating function  $Ez^{S_N}$  is no larger than  $\rho$ . The result (0.2) now follows from Pringsheim's theorem.  $\square$ 

LEMMA 2. Let X be a positive-integer valued random variable with probability generating function  $\varphi(z) = Ez^X$ . If, for some R > 1,

(0.3) 
$$\lim_{n \to \infty} P\{X \ge n\}^{1/n} = 1/R$$

and

(0.4) 
$$\sum_{n\geq 1} R^n P\{X \geq n\} = \infty$$

then  $\varphi(z)$  is finite for  $1 \le z < R$  and is infinite at z = R.

PROOF. This follows by Pringsheim's theorem from the identity

$$\sum_{n=1}^{\infty} z^n P\{X \ge n\} = z \frac{Ez^X - 1}{z - 1}.$$

To prove Theorem 3' we shall compare the growth of the contact processes  $A_t$  and  $A_t^*$  with infection rates  $\lambda$  and  $\lambda_*$ , respectively, by constructing an auxiliary process  $B_t$  whose growth rate is strictly greater than that of  $A_t^*$  and is such that  $B_t \subseteq A_t$ . The construction of these processes uses the augmented percolation structure  $\mathcal P$  described in Section 6 of [1]. Recall that  $\mathcal P$  consists of mutually independent Poisson processes of recovery marks and infection arrows attached to vertices and neighboring pairs of vertices, respectively; the recovery mark processes have intensity 1 and the infection arrow processes have intensity  $\lambda$ . In addition,  $\mathcal P$  attaches to each infection arrow  $\alpha$  a Bernoulli-p random variable  $\xi_{\alpha}$ , where  $\lambda_* = \lambda q/(1+p)$  and q = 1-p. The contact process  $A_t$  is constructed from the system of infection arrows and recovery marks of  $\mathcal P$  in the usual way (see [3]).

The contact process  $A_t^* = A'_{t/(1+p)}$  is constructed in a similar fashion, but using the modified percolation structure  $\mathcal{P}'$  obtained from  $\mathcal{P}$  by changing all infection arrows  $\alpha$  such that  $\xi_{\alpha} = 1$  to recovery marks at their base points. The percolation structure  $\mathcal{P}'$  again consists of independent Poisson processes of recovery marks and infection arrows; the recovery mark processes have intensity 1 + p and the infection arrow processes have intensity  $\lambda q$ . The contact process  $A'_t$  is built using the arrows and recovery marks of  $\mathcal{P}'$  in the usual way. Clearly, if  $A_t$  and  $A_{t'}$  have common initial state  $A_0 = A'_0 = \{e\}$  then  $A_t \subseteq A'_t$  for all  $t \ge 0$ .

The auxiliary process  $B_t$  is constructed using a sequence of independent severed contact processes  $C^1, C^2, \ldots$  A contact process  $C_t$  is severed across an edge  $\varepsilon$  of

the tree if it is constructed in the usual manner using the percolation structure  $\mathcal{P}'$  from which all infection arrows across the edge  $\varepsilon$  have been removed. Let  $x_0, x_1, x_2, \ldots$  be the successive vertices along a geodesic ray  $\gamma$  emanating from  $x_0 = e$ , and let  $x_{-1}$  be a vertex at distance one from e distinct from  $x_1$ . Define  $C_t^1$  to be the contact process with initial state  $C_0^1 = \{e\}$  that is severed across the edge joining  $x_{-1}$  and e. By construction,  $C_t^1 \subseteq A_t'$ . Since  $A_t'$  does not survive strongly, there is a maximal integer v(1) such that vertex  $x_{v(1)}$  is ever infected by the severed contact process, that is,

$$x_{\nu(1)} \in \bigcup_{t \ge 0} C_t^1.$$

Observe that, since  $x_{\nu(1)}$  is the farthest vertex along the geodesic ray  $\gamma$  ever infected by the severed contact process  $C^1$ , there must be a recovery mark  $\rho_1$  at  $x_{\nu(1)}$  terminating the last period of infection at  $x_{\nu(1)}$ .

Define  $C^2, C^3, \ldots$  inductively as follows: First, adjoin to the vertex  $x_{\nu(1)+1}$ , beginning at the time  $\tau_1$  of the recovery  $\rho_1$ , a contact process  $C^2$  that is severed across the edge joining  $x_{\nu(1)}$  to  $x_{\nu(1)+1}$ . Define  $\nu(2) > \nu(1)$  to be the index of the last vertex  $x_{\nu(2)}$  along the geodesic ray  $\gamma$  that is ever infected by  $C^2$ , and let  $\rho_2$  be the recovery mark terminating the last epoch of infection at  $x_{\nu(2)}$ . Adjoin to  $x_{\nu(2)+1}$ , starting at the time  $\tau_2$  of the recovery mark  $\rho_2$ , a contact process  $C^3$  that is severed across the edge joining  $x_{\nu(2)}$  to  $x_{\nu(2)+1}$ . Continue this process indefinitely. Observe that the severed contact processes  $C^1, C^2, C^3, \ldots$  are, modulo translation in time and "space," independent and identically distributed, since each  $C^n$  is built in the part of the percolation structure not used by the preceding contact processes  $C^1, C^2, \ldots, C^{n-1}$ . Thus, the random variables  $\nu(1), \nu(2) - \nu(1), \ldots$  are also independent and identically distributed.

Each of the severed contact processes  $C^n$  reaches a rightmost point  $x_{\nu(n)}$  along the geodesic ray  $\gamma$ , and the last epoch of infection is terminated by a recovery mark  $\rho_n$ . Recovery marks in the percolation structure  $\mathcal{P}'$  are of two types: those that were recovery marks in the percolation structure  $\mathcal{P}$ , and those that were infection arrows in  $\mathcal{P}$ . Let N be the first of the contact processes  $C^n$  whose terminal recovery mark  $\rho_n$  is *not* an infection arrow in  $\mathcal{P}$  pointing from  $x_{\nu(n)}$  to  $x_{\nu(n)+1}$ . Note that N has the geometric distribution with parameter p. Define  $B_t$  to be the process obtained by concatenating the processes  $C^1, C^2, \ldots, C^N$ . Since the concatenations are all across infection arrows in  $\mathcal{P}$ , the process B(t) is dominated by A(t), that is,  $B(t) \subseteq A(t)$  for all  $t \ge 0$ . Consequently, the vertex  $x_{\nu(N)}$  is among the vertices infected by  $A_t$ , and so, in particular,

(0.5) 
$$u_n := P\{x_n \in A_t \text{ for some } t \ge 0\} \ge P\{v(N) \ge n\}.$$

Since  $\beta(\lambda) = \lim_{n \to \infty} u_n^{1/n}$ , it now follows, by Lemmas 1 and 2, that to prove Theorem 3' it suffices to prove the following.

PROPOSITION 3. The random variable v := v(1) is such that

$$\lim_{n \to \infty} P\{v \ge n\}^{1/n} = \beta(\lambda_*)$$

and

(0.7) 
$$\sum_{n=1}^{\infty} P\{\nu \ge n\} / \beta(\lambda_*)^n = \infty.$$

PROOF. Relation (0.6) was proved in [2]. The proof of relation (0.7) will be based on Schonmann's inequality (0.1) and the fact that  $\beta(\lambda) < \sqrt{d}$  implies  $\eta(\lambda) < 1$ . Set  $v_n = P\{v \ge n\}$ ; this is the probability that the contact process  $C^1$  started at the root and severed across the edge joining  $x_{-1}$  and  $x_0 = e$  ever infects the vertex  $x_n$ . Consider the event  $F_n$  that  $A_t$  ever infects  $x_n$ . On this event, there must be an infection trail from e to  $x_n$ , and this trail must visit  $x_{-1}$ , if at all, for a last time, and so, after passing through an infection arrow from  $x_{-1}$  to e for a last time, must then avoid the edge joining  $x_{-1}$  to e altogether. Let  $T(1), T(2), \ldots$  be the times of the infection arrows from  $x_{-1}$  to e in the percolation structure  $\mathcal{P}$ , and let T(0) = 0. Since these are stopping times for the contact process, and since they mark the occurrence times of a rate- $\lambda$  Poisson process, the strong Markov property implies that

$$u_n \le \sum_{k=0}^{\infty} v_n P\{x_{-1} \in A_{T(k)}\} \le v_n \lambda \int_{t=0}^{\infty} P\{x_{-1} \in A_t\} dt.$$

Conditional on the event  $x_{-1} \in A_t$ , the probability that  $x_0 \in A_{t+1}$  is at least  $\varepsilon(\lambda) := (1 - e^{-\lambda})e^{-1}$ , this being the probability that there is at least one infection arrow from  $x_{-1}$  to  $x_0$  and no recovery marks at  $x_0$  in a time interval of duration 1. Consequently,

$$\lambda \int_{t=0}^{\infty} P\{x_{-1} \in A_t\} dt \le \lambda \varepsilon (\lambda)^{-1} \int_{t=1}^{\infty} P\{x_0 \in A_t\} dt$$
$$\le \lambda \varepsilon (\lambda)^{-1} \int_{t=1}^{\infty} \eta^t dt$$
$$:= C' < \infty.$$

Thus, by (0.1),

$$v_n \ge u_n/C' \ge \beta(\lambda)^n/C'Cn$$
,

and inequality (0.7) now follows.  $\square$ 

## **REFERENCES**

[1] LALLEY, S. P. (1999). Growth profile and invariant measures for the weakly supercritical contact process on a homogeneous tree. *Ann. Probab.* **27** 206–225.

- [2] LALLEY, S. P. and SELLKE, T. (1998). Limit set of a weakly supercritical contact process on a homogeneous tree. *Ann. Probab.* **26** 644–657.
- [3] LIGGETT, T. (1985). Interacting Particle Systems. Springer, New York.
- [4] SCHONMANN, R. (1998). The triangle condition for contact processes on homogeneous trees. J. Statist. Phys. **90** 1429–1440.

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