A High Frequency Mutation Starts Sexual Reproduction in Volvox carteri

Bernd Weisshaar, Rainer Gilles, Regina Moka, and Lothar Jaenicke
Institut für Biochemie, Universität zu Köln, An der Botmühle 2, D-5000 Köln 1, Bundesrepublik Deutschland

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Pheromone-initiated sexual reproduction of Volvox carteri f. nagariensis is started by a "spontaneous" generation of sexual male spheroids at a rate of \(10^{-4}\). As previously suggested, these males are mutants (\(sex^e\)) which become independent of pheromone-controlled sex-induction. Unequivocal evidence for a mutative event now comes from the observation that the \(sex^e\) trait is transmitted through cell lineages during embryogenesis, thus confirming earlier morphological analysis. Mutation seems to be restricted to cell division. The high frequency of mutation points to a special enhancing mechanism for which the possibility of a genome rearrangement is discussed.

The multicellular green alga Volvox carteri f. nagariensis multiplies asexually by strictly ordered cleavages of its up to 16 gonidia [1] (Fig. 1). On sexual reproduction the gonidia of male and female strains differentiate to form spheroids with sperm packets and eggs, respectively. This is accomplished by a species-specific high-molecular pheromone released into the medium by disintegrating sperm packets [2]. Since the pheromone is only produced by sexual males, the obvious question is, how the first male becomes sexual. Occasionally, in asexual male cultures, sexual male spheroids appear which, on maturation, release sufficient inducer to turn sexual the spheroids of the next generation. R. C. Starr supposed these "spontaneous" males to be mutants in which a gene controlling the sexual development is affected [3, 4]. The frequency of their occurrence is \(1.8 \times 10^{-4}\) [5], approx. two orders of magnitude higher than that given for other mutations in Volvox [6]. It was therefore assumed that – in addition to mutations – entry into the sexual pathway is due to response either to an unknown environmental signal [5] or to accumulating inducer produced in trace amounts by asexual male cultures [7]. The usual test for a mutation, i.e. to check whether the trait can be established in a clonal culture, is not practicable in this case, since sexual males disintegrate completely, and an intersection with female strains and sperms from one male only is technically very difficult. Thus, the question which mechanism is responsible for "spontaneous" males is still open. In this communication we present evidence for an inducer-independent appearance of the first males in an asexual culture. Our data also indicate that the basic event is on the DNA level and occurs at the early stages of embryogenesis.

Materials and Methods

Culture conditions

The female strain HK-10 and the male strain 69-1b of Volvox carteri f. nagariensis Iyengar were obtained from R. C. Starr (Culture Collection of Algae, University of Texas). The algae were grown axenically in Volvox medium [8] at 28 °C in a light/dark rhythm of 16:8 h (10000 lux).

Screening for sex\(^e\) mutants

Asexual cultures were started with one young sperroid in phase I (see Fig. 1) in 10 ml Volvox medium. After 5 days algae containing the 3rd generation offspring (3300 ± 200) were scored under a dissecting microscope. By screening in phase IV, i.e. before hatching of the young individuals, we were able to deduce in which gonidium the mutation occurred. As an additional method, the inducer found in the medium of male cultures with sex\(^e\) mutants was measured: 1 ml medium of a six day test tube culture was inoculated with 3 HK-10 spheroids (phase IV) for bioassay [8]. Both methods gave corresponding results.

Micrographs were taken with a Leitz Dialux 20 microscope.

Results and Discussion

The rate of spontaneous induced males and females was estimated by scoring a large number of test tube
Fig. 1. Schematic representation of the life cycle of *Volvox carteri*. In the 48 h life cycle (16 h light/8 h dark) the developmental phases (I to V) are indicated [12]. I: growth of young spheroids; II: cell cleavages; III: inversion; IV: growth of adult spheroids; V: release of progeny. — In the lower part, spheroids are shown with only one tier of 4 gonidia rather than with the total of 16. In the upper part, asexual development of one gonidium is depicted up to the 64-cell stage. All the cell stages are drawn with the anterior pole (phialopore) on top. After the 8-celled embryo, all cleavages are shown in one quarter only. According to nomenclature of [13], the 8-celled embryo consists of an anterior (cells $1/3$) and a posterior tier (cells $2/4$). Since the next cleavage is also oblique, 4 extensively overlapping tiers (cells 1, 2, 3, and 4) arise. The following equatorial cleavage sets apart anterior and posterior derivatives. Cells $1a$, $1p$, $2a$, and $2p$ of the 32-celled embryo are gonidial stem cells. They cleave unequally (6th division) to large gonidial initials, labelled in the figure according to their stem cells, and small somatic precursor cells. The gonidial initials cleave unequally additional three times, while all other cells cleave until the total cell number of 2000 is reached. Then the embryo turns inside out (inversion). During inversion, the anterior pole of the embryo becomes the posterior pole of the young spheroid, also represented here. Note: Positioning of the gonidia in the young spheroid is the inverted one of the 64-cell stage. On sexual reproduction, unequal cleavage is delayed one stage (7th division) for the female strain; for the male strain, the last cleavage (7th or 8th division) is unequal for all cells.

Spheroids with 16 gonidia contain 4 quadrangular tiers of 4 gonidia each [9] (Fig. 1). The same holds for the positioning of the young spheroids after cleavage of the gonidia. Four gonidia lying in one quarter of the spheroid arise from one blastomere of the 4-celled embryo; two each of the gonidia have a common precursor cell at the 16-cell stage. Without exception, multiple sexual individuals are in positions which belong to one cell lineage (Fig. 1).

The existence and position of derivatives of spontaneous induced daughter spheroids are compelling arguments to postulate that the initial event occurs during embryogenesis of the preceding generation when the 16 gonidia, now forming the partially induced offspring, arise from a single gonidium. This explains why more than one sexual individual is formed by a single event. If during the 2-cell stage one blastomere is mutated, half of the resulting embryo (8 gonidia) will form sexual offspring in the next generation. If the same happens at the 4-cell stage, four, and at the 8- or 16-cell stage, two spon-
Table I. Data were obtained by screening 410000 male and 220000 female individuals, respectively. The mutation rate is $1.1 \times 10^{-4}$ for both sexes. The ratio of multiple to single spontaneous induced offspring is 16:52 or 1:3.25. Position of sexual daughter spheroids is given as indicated in Fig. 1.

<table>
<thead>
<tr>
<th>Number of sexuals per spheroid</th>
<th>Position of sexuals</th>
<th>Males 69-1 b</th>
<th>Females HK-10</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>8</td>
<td>2x (1a, 1p, 2a, 2p)</td>
<td>3</td>
<td>2</td>
<td>16</td>
</tr>
<tr>
<td>4</td>
<td>1a, 1p, 2a, 2p</td>
<td>2</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>1a, 1p</td>
<td>3</td>
<td>2</td>
<td>16</td>
</tr>
<tr>
<td></td>
<td>2a, 2p</td>
<td>2</td>
<td>1</td>
<td>16</td>
</tr>
<tr>
<td>1</td>
<td>1a</td>
<td>7</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td></td>
<td>1p</td>
<td>7</td>
<td>4</td>
<td>11</td>
</tr>
<tr>
<td></td>
<td>2a</td>
<td>11</td>
<td>4</td>
<td>15</td>
</tr>
<tr>
<td></td>
<td>2p</td>
<td>9</td>
<td>7</td>
<td>16</td>
</tr>
</tbody>
</table>

*a* In two adjacent quarters.

*b* In one quarter.

Fig. 2. Selected male and female spheroids containing multiple spontaneous induced offspring. Upper left: Female spheroids with 8 sexual daughters (containing ca. 32 eggs, each) in two adjacent quarters. Upper right: Male spheroid with 4 sexuals (containing a 1:1 ratio of androgonidia (=prospective sperm-packets) and somatic cells) in one quarter. Lower left: Female spheroid with 2 sexuals in the 1a and 1p position. Lower right: Male spheroid with 2 sexuals in the 2a and 2p position. As demonstrated in Table I, all other possible patterns are also found in both strains. (×50).

Taneous sexuals will be found. Single sexual daughter spheroids can only arise from an event later than the 16-cell stage. In each case the somatic cells are not visibly affected.

This explanation has important consequences for understanding sexual induction in *Volvox*:

(i) The basic event which establishes spontaneous sexuality is hereditary. Though we cannot exclude other possibilities, it is most likely that the event is on the DNA level, *e.g.* a mutation in a gene controlling sexual differentiation. Since chloroplast DNA is replicated prior to embryogenesis [10], nuclear DNA seems to be affected. We can thus consider the sexc trait in male strains as a stable mutation, as was suggested in [4].

(ii) The sex mutation is an exquisite marker of cell lineages during embryogenesis. We now have direct proof for the validity of the correlation between gonidial initials at the 64-cell stage and the gonidia in the adult spheroid (Fig. 1) which was only derived so far from microscopical observations [9]. Even scanning electron microscopy pictures have been incorrectly interpreted [11] with respect to the origin of the gonidia. Our data, in particular the dual spontaneous sexual daughters, unequivocally show that two tiers, *i.e.* cells 1 and 2 of the 16-celled embryo, are gonidial stem cells.

(iii) The reported mode of entry into the sexual pathway is independent of inducer. No pheromone can be detected even in concentrated medium of asexual male cultures (until a sexc male releases inducer). Inducer synthesis seems to be effectively repressed during the vegetative life cycle.
(iv) If the mutation is expressed directly during embryogenesis, mixed spheroids would arise, composed of an asexual part with gonidia and a sexual one with either sperm-packets or eggs. No such chimeras have ever been observed. During inducer-dependent induction, the irreversible decision, whether to enter the sexual or the asexual pathway, is made prior to the first cleavages of the gonidium [12]. The pattern of the sex mutants fits into this picture. Gonidial stem cells are unable to express the sex\(^c\) trait until they go through the critical phases before the next embryogenesis; then, the whole spheroid will become sexual. The time between the phase of decision and the first cleavage has to be very short, indeed; otherwise we would have found spheroids with 16 sexual offspring which will arise if the mutation occurs during this interval.

(v) From the relative frequency of the single and multiple sex\(^c\) mutants we may define the interval in which the mutation takes place. Assuming the probability of mutation as time-proportional we can deduce: During the first five divisions (1 h each) multiple sexuals are formed; a mutation from the 32-cell stage to the next generation cycle (43 h) will lead to a single sexual; this relative frequency (\(F_r = \text{number of cells} \times \text{time}\) should be \(22:80\) or \(1:3.1\). This proportion is not observed (Table I). If we make the restriction that the sex\(^c\) mutation occurs only during cleavage and take into account that gonidial initials divide until the 512-cell stage [13], we calculate for \(F_r \approx 22:80\) or \(1:3.6\), a frequency which is close to that found (1:3.25).

Hence we conclude that the sex\(^c\) mutation may be limited to cell division or is at least highly enhanced during that stage.

The primary event in sexual induction seems to be a high frequency mutation. The pheromone is a secondary synchronizing signal which enables a single sexual male to induce all Volvox carteri spheroids present in a volume of ca. 1 m\(^3\) [12]. Since a sex\(^c\) mutation is vital for regular sexual reproduction, it is very likely that a special mechanism had to be evolved to increase its frequency. In similar needs in other organisms, it is known that genome rearrangements are involved (references in [14]). An important example is the mating type switch in yeast [15] and the antigenic variability in Trypanosomes [16]. DNA rearrangements by movable transposons control gene expression during development of maize, sea urchin, Drosophila, and mice (review in [17]). So far we have no evidence for such rearrangements in Volvox, but normal base-pair exchanges cannot reasonably explain our results. Evaluation of the molecular basis of the sex\(^c\) mutation remains a challenging problem.

**Acknowledgements**

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Note added in proof:

We found a spontaneous pan mutant [5] in the 69-1b strain. In the presence of inducer, the androgomidia of this mutant develop asexually; however, sex\(^c\)-mutants of this strain show only sexual spheroids in all following generations, even in the absence of inducer.