AUTOTOMY AND INDUCED FRAGMENTATION IN THE RED CORAL (*CORALLIUM RUBRUM* L.)

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The red skeletal axis of the mediterranean sea-fan *Corallium rubrum* has been used to make jewellery since prehistoric times, and throughout the centuries, increasingly efficient methods have been devised to harvest natural banks. Today, colonies sufficiently large for jewellery manufacture have become rare and this led to study as how best to manage this marine resource, also taking into account the biology of the red-coral comes from a study conducted over 100 years ago (LACAZE-DUTHIERS, 1864), which describes the classical life cycle of colonial Anthozoans : vegetative growth of the colony branches by asexual replication of polyps, and formation of new colonies from sexually produced larvae. A much more recent study, albeit 20 years old, showed that the red coral has gonochoric larvae and colonies, and a gonadic cycle that is annual for male and biennial for female colonies. During attempts to rear the red-coral in the laboratory (RUSSO *et al.*, 1993), we became aware of a new type of reproductive process, which seems to implicate a reconsideration of the life cycle of the species: the production of new colonies by asexual reproduction of colonies are very rare in Cotcorallia. In the sea-fan *Plexawa* sp., localised constrictions in branches facilitates fragmentation by external disturbances (LASKER, 1984). Moreover, endogenous processes of fragmentation are known only for some soft-corals (Alcyonacea) (TURSCH 4 TURSCH, 1982) and for two species of fragmentation after stolonization in *Briareum asbestinum* (LASKER, 1983), and autotomy of branch ends in *Junceella fragilis*, whereby the coenenchyme is resorbed and the thin, not-living axial core is mechanically broken-off (WALKER & BULL, 1983). The red skeletal axis of the mediterranean sea-fan Corallium rubrum has been used to

In *Corallium rubrum* the autotomy of branch ends seems to occur with the complete resorption of tissues including the calcareous central axes. The process implies a particular dynamics of calcification which might explain the recently reported higher absorption of the colony (ALLEMAND & BENAZET, 1992). Such a specialized mechanism of asexual reproduction as autotomy is the evidence that fragmentation in red-coral is not an accidental consequence of external disturbances, but the result of natural selection with an adaptive meaning. In red-coral colonies, autotomy, occurred after removal from their natural environment to the laboratory and after drastic changes in water temperature and salinity. This indicates that, like in soft-corals (TURSCH & TURSCH, 1982), this asexual process was stress-induced. That is, under adverse conditions, the colonies asexually produce a sufficient number of propagules to ensure population survival. However, the marine environments colonized by red-coral are very stable and are not subject to such drastic changes. Therefore, reproduction by autotomy of branches seems to have a wider ecological meaning than supposed from data obtained in soft-coral. Fragmentation gives rise to colonies that are physiologically distinct but genetic variability recently found in *Corallium rubrum* (ABBIATI *et al.*, 1992, 1993). Furthermore, population spreading by short-living planulae and by passive benthonic dispersal of fragments probably enhances the treproduction faithfully replicates genotypes over many generations; this is because asexual reproduction faithfully replicates mutants which progressively build up in frequency over time (JACKSON & COATES, 1986). This model predictis that genetic relatedness and intensive inbreeding are extremely favourable for long-living organisms in stable environments, which is likely the case of red-coral. Production of new colonies by mechanical fragmentation, on the other hand, has been reported for many stony-corals (Esacorallia, Scleractiniaria) and seems t

the coral diversity and cover" (SHEPPARD, 1980). Our observations confirm for the red coral what is already well documented for reef building scleractinians: survival and growth of fragments are integral parts of the life cycle of the species and, therefore, are much more than an occasional event. This new reproductive feature should be further studied in order to elucidate a number of biological implications as well as to revaluate the effects of harvesting methods on natural banks. Dredging for red-coral is illegal in most Mediterranean countries because it is considered highly destructive for the bottom communities and scarcely efficient for the harvesting activity. However, because dredging, in which many coral fragments are left on the sea bottom, enhances fragmentation, it could play a not secondary role in the recovery of natural red-coral populations. Red-coral harvesting by scuba divers, on the other hand, is permitted because it is believed to be more efficient and less destructive than dredging (MATE, 1984). However, modern diving techniques are so sophisticated as to allow harvesting over almost the entire depth range of the species and, unlike dredging, in cavities. The selective harvesting by divers has a low degree of disturbance for the overall bottom community but, avoiding breakage and loss of commercially valuable parts, could reduce the role of fragmentation in the recovery of red coral population. It is certainly true that the present situation of overexploitation is mostly the effect of an excessive collecting effort in the time, but the attitude of divers to collect all the material available and to avoid leftovers is an aspect of the problem that should not be longer overlooked. An 18th-century Italian prince, believing that red coral colonies were "zoo-phytes", and thus capable of vegetative reproduction, ordered coral fragments to be scattered over the sea bottom to repopulate the natural banks. Our observations give the first evidence of the effective occurrenc

effective occurrence of a vegetative reproduction in red coral colonies and therefore the conceptual basis of that early experiment is surprisingly modern. **REFERENCES** ABBLATI M., SANTANGELO G. & NOVELLI S. 1993. *Mar. Ecol. Progr. Ser.*, 95 : 245-250 ALLEMAND D. & BENAZET S. 1992. *Rapp. Comm. int. Mer. Médit.* 33 : 27 HIGHSMITH R. C., 1980. *Mar. Ecol. Progr. Ser.*, 7 : 207-226 HUGES T. P., AYRE D. & CONNEL J. H., 1992. *Trends Ecol. Evol.*, 7 : 292-295 JACKSON J. B. C. & COATES A. G., 1986. *Phil. Trans. R. Soc. Lond.*, 8-313 : 7-22 LACXZE-DUTHIERS L. 1864. Histoire naturelle du corail. Baillère, Paris, 370 pp. LASKER H. R., 1983. *J. Exp. Mar. Biol. Ecol.*, 72: 157-169 LASKER H. R., 1984. *Mar. Ecol. Progr. Ser.*, 19 : 261-268 MATÉ P., 1984. *FAO Fish. Rep.*, 306: 79-81 RUSSO G.F., ZUPO V., PIRAINO S., ULLANICH L. & CICOGNA F. (1983). *In* : "Red Coral in the Mediterranean Sea : Art, History and Science", F. Cicogna and R. Cattaneo-Vietti (Eds) : 159-179 SHEPPARD C. R. C., 1980. *Mar. Rol. Biol. Biol.*, 70, 721-32 WALKER T. A. & BULL G. D., 1983. *Mar. Ecol. Progr. Ser.*, 12: 137-143 *Rapp. Comm. int. Mar. Mádit.* 34 (1005)

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