

Cryopreservation of Pacific Oyster larvae: a limited but promising survival!

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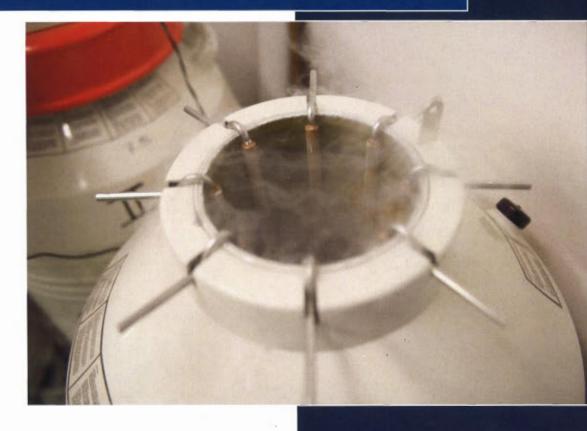
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2) Cryopreservation of Pacific Oyster larvae: a limited but promising survival!

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In aquatic animals, while sperm cryopreservation methods are well established, the preservation of larvae is still far from being achieved. Compared to fish species, mollusk oocytes are generally small and have a lower yolk content. This is favorable to larval cryopreservation. Because of the soaring role of hatcheries, Pacific oysterfarming requires larval cryopreservation.

Larvae were cryopreserved using the following conditions (Microdigitcool, IMV): larval dilution 1V:1V in cryoprotectant (10% ethylene glycol, 1% PVP, 200mM sucrose), 500µl straws, -35°C after 1h30 and then liquid nitrogen, thawing at 37°C. Survival was similar when larvae were cryopreserved at trochophore (13hpf: hours post fertilization) or D-larval stage (24hpf). A higher larval survival was observed when shortening the

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cryopreservation cycle from 1h30 to 45min. After a 3 year rearing period, the growth and reproductive performances of oysters, formely cryopreserved at larval stages, were similar to those observed for unfrozen ones.

The survival yields of thawed larvae (close to 0.5%) can be largely improved in the future but the high quality of thawed larvae shows that this technique can be applicable for the establishment of Pacific oyster larvae cryobanks.

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