Adaptive strategy of *Rapana venosa* (Gastropoda, Muricidae) in the invasive population of the Black Sea

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Introduction

Rapa whelk *Rapana venosa* (Valenciennes, 1846) is one of the brightest examples of global expansion of hydrobionts. The current original name of the species has a great deal of synonyms: *Purpura venosa* (Valenciennes, 1846), *R. thomasianna* (Crosse, 1861), *R. marginata* (Valenciennes, 1846), *R. pachylienis* (Grabau & King, 1928) and *R. pontica* (Nordsieck, 1969). The natural range of the rapa whelk is the water areas of the Yellow, Bohai, East China Seas, and the Sea of Japan. Molecular surveys revealed high genetic diversity among the natural populations and the fact that this is not associated with geographic distance between populations (Chandler et al., 2008; Kim et al., 2012; Shen et al., 2016). Populations of the Far Eastern Seas have been affected by overexploitation and have rapidly decreased (Yang et al., 2008).

For the first time beyond its range, *R. venosa* was recorded in the water area of the Black Sea in 1947. According to some data, this species was brought in on the bottoms of Soviet torpedo boats transferred from the Sea of Japan (Pereladov, 2013). From the Black Sea, the rapa whelk penetrated the Azov Sea already by 1950, and spread to the Sea of Japan, the genetic diversity of rapa whelk is high (Hd = 0.933, π = 0.002). In all the invasive populations of rapa whelk (Black Sea, European and North American), an extremely low level of haplotypic and nucleotide diversity was determined (Hd = 0.0, π = 0.0). Despite low values of genetic diversity, the invasive populations of rapa whelk are characterized by ecological success. We noted stable growth of populations, tolerance to diseases and parasites, effective reproduction, high fertility. This contradicts the main provisions of the population genetics theory of formation of edge of range populations, because usually only a small number of specimens of rapa whelk were introduced. The explanations of the unique condition of the invasive populations of rapa whelk, both from an ecological perspective (high resistance to fluctuations of hydrobiological factors and chemical pollution, absence of enemies, high fertility) and genetic perspective (high selective value of separate haplotypes settled in the new water areas) have not been confirmed.

We determined that a very important factor for the naturalization of the rapa whelk at low genetic diversity is the intra-species morpho-ecological divergence. In the Crimean water area of the Black Sea, *R. venosa* was found to have two morpho-ecological forms associated with the peculiarities of the development of the proportions of the shell. Study of successful invasive species would allow timely and adequate reaction to new cases of introduction.

We conducted molecular-genetic and morphological studies on the veined rapa whelk *Rapana venosa* (Valenciennes, 1846) in the Crimean waters of the Black Sea in order to determine possible reasons of the invasive success of this mollusk. Molecular-genetic tests were performed using COI gene; the surveyed samples reliably identified to *R. venosa* species. We compared the data on initial (from the Far-Eastern seas) and some invasive populations. In the natural conditions of the Yellow, East China Seas, and the Sea of Japan, the genetic diversity of rapa whelk is high (Hd = 0.933, π = 0.002). In all the invasive populations of rapa whelk (Black Sea, European and North American), an extremely low level of haplotypic and nucleotide diversity was determined (Hd = 0.0, π = 0.0). Despite low values of genetic diversity, the invasive populations of rapa whelk are characterized by ecological success. We noted stable growth of populations, tolerance to diseases and parasites, effective reproduction, high fertility. This contradicts the main provisions of the population genetics theory of formation of edge of range populations, because usually only a small number of specimens of rapa whelk were introduced. The explanations of the unique condition of the invasive populations of rapa whelk, both from an ecological perspective (high resistance to fluctuations of hydrobiological factors and chemical pollution, absence of enemies, high fertility) and genetic perspective (high selective value of separate haplotypes settled in the new water areas) have not been confirmed.

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caused significant damage to their fauna. Though there are no field observations of predators which consume rapa whelk in places of introduction, laboratory experiments demonstrated that some crabs can feed on small mollusks (Harding, 2003). In the natural range and newly colonised areas, *R. venosa* demonstrates stability of the main conchological characteristics, therefore it can be easily diagnosed (Pereyaslav, 2013), and the sexual dimorphism in the shell of rapa whelk is absent. In general, a paradoxical situation has developed: one of the most effective and successful invasive species throughout the wide and diverse area of its new range is characterized by extremely low genetic variability (low heterozygosity, extremely low haplotypic and nucleotide diversity, low inter-group subdivision). As a general rule, researchers try to explain the paradox by reproductive and ontogenetic peculiarities of rapa whelk: high fertility, presence of plankton larva (veliger), fast ontogenesis and maturation (Chandler et al., 2008).

Therefore the objective of the present work was the description of genetic diversity of *R. venosa* in the invasive population which lives on the shelf of the Crimean Peninsula of the Black Sea and the search for the reasons for the adaptive success of this mollusk.

**Materials and methods**

The mollusks were collected in 2018 in the Karantin Bay (Sevastopol) under the collectors of the mussel and oyster farm. A total of 10 mature specimens were selected for molecular-genetic analysis. As soon as the specimens were selected for molecular-genetic analysis. As soon as the principal components the vectors were calculated using covariances methods of analysis with Statistica 6.0 software pack. In the analysis of the principal components the vectors were calculated using covariances matrix. The length of vector was assumed to equal 1. The contours of the scatter formed by the individuals on PC1 and PC2 plane were considered to be the boundaries of ontogenetic channel, i.e. the projection of the area of the multidimensional space of characters in which the individual ontogenetic trajectories were located (Mina, 2001).

**Results**

Molecular-genetic analysis. In all 10 individuals of *R. venosa* from the Crimean water area, according to the fragment of 608 bp length, one COI haplotype corresponded to identical fragments from the Yellow, East China Seas and the Sea of Japan (KP136708, KP136707, KP136661, KP136659), massively present in the population of the Turkish water area of the Black Sea (KP136660, KP136661) and Chesapeake Bay of the USA (MH087553, EU250090, EU250111) (Fig. 1, Table 1).

![Neighbor Joining tree of phylogenetic relations of haplotypes of COI of *Rapana venosa* of the Crimean water area of the Black Sea](image_url)

**Fig. 1.** Neighbor Joining tree of phylogenetic relations of haplotypes of COI of *Rapana venosa* of the Crimean water area of the Black Sea (1p-10p), other invasive and initial populations of rapa whelk: haplotypes with letters KP – Turkish water area of the Black Sea, EU – Chesapeake Bay of the USA (except EU250077), the rest haplotypes, including EU250077, – Yellow, East China Seas, and the Sea of Japan; as external groups, *R. besoar* and *Ergalatax margaritica* are indicated; bootstrap supports are indicated in their branching nodes; below the figure the unit of measurement of the lengths of the branches is given – 0.02 nucleotides.

It is important that in rapa whelks from different populations the substitution of nucleotides occurred singularly. At the same time, the substitutions of the transition type definitely dominated. Transversions were usually much rarer (Table 2). The data obtained on the point mutation were confirmed by the analysis of the parameters of the genetic variability. Therefore, the highest values of the haplotype and nucleotide diversity, intra-group differentiation, amount of haplotypes, percentage of mutations were seen in the natural part of the range and the minimum values of these parameters – in the invasive populations (Table 3), the lowest values being observed in the Black Sea, and moreover particular in the Crimean water area.

**Table 1.** Parameters of genetic variability calculated using the DNAASP 5.10 software pack (Llibrado & Rozas, 2009). For comparison, we used the data on haplotypes of COI of the corresponding length of *R. venosa* taken from GenBank, NCBI (NCBI) from the Chesapeake Bay (Chandler et al., 2008; Aguilar et al., 2018), aquatoria of the Black Sea (Chandler et al., 2008; Kim et al., 2012; Sun & Yang, 2016). As the external group, we used ano-
Table 1

Matrix of distances (p) between the sequences of 607 pairs of nucleotides of COI genes of Rapana venosa

| No. | 1   | 2   | 3   | 4   | 5   | 6   | 7   | 8   | 9   | 10  | 11  | 12  | 13  | 14  | 15  | 16  | 17  | 18  | 19  | 20  | 21  | 22  | 23  | 24  | 25  | 26  | 27  | 28  | 29  | 30  | 31  | 32  | 33  | 34  |
|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| A   | 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00|
| B   | 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00| 0.00|

Table 2

Estimation of maximum likelihood of the pattern of nucleotide substitution by the sum of 607 pairs of nucleotides in rapa whelk Rapana venosa of the water area of the Black Sea

| Nucleotides | A   | T   | C   | G   |
|-------------|-----|-----|-----|-----|
| A           | –   | 2.53| 1.07| 11.83*|
| T           | 1.66| –   | 17.87*| 1.33  |
| C           | 1.66| 42.34*| –   | 1.33  |
| G           | 14.77*| 2.53| 1.07| –   |

Table 3

Parameters of genetic variability of the COI gene in some invasive and initial populations of rapa whelk Rapana venosa

| Populations                     | n   | S   | H   | Ht  | π   | K   |
|---------------------------------|-----|-----|-----|-----|-----|-----|
| R. venosa of the Crimean water  | 10  | 0.0 | 1   | 0   | 0   | 0   |
| area of the Black Sea           |     |     |     |     |     |     |
| R. venosa of the Turkish water  | 10  | 0.0 | 1   | 0   | 0   | 0   |
| area of the Black Sea           |     |     |     |     |     |     |
| R. venosa of the Chesapeake     | 10  | 0.0 | 1   | 0   | 0   | 0   |
| Bay of the USA                  |     |     |     |     |     |     |
| R. venosa of the Far-East Seas  | 30  | 40.0| 23  | 0.933| 0.002| 6.206|

Note: n – number of the studied specimens, S – number of polymorphic sites, h – number of haplotypes, Ht – haplotype diversity, π – nucleotide diversity, K – intra-group nucleotide differentiation.

Discussion

Study of nucleotide structure of the COI gene confirmed the conclusion drawn by previous researchers on the mono-morphisation in the populations of the rapa whelk throughout their recently colonised range (Chandler et al., 2008; Shen et al., 2016; Aguilar et al., 2018). Furthermore, if we take into account that the expansion of rapa whelk began from the Black Sea, then the haplotype uniformity of COI in the Crimean water body clearly demonstrates that for more than 70 years of living in the basin, no accumulation of diversity is observed. The data we obtained on the proportion of transitions and transversions indicate...
the stability of the COI gene in rapa whelk and also extremely low mutational potential of this gene in this species. This is especially clearly confirmed by comparison of the rapa whelk from the Turkish and Romanian water areas of the Black Sea.

Table 4

| Morphological characters | Component 1 | Component 2 |
|-------------------------|-------------|-------------|
| 1                       | -5.487      | -2.091      |
| 2                       | -1.216      | -0.518      |
| 3                       | 0.220       | -0.399      |
| 4                       | -1.957      | 0.496       |
| 5                       | 0.429       | 2.008       |
| 6                       | -0.976      | 0.633       |
| 7                       | -3.656      | 1.483       |
| 8                       | 0.547       | -0.403      |
| 9                       | -0.057      | 0.344       |
| 10                      | -1.199      | -1.429      |
| 11                      | -2.063      | -0.406      |
| 12                      | -1.407      | 0.428       |
| 13                      | 2.056       | -0.364      |
| 14                      | 0.199       | 0.239       |
| 15                      | 0.709       | -0.685      |
| 16                      | -0.391      | 0.263       |
| 17                      | 3.643       | -0.357      |
| 18                      | 5.087       | -0.917      |
| 19                      | 0.684       | -0.655      |
| 20                      | -1.871      | 0.906       |
| 21                      | 1.165       | 0.840       |
| 22                      | 0.830       | 0.861       |

At the same time, we should note that the population of rapa whelk of the Black Sea has entered the period of its stabilization characterized by periodic standardized fluctuations of the number (Odum, 1985). At the same time, despite the intense fishing, the population of the Black Sea is quite stable, the density has for many years remained within the range of 0.01–0.05 spec./m² (Revkov, 2009; Bondarev, 2010; Pereladov, 2013).

Many attempts have been made to explain such unique condition of the invasive populations of rapa whelk, both from an ecological perspective (active predation, absence of enemies, good food base, high fertility and fast maturation, plankton stage – veliger, high tolerance to the fluctuations of hydrological-hydrochemical factors and chemical pollutions) (Mann & Harding, 2003; Bondarev, 2010; Pereladov, 2013), and from the genetic perspective (high selective value of separate haplotypes settled in the new water areas) (Chandler et al., 2008; Kim et al., 2012; Zou et al., 2012). The explanation of the latter phenomenon is given from the positions of the principle of the founder and the bottleneck effect (Westphal et al., 2008; Wilson et al., 2009). Actually the only success of the application of this principle regarding the rapa whelk was the confirmation of the fact that most likely introductions and multiple invasions of this mollusk from various places to the Black Sea did not occur (Roman & Darling, 2007). However, such significant genetic monomorphisation of rapa whelk in the Black Sea was not anyhow explained. It was only noted that long term stability of this population is impossible (Sakai et al., 2001), whereas the analysis of ecological and morphological peculiarities of rapa whelk in newly colonized areas, first of all the Black Sea, allowed some authors to state that this population is ecologically flexible and will evolutionally prosper in future (Bondarev, 2010; Pereladov, 2013). The most significant argument for this statement was the analysis of the tendencies of change in growth and peculiarities of the habitus of the shell in the individuals of rapa whelk, representing both different populations and at the intra-populational level. It revealed the existence of several eco-morphological forms (i.e. “typical” and “dwarf”, the latter includes the “Thais” (Stramonita) form) in different regions of the Black Sea (Cesari et al., 1985; Giberto et al., 2006; Bondarev, 2010). The existence of different forms is supported by the data on the peculiarities of diet, and biotope of the rapa whelk. The drilling type of feeding is more characteristic for small forms. At the same, for the natural part of the range the presence of ecomorphs of rapa whelk is not that notable, which is also true for their size-age structure (Razin, 1934).

The determination of two trajectories of development (chreods of ontogenesis) completely confirms the fairness of the presumption of existence of morpho-ecological forms of rapa whelk in the Black Sea. Clearly seen is the formation of ‘scatter’ particularly by the second component, indicating the difference in the programmes of the development directly related to the formation of the proportions of the shell rather than its linear growth (Tissot, 1988). This is confirmed, on the one hand, by the existence of morpho-ecological forms of rapa whelk, and on the other hand connects their formation with the linear growth as presumed by the previous authors (Giberto et al., 2006; Bondarev, 2010). The observed phenotypic divergence implicating the intra-species diversification (Ward & McLennan, 2009; Berner et al., 2010) confirms one of the basic theses of the contemporary theory of bioinvasions that most notably the mechanisms of bioinvasions manifest during the trans-continental transfers (Elton, 1960) and are expressed first of all in the implementation of the adaptive compromise of the epigenetic theory of evolution (Shmalgauzen, 1982).

Conclusions

Thus, R. venosa, 70 years after its introduction to the Black Sea maintains its high abundance, ecological flexibility and significant tolerance to the environmental factors. By COI gene, it was confirmed that in the invasive populations of rapa whelk in different areas of the Global Ocean, its genetic variation is at extremely low level. At the same time, it is assumed that in the populations of the Crimean water area of the Black Sea, there are observed at least two ontogenetic programmes of the development which affect the morphometric proportions of the shell and perhaps manifest in the intra-species diversification of the rapa.
wehkl - formation of two morpho-ecological forms. Therefore, the phenomenon of significant ecological success of the rapa whelk in the invasive populations around the world at extremely low genetic variation is caused first of all by achieving adaptive compromise and formation of several trends of the development.

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