Parasites of the invasive Atlantic Blue Crab *Callinectes sapidus* and native crabs *Carcinus maenas* and *Afruca tangeri* in Merja Zerga Lagoon, Moroccan Atlantic coast

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Introduction

The Atlantic blue crab, *Callinectes sapidus*, is a well-established invasive species along the Mediterranean and the Atlantic coasts of Morocco (Oussellam et al., 2023). The present study aims to (1) explore parasites of the invasive *C. sapidus*, and the native crabs *Carcinus maenas*, and *Afruca tangeri* in the Merja Zerga Lagoon (Atlantic, Morocco); (2) test the enemy release hypothesis in the invasive blue crab, and (3) detect potential parasite spillover or spillback between native and non-native species.

Material and Methods

Overall, 36 Callinectes sapidus, 10 Carcinus maenas, and 43 Afruca tangeri were collected from the Merja Zerga Lagoon (Figure 1) between March and April 2025 and then dissected (Table 1). Hemolymph was microscopically screened for parasites belonging to Hematodinium and internal organs (hepatopancreas, gills, heart, stomach, and ganglion) were examined for other parasites. Genetic analysis of Hematodinium was performed using Sanger sequences from 13 C. sapidus, 3 C. maenas, and 17 A. tangeri. The haplotype network and the phylogenetic tree were created respectively by PopART using the Median-Joining algorithm and MEGA software, with the optimal model Kimura 2-Parameter and Neighbor-Joining method.

Results

Parasites identified are reported in **Figure 2**, their prevalences in **Figure 3** and the infected organs by species in **table 1**.

Genetic analysis, based on the ITS1 rDNA gene from *Hematodinium*, confirmed the sequences as being those of *Hematodinium perezi* (Lamkhalkhal et al., 2024). The haplotype network (Figure 4) and the phylogenetic tree (Figure 5) reveal that Moroccan *H. perezi* sequences are closer to those from England and Greece, but more distant from those in the USA.

Table 1: Morphometric measurements, parasites identified, and infected organs in *C. sapidus*, *C. maenas*, and *A. tangeri* collected from Merja Zerga Lagoon.

Species	CL (cm)		CW (cm)		W (g)		Parasites	Tissues / organs
	Min-Max	Mean±SD	Min- Max	Mean±SD	Min-Max	Mean±SD		
Callinectes	2.1 - 6.7	4.92 ± 0.48	4.1 - 15.3	10.36 ± 2.1	8.2 - 199.6	101.24±46.2	H. perezi	Hemolymph
sapidus							Digenean trematode	Gills - ganglion
							Microsporidian	Ganglion-stomach-hepatopancreas-heart
Carcinus	2.2 - 3.6	2.8 ± 0.3	3 - 4.8	3.78 ± 0.3	5.35-25.45	13.16 ± 3.7	H. perezi	Hemolymph
maenas							Digenean trematode	Gills- hepatopancreas- heart- ganglion
							Haplosporidian	Heart- ganglion
Afruca tangeri	1.1 - 3.7	2.36 ± 0.2	2 - 3.7	3.09 ± 0.3	6.4 - 30.8	17.56 ± 4.8	H. perezi	Hemolymph
							Haplosporidian	Ganglion- heart- gills

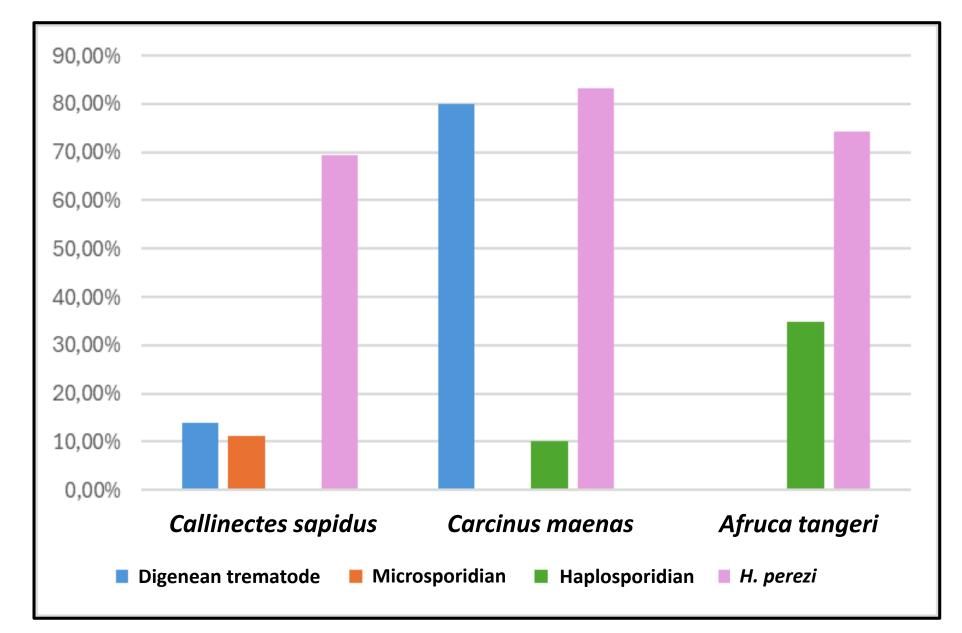


Figure 3: Comparison of parasite prevalence in Callinectes sapidus, Carcinus maenas, and Afruca tangeri.

Figure 4: Haplotype network of *H. perezi* (ITS1 rDNA) from the crab species studied and GenBank sequences, grouped by host and country. Circle size indicates haplotype frequency; vertical lines represent the number of mutations.

Afruca tangeri Morocco

Macrophthalmus abbreviates China
 Hemigrapsus takanoi China

Portunus trituberculatus China

Hemigrapsus penicillatus China

Conclusion

findings support the enemy Ihese hypothesis, indicating that *C.* release sapidus did not arrive in Morocco parasites carrying its instead but experienced parasite spillback from eastern Atlantic species.





References

❖ Oussellam et al. 2023. Bioinvasions Rec. 12: 403–416

❖ Lamkhalkhal et al. 2024. J. Mar. Sci. Eng. 12: 1045.

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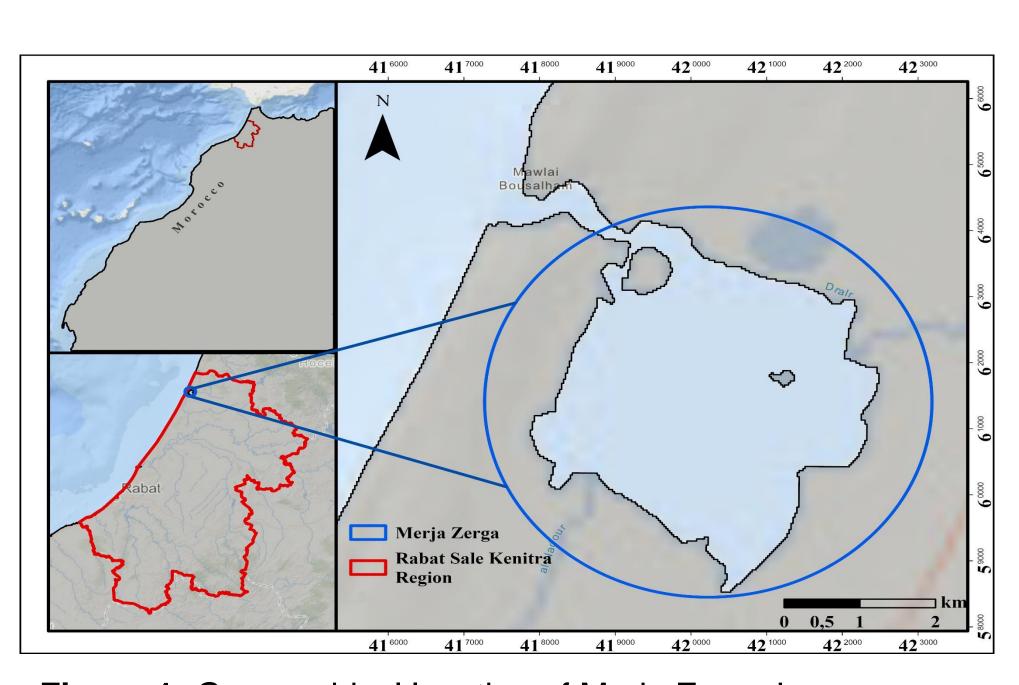
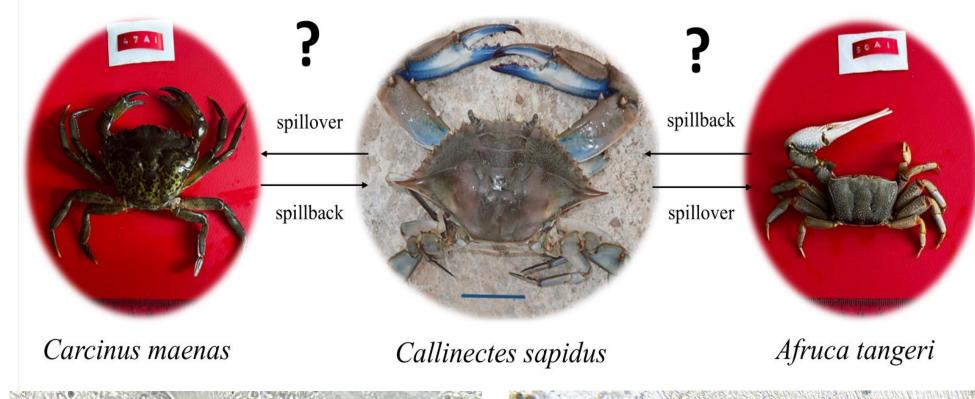
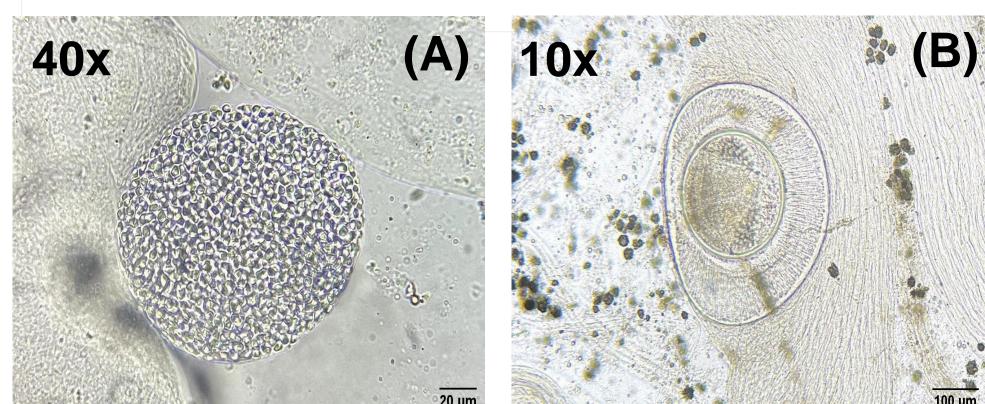
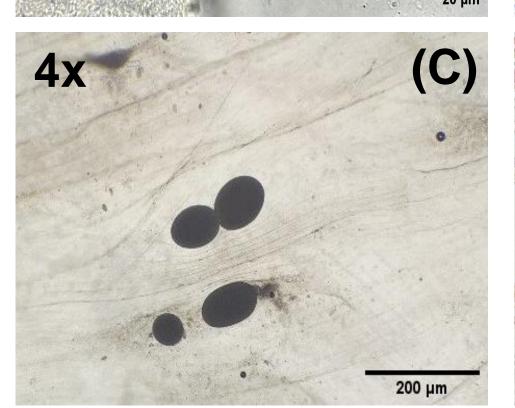


Figure 1: Geographical location of Merja Zerga Lagoon







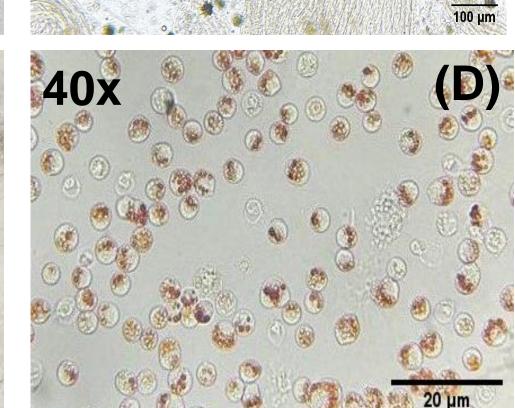


Figure 2: Haplosporidian (A) observed in the ganglion of *Afruca tangeri*, metacercaria (B) and microsporidian (C) in the ganglion of *Callinectes sapidus*; *Hematodinium perezi* (D) in the hemolymph of *C. sapidus*.

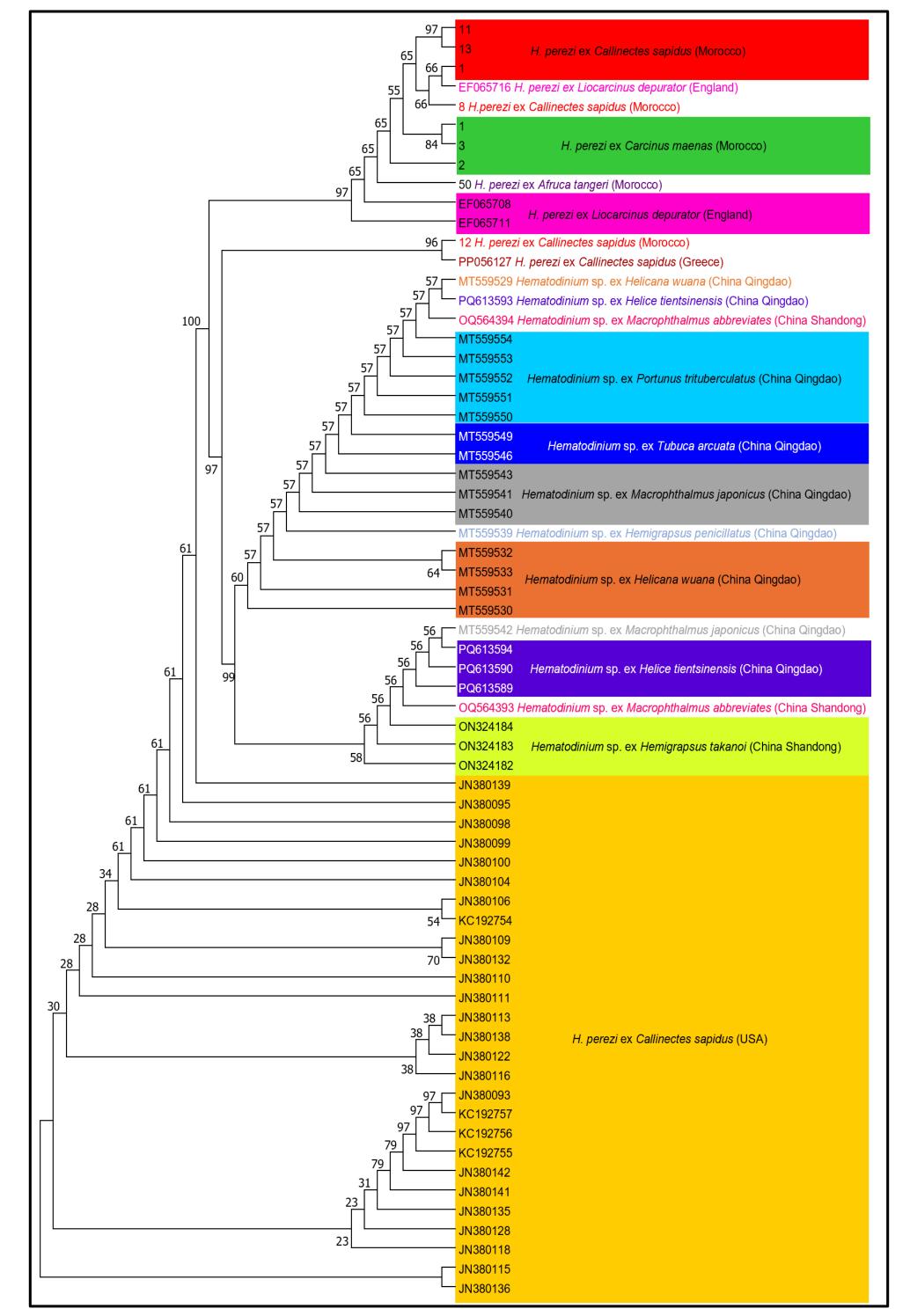


Figure 5: Midpoint-rooted phylogenetic tree of *H. perezi* (ITS1 rDNA) based on sequences from the studied crabs and GenBank data.