

Article

## Scale dependence of drilling predation in the Holocene of the northern Adriatic Sea across benthic habitats and nutrient regimes

Martin Zuschin\* , Rafał Nawrot , Markus Dengg, Ivo Gallmetzer, Alexandra Haselmair, Sandra Wurzer, and Adam Tomašových 

**Abstract.**—Predation has strongly shaped past and modern marine ecosystems, but the scale dependency of patterns in drilling predation, the most widely used proxy for predator–prey interactions in the fossil record, is a matter of debate. To assess the effects of spatial and taxonomic scale on temporal trends in the drilling frequencies (DFs), we analyzed Holocene molluscan assemblages of different benthic habitats and nutrient regimes from the northern Adriatic shelf in a sequence-stratigraphic context. Although it has been postulated that low predation pressures facilitated the development of high-biomass epifaunal communities in the eastern, relatively oligotrophic portion of the northern Adriatic shelf, DFs reaching up to 30%–40% in the studied assemblage show that drilling predation levels are comparable to those typical of late Cenozoic ecosystems. DFs tend to increase from the transgressive systems tract (TST) into the highstand systems tract (HST) at the local scale, reflecting an increase in water depth by 20–40 m and a shift from infralittoral to circalittoral habitats over the past 10,000 years. As transgressive deposits are thicker at shallower locations and highstand deposits are thicker at deeper locations, a regional increase in DFs from TST to HST is evident only when these differences are accounted for. The increase in DF toward the HST can be recognized at the level of total assemblages, classes, and few abundant and widespread families, but it disappears at the level of genera and species because of their specific environmental requirements, leading to uneven or patchy distribution in space and time.

Martin Zuschin, Rafał Nawrot, Ivo Gallmetzer, Alexandra Haselmair, and Sandra Wurzer. Department of Palaeontology, University of Vienna, Althanstrasse 14, 1090 Vienna, Austria. E-mail: [martin.zuschin@univie.ac.at](mailto:martin.zuschin@univie.ac.at), [rafal.nawrot@univie.ac.at](mailto:rafal.nawrot@univie.ac.at), [ivo.gallmetzer@univie.ac.at](mailto:ivo.gallmetzer@univie.ac.at), [alexandra.haselmair@univie.ac.at](mailto:alexandra.haselmair@univie.ac.at), [santwu@gmx.at](mailto:santwu@gmx.at)

Markus Dengg. Centre for Trace Element Analysis, University of Otago, Union Place West, 9010 Dunedin, New Zealand. E-mail: [denma131@student.otago.ac.nz](mailto:denma131@student.otago.ac.nz)

Adam Tomašových. Earth Science Institute, Slovak Academy of Sciences, Dúbravská cesta 9, 84005, Bratislava, Slovakia. E-mail: [adam.tomasovych@savba.sk](mailto:adam.tomasovych@savba.sk)

Accepted: 21 January 2022

\*Corresponding author.

### Introduction

Predation is one of the key factors shaping marine ecosystems today (Paine 1974; Connolly and Roughgarden 1999; Freestone et al. 2011; Harper and Peck 2016) and through the Phanerozoic (Vermeij 1977; Harper 2003) and is hypothesized to be responsible for secular changes in the guild composition and energetics of benthic ecosystems (Bottjer and Ausich 1986; Bambach 1993; McRoberts 2001; Aberhan et al. 2006). Drill holes provide direct evidence of predation and are thus the most often studied predator–prey interaction in the fossil record (Kowalewski 2002). Several authors

argued that the long-term increase in the predation intensity led to the replacement of epifaunal communities with sessile or poorly mobile species by infaunal benthic communities in soft-bottom habitats and that present-day epifaunal communities remain typical of habitats characterized by low predation pressure (Aronson et al. 1997; Gili et al. 2006; Bowden et al. 2011). However, it is unclear how species-level drilling frequencies (DFs) scale to community levels (Kelley and Hansen 2006) and how local estimates scale up to regional or global scales (Hoffmeister and Kowalewski 2001). Given the nonrandom preservation of depositional environments in the

rock record (Holland 2000; Smith 2007) and restricted outcrop area of many ancient successions, understanding these scaling relationships is crucial for interpreting time series of predator–prey interactions and distinguishing evolutionary factors (e.g., coevolution, escalation) from changes in the physical environment as their primary drivers (Leonard-Pingel and Jackson 2016).

First, global compilations demonstrate a distinct Phanerozoic increase in DF at the level of individual prey populations (e.g., Huntley and Kowalewski 2007). However, local-scale dynamics may not directly extrapolate to larger spatial scales: the effects of predator–prey interactions on the food webs can propagate to entire regional ecosystems, but geographic variability in the strength of interactions and source-sink or mass-effect metacommunity dynamics can reduce their importance (Holt and Hoopes 2005). Second, global or regional patterns in local-scale DF are frequently perceived at the level of the total assemblage (Kowalewski et al. 1998), which may reflect trends that are driven by varying relative abundances of prey species and differences in their susceptibility to drilling predation (Leighton 2002). However, most species or genera have constrained environmental and stratigraphic distribution, inhibiting assessments of species-level trends in DF across different habitats and over longer timescales. Aggregating species into higher taxa can thus average out uneven or patchy occurrences of species with narrow niches across samples and can reveal community-level ecological signals that cannot be perceived at the species level (Zuschin et al. 2017).

Here, we assess the congruence between local and regional (i.e., basin-scale) stratigraphic patterns in DFs at different levels of taxonomic resolution and their relationship to environmental proxies using sediment cores from the Holocene succession of the northern Adriatic Sea. This shelf exhibits a marked eastward decline in sea-surface primary productivity and sediment accumulation rate. McKinney (2003) and McKinney and Hageman (2006) suggested that communities dominated by epifaunal suspension feeders in the northern Adriatic Sea are limited to oligotrophic habitats and

that the decline in primary productivity is associated with a decline in predation pressure. In contrast, Zuschin and Stachowitsch (2009) showed that epifaunal communities in the northern Adriatic Sea are not restricted to oligotrophic settings, while predation intensity is similarly high as elsewhere in the Mediterranean Sea (Sawyer and Zuschin 2010). However, as ecological surveys and DFs observed in surface death assemblages are affected by anthropogenic impacts (Smith et al. 2018), assessments of natural predation pressure require analyses of sediment cores that capture Holocene communities inhabiting the northern Adriatic shelf since its flooding ~10,000 years ago. To rigorously assess DFs in the northern Adriatic Sea during the Holocene, high-resolution field data need to be placed within a robust, temporally constrained framework that integrates facies relationships and stratal architecture (Brett 1998; Leonard-Pingel and Jackson 2016; Patzkowsky 2017). Therefore, we evaluated DF patterns in a sequence-stratigraphic framework at different taxonomic resolutions and correlated them with environmental factors, represented by grain size and geochemical proxies. This is the first study on predator–prey interaction in the fossil record that uses such an integrative approach.

We show that stratigraphic trends in DF at regional scales can be confounded by habitat differences in DFs and can be biased by geographic differences in the thickness of systems tracts. We also demonstrate that the frequencies of drilling predation in the oligotrophic parts of the northern Adriatic Sea were not particularly low during the Holocene as previously suggested (McKinney 2007: p. 243) and were similar to average Cenozoic levels (e.g., Baumiller et al. 2006; Klomp maker 2009; Sawyer and Zuschin 2011; Chattopadhyay et al. 2016; Neely et al. 2021) (rather than to Paleozoic or early Mesozoic levels before the onset of the Mesozoic marine revolution; Kowalewski et al. 2005).

One exception is represented by a location that was characterized by seagrass development, indicating that structured biogenic habitats like seagrass meadows can significantly affect predator–prey interactions (e.g., Orth et al. 1984; Leonard-Pingel and Jackson 2016).

## Materials and Methods

**Sediment Cores.**—Piston cores 136–160 cm in length, penetrating the Holocene succession and representing the local scale, were taken in 2013 at four stations with different sediment types and at water depths between 21 and 44 m (Fig. 1). At each station, a separate core was taken to evaluate stratigraphic variability in grain size and chemical composition at the Institute of Marine Sciences in Venice. Ten increments were chosen from each core, three within the uppermost 10 cm (0–2 cm, 4–6 cm, and 8–10 cm) and seven evenly spaced along the rest of the core, at depths slightly deviating between cores from individual stations. Only eight samples in total were analyzed from the Venice core due to its shorter length. For each increment, grain size, total carbon, total organic carbon, total nitrogen, and carbonate content were measured from the sediment (Vidović et al. 2016; Gallmetzer et al. 2019). For faunal analysis, core increments, 4–5 cm in thickness, were sieved through a 1-mm mesh, and molluscan shells were counted and identified to the species level (Gallmetzer et al. 2019).



FIGURE 1. The location of the Venice, Brijuni, Piran 1, and Piran 2 sampling stations in the northern Adriatic Sea. Coordinates of stations are provided in Gallmetzer et al. (2019).

In total, 124 samples containing 70,783 molluscan individuals were analyzed. Two samples were excluded because their number of individuals fell below the threshold of 20 individuals, which resulted in a final dataset of 70,362 individuals. Muricid and naticid gastropods are abundant in our samples and are the most likely producers of predatory drill holes encountered (Bromley 1981; Kelley and Hansen 2003), but no attempt to distinguish between *Oichmus paraboloides* (made by naticids) and *Oichmus simplex* (made by muricids) was made. The mean percentage of naticids per sample is 0.9% with a maximum of 2.7%. For muricids, the mean is 0.7% and the maximum 3%. Naticidae (827 individuals) are strongly dominated by *Euspira nitida* (81.9%), followed by *Euspira macilenta* (15.6%), *Natica stercusmuscarum* (1.7%), and unidentified juvenile naticids (0.8%). Muricidae (495 individuals) are strongly dominated by *Hexaplex trunculus* (77.2%), followed by *Bolinus brandaris* (10.1%), unidentified juvenile muricids (6.5%), *Ocenebra erinaceus* (4.4%), *Ocenebrina helleri* (0.6%), *Trophonospis* sp. 1 (0.8%), and *Typhinellus labiatus* (0.4%). All naticids are apparently capable of shell drilling (e.g., Carriker 1981; Kabat 1990). For the muricids encountered in our samples, shell drilling has been documented for *H. trunculus* (Peharda and Morton 2006; Morton et al. 2007; Sawyer et al. 2009) and *B. brandaris* (Abidli et al. 2012; P. Vasconcelos personal communication August 2021). All juvenile muricids in our samples belong either to *Hexaplex* or *Bolinus*, and although they could not be assigned with confidence to either genus, they therefore still can count as drilling predators. *Ocenebra erinaceus* is also a facultative driller (Hancock 1960). We have no information on the predatory behavior of *O. helleri*, *Trophonospis* sp. 1, and *T. labiatus*, but together they only account for 1.8% of the total number of muricids. All naticids and virtually all muricids in our quantitative material can therefore count as obligatory or facultative shell-drilling predators.

To estimate DF, the total number of drilled shells was divided by the total number of bivalve, gastropod, and scaphopod individuals per core increment. Because only one valve in bivalves is usually drilled, DF was corrected using the equation of Bardhan et al. (2012):

$DF = DV / [(RV + LV) / 2 + A]$ , where DV is the total number of drilled valves, RV the total number of right valves, LV the total number of left valves, and A the number of articulated individuals. As there are only 23 samples with minimum 20 scaphopod shells, we perform class-level analysis only for bivalves and gastropods. Family-level analyses are performed for the eight most abundant families (Table 1), which together account for 64% of the individuals in the total dataset, with a range from 17.4% (Veneridae) to 2.7% (Tellinidae). These families also have a sufficiently high number of samples with minimum 20 individuals. Similarly, a species-level analysis was restricted to the eight most abundant species, which together account for 43% of the individuals in the total dataset, with a range from 10.8% (*Gouldia minima*) to 2.7% (*Striarca lactea*). Note that the families Corbulidae and Noetiidae are each strongly dominated by a single species; therefore, the family- and species-level analyses are almost identical. A genus-level analysis was not performed, because most genera are either monospecific or numerically strongly dominated by one species. Freshwater species, which occur mostly at the base of the cores, were excluded from the analyses.

*Core Geochronology and Stratigraphic Framework.*—The core chronologies are based on extensive dating by radiocarbon-calibrated amino acid racemization of one or two bivalve species occurring at high abundance at each station, with supplementary dates from other taxa (documented in our former studies, see below). For the two Piran cores, *G. minima* (316 shells) and *Corbula gibba* (232 shells) were dated with additional dates from *Arca noae* and *Ostrea* sp. (Mautner et al. 2018; Gallmetzer et al. 2019; Tomašových et al. 2019); for Brijuni, *Timoclea ovata* (305 shells) was used with additional dates from *Glycymeris*, *Aequipecten*, *Chlamys*, *Ostrea*, and a plant remnant (Schneidl et al. 2018; Gallmetzer et al. 2019); and for Venice, *Lucinella divaricata* (341 shells) was used with additional dates from *Aequipecten* (Gallmetzer et al. 2019).

The history of relative sea-level changes since the last glacial maximum in the northern Adriatic is characterized by a rapid deepening during the transgressive phase, followed by a slow

increase in accommodation space over the past 5000 years (Antonioli et al. 2007; Vacchi et al. 2016). Prograding highstand sediment wedges formed near river deltas (Po, Tagliamento, and Isonzo Rivers; e.g., Zecchin et al. 2015; Amorosi et al. 2019) and in bays (e.g., Koper Bay; Novak et al. 2020), whereas transgressive lags formed in current-swept and sediment-starved environments in the Gulf of Venice, in the Gulf of Trieste, and off Istria (Cattaneo et al. 2007; Trobec et al. 2017; Novak et al. 2019). In spite of this along-strike variability in sediment supply (Trincardi et al. 1994; Cătușeanu 2019), the timing of the maximum ingressions and the shift in the stacking from retrogradation to progradation are relatively isochronous in the whole region at millennial timescales. Although beyond the seismic resolution, Holocene sediments in our cores can be subdivided at the centimeter or decimeter scale of resolution into distinct lithologic units that can be correlated (on the basis of geochronological data) with the late-transgressive phase, the phase of maximum flooding, and the highstand phase (Gallmetzer et al. 2019). The starved locations were in the aggradational regime even during the highstand phase due to a slow increase in accommodation space (and locally even under an erosional regime), but the geochronological dating indicates that sediments in the uppermost decimeters of cores collected at Venice, Brijuni, and Piran are dominated by bioclastic (molluscan) particles produced over the past two millennia rather than by skeletal remains reworked from the transgressive sediments (Gallmetzer et al. 2019; Tomašových et al. 2019, 2022). These deposits thus represent thin condensed sediment veneers that are isochronous with thick prograding, highstand wedges in parts of the Northern Adriatic Sea characterized by high terrigenous sediment supply. Distinct shell beds at Piran and Brijuni formed by epifaunal bivalves are located in units that were deposited during the maximum ingressions (~5000–7000 years ago), in accordance with the predicted occurrence of such beds at the maximum flooding surfaces (separating the TST from the HST) due to the lack of dilution and higher productivity of epifauna under slow sedimentation rates (e.g., Kidwell 1988,

TABLE 1. The number of samples and abundance of prey classes, dominant prey families and species, and predatory families per systems tract at the four stations.

Station	Systems tract	Number of samples (n > 20)*	Prey Abundance											
			mollusks	bivalves	gastropods	scaphopods	Veneridae	Cardiidae	Rissoidae	Cerithiidae	Corbulidae	Nassariidae	Noetiidae	Tellinidae
Brijuni	IHST	5	905	509	393	3	90	112	117	115	38	7	19	1
	eHST	14	7442	3883	3539	20	1160	702	900	780	210	72	407	18
	MFZ	6	4834	2521	2310	3	941	307	869	343	72	68	525	7
	TST	8	5986	2986	2998	2	953	278	1276	378	75	129	869	4
Piran1	IHST	2	1799	1026	761	12	424	203	103	401	192	47	4	10
	eHST	2	4263	2188	2017	58	1096	428	419	1015	275	95	13	27
	MFZ	4	8918	4386	4273	259	1886	728	1050	1584	580	547	28	156
	TST	22	9422	5332	3757	333	1255	594	960	1231	1073	628	22	392
Piran2	IHST	2	3725	1941	1773	11	931	432	244	843	292	165	7	22
	eHST	2	3290	1576	1669	45	809	332	273	764	196	160	3	27
	MFZ	4	3719	1778	1866	75	753	294	318	733	221	324	7	108
	TST	24	10506	6383	3853	270	1550	799	736	1003	1315	908	18	386
Venice	IHST	7	958	719	228	11	75	135	35	52	30	20	1	72
	eHST	12	1480	1166	285	29	113	190	82	44	18	16	0	209
	MFZ	3	954	739	183	32	63	80	36	25	12	10	0	156
	TST	5	2161	1554	550	57	144	131	106	139	26	50	2	271
Total		122	70362	38687	30455	1220	12243	4346	7524	9450	4625	3246	1925	1866
Percentage			100	55.0	43.3	1.7	17.4	6.2	10.7	13.4	6.6	4.6	2.7	2.7
number of samples > 20 individuals		122	122	115	23	85	82	77	79	63	42	20	37	
% number of samples > 20 individuals		100.0	100.0	94.3	18.9	69.7	67.2	63.1	64.8	51.6	34.4	16.4	30.3	
*refers to mollusks														

Station	Systems tract	Number of samples (n > 20)*	Predator abundance											
			<i>Gouldia minima</i>	<i>Corbula gibba</i>	<i>Bittium latreilli</i>	<i>Bittium submamillatum</i>	<i>Nassarius pygmaeus</i>	<i>Timoclea ovata</i>	<i>Parvicardium scabrum</i>	<i>Striarca lactea</i>	Naticidae	Naticidae %	Muricidae	Muricidae %
Brijuni	IHST	5	16	38	21	69	7	48	9	19	0	0.00	4	0.44
	eHST	14	72	210	112	526	66	888	130	407	46	0.62	55	0.74
	MFZ	6	86	72	93	199	68	699	89	525	32	0.66	45	0.93
	TST	8	113	75	154	147	128	649	90	869	38	0.63	61	1.02
Piran1	IHST	2	391	192	275	100	47	4	124	4	28	1.56	5	0.28
	eHST	2	1015	275	549	378	95	6	207	13	76	1.78	14	0.33
	MFZ	4	1666	580	841	598	547	9	285	28	142	1.59	62	0.70
	TST	22	884	1072	372	745	628	11	147	22	96	1.02	76	0.81
Piran2	IHST	2	846	292	584	180	165	18	204	7	63	1.69	9	0.24
	eHST	2	748	196	517	176	160	13	153	3	56	1.70	25	0.76
	MFZ	4	623	221	326	332	323	3	127	7	71	1.91	31	0.83
	TST	24	1105	1310	449	433	903	11	221	18	142	1.35	80	0.76
Venice	IHST	7	4	30	24	10	20	7	121	1	4	0.42	17	1.77
	eHST	12	3	18	19	17	16	3	162	0	5	0.34	6	0.41
	MFZ	3	0	12	18	4	10	3	60	0	6	0.63	4	0.42
	TST	5	5	26	63	2	50	2	86	2	22	1.02	1	0.05
Total		122	7577	4619	4417	3916	3233	2374	2215	1925	827	1.18	495	0.70
Percentage			10.8	6.6	6.3	5.6	4.6	3.4	3.1	2.7	1.18	1.18	0.70	0.70
number of samples > 20 individuals		43	63	33	59	40	26	34	20					
% number of samples > 20 individuals		35.2	51.6	27.0	48.4	32.8	21.3	27.9	16.4					
*refers to mollusks														

1993; Kondo et al. 1998; Fürsich et al. 2021). High abundance of large epifaunal mollusks, high degree of skeletal alteration, and taphonomic feedback triggering high abundance of encrusters confirmed that the maximum flooding zone (MFZ) shell bed at Brijuni formed under limited clastic supply that was compensated by high carbonate production of mollusks, leading to temporally invariant long-term sediment accumulation (Tomašových et al. 2022). A similar, although less distinct shell bed with scallops and concretions marks the MFZ at the Venice station (Gallmetzer et al. 2019: fig. 7).

To compare long-term ecological trajectories among stations, core increments were assigned on the basis of grain-size, taphonomic, and paleoecological data to four intervals following the sequence-stratigraphic terminology of model III of Catuneanu et al. (2009), in which the highstand phase terminates before the base-level fall (Hunt and Tucker 1992): (1) a transgressive systems tract (TST) with transgressive lags and mixed assemblages formed by marine, brackish, and freshwater species; (2) an MFZ with densely packed molluscan shell beds and increments rich in concretions; (3) an early highstand systems tract (eHST); and (4) a late highstand systems tract (IHST). The shallower stations at Venice and Piran (21–23 m) were flooded ~9000–10,000 yr BP, and the deeper station at Brijuni (44 m) ~11,300 yr BP (Antonioli et al. 2007; Storms et al. 2008; Vacchi et al. 2016). Thus, all cores generally preserve similar durations of the Holocene record. However, they differ in the thickness of the TST and HST units: the MFZ is located at 90–105 cm at Venice and at 90–120 cm at Brijuni, but only at 16–35 cm at both Piran stations (Fig. 1). We note that the bases and tops of the MFZ shell beds and concretion horizons that separate the TST and the HST at all stations are also geochronologically well defined, effectively represent distinct stratigraphic surfaces. However, all intervals were subjected to intense bioturbation so that transitions between systems tracts are ultimately represented by gradual transitions at the scale of several centimeters or even decimeters. Uppermost increments deposited during the late highstand (effectively anthropogenic) phase (IHST) are separated

from increments deposited during the early highstand (IHST) by geochronological data and by the strong increase in organic and inorganic pollutants at most stations (Gallmetzer et al. 2019: fig. 10).

*Analyses.*—DFs between higher taxa, between stations, and between systems tracts were compared with the Wilcoxon rank-sum test and with the Kruskal-Wallis test (followed by the pairwise Wilcoxon test), using a posteriori Bonferroni correction for multiple comparison. To account for the unequal number of increments per sequence-stratigraphic unit at stations, two increments per station (i.e., the minimum number available) were randomly drawn from each systems tract, resulting in eight increments per unit in total. This subsampling procedure was repeated 10,000 times to obtain mean values and corresponding 95% confidence intervals (CIs) for regional-scale median DF per increment. Only increments with at least 20 individuals for each studied taxon were used in statistical analyses.

To explore the effect of scale on the identification of potential environmental correlates of DFs, the proportion of clay (PC), an indicator of hydrodynamic conditions, total organic carbon (TOC) and total nitrogen content (TN), both proxies for sediment organic enrichment, and CaCO<sub>3</sub> content of the sediment were used in the analyses. The PC in bottom sediments typically explains a large portion of variation in the composition of benthic communities (Sanders 1958; Rosenberg 1995), and in addition to water energy can be also affected by proximity of mud-sourcing river deltas and by bioturbation that determines mud erodibility (Rhoads and Boyer 1982; Brückner et al. 2021). At the regional scale, we estimated the effect of these environmental factors and of the relative abundance of the most likely predators (i.e., muricid and naticid gastropods relative to the total number of molluscan individuals) on the DF of the total assemblage and of the molluscan classes using generalized linear mixed models (GLMM) with station included as a random effect; for the stations we used generalized linear models (GLM) (O'Hara 2009; Bates et al. 2015). We also used generalized least squares (GLS; Pinheiro and Bates 2006) to assess whether per-station

analyses are affected by temporal autocorrelation, with first-order autoregressive correlation structure. TOC and TN were excluded from model fitting because they are strongly correlated with each other and with PC (Supplementary Figure 1). The GLMM was fit using the lme4 R-package (v. 1.-26; Bates et al. 2020). Statistical analyses were performed with the software package R (v. 4.0.3.). All data used in this study are shown in Supplementary Table 1.

## Results

DFs differ between localities and show distinct trends along individual sediment cores (Fig. 2). At the regional scale, the median per-increment DF in the total assemblage is 17%. DF is equally high in bivalves and gastropods (18%) and low in scaphopods (7%) (Fig. 3, Table 2).

A spatial comparison reveals that the median per-station increment-level DF of the total assemblage is similarly high at Brijuni (17%), Piran 1 (19%), and Piran 2 (22%), but is very low at Venice (7%). DFs for bivalves and gastropods range from 15% to 25% at Piran 1, Piran 2, and Brijuni, although in changing rank order (Fig. 4, Table 2). The very low median DF of the total assemblage at Venice is shared by both bivalves (6%) and gastropods (8%) (Fig. 5, Table 2).

At the local scale, a distinct trend emerges at most stations: DFs of the total assemblage are low in the TST and MFZ and high in the eHST and IHST. DFs increase from ~12% in the TST to ~25% in the HST at Brijuni, and from 16%–21% in the TST to 28%–30% in the HST at the Piran stations. At Venice, however, DFs are low across all systems tracts (Fig. 5). The stratigraphic increase in DF holds for bivalves and gastropods (Fig. 5) and for the families Rissoidae and Cerithiidae (Fig. 6). However, it is absent in most families, genera, or species (Figs. 6, 7).

At the regional scale, when data from all stations are considered together, the median DFs hardly differ between systems tracts at the level of the total assemblages (16%–20%) (Fig. 8, Table 2) and of bivalves (17%–19%), gastropods (15%–22%), and scaphopods (5%–12%)

(Fig. 8). Standardizing the number of increments per station and systems tract demonstrates, however, that DFs are significantly higher in the HST than in the TST and MFZ, both for the total assemblage and for individual classes (Fig. 8).

At the regional scale, the abundances of predators, PC, and  $\text{CaCO}_3$  significantly and positively correlate with DFs in the total assemblage (Fig. 9). The results are similar for bivalves and gastropods, although PC has no significant influence on DFs in bivalves and abundance of muricids has no significant influence on DFs in gastropods. In contrast, at the local scale of individual stations, the relationships between DFs and these environmental variables are divergent, mainly owing to between-station differences in the raw relationship between PC and  $\text{CaCO}_3$ . The Spearman rank relation between PC and  $\text{CaCO}_3$  is negative at both Piran stations ( $r = -0.38$ ,  $p = 0.28$  at Piran 1,  $r = -0.9$ ,  $p = 0.0009$  at Piran 2) and at Venice ( $r = -0.6$ ,  $p = 0.1$ ), but it is strongly positive at Brijuni ( $r = 0.77$ ,  $p = 0.01$ ). Differencing these variables leads to low and insignificant correlations between PC and  $\text{CaCO}_3$  at the scale of stations. However, when accounting for temporal autocorrelation in GLS analyses with single predictors, the contrast between the local and regional scales persists: the effects of PC on DFs are negative at both Piran stations (slope =  $-0.07$ ,  $p = 0.0001$  at Piran 1, slope =  $-0.03$ ,  $p = 0.09$  at Piran 2), positive at Brijuni (slope =  $0.04$ ,  $p = 0.08$ ), and negligible at Venice (slope =  $-0.003$ ,  $p = 0.7$ ). The effects of  $\text{CaCO}_3$  on DFs are consistently positive at all stations. Within-station GLM analyses with four predictors show that DFs in the total assemblage at Piran 1 increase with decreasing PC and with increasing abundances of muricids, while at Piran 2 they increase with PC,  $\text{CaCO}_3$ , and abundance of naticids. At Brijuni, DFs of the total assemblage increase with PC, and no significant relations are evident for DFs at Venice. These results are largely similar for bivalves and gastropods (Fig. 9, Table 3). Accounting for temporal autocorrelation in GLS with four predictors, DFs in the total assemblage increase with decreasing PC at Piran 1 and with increasing  $\text{CaCO}_3$  at Piran 2 and Brijuni. However, as mentioned earlier,

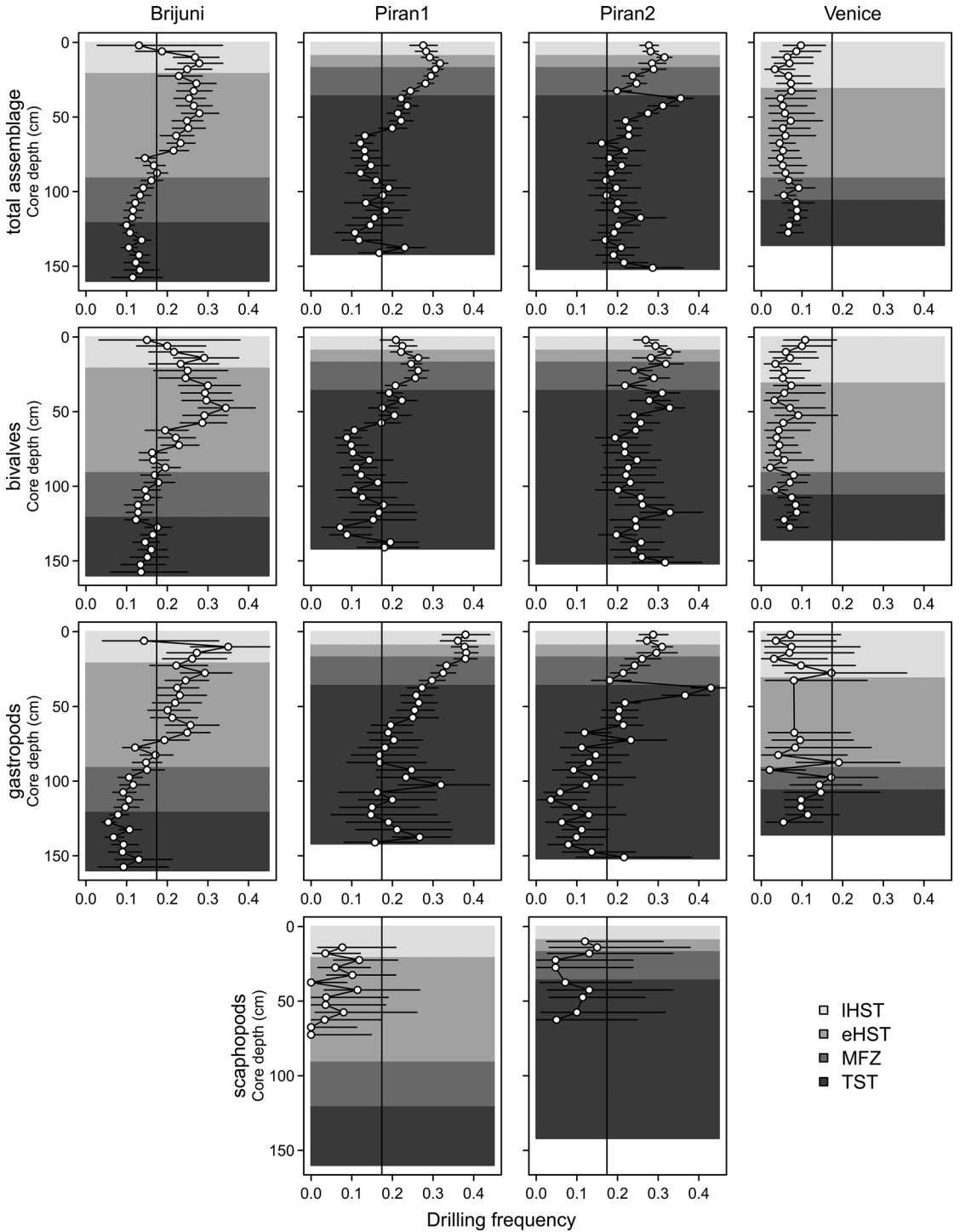


FIGURE 2. Stratigraphic changes in drilling frequencies (DFs) in the total molluscan assemblage and molluscan classes at the four sampling stations with 95% binomial confidence intervals (CIs); vertical lines represent the regional median DF of the total assemblage. Note that Piston cores differ in length between 136 and 160 cm. eHST, early highstand systems tract; IHST, late highstand systems tract; MFZ, maximum flooding zone; TST, transgressive systems tract.



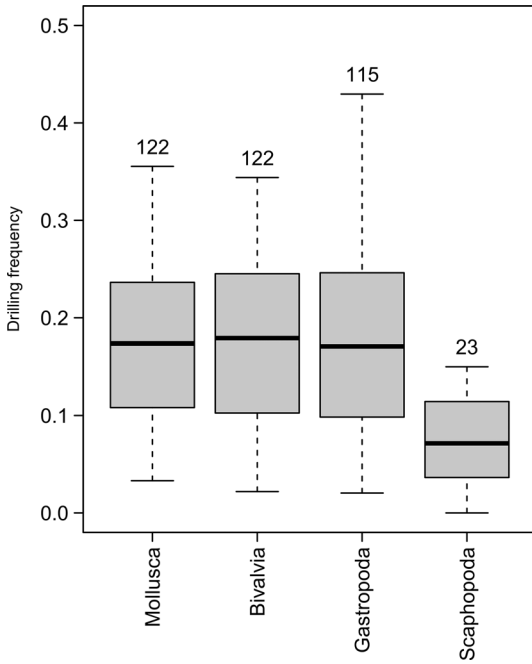


FIGURE 3. Box plots of drilling frequencies (DFs) of the total assemblage and the three classes at the regional scale. Numbers above box plots refer to number of samples with at least 20 individuals.

the within-station collinearity between PC and CaCO<sub>3</sub> affects these analyses and hides the observation that DFs decline with increasing clay content at the Piran stations, whereas they increase with increasing clay content at Brijuni.

**Discussion**

*Differences in DF between Regional and Local Scales.*—Our results demonstrate that spatial heterogeneity in DFs is very high at local scales and between taxa, making extrapolations to entire communities and larger spatial scales problematic. At the local scale, assemblage-level DFs tend to increase during the postglacial sea-level rise (from the TST to HST) at three out of four stations, but regional-scale comparisons suggest that the median DF remained stable within the basin throughout the Holocene. This pattern can be explained by strong habitat variability within systems tracts typical for higher-order sequences like the one studied here (e.g., with the development of seagrass communities at Venice station and level-bottom communities at other stations

TABLE 2. Results of statistical tests for significance of differences in drilling frequencies (DFs). Bold values indicate significance level < 0.01; underlined values indicate significance level < 0.05.

A. Kruskal-Wallis test for differences of DFs for molluscan classes between stations and systems tracts			
	Chi-squared	df	p-value
Classes	29.377	2	<0.001
Stations_Mollusks	67.99	3	<0.001
Stations_Bivalves	79.445	3	<0.001
Stations_Gastropods	38.309	3	<0.001
Systems tracts_Mollusks	0.32033	3	0.956
Systems tracts_Bivalves	0.74315	3	0.863
Systems tracts_Gastropods	3.6923	3	0.297
Systems tracts_Scaphopods	4.094	2	0.129
B. The p-values from pairwise Wilcoxon test, using a posteriori Bonferroni correction for multiple comparison			
Mollusks	Brijuni	Piran1	Piran2
Piran1	1		
Piran2	<u>0.031</u>	0.242	
Venice	<0.001	<0.001	<0.001
Bivalves	Brijuni	Piran1	Piran2
Piran1	0.505		
Piran2	0.001	<0.001	
Venice	<0.001	<0.001	<0.001
Gastropods	Brijuni	Piran1	Piran2
Piran1	0.001		
Piran2	1	<u>0.014</u>	
Venice	0.002	<0.001	<0.001

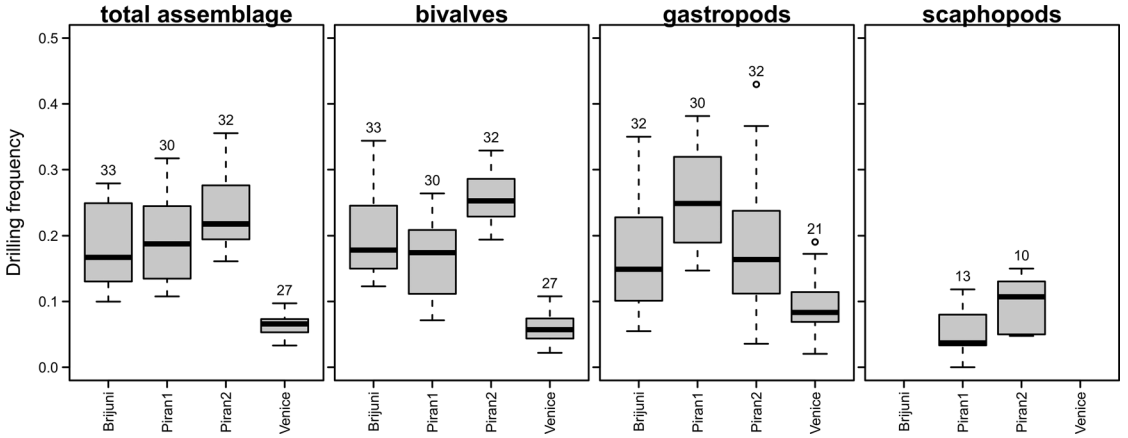


FIGURE 4. Box plots of drilling frequencies (DFs) of the total assemblage and the three classes at stations. Numbers above box plots refer to number of samples with at least 20 individuals.

at similar depths; Scarponi and Kowalewski 2007; Zuschin et al. 2011) and by the differences in thickness of systems tracts between the individual stations.

Spatial distribution and abundance of both predator and prey taxa vary between habitats (e.g., Sawyer and Zuschin 2010; Leonard-Pingel and Jackson 2016), which in turn shift laterally through time in response to changes in sea level and available accommodation space (Holland 2000), producing systematic changes in facies—and consequently DFs—observed at stratigraphic sections. As different stations within the basin differ in their environmental histories and thicknesses of systems

tracts, the local trends in DF can be averaged out regionally and thus do not necessarily scale up to the entire basin.

Such distinct stratigraphic patterns in facies and unit thickness are evident among the studied cores. For example, owing to earlier marine flooding, the HST at Brijuni (fining-upward trend) and Venice (sandy throughout) is expanded, in contrast to the condensed, coarse-grained HST at Piran 1 and 2 (Gallmetzer et al. 2019). Consequently, the HST is represented by many increments with low DFs at Brijuni and especially Venice and few increments with consistently high DFs at Piran. Pooling data from these stations together without accounting for

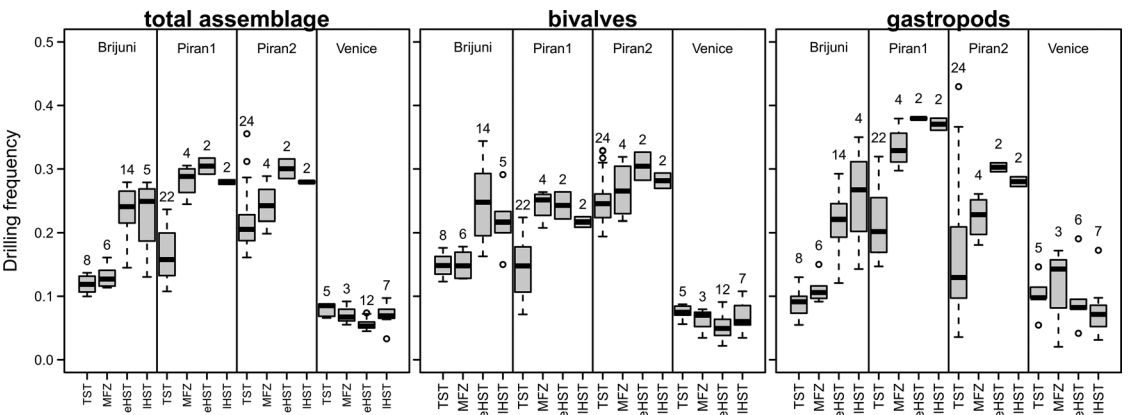


FIGURE 5. Box plots of drilling frequencies (DFs) in systems tracts at stations (i.e., local scale) for the total assemblage and molluscan classes. Numbers above box plots refer to number of samples with at least 20 individuals. eHST, early highstand systems tract; IHST, late highstand systems tract; MFZ, maximum flooding zone; TST, transgressive systems tract.

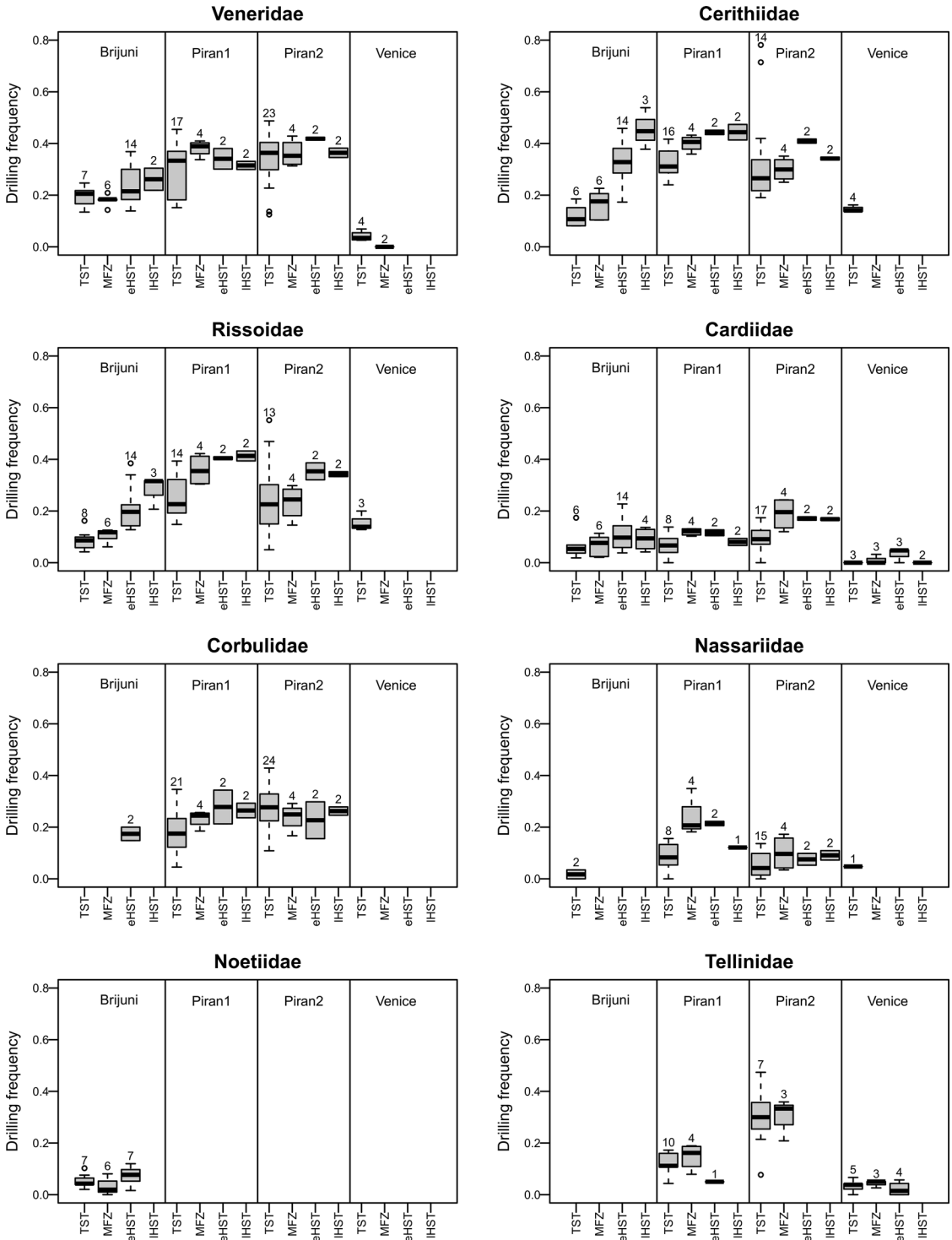


FIGURE 6. Box plots of drilling frequencies (DFs) for the eight most abundant families in systems tracts at stations. Numbers above box plots refer to number of samples with at least 20 individuals. eHST, early highstand systems tract; IHST, late highstand systems tract; MFZ, maximum flooding zone; TST, transgressive systems tract.

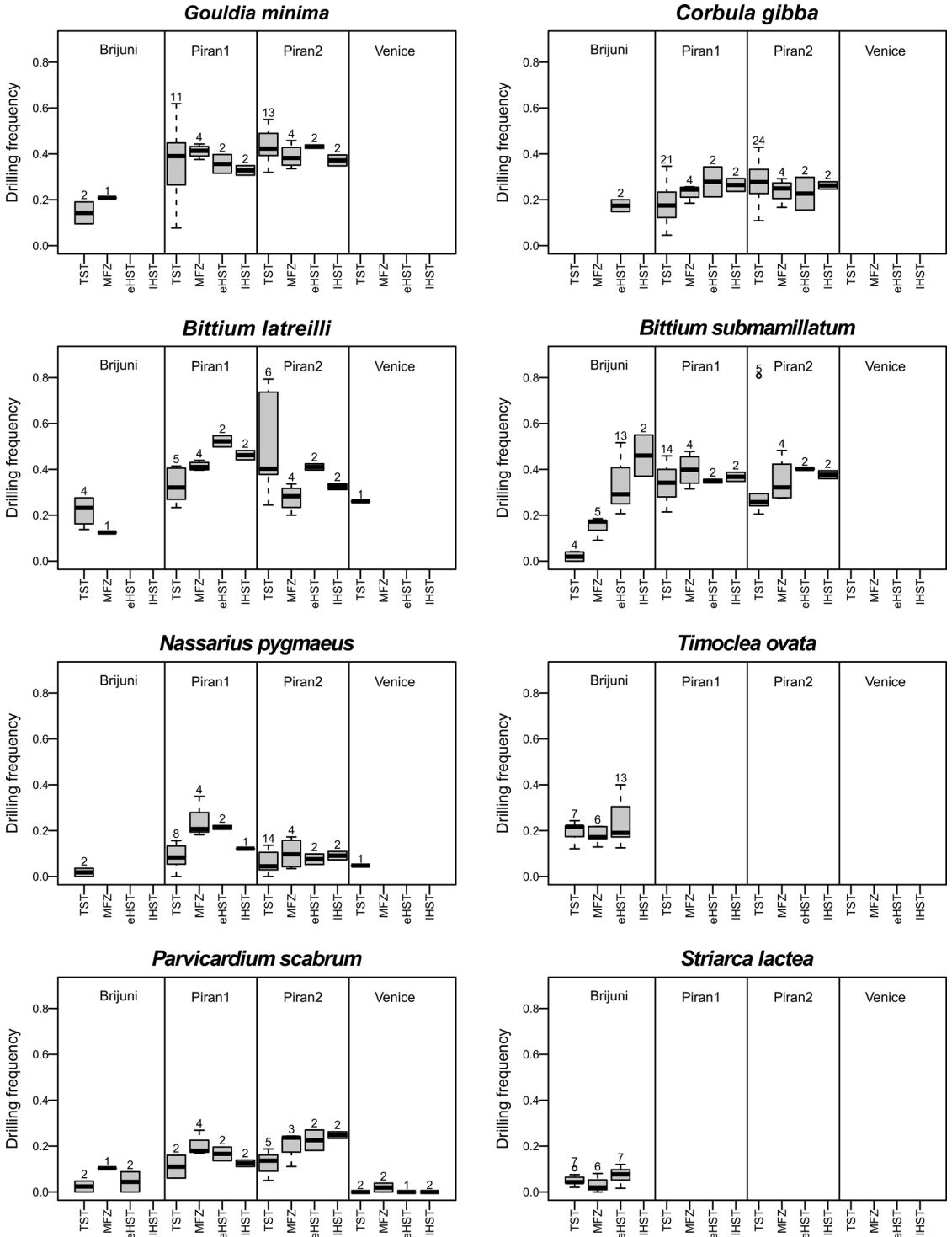


FIGURE 7. Box plots of drilling frequencies (DFs) for the eight most abundant species in systems tracts at stations. Numbers above box plots refer to number of samples with at least 20 individuals. eHST, early highstand systems tract; IHST, late highstand systems tract; MFZ, maximum flooding zone; TST, transgressive systems tract.

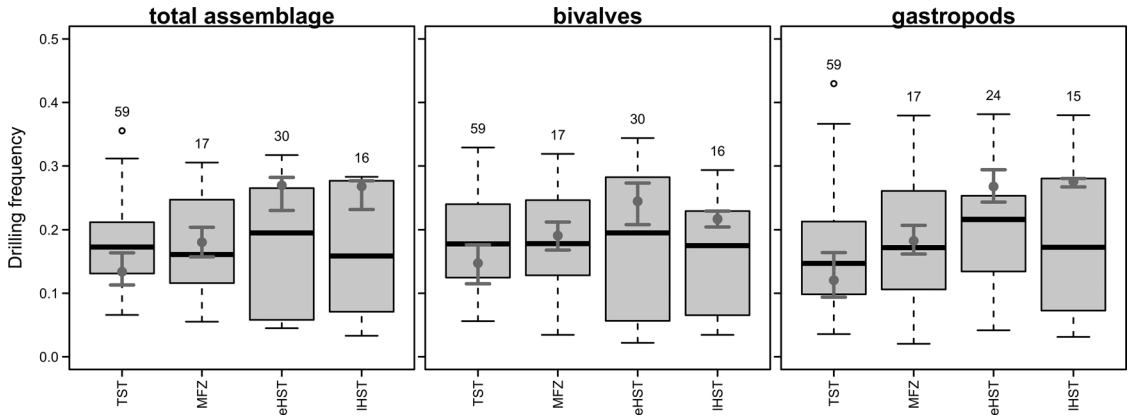


FIGURE 8. Box plots of drilling frequencies (DFs) in systems tracts at the regional scale for the total assemblage and molluscan classes. Results are shown for raw data (box plots) and after resampling to account for different number of samples per systems tracts at individual stations (mean value and 95% confidence intervals [Cis] across 10,000 iterations). Numbers above box plots refer to number of samples with at least 20 individuals. eHST, early highstand systems tract; IHST, late highstand systems tract; MFZ, maximum flooding zone; TST, transgressive systems tract.

the differences in numbers of increments dampens the regional DF. Conversely, the large number of increments combined with relatively high DFs in the stratigraphically expanded TST at the two Piran stations results in higher regional DF. Therefore, even though each station is represented by a similar number of equally spaced samples (Figs. 1, 2), the regional patterns are still biased by the variable thickness of systems tracts and thus by the number of samples representing each stratigraphic unit. As a result, local and regional stratigraphic patterns in DFs appear decoupled. However, a consistent temporal trend of increasing DFs emerges at both scales when the same number of samples from each station and systems tracts is used (Fig. 8). Therefore, integration of information on sequence-stratigraphic architecture and facies changes at individual locations/sections across a basin provides the most useful framework for tracking DF patterns at regional scales.

*The Advantages of Analyzing DF on Assemblage and Higher-Taxon Levels.*—Our results suggest that the regional within-sequence trends in DFs and along environmental gradients manifest primarily at higher taxonomic levels or at the level of whole assemblages, not at the level of species or genera. Lower taxa are typically confined to specific habitats, which vary in their geographic extent through time. Therefore, most species, genera, and even families

do not range through the studied Holocene sequence at sufficient abundance to allow meaningful comparison of DF patterns across the basin or between systems tracts.

Abundances of predators and environmental factors do not correlate with local DFs in a consistent way. At the regional scale, however, DF largely increases with increasing PC (and correlated increase in organic enrichment and/or decrease in hydrodynamic energy driven by the increase in water depth), with increasing predation pressure (as reflected in the relative abundance of naticid and muricid gastropods), and with increasing  $\text{CaCO}_3$  content. These relationships suggest that over the timescales of thousands of years, in which the regional species pool remained largely stable, an increase in organic enrichment and an increase in water depth enhances predator–prey interactions in the metacommunity (e.g., Menge and Sutherland 1987; Leibold 1989).

*No Evidence for Unusually Low Predation Pressure in the Northern Adriatic Sea.*—The northern Adriatic Sea was considered to be an area of overall unusually low predation pressure, a background condition that would allow sessile epibenthos to flourish in the oligotrophic waters of the eastern part of this basin (McKinney and Hageman 2006; McKinney et al. 2007). By pooling all increments at the regional scale, we show, however, that DFs during the

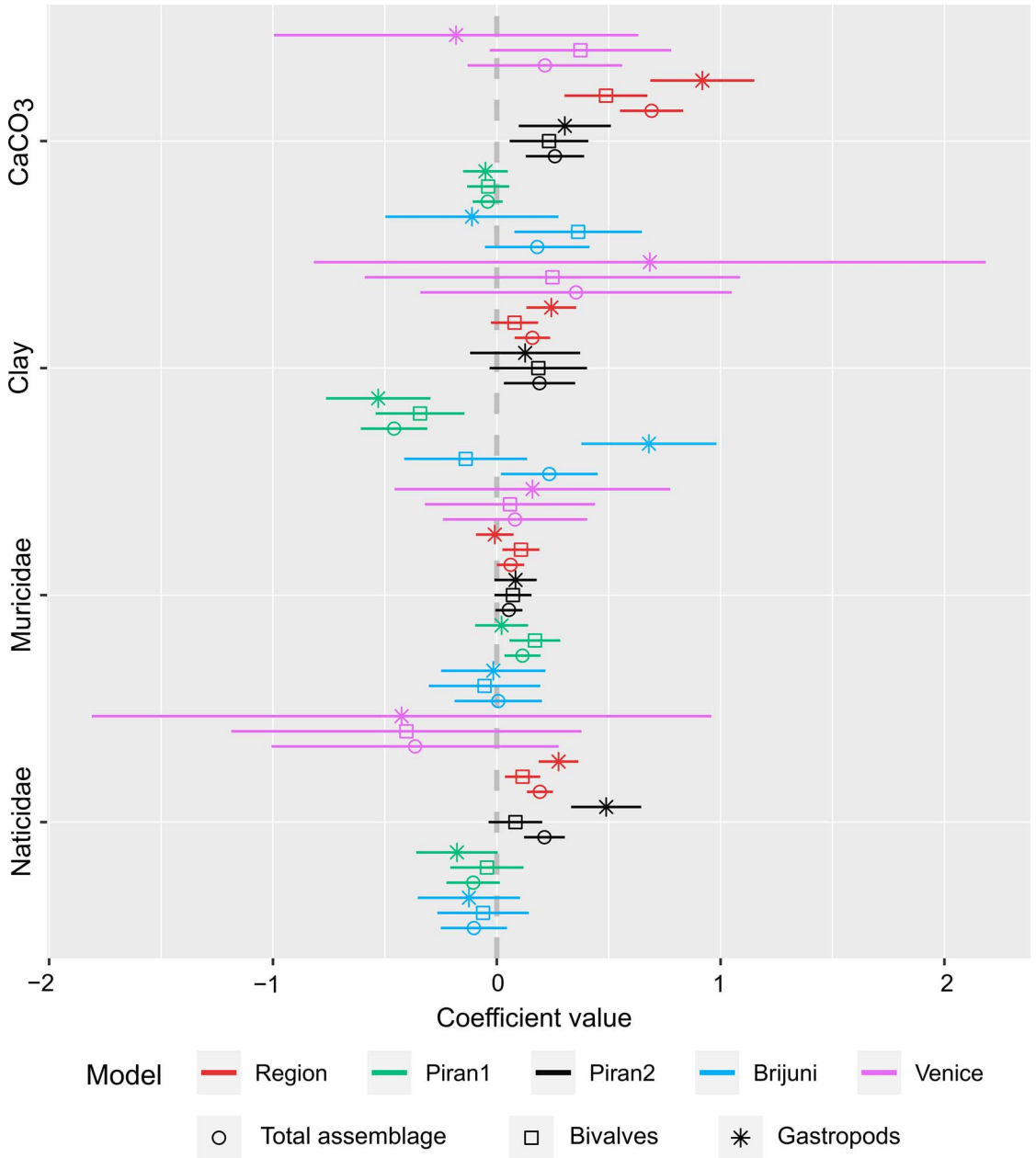


FIGURE 9. Coefficient plot showing results of the fitted generalized linear mixed models (GLMM) and generalized linear models (GLM) for the drilling frequencies (DFs) in the total assemblage and classes in the region and at each station. Effect sizes for predictors are shown with 95% confidence intervals (CIs). For significances and  $R^2$  of the regional GLMM, see Table 3.

Holocene have remained at levels very typical for warm-temperate seas in the Cenozoic (Yochelson et al. 1983; Kelley and Hansen 2003). Similar to other regions, DFs are high for bivalves and gastropods and significantly lower for scaphopods. Our results demonstrate

that DF in the northern Adriatic varies between and within systems tracts, generally increasing toward deeper habitats. Previous studies targeting surface death assemblages documented high environmental variability in DF in that basin as well. For example, Huntley and

TABLE 3. The fitted generalized linear mixed models (GLMM) and generalized linear models (GLM) for the total assemblage and molluscan classes at regional and local scale. Bold values indicate significance level < 0.01, underlined values indicate significance level < 0.05. R2m, marginal R squared; R2c, conditional R squared; AIC = Akaike information criterion.

Molluscs	Region GLMM	Piran 1 GLM	Piran 2 GLM	Brijuni GLM	Venice GLM
Intercept	<b>-1.5829</b>	<b>-1.2680</b>	<b>-1.1189</b>	<b>-1.4143</b>	<b>-2.6187</b>
Naticidae	<b>0.1925</b>	-0.1055	<b>0.2130</b>	-0.1022	-0.3648
Muricidae	<u>0.0617</u>	<b>0.1148</b>	0.0542	0.0064	0.0815
Clay	<b>0.1592</b>	<b>-0.4588</b>	<u>0.1909</u>	<u>0.2346</u>	0.3540
CaCO <sub>3</sub>	<b>0.6916</b>	-0.0409	<b>0.2598</b>	<u>0.1807</u>	0.2152
AIC	382.6	91.588	91.775	78.867	44.036
Theoretical R2m	0.1346				
Theoretical R2c	0.2667				
Bivalves	Region GLMM	Piran 1 GLM	Piran 2 GLM	Brijuni GLM	Venice GLM
Intercept	<b>-1.6055</b>	<b>-1.4545</b>	<b>-0.9936</b>	<b>-1.3528</b>	<b>-2.7352</b>
Naticidae	<b>0.1153</b>	-0.0439	0.0834	-0.0612	-0.4041
Muricidae	<b>0.1078</b>	<b>0.1705</b>	0.0722	-0.0546	0.0590
Clay	0.0794	<b>-0.3428</b>	0.1856	-0.1390	0.2486
CaCO <sub>3</sub>	<b>0.4879</b>	-0.0385	<b>0.2334</b>	<u>0.3634</u>	0.3737
AIC	317.5	87.605	84.529	71.24	44.388
Theoretical R2m	0.0681				
Theoretical R2c	0.2133				
Gastropods	Region GLMM	Piran 1 GLM	Piran 2 GLM	Brijuni GLM	Venice GLM
Intercept	<b>-1.4176</b>	<b>-0.9967</b>	<b>-1.3831</b>	<b>-1.5374</b>	<b>-2.1560</b>
Naticidae	<b>0.2761</b>	-0.1780	<b>0.4888</b>	-0.2006	-0.4253
Muricidae	-0.0089	0.0214	0.0837	-0.0900	0.1594
Clay	<b>0.2435</b>	<b>-0.5297</b>	0.1267	<b>0.8186</b>	0.6837
CaCO <sub>3</sub>	<b>0.9183</b>	-0.0506	<b>0.3041</b>	-0.2793	-0.1819
AIC	330.4	69.342	75.446	63.429	30.951
Theoretical R2m	0.2331				
Theoretical R2c	0.3461				

Scarponi (2015) found that DF is highest in sediment-starved environments north of the Po delta, while it is considerably lower in high-sedimentation environments south of it. Sawyer and Zuschin (2010) demonstrated that DFs in subtidal environments of the Isonzo delta and Bay of Panzano were of typical modern levels, while those from tidal flats were very low. However, subtidal habitats in the northeast Adriatic Sea, now inhabited by abundant epifaunal suspension feeders at Piran and Brijuni, have consistently high DFs (>25%), contradicting the hypothesis that low sea-surface primary productivity typical for these areas is associated with reduced predation intensities. At the same time, our study also suggests the very low DFs can be a persistent feature in subtidal areas with seagrass development (see following section). However, this pattern only further underscores high spatial

variability in drilling predation pressure frequently observed at regional scales (e.g., Hansen and Kelley 1995; Hoffmeister and Kowalewski 2001; Sawyer and Zuschin 2011) and does not qualify low predation intensity as a key condition in the northern Adriatic Sea.

*Reduced Drilling Frequencies in Seagrass Habitats.*—At Venice, DFs are significantly lower than at the other three stations. This unusually low DF is possibly due to pervasive seagrass development at that site throughout the Holocene succession, as indicated by the high abundance of the chemosymbiotic lucinid bivalve *Lucinella divaricata*, which occurs preferentially in sediments of seagrass meadows, and by the occurrence of strictly seagrass-associated rissoid gastropods such as *Rissoa monodonta* and *Rissoa membranacea* (Gallmetzer et al. 2019). The millennial-scale stability of seagrass ecosystems is well proven (e.g.,

Mateo et al. 2010; Leiva-Dueñas et al. 2018; Hyman et al. 2019), but the significance of seagrass development for predator–prey interactions is controversial. Experiments in modern environments suggest that the root mat typically formed by seagrasses protects infauna from burrowing predators (Orth et al. 1984), and this is particularly true for infaunal bivalves (Peterson 1982; Irlandi 1997; Goshima and Peterson 2012). On the other hand, DFs in the Neogene fossil record of the Caribbean were shown to be uniformly higher in biogenic habitats such as seagrass meadows compared with soft sediments (Leonard-Pingel and Jackson 2016). The very low DFs at Venice are in line with the experimental studies and suggest that seagrasses provided mollusks a refuge from drilling predation over the paleoecologically relevant timescale of millennia.

### Acknowledgments

Thanks to the captain of the coring vessel, J. Sedmak, to numerous students for their help with processing of samples, and to D. Chattopadhyay and P. Vasconcelos for sharing information on the ecology of muricid predators. This work was funded by the Austrian Science Fund (FWF): P24901. A.T. was supported by the Slovak Research and Development Agency (APVV 0555-17) and by the Slovak Scientific Grant Agency (VEGA 2/0169-19). Two reviewers provided helpful comments on the article.

### Data Availability Statement

Data available from the Dryad Digital Repository: <https://doi.org/10.5061/dryad.qfttdz0jj>.

Supplementary Figure 1. Correlations between environmental variables (TN, TOC, CaCO<sub>3</sub>, PC).

Supplementary Table 1. All data used for this study.

### Literature Cited

- Aberhan, M., W. Kiessling, and F. T. Fürsich. 2006. Testing the role of biological interactions in the evolution of mid-Mesozoic marine benthic ecosystems. *Paleobiology* 32:259–277.
- Abidli, S., P. Vasconcelos, Y. Lahbib, C. M. Barroso, N. Trigui El Menif, and M. B. Gaspar. 2012. Bivalve prey preference of *Bolinus brandaris* (Gastropoda: Muricidae) and type of prey attack on the bivalves *Cerastoderma glaucum*, *Venerupis decussata* and *Mytilus galloprovincialis* under laboratory conditions. Le 3ème Congrès Franco-Maghrébin de Zoologie et d'Ichtyologie, 6–10 Novembre 2012, Marrakech, Morocco.
- Amorosi, A., G. Barbieri, L. Bruno, B. Campo, T. M. Drexler, W. Hong, V. Rossi, I. Sammartino, D. Scarponi, S. C. Vaiani, and K. M. Bohacs. 2019. Three-fold nature of coastal progradation during the Holocene eustatic highstand, Po Plain, Italy—close correspondence of stratal character with distribution patterns. *Sedimentology* 66:3029–3052.
- Antonoli, F., M. Anzidei, K. Lambeck, R. Auriemma, D. Gaddi, S. Furlani, P. Orrù, E. Solinas, A. Gaspari, S. Karinja, and V. Kovačić. 2007. Sea-level change during the Holocene in Sardinia and in the northeastern Adriatic (central Mediterranean Sea) from archaeological and geomorphological data. *Quaternary Science Reviews*, 26:2463–2486.
- Aronson, R. B., D. B. Blake, and T. Oji. 1997. Retrograde community structure in the late Eocene of Antarctica. *Geology* 25:903–906.
- Bardhan, S., D. Chattopadhyay, S. Mondal, S. S. Das, S. Mallick, A. Roy, and P. Chanda. 2012. Record of intense predatory drilling from Upper Jurassic bivalves of Kutch, India: implications for the history of biotic interaction. *Palaeogeography, Palaeoclimatology, Palaeoecology* 317–318:153–161.
- Bambach, R. K. 1993. Seafood through time: changes in biomass, energetics, and productivity in the marine ecosystem. *Paleobiology* 19:372–397.
- Bates, D., M. Mächler, B. Bolker, and S. Walker. 2015. Fitting linear mixed-effects models using lme4. *Journal of Statistical Software* 67:1–48.
- Bates, D., M. Maechler, B. Bolker, S. Walker, R. H. B. Christensen, H. Singman, B. Dai, F. Scheipl, G. Grothendieck, P. Green, J. Fox, A. Bauer, and P. N. Krivitsky. 2020. lme4: linear mixed-effects models using ‘Eigen’ and S4, R package version 1.1-26. <http://CRAN.R-project.org/package=lme4>, accessed 9 March 2021.
- Baumiller, T. K., M. A. Bitner, and C. C. Emig. 2006. High frequency of drill holes in brachiopods from the Pliocene of Algeria and its ecological implications. *Lethaia* 39:313–320.
- Bottjer, D. J., and W. I. Ausich. 1986. Phanerozoic development of tiering in soft substrata suspension-feeding communities. *Paleobiology* 12:400–420.
- Bowden, D. A., S. Schiaparelli, M. R. Clark, and G. J. Rickard. 2011. A lost world? Archaic crinoid-dominated assemblages on an Antarctic seamount. *Deep-Sea Research, part II (Topical Studies in Oceanography)* 58:119–127.
- Brett, C. E. 1998. Sequence stratigraphy, paleoecology, and evolution: biotic clues and responses to sea-level fluctuations. *Palaio* 13:241–262.
- Bromley, R. G. 1981. Concepts in ichnotaxonomy illustrated by small round holes in shells. *Acta Geológica Hispánica* 16:55–64.
- Brückner, M. M. Z., C. Schwarz, G. Cocco, A. Baar, M. Boechat Albernaz, and M. G. Kleinhans. 2021. Benthic species as mud patrol-modelled effects of bioturbators and biofilms on large-scale estuarine mud and morphology. *Earth Surface Processes and Landforms* 46:1128–1144.
- Carriker, R. M. 1981. Shell penetration and feeding by naticacean and muricacean predatory gastropods: a synthesis. *Malacologia* 20:403–422.
- Cattaneo, A., F. Trincardi, A. Asioli, and A. Correggiari. 2007. The Western Adriatic shelf clinof orm: energy-limited bottomset. *Continental Shelf Research* 27:506–525.
- Catuneanu, O. 2019. Model-independent sequence stratigraphy. *Earth-Science Reviews* 188:312–388.
- Catuneanu, O., V. Abreu, J. P. Bhattacharya, M. D. Blum, R. W. Dalrymple, P. G. Eriksson, C. R. Fielding, W. E. Galloway, M. R. Gibling, K. A. Giles, J. M. Holbrook, R. Jordan, C. G. St. C. Kendall, B. Macurda, O. J. Martinsen, A. D. Miall, J.



- E. Neal, D. Nummedal, L. Pomar, H. W. Posamentier, B. R. Pratt, J. F. Sarg, K. W. Shanley, R. J. Steel, A. Strasser, M. E. Tucker, and C. Winker. 2009. Towards the standardization of sequence stratigraphy. *Earth-Science Reviews* 92:1–33.
- Chattopadhyay, D., M. Zuschin, S. Dominici, and J. A. Sawyer. 2016. Patterns of drilling predation in relation to stratigraphy, locality and sieve size: insights from the Eocene molluscan fauna of the Paris Basin. *Palaeogeography, Palaeoclimatology, Palaeoecology* 459:86–98.
- Connolly, S. R., and Roughgarden, J. 1999. Theory of marine communities: competition, predation, and recruitment-dependent interaction strength. *Ecological Monographs* 69:277–296.
- Freestone, A. L., R. W. Osman, G. M. Ruiz, and M. E. Torchin. 2011. Stronger predation in the tropics shapes species richness patterns in marine communities. *Ecology* 92:983–993.
- Fürsich, F. T., Alberti, M., and D. K. Pandey. 2021. Palaeoecological analysis of maximum flooding zones from the Tithonian (Upper Jurassic) of the Kachchh Basin, western India. *Facies* 67:1–27.
- Gallmetzer I., A. Haselmair, A. Tomašových, A. K. Mautner, S. M. Schnedl, D. Cassin, R. Zonta, and M. Zuschin. 2019. Tracing origin and collapse of Holocene benthic baseline communities in the northern Adriatic. *Palaios* 34:121–145.
- Gili, J. M., W. E. Arntz, A. Palanques, C. Orejas, A. Clarke, P. K. Dayton, E. Isla, N. Teixidó, S. Rossi, and P. J. López-González. 2006. A unique assemblage of epibenthic sessile suspension feeders with archaic features in the high-Antarctic. Deep-Sea Research, part II (Topical Studies in Oceanography) 53:1029–1052.
- Goshima, S., and C. H. Peterson. 2012. Both below- and above-ground shoalgrass structure influence whelk predation on hard clams. *Marine Ecology Progress Series* 451:75–92.
- Hancock, D. A. 1960. The ecology of the molluscan enemies of the edible mollusc. *Journal of Molluscan Studies* 34:123–143.
- Hansen, T. A., and P. H. Kelley. 1995. Spatial variation of naticid gastropod predation in the Eocene of North America. *Palaios* 10:268–278.
- Harper, E. M. 2003. Assessing the importance of drilling predation over the Palaeozoic and Mesozoic. *Palaeogeography, Palaeoclimatology, Palaeoecology* 201:185–198.
- Harper, E. M., and L. S. Peck. 2016. Latitudinal and depth gradients in marine predation pressure. *Global Ecology and Biogeography* 25:670–678.
- Hoffmeister, A. P., and M. Kowalewski. 2001. Spatial and environmental variation in the fossil record of drilling predation: a case study from the Miocene of central Europe. *Palaios* 16:566–579.
- Holland, S. M. 2000. The quality of the fossil record: a sequence stratigraphic perspective. *Paleobiology* 26(Suppl. to No. 4):148–168.
- Holt, R. D., and M. F. Hoopes. 2005. Food web dynamics in a meta-community context: modules and beyond. Pp. 68–94 in M. Holyoak, M. A. Leibold, and R. D. Holt, eds. *Metacommunities: spatial dynamics and ecological communities*. University of Chicago Press, Chicago.
- Hunt, D., and M. E. Tucker. 1992. Stranded parasequences and the forced regressive wedge systems tract: deposition during base-level fall. *Sedimentary Geology* 81:1–9.
- Huntley, J. W., and M. Kowalewski. 2007. Strong coupling of predation intensity and diversity in the Phanerozoic fossil record. *Proceedings of the National Academy of Sciences USA* 104:15006–15010.
- Huntley, J. W., and D. Scarponi. 2015. Geographic variation of parasitic and predatory traces on mollusks in the northern Adriatic Sea, Italy: implications for the stratigraphic paleobiology of biotic interactions. *Paleobiology* 41:134–153.
- Hyman, A. C., T. K. Frazer, C. A. Jacoby, J. R. Frost, and M. Kowalewski. 2019. Long-term persistence of structured habitats: seagrass meadows as enduring hotspots of biodiversity and faunal stability. *Proceedings of the Royal Society of London B* 286:20191861.
- Irlandi, E. A. 1997. Seagrass patch size and survivorship of an infaunal bivalve. *Oikos* 78:511–518.
- Kabat, A. R. 1990. Predatory ecology of naticid gastropods with a review of shell boring predation. *Malacologia* 32:155–193.
- Kelley, P. H., and T. A. Hansen. 2003. The fossil record of drilling predation on bivalves and gastropods. Pp. 113–139 in P. H. Kelley, M. Kowalewski, and T. A. Hansen, eds. *Predator-prey interactions in the fossil record*. Topics in Geobiology 20. Kluwer Academic/Plenum Publishers, New York.
- Kelley, P. H., and T. A. Hansen. 2006. Comparisons of class- and lower taxon-level patterns in naticid gastropod predation, Cretaceous to Pleistocene of the U.S. Coastal Plain. *Palaeogeography, Palaeoclimatology, Palaeoecology* 236:302–320.
- Kidwell, S. M. 1988. Taphonomic comparison of passive and active continental margins: Neogene shell beds of the Atlantic coastal plain and northern Gulf of California. *Palaeogeography, Palaeoclimatology, Palaeoecology* 63:201–223.
- Kidwell, S. M. 1993. Taphonomic expressions of sedimentary hiatuses: field observations on bioclastic concentrations and sequence anatomy in low, moderate and high subsidence settings. *Geologische Rundschau* 82:189–202.
- Klomp maker, A. A. 2009. Taphonomic bias on drill-hole predation intensities and paleoecology of Pliocene mollusks from Langenboom (Mill), the Netherlands. *Palaios* 24:772–779.
- Kondo, Y., S. T. Abbott, A. Kitamura, P. J. J. Kamp, T. R. Naish, T. Kamataki, and G. S. Saul. 1998. The relationship between shellbed type and sequence architecture: examples from Japan and New Zealand. *Sedimentary Geology* 122:109–127.
- Kowalewski, M. 2002. The fossil record of predation: an overview of analytical methods. *Paleontological Society Papers* 8:3–42.
- Kowalewski, M., A. Dulai, and F. T. Fürsich. 1998. A fossil record full of holes: the Phanerozoic history of drilling predation. *Geology* 26:1091–1094.
- Kowalewski, M., A. P. Hoffmeister, T. K. Baumiller, and R. K. Bambach. 2005. Secondary evolutionary escalation between brachiopods and enemies of other prey. *Science* 308:1774–1777.
- Leibold, M. A. 1989. Resource edibility and the effects of predators and productivity on the outcome of trophic interactions. *American Naturalist* 134:922–949.
- Leighton, L. R. 2002. Inferring predation intensity in the marine fossil record. *Paleobiology* 28:328–342.
- Leiva-Dueñas, C., L. López-Merino, O. Serrano, A. Martínez Cortizas, and M. A. Mateo. 2018. Millennial-scale trends and controls in *Posidonia oceanica* (L. Delile) ecosystem productivity. *Global and Planetary Change* 169:92–104.
- Leonard-Pingel, J. S., and J. B. C. Jackson. 2016. Drilling predation increased in response to changing environments in the Caribbean Neogene. *Paleobiology* 42:394–409.
- Mateo M. A., P. Renom, and R. H. Michener. 2010. Long-term stability in the production of a NW Mediterranean *Posidonia oceanica* (L.) Delile meadow. *Palaeogeography, Palaeoclimatology, Palaeoecology* 291:286–296.
- Mautner A-K., I. Gallmetzer, A. Haselmair, S.-M. Schnedl, A. Tomašových, and M. Zuschin. 2018. Long-term environmental shifts and human-induced loss of *Arca* and *Ostrea* shell beds in the north-eastern Adriatic Sea. *Marine Pollution Bulletin* 126:19–30.
- McKinney, F. K. 2003. Preservation potential and paleoecological significance of epibenthic suspension feeder-dominated benthic communities (northern Adriatic Sea). *Palaios* 18:47–62.
- McKinney F. K. 2007. The Northern Adriatic ecosystem: deep time in a shallow sea. Columbia University Press, New York.
- McKinney, F. K., and S. J. Hageman. 2006. Paleozoic to modern marine ecological shift displayed in the northern Adriatic Sea. *Geology* 34:881–884.

- McKinney, F. K., S. J. Hageman, and A. Jaklin. 2007. Crossing the ecological divide: Paleozoic to modern marine ecosystem in the Adriatic Sea. *Sedimentary Record* 5:4–8.
- McRoberts, C. A. 2001. Triassic bivalves and the initial marine Mesozoic revolution: a role for predators? *Geology* 29:359–362.
- Menge, B. A., and J. P. Sutherland. 1987. Community regulation: variation in disturbance, competition, and predation in relation to environmental stress and recruitment. *American Naturalist* 130:730–757.
- Morton, B., M. Peharda, and E. M. Harper. 2007. Drilling and chipping patterns of bivalve prey predation by *Hexaplex trunculus* (Mollusca: Gastropoda: Muricidae). *Journal of the Marine Biological Association of the United Kingdom* 87:933–940.
- Neely, S. H., P. H. Kelley, and M. M. Friedman. 2021. Predator–prey interactions among Pliocene mollusks from the Tjörnes Peninsula, Iceland; across the trans-Arctic invasion. *Lethaia*. <https://doi.org/10.1111/let.12426>.
- Novak, A., A. Šmuc, S. Poglajen, and M. Vrabec. 2019. Linking the high-resolution acoustic and sedimentary facies of a transgressed Late Quaternary alluvial plain (Gulf of Trieste, northern Adriatic). *Marine Geology* 419:106061.
- Novak, A., A. Šmuc, S. Poglajen, B. Celarc, and M. Vrabec. 2020. Sound velocity in a thin shallowly submerged terrestrial-marine Quaternary succession (northern Adriatic Sea). *Water* 12:560.
- O'Hara, B. 2009. How to make models add up—a primer on GLMMs. *Annales Zoologici Fennici* 46:124–137.
- Orth, R. J., K. L. Heck, and J. van Montfrans. 1984. Faunal communities in seagrass beds: a review of the influence of plant structure and prey characteristics on predator–prey relationships. *Estuaries* 7:339–350.
- Paine, R. T. 1974. Experimental studies on the relationship between a dominant competitor and its principal predator. *Oecologia* 15:93–120.
- Patzkowsky, M. E. 2017. Origin and evolution of regional biotas: a deep-time perspective. *Annual Review of Earth and Planetary Sciences* 45:471–495.
- Peharda, M., and B. Morton. 2006. Experimental prey species preferences of *Hexaplex trunculus* (Gastropoda: Muricidae) and predator–prey interactions with the black mussel *Mytilus galloprovincialis* (Bivalvia: Mytilidae). *Marine Biology* 148:1011–1019.
- Peterson, C. H. 1982. Clam predation by whelks (*Busycon* spp.): experimental tests of the importance of prey size, prey density, and seagrass cover. *Marine Biology*, 66:159–170.
- Pinheiro, J., and D. Bates. 2006. Mixed-effects models in S and S-PLUS. Springer Science & Business Media, New York.
- Rhoads, D. C., and L. F. Boyer. 1982. The effects of marine benthos on physical properties of sediments. Pp. 3–52 in P. L. McCall and M. J. S. Tevesz, eds. *Animal–sediment relations*. Topics in Geobiology 100. Springer, Boston.
- Rosenberg, R. 1995. Benthic marine fauna structured by hydrodynamic processes and food availability. *Netherlands Journal of Sea Research* 34:303–317.
- Sanders, H. L. 1958. Benthic studies in Buzzards Bay. I. Animal–sediment relationships 1. *Limnology and Oceanography* 3:245–258.
- Sawyer, J. A., and M. Zuschin. M. 2010. Intensities of drilling predation of molluscan assemblages along a transect through the northern Gulf of Trieste (Adriatic Sea). *Palaeogeography, Palaeoclimatology, Palaeoecology* 285:152–173.
- Sawyer, J. A., and M. Zuschin. 2011. Drilling predation in mollusks from the Lower and Middle Miocene of the Central Paratethys. *Palaios* 26:284–297.
- Sawyer J., M. Zuschin, B. Riedel, and M. Stachowitsch. 2009. Predator–prey interactions from in situ time-lapse observations of a sublittoral mussel bed in the Gulf of Trieste (northern Adriatic). *Journal of Experimental Marine Biology and Ecology* 371:10–19.
- Scarponi, D., and M. Kowalewski. 2007. Sequence stratigraphic anatomy of diversity patterns: late Quaternary benthic mollusks of the Po Plain, Italy. *Palaios* 22: 296–305.
- Schnedl, S.-M., I. Gallmetzer, A. Haselmair, A.-K. Mautner, A. Tomašových, and M. Zuschin. 2018. Molluscan benthic communities at Brijuni Islands (northern Adriatic Sea) shaped by Holocene sea-level rise and recent human eutrophication and pollution. *The Holocene* 28:1801–1817.
- Smith, A. B. 2007. Marine diversity through the Phanerozoic: problems and prospects. *Journal of the Geological Society of London* 164:1–15.
- Smith, J. A., J. C. Handley, and G. P. Dietl. 2018. Effects of dams on downstream molluscan predator–prey interactions in the Colorado River estuary. *Proceedings of the Royal Society of London B* 285:20180724.
- Storms, J. E., G. J. Weltje, G. J. Terra, A. Cattaneo, and F. Trincardi. 2008. Coastal dynamics under conditions of rapid sea-level rise: Late Pleistocene to Early Holocene evolution of barrier–lagoon systems on the northern Adriatic shelf (Italy). *Quaternary Science Reviews* 27:1107–1123.
- Tomašových, A., I. Gallmetzer, A. Haselmair, D. S. Kaufman, B. Mavrič, B., and M. Zuschin. 2019. A decline in molluscan carbonate production driven by the loss of vegetated habitats encoded in the Holocene sedimentary record of the Gulf of Trieste. *Sedimentology* 66:781–807.
- Tomašových, A., Gallmetzer, A., Haselmair, and M. Zuschin. 2022. Inferring time averaging and hiatus durations in the stratigraphic record of high-frequency depositional sequences. *Sedimentology*. <https://doi.org/10.1111/sed.12936>.
- Trincardi, F., A. Correggiari, and M. Roveri. 1994. Late-Quaternary transgressive erosion and deposition in a modern epicontinental shelf: the Adriatic semienclosed basin. *Geo-Marine Letters* 14:41–51.
- Trobec, A., A. Šmuc, S. Poglajen, and M. Vrabec, M. 2017. Submerged and buried Pleistocene river channels in the Gulf of Trieste (northern Adriatic Sea): geomorphic, stratigraphic and tectonic inferences. *Geomorphology*, 286:110–120.
- Vacchi, M., N. Marriner, C. Morhange, G. Spada, A. Fontana, and A. Rovere. 2016. Multiproxy assessment of Holocene relative sea-level changes in the western Mediterranean: sea-level variability and improvements in the definition of the isostatic signal. *Earth-Science Reviews* 155:172–197.
- Vermeij, G. J. 1977. The Mesozoic marine revolution: evidence from snails, predators and grazers. *Paleobiology* 3:245–258.
- Vidović, J., R. Nawrot, I. Gallmetzer, A. Haselmair, A. Tomašových, M. Stachowitsch, V. Čosović, and M. Zuschin. 2016. Anthropogenically induced environmental changes in the northeastern Adriatic Sea in the last 500 years (Panzano Bay, Gulf of Trieste). *Biogeosciences* 13:5965–5981.
- Yochelson, E. L., D. Dockery, and H. Wolf. 1983. Predation of sub-Holocene scaphopod mollusks from southern Louisiana. *U.S. Geological Survey Professional Paper* 1282.
- Zecchin, M., E. Gordini, and R. Ramella. 2015. Recognition of a drowned delta in the northern Adriatic Sea, Italy: stratigraphic characteristics and its significance in the frame of the early Holocene sea-level rise. *The Holocene* 25:1027–1038.
- Zuschin, M., and M. Stachowitsch. 2009. Epifauna-dominated benthic shelf assemblages: lessons from the modern Adriatic Sea. *Palaios* 24:211–221.
- Zuschin, M., M. Harzhauser, and O. Mandic. 2011. Disentangling palaeodiversity signals from a biased sedimentary record: an example from the Early to Middle Miocene of Central Paratethys Sea. In A. McGowan and A. B. Smith, eds. *Comparing the geological and fossil records: implications for biodiversity studies*. Geological Society of London Special Publication 358:123–139.
- Zuschin M., R. Nawrot, M. Harzhauser, O. Mandic, and A. Tomasovych. 2017. Taxonomic and numerical sufficiency in depth- and salinity-controlled marine paleocommunities. *Paleobiology* 43:463–478.