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1	DEMISE AND RECOVERY OF ANTILLEAN SHALLOW MARINE CARBONATE
2	FACTORIES ADJACENT TO ACTIVE SUBMARINE VOLCANOES (LUTETIAN-
3	BARTONIAN LIMESTONES, ST. BARTHOLOMEW, FRENCH WEST INDIES)
4	
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13 ABSTRACT

14 Among other parameters, volcanic activity adjacent to shallow marine environments influences 15 the development of ecosystems and their carbonate-producing communities. Volcaniclastic 16 sediment influx in particular has potential to cause rapid and drastic environmental changes 17 affecting biological systems in their composition and activity, and ultimately leading to changes 18 to and breaks in carbonate sedimentation. Such sedimentary breaks that form in response to 19 volcanic processes are rarely described in detail despite the common occurrence of carbonate platforms adjacent to active volcanoes both in the recent and past geological record. The island of 20 21 St. Bartholomew (St. Barths), French West Indies, exposes sections of middle Eocene limestones 22 intercalated with thick volcaniclastic deposits and lavas. Theses carbonates provide an example 23 of a low-latitude tropical platform where non-framework building biota were important, if not

24	dominant sediment contributors. The carbonate system records the repeated collapse and renewal
21	
25	of carbonate production, as a result of episodic volcaniclastic material input. The discontinuous
26	nature of the carbonate sedimentation is reflected in contrasted depositional systems across
27	sedimentary surfaces and gradational contacts. The studied Eocene deposits provide a
28	sedimentary record of how volcanic events impacted warm-water carbonate factories, both in
29	their disturbance, demise and recovery.
30	
31	Keywords: Eocene, Caribbean, carbonate ramp, skeletal carbonates, volcaniclastics
32	
33	1. Introduction
34	
35	Volcanism may be intuitively regarded as critically detrimental to shallow-water benthic
36	communities, and as such as one environmental parameter impacting the development of
37	carbonate platforms. Lava flows and large pyroclastic eruptions, either subaqueous or reaching
38	subtidal environments from emerged volcanoes, can undeniably result in mass mortality of
39	carbonate producers and therefore cause carbonate production to cease (Tomascik et al., 1996;
40	Vroom and Zgliczynski, 2011). However, recent studies have shown that volcaniclastic influx
41	may also allow carbonate-producing organisms with higher surviving capabilities to thrive, and
42	may promote a change of benthic communities, prolonging the existence of carbonate factories
43	(Wilson and Lokier, 2002; Lokier et al., 2009). These biota prove to be adapted to post-eruption
44	environmental conditions such as reduced luminosity and increased turbidity due to suspended
45	ash, change of substrate, change of nutrient levels and of water chemistry through alteration of
46	volcanic material (Reuter and Piller, 2011). It follows that volcanic events, even of short duration
47	and/or limited in the volume of material delivered to the marine realm, may be diversely

48 expressed in otherwise carbonate-dominated settings and ultimately in outcrops. Expected 49 physical, stratal and facies expressions of volcaniclastic influx include: (1) erosion of the seafloor 50 and of reefal buildups (Wilson, 2000); (2) reduction of or temporarily arrested carbonate 51 production; (3) deposition of volcaniclastic material; and (4) a change in the depositional systems 52 and in the biotic assemblages (Wilson and Lokier, 2002; Dorobek, 2008; Lokier et al., 2009; 53 Reuter et al., 2012). 54 Herein, we propose a depositional model for Eocene tropical carbonates that formed in a 55 volcanically active area, namely the Lesser Antilles Volcanic Arc, West Indies. Here, carbonate 56 and mixed carbonate-volcaniclastic units are intercalated between volcaniclastics and lava flows, 57 pointing to the repeated demise and foundering of carbonate factories, but also to the interplay 58 between volcanic activity and penecontemporaneous carbonate production. This study is the first 59 to document in detail the compositional characteristics of the Antillean carbonate successions on 60 St. Barths Island. 61 62 2. Geological setting 63 64 The Lesser Antilles arc is related to the westward-dipping subduction zone where the oceanic

65 crust of the American plate is underthrusting the Caribbean plate, hence causing important

66 volcanic and seismic activities (Fig. 1A).

North of the Island of La Martinique, the Lesser Antilles arc consists of two divergent volcanic lines, namely the now extinct outer arc active from the early Eocene to middle Oligocene, and the still active inner arc (Bouysse et al., 1990). Owing to this evolution and subsequent uplift, the outer arc system became the locus of abundant carbonate production and accumulation during the late Paleogene and Neogene. It evolved into extended carbonate platforms that correspond to the 72 so-called Limestone Caribbees, whereas to the west, intense volcanic activity led to the 73 construction of an archipelago of islands referred to as the Volcanic Caribbees (Bouysse et al., 74 1990). 75 The Island of St. Barths lies on the northeastern sector of the Lesser Antilles outer arc and 76 belongs to the present-day platform. It consists predominantly of limestone beds intercalated 77 between volcanics (Fig. 1B). The latter includes hyaloclastite deposits generated by hydroexplosions interbedded with andesitic and basaltic submarine lava-flows. Up to six limestone 78 79 sheets from 3 up to 40 m thick have been recognized and previously interpreted as peri-reefal 80 carbonates that were given a late middle Eocene (Lutetian and Bartonian) age on the basis of 81 microfaunal content (Westercamp and Andreieff, 1983a, 1983b; Gradstein et al., 2012). This 82 study concentrates on the limestone units and their under- and overlying contacts with volcanics. 83 3. Methods 84 85 86 3.1. Field and laboratory analysis 87 88 The bulk of information used in this study comes from field observation and description of 13 89 stratigraphic sections along the coasts of St. Barths and on two islets located southwest and 90 southeast off the main island, namely Pain de Sucre and Île Coco, respectively (Figs. 1C, 2). 91 Due to their common occurrence in the carbonate strata and locally in volcaniclastic units, and to

92 their relative abundance in some horizons associated with key sedimentary surfaces, echinoid

- 93 macrofossils were given special attention (Fig. 2). Field characteristics (diversity, abundance,
- 94 orientation and preservation) of spatangoid (Schizaster L. Agassiz, Agassizia L. Agassiz and
- 95 Desor, Eupatagus L. Agassiz, Antillaster Lambert) and oligopygoid (Haimea Michelin) echinoids

96	in particular, were determined prior to being collected where possible for taxonomic
97	identification. Echinoids include epibenthic dwellers and shallow to deep endobenthic burrowers
98	(Kier, 1984; Kanazawa, 1992; Donovan and Rowe, 2000). As such, echinoids are useful
99	indicators of paleoenvironmental conditions, including trophic resources, absence/presence of
100	seagrass beds, and substrate (Fig. 3). Characteristics of other dominant macrofossils, such as
101	corals and pinnids, were also evaluated qualitatively in the field (living position, mechanical
102	erosion, bioinfestation). In this study, the taxonomy of coralline algae uses at best generic names
103	only. Genus circumscription follows Rasser and Piller (1999) and Braga et al. (2009). Family to
104	subfamily names follow Harvey et al. (2003). Coralline algal growth-form terminology follows
105	Woelkerling et al. (1993). The taxonomy of larger foraminifera is restricted to Family level and at
106	best genus level except for those listed in Westercamp and Andreieff (1982, 1983).
107	The Eocene limestone units were analyzed petrographically using 110 thin-sections cut in
108	samples selectively collected below and above sharp, erosive, and burrowed contacts. The aim
109	was to provide estimates of the relative abundances of skeletal and non-skeletal constituents, and
110	to assess facies changes across sedimentary surfaces. Results were converted for the dominant
111	taxa at subfamily level and grain types into abundance indices of rare (one or two grains seen),
112	occasional (about 10%), common (conspicuous throughout, i.e., 20-30%), and abundant (the
113	grain category considered is the dominant skeletal component, i.e., \geq 50%). The data obtained
114	were used to determine the composition of the skeletal associations only, and are not suitable for
115	statistical analysis. These qualitative data were complemented by textural information using the
116	criteria of Embry and Klovan (1971). In addition, a series of specific scales of taphonomic
117	features have been defined for various fossil types observed in thin section to help assess the
118	effects of early sea-floor processes, namely fragmentation, abrasion, bioerosion and encrustation

- 119 (Fig. 4).
- 120
- 121 3.2. Skeletal associations
- 122

123 Because pervasive bioturbation has obliterated, but in a few examples, primary wave- and 124 current-induced sedimentary structures, interpretation of the facies in terms of depositional 125 conditions was based on comparative ecology and sedimentology with ancient and modern 126 shallow marine environments (e.g., Hallock and Glenn, 1986; Hottinger, 1997; Bassi, 1998, 127 2005; Braga and Aguirre, 2001; Hallock, 2001; Hohenegger and Yordanova, 2001; Beavington-128 Penney and Racey, 2004; Hallock et al., 2006; Hohenneger, 2009; Payros et al., 2010). Results of 129 the petrographic analysis highlight a diverse array of biotic constituents and skeletal associations, 130 which can satisfactorily be classified as chlorozoan, chloralgal and foramol (Lees and Buller, 131 1972; Lees, 1975; James, 1997), rhodalgal (Carannante et al., 1988), echinofor and foralgal 132 (Hayton et al., 1995). Although chloralgal and chlorozoan associations dominated by 133 zooxanthellate corals, calcareous green algae and non-skeletal carbonates (e.g., ooids, peloids and 134 oncoids) are robust indicators of relatively oligotrophic clear sea waters inside the tropical 135 climate belt, interpretation of the majority of skeletal associations in terms of temperature, 136 climate and paleoecology needs to integrate additional palaeoenvironmental information such as 137 nature of the substrate, siliciclastic fluxes and trophic resources (e.g., Hallock and Schlager, 138 1986; Hallock, 1985, 1987; Allmon, 1992; Brasier, 1995a, b; Pomar et al., 2004; Wilson and 139 Vecsei, 2005; Lokier et al., 2009; Reuter and Piller, 2011). 140

140

141 3.4. Paleoenvironments and paleodepths

143 Although small isolated coral bioherms (patch-reefs s.l.) and coral carpets are present in some of 144 the limestone units, the absence of major reefal constructions (e.g., barrier coral-reef) and the 145 sedimentological characteristics of facies within the St. Barths limestones suggest that the general 146 depositional setting was a carbonate ramp (i.e., following the terminology of Burchette and 147 Wright, 1992) attached to and/or surrounding the flanks of active submarine volcanoes. The 148 subdivision of the carbonate ramp system into inner, middle and outer ramp depositional settings 149 follows the criteria proposed by Pomar (2001). The term inner ramp refers to depositional 150 environments between upper shoreface (i.e., the zone of breaking waves or lagoonal shorelines) 151 and Fair-Weather Wave Base (FWWB), middle ramp between FWWB and Storm-Wave Base 152 (SWB), and outer ramp below SWB (Burchette and Wright, 1992). The division of photic zones 153 with respect to water depths is based on the criteria of Pomar (2001). The low-light conditions 154 can be inferred from the absence or very rare co-occurrences of miliolids and calcareous green 155 algae (dasycladaleans), which typically are indicative of inner ramp environments when they are 156 the dominant biotic constituents (Geel, 2000; Zamagni et al., 2008). 157 The presence and abundance of taxa indicative of specific depth constraints known from modern 158 environments and interpreted from ancient deposits were only used to discriminate between 159 euphotic and oligophotic conditions (Pomar, 2001). Shallow-water biotic indicators include 160 zooxanthellate corals, pinnid bivalves, miliolids, conical agglutinated foraminifera and seagrass 161 dwellers such as small benthic rotaliids (Geel, 2000; Vecchio and Hottinger, 2007). The 162 estimation of paleodepth using trends in test flattening of larger benthic foraminifera, combined 163 with textural attributes of host sediments, namely siliciclastic content, grain size, and presence of 164 carbonate/argillaceous mud (Hohenegger et al., 2000; Beavington-Penney and Racey, 2004; 165 Cosovic et al., 2004; Hohenegger, 2009), suggests deposition in inner- to outer-ramp settings for 166 amphisteginids- and nummulitids-bearing deposits. Foralgal associations that include coralline

167	algae, such as dominant melobesioids (Lithothamnion, Mesophyllum) and subordinate
168	mastophoroids (Lithoporella) with rare sporolithales (Sporolithon) are generally associated with
169	middle-ramp to outer ramp settings (Nebelsick and Bassi, 2000; Braga and Aguirre, 2001; Pomar
170	et al., 2004). The abundance of flattened orthophragminids and planktic foraminifera points to
171	distal middle ramp to outer ramp settings (Bassi, 2005; Zamagni et al., 2008).
172	Changes in the depositional environments and in water depth thresholds for photodependent biota
173	are however to be expected in the immediate aftermath of volcanic eruptions due to clastic influx
174	and ash suspended in water (Wilson and Lokier, 2002; Lokier et al., 2009; Reuter and Piller,
175	2011; Reuter et al., 2012).
176	
177	4. Description of volcaniclastic deposits
178	
179	Volcaniclastic deposits represent the products of submarine eruptive episodes (Westercamp and
180	Andreieff, 1983a), and include a basal ensemble of arc tholeiites and an upper ensemble of calc-
181	alkaline series (Fig. 1B-C). Volcaniclastic deposits range in thickness from several tens of meters
182	to hundreds of meters. They are dominated by maar-type breccia (Vbr), coarse-grained
183	hyaloclastites (Hy), and medium- to fine-grained tuffs (Tu), in a proximal to distal distance
184	gradient from eruptive centers.
185	Coarse volcaniclastic breccia (Vbr) are massive to crudely bedded and contain heterometric
186	angular clasts ranging in size from gravel to boulder, in a coarse-grained tuffitic matrix. These
187	deposits are generally azoic, but may contain limestone clasts and coral heads reworked from
188	underlying limestone units (Fig. 5A-B).
189	Lithologies of coarse-grained hyaloclastites (Hy) and fine- to medium-grained tuffs (Tu) are
190	highly variable (see Westercamp and Andreieff, 1983b). The former can be subdivided into two

191 major groups, based on bedding structure and clast shape.

The first group (Hy1) consists of horizontal and low-angle tabular, fining-upward, dm-thick beds 193 having erosional bases. The basal contact is overlain by gravel- to cobble-sized (sub-)angular 194 volcanic clasts in a poorly-sorted sandy/gravely matrix that grades into moderately sorted 195 volcanic sands (Fig. 5C). 196 The second group (Hy2) represents reworked hyaloclastites (Fig. 5D-E). The deposits 197 conformably overlie mixed carbonate-volcaniclastic limestones and exhibit alternating 198 poorly/moderately-sorted conglomeratic beds that contain sub-rounded coarse sand- to pebble-199 sized volcanic clasts, and moderately-sorted coarse-grained volcaniclastic sandstones (Fig. 5E). 200 Bioclasts and macrofossils, including bivalves, gastropods, bryozoans, fragmented coral branches 201 and echinoids are common in the deposits. 202 Tuffs (Tu) include cm- and dm-thick tabular beds of silty to sandy, moderately- to well-sorted 203 deposits (Fig. 5F). Tuffs may be bioturbated and fossiliferous, with larger benthic foraminifera, 204 solitary corals, and spatangoids. 205 206 5. Carbonate facies description 207 208 A total of 15 facies have been distinguished that point to various environmental and 209 hydrodynamic conditions (Table 1). Features common to all facies are that none contain ooids or 210 evaporitic features.

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212 5.1. Mixed carbonate-volcaniclastic facies association (Mf1 to Mf5)

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214 The abundance of volcaniclastic, tuffitic, feldspar crystals and quartz grains characterize this facies association. It can be subdivided into five major facies depending on the characteristics of volcaniclastic components and on the biotic/non-skeletal carbonate particle content. Tuffitic sediments, angular/subangular volcaniclastic fine sand to granule sized-debris and quartz fine-to medium sand represent 20% to 50% of the sediment fraction.

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5.1.1. Smaller foraminiferal – dasycladalean - coral photozoan mixed facies (Mf1 and Mf2)
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Two biotic associations comprise this facies. Mf1 is a smaller foraminiferal-dasycladalean
association, and includes moderately- to well-cemented wackestones/floatstones, packstones and
more rarely grainstones. Mf2 includes dasycladalean-coral floatstones and coral boundstones.
Intense bioturbation produces a typical nodular aspect to facies Mf1 and Mf2 beds, though
massive bedding also occurs. Facies Mf1 and Mf2 strata may either overlie volcaniclastic units
across sharp and gradational contacts, or carbonate beds across sharp undulating and erosional
surfaces penetrated by *Thalassinoides* burrows.

229 In general, facies Mf1 is characterized by a low-diversity biota with a relative abundance of 230 smaller benthic foraminifera, and subordinate miliolids and dasycladalean algae (Fig. 6A). 231 Peloids may be common to abundant, and micritized grains are generally present. Agglutinated 232 conical foraminifera and laminar crusts of coralline red algae (Lithoporella) are occasional to 233 common (Fig. 6B). Other organisms, such as echinoids (Eupatagus, Antillaster), thin-walled 234 gastropods, bivalves, branching corals in living position and rare amphisteginids are also present. 235 Macrofossils and microfossils in micritic floatstones and wackestones/packstones show low to 236 moderate degrees of fragmentation and abrasion, low degrees of encrustation and moderate 237 degrees of bioerosion, except for corals which may be intensely bored. Grainstones typically 238 exhibit higher degrees of fragmentation and abrasion, lower encrustation, and similar bioinfestion of skeletal grains.

241 filled moulds in a micritic matrix and constitute coral boundstones that locally build up small 242 patchreef frameworks. Thick-shelled gastropods, dasycladalean algae and miliolids may be 243 common. Large photosymbiont-bearing foraminifera are rare to common in places and include 244 amphisteginids and nummilitids. Agglutinated conical foraminifera and echinoid fragments are 245 present in small amounts. Other fossils include epifaunal spatangoids such as Antillaster. 246 Skeletons in facies Mf2 show low to moderate degrees of fragmentation and abrasion leaving 247 gastropods with their tubercules left intact for example. Corals are commonly partly micritized 248 and encrusted by cyanobacterial micritic laminae.

Facies Mf2 differs from Mf1 facies in being dominated by branching corals, which occur as spar-

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250 5.1.2. Echinoid – Red algal photozoan facies (Mf3)

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Textural attributes of facies Mf3 are variable and range from wackestone to packstone/grainstone
(Fig. 6C). Massive bedding is typical, although nodular bedding also occurs pointing to intense
bioturbation that likely obliterated primary sedimentary structures.

255 Facies Mf3 is characterized by low-diversity biogenics with a relative abundance of echinoids 256 (plates and spine fragments) and fragments of rhodoliths, branches and laminar crusts. Larger 257 foraminifera are unusual, but when present consist predominantly of orthophragminids, and rare 258 nummulitids and amphisteginids. Micritic grains, such as peloids and cortoids, may be abundant. 259 Subordinate biogenics include bivalves and corals. Intraclasts are present and consist of skeletal 260 grains bearing rims of palisadic spar crystals with scalenohedral terminations and echinoid plates 261 with dirty spar syntaxial overgrowths. Facies Mf3 has moderate to high degrees of fragmentation 262 and abrasion, and relatively low to moderate degrees of bioerosion and encrustation.

264 5.1.3. Mixed larger foraminiferal – mollusk photozoan facies (Mf4)

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266 The massive beds of facies Mf4 are sharp-based. Mollusks, larger ovate nummulitids, 267 amphisteginids and agglutinated conical foraminifera with particularly high amounts of coarse 268 sand to granule sized-volcaniclastics dominate these sediments. Volcaniclastic grains are sub-269 angular and poorly- to moderately-sorted. Coral and coralline algal debris, and bryozoans are 270 occasional to common and typically show moderate to high degrees of fragmentation and 271 abrasion, little if any encrustation, and low bioerosional features. Echinoids are generally found 272 fragmented and include sand-dwellers Eupatagus, Meoma and Haimea (Fig. 3). Sedimentary 273 structures vary from planar bedding to sigmoidal and hummocky cross-stratifications. Facies 274 textures are volcaniclastic rudstones with a packstone and/or grainstone matrix.

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276 5.1.4. Mixed Bioclastic – peloidal facies (Mf5)

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278 Beds of facies Mf5 are either massive with a wavy basal contact or nodular grading from 279 underlying volcaniclastic deposits across a transitional interval of carbonate nodules in an 280 argillaceous matrix. Moderately to well-sorted silt and fine sand-sized bioclastic and tuffitic 281 material dominate this facies (Fig. 6D). The carbonate fraction consists predominantly of peloids 282 originating in micritized skeletal fragments, such as red algal debris, and of cortoids in a 283 packstone matrix. Subordinate components include fragmented benthic foraminifera, echinoid 284 plates and spines, and corals. Thin-shelled disarticulated bivalves may be present. Fragmentation 285 and abrasion are moderate, bioerosion is high and encrustation is rare.



310 5.3.1. Larger benthic foraminiferal – rhodolith photozoan facies (LfRh1 to LfRh3)

312 These facies consist of coarse-grained packstones, sparitic rudstones/grainstones and rare 313 bindstones dominated by larger benthic foraminifera and coralline red algae. Decimeter-thick 314 beds have wavy bedding planes and are intensely bioturbated, resulting in a typical nodular 315 texture. Medium-scale (up to 1 m large and dm deep) trough cross-stratifications and low- to 316 high-angle foresets are locally present. The larger foraminiferal assemblage of facies LfRh1 317 consists of common amphisteginids, rare to occasional nummulitids, and lepidocyclinids. Facies 318 LfRh2 is a nummulitid-lepidocyclinid packstone-rudstone. Fragments of fruticose coralline thalli 319 and rhodoliths are also present in both facies and may be common to abundant within the 320 carbonate matrix. In addition, other biogenic components include occasional textulariids, rare 321 miliolids, occasional mollusks dominated by epifaunal bivalves, occasional coral fragments and 322 planktonic foraminifera. Peloids and cortoids range in abundance from rare to common. Coralline 323 bindstones are present locally in facies LfRh2 as intraclasts. The binding structure is composed of 324 laminar coralline thalli 0.5-1.0 mm thick and cyanobacterial laminae with a stromatolithic 325 texture. Mastophoroids (Lithoporella) and sporolithalean (Sporolithon) are present in facies 326 LfRh1, and rare to absent in facies LfRh2 and LfRh3. Facies LfRh3 is dominated by rhodoliths in 327 a bioclastic grainstone matrix composed of common echinoid plates and peloids, occasional 328 larger benthic foraminifera, mollusks, corals, and dasycladalean algae (Fig. 6F). 329 Rhodoliths in facies LfRh 1 and LfRh2 are variable in size with a mean diameter from 1 to 10 330 cm, and are typically dispersed within a coarse-grained grainstone matrix. Coalescence of two 331 rhodoliths has been observed. Rhodoliths in facies LfRh3 are more homogeneous in size ranging 332 from 1-5 cm. The most common rhodolith morphology is sub-spheroidal with a dense inner 333 laminar arrangement of coralline thalli. Epifaunal macrofossils include irregular echinoids 334 Haimea and Antillaster, and large thick-shelled oysters, which may locally be abundant in facies

335 LfRh1.

There is no significant difference between facies LfRh1, LfRh2 and LfRh3 in their degree of taphonomic alteration. Each facies consists of a mixture of poorly to highly altered skeletal material with respect to fragmentation, abrasion and bioerosion. Encrusters include red algae, serpulids, foraminifera, and cyanobacteria.

340

5.3.2. Larger benthic foraminiferal – coralline algal photozoan facies (LfCa1 and LfCa2)
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343 Facies LfCa1 and LfCa2 consist of massive- to nodular-bedded larger foraminiferal-red algal 344 packstones and grainstones. Basal contacts are either gradational or sharp. Underlying strata are 345 mixed carbonate-volcaniclastic and carbonate units. The main components are coralline algal 346 branches, less than 1 cm-sized rhodoliths, and their debris, and larger benthic foraminifera. 347 Among the latter, amphisteginids and subordinate nummulitids dominate facies LfCa1 (Fig. 6G), whereas nummilitids associated with orthophragminids are abundant in facies LfCa2 (Fig. 6H). 348 349 Other dominant components include echinoid plates and spines, smaller benthic foraminifera 350 (rotaliids and occasional textulariids), and peloids and micritized skeletons. Subspheroid 351 rhodoliths (up to 3 cm in diameter) may also be present. Coralline branches and debris consist mostly of melobesioids and occasional mastophoroids (Lithoporella). 352 353 The foraminiferal assemblage is dominated by *Amphistegina* and *Nummulites* in LfCa1, and 354 Nummulites, Discocyclina and Pseudophragmina in LfCa2. 355 Skeletal material in facies LfCa1 has lower degrees of alteration than its counterpart in facies 356 LfCa2. In the latter, coralline thalli and larger foraminifera exhibit a wide range of alteration from 357 low to high for fragmentation, abrasion, bioerosion and encrustation, pointing to the mixing of 358 fresh and older skeletons, i.e., exposed for longer periods of time to destructive processes.

361

Facies EcF1 are well-cemented packstones and grainstones containing abundant echinoid fragments, common peloids and subordinate larger benthic foraminifera (Fig. 6I). Beds of facies EcF1 are massive to nodular with sharp basal contacts. Facies EcF2 are semi-lithified packstones and wackestones rich in small benthic foraminifera (rotaliids), and comprise a complex assemblage of echinoid macrofossils. Decimeter-thick beds of facies EcF2 generally overlie wavy surfaces pervasively bioturbated by *Thalassinoides*.

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369 5.4.1. Echinoid – non-skeletal carbonate photozoan facies (EcF1)

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371 Irregular echinoids, including spatangoids (Haimea) and cassiduloids (Echinolampas) are 372 common in these rocks. Branching zooxanthellate corals in growth position are ubiquitous at 373 some localities. Gastropods and disarticulated bivalves are present. The fine- to medium-sand 374 sized biogenic particles comprise common to abundant echinoid plates (Fig. 6I), occasional larger 375 benthic foraminifera, particularly amphisteginids and nummulitids, molluskan fragments, 376 geniculate and non-geniculate coralline debris, and less common miliolids, small agglutinated 377 foraminifera, coral debris, and bryozoans. Peloids are common to abundant. Frequent presence of 378 micritized skeletons suggests that the majority of peloids may be small micritized grains.

379 Degrees of fragmentation and abrasion are moderate to high, and common micritization indicates380 that bioinfestation is pervasive. Encrustation of coral fragments and mollusks is low to moderate,

and involves cyanobacteria, red algae and foraminifera.

385 The co-occurrence of spatangoids, cassiduloids, and oligopygoids with different living habitats 386 and substrate dependencies is frequent in facies EcF2 deposits. The co-occurrence of Haimea, 387 Schizaster, Agassizia, Eupatagus, and Echinolampas provides an example of such complex 388 diverse echinoid assemblages (Fig. 3) that point to contrasted depositional conditions over time. 389 Results show that about 30% echinoids are in living position, a majority of which (80%) being 390 represented by shallow (Agassizia) and deep (Schizaster) infaunal dwellers. Other macrofossils 391 include large thick-shelled gastropods and pinnid bivalves dispersed in the deposits. Foraminifera 392 comprise the porcelaneous and hyaline smaller benthic majority. Larger forms are occasional. 393 Coralline fragments are rare and tend to be dominated by geniculate forms. Small agglutinated 394 foraminifera, mollusks, miliolids, and calcispheres are occasional. Peloids may be abundant, 395 whereas cortoids and oncoids are unusual. Volcaniclastic content is low.

396 The skeletal fraction has moderate to high degrees of fragmentation and abrasion, low to 397 moderate degrees of bioerosion and encrustation. However, echinoid plates are typically less 398 affected by destructive processes than other skeletons.

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400 5.5. Coral photozoan facies (CoP1 and CoP2)

401

Two facies that reflect different biohermal geometries, hydrodynamic conditions and coral shapes are present. Members of the genera *Astrocoenia*, *Hydnophora*, and *Cladocora* have been identified. Small bioherms with lenticular geometry, a few meters large and up to 2-4 m high, and coral carpets characterize facies CoP1. Branching and platy corals in growth position constitute the coral framework. Corals are enclosed in a peloidal-micritic matrix. Other components include 407 encrusting coralline algae, gastropods and small benthic foraminifera. Meter-sized lenticular 408 bioherms exhibit several growth-phases, each one being laterally correlated to tabular beds of 409 facies Mf2, EcF2, and DaMi (Fig. 7A), a geometry that reflects a strong interaction between 410 biohermal building and ambient sedimentation (Gaillard, 1983). Massive coral framework and 411 biohermal rudstones compose facies CoP2. Individual biohermal structures are up to 6-8 m-high. 412 Massive dendroid corals in living position dominate in the upper half part of the constructional 413 bodies (Fig. 7B). Pebble to cobble-sized reef debris constitute lateral correlatives of the coral 414 build-ups and are deposited as structureless up to 1 m-thick beds intercalated between tabular and 415 nodular beds of facies EcF1 and LfCa1. The base of the coral build-ups appear sharp and possibly 416 erosional (Fig. 7B), though not confirmed by field observation, a discrepancy that may originate 417 in differential compaction and diagenesis (Rusciadelli and Di Simone, 2007; Berra and 418 Carminati, 2012).

419

420 **6. Carbonate facies interpretation**

421

422 The virtual absence of silty micrite-rich planktonic foraminifera-dominated facies indicates that 423 St. Barths Eocene carbonates and mixed carbonate-volcaniclastic sediments were principally 424 deposited in inner to middle ramp settings (Nebelsick et al., 2005). In addition, the rare to 425 occasional occurrence of symbiont-bearing larger benthic foraminifera with flattened shapes 426 (orthophragminids, lepidocyclinids), a morphological adaptation to maximize light collection in 427 otherwise oligophotic conditions below wave influence (Hallock and Glenn, 1986; Beavington-428 Penney and Racey, 2004; Hohenegger, 2009), and of light-independent biota-dominated facies 429 confirm that the studied deposits are representatives of inner ramp and proximal to possibly distal 430 middle ramp facies belts. Figure 8 proposes a reconstruction of the depositional environments

431	envisaged for the St. Barths carbonate facies. We must emphasize that different microfacies
432	associated with identical parts of the carbonate ramp may not have all developed
433	contemporaneously, even though a mosaic of subenvironments, each being characterized by
434	specific ecological attributes in relation to subtle or major changes in substrate, nutrient levels,
435	water depth, salinity and/or other parameters, should be expected in settings influenced by
436	volcanism, subsequent influxes of volcaniclastics and resulting environmental stresses.

438 6.1. Inner ramp: shallow back barrier setting and patch reefs

439

440 Biogenic assemblages dominated by euphotic taxa, sedimentary structures comprising thin and 441 planar beds, and micritic textures of facies Mf1, Mf2, Mf5, DaMi, CoP1 and EcF2 (Table 1) are 442 interpreted to signal shallow-water inner ramp environments. The abundance of smaller benthic 443 foraminifera, including textulariids, the common occurrence of dasycladal algae, miliolids, agglutinated conical foraminifera, and the rarity of hyaline larger foraminifera in facies Mf1 444 445 suggest light-level conditions of the upper euphotic zone (Murray, 1991; Geel, 2000; Zamagni et 446 al., 2008). The presence of delicate laminar crusts of mastophoroid coralline algae, which appear 447 detached from their substrate and commonly exhibit hook and ring shapes (Fig. 6E), together 448 with common calcareous green algae fragments and smaller benthic foraminifera hint for the 449 existence of seagrass beds in the depositional environment (Sola et al., 2013). 450 The local abundance of coral colonies constructing isolated bioherms, the presence of larger 451 foraminifera and the occasional occurrence of smaller benthics suggest normal marine salinities 452 such as in facies Mf2 and DaMi.

Fossils of these facies and their taphonomic attributes, together with wackestone and packstone
textures and sedimentary structures, indicate low-energy hydrodynamic conditions in shallow-

455	water. The abundance of miliolids, together with spatangoids <i>Schizaster</i> and <i>Brissoides</i> (Fig. 3)
456	points to soft fine-sand and muddy substrates. The abundance of volcaniclastics in mixed facies
457	likely resulted in coarser substrates favorable to epifaunal and shallow endofaunal dwellers, such
458	as Antillaster and Eupatagus (Kanazawa, 1992; Donovan and Rowe, 2000). Although the final
459	depositional setting for facies EcF2 sediments is interpreted to be a shallow protected
460	environment (i.e., occurrence of spatangoid dwellers and mud-dominated deposits), the diverse
461	echinoid assemblage pointing to varied palaeoecological modes (i.e., from sand to muddy
462	substrates; Fig. 3) associated with high fragmentation and abrasion indices of the bioclastic
463	fraction implies a complex depositional history from high/moderate- to low-energy
464	hydrodynamic conditions.
465	Abundance of volcaniclastic elements in carbonate strata overlying volcaniclastic units across
466	erosive surfaces (see discussion below) suggests reworking of underlying beds through wave
467	action.
468	
469	6.2. Open inner ramp to middle ramp: wave-dominated settings
470	
471	Open inner ramp and transitional environments to middle ramp settings are dominated by wave
472	action. The influence of currents and high-energy waves can be deduced from the lack of easily
473	winnowed silts and micrite, relatively coarse grain sizes, moderate to high degrees of
474	fragmentation and abrasion of skeletons and lithoclasts, and cross-stratifications. In addition,
475	robust coral morphologies and overall thick-walled skeletons more resistant to wave-induced
476	transportation and to waterborne projectiles constitute useful indicators of strong hydrodynamic
477	conditions (Hughes 1087)
	conditions (flughes, 1987).

479 and CoP2 (Table 1). With the exception of chlorozoan facies (CoP2), most facies are

480 characterized by diverse faunal assemblages with varied degrees of alteration.

481 The dominance of biotic assemblages with robust larger foraminifera (Amphistegina,

482 *Nummulites*, and subordinate *Discocyclina*), coralline algal branches, subspheroid rhodoliths,

483 togehther with the presence of zooxanthellate corals and smaller benthic foraminifera, miliolids

484 and dasycladal algae, indicate deposition within the euphotic zone (Pomar, 2001; Nebelsick et al.,

485 2005; Afzar et al., 2011). The common occurrence of larger foraminifera argues against high

486 nutrient levels, since these foraminifera thrive in mesotrophic to principally oligotrophic waters

487 (Hottinger, 1997; Langer and Hottinger, 2000; Halfar et al., 2004).

488 Constituents of these coarse-grained facies (packstones, grainstones and sparitic rudstones)

489 possess moderate to high degrees of fragmentation and abrasion consistent with relatively high-

490 energy deposition above FWWB (Burchette and Wright, 1992). Particularly, the common

491 occurrence of broken, poorly-preserved and abraded larger foraminifera not only provides

492 evidence of high-energy conditions, but also may indicate repetitive burial and exhumation due to

493 frequent physical reworking (Beavington-Penney, 2004).

494 Low- to high-angle cross-stratifications may represent internal structures of subaqueous skeletal

495 dunes that developed through the reworking of bottom sediments including inner ramp protected

496 smaller foraminiferal – miliolid – dasycladalean sediments and deeper, more open, hyaline

497 foraminiferal – coralline algal deposits. Sigmoidal and hummocky cross-stratifications record

498 wave-dominated hydrodynamic conditions (Dumas and Arnott, 2006; Tinterri, 2011).

499 Similar inner to middle ramp packstones and grainstones dominated by larger foraminifera and

500 coralline algae with comparable sedimentary attributes and occasional coral occurrences have

501 been documented in south-east Asia from Eocene to middle Miocene limestones (Adams, 1965;

502 Wilson et al., 2000; Wilson, 2002; Wilson and Lokier, 2002), in the Indus Basin from Paleogene

limestones (Afzal et al., 2011), and in other Paleogene western Tethyan deposits (e.g., Loucks et
al., 1998) for example. In the Caribbean realm, the middle Eocene Swanswick Formation of the
White Limestone Group of Jamaica contains a diverse larger foraminiferal-algal assemblage with
common echinoderm and molluskan fragments that Mitchell (2004, 2013) similarly interpreted as
open marine high-energy platform deposits.

508

509 6.3. Distal middle ramp facies belt: episodic wave influence

510

511 Middle ramp deposits are represented by foralgal and rhodalgal associations of facies LfRh2, 512 LfRh3, Mf3, and LfCa2 (Table 1). The dominance of wackestones and packstones, and 513 subordinate grainstones are compatible with low-energy conditions allowing settlement of fine 514 particles, alternating with episodes of high-energy wave events (Burchette and Wright, 1992; 515 Nebelsick et al., 2005; Afzal et al., 2011). Abraded rhodoliths and the common occurrence of 516 algal debris, and other bioclastic, material are supportive of current or wave reworking. Similar 517 features have been recognized in open marine platforms influenced by sporadic storms (Lund et 518 al., 2000), and fossil examples have been described from Cenozoic successions (Braga and 519 Martin, 1988; Bassi, 2005; Barattolo et al., 2007; Payros et al., 2010). The regime of waves and 520 currents is an important factor that determines the shape and internal structure of rhodoliths (e.g., 521 Freiwald, 1995; Marrack, 1999). As such, the coalescence of rhodoliths (Fig. 6F) suggests times 522 of relative hydrodynamic quiescence between episodes of turbulence preventing burial by fine 523 sediments. The coralline algal assemblages are dominated by melobesioids with accessory 524 mastophoroids and sporolithaceans. Compared to facies interpreted as inner ramp deposits, there 525 is a taxonomic variation corresponding to a relative decrease in mastopohoroids (*Lithoporella*) 526 and sporolithaleans (Sporolithon), and an increase in melobesioids (Lithothamnion). Such

527	taxonomic changes in coralline assemblages showing a depth gradient have fossil analogues
528	(Rasser, 2000; Braga and Aguirre, 2001; Aguirre et al., 2017) and have been documented in
529	modern settings (Iryu et al., 1995). A distal middle ramp setting is compatible with the
530	foraminiferal assemblage dominated by nummulitids and orthophragminids, with subordinate
531	lepidocyclinids, rare amphisteginids and occasional planktonic foraminifera (Nebelsick et al.,
532	2005). In addition, the occasional occurrence of flattened test morphologies but also of light-
533	dependent organisms, and the abundance of coralline algae suggest depositional conditions below
534	FWWB and above SWB (Hottinger, 1997; Hohenegger et al., 2000; Bassi, 2005; Hohenegger,
535	2009).
536	
537	7. Discussion
538	
539	Specific to carbonate systems is that any significant change of either one of the parameters, (e.g.,
540	trophic resources, depth of light penetration, turbidity, nature of substrate, etc.) influencing
541	carbonate factories will result in either temporary or prolonged halts in carbonate production and
542	accumulation in different sectors of the shelf (Wilson, 2000; Wilson and Lokier, 2002; Wilson
543	and Vecsei, 2005; Lokier et al., 2009; Reuter et al., 2012). St. Barths limestones record the
544	repeated collapse and renewal of carbonate production, as a result of episodic volcaniclastic
545	material input (see below; Figs. 9, 10).
546	
547	7.1. Demise of carbonate factories
548	
549	Submersion and drowning caused by relative sea-level changes, environmental deterioration and
550	burial by clastics are major controls on the demise of tropical carbonate platforms (Schlager,

551 1989). Termination of St. Barths carbonate factories relates to volcaniclastic input with

accumulation rates exceeding rates of skeletal carbonate production and reef building until burial.

553 Erosional and sharp sedimentary surfaces between carbonate deposits below and volcaniclastic

above are the dominant stratigraphic expressions identified in the studied successions for the

555 demise of carbonate factories (Figs. 11A-D, 12).

556 Erosional surfaces are overlain by polymictic matrix-supported breccias (Facies Vbr), and 557 represent syneruptive surfaces generated by the discharge of coarse volcanic material in the 558 carbonate realm (Figs. 2B, 5B, 10A, 11A-B) (Trofimovs et al., 2006). Sharp surfaces represent 559 sedimentation events (Reading, 1996) and coincide with the abrupt transition from carbonate to 560 volcaniclastic sedimentation, dominated by hyaloclastites of facies Hy1, with minor if any 561 evidence of erosion and bioturbation (Figs. 11C, 12A). Such sharp non-erosional surfaces have 562 been interpreted to reflect insufficient skeletal production of carbonate systems compared to the 563 rates of clastic accumulation (Schlager, 1989; Wilson, 2002; Lokier et al., 2009). Suffocation of 564 carbonate factories by volcaniclastics was likely facilitated by the rarity of significant coral reef 565 buildups and the predominance of slow carbonate producing biota (Reuter, 2011), such as 566 foraminifera and coralline red algae, in the St. Barths limestones.

567 Gradational contacts also occur and represent progressive burial of carbonate systems by 568 volcaniclastics of facies Hy1 and Hy2 (Figs. 2C, 11D, 12B). They indicate that prolongation of 569 carbonate production during volcaniclastic input was possible provided a change of benthic 570 communities from soft muddy to gravelly-shelly substrate dwellers having the capability to 571 escape burial such as irregular echinoids (from LfCa1 to facies Mf4; Fig. 11D). Low- to high-572 angle cross-stratifications in the overlying volcanicastic deposits (Figs. 11D, 12B) suggest that 573 volcaniclasts were transferred, considering the proximity of volcanic centers (Fig. 1), downslope 574 via dunes, sand waves or in avalanche flows (Caron et al., 2004; Dorobek, 2008; Puga-Bernabéu, 575 2010).

576

577 7.2. Temporary halts in carbonate production and deposition

578

Within the studied Eocene carbonate successions burrowed surfaces, interpreted as omission
surfaces, are of particular interest since some of them are associated with tuffitic layers (Figs. 9,
10B, 11H-J, 12).

582 Omission surfaces are emphasized by a complex network of *Thalassinoides* burrows reaching 583 their maximum density beneath the surface and decreasing downwards (Figs. 2D, 11I, 12C-H). 584 Most published works relate omission surfaces to stratigraphic condensation during periods of 585 slowed/non-deposition (e.g., Clari et al., 1995; Hillgärtner, 1998; Christ et al., 2012). Sediments 586 by-pass, drained siliciclastic sources, wave- and/or current-induced sediment transfer and 587 submarine erosion are major factors considered to promote non-deposition on the sea floor. Previous authors have tied the formation of omission surfaces to specific portions of relative sea-588 589 level cycles where the position of the sea floor relative to the wave base is considered a key 590 parameter to the onset of omission (Lukasik and James, 2003; Christ et al., 2012). However, an 591 alternative explanation for the genesis of omission surface can be proposed based on the 592 assumption that starved sedimentation may also result from the interruption of *in situ* carbonate 593 production below and above fair-weather wave base. Although some organisms (e.g., Bivalves 594 and Echinoids) have the ability to extricate themselves if buried beneath sediments and others to 595 remove fine particles from their surface (Hinchey et al., 2006; Lokier et al., 2009; Knoll et al., 596 2017), episodic volcanic eruptions delivering high amounts of volcaniclastic material including 597 ash (Fig. 10B) will result in rapid physical burial and consequent mass-mortality of benthic 598 communities (Heikoop et al., 1996; Tomascik et al., 1997; Alve, 1999; Reuter and Piller, 2011).

599 The formation of omission surfaces depends on the duration of sediment starvation, and therefore 600 here the time span between volcanic events and the recovery of carbonate factories. Where 601 volcaniclastics were deposited above wave base, resuspension of their finer portion is likely to 602 result in higher turbidity and in reduced light levels, thereby delaying the reestablishment of 603 photodependent organisms (Hallock and Glenn, 1986). In addition, the potential increase of 604 nutrient input through chemical alteration of volcanic particles may cause eutrophication at the 605 sea floor and prevent colonization by oligotrophic and mesotrophic communities (Hallock and 606 Schlager, 1986; Allmon, 1992; Brasier, 1995a; Hallock, 2001; Reuter and Piller, 2011). The time 607 frame for either full recovery of the benthic communities that existed prior to the volcanic event 608 or the renewal of carbonate systems via a switch to different biotic assemblages is difficult to 609 estimate from the geological record but examples in the literature point to reestablishment over 610 decades and centuries (Heikoop et al., 1996; Tomascik et al., 1996; Pandolfi et al., 2006; Reuter 611 et al., 2011; Vroom and Zgliczynski, 2011). This time frame is compatible with the development 612 of a dense burrowing network and of omission (Dashtgard et al., 2008).

613

614 7.3. Recovery and renewal of carbonate factories (Fig. 13)

615

Besides light conditions and nutrient levels, periods of volcanic quiescence and reduced
availability of volcaniclastic material were likely the principal promoting conditions for the
renewal of carbonate production. In the studied successions, mixed facies (Mf1-Mf3) followed by
open marine foralgal (larger foraminifera-coralline algal facies; LfCa) and rhodalgal facies
(LfRh) occur above erosional surfaces cut in sandy to gravely volcaniclastic substrates (Figs.
11F, 13C). Protected lagoonal foralgal facies (dasycladalean-miliolid facies; DaMi) are found
above omission surfaces capping silty-sandy substrates and above marly concretionary horizons

623 grading from sand-sized volcaniclastics (Figs. 2C, 11E, 13E-F). Echinofor facies (echinoids-624 larger foraminifera; EcF) capped by coral carpets locally represent the initial carbonate 625 development above gradational contacts. A review of foundering photozoan carbonate facies 626 described in the literature shows similar colonizing biota on siliciclastic substrates (e.g., Hendry 627 et al., 1999; Wilson, 2002; Burton, 2004; Lokier et al., 2009; Reuter and Piller, 2011; Reuter et 628 al., 2012). Rhodolith-bearing units similar to those reported here (Fig. 11F), and located above 629 erosional unconformities cut in rocky and clastic substrates, have been interpreted as 630 transgressive deposits sitting on surfaces of marine erosion (e.g., Nalin et al., 2007). Gradational 631 contacts have been inferred to record a progressive decrease of siliciclastic input and subsequent 632 colonization by carbonate producers reflected in transitional mixed facies (e.g., Hendry et al., 633 1999; Burton, 2004; Lokier et al., 2009). 634 Characteristics of the renewal of carbonate production described here suggest that they may result 635 from a combination of factors promoting the reduction of volcaniclastic influx (e.g., changes in 636 sediment transport direction and volcanic quiescence) and relative sea-level changes. Marly 637 horizons at the volcaniclastic-carbonate interface, erosional and sharp-cut surfaces (Fig. 13A-B) 638 are compatible with marine flooding. However, deepening-upward trends in facies above these 639 surfaces have not been observed. This discrepancy might be explained by a slow rise of relative 640 sea-level allowing aggradation without significant change of carbonate-producing biota. 641

642 8. Conclusions

643

644 The main results of the present study and their broader implications for the analysis of carbonate645 successions in volcanically-active settings may be summarized as follows.

646 1 The middle Eocene limestones of St. Bartholomew Island contain marine depositional facies

647 characteristic of inner to distal middle ramp settings affected by episodic volcanic eruptions. The 648 variety of carbonate-producing biota (i.e., chlorozoan, chloralgal, foralgal), in terms of their 649 dependence to light intensity and tolerance to nutrient levels, suggests that carbonate production 650 occurred mostly in euphotic conditions across an unrimmed platform with isolated coral 651 bioherms and shoals. Other biogenic contributors (i.e., rhodalgal, echinofor) are consistent with 652 oligophotic conditions related to resuspended fine material after storms or fine volcaniclastics 653 input following eruption that diminished water transparency. The St. Barths Eocene carbonates 654 provide an example of low-latitude tropical platform where non-framework building biota were 655 important, if not dominant, sediment contributors, compared to other Eocene carbonate deposits. 656 2 This study shows that the effects of volcanism on carbonate factories range from burial and 657 prolonged termination of carbonate factories, to temporarily arrested or slowed carbonate 658 production. When volcaniclastic inputs ceased, carbonate systems reestablished. Foundering 659 biotic communities include euphotic chlorozoan facies, oligophotic foralgal, rhodalgal and 660 echinofor facies mixed in various proportions with volcaniclastics. 661 3 The discontinuous nature of the carbonate sedimentation in relation to volcaniclastic influx is 662 reflected in: (a) erosional and sharp sedimentary surfaces that mark pronounced facies changes; 663 and, (b) gradational contacts. 664 Some sedimentary surfaces and contacts located at volcaniclastic-carbonate boundaries record the 665 renewal of carbonate production above volcaniclastic deposits, while others record either the 666 temporary halt of carbonate production or the demise of carbonate platforms buried beneath thick 667 volcaniclastics. 668 The demise of carbonate factories and their prolonged shut-down in the vicinity of active

volcanoes originated in the propagation of syneruptive pyroclastic flows or in the

670 progradation/migration of wave-reworked volcaniclastic material, eroding and burying carbonate

671	deposits. Temporary shut-down of <i>in situ</i> carbonate production also occurred and was likely a
672	result of environmental stresses caused by fine volcaniclastics, including burial, lowered light
673	levels and eutrophication.
674	The recovery of carbonate factories resulted from an interruption of volcaniclastic influx at the
675	onset of volcanic quiescence promoting the reduction of volcaniclastic influx at the locus of
676	renewed carbonate production.
677	Future investigations should concentrate, in a broader regional context, on the characterization of
678	key sedimentary surfaces and discontinuities in mixed volcaniclastic-carbonate successions of the
679	Volcanic Arc Antillean area. Correctly interpreting the origin and significance of discontinuities
680	in carbonates deposited adjacent to active volcanoes is important to avoid misinterpretation of
681	other key surfaces in a cyclostratigraphic and sequence stratigraphic context.
682	
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689	
690	References
691	
692	Adams, C.G., 1965. The foraminifera and stratigraphy of the Melinau Limestone, Sarawak, and

693 its importance in Tertiary correlation. Quarterly Journal of the Geological Society of London694 121, 283-338.

696	Afzal, J., Williams, M., Leng, M.J., Aldridge, R.J., 2011. Dynamic response of the shallow
697	marine benthic ecosystem to regional and pan-Tethyan environmental change at the
698	Paleocene Eocene boundary. Palaeogeography Palaeoclimatology Palaeoecology 309, 141-
699	160.
700	
701	Aguirre J., Braga J.C., Bassi D., 2017. The role of rhodoliths and rhodolith beds in the rock
702	record and their use in palaeoenvironmental reconstructions. In : Riosmena-Rodriguez R.,
703	Nelson W., Aguirre J. (Eds), Rhodolith/maerl beds: a global perspective. Springer-Verlag,
704	Berlin, special volume, pp. 105-138
705	
706	Allmon, W.D., 1992. Role of temperature and nutrients in extinction of turritelline gastropods:
707	Cenozoic of the northwestern Atlantic and northeastern Pacific. Palaeogeography
708	Palaeoclimatology Palaeoecology 92, 41-54.
709	
710	Alve, E., 1999. Colonization of new habitats by benthic foraminifera: a review. Earth-Science
711	Reviews 46, 167-185.
712	
713	Barattolo, F., Bassi, D., Romano, R., 2007. Upper Eocene larger foraminiferal-coralline algal
714	facies from the Klokova Mountain (southern continental Greece). Facies 53, 361-375.
715	
716	Bassi, D., 1998. Coralline algal facies and their palaeoenvironment in the Late Eocene of
717	Northern Italy (Calcare di Nago, Trento). Facies 39, 179-202.
718	

719	Bassi, D., 2005. Larger foraminiferal and coralline algal facies in an Upper Eocene storm-
720	influenced, shallow-water carbonate platform (Colli Berici, north-eastern Italy).
721	Palaeogeography Palaeoclimatology Palaeoecology 226, 17-35.
722	
723	Beavington-Penney, S.J., 2004. Analysis of the effects of abrasion on the tests of
724	Palaeonummulites venosus: implications for the origin of nummulithoclastic sediments.
725	Palaios 19, 143-155.
726	
727	Beavington-Penney, S.J., Racey, A., 2004. Ecology of extant nummulitids and other larger
728	benthic foraminifera: applications in palaeoenvironmental analysis. Earth-Science Reviews
729	67, 219-265.
730	
731	Berra, F., Carminati, E., 2012. Differential compaction and early rock fracturing in high-relief
732	carbonate platforms : numerical modelling of a Triassic case study (Esino limestone, central
733	southern Alps, Italy). Basin Research 24, 598-614.
734	
735	Bouysse, P., Westercamp, D., Andreieff, P., 1990. The Lesser Antilles island arc. In: Moore, J.C.,
736	Mascle, A. et al. (Eds.) Proceedings of ODP Science Research 110, pp. 29-44.
737	
738	Burchette, T.P., Wright, V.P., 1992. Carbonate ramp depositional systems. Sedimentary Geology
739	79, 3-57.
740	
741	Braga, J.C., Martìn, J.M., 1988. Neogene coralline-algal growthforms and their
742	palaeoenvironments in the Almanzora River Valley (Almeria S.E. Spain). Palaeogeography

Palaeoclimatology Palaeoecology 67, 285-303.

745	Braga, J.C., Aguirre, J., 2001. Coralline algal assemblages in Upper Neogene reef and temperate
746	carbonates in southern Spain. Palaeogeography Palaeoclimatology Palaeoecology 175, 27-
747	41.
748	
749	Braga, J.C., Vescogni, A., Bosellini, F.R., Aguirre, J., 2009. Coralline algae (Corallinales,
750	Rhodophyta) in western and central Mediterranean Messinian reefs. Palaeogeography.
751	Palaeoclimatology Palaeoecology 275, 113-128.
752	
753	Brasier, M.D., 1995a. Fossil indicators of nutrient levels. 1. Eutrophication and climate change.
754	In: Bosence, D.W.J., Allison, P.A. (Eds.), Marine Palaeoenvironmental Analysis from
755	Fossils, Geological Society of London Special Publication 83, pp. 113-132.
756	
757	Brasier, M.D., 1995b. Fossil indicators of nutrient levels. 2. Evolution and extinction in relation
758	to oligotrophy. In: Bosence, D.W.J., Allison, P.A. (Eds.), Marine Palaeoenvironmental
759	Analysis from Fossils, Geological Society of London Special Publication 83, pp. 133-150.
760	
761	Burton, L.M., 2004. Carbonate-siliciclastic Interactions; Tertiary Examples from Spain. PhD
762	Thesis, University of Durham, UK, 396 p.
763	
764	Carannante, G., Esteban, M., Milliman, J.D., Simone, L., 1988. Carbonate lithofacies as
765	paleolatitude indicators: problems and limitations. Sedimentary Geology 60, 333-346.
766	

767	Caron, V., 2011. Contrasted textural and taphonomic properties of high-energy wave deposits
768	cemented in beachrocks (St. Bartholomew Island, French West Indies). Sedimentary
769	Geology 237, 189-208.
770	
771	Caron, V., Nelson, C.S., Kamp, P.J.J., 2004. Contrasted carbonate depositional systems for
772	Pliocene cool-water limestones cropping out in central Hawke's Bay, New Zealand. New
773	Zealand journal of Geology and Geophysics 47, 697-717.
774	
775	Christ, N., Immenhauser, A., Amour, F., Mutti, M., Tomás, S., Always, R., Kabir, L., 2012.
776	Characterization and interpretation of discontinuity surfaces in a Jurassic ramp setting (High
777	Atlas, Morocco). Sedimentology 59, 249-290.
778	
779	Christman, R.A., 1953. Geology of S ^t . Bartholomew, S ^t . Martin, and Anguilla, Lesser Antilles.
780	Bulletin of the Geological Society of America 6, 65-96.
781	
782	Clari, P.A., Dela Pierre, F., Martire, L., 1995. Discontinuities in carbonate successions:
783	identification, interpretation and classification of some Italian examples. Sedimentary
784	Geology 100, 97–121.
785	
786	Cosovic, V., Drobne, K., Moro, A., 2004. Paleoenvironmental model for Eocene foraminiferal
787	limestones of the Adriatic carbonate platform (Istrian Peninsula). Facies 50, 61-75.
788	
789	Dashtgard, S.E., Gingras, M.K., Pemberton, S.G., 2008. Grain-size controls on the occurrence of
790	bioturbation. Palaeogeography Palaeoclimatology Palaeoecology 257, 224-243.

792	Donovan, S.K., Rowe, DA.C., 2000. Spatangoid echinoids from the Eocene of Jamaica. Journal
793	of Paleontology 74, 654-661.
794	
795	Dorobek, S.L., 2008. Carbonate-platform facies in volcanic-arc settings: Characteristics and
796	controls on deposition and stratigtraphic development. Geological Society of America
797	Special Papers 436, 55-90.
798	
799	Dumas, S., Arnott, R.C.W., 2006. Origin of hummocky and swaley cross-stratifications. The
800	controlling influence of unidirectional current strength and aggradation rate. Geology 34,
801	1073-1076.
802	
803	Embry, A.F., Klovan, J.E., 1971. A late Devonian reef tract on northeastern Banks Island.
804	N.W.T. Bulletin of Canadian Petroleum Geology 19, 730-781.
805	
806	Freiwald, A., 1995. Sedimentological and biological aspects in the formation of branched rhodoliths
807	in northern Norway. Beiträge zur Paläontologie 20, 7-19.
808	
809	Gaillard, C., 1983. Les biohermes à spongiaires et leur environnement dans l'Oxfordien du Jura
810	méridional. Document du Laboratoire de Géologie de Lyon 90, 515 pp.
811	
812	Geel, T., 2000. Recognition of stratigraphic sequences in carbonate platform and slope deposits:
813	empirical models based on microfacies analysis of Palaeogene deposits in southeastern
814	Spain. Palaeogeography Palaeoclimatology Palaeoecology 155, 211-238.

816	Gradstein, F.M, Ogg, J.G., Schmitz, M.D., et al., 2012. The Geologic Time Scale 2012: Boston,
817	USA, Elsevier, DOI: 10.1016/B978-0-444-59425-9.00004-4
818	
819	Halfar, J., Godinez-Orta, L., Mutti, M., Valdez-Holguin, J.E., Borges, J.M., 2004. Nutrient and
820	temperature controls on modern carbonate production: An example from the Gulf of
821	California, Mexico. Geology 32, 213-216.
822	
823	Hallock, P., 1985. Why are larger foraminifera large? Paleobiology 11, 195-208.
824	
825	Hallock, P., 1987. Fluctuations in the trophic resource continuum: a factor in global diversity
826	cycles? Palaeoceanography 2, 457-471.
827	
828	Hallock, P., 2001. Coral reefs, carbonate sedimentation, nutrients, and global change. In: Stanley,
829	G.D. (Ed.), The History and Sedimentology of Ancient Reef Ecosystems, Amsterdam
830	(Kluwer), pp. 387-427.
831	
832	Hallock, P., Glenn, E.C., 1986. Larger foraminifera: a tool for palaeoenvironmental analysis of
833	Cenozoic depositional facies. Palaios, 1, 55-64.
834	
835	Hallock, P., Schlager, W., 1986. Nutrient excess and the demise of coral reefs and the demise of
836	coral reefs and carbonate platforms. Palaios 1, 389-398.
837	
838	Hallock, P., Sheps, K., Chapronière, G., Howell, M, 2006. Larger benthic foraminifers of the

839	marion plateau, northeastern Australia (ODP leg 194): comparison of faunas from bryozoan
840	(sites 1193 and 1194) and red algal (sites 1196-1198) dominated carbonate platforms. In:
841	Anselmetti, F.S., Isern, A.R., Blum, P., Betzler, C. (Eds.), Proceedings of the Ocean Drilling
842	Program, Scientific Results 194, pp. 1-31.
843	
844	Harvey, A.S., Broadwater, S.T., Woelkerling, W.J., Mitrovski, P.J., 2003. Choreonema
845	(Corallinales, Rhodophyta): 18S rDNA phylogeny and resurrection of the Hapalidiaceae for
846	the subfamilies Choreonematoideae, Austrolithoideae, and Melobesioideae. Journal of
847	Phycology 39, 988-998.
848	
849	Hayton, S., Nelson, C.S., Hood, S.D., 1995. A skeletal assemblage classification system for non-
850	tropical carbonate deposits based on New Zealand Cenozoic limestones. Sedimentary
851	Geology 100, 123-141.
852	
853	Heikoop, J.M., Tsujita, C.J., Heikoop, C.E., Risk, M., Dickin, A.P., 1996. Effects of volcanic
854	ashfall recorded in ancient marine benthic communities: comparison of a nearshore and an
855	offshore environment. Lethaia 29, 125-139.
856	
857	Hendry, J.P., Taberner, C., Marshall, J.D., Pierre, C., Carey, P.F., 1999. Coral reef diagenesis
858	records pore-fluid evolution and paleohydrology of a siliciclastic basin margin succession
859	(Eocene South Pyrenean foreland basin, northeastern Spain). Geological Society of America
860	Bulletin 111, 395-411.
861	

863	Valanginian, France and Switzerland). Journal of Sedimentary Research 68, 1093-1108.
864	
865	Hinchey, E., Schaffner, L., Hoar, C., Vogt, B., Batte, L., 2006. Responses of estuarine benthic
866	invertebrates to sediment burial: the importance of mobility and adaptation. Hydrobiologia
867	556, 85-98.
868	
869	Hohenegger, J., 2009. Functional shell geometry of symbiont-bearing foraminifera. Research
870	Galaxea, Journal of Coral Reef Studies 11, 81-89.
871	
872	Hohenegger, J., Yordanova, E., 2001. Displacement of larger Foraminifera at the western slope
873	of Motobu Peninsula (Okinawa, Japan). Palaios 16, 53-72
874	
875	Hohenegger, J., Yordanova, E., Hatta, A., 2000. Remarks on West Pacific Nummulitidae
876	(Foraminifera). Journal of Foraminiferal Research 30(1), 3-28.
877	
878	Hottinger, L., 1983. Processes determining the distribution of larger foraminifera in space and
879	time. Utrecht Micropaleontogy Bulletin 30, 239-253.
880	
881	Hottinger, L., 1997. Shallow benthic foraminiferal assemblages as signals for depth of their
882	deposition and their limitations. Bulletin de la Société Géologique de France 168, 491-505.
883	
884	Hughes, T.P., 1987. Skeletal density and growth form of corals. Marine Ecology – Progress
885	Series 35, 259-266.
886	

887	Iryu, Y., Nakamori, T., Matsuda, S., Abe, O., 1995. Distribution of marine organisms and its
888	geological significance in the modern reef complex of the Ryukyu Islands. Sedimentary
889	Geology 99, 243-258.
890	
891	James, N.P., 1997. The cool-water carbonate depositional realm. In: James, N.P., Clarke, J.
892	(Eds.), Cool-Water Carbonates. SEPM Special Publication 56, pp. 1-20.
893	
894	Jeans, C.V., Wray, D.S., Merriman, R.J., Fisher, M.J., 2000. Volcanogenic clays in Jurassic and
895	Cretaceous strata of England and the North Sea Basin. Clay Minerals 35, 25-55.
896	
897	Kanazawa, K., 1992. Adaptation of test shape for burrowing and locomotion in spatangoid
898	Echinoids. Paleontology 35, 733-750.
899	
900	Kier, P.M., 1984. Fossil Spatangoid Echinoids of Cuba. Smithsonian Contribution to
901	Paleobiology 55, 336 pp.
902	
903	Knoll, K., Chamberlain, R.B., Chamberlain, J.A. Jr., 2017. Escape burrowing of modern
904	freshwater bivalves as a paradigm for escape behavior in the Devonian bivalve Archanodon
905	catskillensis. Geosciences 7, 37 pp.
906	
907	Langer, M.R., Hottinger, L., 2000. Biogeography of selected larger foraminifera.
908	Micropaleontology 46, 105-126.
909	

910 Lees, A., 1975. Possible influence of salinity and temperature on modern shelf carbonate

sedimentation. Marine Geology 19, 159-198.

- Lees, A., Buller, A.T., 1972. Modern temperate water and warm water shelf carbonate sediments
 contrasted. Marine Geology 13, M67–M73.
- 915
- Lokier, S.W., Wilson, M.E.J., Burton, L.M., 2009. Marine biota response to clastic sediment
 influx: a quantitative approach. Palaeogeography Palaeoclimatology Palaeoecology 175,
 249-272.
- 919
- Loucks, R.G., Moody, R.T.J., Bellis, J.K., Brown, A.A., 1998. Regional depositional setting and
 pore network systems of the El Garia Formation (Metlaoui Group, Lower Eocene), offshore
 Tunisia. In: Macgregor, D.S., Moody, R.T.J., Clark-lowes, D.D. (Eds.), Petroleum geology
 of North Africa, Geological Society of London Special Publication 132, pp. 355-374.
- 924
- Lukasik, J.J., James, N.P., 2003. Deepening-upward subtidal cycles, Murray Basin, South
 Australia. Journal of Sedimentary Research 73, 653-671.
- 927
- Lund, M., Davies, P.J., Braga, J.C., 2000. Coralline algal nodules off Fraser Island eastern
 Australia. Facies 42, 25-34.
- 930
- Marrack, E.C., 1999. The relationship between water motion and living rhodolith beds in the
 southwestern Gulf of California, Mexico. Palaios 14, 159-171.
- 933
- 934 Mitchell, S.F., 2004. Lithostratigraphy and palaeogeography of the White Limestone Group.

Cainozoic Research 3, 5-29.

936

937 Mitchell, S.F., 2013. Stratigraphy of the White Limestone of Jamaica. Bulletin de la Société
938 Géologique de France 184, 111-118.

939

Murray, J.W., 1991. Ecology and distribution of benthic foraminifera. In: Lee, J.J., Anderson,
R.O. (Eds.), Biology of Foraminifera. Academic Press, London, pp. 221-284.

942

Nalin, R., Massari, F., 2009. Facies and stratigraphic anatomy of a temperate carbonate sequence
(Capo Colonna terrace, Late Pleistocene, Southern Italy). Journal of Sedimentary Research
79, 210-225.

946

947 Nalin, R., Nelson, C.S., Basso, D., Massari, F., 2007. Rhodolith-bearing limestones as
948 transgressive marker beds: fossil and modern examples from North Island, New Zealand.
949 Sedimentology 55, 249-274.

950

- 951 Nebelsick, J.H., Bassi, D., 2000. Diversity, growth forms and taphonomy: key factors controlling
- 952 the fabric of coralline algae dominated shelf carbonates. In: Insalaco, E. Skelton, P.W.,
- 953 Palmer, T.J. (Eds.), Carbonate Platform Systems: components and interactions, Geological
- 954 Society of London Special Publication 178, pp. 89-107.

955

956 Nebelsick J.H., Rasser M., Bassi D. 2005. Facies dynamics in Eocene to Oligocene circumalpine
957 carbonates. Facies, 51, 197-216.

959	Pandolfi, J.M., Tudhope, A.W., Burr, G., Chapell, J., Edinger, E., Frey, M., Steneck, R., Sharma,
960	C., Yeates. A., Jennions, M., Lescinsky, H., Newton. A., 2006. Mass mortality following
961	disturbance in Holocene coral reefs from Papua New Guinea. Geology 34, 949-952.
962	
963	Payros, A., Pujalte, V., Tosquella, J., Orue-Etxebarria, V., 2010. The Eocene storm-dominated
964	foralgal ramp of the western Pyrenees (Urbasa-Andia Formation): An analogue of future
965	shallow-marine carbonate systems? Sedimentary Geology 228, 184-204.
966	
967	Perry, C.T., 1998. Grain susceptibility to the effects of microboring: implications for the
968	preservation of skeletal carbonates: Sedimentology 45, 39-51.
969	
970	Pomar, L., 2001. Ecological control of sedimentary accommodation: evolution from carbonate
971	ramp to rimmed shelf, Upper Miocene, Balearic Islands. Palaeogeography Palaeoclimatology
972	Palaeoecology 175, 249-272.
973	
974	Pomar, L., Brandano, M., Westphal, H., 2004. Environmental factors influencing skeletal grain
975	sediment associations: a critical review of Miocene examples from the western
976	Mediterranean. Sedimentology 51, 627-651.

978	Puga-Bernabéu, Á., Martín, J.M., Braga, J.C., Sánchez-Almazo, I.M., 2010. Downslope-
979	migrating sandwaves and platform-margin clinoforms in a current-dominated, distally
980	steepened temperate-carbonate ramp (Guadix Basin, Southern Spain). Sedimentology 57,
981	287-311.
982	
983	Rasser, M.W., 2000. Coralline red algal limestones of the Late Eocene Alpine Foreland Basin in
984	Upper Austria: component analysis, facies and paleoecology. Facies 42, 59-92.
985	
986	Rasser, M.W., Piller, W.E., 1999. Application of neontological taxonomic concepts to Late
987	Eocene coralline algae (Rhodophyta) of the Austrian Molasse Zone. Journal of
988	Micropalaeontology 18, 67-80.
989	
990	Reading, H.G., 1996. Sedimentary environments: Processes, facies and stratigraphy. Blackwell
991	Science, Oxford.
992	
993	Reuter, M., Piller, W.E., 2011. Volcaniclastic events in coral reef and seagrass environments:
994	evidence for disturbance and recovery (Middle Miocene, Styrian Basin, Austria). Coral
995	Reefs 30, 889-899.
996	
997	Reuter, M., Piller, W.E., Erhart, C., 2012. A Middle Miocene carbonate platform under silici-
998	volcaniclastic sedimentation stress (Leitha Limestone, Styrian Basin, Austria) - Depositional
999	environments, sedimentary evolution and palaeoecology. Palaeogeography
1000	Palaeoclimatology Palaeoecology 350-352, 198-211.

1002	Rusciadelli, G., Di Simone, S., 2007. Differential compaction as control on depositional
1003	architectures across the Maiella carbonate platform margin (central Appenines, Italy).
1004	Sedimentary Geology 196, 133- 155.
1005	
1006	Schlager, W., 1989. Drowning unconformities on carbonate platforms. In: Crevello, P.D.,
1007	Wilson, J.L., Sarg, J.F., Read, J.F. (Eds.), Controls on Carbonate Platforms and Basins
1008	Development. SEPM Special Publication 44, pp. 15-25.
1009	
1010	Sola, F., Braga, J.C., Aguirre, J., 2013. Hooked and tubular coralline algae indicate seagrass beds
1011	associated to Mediterranean Messinian reefs (Poniente Basin, Almería, SE Spain).
1012	Palaeogeography Palaeoclimatology Palaeoecology 374, 218-229.
1013	
1014	Tinterri, R., 2011. Combined flow sedimentary structures and the genetic link between
1015	sigmoidal- and hummocky- cross stratification. GeoActa 10, 43-85.
1016	
1017	Tomascik, T., van Woesik, R., Mah, A.J., 1996. Rapid coral colonization of a recent lava flow
1018	following a volcanic eruption, Banda Islands, Indonesia. Coral Reefs 15, 169-175.
1019	
1020	Trofimovs, J., Amy, L., Boudon, G., Deplus, C. Doyle, E., Fournier, N., Hart, M.B., et al. 2006.
1021	Submarine pyroclastic deposits formed at the Soufrière Hills volcano, Montserrat (1995-
1022	2003): What happens when pyroclastic flows enter the ocean? Geology 34, 549-552.
1023	
1024	Vecchio, E., Hottinger, L., 2007. Agglutinated conical foraminifera from the Lower-Middle

Eocene of the Trentinara Formation (southern Italy). Facies 53, 509-533.

1026

1027 Vroom, P.S., Zgliczynski, B.J., 2011. Effects of volcanic ash deposits on four functional groups
1028 of a coral reef. Coral Reefs 30, 1025-1032.

1029

1030 Westercamp, D., Andreieff, P., 1983a. St-Barthélémy et ses îlets, Antilles françaises:

1031 stratigraphie et evolution magmato-structurale. Bulletin de la Société Géologique de France
1032 25, 873-883.

1033

1034 Westercamp, D., Andreieff, P., 1983b. Saint-Barthélémy et ses îlets - Carte géologique à
 1035 1/20000: Notice explicative. Service Géologique National, Orléans, France, 38 pp.

1036

1037 Wilson, M.E.J., 2002. Cenozoic carbonates in Southeast Asia: implications for equatorial
1038 carbonate development. Sedimentary Geology 147, 295-328.

1039

Wilson, M.E.J., Bosence, D.W.J., Limbong, A., 2000. Tertiary syntectonic carbonate platform
development in Indonesia. Sedimentology 47, 395-419.

1042

Wilson, M.E.J., Lokier, S.W., 2002. Siliciclastic and volcaniclastic influences on equatorial
carbonates: insights from the Neogene of Indonesia. Sedimentology 49, 583-601.

1045

Wilson, M.E.J., Vecsei, A., 2005. The apparent paradox of abundant foramol facies in low
latitudes: their environmental significance and effect on platform development. EarthScience Reviews 69, 133-168.

1050	Zamagni, J., Mutti, M., Kosir, A., 2008. Evolution of shallow benthic communities during the
1051	Late Paleocene-earliest Eocene transition in the Northern Tethys (SW Slovenia). Facies 54,
1052	25-43.
1053	
1054	Figure captions
1055	
1056	Fig. 1. (A) Location of St. Bartholomew relative to the main geological features of the Lesser
1057	Antilles in the eastern Caribbean. (B) Simplified lithostratigraphy of the Middle Eocene of St
1058	Bartholomew showing the main volcanic and limestone units. Not to scale. (C) Map showing
1059	outcropping distribution of limestone units, key stratigraphical sections (arrowed) and inferred
1060	location of eruptive centers (based on Westerkamp and Anfreieff, 1983b).
1061	
1062	Fig. 2. Examples of stratigraphic columns logged in the studied Eocene limestones. See Fig. 1 for
1063	locations.
1064	
1065	Fig. 3. Paleoecological modes of the most frequent irregular echinoids found in the St.
1066	Bartholomew middle Eocene limestones (based on Kanazawa, 1992).
1067	
1068	Fig. 4. Taphonomic grading system established for several skeletal categories. The scale of
1069	fragmentation features for benthic foraminifers derives from that proposed by Beavington-
1070	Penney (2004). The scale of mechanical alteration for calcareous green algae is modified from
1071	Caron (2011). The importance of test infestation follows the visual chart presented in Perry
1072	(1998).



1096 reef possibly exhibiting several growth-phases (arrows), each one being laterally correlated to

1097	limestone beds. Location: Anse des Cayes. (B) Massive coral buildup, 8 m thick (dashed line),
1098	interpreted as an isolated coral bioherm. Coral heads are in living position in the upper part of the
1099	structure whereas it is not as clear in its lower part. Location: Pain de sucre.
1100	
1101	Fig. 8. Hypothetical paleoenvironmental reconstruction and depositional profile of the ramp
1102	attached to submarine volcanoes for the middle Eocene limestones of St. Bartholomew.
1103	
1104	Fig. 9. Anse des Cayes section (Limestone unit 3 - see Fig. 1 for location and Fig. 12 for field
1105	photograph) showing the interplay between volcaniclastic and carbonate sedimentation. t. arr. :
1106	temporarily arrested.
1107	
1108	Fig. 10. Consequences of the onset of volcanic activity on inner- to middle-ramp carbonate
1109	factories. A) Demise due to burial following emplacement of coarse-grained pyroclastic deposits
1110	(facies Vbr and Hy; Table 1); B) Collapse due to burial by volcanic ash (facies Tu; Table 1).
1111	
1112	Fig. 11. Detailed portions of lithologic logs (see Fig. 2) showing the vertical facies successions
1113	within the deposits associated with carbonate production breaks and renewal.
1114	
1115	Fig. 12. Sedimentary surfaces recording interruption of carbonate production. See Figure 1 for
1116	locations. (A) Sharp contact (sc) between lagoonal facies below and sand-sized volcaniclastics
1117	above. Hammer: 30 cm. Location: Pointe Mangin. (B) Gradational contact (gc) between mixed
1118	carbonate-volcaniclastic deposits (Mf) below and cross-bedded hyaloclastites above (Hy2). The
1119	contact with the limestone unit (Lu) above is described in Figure 11E. Location: Anse des
1120	Flamands. (C, D) Burrowed omission surface (bs). Close-up view in D. Scale: 12 cm. (E)

Burrowed surface (bs). Iron mineralization is pervasive in the 50 cm-thick layer above. Location:
Anse Gouverneur. (F) Burrowed omission surface seen in bedding plane on the right and
overlying iron-stained deposit on the left. Arrow points to iron mineralization of burrow walls.
Scale: 12 cm. (G) Close-up view of iron-stained deposit in E and F. Irregular echinoids (Ec) of
different paleoecological modes (see text for details) are abundant together with pinnid fragments
(Pi). Scale: 12 cm.

1127

1128 Fig. 13. Sedimentary surfaces recording the renewal of carbonate production after prolonged 1129 interruption due to volcanic activity. See Figure 1 for locations. (A) Sharp (sc), slightly 1130 undulating, weakly burrowed surface at volcaniclastic-carbonate boundary. Location: Anse des 1131 Cayes. (B) Lateral correlative surface seen in B. (C) Erosional surface (es) with a relief up to 10 1132 cm. From the surface, fine-grained volcaniclastic deposits below are densely bioturbated by 1133 Thalassinoides (See D). Location: Pointe Mangin. (D) Close up view of Thalassinoides burrows 1134 beneath the erosional surface in C. Scale: 12 cm. (E-F) Gradational contact (gc) from argillaceous 1135 tuffitic deposits below to well-lithified grey limestone beds above with a transitional 1136 concretionary layer in between. Close-up view in F. Scale: 20 cm. Location: Pointe Lézarde. 1137 1138 Table 1. Summary of facies in middle Eocene limestone units of St. Bartholomew island. 1139 Texture : 'Ms' correspond to mudstone ; 'Ws' to wackestone ; 'Ps' to packstone ; 'Gs' to 1140 grainstone; 'Fs' to floatsone; 'Rs' to rudstone; 'Bs' to boudnstone; and 'Bis' to bindstone. 1141 Taphonomy : 'l' correspond to low ; 'm' to moderate, 'h' to high degree of alteration. 1142 1143









firm substrate

sandy-shelly substrate

soft muddy substrate



H.: Haimea; E.: Eupatagus; An: Antillaster; Ag.: Agassizia; B.: Brissoides; S.: Schyzaster.















foralgal (larger foraminifer-coralline algae photozoan): LfCa1, 2 rhodalgal (rhodolith photozoan): LfRh1, 2, 3 chloralgal (dasycladacean-miliolid photozoan): Mf1, 2 - DaMi chlorozoan (coral photozoan): CoP1, 2 echinofor (echinoid-larger foraminifer-coralline algae photozoan): EcF1,2; Mf3 foramol (larger foraminifer-mollusk photozoan): Mf4







B. Recovery of carbonate factories



C. Temporarily arrested or slowed carbonate production







Table 1. Lithofacies in the Eocene limestones of St Bartholemew.								
Facies code	Facies after James (1997)	Biotic constituents and skeletal association	Texture	Main skeletal and non skeletal components	Subordinate skeletal and non skeletal components	Taphonomy	Depositional environnement	
Mixed facie	es							
MF1	Smaller foraminiferan – dasycladalean – coral photozoan	Foralgal 1 Chloralgal 2 Chlorozoan 3	Ws/Fs, Ps and rarely Gs	Smaller benthic foraminifera. Peloids and micritized grains (cortoids). Agglutinated conical foraminifera and laminar cruts of coralline red algae (<i>Lithoporella</i>).	Miliolids and dasyclad algae. Echinoid (<i>Eupatagus, Antillaster</i>), ostracods, gastropods, bivalves, braching corals and amphisteginids.	Fragmentation : l to m Abrasion : t to m Encrustation : l Bioerosion : m	Inner ramp : shallow back shoal, lagoonal	
MF2	Dasycladalean- coral photozoan	Chlorozoan 1 Chloralgal 2 Foralgal 3	Bs Matrix : Fs	Same as MF1 but dominated by branching corals. Gastropods, dasyclad algae and miliolids.	Amphisteginids and nummulitids. Echinoid spatangoids (<i>Antillaster</i>) and agglutinated conical formainifera.	Fragmentation : 1 to m Abrasion : 1 tp m Encrustation : m Bioerosion : m	Inner ramp : shallow back shoal, lagoonal	
MF3	Echnoid – Red algal photozoan	Echinoforalgal	Ws to Ps/Gs	Echinoids and non-geniculate, small, rhodoliths, coralline branches and laminar crusts Peloids and cortoids.	Large foraminifera (orthophragminids, rare nummulitids and amphisteginids). Bivalves and corals	Fragmentation : m to h Abrasion : m to h Encrustation : l to m Bioerosion : l to m	Distal middle ramp : episodic wave influence	
MF4	Larger foraminiferan – molluscan photozoan	Foramol	Ps and/or Gs – volcaniclastic Rs.	Larger nummulitids, amphisgenitds and agglunatinated conicals.	Cora land coraline algal, mollusks. Echinoids are found fragented (<i>Eupatagus, Meoma</i> and <i>Haimea</i>). Peloids, bryozoans.	Fragmentation : m to h Abrasion : m to h Encrustation : l Bioerosion : l	Open inner ramp to middle ramp : wave- dominated	
MF5	Bioclastic – peloidal <u>photozoan</u>	Chloralgal	Ps	Peloids and cortoids.	Benthic formaminifera, echinoid plates and spines, and corals. Thin-shelled mollusks.	Fragmentation : m to h Abrasion : m to h Encrustation : l Bioerosion : h	Inner ramp : shallow back shoal, lagoonal	
Carbonate-	dominated facies					Diocrosion . II		
DaMi	Dasycladalean algae – miliolid photozoan association	Foralgal 1 Chloralgal 2 Chlorozoan 3	Ws and Ps	Similar to MF1 and MF2. Smaller benthic foraminifera, associated with miliolids and peloids. Dasyclad algae and ostracods. Branching corals and articulated bivales.	Cortoids, bivalves, agglutinated conicals (<i>Heterodictyoconus</i>) and amphisteginids. Melobesioids.	Fragmentation : 1 Abrasion : 1 Encrustation : m to h Bioerosion : m to h	Inner ramp : shallow back shoal, lagoonal	
LfRh1	Large bentihc foraminifera – rhodolith photozoan	Rhodalgal 1 Foralgal 2	Ps, Rs/Gs and rare Bis	Amphisteginids. Mastophoroids (<i>Litophorella</i>) and sporolithecean (<i>Sporolithon</i>). Melobesioids (<i>Mesophyllum</i> , <i>Lithothamnion</i>). Peloids, oncoids and cortoids. Rhodoltihs mean diameter range in size from 1-10 cm. Echinoids (<i>Haimea</i> , <i>Antillaster</i>) and large thick-shelled ovsters.	Nummulitids (<i>Nummulites</i>) and lepidocyclinids (<i>Polylepidina</i>). Textulariids, mollusks (epifaunal bivalves), small benthic foraminifers (textulariids, miliolids), encrusting foraminifera (acervulinids), bryozoan, molluks, and corals.	Fragmentation : l to h Abrasion : l to h Encrustation : l to h Bioerosion : l to h	Open inner ramp to middle ramp : wave- dominated	
LfRh2	Large bentihc foraminifera – rhodolith photozoan	Rhodalgal 1 Foralgal 2	Ps, Rs.	Nummulitids (<i>Nummulites</i>) and lepidocyclinids (<i>Polylepidina</i>). Rhodoltihs mean diameter range in size from 1-10 cm.	Amphisteginids, textulariids, mollusks (epifaunal bivalves dominate), coral fragments and planktonic foraminifera. Rare to common peloids and cortoids.	Fragmentation : l to h Abrasion : l to h Encrustation : l to h Bioerosion : l to h	Middle ramp : episodic wave influence	
LfRh3	Large bentihc foraminifera – rhodolith photozoan	Rhodalgal	Gs	Small rhodoliths. Echinoids and peloids. Rhodoltihs mean diameter range in size from 1-5 cm.	Larger benthic foraminifera, mollusks, corals, dasyclad algae and planktonic foraminifera.	Fragmentation : l to h Abrasion : l to h Encrustation : l to h Bioerosion : l to h	Middle ramp : episodic wave influence	
LfCa1	Larger benthic foraminiferan – coralline algal photozoan	Foralgal	Ps, Gs	Amphisteginids (<i>Amphistegina</i>). Echinoid, Peloids, cortoids and smaller bentic foraminifera (rotaliids). Subspherical rhodoliths (up to 3 cm diameter). Melobesioids and mastophoroids (<i>Lithoporella</i>).	Nummulitids (<i>Nummulites</i>). Smaller bentic foraminifera (textulariids). Lepidocyclinids (<i>Polylepidinia</i>).	Fragmentation : l to m Abrasion : l to m Encrustation : m Bioerosion : l to m	Open inner ramp to middle ramp : wave- dominated	
LfCa2	Larger benthic foraminiferan – coralline algal	Foralgal	Ps, Gs	Nummulitids (<i>Nummulites</i>) and orthophragminids (<i>Discocyclina, Pseudophragmina</i>).	Lepidocyclinids (Polylepidinia).	Fragmentation : l to h Abrasion : l to h Encrustation : l to h	Middle ramp : episodic wave influence	

	photozoan					Bioerosion : 1 to h	
EcF1	Echnoid – non- skeletal	Echinofor Echinoforalgal	Ps, Gs	Echinoid spatangoids (<i>Haimea</i>) and cassiduloids (<i>Echinolampas</i>). Peloids and	Corals. Bryozoans. Mollusks. Larger benthic foraminifera (amphisteginids	Fragmentation : m to h Abrasion : m to h	Open inner ramp to middle ramp : wave-
	carbonate photozoan			cortoids.	and nummulitids), red algal debris and miliolids.	Encrustation : 1 to m Bioerosion : h	dominated
EcF2	Echnoid – smaleer benthic foraminiferan phtozoan	Echinoforalgal	Ps, Ws	Infaunal and epifaunald echinoids are abundant. Co-occurrence of <i>Haimea</i> , <i>Schizaster</i> , <i>Agassizia</i> , <i>Eupatagus</i> and <i>Echinolampas</i> . Small benthic foraminifera (rotaliids). Mollusks. Peloids.	Large benthic foraminifera. Red algal fragments. Small agglutinated foraminifera, miliolids and calcispheres. Cortoids and oncoids.	Fragmentation : m to h Abrasion : m to h Encrustation : l to m Bioerosion : l to m	Inner ramp : shallow back shoal, lagoonal
CoP1	Coral photozoan	Chlorozoan	Bs Matrix : Ws, Ps, Gs .	Branching and platy corals. Peloids.	Encrusting coralline algae, gastropods, and small benthic foraminifera.	Fragmentation : l to m Abrasion : l to m Encrustation : m Bioerosion : m	Inner ramp : shallow back barrier, lagoonal. Lenticular coral bioherms.
CoP2	Coral photozoan	Chlorozoan	Bs Matrix : Rs	Dendroid corals. Peloids, amphisteginids, nummulitids and orthophragminids, mollusks, coralline red algae.	Dasyclad algae and smaller benthic foraminifera.	Fragmentation : m Abrasion : m Encrustation : m Bioerosion : m to h	Open inner ramp to middle ramp : wave- dominated. Domal coral bioherms