Surface sediment samples from early age of seafloor exploration can provide a late 19th century baseline of the marine environment

Marina C. Rillo [1,2,*], **Michal Kucera** [3], **Thomas H. G. Ezard** [1] **and C. Giles Miller** [2,*]

[1] *Ocean and Earth Science, National Oceanography Centre Southampton, University of Southampton, Southampton, UK*
[2] *Department of Earth Sciences, Natural History Museum, London, UK*
[3] *MARUM Center for Marine Environmental Sciences, University of Bremen, Bremen, Germany*

Correspondence*:
Marina C. Rillo
Ocean and Earth Science, National Oceanography Centre Southampton, University of Southampton Waterfront Campus Southampton SO143ZH, UK
marina.rillo@evobio.eu

C. Giles Miller
Department of Earth Sciences, Natural History Museum, Cromwell Road, London SW7 5BD, UK
g.miller@nhm.ac.uk

2 ABSTRACT

Ocean-floor sediment samples collected up to 150 years ago represent an important historical
archive to benchmark global changes in the seafloor environment, such as species' range shifts
and invasions and pollution trends. Such benchmarking requires that the historical sediment
samples represent the state of the environment at − or shortly before the time of collection.
However, early oceanographic expeditions sampled the ocean floor using devices like the
sounding tube or a dredge, which potentially disturb the sediment surface and recover a
mix of Holocene (surface) and deeper, Pleistocene sediments. Here we use climate-sensitive
microfossils as a fast biometric method to assess if historical seafloor samples contain a mixture
of modern and glacial sediments. Our assessment is based on comparing the composition of
planktonic foraminifera (PF) assemblages in historical samples with Holocene and Last Glacial
Maximum (LGM) global reference datasets. We show that eight out of the nine historical samples
contain PF assemblages more similar to the Holocene than to the LGM PF assemblages, but the
comparisons are only significant when there is a high local species' temporal turnover (from the
LGM to the Holocene). When analysing temporal turnover globally, we show that upwelling and
temperate regions had greatest species turnover, which are areas where our methodology would
be most diagnostic. Our results suggest that sediment samples from historical collections can
provide a baseline of the state of marine ecosystems in the late 19th century, and thus be used
to assess ocean global change trends.

1 INTRODUCTION

Late nineteenth and early twentieth century oceanographic expeditions set out to explore the vast and then widely unknown deep ocean. The voyage of the HMS *Challenger* is a notable example. As she sailed around the globe between 1872−76, researchers mapped for the first time the shape of the ocean basins and described over 4,500 new species of marine life (Manten, 1972). These early expeditions have important historical significance, as they mark the beginning of modern oceanography and stimulated further ocean exploration (Wüst, 1964).

From a scientific perspective, the observations and material acquired by these historical expeditions have great potential for global change research (Lister and Group, 2011; Johnson et al., 2011), as they provide a pre-1900 baseline of the marine environment (e.g., Roemmich et al. 2012; Gleckler et al. 2016). Yet historical seafloor sediment samples remain largely underutilised, because early seafloor sampling techniques involved collecting surficial sediments with instruments like the sounding tube, dredge or even the anchor (Thomson and Murray, 1891). All these instruments can penetrate below the surface and disturb the top layer of the sediment. As a result, such historical sediment samples might contain surface (Holocene) sediments mixed with deeper, glacial material (Hayward and Kawagata, 2005), potentially hindering their use as a historical baseline of the modern marine environment. Coring techniques provide more accurate sediment chronology (e.g., Röhl et al. 2000); however, historical samples represent the seafloor environment as much as 50 years earlier than the earliest core samples collected (Wüst, 1964), and thus contain sediments without any objects deposited after 1900. These uncontaminated historical samples can be useful for chemical analyses of the seafloor (e.g., pollution trends; Dekov et al. 2010), single-specimen analysis (e.g., Reichart et al. 2003; Wit et al. 2010) and investigations of species range shifts and invasions in the past century (e.g., Hoeksema et al. 2011). Therefore, it is important to assess the degree to which historical sediment samples represent Holocene or mixed-Pleistocene sediments.

One way to assess the degree of glacial mixing in the historical material would be to determine its absolute age using the radiocarbon dating technique, or to use glacial material proxies (e.g., Mg/Ca, oxygen isotopes). However, in cases where the extent of mixing is small, the exponentially decaying nature of the radiocarbon analysis can cause an ambiguous dating, and the isotopic analysis would require a large enough number of specimens to correctly represent the extent of the glacial mixing. Here we propose a complimentary method that uses planktonic foraminifera assemblage composition as a climate-sensitive fingerprint of the sediment age. Planktonic foraminifera (PF) are single-celled zooplankton that produce calcium carbonate shells and, upon death, accumulate in great numbers on the ocean floor (Hemleben et al., 1989). PF assemblage composition is sensitive to sea-surface temperature (Morey et al., 2005; Fenton et al., 2016) and its change between glacial and interglacial times has been used to determine the magnitude of glacial ocean cooling (MARGO Project Members, 2009).

In this brief report, we make use of the temperature sensitivity of PF and compare the composition of their assemblages in nine historical (> 100 years old) samples against reference PF assemblages from the Holocene (Siccha and Kucera, 2017) and the Last Glacial Maximum (Kucera et al., 2005a). We test whether it is possible to recover the extent of glacial mixing in historical seafloor sediment samples using PF assemblage composition. This new biometric method contributes to a more multidisciplinary approach to dating historical sediments.

2 MATERIAL AND METHODS

2.1 Historical samples

Historical samples were retrieved from the Ocean-Bottom Deposits (OBD) Collection held by The Natural History Museum in London. The OBD Collection holds about 40,000 historical samples from all the world's oceans (https://doi.org/10.5519/0096416), including most of the sediment samples collected by the HMS *Challenger* and the British Royal Navy survey ships (Kempe and Buckley, 1987). The OBD samples are kept sealed in their original glass jars and tubes and are usually dry as the result of the long (over 100 years) storage. We selected nine samples collected between 1874 and 1905, chosen to cover different oceans, latitudes and historical marine expeditions (Table 1). Half of the amount available in the OBD containers was further split into two equal parts, leaving an archive sample and a sample to be processed. The sample processing consisted of weighing, wet washing over a 63μm sieve and drying in a $60°$C oven. The residues were further dry sieved over a 150μm sieve and the coarser fraction was split with a microsplitter as many times as needed to produce a representative aliquot containing around 300 PF shells (see Al-Sabouni et al. 2007). All PF specimens in each of the nine final splits were picked, glued to a micropaleontology slide and identified under a stereomicroscope to species level, resulting in a total of 2,611 individuals belonging to 31 species (Table 1, Table S1).

2.2 Holocene and Last Glacial Maximum data

We tested whether the composition of PF assemblages in the historical samples is more similar to assemblages of the Holocene (last 11,700 years, Walker et al. 2008) or the Last Glacial Maximum (LGM, 21,000 years ago, MARGO Project Members 2009). The Holocene census dataset (i.e., marine surface sediment samples taken by coring methods after 1945) was recently curated and published as the ForCenS dataset, comprising 4,205 assemblage counts from unique sites (Siccha and Kucera, 2017). Three LGM datasets from the MARGO project (Kucera et al., 2005a,b,c; Barrows and Juggins, 2004) were merged following the taxonomic standardisation of Siccha and Kucera (2017). This merged LGM final dataset included 1165 counts from 389 unique sites. Moreover, local estimates of open-ocean sedimentation rates are available for 156 samples in the LGM datasets, and were used to analyse the results.

The assemblage compositions of the nine historical samples were then compared to the samples from the geographically nearest site in the Holocene and LGM datasets (Fig. S1A). The distances between sites were calculated using the World Geodetic System of 1984 (WGS84, Hijmans 2015). We then compiled annual mean and standard deviation values of sea surface temperature (SST) from the World Ocean Atlas 2013 (WOA13, 0 meters depth, Locarnini et al. 2013) for each of the 27 sites (historical, Holocene and LGM) to evaluate whether the neighbouring sites are at similar SST ranges. Most neighbouring sites had similar SST values except the two most southern samples (Fig. S1B). The exceptional Holocene nearest sample neighbouring M.8780 was substituted by the fourth nearest neighbour, 66 km farther but more similar in SST (Fig. S1B). The mean distance between our nine sites and their nearest neighbour set was 253 km in the Holocene data and 415 km in the LGM data.

2.3 Compositional similarity

Assemblage similarity was expressed using the Morisita-Horn index (Morisita, 1959; Horn, 1966), which is an abundance-based overlap measure that preserves essential properties of similarity measurements (Jost et al., 2011). The Morisita-Horn calculates the compositional similarity by pairwise comparison of the relative abundance of each species, and is robust to under-sampling (i.e., rare species occurrence) (Jost et al., 2011). The index was calculated using the under-sampling bias correction (Chao et al., 2006),

bootstrap confidence intervals based on 100 replicates and the R package *SpadeR* (version 0.1.1, Chao et al.
2016).

For each of the nine historical assemblages, we calculated the Morisita-Horn index three times: between
(1) historical and neighbouring Holocene assemblages, (2) historical and neighbouring LGM assemblages
and (3) neighbouring Holocene and LGM assemblages. This third comparison gives us a baseline index
value of how much the PF assemblage composition changed locally since the LGM. If historical samples
are representative of surface sediments, the similarity index calculated between historical and Holocene
samples should be higher than between historical and LGM samples, with non-overlapping confidence
intervals. However, if historical samples are a mixture of Holocene and LGM material, confidence intervals
of the historical-Holocene and historical-LGM comparisons overlap. Confidence intervals might also
overlap if the baseline index value calculated between the Holocene and the LGM reference samples is
high. A high baseline value means that the local Holocene and LGM assemblages are similar and thus
there is less statistical potential to detect whether a historical sediment is a surface or mixed-glacial sample.
To understand where our biometric methodology would be most diagnostic, we compiled a world map
of species turnover since the LGM, by calculating the Morisita-Horn index for each LGM sample and its
nearest Holocene neighbour. Compositional similarity indexes were averaged per site. Distances between
the Holocene sites and the 389 LGM sites were calculated using the WGS84, and averaged 52 km (median
1.5 km).

3   RESULTS AND DISCUSSION

In eight out of the nine samples, the compositional similarity was higher between the historical and
Holocene assemblages than between historical and LGM assemblages (Fig. 1A). However, non-overlapping
confidence intervals were only present in two samples (M.25 and M.5246), inferring a Holocene age
for these historical sediments. These two samples also showed the lowest similarities in assemblage
composition between neighbouring LGM and Holocene assemblages (Fig. 1A, grey dots), meaning that
there was a greater change in PF assemblage composition since the LGM in these two locations. The main
differences in species compositions were the high relative abundance of *Globigerina bulloides* in the LGM
neighbouring sample of M.25, and of *Globoconella inflata* in the LGM neighbouring sample of M.5246
(Tables S3,S5).

Compositional similarity index between historical and Holocene samples was always above 0.75, reaching
maximum similarity in five samples (Fig. 1A). In six samples, the confidence intervals of the three
comparisons overlapped, and all the similarity indexes were above 0.75. Since LGM and Holocene
PF assemblages showed higher similarity at these six sites, our biometric test is less diagnostic. The
historical sample M.8780 showed no overlap between historical-Holocene and Holocene-LGM, but the
historical-LGM comparison overlapped with both comparisons, suggesting that either this sample has a
mix of Holocene and glacial material, or the SST differences among these neighbouring samples prevents
appropriate compositional comparisons (Fig. S1B). M.8780 had more *Neogloboquadrina pachyderma*
than both the Holocene and LGM neighbours, and had *G. bulloides* abundances above 20%, similar to the
LGM neighbour (Table S3). Moreover, differences in SST among the neighbouring samples of M.192 (Fig.
S1B) did not seem to influence their compositional similarities (Fig. 1A). Finally, the historical sample
M.7487 was the only one that showed higher similarity with LGM than Holocene assemblages, suggesting
sediment mixing. The higher relative abundances of *Trilobatus sacculifer*, *N. incompta* and *Globigerinita*
glutinata were responsible for this pattern (Table S5).

The global comparison between the Holocene and the LGM reference datasets shows the magnitude
of PF assemblages turnover since the LGM (i.e., temporal beta-diversity, Fig. 1B). In general, upwelling
(eastern boundary currents and equatorial regions) and temperate sites had greatest species turnover. Our
methodology would be most diagnostic in these settings. Open-ocean sedimentation rates available for
156 sites averaged 6.8 centimetres per thousand of years (cm/ky). Therefore, historical sampling devices
would have had to penetrate on average 142.8 cm (6.8 cm/ky times 21 ky) into the sediment to contaminate
the surface seafloor sample with glacial material. Nevertheless, comparing sedimentation rates to the
compositional similarity between LGM samples and their Holocene neighbours reveals that the greatest
temporal turnover in species composition happened at sites of lower sedimentation rates (Fig. S2), where
sediment mixing during historical sampling would be most likely. Furthermore, six LGM neighbours of our
nine historical samples had local estimates of sedimentation rate, which varied from 1.0 to 3.4 cm/ky (mean
1.8 cm/ky, Tables S3,S4,S5). Thus, considering our historical samples only, depths of $21-71.4$ cm into
the sediment would have already reached glacial age material. Historical ocean-floor sampling methods
potentially disturbed the surface and led to recovery of a mix of Holocene and deeper sediments, especially
at sites of lower sedimentation rates. Our biometric method would be most useful at these sites with similar
sedimentation settings, which also show the greatest temporal turnover in species composition (Fig. S2).

4   CONCLUSION

Our results indicate that historical ocean-floor sediment samples (collected more than 100 years ago) can
represent surface (Holocene) sediments, despite the use of technology not designed to preserve undisturbed
sediments. We show that the temporal turnover in species occurrence since the LGM varies in space making
our biometric method particularly suitable for upwelling and temperate areas with low sedimentation rates.
The new method allows a non-destructive preliminary assessment of glacial contamination of historical
samples. Independent proxy records (e.g., Mg/Ca) and/or radiocarbon dating would be valuable to validate
the success of our technique. Ideally, the biometric approach would compliment chemical-based techniques
to date historical sediments, aiming for more robust results as multiple alternative lines of evidence are
presented. Our results highlight the scientific potential of historical seafloor sediment collections (Table
S2). As human activities increasingly modify the marine environment, these historical collections contain
important information on the pre-1900 state of marine ecosystems.

CONFLICT OF INTEREST STATEMENT

169 The authors declare that the research was conducted in the absence of any commercial or financial
170 relationships that could be construed as a potential conflict of interest.

AUTHOR CONTRIBUTIONS

171 All authors designed the research question. MR processed the historical sediments, with input from GM.
172 MR and MK identified all specimens. MR designed and performed the analysis, with input from MK and
173 TE. MR wrote the initial draft, and all authors reviewed and edited the final manuscript.

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DATA AVAILABILITY STATEMENT

181 Upon acceptance of the manuscript, the datasets used and generated for this study will be available online
182 at the NHM Data Portal (with DOI number), as well as the R Code used for the analysis and figures.

TABLE

Table 1. Information about the historical sediment samples from the Ocean Bottom Deposits (OBD) Collection at The Natural History Museum, London (NHMUK). Columns: NHMUK sample number; vessel that collected the sample (HMS stands for Her/His Majesty's Ship); date that sample was collected; method used for sampling; ocean where sample was collected; latitude and longitude in decimal degrees; water depth in meters (transformed from fathoms); sampled mass before processing, in grams; number of splits to sample around 300 planktonic foraminifera (PF) individuals; number of PF individuals identified; number of PF species identified.

Sample	Vessel HMS	Date	Method	Ocean	Lat	Long	Depth	m(g)	N(spl)	N(ind)	N(sp)
M.25	*Challenger*	21/02/1873	Sounding	Atlantic	24.33	-24.47	5011	2.73	5	260	18
M.192	*Challenger*	07/03/1874	Sounding	Indian	-50.02	123.07	3976	0.19	5	318	7
M.284	*Challenger*	11/03/1875	Sounding	Pacific	-0.70	147.00	2012	1.98	7	331	16
M.408	*Challenger*	21/03/1876	Dredge	Atlantic	-21.25	-14.03	3639	9.35	7	265	21
M.3787	*Egeria*	31/10/1887	Sounding	Indian	-19.57	64.63	2869	1.23	5	376	24
M.4080	*Egeria*	26/08/1889	Sounding	Pacific	-15.65	-179.06	2579	2.42	8	300	13
M.5246	*Penguin*	16/04/1891	Sounding	Indian	-26.94	111.18	3676	1.49	5	279	18
M.8780	*Waterwitch*	22/01/1895	Sounding	Indian	-40.45	49.82	3780	1.51	6	177	11
M.7487	*Sealark*	09/11/1905	Sounding	Indian	-7.59	61.48	3758	2.86	8	305	18
Total										2,611	31

FIGURE

Figure 1. (A) Compositional similarity (Morisita-Horn index) between planktonic foraminiferal assemblages from historical sediment samples and assemblages from surface sediments (Holocene, brown triangle) and from the Last Glacial Maximum (LGM, green squares); and between Holocene and LGM assemblages (grey dots, baseline value of local temporal turnover). 0 means that the two assemblages share no species; 1 means that the same species were present in both samples at statistically indistinguishable proportions. Lines represent confidence intervals based on 100 bootstrap replicates. The x-axis shows the historical sample number. **(B)** Black triangles: historical samples (nine in total). Coloured dots: LGM samples from 389 sites worldwide. The colours represent the Morisita-Horn similarity index between the LGM sample and its neighbouring Holocene sample (i.e., temporal turnover). Red to orange dots indicate low similarity (i.e., high species turnover), whereas blue dots indicate similar Holocene and LGM planktonic foraminifera assemblages.

REFERENCES

Al-Sabouni, N., Kucera, M., and Schmidt, D. N. (2007). Vertical niche separation control of diversity and size disparity in planktonic foraminifera. *Marine Micropaleontology* 63, 75–90. doi:https://doi.org/10.1016/j.marmicro.2006.11.002

[Dataset] Barrows, T. T. and Juggins, S. (2004). Compilation of planktic foraminifera LGM assembages from the Indo-Pacific Ocean. doi:https://doi.org/10.1594/PANGAEA.227319

Chao, A., Chazdon, R. L., Colwell, R. K., and Shen, T.-J. (2006). Abundance-based similarity indices and their estimation when there are unseen species in samples. *Biometrics* 62, 361–371. doi:https://doi.org/10.1111/j.1541-0420.2005.00489.x

Chao, A., Ma, K. H., Hsieh, T. C., and Chiu, C.-H. (2016). *SpadeR: Species-Richness Prediction and Diversity Estimation with R*. R package version 0.1.1, https://CRAN.R-project.org/package=SpadeR

Dekov, V. M., Cuadros, J., Kamenov, G. D., Weiss, D., Arnold, T., Basak, C., et al. (2010). Metalliferous sediments from the H.M.S. *Challenger* voyage (1872–1876). *Geochimica et Cosmochimica Acta* 74, 5019–5038. doi:https://doi.org/10.1016/j.gca.2010.06.001

Fenton, I. S., Pearson, P. N., Jones, T. D., and Purvis, A. (2016). Environmental predictors of diversity in Recent planktonic foraminifera as recorded in marine sediments. *PloS ONE* 11, e0165522. doi:https://doi.org/10.1371/journal.pone.0165522

Gleckler, P. J., Durack, P. J., Stouffer, R. J., Johnson, G. C., and Forest, C. E. (2016). Industrial-era global ocean heat uptake doubles in recent decades. *Nature Climate Change* 6, 394–398. doi:http://dx.doi.org/10.1038/nclimate2915

Hayward, B. W. and Kawagata, S. (2005). Extinct foraminifera figured in Brady's *Challenger* Report. *Journal of Micropalaeontology* 24, 171–175. doi:https://doi.org/10.1144/jm.24.2.171

Hemleben, C., Spindler, M., and Anderson, O. R. (1989). *Modern Planktonic Foraminifera* (Springer-Verlag New York Inc.)

Hijmans, R. J. (2015). *geosphere: Spherical Trigonometry*. R package version 1.5-7, https://CRAN.R-project.org/package=geosphere

Hoeksema, B. W., van der Land, J., van der Meij, S. E., van Ofwegen, L. P., Reijnen, B. T., van Soest, R. W., et al. (2011). Unforeseen importance of historical collections as baselines to determine biotic change of coral reefs: the Saba Bank case. *Marine Ecology* 32, 135–141. doi:https://doi.org/10.1111/j.1439-0485.2011.00434.x

Horn, H. S. (1966). Measurement of "overlap" in comparative ecological studies. *The American Naturalist* 100, 419–424. doi:https://www.jstor.org/stable/2459242

Johnson, K. G., Brooks, S. J., Fenberg, P. B., Glover, A. G., James, K. E., Lister, A. M., et al. (2011). Climate change and biosphere response: unlocking the collections vault. *BioScience* 61, 147–153. doi:https://doi.org/10.1525/bio.2011.61.2.10

Jost, L., Chao, A., and Chazdon, R. L. (2011). *Compositional similarity and β (beta) diversity (in Biological diversity: frontiers in measurement and assessment)* (Oxford University Press), chap. 6. 1 edn., 66–84

Keltie, J. S. (1898). The German Deep-Sea Expedition. *The Geographical Journal* 12, 494–496. doi:https://archive.org/stream/geographicaljou33britgoog#page/n528

Kempe, D. R. C. and Buckley, H. A. (1987). Fifty years of oceanography in the Department of Mineralogy, British Museum (Natural History). *Bulletin of the British Museum (Natural History)* 15, 59–97

Kucera, M., Rosell-Mele, A., Schneider, R., Waelbroeck, C., and Weinelt, M. (2005a). Multiproxy approach for the reconstruction of the glacial ocean surface (margo). *Quaternary Science Reviews* 24, 813–819. doi:https://doi.org/10.1016/j.quascirev.2004.07.017

[Dataset] Kucera, M., Weinelt, M., Kiefer, T., Pflaumann, U., Hayes, A., Weinelt, M., et al. (2005b). Compilation of planktic Foraminifera census data, LGM from the Atlantic Ocean. doi:https://doi.org/10.1594/PANGAEA.227329

[Dataset] Kucera, M., Weinelt, M., Kiefer, T., Pflaumann, U., Hayes, A., Weinelt, M., et al. (2005c). Compilation of planktic Foraminifera census data, LGM from the Pacific Ocean. doi:https://doi.org/10.1594/PANGAEA.227327

Lister, A. M. and Group, C. C. R. (2011). Natural history collections as sources of long-term datasets. *Trends in Ecology & Evolution* 26, 153–154. doi:https://doi.org/10.1016/j.tree.2010.12.009

Locarnini, R. A., Mishonov, A. V., Antonov, J. I., Boyer, T. P., Garcia, H. E., Baranova, O. K., et al. (2013). World Ocean Atlas 2013, Volume 1: Temperature. *NOAA Atlas NESDIS* 73, 1–40

Manten, A. A. (1972). C. Wyville Thomson, J. Murray, and the "Challenger" expedition. *Earth-Science Reviews* 8, 255–266. doi:https://doi.org/10.1016/0012-8252(72)90108-0

MARGO Project Members (2009). Constraints on the magnitude and patterns of ocean cooling at the Last Glacial Maximum. *Nature Geoscience* 2, 127–132. doi:http://dx.doi.org/10.1038/ngeo411

Morey, A. E., Mix, A. C., and Pisias, N. G. (2005). Planktonic foraminiferal assemblages preserved in surface sediments correspond to multiple environment variables. *Quaternary Science Reviews* 24, 925–950. doi:https://doi.org/10.1016/j.quascirev.2003.09.011

Morisita, M. (1959). Measuring of interspecific association and similarity between communities. *Memoirs of the Faculty of Science, Kyushu University, Series E (Biology)* 3, 65–80

Reichart, G.-J., Jorissen, F., Anschutz, P., and Mason, P. R. (2003). Single foraminiferal test chemistry records the marine environment. *Geology* 31, 355–358

Roemmich, D., Gould, W. J., and Gilson, J. (2012). 135 years of global ocean warming between the *Challenger* expedition and the Argo Programme. *Nature Climate Change* 2, 425. doi:http://dx.doi.org/10.1038/nclimate1461

Röhl, U., Bralower, T., Norris, R., and Wefer, G. (2000). New chronology for the late Paleocene thermal maximum and its environmental implications. *Geology* 28, 927

Sandeman, G. (1900). The German deep sea expedition on the ship "Valdivia". *Scottish Geographical Magazine* 16, 416–418. doi:https://archive.org/stream/scottishgeograp00unkngoog#page/n446

Siccha, M. and Kucera, M. (2017). ForCenS, a curated database of planktonic foraminifera census counts in marine surface sediment samples. *Scientific Data* 4, sdata2017109. doi:http://dx.doi.org/10.1038/sdata.2017.109

Thomson, C. W. and Murray, J. (1891). *Report on the Scientific Results of the Voyage of HMS Challenger during the years 1872−76. Deep-Sea Deposits.* (United Kingdom: HM Stationery Office). doi:http://epic.awi.de/38636/

Walker, M., Johnsen, S., Rasmussen, S. O., Steffensen, J.-P., Popp, T., Gibbard, P., et al. (2008). The Global Stratotype Section and Point (GSSP) for the base of the Holocene Series/Epoch (Quaternary System/Period) in the NGRIP ice core. *Episodes* 31, 264–267

Wit, J., Reichart, G.-J., A Jung, S., and Kroon, D. (2010). Approaches to unravel seasonality in sea surface temperatures using paired single-specimen foraminiferal $\delta18o$ and mg/ca analyses. *Paleoceanography and Paleoclimatology* 25

Wüst, G. (1964). The major deep-sea expeditions and research vessels 1873-1960: a contribution to the history of oceanography. *Progress in Oceanography* 2, 1–52. doi:https://doi.org/10.1016/0079-6611(64)90002-3

SUPPLEMENTAL DATA

Table S1. Planktonic foraminifera species counts of the nine historical samples (see Table 1). IRN (OBD) stands for the internal registration number of the Ocean-Bottom Deposits collection at the Natural History Museum in London. Individuals were identified to the species level by M. Rillo and M. Kucera together.

Historical samples									
Museum number	M.192	M.8780	M.5246	M.408	M.3787	M.4080	M.7487	M.284	M.25
IRN (OBD)	32657	38482	36053	34991	34671	34993	37148	33668	33286
Latitude	-50.02	-40.45	-26.94	-21.25	-19.57	-15.65	-7.59	-0.70	24.33
Longitude	123.07	49.82	111.18	-14.03	64.63	-179.06	61.48	147	-24.47
Species									
Beella digitata	0	0	0	0	2	1	0	0	0
Berggrenia pumilio	0	0	0	0	0	0	0	0	0
Candeina nitida	0	0	0	0	1	0	0	0	0
Globigerina bulloides	175	38	13	4	2	3	8	28	6
Globigerina falconensis	0	4	11	17	2	6	0	12	0
Globigerinella calida	1	1	6	8	11	15	0	12	3
Globigerinella siphonifera	0	0	5	4	29	10	16	14	2
Globigerinita glutinata	5	5	52	18	24	63	77	65	2
Globigerinoides conglobatus	0	0	1	3	12	8	6	2	2
Globigerinoides ruber pink	0	0	0	11	0	0	0	0	0
Globigerinoides ruber white	0	3	73	84	196	131	53	115	68
Globigerinoides tenellus	0	0	0	20	9	19	0	6	0
Globoconella inflata	96	58	10	31	2	0	0	0	35
Globoquadrina conglomerata	0	0	2	0	1	0	0	1	0
Globorotalia crassaformis	0	0	2	2	1	0	0	0	5
Globorotalia hirsuta	0	0	0	1	0	0	0	0	0
Globorotalia menardii	0	0	37	0	1	0	30	1	23
Globorotalia scitula	0	0	0	2	1	0	5	0	0
Globorotalia theyeri	0	0	0	0	0	0	1	0	0
Globorotalia truncatulinoides	14	10	6	12	3	0	1	0	10
Globorotalia tumida	0	0	0	2	0	0	15	2	7
Globorotalia ungulata	0	0	3	2	2	0	4	0	0
Globorotaloides hexagonus	0	0	0	0	1	0	0	0	0
Globoturborotalita rubescens	0	0	0	3	2	4	4	4	0
Neogloboquadrina dutertrei	0	0	34	1	11	4	21	9	7
Neogloboquadrina incompta	8	17	4	1	0	0	33	10	33
Neogloboquadrina pachyderma	19	38	0	0	0	0	0	0	0
Orbulina universa	0	1	1	6	3	4	0	0	1
Pulleniatina obliquiloculata	0	0	2	3	7	0	16	22	5
Sphaeroidinella dehiscens	0	0	0	0	1	0	1	0	2
Trilobatus sacculifer	0	0	17	30	52	32	13	28	46
Turborotalita humilis	0	0	0	0	0	0	1	0	3
Turborotalita quinqueloba	0	2	0	0	0	0	0	0	0
Total count	318	177	279	265	376	300	305	331	260

Figure S1. (A) Map of historical sample sites (red triangles) and corresponding neighbouring sites of the Holocene (green squares, ForCenS database) and Last Glacial Maximum (blue dots, LGM, MARGO datasets). Distances between each historical site and its corresponding Holocene and LGM neighbours can be seen in Tables S3,S4,S5. (B) Mean annual sea surface temperature (SST, in degree Celsius) and annual standard deviations (bars) at each sample site (WOA13 data). Note that historical sample M.8780 has two values for the Holocene neighbour. The nearest Holocene neighbour (SST 7.8°C, distance of 374 km) was substituted by the fourth nearest neighbour, which had more similar SST values (17.8°C, distance of 440 km, see map above).

Figure S2. Compositional similarity (Morisita-Horn index) between planktonic foraminifera assemblages in the LGM sample and its neighbouring Holocene sample, plotted against local estimates of open-ocean sedimentation rates (centimetres per thousand of years). Low similarity means a greater temporal turnover in local assemblage composition. In total there are 150 sedimentation rate estimates; six samples with sedimentation rate higher than 20 cm/ky and compositional similarity above 0.75 are not shown.

Table S2. Historical surface sediments collections

Museum	Expedition	Spatial coverage	Period
Natural History Museum in London	*Challenger*[1]	Atlantic, Antarctic, South Indian, Pacific	1872−76
Muséum National d'histoire Naturelle in Paris	*Travailleur, Talisman*[2]	Atlantic, Mediterranean	1880−83
Museum für Naturkunde in Berlin	German Deep-Sea[2] (*Valdivia*)	Atlantic, Antarctic, Indian, Mediterranean	1898−99

[1] https://www.hmschallenger.net/
[2] https://gallica.bnf.fr/ark:/12148/bpt6k65190274
[3] Keltie 1898; Sandeman 1900

Table S3. Relative abundance of planktonic foraminifera species in the historical samples M.192, M.8780 and M.5246, and their corresponding neighbouring samples in the Holocene and Last Glacial Maximum reference datasets. Species not shown had 0% abundance in all samples. Local sedimentation rate is given by the LGM datasets (see Methods) and expressed in centimetres per thousand of years.

Sample	Historical	Holocene	LGM	Historical	Holocene	LGM	Historical	Holocene	LGM
Sample/core number	M.192	CLIMAP_0078	E45-078	M.8780	BUFD_1027	RC14-009	M.5246	MARGO_2820	RC11-145
Latitude	-50.02	-47.77	-45.04	-40.45	-36.50	-39.02	-26.94	-26.97	-25.48
Longitude	123.07	123.10	114.35	49.82	49.57	47.88	111.18	111.33	110.02
Distance (km)	0	250.11	857.04	0	440.25	229.83	0	15.42	199.38
Sedimentation rate (cm/ky)			1.5			3.4			1.4
Species									
Beella digitata	0	0	0	0	0.48	0	0	0	0
Globigerina bulloides	55.03	60.71	51.67	21.47	7.88	25.58	4.66	8.85	1.81
Globigerina falconensis	0	0	0	2.26	15.75	0.66	3.94	0.66	3.32
Globigerinella calida	0.31	0	0	0.56	4.06	0	2.15	0	0.30
Globigerinella siphonifera	0	0	0	0	3.34	0.33	1.79	5.53	1.21
Globigerinita glutinata	1.57	1.53	3.67	2.82	3.58	1.99	18.64	19.25	18.73
Globigerinoides conglobatus	0	0	0	0	0	0	0.36	0.33	0.60
Globigerinoides ruber white	0	0	0	1.69	7.64	1.00	26.16	28.10	16.92
Globigerinoides tenellus	0	0	0	0	0	0	0	5.20	0.30
Globoconella inflata	30.19	20.66	17.33	32.77	35.56	11.63	3.58	6.19	47.73
Globoquadrina conglomerata	0	0	0	0	0	0	0.72	0	0
Globorotalia crassaformis	0	0	0.33	0	0	0	0.72	0	0.60
Globorotalia hirsuta	0	0	0	0	1.43	0.66	0	1.00	0
Globorotalia menardii	0	0	0	0	0	0	13.26	9.85	0.30
Globorotalia scitula	0	0	0.33	0	0.48	0	0	0	0
Globorotalia truncatulinoides	4.40	6.12	1.33	5.65	2.86	1.66	2.15	1.66	1.51
Globorotalia ungulata	0	0	0	0	0	0	1.08	0	0
Globoturborotalita rubescens	0	0	0	0	0	0.33	0	0.33	0
Neogloboquadrina dutertrei	0	0	0	0	1.19	1.00	12.19	3.87	3.93
Neogloboquadrina incompta	2.52	3.32	4.67	9.60	10.02	40.53	1.43	1.00	0.91
Neogloboquadrina pachyderma	5.97	6.63	14.33	21.47	0.24	5.65	0	0	0
Orbulina universa	0	0.51	0.33	0.56	0.72	0.33	0.36	0.66	0.60
Pulleniatina obliquiloculata	0	0	0	0	0	0	0.72	1.66	0
Trilobatus sacculifer	0	0	0	0	0.48	0	6.09	5.86	0
Turborotalita humilis	0	0	0.33	0	0	0	0	0	0
Turborotalita quinqueloba	0	0	5.67	1.13	0.24	8.64	0	0	0

Table S4. Relative abundance of planktonic foraminifera species in the historical samples M.408, M.3787 and M.4080, and their corresponding neighbouring samples in the Holocene and Last Glacial Maximum reference datasets. Species not shown had 0% abundance in all samples. Local sedimentation rate is given by the LGM datasets (see Methods) and expressed in centimetres per thousand of years.

Sample	Historical	Holocene	LGM	Historical	Holocene	LGM	Historical	Holocene	LGM
Sample/core number	M.408	BUFD_0336	GeoB1417-1	M.3787	CLIMAP_0211	V20-175	M.4080	MARGO_1184	RC13-038
Latitude	-21.25	-24.07	-20.37	-19.57	-20.58	-22.30	-15.65	-15.15	-14.52
Longitude	-14.03	-15.12	-12.71	64.63	63.53	68.00	-179.06	-176.49	177.10
Distance (km)	0	332.71	169.48	0	161.12	463.76	0	281.64	431.21
Sedimentation rate (cm/ky)			NA			1.5			1.9
Species									
Beella digitata	0	2.08	0.38	0.53	0	1.82	0.33	0.10	0.61
Candeina nitida	0	0	0.19	0.27	0.27	0.36	0	0	0
Dentigloborotalia anfracta	0	0	0.32	0	0	0	0	0	0
Globigerina bulloides	1.51	1.33	17.75	0.53	2.73	2.55	1.00	0.50	2.66
Globigerina falconensis	6.42	3.60	2.02	0.53	2.19	0.73	2.00	0.70	1.43
Globigerinella adamsi	0	0	0	0	0	0	0	0.10	0
Globigerinella calida	3.02	2.84	0.78	2.93	1.91	5.84	5.00	4.73	3.89
Globigerinella siphonifera	1.51	5.30	2.48	7.71	2.46	1.82	3.33	4.53	4.09
Globigerinita glutinata	6.79	12.88	8.62	6.38	9.29	18.25	21.00	19.32	31.49
Globigerinoides conglobatus	1.13	0.76	0.89	3.19	0.55	1.82	2.67	5.03	2.45
Globigerinoides ruber pink	4.15	1.33	0.81	0	0	0	0	0	0
Globigerinoides ruber white	31.70	40.15	24.42	52.13	56.56	44.16	43.67	38.23	40.90
Globigerinoides tenellus	7.55	4.55	2.50	2.39	0.55	4.38	6.33	4.02	2.86
Globoconella inflata	11.70	2.08	10.20	0.53	0.82	0.36	0	0	0
Globoquadrina conglomerata	0	0	0	0.27	0	0	0	0	0
Globorotalia crassaformis	0.75	0.38	1.82	0.27	0	0.36	0	0	0.41
Globorotalia hirsuta	0.38	0.57	0	0	0	0.73	0	0	0
Globorotalia menardii	0	2.65	0.04	0.27	0.82	0.73	0	0	0.20
Globorotalia scitula	0.75	1.70	1.81	0.27	1.37	1.09	0	0	0.20
Globorotalia truncatulinoides	4.53	5.49	6.34	0.80	1.37	6.57	0	0	0
Globorotalia tumida	0.75	0.38	0.02	0	0.82	0	0	0.20	1.02
Globorotalia ungulata	0.75	0	0	0.53	0	0	0	0	0
Globorotaloides hexagonus	0	0	0	0.27	0	0	0	0.30	0
Globoturborotalita rubescens	1.13	2.08	4.73	0.53	1.64	3.65	1.33	3.82	1.23
Neogloboquadrina dutertrei	0.38	0.19	0.89	2.93	2.73	1.82	1.33	0.10	2.25
Neogloboquadrina incompta	0.38	0	3.97	0	0.82	0	0	0	0.41
Neogloboquadrina pachyderma	0	0	0	0	0.27	0	0	0	0
Orbulina universa	2.26	3.22	0.96	0.80	2.73	1.46	1.33	2.41	2.66
Pulleniatina obliquiloculata	1.13	0	0	1.86	0	0.36	0	0	0.61
Sphaeroidinella dehiscens	0	0	0	0.27	0.27	0.36	0	0	0
Tenuitella iota	0	0	0.17	0	0	0	0	0	0
Trilobatus sacculifer	11.32	5.30	7.33	13.83	9.56	0	10.67	11.37	0
Turborotalita humilis	0	0	0.35	0	0	0.36	0	0.20	0
Turborotalita quinqueloba	0	0	0.16	0	0	0.36	0	4.33	0.61

Table S5. Relative abundance of planktonic foraminifera species in the historical samples M.7487, M.284 and M.25, and their corresponding neighbouring samples in the Holocene and Last Glacial Maximum reference datasets. Species not shown had 0% abundance in all samples. Local sedimentation rate is given by the LGM datasets (see Methods) and expressed in centimetres per thousand of years.

Sample	Historical	Holocene	LGM	Historical.1	Holocene.1	LGM.1	Historical.2	Holocene.2	LGM.2
Sample/core number	M.7487	BUFD_1236	V29-048	M.284	BUFD_0402	KH92-1-5a	M.25	ATL947_0944	V23-100
Latitude	-7.59	-6.27	-6.27	-0.70	1.27	3.53	24.33	22.70	21.30
Longitude	61.48	63.43	63.43	147.00	149.20	141.86	-24.47	-25.33	-22.68
Distance (km)	0	261.13	260.62	0	328.48	740.79	0	202.35	384.21
Sedimentation rate (cm/ky)			1.0			NA			NA
Species									
Beella digitata	0	0.26	1.11	0	0.11	0	0	0.50	0
Candeina nitida	0	0	0	0	0.22	0.50	0	0	0
Globigerina bulloides	2.62	5.03	3.33	8.46	4.20	5.50	2.31	2.30	9.64
Globigerina falconensis	0	0	0	3.63	0.82	1.50	0	0	1.79
Globigerinella calida	0	1.32	3.33	3.63	2.78	0	1.15	0.20	0.30
Globigerinella siphonifera	5.25	4.76	5.56	4.23	5.94	18.50	0.77	2.90	0.30
Globigerinita glutinata	25.25	6.61	9.63	19.64	17.18	25.50	0.77	2.90	3.78
Globigerinoides conglobatus	1.97	3.44	4.07	0.60	1.42	0.50	0.77	1.80	0
Globigerinoides ruber	0	0	0	0	0	0	0	0.50	0
Globigerinoides white	17.38	24.07	22.22	34.74	28.57	31.00	26.15	15.18	7.36
Globigerinoides tenellus	0	0.26	0.74	1.81	1.42	1.50	0	0.20	0.50
Globoconella inflata	0	0	0	0	0	0	13.46	16.28	0.30
Globoquadrina conglomerata	0	4.76	0.37	0.30	0.87	0	0	0	0
Globorotalia crassaformis	0	0	0.74	0	0.44	0	1.92	1.80	46.42
Globorotalia hirsuta	0	0	0	0	0.11	0	0	0	0
Globorotalia menardii	9.84	14.02	14.44	0.30	2.24	1.75	8.85	13.59	0
Globorotalia scitula	1.64	0.79	1.48	0	0.22	0	0	0.20	0
Globorotalia theyeri	0.33	0	0	0	0	0	0	0	0
Globorotalia truncatulinoides	0.33	0	0	0	0.49	0	3.85	7.89	1.49
Globorotalia tumida	4.92	1.85	1.48	0.60	1.58	1.75	2.69	4.30	0
Globorotalia ungulata	1.31	0	0	0	0	0	0	0	0
Globorotaloides hexagonus	0	3.44	2.22	0	0.11	0	0	0	0
Globoturborotalita rubescens	1.31	0	0.74	1.21	2.29	0.50	0	0	0
Neogloboquadrina dutertrei	6.89	7.94	17.41	2.72	4.74	3.50	2.69	5.69	1.99
Neogloboquadrina incompta	10.82	0.26	4.44	3.02	0.27	0	12.69	4.80	0
Neogloboquadrina pachyderma	0	0	0	0	0.05	0	0	0	5.07
Orbulina universa	0	0.26	1.11	0	0.11	0	0.38	0.20	3.08
Pulleniatina obliquiloculata	5.25	3.17	5.19	6.65	12.21	4.50	1.92	0.70	13.72
Sphaeroidinella dehiscens	0.33	0.79	0.37	0	0.22	0	0.77	0.90	0
Trilobatus sacculifer	4.26	15.87	0	8.46	10.69	3.50	17.69	17.18	3.28
Turborotalita humilis	0.33	0	0	0	0	0	1.15	0	0.99
Turborotalita quinqueloba	0	0	0	0	0	0	0	0	0.30