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The longest-lived Pacific hotspots reveal a plume tail for the largest oceanic plateau

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20 Volcanic hotspots are thought to initially form by melting in an upwelling mantle plume head
21 followed by melting of the plume tail. Plate motion then generates an age progressive volcanic
22 track originating from a large igneous province that connects to an active hotspot. However,
23 the most voluminous large igneous province, the ~120 Ma Ontong-Java Nui Plateau (OJP-
24 Nui) in the mid-Pacific, appears to lack such a volcanic track. Although the Louisville hotspot
25 track was originally proposed as a candidate, limited constraints for Pacific absolute plate
26 and plume motion prior to 80 Ma suggest a mismatch¹. Existing Pacific models rely on age-
27 distance data from the continuous Hawaii-Emperor and Louisville volcanic tracks, but their
28 seamounts older than ~80 Ma are now subducted, and elsewhere on the Pacific plate only
29 discontinuous and sparse seamount tracks can be found that formed prior to 80 Ma²⁻⁷. These
30 existing models require ~1,200 km of latitudinal motion for the Louisville plume to also erupt
31 the OJP-Nui¹, yet paleolatitude estimates from ~70 Ma to today remain within error of its
32 present location^{8,9} and suggest that any major amount of Louisville plume motion should
33 precede that time. Here we provide evidence from geochemistry and eruption ages⁹⁻¹⁴
34 demonstrating that Samoa and Rurutu-Arago are the longest-lived Pacific hotspots that can
35 be traced back to ~120 Ma (and older) in the West Pacific where they subduct into the
36 Mariana Trench. These newly defined tracks provide for an alternative Pacific absolute plate
37 motion model, with better constraints for a plate rotation between 80-100 Ma, and allow us
38 to establish Louisville as the missing volcanic track for OJP-Nui without requiring major
39 plume motion.

40 Plume-fed hotspots exhibit age-progressive volcanic tracks that often originate from a large
41 igneous province (LIP), which marks a hotspot's inception¹⁵⁻¹⁸. Eruption of a LIP is thought to
42 correspond to the arrival of the head of a deeply-rooted mantle plume, causing extensive melting
43 in the upper mantle and unusually voluminous eruption comprising the LIP^{3,19,20}. The Ontong-Java
44 Plateau (OJP; Fig. 1) once formed greater Ontong-Java Nui (OJP-Nui), together with the Manihiki
45 and Hikurangi plateaus²¹. Despite a proposed plume origin^{22,23} there is no obvious age-progressive
46 volcanic track emerging from the plateau. Although the Louisville hotspot was proposed to be
47 related²⁴, later paleomagnetic and geochemistry data argued against a Louisville hotspot
48 connection with the OJP²⁵⁻²⁷. Recently obtained Louisville paleomagnetic latitudes are constant
49 within error up to ~70 Ma⁸, suggesting that any Louisville plume motion should precede 70 Ma.
50 In that timeframe, existing absolute plate motion (APM) models require ~1,200 km of plume

51 motion to place Louisville and OJP in the same original eruptive location¹. However, more recent
52 geochemical data revealing similar Nd-Pb-Sr systematics between the main phase of the OJP-Nui
53 system and the Louisville Seamount track are permissive of the Louisville-OJP connection²⁸ (Fig.
54 2). Additionally, here we show that the two longest-lived Pacific hotspot tracks —Samoa and
55 Rurutu-Arago^{9,14} (see below)—yield a new APM model that supports a genetic link between the
56 Louisville hotspot track and OJP-Nui, the largest LIP preserved in the geologic record.

57 Most existing Pacific APM models rely significantly on the Hawaii-Emperor and
58 Louisville hotspots back to 80 Ma, while unrelated discontinuous volcanic structures are used for
59 Pacific plate motion prior to 80 Ma^{2,3,7,18,29,30} (Fig. 1, Extended Data Fig. 1). These structures
60 include the Shatsky Rise, Hess Rise, Mid-Pacific Mountains, Line Islands, Liliuokalani
61 Seamounts, Musician Seamounts, Wake Seamounts, Marshall Islands, and Magellan Seamounts.
62 From these, the Mid-Pacific Mountains and Shatsky Rise erupted near all-ridge triple junctions,
63 while Hess Rise and Musicians Seamounts erupted near-ridge^{29–32} (Extended Data Fig. 2). Plume-
64 ridge interactions could significantly displace the upwelling mantle plume³³ such that resulting
65 plume motion is inadvertently included in the APM models. Moreover, the Line Islands lack a
66 clear age progression³⁴, the Musician Seamounts are complicated by overprinting of deformation-
67 related volcanism³³, and Shatsky Rise was recently suggested to be controlled by seafloor
68 spreading³⁵. In summary, all of these structures are unlikely to exclusively represent absolute plate
69 motion, yet the Wake, Marshall, and Magellan seamounts appear to be the only truly intra-plate
70 Pacific hotspots prior to 80 Ma (Extended Data Fig. 2). As we will demonstrate, they represent the
71 Cretaceous portions of two long-lived Pacific hotspots—Samoa and Rurutu-Arago—that can
72 extend Hawaii-Emperor and Louisville anchored Pacific APM models to the 80 to 120 Ma period.

73 The clearly defined Hawaii-Emperor and Louisville hotspot tracks provide a continuous
74 hotspot record for APM modeling back to approximately 80 Ma, where they subduct into the
75 Kamchatka and Tonga trenches, respectively^{8,36} (Fig. 1; Extended Data Fig. 1). However, changing
76 paleomagnetic latitudes³⁷, predictions of hotspot motion based on Indo-Atlantic hotspots³⁸, inter-
77 hotspot differences in age progressions and distances^{9,39,40}, and disparity in the timing of the
78 Hawaii-Emperor bend compared to any known large global plate reorganization⁴¹, all suggest
79 relative motion between these two plumes and the Pacific plate. Supporting this contention, models
80 using global mantle flow to predict plume motions provide a better fit to the actual hotspot tracks

81 across ocean basins than fixed plume models⁶. The improved fit incorporating these geodynamic
82 models suggests that hotspots provide a combined record of plate motion and plume motion^{3,5,18}.

83 On the Pacific plate, the Rurutu-Arago volcanic track—defined from young to old by the
84 Cook-Austral Islands, Tuvalu Islands, the Marshall Islands and Wake Seamounts—represents the
85 third long-lasting hotspot in addition to Hawaii-Emperor and Louisville (Fig. 1). Changes in the
86 distance between these hotspots from 60 to 50 Ma suggest that the Hawaii-Emperor hotspot moved
87 more significantly and mostly independently from both Louisville and Rurutu-Arago⁹. However,
88 more critical here is the fact that the Rurutu-Arago and Samoan hotspots can be traced further back
89 into the Cretaceous than Hawaii and Louisville. The older portions of the Rurutu-Arago and Samoa
90 hotspots both extend into the West Pacific, where tracing their tracks through the high density of
91 Cretaceous seamounts has required the mapping out of well-defined age progressions combined
92 with unique geochemical signatures^{4,12,14} (Fig. 1, Extended Data Fig. 7, 8).

93 To this end, we present new isotope geochemistry from seamounts around Wake Island, a
94 critical geological nexus that links the older (>80 Ma) segment of the long-lived Rurutu-Arago
95 hotspot track to the younger (< 80 Ma) portion originating in the Cook-Austral Islands^{4,12,14,42,43}.
96 We use this hotspot track, together with the long-lived Samoan hotspot track, to generate a new
97 APM stage pole between 80-100 Ma that resolves the apparent disconnect between OJP-Nui and
98 the Louisville hotspot.

99 The extreme and distinct hotspot compositions originating in the central Pacific⁴² have
100 generated clearly traceable hotspot tracks (Fig. 1, Extended Data Fig. 8). We reveal these tracks
101 by color coding (Fig. 1) based on the ⁸⁷Sr/⁸⁶Sr-¹⁴³Nd/¹⁴⁴Nd-²⁰⁶Pb/²⁰⁴Pb isotopic compositions of
102 lavas at each volcano along the hotspot tracks^{9-14,44,45}. In particular, high ²⁰⁶Pb/²⁰⁴Pb compositions
103 are an identifying feature of the Rurutu-Arago hotspot and can be traced back into the Western
104 Pacific along an age progression (blue, Fig. 1; Extended Data Fig. 8). However, north of the
105 Marshall Islands, a data gap in the Rurutu-Arago hotspot track separated these ≤80 Ma volcanoes
106 from the ≥100 Ma Wake Seamounts, complicating efforts to trace the Rurutu-Arago track prior to
107 80 Ma. Recent sampling of seamounts around Wake Island fills this gap and new Pb-Sr-Nd isotopic
108 compositions (Methods) of these samples match those of the rest of the Rurutu-Arago hotspot track
109 (Extended Data Fig. 8, Extended Data Fig. 9).

110 Further west, the Cretaceous Magellan seamounts also span the ~80-100 Ma age range⁴
111 and overlap with the Samoa hotspot in composition. Critically, the age and location of these
112 seamounts matches the predicted track of Samoa (Fig. 1; Extended Data Fig. 8) that can be traced
113 from the present-day location near the northern terminus of the Tonga Trench to ~25 Ma Alexa
114 Bank, west of which the trace of the Samoan hotspot is lost in the Vitiiaz Trench. The predicted
115 Samoan hotspot track passes through the OJP, where its large lithospheric thickness²⁰ likely
116 suppressed Samoan plume melting and volcanic construction⁴⁶. The predicted hotspot therefore is
117 expected to emerge on the north side of the OJP as 80-100 Ma volcanoes, which is consistent with
118 the age and location of the Magellan Seamounts. Further supporting a link to the Samoan hotspot,
119 the Magellan seamounts feature the same unique combination of intermediate ²⁰⁶Pb/²⁰⁴Pb and
120 extreme radiogenic ⁸⁷Sr/⁸⁶Sr isotope compositions observed in Samoan shield lavas¹⁰ (red, Fig. 1),
121 and are capped by late-stage volcanism with a characteristic low ²⁰⁶Pb/²⁰⁴Pb and elevated ⁸⁷Sr/⁸⁶Sr
122 isotope composition^{45,47} (green, Fig. 1, Extended Data Fig. 3). Magellan seamount compositions
123 and ages⁴ are reassessed here, revealing a Samoan shield to rejuvenated stage isotopic sequence⁴⁷
124 at Hemler and Vlinder Seamounts, while samples obtained from the younger Magellan Seamounts
125 are similar to Samoan late stage rejuvenated compositions (Extended Data Fig. 3).

126 APM models for the predicted tracks of the high ²⁰⁶Pb/²⁰⁴Pb Rurutu-Arago hotspot and the
127 high ⁸⁷Sr/⁸⁶Sr Samoa hotspot suggest these two hotspot tracks should continue into the Mariana
128 Trench^{7,48} (Fig. 1). Geochemical data on Mariana arc volcanoes provides further support that the
129 Rurutu-Arago and Samoa hotspot tracks are subducted near the region proposed by APM models.
130 When the isotopic compositions of the Mariana arc volcanoes are examined, an unusual
131 geographically constrained high ²⁰⁶Pb/²⁰⁴Pb anomaly^{46,47,48,49} and an adjacent high ⁸⁷Sr/⁸⁶Sr
132 anomaly (which overlaps, but is shifted slightly to the south) are found where the Rurutu-Arago
133 and Samoa hotspot tracks are predicted to subduct based on the revised APM model (Fig. 1,
134 Extended Data Fig. 4). Consistent with the hypothesis that the subduction of seamounts impacts
135 the chemistry of nearby arc volcanoes, the ²⁰⁶Pb/²⁰⁴Pb and ⁸⁷Sr/⁸⁶Sr anomalies in the Mariana arc
136 provide supporting evidence for the presence of the long-lived Rurutu-Arago and Samoan hotspot
137 tracks in the western Pacific during the Early Cretaceous. Furthermore, their offset position in the
138 Mariana arc also supports the presence of a more northerly Rurutu-Arago hotspot track in the
139 western Pacific region predicted by our new APM, which is different from predictions by prior
140 APM models (Fig. 1, Extended Data Fig. 4). Consequently, we now can explore Rurutu-Arago

141 and Samoa as a pair of long-lived parallel (but offset) hotspot tracks, providing a multi-hotspot
142 continuation of APM constraints beyond the 80 Ma limit that is provided by the Hawaii-Emperor
143 and Louisville hotspot tracks.

144 While the Rurutu-Arago hotspot track is defined by seamounts with related geochemical
145 fingerprints that closely follow APM models back to 80 Ma, existing APM models⁷ predict the
146 track to significantly bend to the west around the Wake Seamounts (Extended Data Fig. 4). This
147 region is only sparsely populated by seamounts when compared to the abundance of seamounts
148 around Wake Island and the Wake Seamounts, which provide a clearer morphological continuation
149 of the hotspot track, but more importantly a geochemical match to the Rurutu-Arago hotspot^{29,48}
150 (Extended Data Fig. 4). The clear impact of Rurutu-Arago hotspot on Mariana arc
151 compositions^{48,49} (Extended Data Fig. 4) suggests that, in contrast to existing APM models, the
152 hotspot followed a more northerly track between 120 and 80 Ma—offset by >1200 km from
153 existing models. This alternative “northern path” (blue path in the Wake Seamounts in Fig. 1, “this
154 study” in Extended Data Fig. 4) for the pre-80 Ma Rurutu-Arago hotspot track provides a unified
155 explanation for all these observations, suggesting that a reconsideration of the APM models is
156 warranted.

157 The period of plate motion covered by our new data is constrained by the ~80-100 Ma
158 Marshall Island volcanoes and Wake Island area seamounts. Critically, existing ages for the two
159 volcanic chains correspond closely with the 80-100 Ma single plate rotation (stage) of the existing
160 K01 plate motion model¹⁸. For the period prior to 80 Ma, modern APM models⁵⁰ still define motion
161 with the large time-step plate rotations of early models^{2,18}. Given the limited availability of ages
162 around Wake Island, we resort to a similar approach as used in the K01 model³ and calculate plate
163 motion using large time-steps rather than applying a more continuous age models such as WK08³⁹.
164 For the 80-100 Ma time interval, we modeled the plate motion by finding the stage pole (5°N,
165 306°E) that minimizes the difference in Euler pole distance between the actual seamounts and the
166 predicted age progressive paths for the Samoa and Rurutu-Arago hotspot tracks (Methods). Using
167 the new modified APM model, K01m, our results show that the predicted track projects Rurutu-
168 Arago through the Wake area seamounts (blue path in Fig. 1), whereas prior models failed to
169 capture the Wake Islands and seamounts (Extended Data Fig. 4). Due to the southeasterly shift in
170 location of the K01m rotation pole compared to K01, the relative angular distances from the Euler
171 pole to Samoa and Rurutu-Arago have changed in a manner where the resulting predicted path

172 length for Samoa remains similar to older models, while that of Rurutu-Arago lengthens
173 significantly.

174 The application of this new 80-100 Ma rotation pole to the Louisville hotspot has profound
175 implications for linking the Louisville track to OJP-Nui. While prior plate motion models placed
176 the Louisville hotspot far from OJP at the time of eruption¹, the new 80-100 Ma rotation increases
177 the north-south component of the predicted Louisville path (Fig. 3). As a result, the new K01m
178 model using long-lived intraplate hotspots traces the predicted Louisville hotspot track directly
179 into the center of OJP-Nui by ~120 Ma, the modeled age of formation for the superplateau^{21,51}.
180 This is in contrast to existing APM models^{3,6,7,50} that all predict a more southerly track (Fig. 3) that
181 falls short of the OJP, most likely because these models incorporate near-ridge large structures
182 (rises) and seamounts in their datasets. An assessment of the fit of individual track segments using
183 backtracking with different APM models confirms this result (Methods; Extended Data Fig. 5).
184 The formation age of OJP-Nui has been estimated based on decades old lower quality ⁴⁰Ar/³⁹Ar
185 ages and plate reconstructions to be around 120 Ma^{21,51}, yet recent high-precision geochronology
186 suggests that the main phase of OJP volcanism began around 116 Ma⁵². This makes the
187 reconstructed link between Louisville and OJP-Nui even more likely.

188 However, in the formation of the Louisville hotspot track from OJP-Nui until present day
189 (Fig. 3), there are two time periods with poor constraints. First, the 80-90 Ma Louisville Seamounts
190 all have subducted into the Tonga Trench, interrupting the direct connection between the OJP and
191 the Louisville seamount track. Second, the location of the earliest portion of the Louisville
192 seamount track between 90-116 Ma depends on whether or not the Manihiki or Hikurangi Plates
193 (and their associated plateaus) rotated during the break-up¹ of OJP-Nui (Fig. 3), and whether there
194 exists an eruptive age progression throughout OJP-Nui given that Manihiki may predate OJP by
195 up to 6 Myr⁵². In either case, the eruptive location of the earliest Louisville track is within the OJP
196 outline and, importantly, we show for the first time that 90-115 Ma seamounts in the Ellice Basin
197 agree with the expected age progression, location, and isotopic signature of Louisville hotspot-
198 related volcanoes in the region¹⁴ (Fig. 2). These seamounts therefore provide the “missing link”
199 connecting the OJP to the start of the Louisville track. The Louisville Seamounts and Ellice Basin
200 seamounts are geochemically similar and plot along the array defined by Manihiki’s Danger
201 Islands and OJP melts (Fig. 2; Methods). Therefore, the Louisville and Ellice Basin seamounts are
202 geochemically and temporally linked to OJP-Nui.

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203 Our demonstrated fit of OJP-Nui to the Louisville hotspot track (Fig. 3; Extended Data Fig.
204 5) requires no significant plume motion, which agrees with estimates of limited latitudinal motion
205 of this hotspot between $\sim 70\text{-}0\text{ Ma}^8$. Our new data, and new interpretation of existing data, thus
206 present a simple argument for the genetic connection between Louisville and OJP-Nui. This
207 implies a plume-driven origin for the largest LIP in the geologic record, in contrast to recent
208 arguments for a major role for seafloor spreading in the construction of LIPs, such as Shatsky
209 Rise³⁵. Renewed consideration for other Cretaceous “orphan” LIPs—that may still lack obvious
210 volcanic trails—is warranted, in light of the suggested revision to the 80-100 Ma stage pole that
211 may serve to further address identified uncertainties in Pacific APM during the Cretaceous.

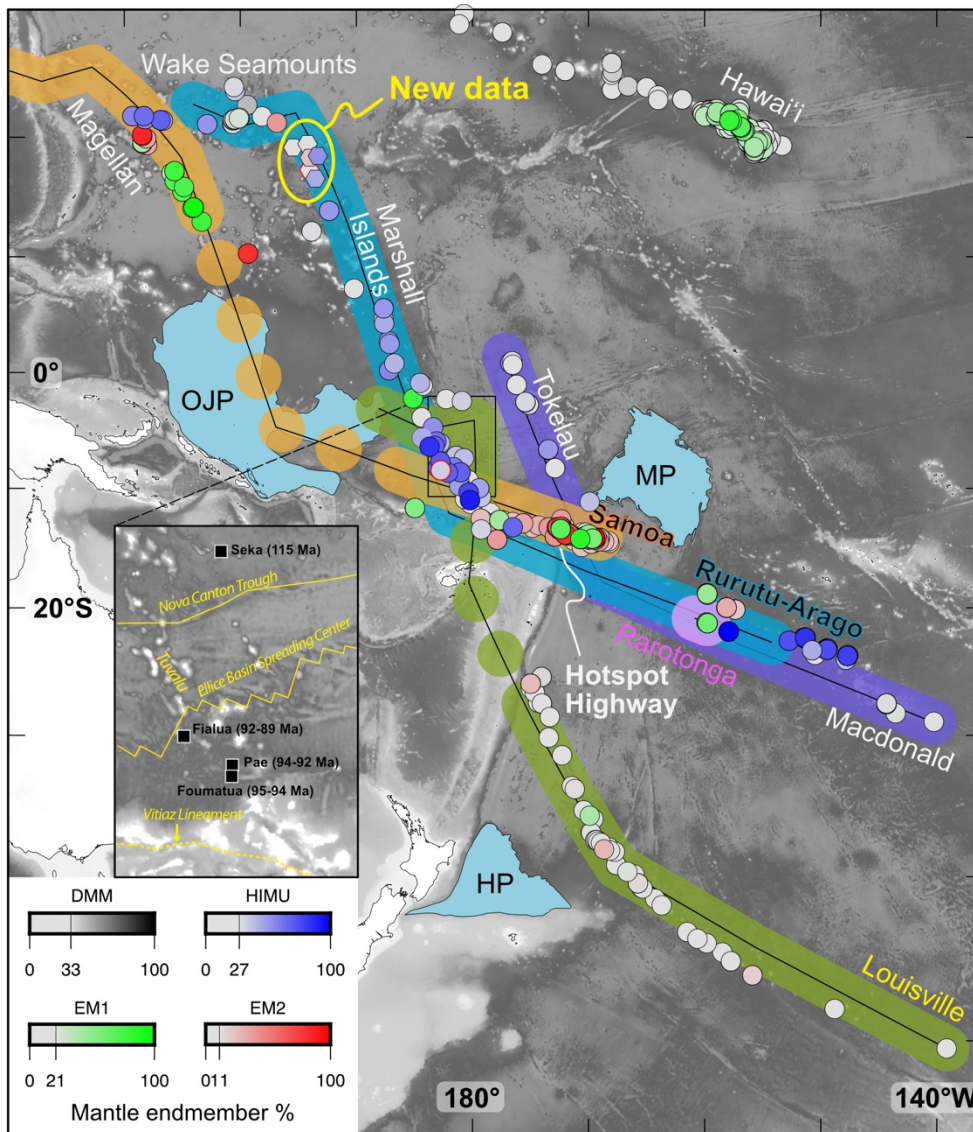
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219

220 FIGURES

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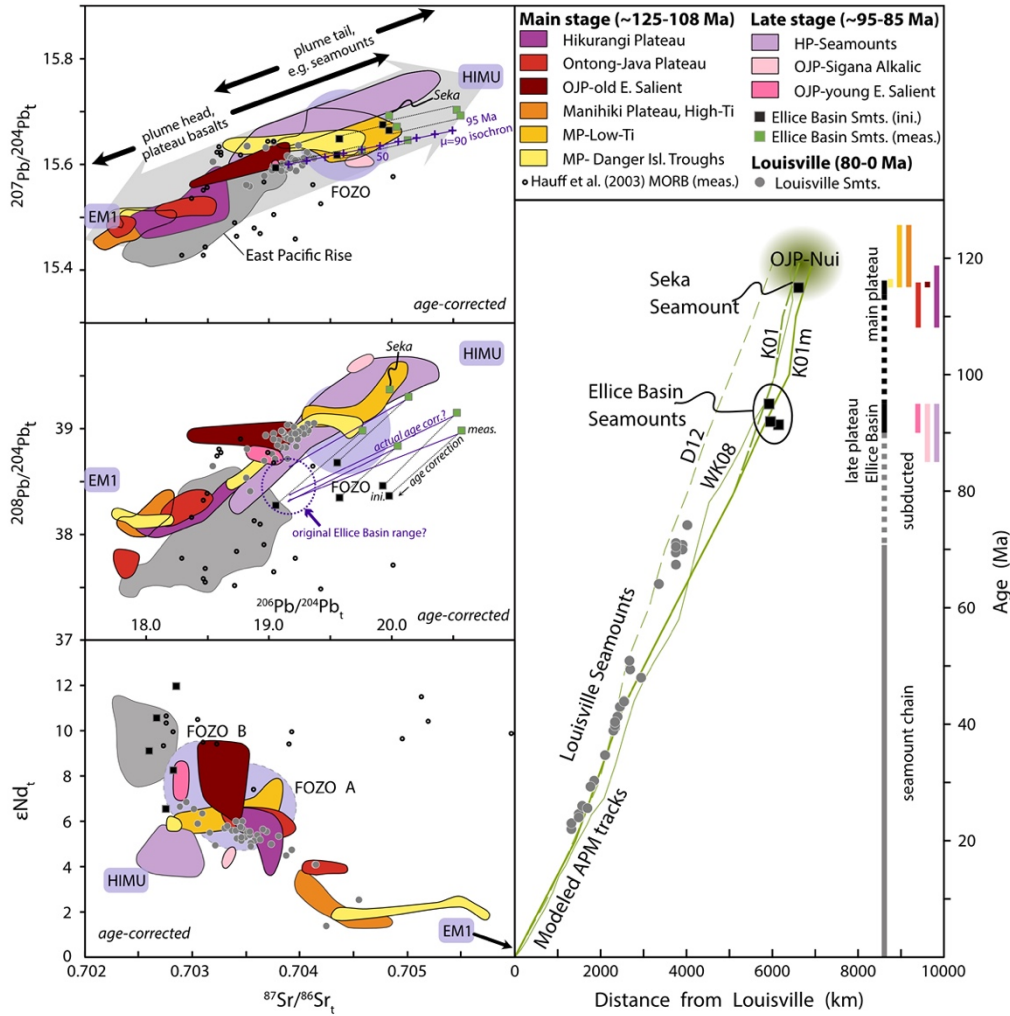
224 **Figure 1.** Pacific hotspot tracks and color-coded geochemical lava flow compositions (bottom
225 right inset, Methods) demonstrate unique signatures for Rurutu-Arago (blue circles) and Samoa
226 (red and green) and thus allow for the effective geochemical tracing of these hotspots, when
227 combined with seamount ages. Ellice Basin Group and the New Wake samples with a red outline
228 are confirmed phosphatized (See Alteration section in Methods) and not considered in discussions
229 of $^{143}\text{Nd}/^{144}\text{Nd}$ behavior. The known extents of the Rarotonga and Macdonald hotspot tracks are

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230 *provided for reference. The colored tracks show predicted hotspot tracks following the new APM*
231 *(specifically, 80-100 Ma stage represented by the Wake-Marshall and Magellan areas). The*
232 *dashed portions of the predicted Louisville hotspot track mark where the track was subducted. For*
233 *Louisville (green), the updated APM predicts an origin at Ontong-Java Plateau (OJP). The new*
234 *APM modeling relies on Wake-Marshall and Magellan seamounts, omitting the Shatsky and Mid-*
235 *Pac groups, and for the first time directly relates the Louisville hotspot to OJP. Hawaiian-Emperor*
236 *hotspot sample data are shown for reference. See Extended Methods for data sources. **Inset:***
237 *Locations and ages of all Ellice Basin Seamounts and Seka Seamount, as confirmed by*
238 *geochronology, as black squares. The Tuvalu chain, Nova Canton Trough, Ellice Basin Spreading*
239 *Center, and Vitiaz Lineament are shown in yellow.*

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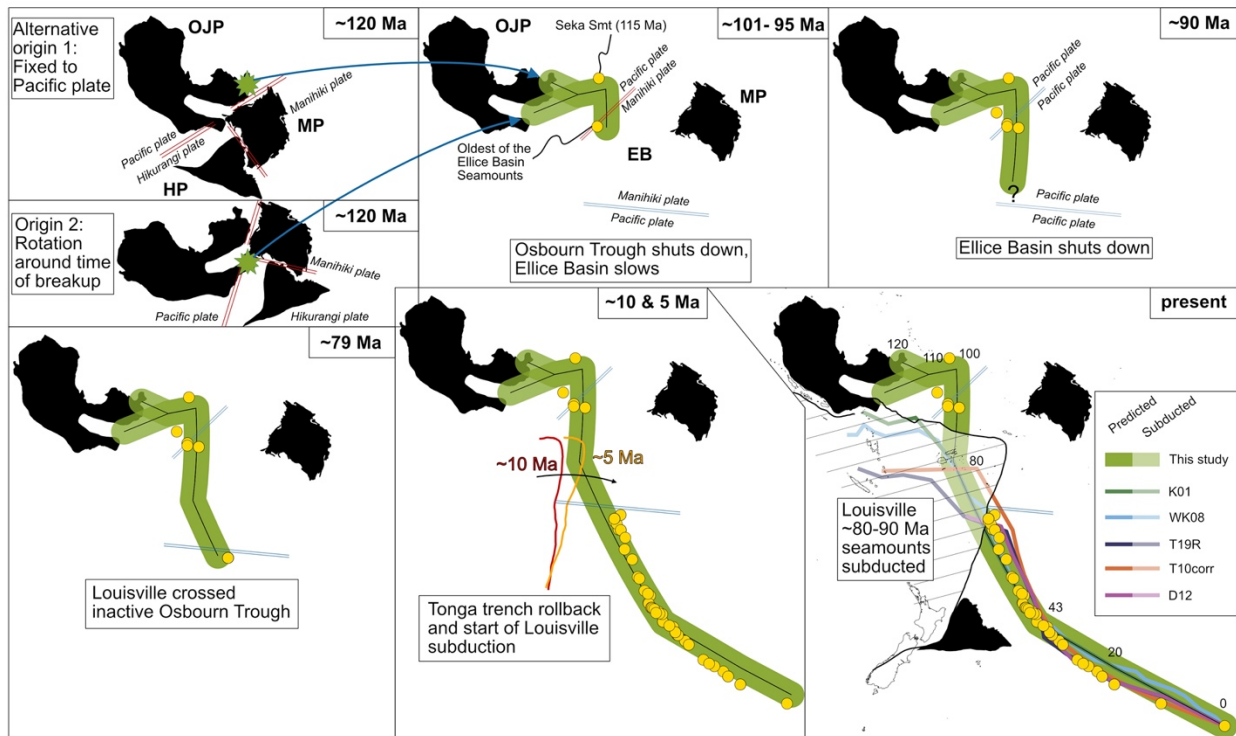
243 **Figure 2.** Right: Age progression for the Louisville hotspot track (symbols same as in plots). Ellice
 244 Basin Seamounts¹⁴ and Seka Seamount⁴⁵ are shown. Top: Age-corrected $^{207}\text{Pb}/^{204}\text{Pb}$ vs.
 245 $^{206}\text{Pb}/^{204}\text{Pb}$ isotope compositions for Ontong Java Nui, and related structures. Middle: Age-
 246 corrected $^{208}\text{Pb}/^{204}\text{Pb}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$ isotope compositions for Ontong Java Nui, and related
 247 structures. Bottom: $\epsilon^{143}\text{Nd}$ vs. age-corrected $^{87}\text{Sr}/^{86}\text{Sr}$ for Ontong-Java Nui and related structures.
 248 Pb-isotope data for Ontong-Java (OJP), Manihiki (MP), and Hikurangi plateaus (HP), which
 249 represent the main pulses of voluminous plateau-building volcanism (~120 and ~90 Ma), are
 250 frequently less radiogenic, but extend to radiogenic values in some settings. By contrast, smaller
 251 structures such as seamounts are confined to intermediate-to-radiogenic values. Louisville and
 252 Ellice Basin seamounts exhibit more radiogenic Pb-isotopes, with Ellice Basin being particularly
 253 radiogenic due to seawater U influence, but likely a mixture of FOZO and DMM that originally
 254 resembled slightly depleted Louisville-type compositions (see Methods). Green squares are

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255 *present-day Ellice Basin compositions, compared to corresponding age-corrected (where*
256 *possible) compositions. Louisville seamounts overlap directly with some OJP basalts in Pb-Nd-Sr*
257 *isotope space, and are well within the array of compositions found in the greater combined set of*
258 *plateaus and structures from OJP, MP and HP. Furthermore, Ellice Basin and Seka seamounts*
259 *are spatiotemporally linked to the OJP. See Methods for description of isotopic data compared to*
260 *variations in plume head and plume tail stages of volcanic activity, analysis of Pb-Sr-Nd isotopic*
261 *systematics in the Ellice Basin Samples, and data sources.*

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263



264

265 **Figure 3.** Cartoon of Louisville hotspot (green swath) evolution. The revised model predicts that
 266 the Louisville hotspot initiated within Ontong Java Plateau (OJP), however the exact initial
 267 location is sensitive to whether the plateaus rotated around the time of OJP-Nui breakup⁵¹, so two
 268 alternatives are shown for ~120 Ma. By ~95 Ma, the Ellice Basin (EB) had (mostly) opened
 269 between OJP and Manihiki Plateau (MP), as suggested by the age of the oldest Ellice Basin
 270 Seamounts (95.0 Ma¹⁴)—located centrally in the basin—which we link to the Louisville hotspot.
 271 Seka Seamount (115.0 Ma⁴⁵), located at the north end of the basin, is isotopically similar to the
 272 Ellice Basin seamounts (suggesting significant Ellice Basin spreading occurred by 115 Ma) and
 273 is also linked to the Louisville hotspot. The new model predicts Louisville to be present in the Ellice
 274 Basin at this time, progressing to younger ages to the south. The oldest Louisville Seamount
 275 erupted at ~79 Ma^{8,36}, located south of the Osbourn Trough, represents the oldest part of the
 276 continuously defined Louisville hotspot track. In the past 5 to 10 Ma, roll-back of the Tonga Trench
 277 caused the subduction of any Louisville Seamounts with ages between 79 to 90 Ma, leaving only
 278 the older (>90 Ma) Louisville hotspot-related structures in the Ellice Basin and OJP areas
 279 preserved. Consequently, the relationship between >79 Ma Louisville hotspot volcanism and the

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280 *paleoridge represented by the Osbourn Trough is unclear. Predicted hotspot tracks show that only*
281 *the new model tracks Louisville hotspot back to OJP-Nui, specifically the OJP. See Methods for*
282 *details on this model.*

283

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548 **METHODS**

549 **Sample analysis**

550 Samples were collected with an ROV, aboard the NOAA exploration vessel *Okeanos Explorer*,
551 during expedition EX1606 to the Wake Island Unit of the U.S. Pacific Remote Island Marine
552 National Monument. Dive samples were cut open and the six least altered samples were selected.
553 Pb-Sr-Nd isotopic compositions were obtained following mechanical and chemical procedures to
554 remove seafloor alteration contributions⁴ and previously detailed elemental separation
555 procedures⁵³. In short, small piece from the center of each sample was crushed, altered pieces
556 were removed by hand-picking, the clean fraction was acid-leached, and subsequently dissolved
557 for Pb separation using Eichrom resin. Isotope measurements were performed on a Nu Plasma
558 HR MC-ICP-MS at University of Hawai'i at Mānoa (Pb) and a ThermoFinnegan NeptunePlus at
559 University of South Carolina (Sr, Nd, and Hf⁵⁴). Additional analytical details, results and brief
560 sample characterizations, including assumed age-corrected isotopic ratios for the EX1606
561 samples, are reported in the Extended Data Table 1. Isotopic behavior shown in Extended Data
562 Fig. 8 and Extended Data Fig. 9 demonstrate the compositional agreement between the new
563 samples characterized here, and published data for the Rurutu-Arago hotspot track.

564 Trace elements were collected using a ThermoFinnegan Element2 at University of South
565 Carolina on aliquots of powders lightly leached with 0.1N HCl in a sonic bath for 20 minutes to
566 mitigate any low-temperature overprinting by seawater without compromising the primary bulk
567 composition.

568 **Color coding**

569 In order to show the compositional distinction between the hotspot tracks in the western Pacific,
570 as well as to show how compositions backtrack around present-day hotspots, the radiogenic
571 isotope composition of the tracks are color-coded^{14,17,45}. The technique focuses on use of
572 ⁸⁷Sr/⁸⁶Sr-¹⁴³Nd/¹⁴⁴Nd-²⁰⁶Pb/²⁰⁴Pb isotope compositions, because principal component analysis
573 shows these represent the largest compositional variations in hotspots^{55,56}. In this space, hotspots
574 scatter between four extreme end-member compositions: HIMU, EMI, EMII, and DMM⁵⁷. Each
575 hotspot's samples define an elongated, prolate ellipsoid, and these ellipsoids (32 in the global
576 hotspot database) radiate from a central region, known as FOZO⁵⁶ or C⁵⁸. The color-coding⁴⁵

577 assigns a level of color-saturation based on its distance from this center, while the color depends
578 on the end-member that a particular sample trends closest to (Fig. 1 inset). Compositions that are
579 on the trend toward an end-member, but cannot be resolved from the central FOZO⁵⁶ or C⁵⁸
580 component are colored grey (note that the cutoff percentages for each end-member are shown on
581 colorbars in the Fig. 1 inset). Using this color coding approach results in a “blue” color for
582 symbols, representing HIMU compositions, which dominate hotspot track that run from the
583 Cook-Austral Islands in the central equatorial Pacific to seamount tracks in the Western Pacific.
584 This HIMU compositional “signature band” in map view (i.e., as the blue symbols the represent
585 the Rurutu-Arago HIMU hotspot track form a “band” across the south Pacific and into the
586 Western Pacific Seamount Province) is consistent with predicted absolute plate motion (APM) in
587 the Pacific until at least 80 Ma, and forms the basis for our investigation that links the Wake area
588 seamounts in the Western Pacific to the long-lived Rurutu/Arago hotspot that also includes the
589 Tuvalu islands and the Gilbert Ridge seamount track. In addition, the red (representing EM2) and
590 green (EM1) compositional color coding for Samoa appears in young Samoan hotspot lavas and
591 also appears in the far western Pacific in the Cretaceous Magellan seamounts. Recent work in
592 Samoa has shown a temporal compositional evolution from red (EM2) to green (EM1),
593 representing the evolution from shield to rejuvenated stage of volcanism at Samoa, and the same
594 is observed in the Western Pacific when compositions and ages are assessed (Hemler and
595 Vlinder seamounts^{4,59}; Extended Data Fig. 5).

596 **Absolute Plate Motion Modeling**

597 Although initial models^{2,60} already showed that simple plate rotations do fit the shapes of several
598 hotspot tracks, recent models³⁹ have evolved to include more sophisticated techniques to find the
599 best-fit absolute plate motion models by using multiple hotspots and by resolving continuous
600 plate rotations from both seamount locations and ages. In addition, some models include large
601 scale mantle flow, comparisons between ocean basins, and plume motion^{6,38,61}. For the Pacific
602 plate >80 Ma, little data are available as neither Hawaii nor Louisville continue past 80 Ma^{88,62},
603 and plate circuits that would allow comparison with the Indo-Atlantic hotspot do not extend
604 beyond this time either⁶. In most existing models, the time period prior to 80 Ma is populated
605 with data from Shatsky Rise and the Mid-Pacific Mountains, yet their respective structures were
606 within ~1,000 km of the ridge system⁶³ (Extended Data Fig. 1) and during their formation these

607 tracks are likely to have been influenced by ridge interaction⁵⁵. However, the Rurutu-Arago and
608 Samoa hotspots do define tracks in this age range that are clearly distinct in their isotopic
609 compositions (Fig. 1; see Section on Color Coding) and that are truly intraplate prior to 80 Ma
610 (Extended Data Fig. 1). We conclude that Rurutu-Arago and Samoa therefore are a more faithful
611 reflection of APM for the Pacific plate.

612 The presence of two hotspot tracks—Samoa and Rurutu-Arago—provides a means to fit a plate
613 rotation for 80-100 Ma that fits both hotspots, thereby relying on their common motion to define
614 an APM pole for that period. Unlike younger volcanic tracks, Samoa and Rurutu-Arago provide
615 enough data to outline the overall hotspot track; however, the sample and data density are low,
616 such that constraints are lacking for a high-resolution model³⁹. Instead, the available data here
617 are used to identify an 80-100 Ma section in both volcanic tracks (Extended Data Fig. 6;
618 Extended Data Tables 2, 3), after which a least-squares method is used to find the best-fit
619 rotation pole for both hotspots. The method¹⁸ consists of finding the best-fit pole that minimizes
620 the variance in calculated angular (seamount-pole) distances, in the least-squares sense. The
621 solution is found with a grid-search algorithm that tests

$$622 \quad \text{var}(d_{ij}) = \sum_{j=1}^M \sum_{i=1}^N \frac{(d_i^j - d_{mean}^j)^2}{N}$$

623 which calculates the variance in distance (d) for all seamounts in the 80-100 Ma track segment
624 relative to the mean distance for a given hotspot (d_{mean}). The variation in distance per hotspot is
625 normalized by the number of seamounts (N) per hotspot. This technique suggests a best-fit stage
626 pole west of the Samoa-related seamounts, near the equator (5.0°N, 54.0°W, rotation rate:
627 0.85°/Myr; Extended Data Table 2). The resulting K01m APM model is a much better fit to the
628 two hotspot tracks, particularly for the Rurutu-Arago hotspot, where the predicted track now
629 bends to the west at ~100 Ma, just northwest of Wake Island (Fig. 1, Extended Data Fig. 4, 5).
630 Due to the shift in the rotation pole compared to prior models, the angular distance from the pole
631 to the Rurutu-Arago track and the new rotation rate provide a better fit to the seamount track.
632 Despite these improvements, several issues must be considered before the new model can be
633 applied to the Louisville hotspot track.

634 Plate motion southeast of OJP involved more than just the Pacific plate. During the break-up of
635 OJP-Nui, the Manihiki and Hikurangi plates formed, while the Chazca plate resided immediately
636 to the east⁵⁰. It is important to evaluate whether these plates moved independently with respect to
637 the Pacific plate while the Ellice Basin was located over the Louisville hotspot. The relevant
638 plates/microplates are associated with the break-up of OJP-Nui. The oldest Louisville Seamounts
639 (ranging in age from 1 to 79 Ma^{8,36}) are located south of the Osbourn Trough and west of the
640 Wishbone Scarp, representing a small plate forming as Hikurangi separated from OJP. However,
641 true spreading on the Osbourn Trough may have ceased between 101-100 Ma^{64,65}. Any further
642 rotation of this plate ceased between ~84 Ma-79 Ma^{66,67}, so Louisville Seamounts younger than
643 79 Ma formed after this plate was already part of the Pacific plate. Thus, Louisville Seamounts
644 after 79 Ma properly represent Pacific plate motion with respect to the Louisville hotspot. The
645 only other area where older volcanoes related to Louisville may be exposed is in the area of the
646 Ellice Basin between OJP and Manihiki¹⁴. However, the exact timing of cessation of motion
647 between the separating plateaus is not well constrained²¹. The sparse age information from the
648 Ellice Basin includes the 95 Ma Foumatua seamount located on the fossil ridge system¹⁴,
649 constraining a minimum age of ~95 Ma for when seafloor spreading in the Ellice Basin shut
650 down, as suggested by models based on seafloor fabric⁶⁸. Critically, Foumatua Seamount is part
651 of a group of similarly aged “Ellice Basin Seamounts” that have an isotopic composition that
652 overlaps with that of the younger Louisville Seamounts¹⁴ (Fig. 2). These seamounts are located
653 where our APM model predicts the Louisville hotspot around ~95 Ma, and the seamounts are
654 also age-progressive with the Louisville seamounts (Fig. 1, 2). The 115 Ma Seka Seamount,
655 located north of the Ellice Basin fossil ridge and near the southern terminus of Gilbert Ridge
656 several hundred kilometers east of the OJP, places the underlying seafloor – and westward
657 motion of the OJP fragment relative to the plume – at a minimum age of 115 Ma, thus indicating
658 that early stage Ellice Basin opening was rapid.

659 Due to the known wide variation in paleomagnetic latitude estimates for OJP⁵¹, any microplate
660 considerations prior to ~100 Ma are underconstrained at best. Either the plateaus rotated around
661 the time of break-up⁵¹, or OJP was already fixed with respect to the Pacific plate⁶⁷. Regardless,
662 the sheer size of the entire Ontong Java Nui combined structure is so large that neither a rotation
663 of the plateaus, nor ongoing spreading in the Ellice Basin until ~100 Ma, would place the
664 predicted Louisville hotspot outside of the outline of the combined OJP-Nui plateau (or even

665 OJP by itself) at 120 Ma (Fig. 3). This is mainly due to the final east-west length of the 100-120
666 Ma predicted Louisville hotspot track being shorter than the width of OJP, which is thought to
667 have largely moved east-west^{21,68} (Fig. 1). In Fig. 3, the approximate eruptive locations are also
668 shown assuming OJP either was fixed from ~120 Ma, or only rotated around a pole internal to
669 the plateau⁵¹. Intriguingly, the rotation of the plateaus places the eruptive location at ~120 Ma
670 near the triple junction between the three plateaus.

671 **Backtracking**

672 Individual volcanoes that have been age-dated can be backtracked to their original eruptive
673 location, by rotating present-day volcano locations in the opposite direction of plate motion
674 according to the sequence of APM model rotations: Existing APM models are used as stage pole
675 rotations to accommodate volcanic ages in between supplied finite rotations. Instead of each
676 volcano backtracking to the same single hotspot location, this process generates clusters due to
677 various geologic and measurement uncertainties, such as: (1) volcanic age precision and
678 accuracy, (2) extended eruption of single volcanoes (up to 7 Ma⁶⁹), (3) offset between sampled
679 rift zone eruptions and the central crater, and (4) lithospheric structure offsetting volcano from
680 mantle source^{4,70}. Cumulatively, these uncertainties could cause hundreds of km of scatter in
681 backtracked locations that cluster around the present-day hotspot⁴⁵.

682 Two categories of structures were backtracked here: (1) individual volcanoes of Rurutu-Arago,
683 Samoa and Louisville, and (2) the OJP as a single structure. For the individual hotspot volcanoes
684 (Extended Data Table 3), published ⁴⁰Ar/³⁹Ar ages were used^{9,71-75}, while symbols for
685 reconstructed volcanoes are colored (see Color Coding) based on their isotopic composition<sup>10-
686 12,14,44</sup>. This highlights the compositional groups by color, matching present-day hotspot locations
687 at Samoa and Rurutu-Arago, as well as Macdonald and Rarotonga for reference. Moreover, the
688 probability density function of backtracked locations (constructed with a Gaussian kernel) shows
689 four peaks in the distribution that also correspond to the same four present-day hotspots
690 (Extended Data Fig. 5). In addition to the density estimates, a running mean can also be
691 calculated for the backtracked locations and related sample ⁴⁰Ar/³⁹Ar ages. These “age tracks”
692 represent a smoothed estimate of how the hotspot source moved in time, with respect to the
693 applied APM model, which can be thought of a proxy for plume motion in that reference frame.

694 The results for the various models emphasize the tighter scattering (in blue HIMU-composition
695 around Rurutu-Arago, red-green around Samoa) and shorter plume motion tracks for the new
696 K01m model. This result implies the least amount of plume motion is required for these models,
697 while models that allow for plume motion require a significant amount, but also match predicted
698 plume motion^{6,61} back to ~50 Ma for Hawaii and Louisville (Extended Data Fig. 5). Since there
699 are no plate circuits >80 Ma to enable comparison to the Indo-Atlantic hotspot reference frame³⁸,
700 the modeling results can only be tested against Pacific hotspots.

701 While testing the model against Rurutu-Arago and Samoa constraints is potentially circular,
702 backtracking the OJP constitutes a more interesting test, as it was previously shown to backtrack
703 1,200 km away from the closest hotspot at 120 Ma, i.e., Louisville¹. By defining the outline of
704 the plateau as a series of individual points⁷⁶, each individual point is backtracked with the
705 updated K01m APM model. The result shows that the backtracked OJP plateau outline is located
706 directly over the present-day Louisville hotspot (Extended Data Fig. 5). Similarly, the
707 backtracked individual Louisville seamounts are also located over the present-day Louisville
708 hotspot. The updated K01m model thus resolves the north-south discrepancy between modeling
709 predictions of the eruptive location of the OJP and the track of Louisville. The conclusions for a
710 120 Ma OJP formation also hold for a slightly younger age of formation, recently suggested to
711 be ~116 Ma⁵². In this case, the modeled Louisville hotspot and Eastern Salient of the OJP are
712 still closely spatially associated with each other at ~116 Ma. The revised OJP formation age is
713 similar to that of Seka Seamount, which would further suggest that OJP-Nui formation and
714 initiation of Ellice Basin rifting were simultaneous events. This model represents a fixed hotspot
715 model, while multiple lines of evidence suggest hotspot sources to be mobile⁵. However, motion
716 estimates for Louisville hotspot are within error of its location until at least ~70 Ma^{8,9}, and older
717 data for greater Ontong-Java Nui are too variable to constrain any possible plume motion.
718 Regardless of a lack of tight paleolatitude constraints, the size of greater Ontong-Java Nui allows
719 for hundreds to thousands of kilometers of plume motion superimposed on our new model, while
720 still maintaining the “end” of the Louisville hotspot track within its outline.

721 **Background Data Sources for Fig. 1**

722 Most of the previously published data used in Fig. 1 were sourced from GEOROC
723 (<https://georoc.eu/>) precompiled files for ocean islands and seamounts, which were filtered to

724 remove sample data known to be affected by contaminants and/or analytical problems. Non-
725 igneous lithologies were also filtered out. Additional isotope and/or geochronological data were
726 compiled for the Northwest Hawaiian Ridge⁷⁷⁻⁷⁹, Vlinder Seamount⁸⁰, Macdonald Seamount⁸¹,
727 Arago Seamount⁸², and Moki Seamount and Rose Atoll⁸³.

728

729 **Isotope data groupings used in Fig. 2**

730 The isotopic compositional range spanned by the plateau and smaller structures follow a
731 general temporal pattern. The East Pacific Rise field⁸⁴ is given for comparison. The majority of
732 the plateau data from OJP, MP, and HP plots between EM1⁵⁷ and a central geochemical
733 component, like FOZO⁵⁶. The OJP data⁸⁵⁻⁸⁸ consist of the major basement Kwaimbaita and
734 Kroenke components, which represent a relatively high degree of mantle melting with a
735 composition between EM1-FOZO, while the Singgalo component represents a later phase of
736 plateau construction, with a more EM1-rich composition⁸⁹ during the LIP stage of the plume.
737 These OJP stages are shown as red fields in Fig. 2, and they are followed by a much later and
738 smaller ~91 Ma alkalic stage⁸⁶ (Sigana Formation) that shifted to HIMU⁵⁷ compositions (pink
739 field in Fig. 2) in the later stages of OJP activity. Data from MP show basement rock
740 compositional groups similar to the Kwaimbaita/Kroenke and Singgalo lavas at OJP, but
741 recognized⁹⁰ as high-Ti EM1-type lavas (dark orange field in Fig. 2), while low-Ti lavas have
742 FOZO-type compositions incorporating potentially a small amount of a HIMU-type source (light
743 orange field in Fig. 2). This HIMU component is also argued to be the primary source for MP
744 late-stage lavas, just as it is at OJP; similarly at HP, the EM1-FOZO main plateau lavas (dark
745 purple, Fig. 2) are followed by the late stage HIMU HP Seamounts^{28,86,89,91} (light purple, Fig. 2).
746 Essentially, EM1-FOZO melts were dominant during the voluminous plateau-building stages,
747 while the later, less voluminous stages of volcanic activity shifted to dominantly HIMU-FOZO
748 melts. It is this full range of compositions and transition from plume head (plateau-building) to
749 plume tail (seamount chain-forming) activity that provides the critical context for the potential
750 chemical connection between Louisville Seamounts and OJP-Nui lavas²⁸, as well as the Ellice
751 Basin seamounts.

752 Given the well-known challenges involved in linking plateau-forming and seamount-
753 forming phases of mantle plume activity using plate motion models, the Ellice Basin Seamounts

754 and Seka Seamount offer an alternative approach: Their proximity to the OJP and their 90-115
755 Ma age range provide evidence of a temporal relationship to OJP-Nui (and specifically the OJP),
756 while representing smaller, seamount-forming stages of volcanic activity more directly
757 comparable to the Louisville hotspot track than the plateau-forming melts. The observation that
758 the 90-115 Ma Ellice Basin and Seka seamounts follow the Louisville age progression (Figure 2)
759 and fill in a critical gap along the age progression between Louisville (1 to 79 Ma) and OJP (120
760 Ma) lends strong support to the hypothesis that these seamounts provide the “missing link”
761 between the Louisville hotspot and OJP.

762 Ellice Basin Seamounts and Seka Seamount exhibit varying degrees of seawater U
763 enrichment consistent with patterns identified in some Jurassic Pacific MORB⁹². This alteration
764 affects uraniumogenic ²⁰⁶Pb and to a lesser extent ²⁰⁷Pb, but not thorogenic ²⁰⁸Pb as Th is largely
765 immobile during low-T alteration processes. Thus, seawater U alteration is detectable when
766 samples plot right of the NHRL in ²⁰⁸Pb/²⁰⁴Pb versus ²⁰⁶Pb/²⁰⁴Pb, which over time develops into
767 a broadly horizontal array away from analogous unaltered samples that have retained their
768 primary isotopic composition (Fig. 2). In ²⁰⁷Pb/²⁰⁴Pb versus ²⁰⁶Pb/²⁰⁴Pb, however, affected
769 samples will plot close or slightly under the NHRL, with ingrowth following an isochron
770 consistent with the time of alteration (Fig. 2, top right panel). Samples affected in this manner
771 may, however, remain under-corrected (too radiogenic) as the measured U abundances may not
772 adequately reflect the true ingrowth rate⁹². The Ellice Basin Seamounts scatter right of the FOZO
773 region in ²⁰⁸Pb/²⁰⁴Pb versus ²⁰⁶Pb/²⁰⁴Pb space, while ²⁰⁸Pb/²⁰⁴Pb overlaps with the least
774 radiogenic samples from the Louisville hotspot track^{28,93}. Further, in ²⁰⁷Pb/²⁰⁴Pb versus
775 ²⁰⁶Pb/²⁰⁴Pb, the Ellice Basin samples define an array consistent with a 95 Ma isochron from a
776 median Louisville initial Pb composition (Fig. 2) that intersects with age-corrected Louisville
777 hotspot track samples and thus suggests, despite undercorrection for ²⁰⁶Pb ingrowth, the two
778 groups share a genetic link.

779 The ¹⁴³Nd/¹⁴⁴Nd versus ⁸⁷Sr/⁸⁶Sr, the Ellice Basin Seamounts (and Seka) are depleted^{14,45},
780 resembling varying mixtures of FOZO and DMM consistent with our conclusions about the
781 original Pb isotope composition of the Ellice Basin and Seka seamounts (Fig. 2). The least
782 radiogenic (age-corrected) Ellice Basin-type samples overlap with the most radiogenic
783 compositions in the Louisville hotspot track field, further corroborating a genetic link between

784 the two groups. The Ellice Basin Seamounts are similar in age to crustal ages inferred for the
785 Ellice Basin itself⁶⁸; thus the Ellice Basin Seamount mantle source melted below young, thin
786 lithosphere. As a result, the Ellice Basin seamount melts likely incorporated a larger proportion
787 of a DMM-type component than the younger Louisville seamounts that erupted below older,
788 thicker lithosphere⁹³, not unlike increasing proportions of DMM observed in the older stages of
789 the Hawaiian and Louisville hotspot tracks as a function of proximity to a ridge^{94,95} (Extended
790 Data Fig. 9). The combination of 1) overlapping componentry between Ellice Basin and Seka
791 seamounts and the Louisville hotspot track, 2) spatiotemporal proximity of the Ellice Basin and
792 Seka seamounts to the OJP, and 3) excellent agreement of Ellice Basin and Seka to our new
793 APM model reconstruction of the Louisville hotspot track (Fig. 1) and age progression
794 (Extended Fata Fig. 8) provide the first robust evidence of the long-proposed link between
795 Louisville and OJP-Nui.

796 **Age corrections to isotope data**

797 Data in Fig. 2 were age corrected to initial isotopic ratios in order to compare isotopic
798 compositions of different temporal groups and/or distinct compositional groups characterized in
799 the OJP, MP, and HP as well as the Louisville hotspot track and Ellice Basin Seamounts. Where
800 possible, ages obtained on specific samples were used to correct their isotopic ratios; for suites
801 with known age constraints, recommended age estimates²⁸ were used for samples without age
802 determinations. Parent-daughter (P/D) ratios used to calculate the time-integrated ingrowth
803 relative to ²⁰⁴Pb were determined from trace element data and using decay constants of
804 $1.55125 \times 10^{-10} \text{ yr}^{-1}$ for ²³⁸U, $9.8571 \times 10^{-10} \text{ yr}^{-1}$ for ²³⁵U, and $4.948 \times 10^{-11} \text{ yr}^{-1}$ for ²³²Th.

805 All isotope data for the new Wake samples were also age-corrected using the same decay
806 parameters as for the Louisville data, and assumed ages of 100 Ma for the six seamounts to
807 simulate a reasonable estimate of initial ratios. Initial ratios were then forward-modeled to
808 present-day ratios using estimates of parent-daughter ratios of the HIMU mantle source for Pb,
809 Nd, and Hf isotopes⁹⁶ and (because Nebel et al.⁹⁶ do not provide Rb/Sr ratios) primitive mantle
810 parent-daughter ratios for Sr isotopes⁹⁷, following parameters of an identical exercise¹⁴. These
811 data are reported in Extended Data Table 1.

812 For most of the new Wake samples, age-corrections to $^{87}\text{Sr}/^{86}\text{Sr}$, $^{206}\text{Pb}/^{204}\text{Pb}$, and
813 $^{143}\text{Nd}/^{144}\text{Nd}$ isotope ratios are large, producing initial isotopic ratios significantly less radiogenic
814 than expected for the modern mantle source. Forward-modeling those initial ratios back to
815 present-day estimates, however, produces model ratios comparable to most of the measured
816 sample data (Extended Data Table 1). Prior modeling on 100 Ma alkali ocean island basalts
817 found similar disagreement between age-corrected and modern isotope ratios, particularly when
818 parent-daughter ratios are high (e.g., Pb isotopes in HIMU melts), yet also good agreement
819 between forward-modeled and measured ratios⁹⁸. This suggests that the utility of age corrections
820 for old alkaline basalts with high P/D ratios may be questionable when the intent is to compare
821 isotopic compositions with their modern counterparts that have undergone little radiogenic
822 ingrowth.

823 Figure 1 and Extended Data Figure 9 provide examples of the comparability of non-age
824 corrected (i.e., measured) Cretaceous and modern lavas from the Rurutu-Arago hotspot;
825 measured $^{143}\text{Nd}/^{144}\text{Nd}$ ratios from Cretaceous parts of the hotspot track, for example, closely
826 resemble $^{143}\text{Nd}/^{144}\text{Nd}$ ratios from the youngest part of the track. Furthermore, trace element data
827 are not available for all previously published isotope data used in this study, and thus age
828 corrections are not possible for all data. For completeness, only measured isotopic ratios are
829 shown in figures discussing seamount geochemistry. The exception is Fig. 2, where a broad
830 range of OJP-Nui-Louisville melts, which represent a range of high and low degree melts and
831 mantle source compositions produced from 120 Ma to present, are shown.

832 **Alteration**

833 Cretaceous seafloor rocks are nearly ubiquitously altered by seawater and various secondary
834 mineralization processes. Abundances of fluid-mobile elements, and correspondingly some
835 parent-daughter ratios of traditional radiogenic isotope systems used to identify mantle sources
836 of intraplate volcanoes, may be modified by these processes. Alternatively, dissolved trace
837 metals in seawater (e.g., Sr, Nd) can overprint the magmatic isotopic signature of a lava. Strong
838 acid leaching is employed to remove secondary isotopic overprints obscuring primary magmatic
839 signatures²⁹. However, cryptic alteration of some isotopes and/or time-integrated primary

840 isotopic ratios resulting from modified parent-daughter ratios may also be present, and cannot be
841 restored via leaching.

842 Trace element ratios such as Th/U and Y/Y* can be used to proxy U mobility and
843 phosphatization processes, respectively. In Extended Data Fig. 10, elevated Th/U correlates with
844 $^{208}\text{Pb}/^{204}\text{Pb}$ of the Wake samples, but no correlation exists with $^{206}\text{Pb}/^{204}\text{Pb}$ (but we note that a
845 lack of correlation is not necessarily related to alteration). Regardless, the new Pb isotopic ratios
846 reported here on Wake lavas are consistent with the range defined by younger segments of the
847 Rurutu-Arago hotspot track (Extended Data Fig. 7), so any isotopic modification by seafloor
848 alteration does not cause the new Wake samples to plot outside of the known Rurutu-Arago
849 isotope field. No correlation with Th/U is apparent with $^{143}\text{Nd}/^{144}\text{Nd}$ or $^{87}\text{Sr}/^{86}\text{Sr}$. When Y/Y* (a
850 proxy for phosphatization) is plotted (Extended Data Fig. 10), the most radiogenic $^{143}\text{Nd}/^{144}\text{Nd}$
851 occurs with the strongest positive Y/Y* anomaly, but no correlation with Pb isotopes is
852 observed. The same pattern is present in the Ellice Basin Group samples (where characterized for
853 bulk compositional data¹⁴), suggesting that samples with sufficiently high levels of
854 phosphatization develop DMM-like $^{143}\text{Nd}/^{144}\text{Nd}$, which remains unusually radiogenic even after
855 age correction. Based on trace element evidence, we exclude $^{143}\text{Nd}/^{144}\text{Nd}$ of phosphatized Ellice
856 Basin and Wake samples from further discussion; data for samples suspected of phosphatization
857 are shown in plots but are distinguished from the other samples. Ellice Basin samples lacking
858 bulk composition data are also flagged in relevant figures because the extent of phosphatization
859 cannot be evaluated for those samples.

860

861 **Plate motion history and model shown in Fig. 3**

862 The exact plate motions of the pieces of greater Ontong Java Nui are still not well
863 established, however paleomagnetic data of the plateaus, as well as seafloor morphology and
864 ages between the plateaus has led to different suggestions^{21,51,67,68}. In the scenario where the OJP
865 was fixed with respect to the Pacific Plate around ~120 Ma, the new model predicts an ~120 Ma
866 eruptive location on the north side. Alternatively, the plateaus rotated around the time of break-
867 up around a rotation axis internal to OJP⁵¹, and the 120 Ma Louisville eruptive location would be
868 approximately central between the three plateaus that separated. Critically, when employing our

869 new absolute plate motion model, the large size of greater Ontong Java Nui contains the original
870 eruptive location of Louisville, regardless of the uncertainties regarding these motions. By ~95
871 Ma, HP had rifted south, and the Ellice Basin between OJP and MP had mostly opened⁶⁷. The
872 oldest Ellice Basin Seamount (95 Ma) is located on the fossil ridge^{14,68} and thus defines a
873 minimum age for final Ellice Basin spreading. An older seamount (Seka, 115 Ma⁴⁵), located
874 near the southern Gilbert Ridge, has a similar isotopic composition to the Ellice Basin
875 Seamounts and is likely genetically related (Fig. 2), a hypothesis supported by the observation
876 that Seka both lies on the new reconstructed hotspot track for Louisville and lies on the
877 Louisville age progression. The presence of this seamount further supports early, rapid opening
878 of the Ellice Basin by the time the main plateau-forming phase of OJP-Nui was waning. By this
879 time, MP had been captured by the Pacific plate⁶⁷, so a Louisville track prediction using Pacific
880 plate motion is appropriate. Around ~90 Ma, the exact relationship of the Louisville hotspot to
881 the Osbourn ridge is unclear; volcanoes of this age range have been subducted, and the lack of
882 seafloor magnetic anomalies around the Osbourn Trough makes establishing exact spreading
883 rates within a 95-80 Ma time span difficult. However, motion may still have continued on the
884 Osbourn plate boundary until 79 Ma, after which the entire area was captured by the Pacific
885 plate⁶⁷. At ~79 Ma, the oldest Louisville seamount^{8,36} erupted south of the Osbourn Trough, so
886 all of the Louisville seamounts that erupted from that time onward were subjected to Pacific
887 plate motion. Much later, the Tonga trench experienced roll-back, while the Pacific plate
888 continued to subduct, initiating Louisville Seamount subduction around ~5 Ma⁹⁹ and continuing
889 today. The “present” panel of Fig. 3 also shows alternative APM model predictions, all
890 reconstructing south of OJP; K01¹⁸; WK08³⁹; T19R: model R⁵⁰; T10corr¹⁰⁰: subsequently
891 corrected⁵⁰; D12⁶.

892

893

894

895 **Data availability**

896 All data generated during this study are included in this published article (and its
897 supplementary information files), and are available in the EARTHCHM repository (doi will be
898 supplied).

899

900 **Code availability**

901 The best-fit plate rotation Matlab code is available upon request from the corresponding
902 author.

903 **Author contribution statement**

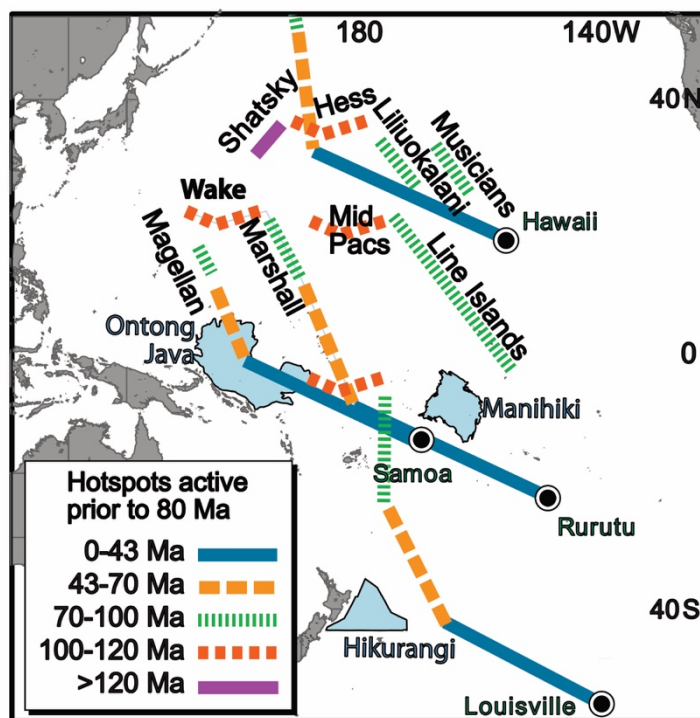
904 JK and VAF contributed equally to this study. Conceptualization: JK, VAF, MGJ, AAPK. Field
905 expedition and sampling: CK, JK. Sample preparation and data collection: AA, JK, MB.
906 Modeling: JK, PW, AAPK. Writing, editing, and figures (original draft): All authors. Writing,
907 editing, modeling, and figures (revised version): VAF, KK, MGJ, AAPK.

908 **EXTENDED DATA**

909 **Extended Data Figures 1-8**

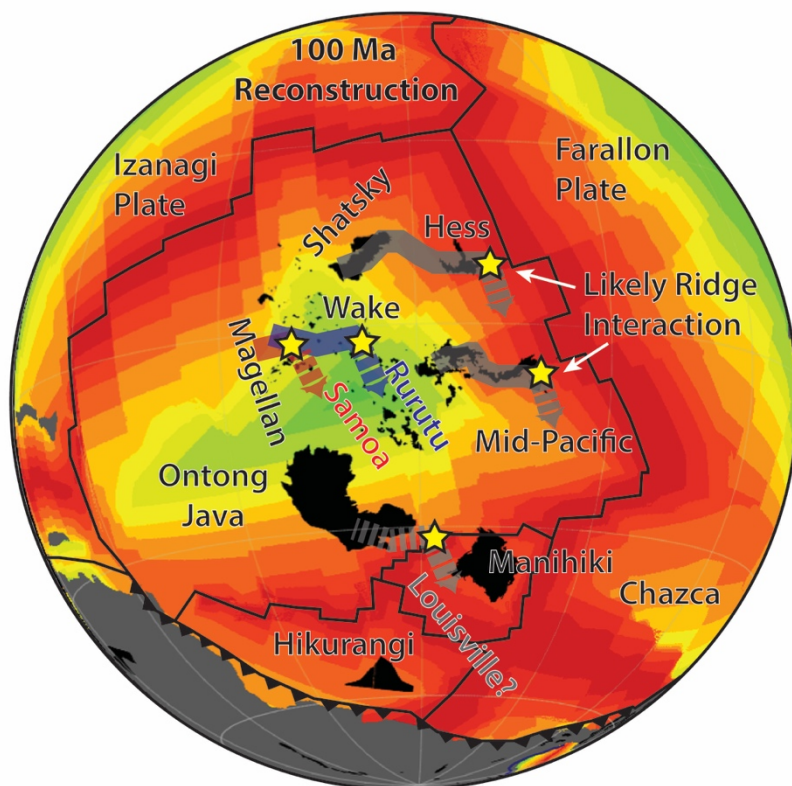
910 **Extended Data Tables 1-3**

This is a preprint of an article that has been submitted to Nature, and revised based on one round of peer review, but has yet to be formally accepted for publication.

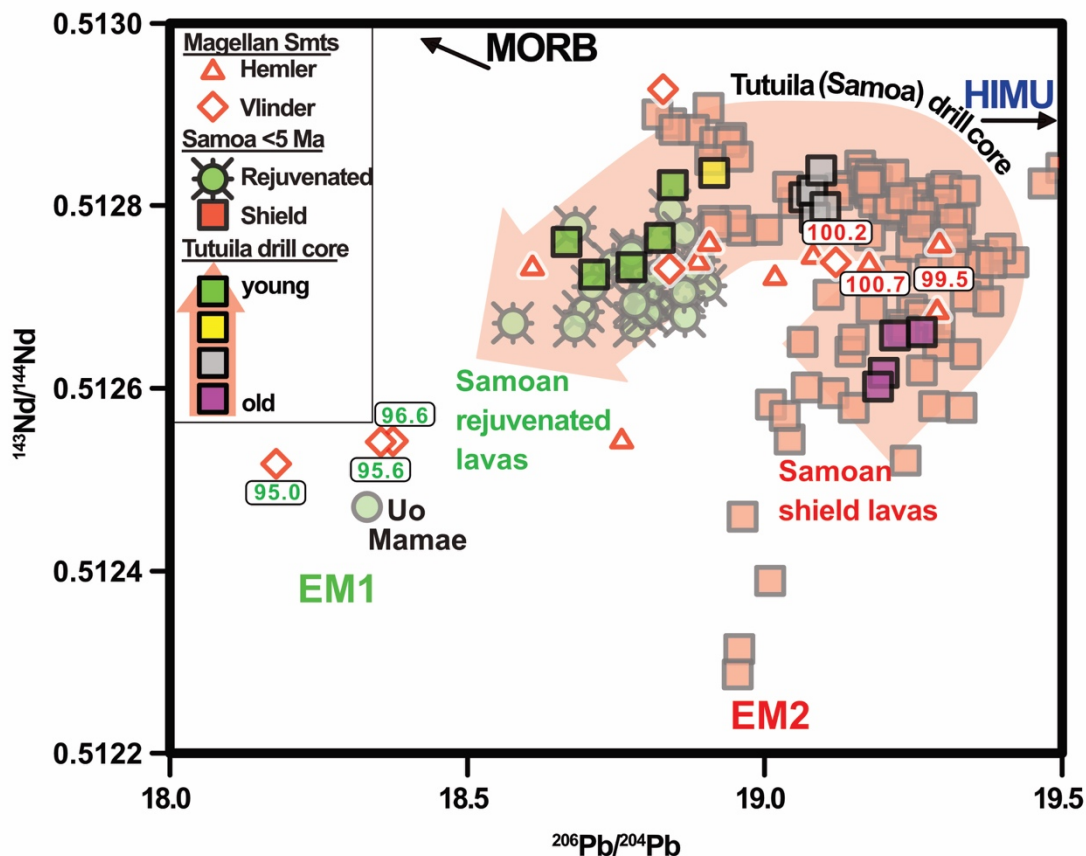


Extended Data Figure 1. Idealized map of the Pacific Ocean basin, showing relevant hotspot tracks, anchored at present-day locations indicated with black dots. Different sections of the hotspot tracks are color-coded by approximate range in eruption ages¹⁸. In the West Pacific, many sections are represented by seamount groups known by their own names. In light blue, the three oceanic plateaus are shown that are thought to have made up Ontong-Java Nui together.

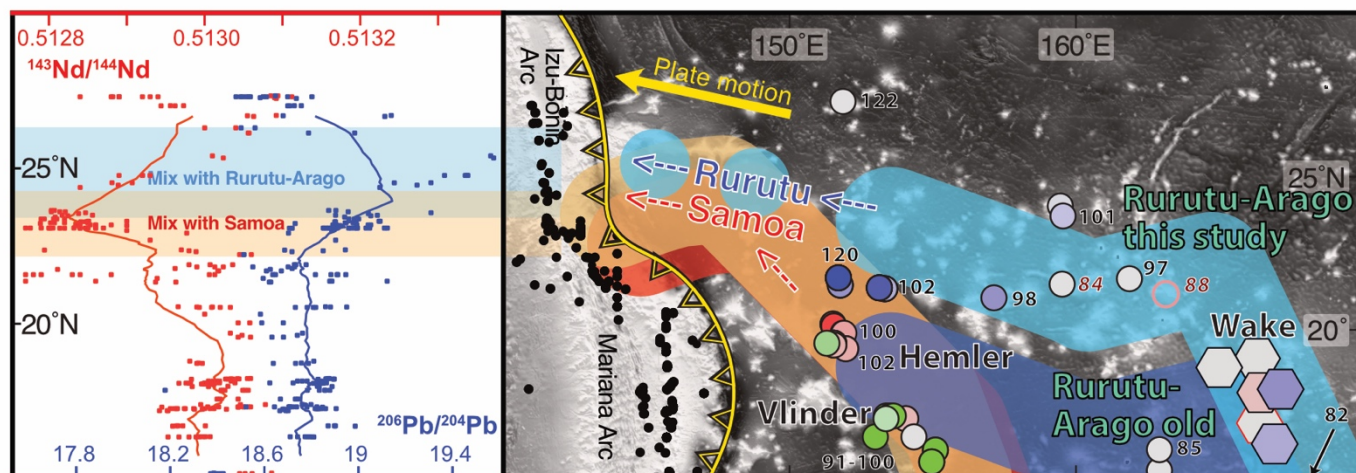
This is a preprint of an article that has been submitted to Nature, and revised based on one round of peer review, but has yet to be formally accepted for publication.



Extended Data Figure 2. 100 Ma plate configuration and relative position of volcanic structures relevant to APM modeling (global projection from GPlates¹⁰¹). Shatsky Rise and Mid-Pacific Mountains erupted near a spreading center (likely causing plume motion⁵), while Rurutu-Arago and Samoa erupted within the growing Pacific plate. Louisville's position shows the approximate modeled track for the updated model rotation.

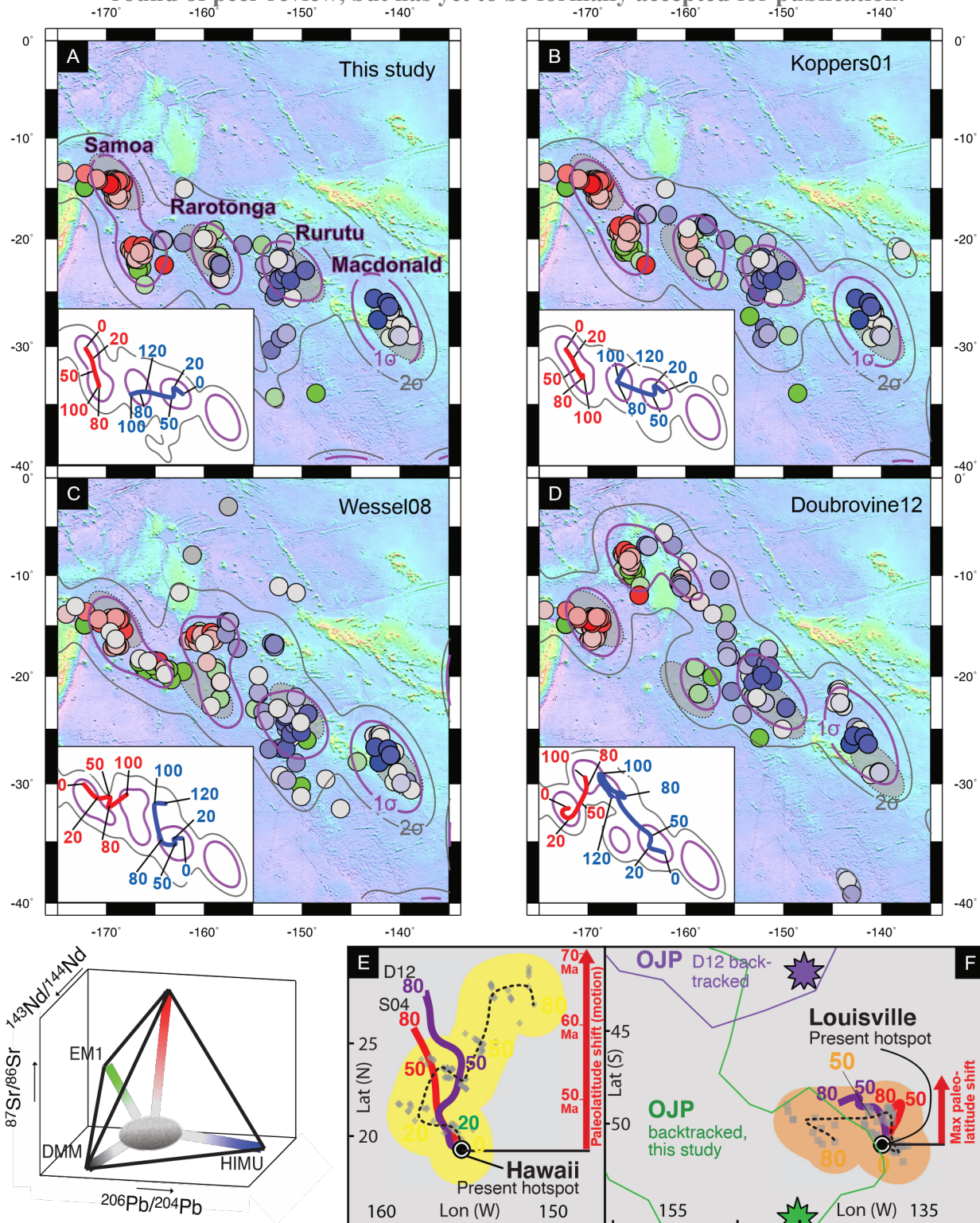


Extended Data Figure 3. Compositional overlap between modern Samoa (< 5 Ma) and Cretaceous Samoan volcanoes—Hemler and Vlinder seamounts—in the West Pacific. Most of the geochemical evolution of Samoan volcanoes through time is reflected in a drill core into Tutuila Island (Samoa⁴⁷). This shows (red arrow) a change in lava compositions from shield to rejuvenated lavas, with volcanoes active over the past 5 Ma⁷². The older samples^{29,59} (ages shown in white text bubbles) from Hemler and Vlinder (~100 Ma) mainly plot around the shield lavas in Samoa while the youngest samples (~95 Ma) continue through the rejuvenated lavas represented by the most extreme EM-1 type composition in the Samoan area (Uo Mamae¹⁰²).



Extended Data Figure 4. Map of West Pacific seamounts, showing Rurutu-Arago and Samoa predicted hotspot tracks (K01¹⁸ in dark shade, this study in light shade). Black numbers with white outline represent ages^{29,73}, while numbers in red represent likely unrelated volcanic ages, given large age difference with expected age along the hotspot track and difference in composition (pink circle²⁹). New Pb isotope data (hexagons) provides the “missing link” that suggests that Rurutu hotspot continues through the seamounts west of Wake Island. The samples outlined with red has $^{143}\text{Nd}/^{144}\text{Nd}$ affected by phosphatization. The existing (dark blue¹⁸) APM model track for the Rurutu-Arago hotspot is devoid of major seamounts, while the new track (light blue) continues the unusual isotopic composition and morphological chain to the Izu-Bonin-Mariana trench. A similar prediction for Samoa (red: K01¹⁸; orange: updated) shows both hotspots have a corresponding unusual spike in isotopic compositions in the arc (left panel; lines represent running means), indicating prior subduction of a continuing chain of similar composition, mixing Rurutu-Arago (HIMU) or Samoa (EM2) hotspot material into the mantle wedge.

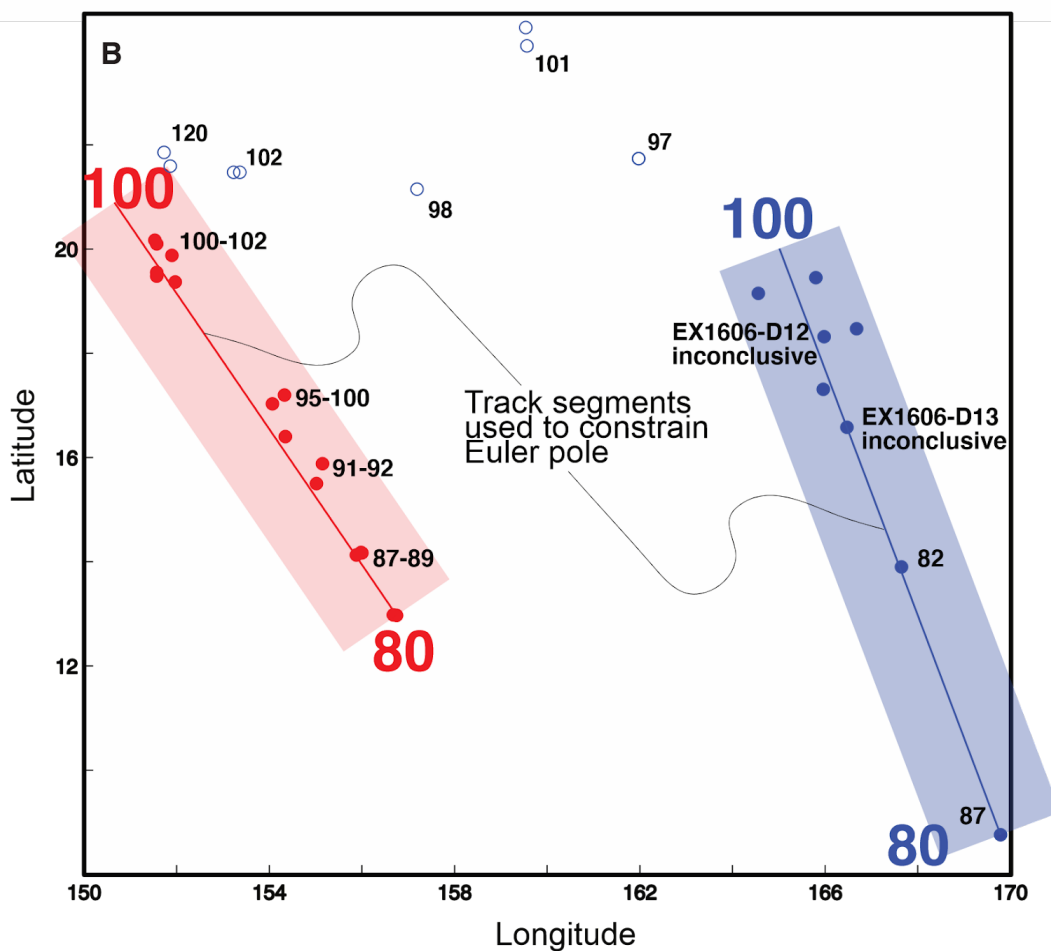
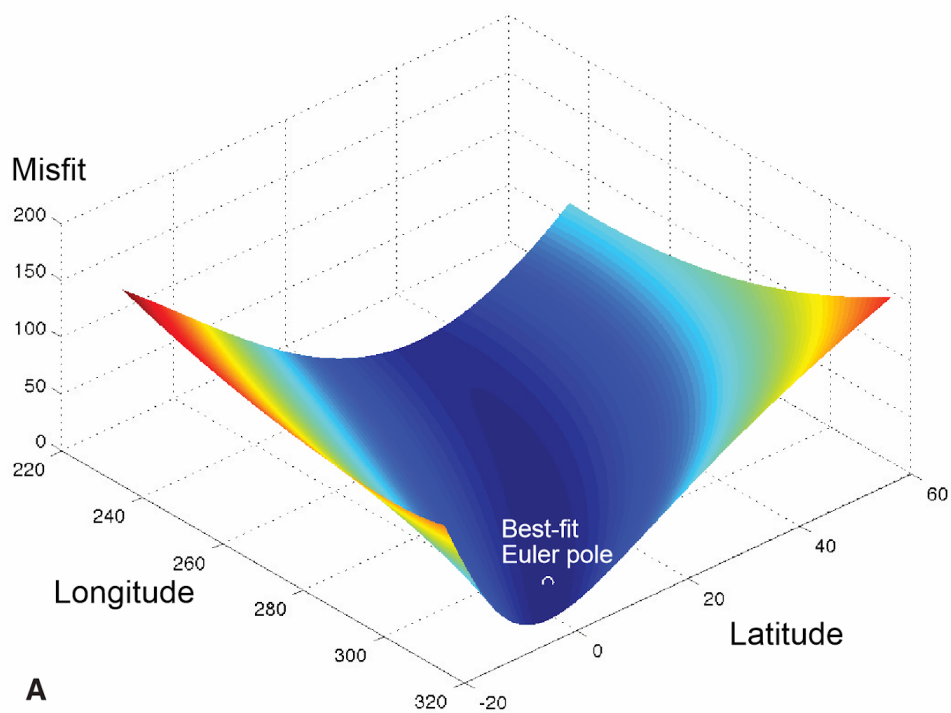
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Extended Data Figure 5. Backtracked original eruptive locations for Samoa and Cook-Austral-related volcanoes (Cook-Austral, Samoa, Tuvalu, Gilbert, Wake, Magellan, Tokelau), using various absolute plate motion models (a. this study; b. Koppers01¹⁸; c. Wessel08; d. Dobrovine12⁶). Backtracked seamounts color-coded (lower left panel) for their isotopic compositions show that at present-day

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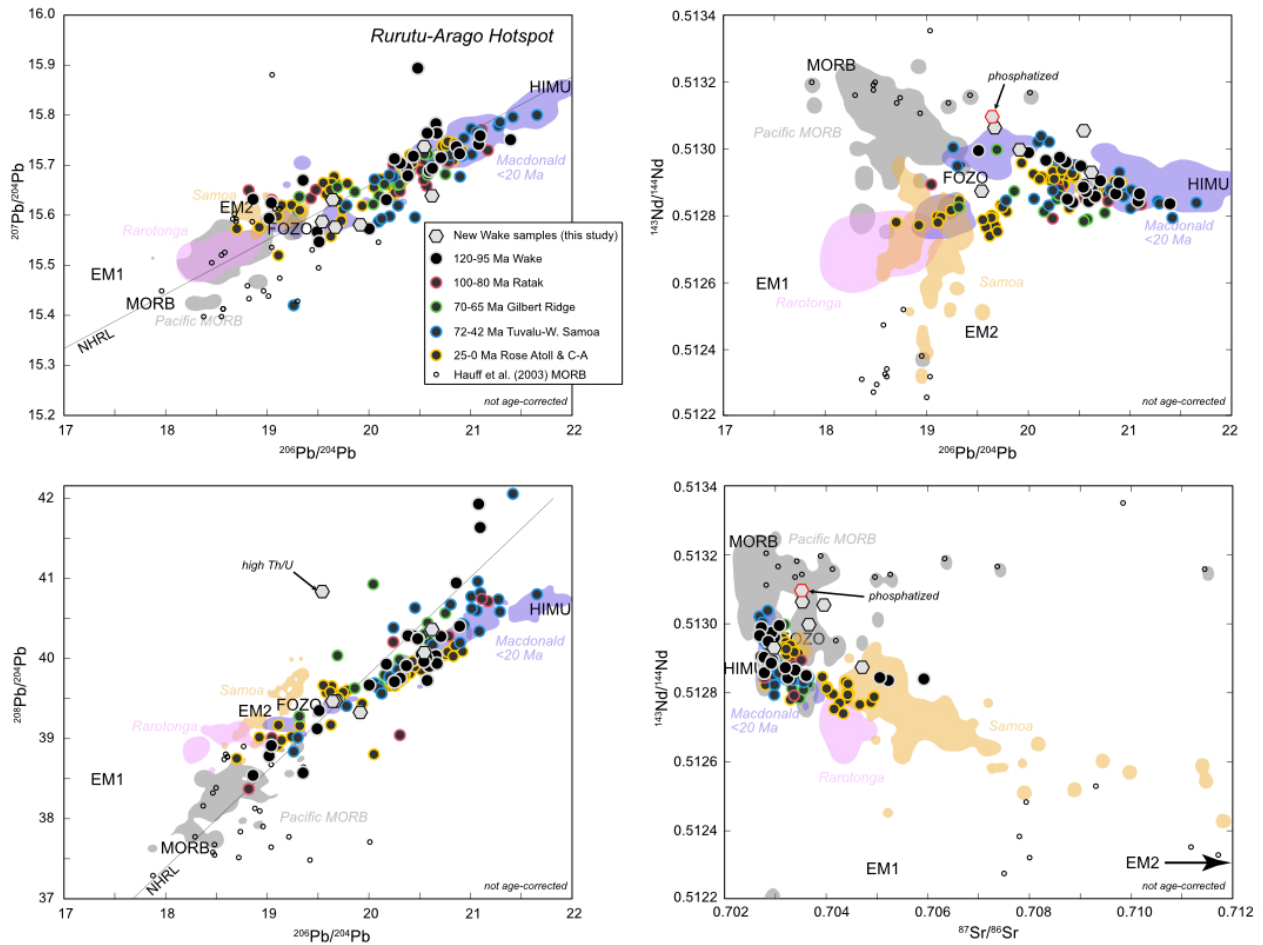
hotspots (purple text) are the focus of clusters of consistent geochemical compositions, defined by grey (2) and purple (1) contours for Gaussian-kernel probability density estimates for the backtracked locations. Insets show these density contours and the running mean of backtracked location and age to estimate plume motion from model mismatch from fixed hotspots (a-d). Backtracked (grey symbols⁶) locations for Hawaii (e) and Louisville (f) and their smoothed age-track (wide yellow or orange) highlight a deviation from predicted plume motion (S04/red⁶¹ and D12/purple⁶) beyond 50 Ma, indicating a mismatch for plate motion models into the Cretaceous. Outlines show Ontong-Java Plateau (OJP) backtracked with the new model overlap with Louisville, where stars show approximate center of Ontong-Manihiki-Hikurangi. Red arrows show smoothed tracks match derived latitudinal plume motion from paleomagnetic latitudes^{13,103}. Backtracking of OJP assumes fixed relationship to the Pacific plate, with no rotation of the plateau¹.



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Extended Data Figure 6. Rotation (Euler) pole modeling is accomplished by a grid search for the best-fit pole (5°N , 306°E), shown as the latitude-longitude location for which the minimum misfit is found (white dot). Constraints for the modeled rotation are the colored seamount locations (blue and red, selected based on their apparent fit in composition), and the approximate age range for these volcanic tracks (80 -100 Ma), based on model-specific and adjacent seamount volcanic ages^{29,59} (black numbers next to seamount markers). Only two samples from EX1606 had material suitable for age dating, but results lacked a statistically robust plateau and were therefore inconclusive. Open circles represent Wake area seamounts and their ages, predating the 80 – 100 Ma time period modeled.

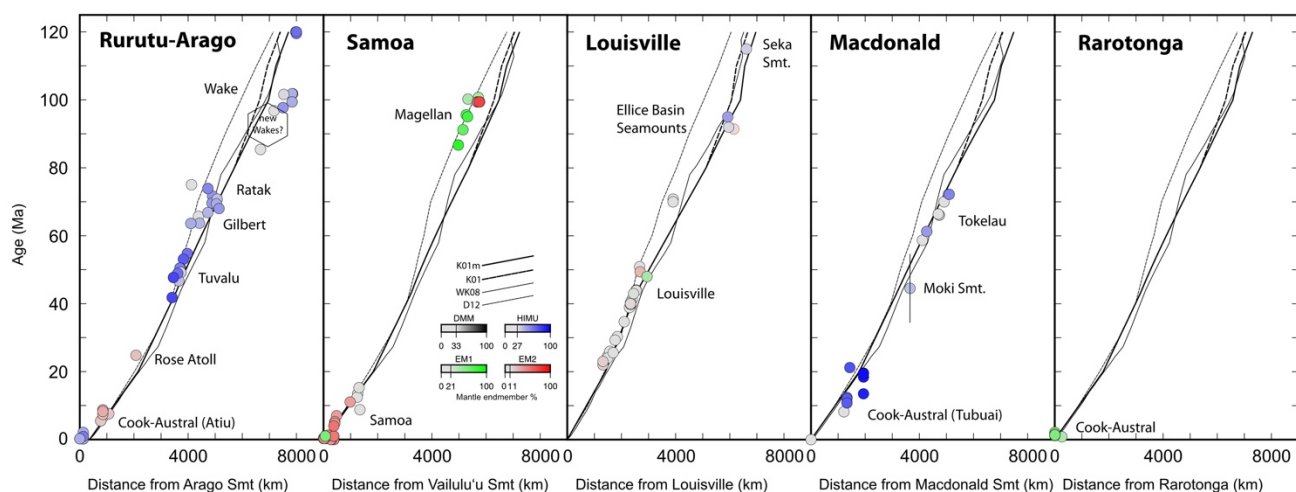
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Extended Data Figure 7. Non-age-corrected plots of **A)** $^{207}\text{Pb}/^{204}\text{Pb}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$, **B)** $^{208}\text{Pb}/^{204}\text{Pb}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$, **C)** $^{143}\text{Nd}/^{144}\text{Nd}$ vs. $^{206}\text{Pb}/^{204}\text{Pb}$, and **D)** $^{143}\text{Nd}/^{144}\text{Nd}$ vs. $^{87}\text{Sr}/^{86}\text{Sr}$ of our new Wake seamount samples (hexagons) compared to published data for the Tuvalu, Gilbert Ridge, Marshall/Ratak, and Wake Seamount portions of the Rurutu-Arago track (circles). Background data for young segments of the Cook-Austral plumes (Macdonald, Rurutu-Arago, Rarotonga), the Samoan plume, and Pacific MORB are given as 2SD contours of KDEs of their respective datasets. Jurassic Pacific MORB data with seawater U alteration⁹² is also shown as small black open circles. A 2SD contour of the KDE for published Wake seamounts is shown as an open grey contour. In both plots, the new Wake data plots within the known compositional range for the Wake Seamounts and the greater extent of the track. One of the new samples has unusually high $^{208}\text{Pb}/^{204}\text{Pb}$; this is a signature occasionally expressed in older Rurutu-Arago lavas and persists after strong leaching¹⁴. Here, $^{208}\text{Pb}/^{204}\text{Pb}$ is positively correlated with

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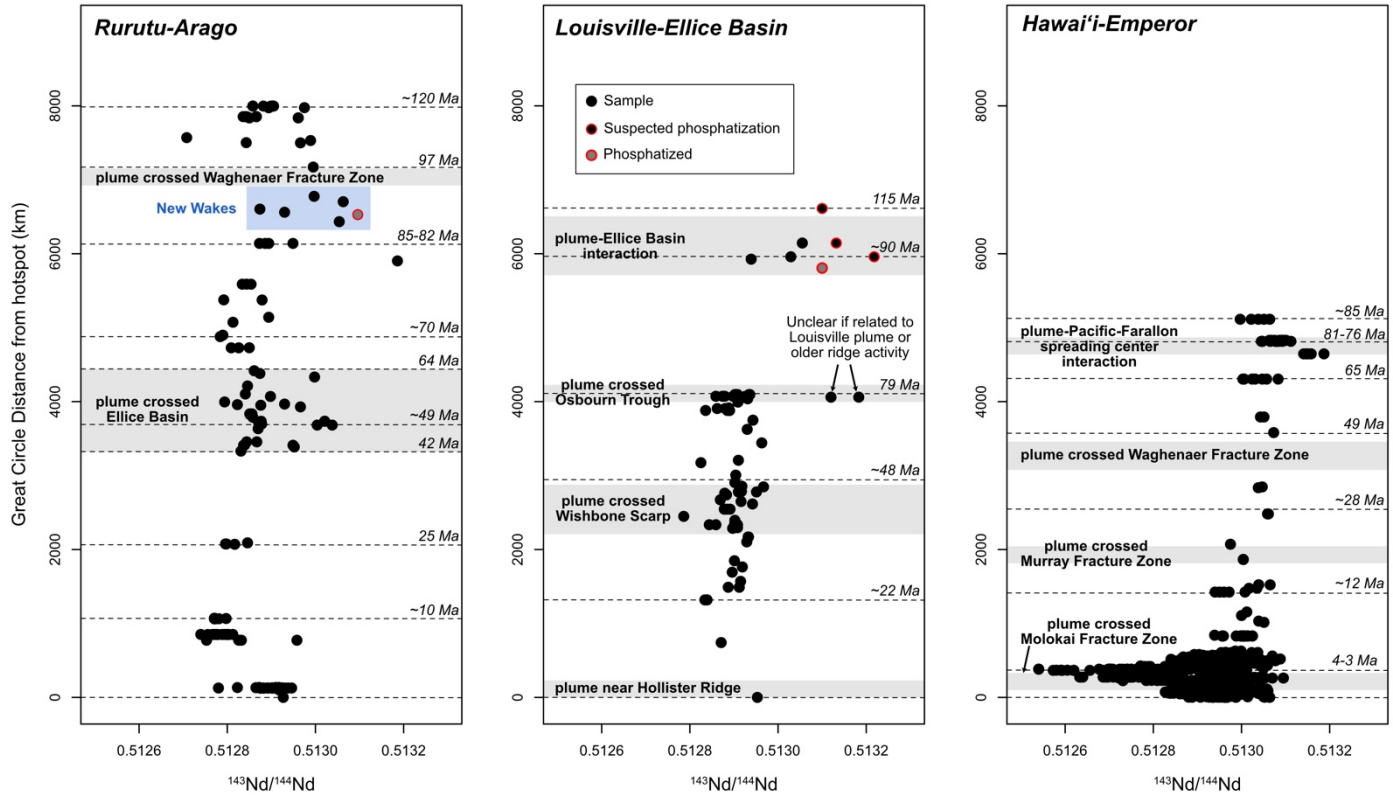
Th/U (see Extended Data Table 1), indicating *U* loss during alteration. Data are not corrected for post-eruptive radiogenic ingrowth (see Methods).



Extended data Figure 8. Age-distance relationships of the Rurutu-Arago, Samoan, Louisville, Macdonald, and Rarotonga plumes (see Background Data Sources For Figures in Methods). Data shown here are only samples with age determinations or published age estimates (where well constrained by nearby volcanoes^{14,28} and Sr-Pb-Nd data to permit color coding. APM models are also included for reference (WK08³⁹ = Wessel and Kroenke, 2008; D12⁶ = Doubrovine et al., 2012 without plume drift correction; K01¹⁸ = Koppers et al., 2001; K01m = modified Koppers et al., 2001 from this study). The oldest portions of the Hotspot Highway are in good agreement with K01m model predictions; some scatter occurs as a function of plume drift (e.g., Cretaceous portion of the Samoa hotspot; see Extended Data Figure 5). For four of the five hotspots shown here, the data are consistent with age progressions that can be traced back into the Cretaceous. Rurutu-Arago and Macdonald have HIMU to FOZO-like compositions, while Samoan volcanoes are EM-type to FOZO in composition. While the Macdonald track appears to have not existed prior to 72 Ma, the Rurutu-Arago age progression can be clearly traced into the Wakes and back to ~120 Ma. The Samoan plume was active during the Cretaceous, forming the Magellan chain in the West Pacific where EM2 and EM1 compositions consistent with those found in Samoan shield and rejuvenated volcanoes, respectively (see Extended Data Figure 3 for details). The FOZO Louisville hotspot track and older Ellice Basin

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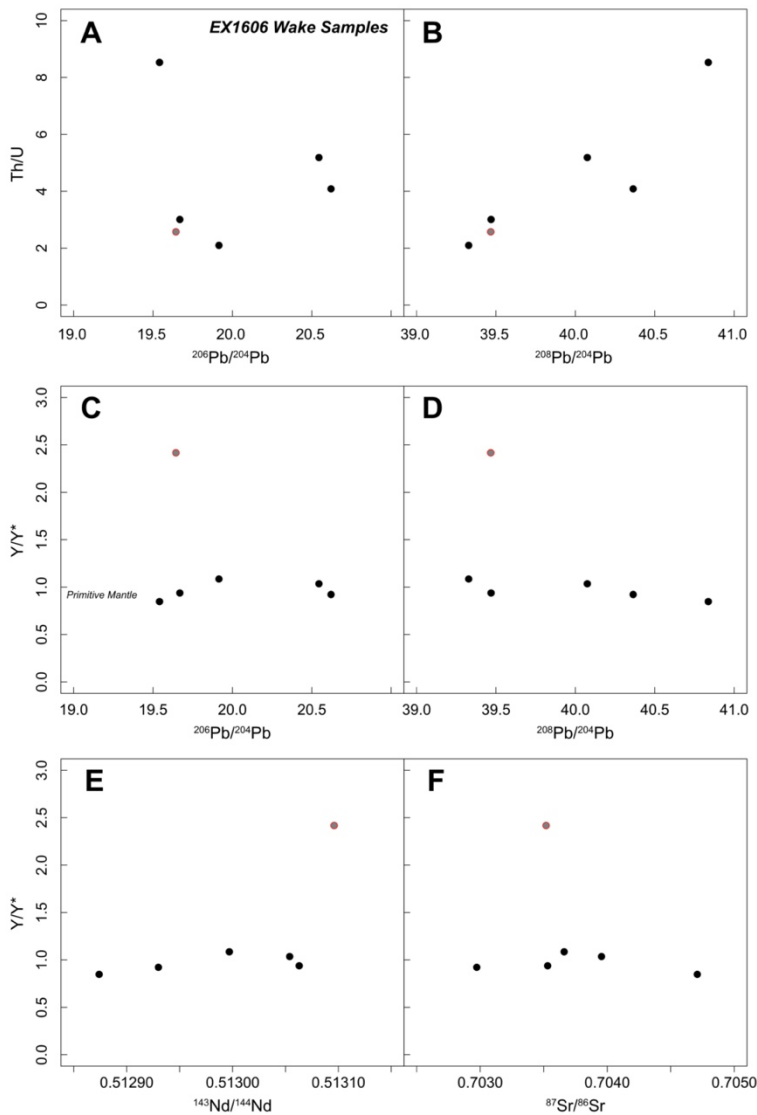
Seamounts as well as Seka Seamount; which are likely FOZO-to-DMMLike with Pb isotopes partly overprinted by seawater U ingrowth, are also age-progressive. The Rarotonga plume is only known from two young volcanoes in the Cook-Austral.



Extended Data Figure 9. Great circle distance (km) from active hotspot center vs. $^{143}\text{Nd}/^{144}\text{Nd}$ (not age-corrected) for the (left) Rurutu-Arago, (center) Louisville-Ellice Basin Seamounts, and (right) Hawaii-Emperor hotspot tracks. Isochrons are provided as dashed lines, and light grey fields mark where the plumes interacted with ridges or fracture zones. All three hotspot tracks record significant variability in isotopic composition over time that correspond to interaction of the plume with major lithospheric structural features. Rurutu-Arago, which was a true-intraplate hotspot for the entirety of its documented history, produced episodes of depleted (high $^{143}\text{Nd}/^{144}\text{Nd}$) melts that coincide spatially with major lithospheric structures, but otherwise maintains a fairly constant $^{143}\text{Nd}/^{144}\text{Nd}$ over time. Samples with evidence of phosphatization (high Y/Y^* and/or P_2O_5 ; see Extended Data Figure 10) are shown as light grey circles with red outlines. Black circles with red outlines are samples that may have been phosphatized ($^{143}\text{Nd}/^{144}\text{Nd} > \sim 0.5131$) but cannot be confirmed due to lack of available major and trace element data. By contrast, the Louisville hotspot track, including the Ellice Basin Seamounts,

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records a long-term trend of enrichment with time that records its transition from plume-ridge interaction to true intraplate. Deviations also occur when the plume crossed the Osbourn Trough (however, it remains unclear whether this is related to the Louisville plume), and later the Wishbone Scarp (attributed to source mantle heterogeneity; Beier et al., 2011; Vanderkluyesen et al., 2014). The broad enrichment trend in Louisville is similar to long-term enrichment observed in the Hawaiian plume, which also interacted with a ridge in the Cretaceous before transitioning to a true-intraplate plume system (Regelous et al., 2003). Data sources are the same as in Fig. 1.”



Extended Data Figure 10. Trace element alteration proxies versus Sr-Nd-Pb isotope compositions of the new Wakes (EX1606) samples. **A)** Th/U versus measured $^{206}\text{Pb}/^{204}\text{Pb}$, **B)** Th/U versus measured $^{208}\text{Pb}/^{204}\text{Pb}$, and **C-F)** Y/Y* versus radiogenic isotope compositions. Versus Th/U, $^{206}\text{Pb}/^{204}\text{Pb}$ (**A**) exhibits only a weak correlation with Th/U, while $^{208}\text{Pb}/^{204}\text{Pb}$ (**B**) is much more strongly correlated,

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indicating modification of U abundances in the EX1606 samples. $^{143}\text{Nd}/^{144}\text{Nd}$ (E) exhibits some correlation versus Y/Y^ , a proxy for phosphatization, becoming more radiogenic at high Y/Y^{**} , while Pb and Sr isotopes show no correlation.*

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Extended Data Table 1. Geochemical compositions for Wake area seamounts

	D5-3		D6-4		D7-1		D10-1		D12-1		D13-1		
Seamount name	McDonnell Guyot		unnamed W		Lafayette Smt		unnamed S		unnamed SW		Batfish Smt		
Latitude	19.15152		19.44748		17.30973		18.46733		18.31973		16.58102		
Longitude	164.55802		165.79677		165.96018		166.67875		165.97903		166.47195		
Majors Elements (LIBS)													
SiO ₂	53.3		54		47.2		53.8		47		58.5		
TiO ₂	3.0		2.1		3.3		2.8		3.3		1.7		
Al ₂ O ₃	16.8		16.9		17.1		16.7		15.5		18.8		
Fe ₂ O ₃	8.4		8.8		11.7		8.9		12.4		4.5		
MgO	3.5		3.8		4.7		3.6		6.2		2		
CaO	9.6		8.2		12.8		8.4		11.9		8.3		
Na ₂ O	3.5		4.2		2.5		3.7		2.7		4		
K ₂ O	1.9		2.1		0.6		2.2		1.1		2.1		
Total	100		100.1		99.9		100.1		100.1		99.9		
Trace Elements (LA-ICP-MS)													
	units		RSD		RSD		RSD		RSD		RSD		
Y	ppm (ug/g)	20.84	1.5	12.95	1.3	118.3	3.4	34.23	1.6	29.69	1.8	24.52	1.4
Zr	ppm (ug/g)	176.3	1.9	127.9	1.5	195.9	2.7	513.7	0.9	276.4	1.6	209.4	1.2
Yb	ppm (ug/g)	1.455	1.4	0.881	3.0	4.456	3.7	2.529	1.0	1.728	2.3	2.374	2.0
W	ppm (ug/g)	0.436	4.0	0.417	2.9	0.795	3.6	0.555	2.5	0.698	3.2	0.418	1.4
Rb	ppm (ug/g)	54.09	2.4	44.97	1.2	49.33	2.8	29.32	2.3	14.97	1.7	62.52	1.1
Sr	ppm (ug/g)	300.6	2.1	295.6	2.0	413.5	3.3	1134	1.5	182.3	1.4	310.0	1.0
Nb	ppm (ug/g)	33.06	1.4	19.21	2.2	40.52	3.2	150.3	1.0	77.61	1.3	72.21	1.2
Mo	ppm (ug/g)	0.911	2.5	1.611	2.0	2.029	3.6	2.521	1.9	2.791	0.9	0.463	1.1
Mo	ppm (ug/g)	0.951	1.5	1.625	3.4	2.031	3.0	2.558	1.4	2.854	1.3	0.473	1.9
Cd	ppm (ug/g)	0.118	4.5	0.117	9.1	0.140	2.9	0.280	3.3	0.236	4.6	0.097	4.4
Sn	ppm (ug/g)	1.656	2.2	1.127	3.3	1.677	3.5	2.546	2.4	2.323	1.3	1.900	0.8
Sb	ppm (ug/g)	3.512	1.5	2.940	2.2	7.086	3.4	1.434	1.0	2.646	1.2	0.583	2.0
Cs	ppm (ug/g)	2.364	1.8	1.287	2.0	2.012	4.5	0.634	0.6	0.404	2.2	0.533	1.0
Ba	ppm (ug/g)	125.8	1.7	129.3	3.1	248.5	3.7	574.7	0.9	198.7	2.2	675.0	1.3
La	ppm (ug/g)	16.71	1.7	11.34	2.3	90.09	3.2	84.21	1.0	39.78	1.9	27.51	1.7
Ce	ppm (ug/g)	31.35	1.6	24.89	1.8	43.30	2.2	168.6	1.1	87.34	2.1	50.58	1.5

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Pr	ppm (ug/g)	4.348	2.3	3.135	1.7	10.64	3.1	18.16	1.2	11.37	1.7	5.732	2.0
Nd	ppm (ug/g)	18.10	1.8	13.25	1.8	42.58	4.4	66.45	1.5	48.76	3.1	21.24	2.2
Sm	ppm (ug/g)	3.948	1.5	3.022	2.3	7.121	3.8	11.24	1.2	10.74	1.6	4.167	1.6
Eu	ppm (ug/g)	1.429	1.5	1.171	2.3	2.363	3.7	3.546	1.8	3.181	1.8	1.933	1.7
Gd	ppm (ug/g)	4.070	1.6	3.084	1.2	9.366	3.8	10.62	1.6	10.23	2.0	4.352	1.6
Tb	ppm (ug/g)	0.590	2.0	0.442	2.5	1.233	4.2	1.331	1.3	1.336	1.8	0.650	2.2
Dy	ppm (ug/g)	3.316	2.3	2.441	1.7	7.653	4.6	6.641	1.1	6.560	1.5	3.951	1.7
Ho	ppm (ug/g)	0.649	1.7	0.456	2.0	1.829	3.9	1.212	0.8	1.092	1.5	0.829	1.4
Er	ppm (ug/g)	1.722	1.7	1.132	2.6	5.372	4.1	3.081	1.2	2.494	1.9	2.403	1.5
Tm	ppm (ug/g)	0.237	2.0	0.149	2.0	0.728	4.2	0.414	1.4	0.300	2.0	0.362	1.7
Lu	ppm (ug/g)	0.212	1.9	0.119	1.1	0.728	4.0	0.351	1.9	0.228	1.6	0.364	1.1
Hf	ppm (ug/g)	4.221	2.1	3.111	1.8	4.652	4.9	9.905	0.9	6.337	1.7	4.449	1.6
Ta	ppm (ug/g)	2.068	2.3	1.124	1.5	2.346	3.2	8.855	1.5	4.404	1.6	4.013	0.6
Pb	ppm (ug/g)	1.623	1.9	1.043	2.1	4.602	4.0	6.936	1.3	3.604	2.1	2.440	1.5
Bi	ppm (ug/g)	0.020	1.9	0.012	11	0.020	4.8	0.048	2.9	0.025	5.1	0.019	2.2
Th	ppm (ug/g)	1.787	1.3	1.122	1.6	2.067	4.1	11.07	1.2	5.851	1.9	3.578	1.5
U	ppm (ug/g)	0.852	1.4	0.373	1.9	0.803	5.1	2.711	1.7	0.686	2.1	0.690	1.4
MgO	wt %	1.835	1.6	1.241	1.6	1.287	0.9	2.965	2.4	6.997	1.5	0.504	3.3
Al ₂ O ₃	wt %	13.15	1.7	14.05	1.1	12.74	1.8	17.60	2.5	11.59	1.7	18.36	3.0
CaO	wt %	3.920	1.6	4.939	1.8	6.371	1.3	5.124	2.3	11.02	1.6	4.456	3.5
Sc	ppm (ug/g)	18.18	1.4	21.09	2.0	18.38	1.4	7.480	1.1	23.54	1.4	4.682	3.1
TiO ₂	wt %	3.112	1.3	2.105	2.5	2.658	1.3	2.968	1.6	3.212	1.4	1.417	3.2
V	ppm (ug/g)	124.7	2.3	117.9	1.8	200.5	0.9	140.7	1.7	277.9	1.9	18.16	3.4
Cr	ppm (ug/g)	41.44	3.1	308.4	1.7	386.6	0.8	7.815	4.1	696.1	2.2	1.194	10.3
MnO	wt %	0.066	1.4	0.138	1.1	0.092	0.8	0.094	0.9	0.118	2.3	0.068	3.1
Fe ₂ O ₃	wt %	9.893	2.1	11.18	1.8	11.42	1.8	8.456	1.3	13.57	2.7	4.479	2.4
Co	ppm (ug/g)	22.28	1.5	56.30	1.6	28.56	2.0	21.62	1.2	35.65	2.3	5.295	3.2
Ni	ppm (ug/g)	41.08	1.2	86.20	2.0	60.54	1.7	22.76	2.5	177.9	1.8	4.571	4.8
Cu	ppm (ug/g)	114.3	1.2	82.69	1.2	168.6	0.9	30.54	2.1	64.58	2.0	14.07	4.6
Zn	ppm (ug/g)	191.2	1.6	158.1	2.0	175.9	2.0	288.0	2.1	189.2	1.2	90.73	3.8
Th/U		2.10		3.00		2.57		4.09		8.53		5.19	
Y/Y*		1.09		0.94		2.42		0.92		0.85		1.04	
Isotope Data (MC-ICP-MS)			2SE		2SE		2SE		2SE		2SE		2SE
²⁰⁶ Pb/ ²⁰⁴ Pb		19.915	4E-04	19.669	5E-04	19.644	6E-04	20.621	3E-04	19.541	1E-03	20.545	4E-04
²⁰⁷ Pb/ ²⁰⁴ Pb		15.581	3E-04	15.576	5E-04	15.631	6E-04	15.639	3E-04	15.587	1E-03	15.737	4E-04
²⁰⁸ Pb/ ²⁰⁴ Pb		39.328	9E-04	39.469	1E-03	39.467	2E-03	40.364	9E-04	40.837	2E-03	40.075	1E-03
¹⁴³ Nd/ ¹⁴⁴ Nd		0.512997	7E-06	0.513063	7E-06	0.513096	4E-06	0.512930	3E-06	0.512874	5E-06	0.513054	6E-06

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$^{87}\text{Sr}/^{86}\text{Sr}$	0.703661	7E-06	0.703531	6E-06	0.703519	7E-06	0.702973	6E-06	0.704709	7E-06	0.703955	7E-06
$^{176}\text{Hf}/^{177}\text{Hf}$	0.282956	3E-06	0.283030	5E-06	0.282948	3E-06	0.282974	3E-06	0.282929	3E-06	0.282931	3E-06
Initial Ratios (100 Ma)												
$^{206}\text{Pb}/^{204}\text{Pb}_i$	19.333		19.272		19.451		20.188		19.330		20.232	
$^{207}\text{Pb}/^{204}\text{Pb}_i$	15.553		15.557		15.622		15.618		15.577		15.722	
$^{208}\text{Pb}/^{204}\text{Pb}_i$	38.938		39.088		39.308		39.798		40.262		39.556	
$^{143}\text{Nd}/^{144}\text{Nd}_i$	0.51291		0.51297		0.51303		0.51286		0.51278		0.51297	
$^{87}\text{Sr}/^{86}\text{Sr}_i$	0.70294		0.70292		0.70304		0.70287		0.70438		0.70315	
$^{176}\text{Hf}/^{177}\text{Hf}_i$	0.28294		0.28302		0.28291		0.28296		0.28292		0.28291	
Calculated P/D Ratios												
$^{238}\text{U}/^{204}\text{Pb}$	37.221		25.399		12.376		27.713		13.488		20.041	
$^{235}\text{U}/^{204}\text{Pb}$	0.270		0.184		0.090		0.201		0.098		0.145	
$^{232}\text{Th}/^{204}\text{Pb}$	78.632		76.837		32.077		114.039		115.953		104.740	
$^{147}\text{Sm}/^{144}\text{Nd}$	0.1373		0.1437		0.1053		0.1065		0.1387		0.1236	
$^{87}\text{Rb}/^{86}\text{Sr}$	0.51		0.43		0.34		0.07		0.23		0.57	
$^{176}\text{Lu}/^{177}\text{Hf}$	0.00700		0.00535		0.02181		0.00494		0.00501		0.01139	
Forward-modeled Ratios												
$^{206}\text{Pb}/^{204}\text{Pb}_f$	19.780		19.719		19.898		20.635		19.777		20.679	
$^{207}\text{Pb}/^{204}\text{Pb}_f$	15.575		15.579		15.644		15.641		15.599		15.744	
$^{208}\text{Pb}/^{204}\text{Pb}_f$	39.516		39.666		39.886		40.377		40.840		40.134	
$^{143}\text{Nd}/^{144}\text{Nd}_f$	0.51305		0.51311		0.51317		0.51300		0.51293		0.51312	
$^{87}\text{Sr}/^{86}\text{Sr}_f$	0.70306		0.70304		0.70316		0.70299		0.70450		0.70327	
$^{176}\text{Hf}/^{177}\text{Hf}_f$	0.28300		0.28308		0.28297		0.28302		0.28298		0.28297	

Major elements collected by laser induced breakdown spectroscopy, using a Nd-YAG 20 mJ pulsed laser and a Catalina Scientific EMU-120 echelle spectrometer. Major element compositions were calibrated against 36 different standards, using partial least squares (PLS) regression, in Matlab. Predicted compositions of unknown samples using PLS show typically <2 wt.% variability on repeat analyses, after each sample is re-normalized to 100% total values. Trace element data were obtained using a ThermoFinnegan Element2 ICP-MS (University of South Carolina). The sample powders were dissolved in Teflon-distilled HF:HNO₃ mixture, subsequently dissolved in 2 wt.% HNO₃, and spiked with In at 2 ppb concentration in the solution. The USGS reference material BHVO-2 was used as a standard and the reference materials BCR-2 and JB-2 were run as unknowns along the samples. The calculated concentrations for the unknowns agree well within 5% for most elements relative to the recommended concentrations from GEOREM

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(accessed, August 2022¹⁰⁴). Pb isotope compositions were acquired with a Nu Plasma HR MC-ICP-MS (University of Hawai‘i at Mānoa), using sample standard bracketing, and Tl doping to monitor fractionation. Pb was separated and purified with Sr resin and AG1 resin⁵³. Repeatability of Pb isotope analysis of a given sample is typically about +/- 1e-3. Compositions are normalized to values¹⁰⁵ for NIST 981 Pb. The Sr and Nd fractions were further processed at the University of South Carolina. Sr samples were purified with Sr-Spec resin in HNO₃. For Nd, the REE fractions were separated from major element matrix via TRU-spec resin in HNO₃ and HCl media, and Nd separated from the other REE on an LN-resin using 0.25N HCl. Hf was isolated from the matrix on LN resin from the fractions recovered from the washes of the TRU spec column¹⁰⁶. Sr, Nd and Hf isotope ratios were acquired using a ThermoFinnegan Neptune multicollector. Repeated analyses of SRM987 dispersed with the samples gave $^{87}\text{Sr}/^{86}\text{Sr} = 0.710321 \pm 0.000007$, (2 stdev, n=11). The data are reported relative to the recommended SRM987 $^{87}\text{Sr}/^{86}\text{Sr}_{\text{SRM987}} = 0.71025$. Nd isotopes were corrected for fractionation using $^{146}\text{Nd}/^{144}\text{Nd} = 0.7219$ and repeated analyses of the JNdi-1 reference material yielded an average $^{143}\text{Nd}/^{144}\text{Nd} = 0.512102 \pm 0.000005$ (2 σ , n=11). The Nd data are reported relative to the accepted value for JNdi-1 of $^{143}\text{Nd}/^{144}\text{Nd} = 0.512115$. Hf isotopes were corrected for fractionation using $^{179}\text{Hf}/^{177}\text{Hf} = 0.7325$. An in-house Hf standard solution was determined at $^{176}\text{Hf}/^{177}\text{Hf} = 0.282142 \pm 0.000006$ (2 σ , n=8), which corresponds to the original JMC 475 solution value of $^{176}\text{Hf}/^{177}\text{Hf} = 0.282163$. The data is reported relative to the accepted JMC 475 value of $^{176}\text{Hf}/^{177}\text{Hf} 0.282160$ ¹⁰⁷. BCR-2 duplicates are duplicate analyses of the same digestion.

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Extended Data Table 2. Stage poles for the Modified Pacific Hotspot Reference Frame Absolute Plate Motion Model (italics from K01¹⁸)

Stage start (Ma)	Stage end (Ma)	Latitude (°N)	Longitude (°E)	Rotation Rate (°/Myr)	Standard Deviation
<i>0</i>	<i>20</i>	<i>70.1</i>	<i>302.0</i>	<i>0.88</i>	<i>0.016</i>
<i>20</i>	<i>43</i>	<i>67.1</i>	<i>294.5</i>	<i>0.58</i>	<i>Not given</i>
<i>43</i>	<i>80</i>	<i>18.8</i>	<i>253.6</i>	<i>0.66</i>	<i>0.029</i>
<i>80</i>	<i>100</i>	<i>5.0</i>	<i>306.0</i>	<i>0.85</i>	<i>0.13</i>
<i>100</i>	<i>110</i>	<i>75.1</i>	<i>44.8</i>	<i>0.44</i>	<i>Not given</i>
<i>110</i>	<i>125</i>	<i>65.3</i>	<i>273.2</i>	<i>0.45</i>	<i>0.042</i>

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Extended Data Table 3. Locations used to constrain the revised 100-80 Ma stage pole.

	Latitude (°N)	Longitude (°E)
Magellan Seamounts:		
1	20.17	151.53
2	19.48	151.57
3	19.55	151.57
4	20.1	151.57
5	19.88	151.9
6	19.37	151.97
7	17.03	154.07
8	17.2	154.33
9	16.4	154.35
10	15.5	155.02
11	15.88	155.15
12	14.13	155.88
13	14.18	155.98
14	14.17	156
15	12.98	156.68
16	12.97	156.75
Marshall Seamounts:		
17	19.15	164.56
18	19.45	165.8
19	17.31	165.96
20	18.32	165.98
21	16.58	166.47
22	18.47	166.68
23	13.9	167.65
24	8.76	169.79

Extended Data Table 4. Standard data.

	Quality Control			duplicate	duplicate	Quality Control		
	BCR-2	n=2				BCR-2	BCR-2	JB-2
Trace Elements (LA-ICP-MS)	[Lit]	[Calculated]	%Recovery			[Lit]	[Calculated]	%Recovery
Y	36.07	36.57	101%			23.56	25.12	107%
Zr	186.5	190.1	102%			48.25	46.69	97%
Yb	3.392	3.269	96%			2.529	2.481	98%
W	0.465	0.507	109%			0.308	0.309	100%
Rb	46.02	50.22	109%			6.4	7.065	110%
Sr	337.4	349.5	104%			175.2	192.5	110%
Nb	12.44	12.36	99%			0.565	0.477	84%
Mo	250.6	251.9	101%			1.014	0.960	95%
Mo	250.6	253.3	101%			1.014	0.981	97%
Cd	0.69	0.620	90%			0.3	0.114	38%
Sn	2.28	2.153	94%			0.635	0.615	97%
Sb	0.302	0.256	85%			0.261	0.209	80%
Cs	1.16	1.134	98%			0.8	0.868	108%
Ba	683.9	683.6	100%			218.1	218.7	100%
La	25.08	25.04	100%			2.281	2.254	99%
Ce	53.12	53.04	100%			6.552	6.354	97%
Pr	6.827	6.854	100%			1.129	1.138	101%
Nd	28.26	28.00	99%			6.392	6.074	95%
Sm	6.547	6.379	97%			2.266	2.177	96%
Eu	1.989	1.998	100%			0.836	0.825	99%
Gd	6.811	6.781	100%			3.123	2.886	92%
Tb	1.077	1.044	97%			0.586	0.554	95%
Dy	6.424	6.328	99%			3.868	3.923	101%
Ho	1.313	1.287	98%			0.863	0.860	100%
Er	3.67	3.560	97%			2.537	2.517	99%
Tm	0.534	0.513	96%			0.393	0.376	96%
Lu	0.505	0.481	95%			0.389	0.372	96%
Hf	4.972	4.788	96%			1.487	1.396	94%
Ta	0.785	0.755	96%			0.040	0.037	94%
Pb	10.59	10.49	99%			5.25	5.01	95%
Bi	0.05	0.06	116%			0.031	0.03	99%
Th	5.828	5.76	99%			0.258	0.25	96%
U	1.683	1.56	93%			0.153	0.14	89%
MgO	3.599	3.82	106%			4.43	5.58	126%
Al ₂ O ₃	13.48	13.98	104%			14.62	16.30	111%
CaO	7.114	7.59	107%			9.852	10.74	109%
Sc	33.53	35.52	106%			54.08	62.56	116%
TiO ₂	2.265	2.34	103%			1.167	1.17	100%
V	417.6	436.3	104%			572.4	607.9	106%
Cr	15.85	15.54	98%			26.65	24.85	93%
MnO	0.200	0.21	103%			0.213	0.23	109%
Fe ₂ O ₃	13.77	14.24	103%			14.28	14.91	104%
Co	37.33	39.30	105%			37.57	38.20	102%
Ni	12.57	12.07	96%			14.77	13.52	92%
Cu	19.66	18.08	92%			222.1	236.2	106%
Zn	129.5	149.76	116%			110.4	122.2	111%

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Isotope Data (MC-ICP-MS)

²⁰⁶Pb/²⁰⁴Pb	18.758	3E-04				
²⁰⁷Pb/²⁰⁴Pb	15.618	3E-04				
²⁰⁸Pb/²⁰⁴Pb	38.725	7E-04				
¹⁴³Nd/¹⁴⁴Nd	0.512643	6E-06	0.512646	6 E-06	0.512643	5E-06
⁸⁷Sr/⁸⁶Sr	0.705014	6E-06	0.705005	7E-06	0.705010	9E-06
¹⁷⁶Hf/¹⁷⁷Hf	0.282868	5-06	0.282869	7E-06	0.282870	4E-06