

Data report: strontium isotope analyses of pore fluids from the CRISP-A transect drilled during Expeditions 334 and 344¹

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Abstract

We report on the Sr isotopic composition of pore fluids recovered during Integrated Ocean Drilling Program Costa Rica Seismogenesis Project Expeditions 334 and 344. Pore fluid samples were acidified and loaded directly onto columns containing EICHRON Sr-Spec resin, followed by analyses using a NU multicollector inductively coupled plasma–mass spectrometer at the Keck Collaboratory, Oregon State University.

Sites drilled on the upper plate exhibited a decrease in the $^{87}\text{Sr}/^{86}\text{Sr}$ ratio to values as low as 0.70762 in the upper 200–300 m of the sediment column, suggesting the role of ash alteration. The unconformity between the slope sediment sequence and underlying framework rock, sampled at Sites U1378–U1380, is characterized by intense fracturing. The fluids sampled within the unconformity at these sites have $^{87}\text{Sr}/^{86}\text{Sr}$ ratios ranging from 0.70834 to 0.70844.

Fluids sampled from the incoming plate sediment show evidence of ash alteration and carbonate diagenesis, but the deepest sediment here may also be modified by diffusion of fluids from the underlying oceanic crust. Intense carbonate recrystallization in the lower 66 m of Site U1414 results in Sr concentrations that are almost an order of magnitude higher than those measured at the base of Site U1381. The deepest pore fluid sample measured at both Sites U1381 and U1414 has a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.7086.

The strontium systematics in this margin, used in the context of other ongoing studies, may be used to unravel processes at this margin, such as carbonate diagenesis, ash alteration, fluid migration, and diffusive exchange with the underlying oceanic crust.

Introduction

The Costa Rica Seismogenesis Project (CRISP) was designed to understand the processes that control fault zone behavior during earthquake nucleation and rupture propagation at erosional subduction zones. The CRISP study area, located offshore the Osa Peninsula of Costa Rica, is part of the active and long-lived subduction erosion from Guatemala to Costa Rica (Ranero et al., 2000a, 2000b; Vannucchi et al., 2004). This area is characterized by low sediment supply, fast convergence rate, abundant plate interface seismicity, and change in subducting plate relief along

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strike (Expedition 334 Scientists, 2012a; see also the “[Expedition 344 summary](#)” chapter [Harris et al., 2013a]). Arcward of the trench, the lower slope consists of a 10–12 km wide frontal prism, where a modern sediment apron overlies older sediment that may have been deposited in a fore-arc basin setting (A. Sakaguchi, pers. comm., 2013). The first phase of the CRISP focuses on sampling sediment, fluid, and crustal rock because fluid and associated diagenetic reactions affect hydrological parameters (e.g., permeability and pore pressure) and may regulate the mechanical state of the plate interface at depth. To this aim, seven sites were drilled on this margin during Integrated Ocean Drilling Program (IODP) Expeditions 334 and 344, which targeted sites on the upper and incoming plates (Fig. F1).

Because the isotopic composition of strontium does not undergo biological fractionation (Mook, 2001), it has proven to be valuable in establishing fluid-rock reactions, sources, and fluid mixing (e.g., Teichert et al., 2005; Torres et al., 2004; Solomon et al., 2009; Joseph et al., 2012). Potential source materials that may alter a de facto seawater $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of interstitial fluids are continental detritus ($^{87}\text{Sr}/^{86}\text{Sr} = \sim 0.7119\text{--}0.7133$), biogenic calcite ($^{87}\text{Sr}/^{86}\text{Sr} = \sim 0.7075\text{--}0.7092$), volcanic ash ($^{87}\text{Sr}/^{86}\text{Sr} = \sim 0.706\text{--}0.704$), and oceanic crust ($^{87}\text{Sr}/^{86}\text{Sr} = \sim 0.703$) (Veizer, 1989). Here we report on pore water strontium isotopic composition of subsamples collected from the seven sites drilled during Expeditions 334 and 344.

Study sites

Five sites were drilled on the overriding plate and sampled the upper and middle slope regions and the prism toe. Two additional sites were drilled in the incoming Cocos plate (Fig. F1). Complete descriptions of the drilling results are given in Expedition 334 Scientists (2012a) and in the “[Expedition 344 summary](#)” chapter (Harris et al., 2013a).

The shallowest site along the CRISP transect, Site U1379, was drilled into the upper slope of the Costa Rica margin, 34 km offshore the Osa Peninsula. This site is thought to overlie the locked portion of the subduction zone seismogenic zone in an area where the plate boundary is 4.5 km below seafloor. Sediment from Site U1379 is divided into five lithostratigraphic units (Fig. F2A).

Site U1378 was drilled into the middle slope of the Costa Rica margin, 41 km offshore the Osa Peninsula (Fig. F1). This site is located above the unlocked portion of the plate boundary, as indicated by interplate earthquake relocation and geodetic measurements (LaFemina et al., 2009). The margin here consists of

an upper plate framework wedge underlying ~550 m of slope sediment. Drilling at this site penetrated a landward-dipping reflector interpreted as a normal fault cutting through the whole upper plate, which may be directly linked with the plate boundary. The sediment at Site U1378 is dominantly composed of a monotonous sequence of silty clay to clay that alternates with widely interspersed centimeter-scale sandy layers, and was divided into three main lithostratigraphic units (Fig. F2).

Site U1380 is a complementary site to nearby Site U1378 where only the upper part (~400 [meters below seafloor [mbsf]] of the slope sediment was drilled during Expedition 334. Site U1380 was drilled to investigate the deeper portions of the upper slope sequence and underlying wedge sediment. The site was cored between ~395 and 480 mbsf during Expedition 334 but was abandoned before science goals were achieved because of hole instability. Hole U1380C was cored during Expedition 344. The sedimentary succession recovered from the framework wedge revealed an alternating terrestrially sourced, turbiditic upper slope (Units I and III) to shelf (Unit II) sequence, eventually being influenced by deltaic-derived sediment (Fig. F2).

Farther north, Site U1413 was also drilled in the middle slope region within a 3-D seismic volume collected in 2013 (Bangs et al., 2013). Three lithostratigraphic units (Fig. F3) can be distinguished in the sediment of Site U1413. The frontal sedimentary prism at the base of the slope was drilled at Site U1412 (Fig. F1). The primary goal of this site was to penetrate the décollement and investigate the fluid flow regime within the sediment as well as the oceanic crust. Three units were distinguished in the sedimentary rock (Fig. F3). Unfortunately, due to hole instability, the décollement and underthrust sediment were not sampled at Site U1412.

To understand seismogenic processes in convergent margins we need to fully characterize the sediment and oceanic crust entering the seismogenic zone. To this aim, two sites were drilled on the subducting aseismic Cocos Ridge. Incoming sediment thickness at Site U1414 is 380 m, and a much thinner section (<100 m) was recovered at Site U1381 located higher on Cocos Ridge. The upper 50 m of Site U1381 is composed of a predominantly monotonous sequence of silty clay to clay (Unit I), which is underlain by a more pelagic sequence (Unit II) characterized by abundant biogenic components (Fig. F4). The contact between the basement and the overlying sediment was recovered at ~95 mbsf in Hole U1381B and at ~104 mbsf in Hole U1381C.

Analytical methods

Pore fluid was collected from whole-round cores that were cut on the catwalk immediately after recovery, capped, and taken to the laboratory for processing using a titanium squeezer, modified after the stainless-steel squeezer of Manheim and Sayles (1974). Gauge pressures up to 30 MPa were applied using a laboratory hydraulic press to extract pore water. Pore fluid was passed through a prewashed Whatman No. 1 filter fitted above a titanium screen, filtered through a 0.2 μm Gelman polysulfone disposable filter, and subsequently extruded into a precleaned (10% HCl), 60 mL plastic syringe attached to the bottom of the squeezer assembly. Details of this procedure are given Expedition 334 Scientists (2012b) and in the “**Methods**” chapter (Harris et al., 2013d).

Strontium concentrations in pore fluids collected during Expedition 334 were measured postcruise by inductively coupled plasma–optical emission spectrometry (ICP-OES) at Oregon State University, as reported by Torres et al. (2014). During Expedition 344, pore fluid Sr concentrations were measured on board by inductively coupled plasma–atomic emission spectroscopy (ICP-AES), and the data are reported in the “**Expedition 344 summary**” chapter (Harris et al., 2013a).

High-precision chloride concentrations were acquired on board during both Expeditions 334 and 344 using a Metrohm 785 DMP autotitrator and silver nitrate (AgNO_3). International Association for the Physical Sciences of the Oceans seawater was used as the standard (Expedition 334 Scientists, 2012b; see also the “**Methods**” chapter [Harris et al., 2013d]).

During Expedition 334, sulfate data were obtained using the shipboard ICP-AES, and the data are reported by Vannucchi, Ujiie, Stroncik, and the Expedition 334 Scientists (2012). During Expedition 344, sulfate concentrations were determined with a Metrohm 861.004 advanced compact ion chromatograph (IC). Details of the procedure and results are given in the “**Methods**” chapter (Harris et al., 2013d).

For analysis of the $^{87}\text{Sr}/^{86}\text{Sr}$ ratios, aliquots of acidified pore fluid samples containing ~ 300 ng Sr were directly loaded onto chromatographic columns. Samples were collected in acid-washed Nalgene bottles and acidified with ultrapure HNO_3 at a ratio of 20 μL acid per milliliter of sample. Strontium separation was carried out using 50 μL Sr-Spec resin from EICHROM. Isotopic analysis was performed using the NU multicollector inductively coupled plasma mass spectrometer (MC-ICPMS) housed in the W.M.

Keck Collaboratory for Plasma Spectrometry in the College of Earth, Ocean and Atmospheric Science at Oregon State University (USA). Instrument mass bias was corrected using an $^{88}\text{Sr}/^{86}\text{Sr}$ ratio of 8.375209, and $^{87}\text{Sr}/^{86}\text{Sr}$ ratio data were normalized to the NBS 987 standard, with a reported $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.71025, with an internal error of ± 0.00005 ($2\sigma_{\text{mean}}$; $n = 91$). Replicate analysis of an in house standard yielded a $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.70819 ± 0.00006 ($2\sigma_{\text{mean}}$; $n = 79$), representing our external error.

Results and discussion

A total of 187 pore fluid samples were analyzed, and the data are listed in Table T1. Figures F2, F3, and F4 show downhole Sr concentration and $^{87}\text{Sr}/^{86}\text{Sr}$ ratio depth profiles, in the context of the corresponding lithology and supporting pore fluid data.

Overriding plate

The upper sediment sections cored at all sites drilled on the upper plate show a significant decrease in the Sr isotopic ratios consistent with in situ alteration of tephra.

Site U1379

At Site U1379, ash alteration is more pronounced within the sediment of lithostratigraphic Subunit IIC and corresponds to an increase in strontium concentrations (Fig. F2). Release of nonradiogenic Sr during alteration of ash is well documented, and this reaction explains the decrease in Sr isotopic values at this site from a near-seawater ratio of 0.70908 to a minimum value of 0.70757 at 369 mbsf. The isotopic ratios within the remaining fluids of Unit II range from 0.70757 to 0.70829. Unit III is dominated by coarser grained sediment as well as several fault zones below ~ 600 mbsf that host freshened fluids, as exhibited by the broad interval of low Cl, which decreases sharply from ~ 560 mbsf and reaches its lowest concentration at ~ 680 mbsf (Expedition 334 Scientists, 2012c; Torres et al., 2013). Concomitant with the Cl decrease, dissolved Sr increases from ~ 41 μM at 560 mbsf to a broad maximum of ~ 70 μM , in spite of the overall dilution trend evidenced in the Cl profile. In this zone, Sr isotopes have a less radiogenic component.

Site U1378/U1380

At Site U1378, drilled in the middle slope, Cl shows a more pronounced decrease relative to Site U1379. Site U1380 is a continuation of the sequence drilled at nearby Site U1378, and pore fluid data obtained

here show discrete anomalies that mark the unconformity separating the slope apron from the underlying wedge material (Torres et al., 2013) (Fig. F2). The Sr isotopic data in fluid from Site U1378 show a steady decrease from seawater value, which points to alteration of tephra throughout Unit II. This process is more pronounced within the sediment interval from 360 to 425 mbsf, where $^{87}\text{Sr}/^{86}\text{Sr}$ ratios show a broad minimum ranging from 0.70802 to 0.70791. Immediately below the zone displaying this broad minimum, $^{87}\text{Sr}/^{86}\text{Sr}$ ratios increase rapidly in samples collected just above the framework sediment, from 450 to 500 mbsf. The Sr isotopic composition in this zone ranges from 0.70834 to 0.70844, values that are similar to those measured within the unconformity drilled at Site U1379. The Sr concentration within the unconformity at Site U1380 is lower than that measured at Site U1379, reflecting the larger degree of dilution by fluid freshening in the middle slope region. Below the unconformity, Sr isotopic values show a steady increase with depth to a value of 0.70868 (at 552 mbsf), accompanied by a general increase in dissolved Sr concentration.

Site U1413

At Site U1413, Ca and Mg concentrations decrease from seawater values at the seafloor to a minimum of 1.1 and 39.8 mM, respectively, at the Unit I/II boundary (see the “[Upper slope Site U1413](#)” chapter [Harris et al., 2013e]), possibly reflecting precipitation of authigenic carbonates, an observation that is consistent with the sharp decrease in dissolved Sr concentration in these shallow sequences (Fig. F3). Strontium, however, is not fractionated during carbonate precipitation, and therefore the observed steady decrease in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from a near-seawater value of 0.70916 in the shallowest sample analyzed (4.4 mbsf) to 0.70899 at 24.4 mbsf indicates a component of ash alteration. Tephra were commonly observed in Unit I to a depth of ~35 mbsf. In Unit II, both the Sr concentration and its isotopic composition remain relatively constant in the depth interval from 24.4 to 135 mbsf, below which there is a marked decrease in $^{87}\text{Sr}/^{86}\text{Sr}$ ratio, which sharply declines from 0.70890 to 0.70824 at 178 mbsf. There is only a single, thin tephra layer reported in sediment from 40 to 140 mbsf, but increased occurrence was noted from 135 to 180 mbsf. These observations suggest that the alteration of reactive ash is controlling the Sr isotopic composition. Although not apparent in lithology, the Cl and Sr profiles display a different behavior below 200 mbsf. Cl shows a steady decrease, whereas Sr remains relatively constant. $^{87}\text{Sr}/^{86}\text{Sr}$ ratios also remain fairly constant to the bottom of the site at 578 mbsf.

Site U1412

At Site U1412, drilled on the prism toe, borehole instability precluded sampling the décollement and recovering the underthrust sediment and igneous basement. The geochemistry of pore fluids largely indicates in situ diagenetic reactions. Because of the poor sample coverage, we grouped the Sr isotope data into three regions, constrained by sample recovery. In the shallowest section, the Sr isotopic values steadily decrease with depth, from the near-seawater value of 0.70918 at 1.4 mbsf to 0.70888 at 111 mbsf, likely reflecting volcanic ash alteration. Only three samples were analyzed between 176.4 and 196 mbsf, with values ranging from 0.70843 to 0.70856, and correspond to the Unit I/II boundary. Only five samples were analyzed from the deepest sections, 328.8–369.02 mbsf, and these display the lowest measured values at this site, from 0.70887 to 0.70837.

Incoming plate

Site U1381

Site U1381 was cored during both Expeditions 334 and 344. The data collected during both expeditions show highly consistent values (Fig. F4). The shallowest sample (1.45 mbsf) analyzed at this site has an $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.70918, which is within the uncertainty of the seawater value of 0.70917. The geochemical profiles obtained shipboard at Site U1381 indicate that the deep sediment at this site are influenced by diffusional communication with a fluid in the igneous basement (Expedition 334 Scientists, 2012d; see also the “[Input Site U1381](#)” chapter [Harris et al., 2013c]). Most characteristic of this effect are sulfate profiles that show a reversal to seawater-like values at the bottom of the hole as observed here as well as in sites drilled offshore Nicoya during Ocean Drilling Program Legs 170 and 205 (Silver et al., 2000; Morris, Villinger, Klaus, et al., 2003). At Site U1381, Sr concentration increases with depth, concomitant with a decrease in the Sr isotopic values, corroborating the inference that the pore fluid within the upper basement has been modified by alteration of the basaltic crust.

Site U1414

The pore fluid composition in the uppermost 80 m of Site U1414 shows trends in alkalinity, sulfate, and ammonium characteristic of organic matter remineralization (see the “[Input Site U1414](#)” chapter [Harris et al., 2013b]). In addition, the alteration of ash layers commonly observed throughout the sediment sequence lead to the observed decrease in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios, from 0.7091 measured in the shallowest sample analyzed (0.56 mbsf) to 0.70874 at ~150 mbsf, at the

lithostratigraphic Unit I/II boundary. The dissolved Sr concentration remains relatively constant within this interval, suggesting that the main reaction involving strontium in Unit I is ash alteration, and there is very little carbonate precipitation. The large increase in Sr concentrations within Unit II, which is dominated by nannofossil-rich calcareous ooze, is the result of carbonate recrystallization, as has been observed in other pelagic carbonate-rich sediment (e.g., Site 1039 off the Nicoya Peninsula of Costa Rica; Shipboard Scientific Party, 1997). At the Unit II/III boundary (~300 mbsf) there is a sharp increase in Sr concentrations, with a maximum value of 941 μM at 327 mbsf (~10 times seawater value), suggesting more intense diagenetic reactions. This concentration change coincides with a seismic reflector and the onset of carbonate cementation (see the “**Input Site U1414**” chapter [Harris et al., 2013b]). The lowermost 66 m of the sediment column at Site U1414 (Unit III) is characterized by a sequence of lithified, calcareous, and siliceous cemented silt- and sandstone that probably lost its biogenic components as a result of diagenetic recrystallization. Fluids in this deepest sediment display the largest changes in Sr concentration as well as in its isotopic composition, which decreases from 0.70884 at 308 mbsf to 0.70861 measured in the deepest sample analyzed at ~337 mbsf. Basement was contacted at ~375 mbsf, but the deepest sediment was too cemented to obtain pore fluid samples. The $^{87}\text{Sr}/^{86}\text{Sr}$ ratio in the deepest sample analyzed is less radiogenic than the one measured at Site U1381 (0.70860). Intense carbonate recrystallization of these deepest sediments results in Sr concentrations that are almost an order of magnitude higher than those measured at the base of Site U1381.

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Figure F1. A. Location map of sites drilled during IODP CRISP program. B. Location of Sites U1378–U1381 drilled during Expedition 334 along the A-A' transect from the incoming plate to the upper slope. During Expedition 344, new holes were drilled at Sites U1380 and U1381, as well as at Sites U1412–U1414. C. Schematic transect showing locations of sites along the A-A' transect and indicating projected locations for Sites U1413 and U1414 (Vannucchi, Ujiie, Stroncik, and the Expedition 334 Scientists, 2012; see also the “[Expedition 344 summary](#)” chapter [Harris et al., 2013a]).

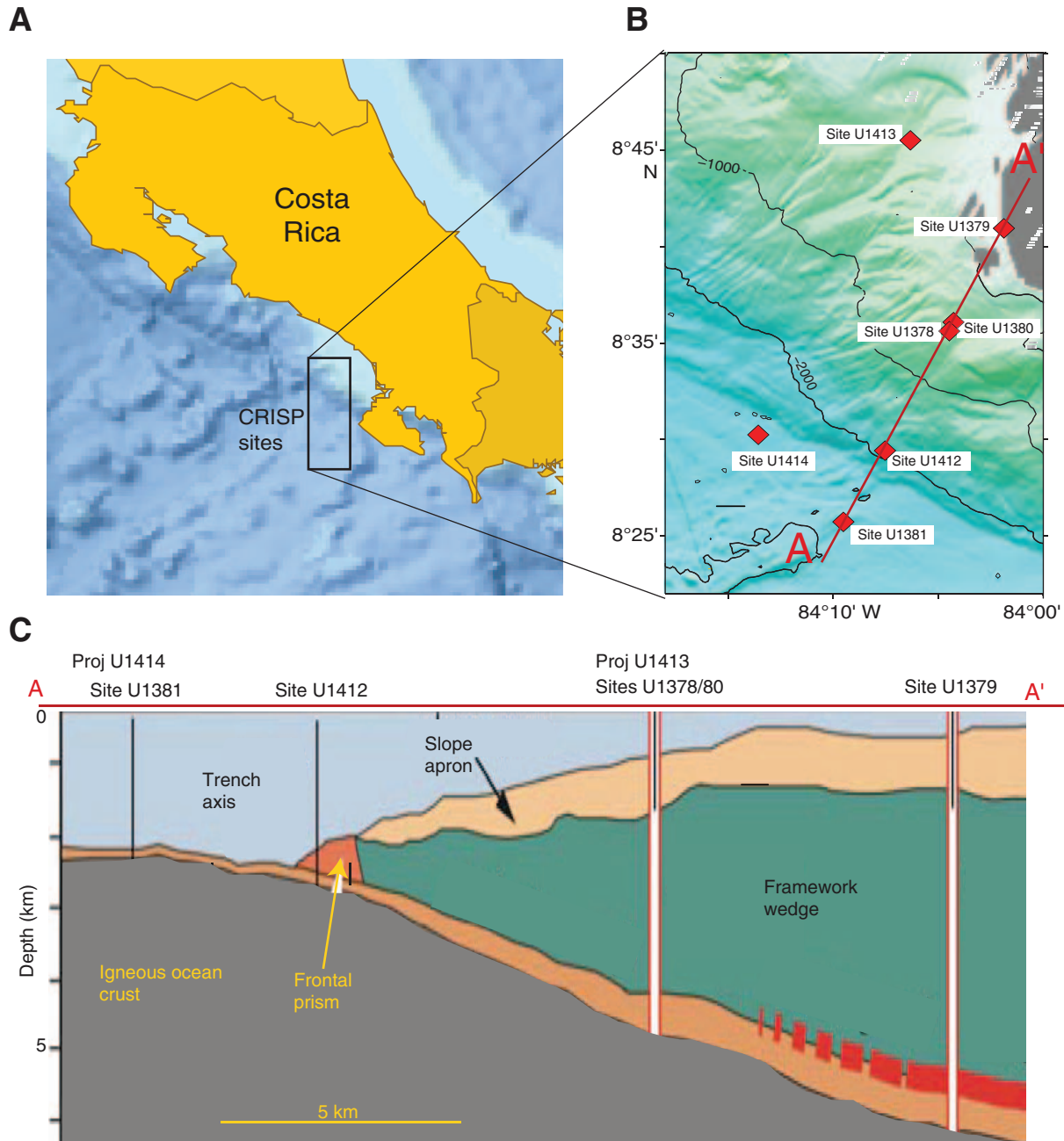


Figure F2. Downcore profiles for $^{87}\text{Sr}/^{86}\text{Sr}$, Sr, and Cl in pore fluids from (A) Site U1379 (in the upper slope) and (B) Site U1378/U1380 drilled in the middle slope. Lithostratigraphic units are from Vannucchi, Ujiie, Stroncik, and the Expedition 334 Scientists (2012) and the “Expedition 344 summary” chapter (Harris et al., 2013a). Blue circles = samples collected during Expedition 334, red circles = samples from Expedition 344. Blue arrows = seawater values, yellow zone = a highly fractured horizon that marks the unconformity between the upper slope and framework rock at Sites U1378/U1380.

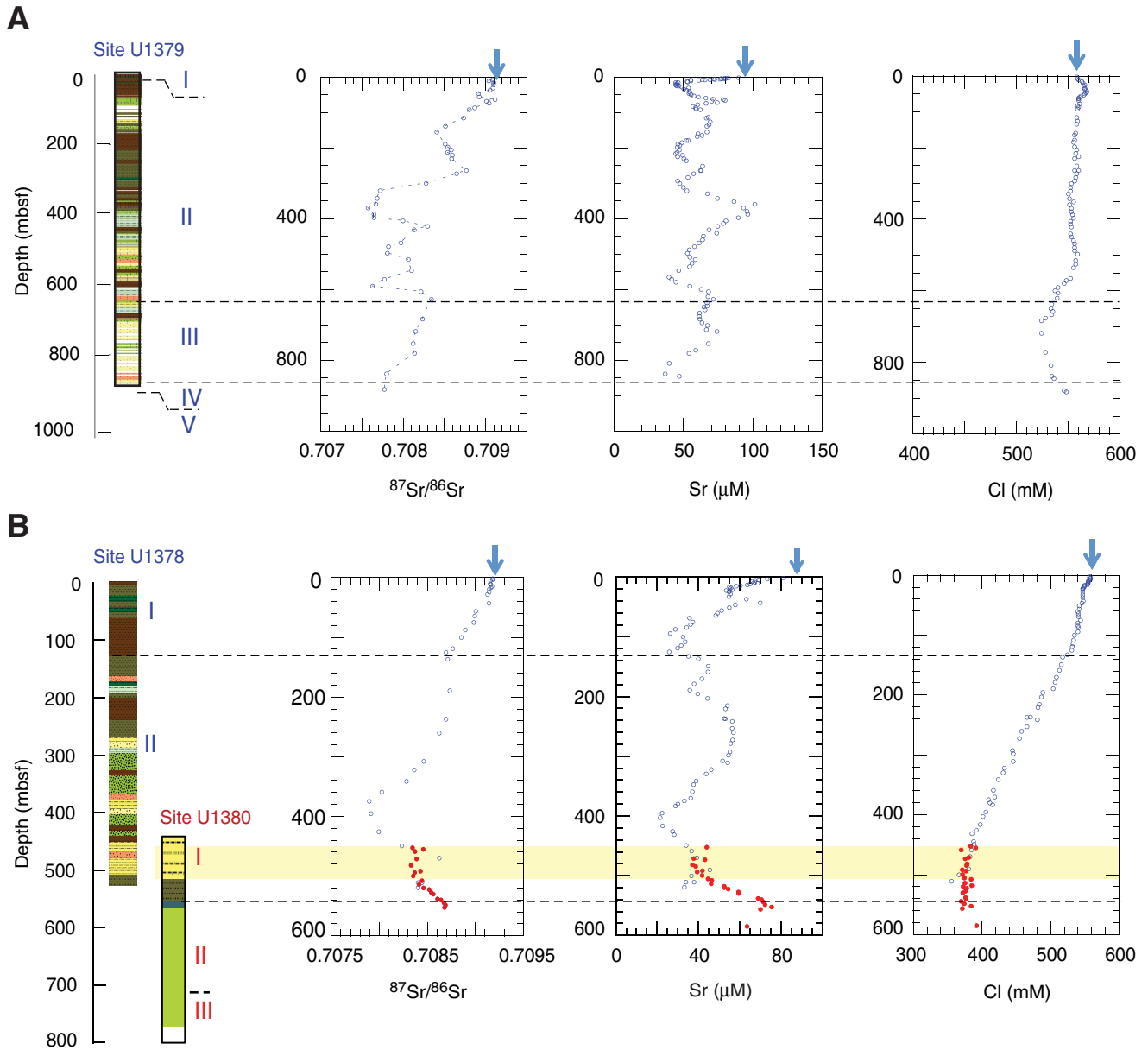


Figure F3. Downcore profiles for $^{87}\text{Sr}/^{86}\text{Sr}$, Sr, and Cl in pore fluids from Sites U1412 (at the prism toe) and U1413 (middle slope). Lithostratigraphic units are from the “[Expedition 344 summary](#)” chapter (Harris et al., 2013a). Blue arrows = seawater values.

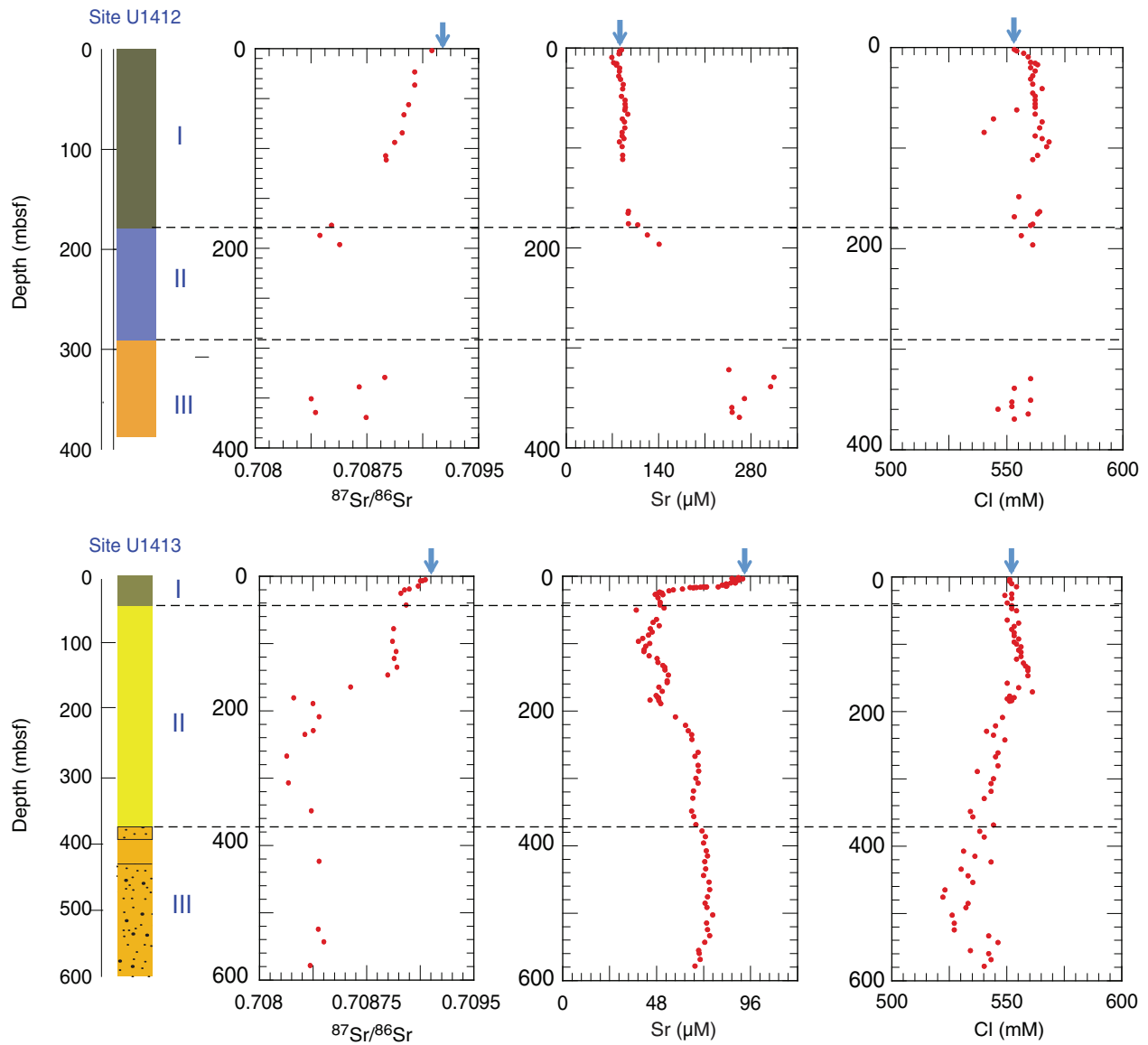


Figure F4. Downcore profiles for $^{87}\text{Sr}/^{86}\text{Sr}$, Sr, and SO_4 in pore fluids from Sites U1381 and U1414, drilled in the incoming Cocos plate. Lithostratigraphic units are from Vannucchi, Ujiie, Stroncik, and the Expedition 334 Scientists (2012) and the “**Expedition 344 summary**” chapter (Harris et al., 2013a). Blue circles = samples collected during Expedition 334, red circles = samples from Expedition 344. Blue arrows = seawater values.

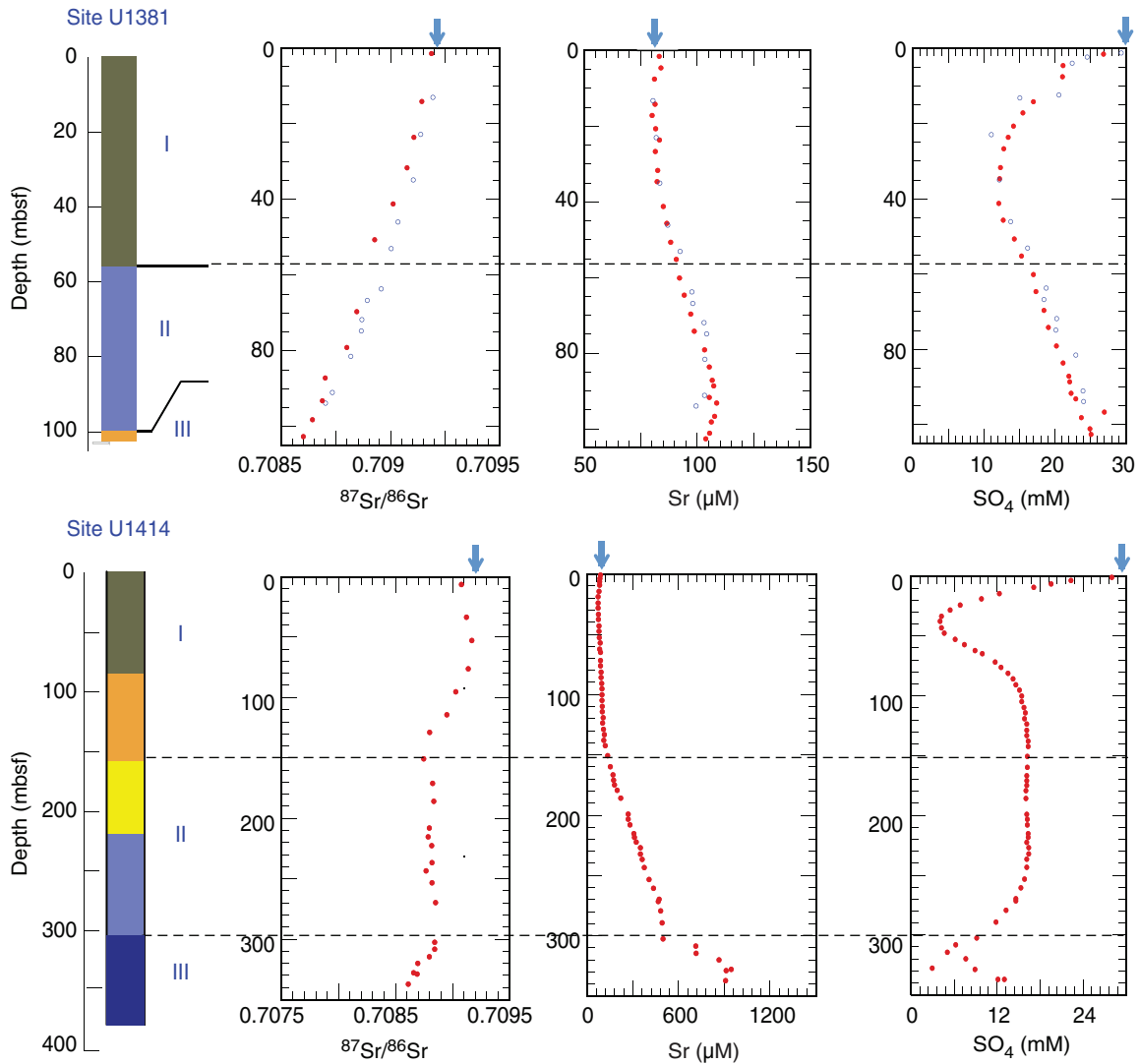


Table T1. Strontium concentrations and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in pore fluid, Expeditions 334 and 344. (Continued on next page).

Core, section, interval (cm)	Depth (mbsf)	Sr (μM)	$^{87}\text{Sr}/^{86}\text{Sr}$	Core, section, interval (cm)	Depth (mbsf)	Sr (μM)	$^{87}\text{Sr}/^{86}\text{Sr}$
334-U1378B-				50X-5, 104–132	405.3	81.8	0.70799
1H-1, 138–150	1.4	81.1	0.70918	52X-2, 118–118	420.2	76.0	0.70829
1H-2, 138–150	2.9	73.2	0.70917	53X-3, 118–118	430.5	69.7	0.70813
1H-4, 46–58	5.0	67.3	0.70916	57X-3, 118–150	467.7	62.6	0.70797
2H-3, 138–150	9.7	66.5	0.70915	58X-4, 90–90	477.7	59.2	0.70782
2H-4, 138–150	11.2	64.9	0.70916	60X-4, 118–150	496.5	54.1	0.70781
2H-6, 138–150	14.2	60.2	0.70917	62X-4, 68–100	514.6	59.8	0.70806
3H-1, 138–150	16.2	54.8	0.70916	65X-6, 78–114	545.3	47.9	0.70810
3H-3, 138–150	19.2	55.0	0.70914	68X-4, 88–120	570.5	42.9	0.70777
3H-6, 133–145	23.7	54.8	0.70914	70X-4, 92–92	589.9	55.9	0.70762
4H-3, 138–150	27.9	55.3	0.70912	72X-2, 88–120	605.4	69.4	0.70821
5H-6, 138–150	42.7	69.8	0.70914	74X-4, 74–107	626.5	72.9	0.70834
7H-2, 138–150	55.7	51.1	0.70900	81X-2, 100–132	682.2	63.1	0.70823
8H-2, 138–150	63.9	48.2	0.70899	85X-1, 77–112	717.3	75.4	0.70815
9H-2, 138–150	74.7	37.1	0.70898	89X-2, 0–40	751.8	69.2	0.70812
10H-4, 138–150	87.2	28.7	0.70889	92X-3, 94–131	779.1	55.3	0.70813
12H-3, 138–150	99.9	32.4	0.70885	98X-4, 79–115	837.4	38.1	0.70779
15H-2, 118–130	118.5	29.7	0.70876	103X-2, 34–60	881.4		0.70777
16H-3, 133–145	124.8	25.7	0.70869	344-U1380C-			
18X-3, 113–130	136.4	39.8	0.70871	3R-3, 114–144	451.9	43.8	0.70834
23X-6, 109–126	188.9	35.6	0.70873	3R-5, 112–142	454.7	—	0.70846
29X-3, 45–83	236.9	52.9	0.70869	4R-1, 51–81	458.1	—	0.70836
32X-3, 73–105	260.0	56.7	0.70862	5R-3, 107–137	470.7	37.6	0.70838
37X-2, 91–123	307.4	51.5	0.70846	6R-3, 122–153	481.3	36.9	0.70833
39X-5, 118–150	321.8	46.0	0.70836	7R-3, 120–150	491.0	41.8	0.70842
41X-5, 104–136	340.9	38.7	0.70828	7R-5, 87–107	493.4	39.2	0.70837
43X-5, 55–87	358.9	36.9	0.70802	8R-2, 116–146	499.1	41.6	0.70835
46X-2, 115–150	374.4	33.1	0.70789	9R-1, 102–132	507.2	46.3	0.70844
49X-3, 106–138	394.8	22.2	0.70791	9R-6, 79–109	513.3	46.0	0.70841
52X-5, 70–102	425.2	27.3	0.70799	10R-4, 53–78	519.5	52.4	0.70846
55X-2, 81–117	449.0	33.9	0.70823	10R-6, 64–90	521.9	54.3	0.70851
57X-3, 63–99	469.5	39.1	0.70862	11R-2, 59–89	526.9	59.2	0.70854
61X-4, 102–138	510.3	33.8	0.70844	11R-4, 87–117	529.5	59.3	0.70856
62X-1, 54–81	510.4	37.7	0.70840	12R-2, 79–113	537.6	68.7	0.70860
63X-4, 84–122	518.9	33.0	0.70840	12R-4, 78–108	539.6	70.3	0.70864
334-U1379C-				12R-8, 62–92	543.7	71.2	0.70867
1H-4, 140–140	6.0	70.4	0.70908	13R-3, 50–80	548.0	72.1	0.70868
2H-2, 140–140	9.7	56.8	0.70903	13R-6, 58–88	551.9	75.3	0.70868
2H-5, 140–140	14.2	47.6	0.70909	334-U1381A-			
3H-3, 138–138	20.6	45.6	0.70909	3R-1, 61–71	13.1	80.2	0.70919
4H-4, 92–92	29.7	54.9	0.70908	4R-1, 46–56	22.8	81.7	0.70914
5H-4, 149–149	36.2	53.6	0.70905	5R-2, 135–150	34.8	83.2	0.70910
7H-2, 138–138	45.4	55.3	0.70891	6R-3, 135–150	45.9	86.8	0.70903
8H-4, 130–130	55.0	61.8	0.70892	7R-2, 60–75	52.9	92.3	0.70900
10H-1, 88–88	62.4	79.8	0.70911	8R-2, 135–150	63.6	97.5	0.70895
11H-3, 78–78	67.3	71.7	0.70901	8R-4, 135–150	66.6	97.9	0.70889
12H-4, 103–103	73.3	67.7	0.70904	9R-1, 135–150	71.8	103	0.70887
15H-3, 100–100	86.2	59.9	0.70886	9R-3, 135–150	74.8	104	0.70886
17H-1, 76–76	91.0	63.5	0.70880	10R-1, 145–158	81.4	103	0.70882
20X-3, 137–137	115.1	67.9	0.70873	11R-1, 135–150	90.9	103	0.70873
22X-6, 138–138	138.1	68.1	0.70851	11R-3, 120–135	93.8	99.2	0.70870
24X-3, 133–133	154.3	68.2	0.70840	344-U1381C-			
27X-6, 113–130	188.0	47.3	0.70850	1H-1, 140–150	1.5	83.0	0.70919
28X-6, 133–133	196.8	47.0	0.70854	2H-4, 140–150	14.1	81.2	0.70914
29X-4, 120–137	204.6	48.3	0.70858	3H-4, 140–150	23.6	83.1	0.70910
30X-2, 133–133	211.6	46.7	0.70853	4H-3, 140–150	31.6	82.3	0.70907
31X-2, 133–133	219.4	50.6	0.70859	5H-3, 140–150	41.1	84.8	0.70901
32X-2, 133–133	229.2	51.5	0.70858	6H-3, 135–150	50.6	88.0	0.70892
33X-5, 78–100	262.6	64.1	0.70876	7H-3, 135–150	60.0	92.0	0.70862
36X-5, 128–128	271.8	58.6	0.70865	8H-3, 136–157	69.6	96.9	0.70884
39X-3, 129–129	299.3	48.7	0.70827	9H-3, 135–150	79.0	103	0.70880
41X-4, 128–128	320.4	53.7	0.70772	10H-2, 140–150	87.1	106	0.70870
43X-6, 81–107	341.5	75.8	0.70768	10H-6, 140–150	93.1	108	0.70868
45X-4, 118–140	358.5	103	0.70767	11H-3, 140–150	98.1	106	0.70864
46X-5, 75–100	368.6	94.6	0.70757	11H-6, 140–150	102.6	104	0.70860
48X-6, 63–63	386.4	97.6	0.70764				
49X-6, 75–100	396.0	90.9	0.70764				

Table T1 (continued).

Core, section, interval (cm)	Depth (mbsf)	Sr (μM)	$^{87}\text{Sr}/^{86}\text{Sr}$	Core, section, interval (cm)	Depth (mbsf)	Sr (μM)	$^{87}\text{Sr}/^{86}\text{Sr}$
344-U1412A-				344-U1413C-			
1H-1, 138–150	1.4	82.9	0.70918	3R-1, 90–122	188.6	49.8	0.70837
3H-5, 138–150	22.8	79.9	0.70907	5R-1, 116–146	208.3	57.4	0.70841
5H-3, 118–135	36.2	85.7	0.70907	7R-2, 77–107	228.8	64.0	0.70837
7H-5, 103–120	55.9	88.3	0.70903	7R-6, 101–131	234.5	65.6	0.70831
9H-2, 128–150	65.8	92.7	0.70900	11R-1, 114–151	266.4	67.3	0.70819
11H-2, 130–152	83.9	83.7	0.70898	15R-2, 103–135	306.1	68.9	0.70820
13H-2, 124–146	93.6	79.7	0.70893	19R-5, 66–96	347.9	65.7	0.70836
15H-2, 93–115	106.8	85.1	0.70887	27R-2, 120–150	423.1	72.3	0.70841
16X-2, 126–148	111.0	85.1	0.70888	37R-5, 61–96	523.6	73.7	0.70841
344-U1412B-				39R-4, 116–151			
5X-1, 120–140	176.4	108	0.70851	43R-2, 73–105	577.7	67.3	0.70835
6X-2, 99–125	186.7	122	0.70843	344-U1414A-			
7X-2, 51–73	196.0	140	0.70856	1H-1, 51–61	0.6	84.5	0.70916
344-U1412C-				2H-3, 131–141	6.1	78.6	0.70907
5R-1, 0–36	328.8	314	0.70887	3H-2, 131–141	14.2	73.5	0.70913
6R-1, 9–20	338.6	309	0.70870	5H-2, 131–141	33.2	70.9	0.70912
7R-2, 96–112	350.3	270	0.70837	7H-2, 140–150	52.4	75.8	0.70916
8R-5, 59–95	364.1	251	0.70840	9H-2, 140–150	71.4	84.5	0.70908
9R-2, 58–88	369.0	262	0.70874	9H-5, 140–150	75.9	85.3	0.70913
344-U1413A-				11H-5, 140–150	94.9	94.9	0.70902
1H-4, 110–122	5.6	88.0	0.70914	13H-5, 140–150	113.9	95.8	0.70894
2H-5, 138–150	14.0	83.2	0.70911	15H-2, 140–150	128.4	104	0.70880
3H-6, 140–152	24.4	50.5	0.70899	17H-4, 135–150	150.3	131	0.70874
5H-5, 135–150	42.2	49.5	0.70902	19H-6, 135–150	170.7	171	0.70882
9H-5, 135–155	77.2	44.6	0.70893	21H-3, 135–150	185.5	217	0.70883
11H-5, 130–152	96.1	38.5	0.70893	23X-5, 130–150	207.5	279	0.70879
13H-4, 130–152	111.3	41.2	0.70895	24X-4, 94–114	214.9	304	0.70879
15H-2, 130–152	121.4	48.0	0.70894	25X-2, 112–132	222.0	318	0.70881
17H-4, 128–150	134.6	51.9	0.70896	26X-5, 130–150	236.3	358	0.70882
19X-4, 97–122	145.9	53.7	0.70890	27X-3, 130–150	242.9	372	0.70881
21X-3, 125–150	164.1	48.9	0.70863	28X-4, 67–97	252.9	402	0.70882
25X-1, 48–84	179.9	48.7	0.70824	30X-2, 79–109	269.4	468	0.70885
344-U1413B-				34X-1, 0–14	302.2	494	0.70884
1H-3, 140–150	4.4	87.9	0.70916	35X-1, 15–39	308.0	710	0.70884
1H-5, 60–70	6.2	87.3	0.70912	36R-2, 67–92	314.0	711	0.70879
2H-5, 140–150	14.0	83.6	0.70913	37R-3, 74–103	319.5	861	0.70869
3H-2, 144–154	18.2		0.70905	38R-1, 123–145	327.4	942	0.70865
3H-3, 142–152	19.7	56.2	0.70901	38R-2, 66–102	328.4	908	0.70869
				39R-1, 80–94	336.7	904	0.70861