

Dear Prof. Oka

Thank you very much for your review and comments. I modified the manuscript based on your comments and suggestions as well as other reviewer's comments.

1. While the original manuscript insisted that the observed transport of radiocaesium suggests a new pathway of STMW circulation, the revised manuscript is more descriptive, focusing mainly on the radiocaesium transport from the East China Sea to the Japan Sea. I felt that this revision was adequate and suggest further revision to concentrate on the ECS to JS transport processes, because the authors are not showing any analysis nor discussion on the transport processes from the North Pacific to the ECS, particularly in Sec. 3.3.

The modified part was as follows:

[P6 L12-16]

We describe the possible pathway of FNPP1-¹³⁷Cs from the Pacific Ocean to the SOJ by ECS. Taking into account the physical ocean circulation, FNPP1-¹³⁷Cs in the STMW would be transported to southwestward by the westward undercurrent as reported by Oka (2009). The STMW containing ¹³⁷Cs would reach the western boundary at lower latitudes, is entrained into the Kuroshio (Oka and Qiu, 2012), and is northward transported to the west of Kyushu by the TWC bifurcated from the Kuroshio.

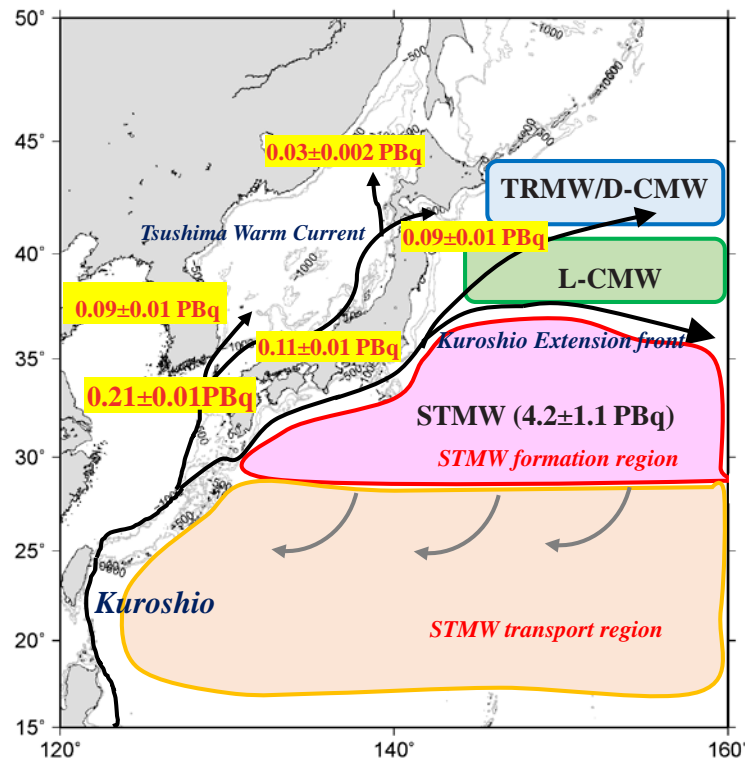
[P7 L 1-6]

There were found several signatures that (i) The maximum ¹³⁷Cs activity concentrations were observed in the subsurface layer in comparison with those in the surface seawater, (ii) This higher ¹³⁷Cs activity concentrations were observed in the STMW based on the potential density data, (iii) In the ECS, the potential density in ¹³⁷Cs existed seawater was almost same value those in the STMW, suggesting the signature of FNPP1-¹³⁷Cs transport into the ECS from the STMW in the NPSJ, (iv) The ¹³⁷Cs activity concentrations in the northern ECS (>30°N) were higher than those in the southern ECS (<30°N) (Fig. SII2). In this study, however, the data was not enough to clarify the transport route in more detail.

2. The schematic in Fig. 8 still needs to be revised. The Kuroshio path is too much simplified. Its pathway from the ECS to the NP via the Tokara Strait is not expressed at all. STMW is formed as deep winter mixed layers north of ~28N and is subducted southward across the mixed layer front at ~28N. Then, it is transport

westward and reaches the western boundary at ~25N or further south (e.g., Oka, 2009).

A. Fig. 8 was modified as shown. The Kuroshio current path was shown in more detail. In added to this, I added the STMW (formation region and transport region), CMW, and TRCMW.



3. As I commented to the original manuscript, I believe that STMW with radiocaesium does NOT “enter the ECS from the region between Kagoshima and Tokara” (p.7, 1.1415) as shown by blue arrows in Fig. 8. If the authors insist so, please show the evidence.

A. Because our data was very limited, the description of the entrance region of FNPP1-¹³⁷Cs would be overstatement. The last sentence in Section 4.1 “FNPP1-radiocaesium might have entered the ECS from the region between Kagoshima and Tokara” was deleted.

[P6 L12-16]

We describe the possible pathway of FNPP1-¹³⁷Cs from the Pacific Ocean to the SOJ by ECS. Taking into account the physical ocean circulation, FNPP1-¹³⁷Cs in the STMW would be transported to southwestward by the westward undercurrent as reported by Oka (2009). The STMW containing ¹³⁷Cs would reach the western boundary at lower latitudes, is entrained into the Kuroshio (Oka and Qiu, 2012), and is northward transported to the west of Kyushu by the TWC bifurcated from the Kuroshio.

Other comments:

1. Title: What does “rapid” mean? Given the timescale of 2-3 years for STMW circulation shown by previous studies, I think the time period for radiocaesium transport shown by the present study is reasonable but not “rapid”.
- A. The term of “rapid” was deleted from the title. The time scale of FNPP1-¹³⁷Cs transport from ECS to SOJ was consistent with those of nutrient species (e.g., Kodate et al., 2016).

The title was modified as follows.

[P1. L1-2]: Transport of FNPP1-derived radiocaesium from subtropical mode water in the western North Pacific Ocean to the Sea of Japan

2. P.1, l.16-18, “A time lag of the maximum .. in the SOJ.”: I do not understand what each sentence mean and do not understand the relation between the two sentences.
- A. This sentence was modified as follows.

[P 1, L 15-21]

In the East China Sea (ECS), the clear increase in the ¹³⁷Cs activity concentration started at 140 meters depth, density=25.2 kgm⁻³, in April 2013, and propagated to the surface layers at approximately 0-50 meters depth, and showing a maximum in 2015 and decreasing in the following years. In the ECS, the maximum Fukushima-radiocaesium activity concentration in surface water was observed in 2014/2015, while it in the SOJ was observed in 2015/2016. Therefore, the propagation of Fukushima-derived radiocaesium in the surface seawater was approximately one year from ECS into the SOJ.

3. p.2, 1.31-32, “The SOJ is connected to the Pacific Ocean at its southwest through the Tsushima Straits”: This is incorrect. The SOJ is connected to the East China Sea.

A. This description was unsuitableness. This sentence was also modified as follows.

[P 2, L 30-31]

The SOJ is connected to the East China Sea at its southwest through the Tsushima Straits and connected to the North Pacific Ocean at its northeast through the Tsugaru Straits.

4. p.2, 1.33, “this current splits into three paths.”: Please add references

A. There are several currents after entering the Sea of Japan such as East Korean Warm Current, North Korean Cold Current, the first and second branches of Tsushima Warm Current. Although several current in the SOJ, Tsushima Warm Current flows to the northeast through the Tsushima Strait from the ECS and mainly separated into two (west of TWC and east of TWC). Therefore, the description was modified as follows.

This part was modified as follows.

[P 2, L 29-39]

The SOJ is located between the Eurasian continent and the Japanese archipelago. The area is 1008000 km², and the mean depth is 1667 m (Menard and Smith, 1966). The SOJ is connected to the East China Sea at its southwest through the Tsushima Straits and connected to the North Pacific Ocean at its northeast through the Tsugaru Straits. Warm and saline seawater passes through the Tsushima Strait as the Tsushima Warm Current (TWC), and this current splits into mainly two paths: One is the nearshore current along the west coast of Honshu Island, Japan, and the seawater passes through the Tsugaru Straits and enters to the Pacific Ocean again. The seawater transported to the northward passes thought the Soya Strait and connected to the Okhotsk Sea. Another current flows north of the Korean Peninsula. This current meets the North Korean Cold Current, which is the prolongation of the Liman Cold Current. Therefore, this northward warm subtropical waters and southward cold subarctic waters form the Polar Front meet at approximately 40°N, and the SOJ is largely divided into two regions (Prants et al., 2015). The increased FNPP1-¹³⁷Cs radioactivity concentrations were observed in the west coast of Honshu island (Aoyama et al., 2017).

5. In the last paragraphs of Sec. 1, the authors need to explain more clearly what advances from the preliminary study of Aoyama et al. (2017) the authors are going to show in this study.

A. Based on the preliminary results, Aoyama et al. (2017) reported the increased of ¹³⁷Cs activity concentrations in the surface seawater around the Japanese islands and ECS in winter 2015/2016. In this study, the author described the temporal variation of ¹³⁷Cs

activities after the 1960's in the SOJ, and estimate the FNPP1-derived ^{137}Cs activity concentrations. The author also investigated the vertical distributions of the ^{137}Cs activity concentrations. The authors also discussed about FNPP1- ^{137}Cs transport route from the NPSJ to the SOJ, estimate of transport amount with uncertainty of FNPP1- ^{137}Cs into the SOJ as well as return amount to North Pacific Ocean from 2012 to 2016.

The modified part was as follows;

[P 2, L 40~ P3 L3]

Aoyama et al. (2017) reported the increased of ^{137}Cs activity concentrations in the surface seawater around the Japanese islands and ECS in winter 2015/2016 as a preliminary result. This research was investigated the temporal variation of ^{137}Cs activities after the 1960's in the SOJ, and estimated the FNPP1-derived ^{137}Cs activity concentrations after the year of 2011. We also investigated the vertical distributions of the ^{137}Cs activity concentrations in the NPSJ, ECS, and SOJ. We also discussed about FNPP1- ^{137}Cs transport route from the North Pacific South of Japan (NPSJ) to the SOJ, estimation of transport amount with uncertainty of FNPP1- ^{137}Cs into the SOJ as well as return amount of FNPP1- ^{137}Cs into the North Pacific Ocean during the period from 2012 to 2016.

6. p.5, l.20: The wording “the southern Japan (SJPN)” does not make sense. I would call the region “North Pacific south of Japan (NPSJ)”.

A. The term “the southern Japan (SJPN)” was modified to “*North Pacific south of Japan (NPSJ)*”.

The modified part was as follows;

[P 5, L 22]

Propagation of radiocaesium from the upstream region (NPSJ and ECS) to the downstream region (the SOJ)

7. p.5, l.34-35, “Fig. 7 displays the latitude-time cross-sections ... along the eastern TWC.”: How did the authors distinguish the eastern and western TWC? Please explain.

A. At first, the cross sections along the western TWC (off Korean Peninsula) and the eastern TWC (along the Honshu islands in Japan) was depicted in Fig. 7. However, the data along with the western TWC was very limited and omitted in the revised version. The term “latitude-time cross-sections” was also modified as “Hovmoller diagrams”.

The modified part was as follows.

[P 5, L 38-39] Fig. 7 displays the Hovmoller diagrams of the ^{137}Cs activity concentrations at potential temperatures of $25.2\pm 0.5 \text{ kgm}^{-3}$ along the TWC in the ECS and coastal site of eastern SOJ.

8. p.6, l.6: What does “the other layers” mean?

A. As described above, the Hovmoller diagrams of the ^{137}Cs activity concentrations along the western TWC (off Korean Peninsula) was deleted. Then the description (P6, L5-10) was also deleted.

Revised manuscript OS-2017-90 entitled “Transport of FNPP1- derived radiocaesium from subtropical mode water in the western North Pacific Ocean to the Sea of Japan” by Inomata et al.

The authors appreciate to the 2nd review and positive comments. We modified the manuscript based on your comments.

Major points:

- The authors may have improved the English of the revised manuscript but not of the response letter which contains numerous mistakes and is, again, very difficult to read and understand. Some co-authors of this paper have extensive experience in publishing scientific papers in English; I am puzzled by the fact they have co-signed such English text (e.g. did they really read that paper and that response letter at least once?).

A. The English was carefully checked and modified in the revised version and response letter.

- I am convinced by the authors that the Cs137 signal in the SOJ first originates at the subsurface (e.g. 140 and 200 m) and then propagate to the surface, thus supporting their hypothesis of “mode water transport”. However, I am not convinced by the first entrance through the Tsushima strait: indeed, on Fig. 3 c/d and Fig. 5 a/b, one can see that the subsurface peak of Cs137 occurred first (mid 2013) at station 105-11 (within the SOJ) and 1 year later at station 304-01 (ECS, e.g. upstream). Why?

A. X-axis in Figure 5a and b show different periods. The author apologized that this might be confusing the reviewer. The x-axis in Figure 5 a and 5b was modified.

The increase of ¹³⁷Cs activity concentrations was observed in the year of 2013 in the stations of 314-01 (140 m depth) and 105-11 (200 m depth). The ¹³⁷Cs activity concentrations in the year of 2013 was almost same in the stations of 314-01 (2.1 ± 0.16 , 2.1 ± 0.19 Bqm⁻³) and 105-11 (2.4 ± 0.18 Bqm⁻³) (see Figure 5) or slightly higher in the station 105-11. The increase of ¹³⁷Cs activity concentrations at subsurface layer occurred in 2013 and this was approximately 1 year earlier than that in the shallower layers. Difference of ¹³⁷Cs activity concentrations among the two stations would strongly reflect whether the seawater locate the center of mode water transport or not. As shown in Fig. 6(e) and (f), the ¹³⁷Cs existed in the denser layer than those in the subtropical mode water, resulting with the cooling in the surface layer by southward transported colder seawater after the sea water was transported from the Tsushima Strait. This effect would also influence the slight difference of ¹³⁷Cs activity concentrations in the stations of 314-01 and 105-11.

The modified parts were as follows ;

[P5, L 9-12] : Fig. 5 show the temporal variation in the ¹³⁷Cs activity concentrations at different depths at station 314-01 in the ECS (Fig. 5a; 0-140 m) and station 105-11 in the SOJ (Fig. 5b; 0-2000 m). The increase in the ¹³⁷Cs activity concentrations at the subsurface layer (140 m at the station 314-01, 200 m at the station 105-11) occurred in 2013 and this was approximately 1 year earlier than that in the shallower layers.

[P 7, L 30-38]

As shown in Fig. 6e, the ¹³⁷Cs activity concentrations at station 105-11, located along the western TWC in the SOJ, were maximum at the surface and gradually decreased with increasing depth. This vertical distribution is different from those in the NPSJ (Fig. 6a) and ECS (Fig. 6c). Particularly, the subsurface

peak observed in the NPSJ and ECS did not appear at station 105-11. At station 105-11, most of the ^{137}Cs existed in seawater with a potential density of $25.7\text{-}27.3\text{ kg m}^{-3}$ (Fig. 6f), which was located in a higher potential density layer than that in the NPSJ and ECS. A similar vertical distribution was also observed at the western coast of the Japanese Islands along the eastern TWC (Fig. S111). These distributions were due to cooling in the surface layer after water was transported from the Tsushima Strait. Physical processes such as the convergence and subduction of surface water inside the eddies are important mechanisms of downward transport of radiocaesium (Miyao et al., 1998; Budyansky et al., 2015).

- The “blob” of high Cs137 ($8\text{-}12\text{ Bq/m}^3$) reported on Fig. 6a, b: could it be related to “exceptional sampling sites”; e.g. sampling within the core of an eddy that transport Cs137? (e.g. Prants et al. 2017 Ocean Sci. Discuss., doi:10.5194/os-2016-103, 2017).

A. Prants et al. (2017) identified the high activity of radioactivity in the mesoscale eddies by lagrangian methodology. However, the high Cs137 ($8\text{-}12\text{ Bq/m}^3$) reported on Fig. 6a, 6b were observed in the region south of Kuroshio based on the temperature profile. Furthermore, based on the analysis by AVISO, these data were observed in the region far from the eddies. I think that these data did not show the signature of direct transport across Kuroshio by the eddy effect.

- Please read better section 3.3. “Vertical distribution of Cs-137 and interior pathways” of Rossi et al. 2013 and please summarize their findings better in your introduction. The corrigendum concerns only the concentrations simulated (1 order of magnitude lower than reported), not the validity of the interior (and surface) pathways and subduction process.

A. The author described the penetration of FNPP1-derived ^{137}Cs in the ocean interior by numerical model simulations (Rossi et al., 2013).

The modified parts were as follows ;

[P 2, L 7-11]; Rossi et al. (2013) reported that simulated FNPP1-derived ^{137}Cs were penetrated into the ocean interior by intense subduction and vertical mixing on winter season around the formation regions of mode waters as described by Oka and Qiu (2012). In their results, the higher ^{137}Cs activity concentrations associated by the subduction was also reproduced in the STMW, Denser, and Lighter CMW. About 43% of FNPP1- ^{137}Cs was transported downward below the mixed layer by eddy processes (Kamidaira et al., 2018).

- Figures are still of poor quality. For instance, labels have been misplaced in Fig. 7; labels are not readable on Fig. 8; etc...
- Fig. 7: those latitude-time plots are usually called “Hovmoller diagram”.

In the revised version, the author selected only one Figure 7a because the data in Figure 7b and c were very limited. The quality of new Figure 7 is improved. Figure 8 is re-depicted by the suggestions of reviewer 1. I also take into account to be higher quality for all Figures.

A. The plots were modified as “Hovmoller diagram”.

The modified parts were as follows ;

[P 5, L 38-39]; Fig. 7 displays the Hovmoller diagrams of the ^{137}Cs activity concentrations at potential temperatures of $25.2\pm 0.5\text{ kgm}^{-3}$ along the TWC in the ECS and coastal site of eastern SOJ.

Caption ; Figure 7: Hovmoller diagrams of the ^{137}Cs activity concentrations at a potential density along with an eastern TWC at a potential density of $25.2 \pm 0.5 \text{ kg m}^{-3}$. The ECS stations described on the x-axis are Kagoshima and Saga stations. Southwestern SOJ includes the monitoring stations Shimane, Fukui, Ishikawa, and Niigata. The northwestern SOJ includes the monitoring stations Aomori and Tomari. Color indicates the ^{137}Cs activity concentrations (Bq m^{-3}).

- Title of section 3.2 (p 7): upstream sites are SJPN and ECS, right?
- A. The author considered the NPSJ (modified from SJPN according to the reviewer 1's comment) and ECS are upstream sites.

The modified part was as follows;

[P 5, L 22]; 3.2 Propagation of radiocaesium from the upstream region (NPSJ and ECS) to the downstream region (the SOJ).

- Reference incomplete p 10 line 41.
- A. The reference was modified as follows.
[P 10, L 12-13] ; Busseler, K., Aoyama, M., Fukasawa, M. Impacts of Fukushima Nuclear Power Plants on Marine Radioactivity, Environ. Sci. Technology, 45, 9931-9935, 2011.

- Abstract line 16-18: a sentence was repeated.

A. This part was modified as follows.

[P 1, L 15-20] ; In the East China Sea (ECS), the clear increase in the ^{137}Cs activity concentration started at 140 meters depth, density= 25.2 kg m^{-3} , in April 2013, and propagated to the surface layers at approximately 0-50 meters depth, and showing a maximum in 2015 and decreasing in the following years. In the ECS, the maximum Fukushima-radiocaesium activity concentration in surface water was observed in 2014/2015, while it in the SOJ was observed in 2015/2016. Therefore, the propagation of Fukushima-derived radiocaesium in the surface seawater was approximately one year from ECS into the SOJ.

Transport of FNPP1-derived radiocaesium from subtropical mode water in the western North Pacific Ocean to the Sea of Japan

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Abstract.

The ¹³⁷Cs activity concentration in the Sea of Japan (SOJ) has been increasing since 2012/2013 and reached a maximum in 10 2015/2016 of approximately 3.4 Bq m⁻³, while the pre-Fukushima accident ¹³⁷Cs activity concentration was approximately 1.5 Bq m⁻³. It was also observed that the ¹³⁴Cs/¹³⁷Cs activity ratios ranged from 0.36 to 0.51 in 2016. Taking into account that the radioactive decay and ocean mixing, these ¹³⁴Cs/¹³⁷Cs activity ratios are evidence that the increase in the ¹³⁷Cs activity concentration was derived from the Fukushima accident. In the North Pacific South of Japan (NPSJ), the higher ¹³⁷Cs activity was observed in the 300 m depths, which corresponds to the subtropical mode water (STMW) based on the potential 15 density, in 2012/2013. In the East China Sea (ECS), the clear increase in the ¹³⁷Cs activity concentration started at 140 meters depth, density=25.2 kgm⁻³, in April 2013, and propagated to the surface layers at approximately 0-50 meters depth, and showing a maximum in 2015 and decreasing in the following years. In the ECS, the maximum Fukushima-radiocaesium activity concentration in surface water was observed in 2014/2015, while it in the SOJ was observed in 2015/2016. Therefore, the propagation of Fukushima-derived radiocaesium in the surface seawater was approximately one year from ECS into the 20 SOJ. These temporal changes in the ¹³⁷Cs activity concentration and ¹³⁴Cs/¹³⁷Cs activity ratio indicated that a part of the ¹³⁷Cs and ¹³⁴Cs derived from the Fukushima accident (FNPP1-¹³⁷Cs and FNPP1-¹³⁴Cs) was transported to the ECS and then to the SOJ from the STMW in the NPSJ within several years. The integrated amount of FNPP1-¹³⁷Cs that entered the SOJ until 2016 was estimated to be 0.21±0.01 PBq, which corresponds to 5.0 % of the estimated total amount of FNPP1-¹³⁷Cs in the STMW. The integrated amount of FNPP1-¹³⁷Cs that returned to the North Pacific Ocean through the Tsugaru Strait was 25 estimated to be 0.09±0.01 Bq, which corresponds to 43 % of the total amount of FNPP1-¹³⁷Cs transported to the SOJ and 2.1 % of the estimated total amount of FNPP1-¹³⁷Cs in the STMW.

1 Introduction

The Fukushima Daiichi Nuclear Power Plant (FNPP1) accident in March 2011 released radiocaesium (¹³⁷Cs (T_{1/2} of 30.07 yr) and ¹³⁴Cs (T_{1/2} of 2.06 yr)) by atmospheric release to the air and the direct discharge of contaminated water to the 30 ocean, mostly in March and April 2011, and 80 % of atmospheric released radiocaesium deposited ocean surface (Aoyama et al., 2016a; Buesseler et al., 2016; Hirose, 2016; Tsumune et al., 2013). The ¹³⁷Cs activity concentration in the surface seawater of the North Pacific Ocean after the FNPP1 accident ranged from a few Bq m⁻³ to ca. 1 kBq m⁻³ (e.g., Aoyama et al., 2012, 2013; Honda et al., 2012; Kaeriyama et al., 2013, 2014). A basin-scale measurement indicated that the FNPP1-derived radiocaesium was observed in a region from 25°N to 50°N and from 135°E to 135°W in April and May of 2011 (e.g., 35 Aoyama et al., 2013; 2016a; Inomata et al., 2016; Tsubono et al., 2016). The radiocaesium moved eastward to approximately 40°N with the North Pacific current systems, such as the Kuroshio and Kuroshio Extension, at a rate of approximately 8 cm sec⁻¹ and reached 165°E in July-September 2011 and the 180th meridian in January-March 2012 (Aoyama et al., 2012, 2013). FNPP1-derived radiocaesium was also found further south in the North Pacific Ocean, with a subsurface maximum at a depth of approximately 300 m (Kaeriyama et al., 2013, 2014, 2016; Kumamoto et al., 2013, 2017). These subsurface 40 maxima of ¹³⁷Cs were found in the subtropical mode water (STMW; potential density (σ_θ)=25.0-25.6 kgm⁻³; Masuzawa,

1969) and central mode water (CMW; $\sigma_{\theta}=26.0-26.5 \text{ kgm}^{-3}$) (Aoyama et al., 2016a; Kumamoto et al., 2014, 2015). It was deduced that atmospherically-deposited ^{137}Cs south of the Kuroshio Extension subducted into the STMW and transported southward within 10 months after the FNPP1 accident.

Ocean circulated models captured that the FNPP1-derived radiocaesium were transported in the North Pacific Ocean with advecting and diluting in the surface seawater (Tsubono et al., 2016; Rossi et al., 2014). It was also revealed that dilution due to horizontal and vertical dispersion in the vicinity of Kuroshio led to a rapid decrease of the radiocaesium activity concentrations (Behrens et al., 2012). Rossi et al. (2013) reported that simulated FNPP1-derived ^{137}Cs were penetrated into the ocean interior by intense subduction and vertical mixing on winter season around the formation regions of mode waters as described by Oka and Qiu (2012). In their results, the higher ^{137}Cs activity concentrations associated by the subduction was also reproduced in the STMW, Denser CMW, and Lighter CMW. About 43% of FNPP1- ^{137}Cs was transported downward below the mixed layer by eddy processes (Kamidaira et al., 2018).

^{137}Cs was injected into the environment by the large-scale atmospheric nuclear weapons testing that occurred in the late 1950s and early 1960s and the Chernobyl accident in 1986, therefore before the FNPP1 accident, ^{137}Cs existed in the North Pacific Ocean and its marginal seas (Aoyama et al., 2006; Inomata et al., 2009). Furthermore, the dumping of radioactive wastes by Russia and the former USSR occurred in the northern part of the Sea of Japan (SOJ), although this caused no significant increase in the activity concentrations (Miyao et al., 1998). In the 2000s, the ^{137}Cs activity concentrations had an almost homogenous distribution in the Pacific Ocean, although relatively high activity was observed in the western part of the subtropical gyre in the North Pacific Ocean (exceeding 2 Bq m^{-3}) and the South Pacific Ocean (exceeding 1.5 Bq m^{-3}) (Aoyama et al., 2012). In March-April 2011, FNPP1-derived ^{137}Cs and ^{134}Cs (FNPP1- ^{137}Cs , FNPP1- ^{134}Cs) were released into the North Pacific Ocean. There have been two sources of ^{137}Cs in the North Pacific Ocean and its marginal seas, the ^{137}Cs released by global fallout (global fallout- ^{137}Cs) and FNPP1- ^{137}Cs . In contrast, the ^{134}Cs derived from global fallout and the Chernobyl accident decayed out at the measurement in 1993 due to its shorter lifetime (Miyao et al., 1998). Therefore, ^{134}Cs is regarded as an adequate chemical tracer of the radiocaesium derived from the FNPP1 accident.

In the northeastern part of the SOJ, atmospherically deposited radiocaesium were caused to approximately 1-2 times higher activity concentrations of radiocaesium in May 2011 ($1.5-2.8 \text{ Bqm}^{-3}$) in northeastern SOJ than those of before the FNPP1 accident ($\sim 1.5 \text{ Bqm}^{-3}$) (Fig. 1). By the end of 2011, the ^{137}Cs activity concentrations in the northeastern part of the SOJ had rapidly decreased to almost the same levels as those before the FNPP1 accident (Inoue et al., 2012). However, several sets of seawater monitoring data have indicated an increase in the ^{137}Cs activity concentration after 2012.

The SOJ is located between the Eurasian continent and the Japanese archipelago. The area is 1008000 km^2 , and the mean depth is 1667 m (Menard and Smith, 1966). The SOJ is connected to the East China Sea at its southwest through the Tsushima Straits and connected to the North Pacific Ocean at its northeast through the Tsugaru Straits. Warm and saline seawater passes through the Tsushima Strait as the Tsushima Warm Current (TWC), and this current splits into mainly two paths: One is the nearshore current along the west coast of Honshu Island, Japan, and the seawater passes through the Tsugaru Straits and enters to the Pacific Ocean again. The seawater transported to the northward passes through the Soya Strait and connected to the Okhotsk Sea. Another current flows north of the Korean Peninsula. This current meets the North Korean Cold Current, which is the prolongation of the Liman Cold Current. Therefore, this northward warm subtropical waters and southward cold subarctic waters form the Polar Front meet at approximately 40°N , and the SOJ is largely divided into two regions (Prants et al., 2015). The increased FNPP1- ^{137}Cs radioactivity concentrations were observed in the west coast of Honshu island (Aoyama et al., 2017).

Aoyama et al. (2017) reported the increased of ^{137}Cs activity concentrations in the surface seawater around the Japanese islands and ECS in winter 2015/2016 as a preliminary result. This research was investigated the temporal variation of ^{137}Cs activities after the 1960's in the SOJ, and estimated the FNPP1-derived ^{137}Cs activity concentrations after the year of 2011. We also investigated the vertical distributions of the ^{137}Cs activity concentrations in the NPSJ, ECS, and SOJ. We also

discussed about FNPP1-¹³⁷Cs transport route from the North Pacific South of Japan (NPSJ) to the SOJ, estimation of transport amount with uncertainty of FNPP1-¹³⁷Cs into the SOJ as well as return amount of FNPP1-¹³⁷Cs into the North Pacific Ocean during the period from 2012 to 2016.

5 2 Sampling, measurements and methods

2.1. Data sources

After the FNPP1 accident, many radiocaesium measurements were taken in the SOJ and western North Pacific Ocean (Fig. 1). To elucidate the temporal and spatial distributions of the radiocaesium activity concentrations, we use as many data points as possible. We, therefore, compiled all available data from the literature and reported studies. Most of the data before 10 the FNPP1 accident was included in the database, “Historical Artificial Radionuclides in the Pacific Ocean and its Marginal Seas (HAM database)” (Aoyama and Hirose, 2003 and their updated version). The data observed after the FNPP1 accident were shown in Aoyama et al. (2016a). The term “surface seawater” used in this study defines a sample collected shallower than 10 m depth.

We also focused on the Japanese government’s monitoring data at Tomari (42.98-43.17°N, 140.21-140.30°E), Aomori 15 (41.13-41.22°N, 141.50-141.67°E), Niigata (37.62-38.10°N, 138.38-138.84°E), Ishikawa (36.87-37.29°N, 136.43-136.47°E), Fukui (35.75-36.09°N, 135.50-135.83°E), Shimane (35.67-35.80°N, 132.87-133.2°E), Saga (33.57-33.62°N, 129.73-129.98°E), and Kagoshima (31.58-31.93°N, 130.02-130.15°E) (Marine Ecology Research Institute, 2011, 2012, 2013, 2014, 2015, 2016) (Fig. 1). These measurements were taken once a year (from the middle of May to early June). Near the Aomori sites, offshore monitoring was also conducted twice a year (May and October) at the Rokkasho Reprocessing Plant (39.5- 20 41.4°N, 141.5-142.3°E). Seawater was sampled from 0-664 m with different depths at each monitoring site. Monitoring data (304-01(33.0°N, 127.7°E), 105-11(37.3°N, 131.3°E)) from the Korean government was also used in this analysis (Korea Institute of Nuclear Safety, 2011, 2012, 2013, 2014, 2015, 2016) (Fig. 1). At these monitoring sites, the surface seawater (0 m) measurements were taken four times (February, April, August, October) a year.

25 2.2. Measurement of the ¹³⁴Cs activity concentration

Because the ¹³⁴Cs activity concentrations in most of the monitoring datasets were less than the detection limit due to low measurement capabilities, we also collected seawater samples to investigate the ¹³⁴Cs activity concentrations. The sample volume ranged from a few litres to ten litres. The radiocaesium in the sampled seawater was extracted by the ammonium phosphomolybdate (AMP)/Cs compound method improved by Aoyama and Hirose (2008b). The AMP/Cs compound was 30 measured by using ultra-low-background gamma-ray detectors in the Low Level Radioactivity Concentration Laboratory, Kanazawa University (Hamajima and Komura, 2004) to measure ¹³⁴Cs and ¹³⁷Cs activity concentrations. The measurement details are described in Aoyama and Hirose (2008b) and Aoyama et al. (2016a). All radioactivity concentrations shown in this research were decay corrected at the time of sample collection. In addition, we also determined the ¹³⁴Cs/¹³⁷Cs activity ratio decay corrected to 11 March 2011 at the time of the FNPP1 accident.

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2.3. Estimation of the FNPP1-¹³⁷Cs activity concentration

In order to investigate the contribution of FNPP1-¹³⁷Cs, increased ¹³⁷Cs activity concentrations derived from the FNPP1 accident were estimated in this study. As shown in Fig. 2., the ¹³⁷Cs activity concentrations in the SOJ exponentially decreased, and the decreasing rate from 2000 to 2010 become to slow. By assuming that the apparent half-residence time of 40 the global fallout-¹³⁷Cs was approximately the same before and after the FNPP1, the global fallout-¹³⁷Cs activity concentrations were estimated as shown in Fig. SI1. Then, the FNPP1-¹³⁷Cs was estimated from the difference between the measured and the extrapolated value of the half-year average in the SOJ during the period from 2000 to 2010 (Fig. SI1).

2.4. Estimation of the ^{137}Cs transport amounts during the period from 2012 to 2016

We estimated the transported amounts of FNPP1- ^{137}Cs in each region during the period from 2012 to 2016 by using the following equation (1):

$$\text{Transport amount} = \sum_{i=2012}^n [\text{FNPP1-}^{137}\text{Cs activity concentration in year } i] \times$$
$$[\text{annual average seawater transport volume in year } i] \quad (1)$$

where $n = 2016$, the annual average FNPP1- ^{137}Cs activity concentrations at the station Saga and the station 304-11 were used for the ECS, the FNPP1- ^{137}Cs activity concentrations at the station Shimane were used for the eastern TWC, the FNPP1- ^{137}Cs activity concentrations at the station 105-01 were used for the western TWC, and the FNPP1- ^{137}Cs activity concentrations at the station Aomori were used for the transported amount to the North Pacific Ocean via the Tsugaru Strait. The amount transported FNPP1- ^{137}Cs northward along the western coast of Hokkaido was estimated from the difference between the total inflow of FNPP1- ^{137}Cs at the entrance and the outflow that passed through the Tsugaru Strait. The annual average volume of seawater transported through the TWC was used the data obtained from Fukudome et al. (2010) and Han et al. (2016). The annual average seawater transport volumes of the TWC into the SOJ ranged from $2.59 \pm 0.50 \times 10^6 \text{ m}^3 \text{ s}^{-1}$ to $2.82 \pm 0.58 \times 10^6 \text{ m}^3 \text{ s}^{-1}$, of which the annual average seawater transport volume through the east and west channels of the Tsushima Strait were 1.08 ± 0.26 - $1.26 \pm 0.27 \times 10^6 \text{ m}^3 \text{ s}^{-1}$ and 1.43 ± 0.38 - $1.56 \pm 0.47 \times 10^6 \text{ m}^3 \text{ s}^{-1}$ during the period from 2012 to 2016, respectively. The volume transported through the Tsugaru Strait was estimated at 1.0 - $1.3 \times 10^6 \text{ m}^3 \text{ s}^{-1}$ by Nishida et al. (2003), and this was used because we did not have *in situ* monitoring data for the seawater transport volume. The evaluation of seasonal variations was difficult in this study because of the limited monitoring data, which were measured once a year at the Japanese monitoring site and four times a year at the Korean monitoring sites.

3 Results

3.1 Increased ^{137}Cs activity concentrations in the ECS and the SOJ

Fig. 2 shows the temporal variation in the ^{137}Cs activity concentrations in the surface seawater of the SOJ after the 1960s. The half-year average values of the ^{137}Cs activity concentrations were also plotted. The ^{137}Cs activity concentrations exponentially decreased from 1960-1970; these concentrations originated from atmospheric nuclear weapons testing (Aoyama, 2009). The increase in the ^{137}Cs activity concentrations in 1986 was derived from the Chernobyl accident (Miyao et al., 1998). The apparent half-residence time during the period from 1970 to 1990 was estimated to be 16.3 ± 0.5 years (Inomata et al., 2009). In the 2000s, the decreasing rate of ^{137}Cs activity concentrations were small and the ^{137}Cs activity concentrations in the surface seawater reached low, 1.5 - 2.4 Bq m^{-3} as for half-year average value. Just several months after the FNPP1 accident, high activity concentrations of ^{137}Cs , up to 3.3 Bq m^{-3} , were observed. As shown in Fig. 1, higher ^{137}Cs activity concentrations were observed in the northeastern part of the SOJ than those in the southwestern part of the SOJ. These high ^{137}Cs activity concentrations were associated with the wet deposition of ^{137}Cs released into the atmosphere during the FNPP1 accident in March-April 2011. Other than the increase in the ^{137}Cs activity concentrations just after the FNPP1 in 2011, the ^{137}Cs activity concentrations gradually increased in the SOJ, and the ^{137}Cs activity concentrations in 2016 reached 2.6 Bq m^{-3} , which is the same level as those in 1998.

Fig. 3 shows the FNPP1- ^{137}Cs activity concentrations measured in the ECS and SOJ by the Japanese and Korean governments. These monitoring stations are located along a branch of the Kuroshio in the ECS and along the western and eastern TWC in the SOJ. The long-term variations in the ^{137}Cs activity concentrations and FNPP1- ^{137}Cs at each monitoring station are also shown in Fig. SI2-9. Excluding the high activity concentrations at Aomori in 2011 caused by the wet deposition of FNPP1- ^{137}Cs released to the atmosphere, an increase in the FNPP1- ^{137}Cs in the SOJ started in 2012/2013, and the FNPP1- ^{137}Cs gradually increased up to 1.3 Bq m^{-3} until 2016 (Fig. 3a, b). The lower activity concentrations at Niigata (Fig. 3b) resulted from the measurements at another station near the Sado Island, off Niigata. At these monitoring stations, there was no significant decreasing trend of the FNPP1- ^{137}Cs activity concentrations in 2015 and 2016. On the other hand,

the FNPP1-¹³⁷Cs activity concentrations at stations located upstream from the SOJ, in the ECS (Fig. 3c; Kagoshima, Saga, and station 314-01), also increased from 2012/2013. Particularly, the increase was clearly observed after 2014 in the ECS. We should note that the FNPP1-¹³⁷Cs activity concentrations in 2016 tended to slightly decrease compared to those in 2015. At the station 105-11, which is located off South Korea (Fig. 1), marked increases in the FNPP1-¹³⁷Cs activity concentrations were observed in 2014, and the FNPP1-¹³⁷Cs activity concentrations reached approximately 1.4 Bq m⁻³ in 2015 (Fig. 3d). In 2016, the FNPP1-¹³⁷Cs concentrations were decreased relative to those in the preceding years.

The ¹³⁴Cs/¹³⁷Cs activity ratios in 2016 ranged from 0.27 to 0.51 (Fig. 4). There was no significant difference in the ¹³⁴Cs/¹³⁷Cs activity ratio among the stations in the ECS and the SOJ.

Fig. 5 show the temporal variation in the ¹³⁷Cs activity concentrations at different depths at station 314-01 in the ECS (Fig. 5a; 0-140 m) and station 105-11 in the SOJ (Fig. 5b; 0-2000 m). The increase in the ¹³⁷Cs activity concentrations at the subsurface layer (140 m at the station 314-01, 200 m at the station 105-11) occurred in 2013 and this was approximately 1 year earlier than that in the shallower layers. At station 314-01, the ¹³⁷Cs activity concentrations at 140 m depth were higher than those at 0 m depth beginning in 2013, and the ¹³⁷Cs activity concentrations increased up to 3.2±0.28 Bq m⁻³ in 2014 and then tended to decrease after 2015. At shallower depths (0-50 m), increased ¹³⁷Cs activity concentrations were found in 2014, and there was no significant decrease in the ¹³⁷Cs activity concentrations in 2015 and 2016. At the station 105-11, the increase in the ¹³⁷Cs activity concentrations started earlier in the subsurface seawater (200 m depth) in 2013. In 2014, the ¹³⁷Cs activity concentrations in surface seawater also increased and reached 2.8±0.2 Bqm⁻³ in 2015. Decreases in the ¹³⁷Cs activity concentrations at 200 m and 0 m depth were observed after 2015, and a subsurface peak in the ¹³⁷Cs activity concentration at 200 m depth was not observed after 2015. Similar variations, in which the increase in the ¹³⁷Cs activity concentration started from the ocean interior, were observed at the Kagoshima and Saga monitoring stations (Fig. SI10).

3.2 Propagation of radiocaesium from the upstream region (NPSJ and ECS) to the downstream region (the SOJ)

The vertical distributions of the ¹³⁷Cs activity concentration over the depth and potential density (σ_θ , kg m⁻³) profiles in the North Pacific South of Japan (NPSJ), in the ECS, and at the station 105-11 are shown in Fig. 6. In the NPSJ, the ¹³⁷Cs activity concentrations showed subsurface maxima at approximately 300 m depth in 2012-2013, with activities of 8.2-12.3 Bqm⁻³ as shown in Fig. 6a. These high ¹³⁷Cs activity concentrations were measured in the region between 136-138°E and 26-30°N. After 2014, a subsurface peak in the ¹³⁷Cs activity concentration was not observed. These subsurface peaks in the ¹³⁷Cs activity concentrations were found in the layer corresponding to a potential density of 25.2 kg m⁻³ (Fig. 6b). This is consistent with previous findings of an ¹³⁷Cs activity maximum in STMW with a potential density of 25.0-25.6 kg m⁻³ (Kaeriyama et al., 2014; Kumamoto et al., 2014). In the ECS, the ¹³⁷Cs activity concentrations gradually increased beginning in 2012 and attained the maximum activity concentrations (2.9±0.24 Bqm⁻³) in 2015, following by a decreasing trend in the ¹³⁷Cs activity concentrations in 2016, as shown in Fig. 6c. The higher ¹³⁷Cs activity concentrations above 2 Bq m⁻³ in the ECS were found in the layer corresponding to a potential density of 23.6-25.2 kg m⁻³, as shown in Fig. 6d. On the other hand, the ¹³⁷Cs activity concentrations at the station 105-11, located in the western SOJ, decreased with increasing depth until 500 m (Fig. 6e). The higher ¹³⁷Cs activity concentrations at the station 105-11 were measured in 2014 and 2015, and the ¹³⁷Cs activity concentrations decreased in 2016. The ¹³⁷Cs activity concentrations above 2 Bq m⁻³ at the station 105-11, except one sample measured at 2015, were located in the layer with a potential density of 25.8-27.1 kg m⁻³ (Fig. 6f).

Fig. 7 displays the Hovmoller diagrams of the ¹³⁷Cs activity concentrations at potential temperatures of 25.2±0.5 kgm⁻³ along the TWC in the ECS and coastal site of eastern SOJ. These potential density surfaces were selected to show the maximum ¹³⁷Cs activity concentrations observed in the ECS. The vertical distributions of the ¹³⁷Cs activity concentrations with depth and potential density at each monitoring station are displayed in Fig. SI11. Note that in the SOJ, the vertical distributions of the ¹³⁷Cs activity concentrations below 250 m were almost constant, and the subsurface peak of ¹³⁷Cs was not found at the monitoring stations along the eastern TWC (Fig. SI11). It is noted that the ¹³⁷Cs activity concentrations before

the FNPP1 accident were approximately 1.5 Bq m^{-3} . As shown in Fig. 7, in the ECS, the ^{137}Cs activity concentrations gradually increased and attained the maximum in 2014/2015. The ^{137}Cs activity concentrations in the ECS tended to decrease in 2016 in a layer with a density of $25.2 \pm 0.5 \text{ kgm}^{-3}$. In the southwestern part of the SOJ (Shimane, Fukui, Ishikawa, Niigata), the ^{137}Cs activity concentrations also gradually increased beginning in 2012, and the activity concentrations attained a maximum of 2.5 Bq m^{-3} in 2015/2016; these trends were almost the same as those at the monitoring stations in the ECS. In the northwestern SOJ (Aomori and Tomari), the ^{137}Cs activity concentrations were slightly increased and higher activity concentrations above 2 Bq m^{-3} were observed in 2016. These results revealed that the propagation of FNPP1- ^{137}Cs occurred within one year from the ECS ($32\text{-}34^\circ\text{N}$) to the SOJ ($35\text{-}38^\circ\text{N}$) along the TWC as shown in Fig. 7.

10 3.3. Transport process and total amount of FNPP1- ^{137}Cs transported into the SOJ during the period from 2012 to 2016

We describe the possible pathway of FNPP1- ^{137}Cs from the Pacific Ocean to the SOJ by ECS. Taking into account the physical ocean circulation, FNPP1- ^{137}Cs in the STMW would be transported to southwestward by the westward undercurrent as reported by Oka (2009). The STMW containing ^{137}Cs would reach the western boundary at lower latitudes, is entrained into the Kuroshio (Oka and Qiu, 2012), and is northward transported to the west of Kyushu by the TWC bifurcated from the Kuroshio. Then, FNPP1- ^{137}Cs was entered the SOJ through the Tsushima Strait by the TWC. FNPP1- ^{137}Cs was divided into western (coastal station in Korea) and eastern (eastern coast of the Japanese Islands) portions along with the TWC. In addition, the remaining FNPP1- ^{137}Cs in the eastern TWC was transported northward to the western coast of Hokkaido (Fig. 8). Table 1 shows the total amount of FNPP1- ^{137}Cs during the period from 2012 to 2016. The integrated FNPP1- ^{137}Cs in the ECS during the period from 2012 to 2016 was estimated to be $0.21 \pm 0.01 \text{ PBq}$, which corresponds to 5.0 % of the FNPP1- ^{137}Cs in the STMW ($4.2 \pm 1.1 \text{ PBq}$) estimated by Kaeriyama et al. (2016). Connected with the separation of the TWC, the transported amount of FNPP1- ^{137}Cs in the eastern channel was estimated to be $0.11 \pm 0.01 \text{ PBq}$, which corresponds to 2.6 % of the FNPP1- ^{137}Cs into the STMW. On the other hand, the transported amount in the western channel was estimated to be $0.09 \pm 0.01 \text{ PBq}$, which corresponds to 2.1 % of the FNPP1- ^{137}Cs injected into the STMW. The eastern and western outflow amounts of FNPP1- ^{137}Cs were consistent with the range of uncertainty for the inflow amounts, $0.21 \pm 0.01 \text{ PBq}$. A small part of the transported FNPP1- ^{137}Cs ($0.09 \pm 0.01 \text{ PBq}$; 2.1 % of FNPP1- ^{137}Cs in STMW) passed through the Tsugaru Strait and has already returned to the North Pacific Ocean. In other words, the amount of FNPP1- ^{137}Cs that has returned to the North Pacific Ocean corresponds to 43 % of the total amount of FNPP1- ^{137}Cs transported to the SOJ and 2.1 % of the estimated total amount of FNPP1- ^{137}Cs in the STMW. The remaining $0.03 \pm 0.002 \text{ PBq}$ (14% of the transported FNPP1- ^{137}Cs in SOJ; 0.7 % of the transported FNPP1- ^{137}Cs in STMW) should be transported to the northern part of the SOJ (west of Hokkaido) or be partially transported to the deep region associated with deep convection and surface mixing.

4 Discussion

4.1. Signature of FNPP1- ^{137}Cs inflow in the SOJ by using $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratio

According to the atmospheric model simulations, atmospheric deposition of FNPP1- ^{137}Cs occurred over a wide region in the western North Pacific Ocean ($30\text{-}55^\circ\text{N}$, $140^\circ\text{E}\text{-}180^\circ$) (e.g., Aoyama et al., 2016). One of the region in which ^{137}Cs was deposited in south of Kuroshio and Kuroshio Extension regions corresponded to the STMW formation region (Aoyama et al., 2016; Oka et al., 2012, 2013). Because the release of FNPP1-radiocaesium in the atmosphere occurred at the end of March and early April, the deposition of FNPP1- ^{137}Cs into the North Pacific Ocean was regarded as a point source associated with time. This means that FNPP1 radiocaesium is a very useful tracer to investigate the transport in the North Pacific Ocean. As shown in Fig. 6, the vertical distributions of the ^{137}Cs activity concentrations in the NPSJ indicate that FNPP1- ^{137}Cs included in the STMW.

There were found several signatures that (i) The maximum ^{137}Cs activity concentrations were observed in the subsurface layer in comparison with those in the surface seawater, (ii) This higher ^{137}Cs activity concentrations were observed in the STMW based on the potential density data, (iii) In the ECS, the potential density in ^{137}Cs existed seawater was almost same value those in the STMW, suggesting the signature of FNPP1- ^{137}Cs transport into the ECS from the STMW in the NPSJ, (iv) The ^{137}Cs activity concentrations in the northern ECS ($>30^\circ\text{N}$) were higher than those in the southern ECS ($<30^\circ\text{N}$) (Fig. SI12). In this study, however, the data was not enough to clarify the transport route in more detail.

Fig. 9 shows the latitudinal distribution of the $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratio, which was decay-corrected to 11 March, 2011. The $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratio ranged from 0.1 to 0.72, and the ratios at Ogasawara region and in the ECS were almost the same as those in the SOJ (Fig. 9a). Considering that the activity ratio of $^{134}\text{Cs}/^{137}\text{Cs}$ that originated from the FNPP1 accident was almost 1 (Busseler et al., 2011), variations in the ratio indicate that FNPP1-labelled seawater mixed with global fallout-labelled seawater (no contamination by FNPP1- ^{137}Cs) during transport. The highest activity concentration ratio (0.72), which was found near Kagoshima, was a signal of the larger contributions of the FNPP1-radiocaesium. In contrast, a relatively low $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratio in the region between 30 and 32°N suggests the small contribution of FNPP1-radiocaesium at the Pacific Ocean site. In the horizontal distributions in the ECS, the ^{137}Cs activity concentrations tended to increase northward (Fig. SI12). The ^{137}Cs activity concentrations measured at Okinawa located in the southern ECS (<http://search.kankyo-hoshano.go.jp/servlet/search.top?pageSID=113836570>), were approximately 1.7 ± 0.47 (~ 2.8) Bq m^{-3} , which are slightly higher rather than those measured before the FNPP1 accident (Fig. SI13). This proves that less FNPP1 radiocaesium was transported in the southern ECS than in the northern ECS as discussed above.

4.2. Advection and vertical mixing of FNPP1- ^{137}Cs in the SOJ

In this study, we revealed that FNPP1- ^{137}Cs entered the SOJ via the ECS; then, FNPP1- ^{137}Cs was transported northward with the TWC. In the SOJ, a time lag of the propagation of FNPP1-radiocaesium of approximately one year was observed (Fig. 7). Based on measurements of phosphate, one of the dominant seawater nutrients, Kodama et al. (2016) revealed that the phosphate concentrations in surface seawater during winter were significantly positively correlated with the concentrations in the saline ECS seawater in the preceding summer, and the surface water of the southern SOJ was almost entirely replaced by the ECS seawater during May–October. They also suggested that the transport of water-soluble constituents from the ECS to the SOJ takes at least approximately 0.5 years. The propagation of FNPP1-radiocaesium in the SOJ was consistent with the propagation time scale of nutrients concentration change from the ECS to the SOJ (Kodama et al., 2016).

As shown in Fig. 6e, the ^{137}Cs activity concentrations at station 105-11, located along the western TWC in the SOJ, were maximum at the surface and gradually decreased with increasing depth. This vertical distribution is different from those in the NPSJ (Fig. 6a) and ECS (Fig. 6c). Particularly, the subsurface peak observed in the NPSJ and ECS did not appear at station 105-11. At station 105-11, most of the ^{137}Cs existed in seawater with a potential density of $25.7\text{--}27.3$ kg m^{-3} (Fig. 6f), which was located in a higher potential density layer than that in the NPSJ and ECS. A similar vertical distribution was also observed at the western coast of the Japanese Islands along the eastern TWC (Fig. SI11). These distributions were due to cooling in the surface layer after water was transported from the Tsushima Strait. Physical processes such as the convergence and subduction of surface water inside the eddies are important mechanisms of downward transport of radiocaesium (Miyao et al., 1998; Budyansky et al., 2015). There is a possibility of the seasonality of downward transport of radiocaesium. As expected, the past global fallout- ^{137}Cs would be already penetrated and accumulated in the deeper layers of the SOJ.

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4.4. Mass balance of FNPP1- ^{137}Cs in the SOJ

As for the most reliable estimate, the amount of FNPP1- ^{137}Cs deposited in the North Pacific Ocean via atmospheric release was $11.7\text{--}14.8$ PBq (Aoyama et al., 2016b; Tsubono et al., 2016) and the amount of FNPP1- ^{137}Cs directly released to

the Ocean was 3.5 ± 0.7 PBq (Tsumune et al., 2012, 2013). The FNPP1- ^{137}Cs inventory in the North Pacific Ocean was estimated to be 15.2-18.3 PBq (Aoyama et al., 2016b), 15.3 ± 2.6 PBq by optimum interpolation analysis (Inomata et al., 2016), and 16.1 ± 1.4 PBq by model simulation (Tsubono et al., 2016). According to the estimation by Kaeriyama et al. (2014), the ^{134}Cs amount was approximately 4.2 ± 1.1 PBq in the STMW in 2012. Taking into account this estimation by Kaeriyama et al. (2014), the FNPP1- ^{137}Cs transported into the SOJ from 2012 to 2016 accounted for 5.0 % of the FNPP1- ^{137}Cs inventory in the STMW. Of this, approximately 60-65 % of the total inflow of FNPP1- ^{137}Cs in the SOJ was transported in 2015 and 2016. After entering the SOJ, the FNPP1- ^{137}Cs followed a divergent path. The amounts of FNPP1- ^{137}Cs transported eastward (2.6 % of the FNPP1- ^{137}Cs inventory in the STMW) and westward (2.1 % of the FNPP1- ^{137}Cs inventory in the STMW) along the TWC were similar or slightly larger along the eastern TWC. Of the FNPP1- ^{137}Cs transported in the SOJ, approximately 43 % returned to the North Pacific Ocean through the Tsugaru Strait during 2012 to 2016.

Since 5 % of the FNPP1- ^{137}Cs in the STMW was transported into the SOJ, 95 % of that amount might remain in the STMW. Based on long-term measurements of the ^{137}Cs activity concentration adjacent to the FNPP1, Tsumune et al. (2017) estimated that the direct release rate of FNPP1- ^{137}Cs was 2.2×10^{14} Bq day $^{-1}$ until November 2011, after which it decreased exponentially with time to 3.9×10^9 Bq day $^{-1}$ on 26 October 2015. Assuming that the rate of decrease in the ^{137}Cs activity concentrations remained the same from 26 October 2015 to 31 December 2016, the total amount of ^{137}Cs directly released from the FNPP1 site during the period from 2012 to 2016 was estimated to be 0.03 PBq (Tsumune et al., 2016). However, not all of the directly released ^{137}Cs subducted into the STNW, 0.03 PBq corresponds to 0.7 % of ^{137}Cs inventory in the STMW. We can conclude that the FNPP1- ^{137}Cs in the STMW was decreasing during the study period.

Based on an analysis of historical data since 1950, the ^{137}Cs activity concentrations in the western North Pacific Ocean were exponentially decreased. However, the ^{137}Cs activity concentrations after 2000 have remained almost constant at 1.5-2 Bq m $^{-3}$ (Inomata et al., 2009), which indicates a source of ^{137}Cs . Considering that the contribution of global fallout- ^{137}Cs deposition amount was scarcely any after the 1970s, the ^{137}Cs activity concentrations in the seawater of the North Pacific Ocean were controlled by physical oceanographic processes such as advection and diffusion until before the FNPP1 accident (Inomata et al., 2009). Several studies reported that the subsurface maxima of ^{137}Cs derived from global fallout existed in layers corresponding to the potential densities of the STMW and CMW (e.g., Aoyama et al., 2008a). A possible explanation for the constant ^{137}Cs after 2000 is that the ^{137}Cs subducted in the STMW and CMW in the 1960s might have returned to the surface after a few decades. However, the increase in the FNPP1- ^{137}Cs activity concentrations in the SOJ derived from the FNPP1 accident began in 2012/2013. This would be existence of the faster transport route from the North Pacific Ocean to the SOJ along with the subtropical gyre.

5. Conclusion

The ^{137}Cs activity concentration in the Sea of Japan (SOJ) has been increasing since 2012/2013 and reached a maximum in 2015/2016 of approximately 3 Bq m $^{-3}$, which is above the pre-Fukushima accident level of the ^{137}Cs activity concentration, 1.5 Bq m $^{-3}$. The $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratios that ranged from 0.36 to 0.51 in the ECS and the SOJ are evidence that the increase in the ^{137}Cs activity concentration was derived from the Fukushima accident.

The increase in the FNPP1- ^{137}Cs activity concentrations was first observed in the subsurface layer with a density of approximately around 25.2 kg m $^{-3}$, corresponding to the STMW, in the southern part of the Kuroshio Current in the NPSJ in 2012-2013, and decreases in the ^{137}Cs activity concentrations in the NPSJ have been occurring since 2014. In the ECS, an increase in FNPP1- ^{137}Cs was observed in 2014, and decreases in the ^{137}Cs activity concentrations in the ECS have been observed since 2016. Note that the vertical distributions of the ^{137}Cs activity concentrations in the ECS were almost constant or contained a subsurface peak. These results indicated that there was a 1-2 year time lag between when the high ^{137}Cs activity concentrations occurred in the NPSJ and ECS. The increase in the ^{137}Cs activity concentrations in the SOJ began in

2012 and continued until 2016. A time lag in the propagation of FNPP1-¹³⁷Cs of approximately one year was observed in the SOJ. The similar ¹³⁴Cs/¹³⁷Cs activity ratios in the surface layers of the ECS, SOJ, and Ogasawara region support an identical source of radiocaesium in these regions.

5 The total amount of recirculated FNPP1-¹³⁷Cs from 2012 to 2016 was estimated to be 0.21±0.01 PBq, which corresponds to 5.0 % of the FNPP1-¹³⁷Cs in the STMW. Of this, the amounts of FNPP1-¹³⁷Cs transported to the west and east along the TWC were almost the same or slightly larger along the eastern TWC. Of the FNPP1-¹³⁷Cs transported into the SOJ, approximately 43 % (2.1 % of the total amount in STMW) passed through the Tsugaru Strait and back into the North Pacific Ocean. The FNPP1-¹³⁷Cs transported northward was estimated to 14 % of the amount transported into the SOJ (0.7 % of the total amount in STMW).

10 The most important result of this study was the determination that FNPP1-¹³⁷Cs was rapidly transported into the SOJ within several years after the FNPP1 accident.

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30 References

- Aoyama, M., and Hirose, K.: Artificial Radionuclides database in the Pacific Ocean:HAM database, *Sci. World J.*, 4, 200-2015, 2004.
- Aoyama, M., Hirose, K. and Igarashi, Y.: Re-construction and updating our understanding on the global weapons tests ¹³⁷Cs fallout, *J. Environ. Monit.*, 8, 431-438, 2006.
- 35 Aoyama, M., Hirose, K., Nemoto, K., Takatsuki, Y. and Tsumune D.: Water masses labeled with global fallout ¹³⁷Cs formed by subduction in the North Pacific, *Geophys. Res. Lett.*, 35, L01604, 2008a.
- Aoyama, M. and Hirose, K.: Radiometric determination of anthropogenic radionuclides in seawater. *Radioactivity in the Environment*, P. P. Pavel, Elsevier. Volume 11: 137-162, 2008b.
- Aoyama, M., Tsumune, D. and Hamajima, Y.: Distribution of ¹³⁷Cs and ¹³⁴Cs in the North Pacific Ocean: impacts of the
40 TEPCO Fukushima-Daiichi NPP accident, *J. Radioanal. Nucl. Chem.*, 1-5, 2012.
- Aoyama, M., Uematsu, M., Tsumune, D. and Hamajima, Y.: Surface pathway of radioactive plume of TEPCO Fukushima NPP1 released ¹³⁴Cs and ¹³⁷Cs, *Biogeosci.*, 10, 3067-3078, 2013.

- Aoyama, M., Hamajima, Y., Hult, M., Uematsu, M., Oka, E., Tsumune, D., Kumamoto, Y.: ^{134}Cs and ^{137}Cs in the North Pacific Ocean derived from the March 2011 TEPCO Fukushima Dai-ichi Nuclear Power Plant accident, Japan. Part one: surface pathway and vertical distributions, *J. Oceanogr.*, 72, 53-65, 2016a.
- Aoyama, M., Kajino, M., Tanaka, T.Y., Sekiyama, T.T., Tsumune, D., Tsubono, T., Hamajima, Y., Inomata, Y., Gamo.: ^{134}Cs and ^{137}Cs in the North Pacific Ocean derived from the March 2011 TEPCO Fukushima Dai-ichi Nuclear Power Plant accident, Japan. Part two: estimation of ^{134}Cs and ^{137}Cs inventories in the North Pacific Ocean, *J. Oceanogr.*, 72, 67-76, 2016b.
- Aoyama, M., Hamajima, Y., Inomata, Y., Oka, E.: Recirculation of FNPP1-derived radiocaesium observed in winter 2015/2016 in coastal regions of Japan, *Applied Radi. Isotopes.*, 126, 83-87, 2017.
- Behrens, E.; Schwarzkopf, F. U.; Lubbecke, J. F.; Boning, C. W. Model simulations on the long-term dispersal of ^{137}Cs released into the Pacific Ocean off Fukushima. *Environ. Res. Lett.*, 7, 034004, 2012.
- Busseler, K., Aoyama, M., Fukasawa, M. Impacts of Fukushima Nuclear Power Plants on Marine Radioactivity, *Environ. Sci. Technology*, 45, 9931-9935, 2011.
- Buesseler, K., Dai, M., Aoyama, M., Benitez-Nelson, C., Charmasson, S., Higley, K., Maderich, V., Masque, P., Oughton, D., Smith, J. N.: Fukushima Daiichi-Derived Radionuclides in the Ocean: Transport, Fate, and Impacts, *Annu. Rev. Mar. Sci.*, 9, 1-31, 2016.
- Budyansky, M.V., V.A. Goryachev, D.D. Kaplunenko, V.B. Lobanov, S.V. Prants, A.F. Sergeev, N.V. Shlyk, M.Yu. Uleysky, Role of mesoscale eddies in transport of Fukushima-derived cesium isotopes in the ocean, *Deep Sea Research Part I: Oceanographic Research Papers*, Volume 96, 2015, Pages 15-27, <https://doi.org/10.1016/j.dsr.2014.09.007>.
- Fukudome, K., Yoon, J-H., Ostrovskii, A., Takikawa, T., and Han. I-S.: Seasonal volume transport variation in the Tsushima warm current through the Tsushima Straits from 10 Years of ADCP Observations, *J. Oceanogr.*, 66, 539–551, 2010.
- Hamajima, Y. and Komura, K.: Background components of Ge detectors in Ogoya underground laboratory, *Appl. Radiat. Isot.* 61, 179-183, 2004.
- Han, S., Hirose, N., Usui, N., Miyazawa, Y.: Multi-model ensemble estimation of volume transport through the straits of the East/Japan Sea, *Ocean Dynamic.*, 66, 59-76, 2016.
- Hirose, K.: Fukushima Daiichi Nuclear Plant accident: Atmospheric and oceanic impacts over the five years, *J. Environ. Radioactiv.*, 157, 113-130, 2016.
- Honda, M. C., Aono, T., Aoyama, M., Hamajima, Y., Kawakami, H., Kitamura, M., Masumoto, Y., Miyazawa, Y., Takigawa, M. and Saino, T. Dispersion of artificial caesium-134 and-137 in the western North Pacific one month after the Fukushima accident, *Geochemical J.*, 46, e1-e9, 2012.
- Inomata, Y., Aoyama, M., Hirose, K.: Record of surface ^{137}Cs concentrations in the global ocean using the HAM-global database, *J. Environ. Monitor.*, 11, 116–125, 2009.
- Inomata, Y., Aoyama, M., Tsumune, D., Motoi, T., Nakano, H.: Optimum interpolation analysis of basin-scale ^{137}Cs transport in surface seawater in the North Pacific Ocean, *J. Environ. Monitor.*, 14, 3146–3155, 2012.
- Inomata, Y., Aoyama, M., Tsubono, T., Tsumune, D., and Hirose, K.: Spatial and temporal distributions of ^{134}Cs and ^{137}Cs derived from the TEPCO Fukushima Daiichi Nuclear Power Plant accident in the North Pacific Ocean by using optimal interpolation analysis, *Environ. Sci.: Processes Impacts*, 18, 126-136, 2016.
- Inoue, M., Kofuji, H., Nagao, S., Yamamoto, M., Hamajima, Y., Yoshida, K., Fujimoto, K., Takada, T., Isoda, Y.: Lateral variation of ^{134}Cs and ^{137}Cs concentrations in surface seawater in and around the Japan Sea after the Fukushima Dai-ichi Nuclear Power Plant accident, *J. Environ., Radioact.*, 109, 45-51, 2012.
- Kaeriyama, H., Ambe, D., Shimizu, Y., Fujimoto, K., Ono, T., Yonezaki, S., Kato, Y., Matsunaga, H., Minami, H., Nakatsuka, S., and Watanabe, T.: Direct observation of ^{134}Cs and ^{137}Cs in surface seawater in the western and central North Pacific after the Fukushima Dai-ichi nuclear power plant accident, *Biogeosci.*, 10, 4287–4295, 2013.

- Kaeriyama, H., Shimizu, Y., Ambe, D., Masujima, M., Shigenobu, Y., Fujimoto, K., Ono, T., Nishiuchi, K., Taneda, T. and Kurogi, H.: Southwest intrusion of ^{134}Cs and ^{137}Cs derived from the Fukushima Dai-ichi Nuclear Power Plant accident in the Western North Pacific, *Environ. Sci. Technol.*, 48, 3120-3127, 2014.
- Kaeriyama, H., Shimizu, Y., Setou, T., Kumamoto, Y., Okazaki, M., Ambe, D., and Ono, T.: Intrusion of Fukushima-derived radiocaesium into subsurface water due to formation of mode waters in the North Pacific, *Sci. Rep.*, 6, 22010, DOI: 10.1038/srep22010, 2016.
- Kamidaira, Y., Uchiyama, Y., Kawamura, H., Kobayashi, T., Furuno, A.: Submesoscale Mixing on Initial Dilution of Radionuclides Released From the Fukushima Daiichi Nuclear Power Plant, *J. Geophys. Res.*, DOI:10.1002/2017JV013359, 2018.
- 10 Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2011.
Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2012.
Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2013.
Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2014.
Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2015.
- 15 Korea Institute of Nuclear Safety: Marine Environmental Radioactivity Survey, 2016.
Kumamoto, Y., Murata, A., Kawano, T. and Aoyama, M.: Fukushima-derived radiocesium in the northwestern Pacific Ocean in February 2012, *Appl. Radiat. Isoto.*, 81, 335-339, 2013.
Kumamoto, Y., Aoyama, M., Hamajima, Y., Aono, T., Kouketsu, S., Murata, A. and Kawano, T.: Southward spreading of the Fukushima-derived radiocesium across the Kuroshio Extension in the North Pacific, *Sci. Rep.*, 4, 2014.
- 20 Kumamoto, Y., Aoyama, M., Hamajima, Y., Murata, and A., Kawano, T.: Impact of Fukushima-derived radiocesium in the western North Pacific Ocean about ten month after the Fukushima Dai-ichi nuclear power plant accident, *J. Environ. Radioactiv.*, 140, 114-122, 2015.
Kumamoto, Y., Aoyama, M., Hamajima, Y., Nagai, H., Yamagata, T., Kawai, Y., Oka, E., Yamaguchi, A., Imai, K., and Murata, A.: Fukushima-derived radiocesium in the western North Pacific in 2014, *J. Radioanal. Nucl. Chem.*, 311, 1209-25
1217, 2017.
- Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2011.
Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2012.
Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2013.
Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2014.
- 30 Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2015.
Marine Ecology Research Institute: Radiation level survey in marine organizations and their environment, 2016.
Menard, H. W. and Smith, M.: Hypsometry of ocean basin provinces, *J. Geophys. Res.*, 71, 4305-4325.
Masuzawa, J.: Subtropical Mode Water, *Deep-Sea Res.*, 16, 463-472, 1969.
Miyao, T., Hirose, K., Aoyama, M., Igarashi, Y.: Temporal Variation of ^{137}Cs and $^{239,240}\text{Pu}$ in the Sea of Japan, *J. Environ. Radioactiv.*, 40, 239-250, 1998.
- 35 Nishida, Y., Kanomata, I., Tanaka, I., Sato, S., Takahashi, S., Matsubara, H.: Seasonal and interannual variations of the volume transport through the Tsugaru Strait, *Oceanography in Japan*, 12, 487-499, 2003.
Oka, E.: Seasonal and interannual variation of North Pacific Subtropical Mode Water in 2003–2006, *J. Oceanogr.* 65, 151-164, 2009.
- 40 Oka, E., Qiu, B., 2012. Progress of North Pacific mode water research in the past decade. *J. Oceanogr.* 68, 5–20, <http://dx.doi.org/10.1007/s10872-011-0032-5>, 2012.
Oka, E., Qiu, B., Kouketsu, S., Uehara, K., Suga, T., 2012. Decadal seesaw of the Central and Subtropical Mode Water formation associated with the Kuroshio Extension variability. *J. Oceanogr.* 68, 355–360.

Prants, S.V., M.V. Budyansky, V.I. Ponomarev, M.Yu. Uleysky, P.A. Fayman. Lagrangian analysis of the vertical structure of eddies simulated in the Japan Basin of the Japan/East Sea. *Ocean Modelling*. V.86 pp.128-140 (2015). <http://dx.doi.org/10.1016/j.ocemod.2014.12.010>.

Rossi, V.; Van Sebille, E.; Sen Gupta, A.; Garçon, V.; England, M. H. Multi-decadal projections of surface and interior pathways of the Fukushima Cesium-137 radioactive plume. *Deep Sea Res., Part I* 2013, 80, 37–46.

Rossi, V.; Van Sebille, E.; Sen Gupta, A.; Garçon, V.; England, M. H. Corrigendum to “Multi-decadal projections of surface and interior pathways of the Fukushima cesium-137 radioactive plume” [*Deep-Sea Res.* I80(2013)37–46], *Deep Sea Res., Part I* 2014, 93, 162–164.

Tsubono, T., Misumi, K., Tsumune, D., Bryan, F.O., Hirose, K., and Aoyama, M.: Evaluation of radioactive cesium impact from atmospheric deposition and direct release fluxes into the North Pacific from the Fukushima Daiichi nuclear power plant, *Deep Sea Res. Part I: Oceano. Res. Papers*, 115, 10-21, 2016.

Tsumune, D., Tsubono, T., Aoyama, M., and Hirose, K.: Distribution of oceanic ^{137}Cs from the Fukushima Dai-ichi Nuclear Power Plant simulated numerically by a regional ocean model, *J. Environ. Radioactiv.*, 111, 100–108, 2012.

Tsumune, D., Tsubono, T., Aoyama, M., Uematsu, M., Misumi, K., Maeda, Y., Yoshida, Y., and Hayami, H.: One-year, regional-scale simulation of ^{137}Cs radioactivity in the ocean following the Fukushima Daiichi Nuclear Power Plant accident, *Biogeosci.*, 10, 5601-5617, 2013.

Tsumune, F., Aoyama, M., Hirose, K., Tsubono, T., Misumi, K., Tateda, Y.: Estimation of direct release rate of ^{137}Cs , ^{90}Sr and ^3H from the Fukushima Dai-ichi Nuclear Power Plant for four-and-a-half years, in: *Proceedings of the 4th International Conference on Environmental Radioactivity*, Vilnius, Lithuania, 29 May-2 June 2017, 24-28, 2017.

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Figure 1: Location of the sampling points after the FNPP1 accident. Large black circles are sites monitored by the Japanese government. Blue circles are sites monitored by the Korean government. Black triangles are sites with measured vertical profiles. Circle colors corresponds to the ^{137}Cs activity concentrations measured in 2011. Open circles are sites measured after the FNPP1-accident. The area around Japan was divided into 3 regions: the SOJ, ECS, and NPSJ (<141.5°E). The locations of monitoring stations are also plotted in the upper Figure.

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Figure 2: Temporal variations in the ^{137}Cs activity concentrations in surface seawater in the Sea of Japan during the period from 1960 to 2016. Circles indicate measured values, and red circles indicate data measured after the accident (2011 March 11). Black circles indicate the 0.5-year average value. Standard deviations of data were removed to clearly show the temporal variation.

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Figure 3: Temporal variations in the FNPP1- ^{137}Cs activity in the surface seawater at the monitoring sites in Japan after 2011. FNPP1- ^{137}Cs : (a) Tomari, Aomori, (b) Niigata, Ishikawa, Fukui, Shimane, (c) Saga, Kagoshima, 314-01, and (d) 105-11.

Figure 4: Temporal variations in the $^{134}\text{Cs}/^{137}\text{Cs}$ activity concentration ratio in the surface seawater at the monitoring stations in Japan from Jan 2016 to Jun 2016. (a) Tomari, Aomori, (b) Niigata, Ishikawa, Fukui, Shimane, and (c) Saga, Kagoshima.

35 Figure 5: Temporal variations in the ^{137}Cs activity concentrations at stations (a) 314-01 and (b) 105-11.

Figure 6: Vertical distributions of the ^{137}Cs activity concentrations over the (a) depth profile in the NPSJ, (b) potential density profile in the NPSJ, (c) depth profile in the ECS, (d) potential density profile in the ECS, (e) depth profile at 105-01 along the WTWC, and (f) potential density profile at 105-01 along the western TWC. Colour indicates the collection time (year).

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Figure 7: Hovmoller diagrams of the ^{137}Cs activity concentrations at a potential density along with an eastern TWC at a potential density of $25.2 \pm 0.5 \text{ kg m}^{-3}$. The ECS stations described on the x-axis are Kagoshima and Saga stations. Southwestern SOJ includes the monitoring stations Shimane, Fukui, Ishikawa, and Niigata. The northwestern SOJ includes the monitoring stations Aomori and Tomari. Color indicates the ^{137}Cs activity concentrations (Bq m^{-3}).

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Figure 8: Schematic diagram of the FNPP1- ^{137}Cs transport in the North Pacific Ocean. The bold line indicates the Kuroshio pathway. The thin lines indicate the flow pathway of the Tsushima Warm Current. The circle with red dotted line are STMW formation area. The dotted green circle indicates the CMW formation area. Unfilled blue arrows indicate the FNPP1- ^{137}Cs transport route deduced in this study. The red circles indicate the entrance region in the ECS. The estimated accumulated

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flux during the period from 2012 to 2016 at each section is shown in the parenthesis. The inventory in the STMW (*) was deduced by Kaeriyama et al. (2016).

5 Figure 9: Latitudinal and horizontal distributions of the $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratios measured at the coastal sites of the SOJ and ECS in 2015-2016. The values were radioactive decay corrected to 11th March, 2011. The data measured in the Ogasawara area (red circles in (b)) were also added. (a) latitudinal distribution, (b) horizontal distribution.

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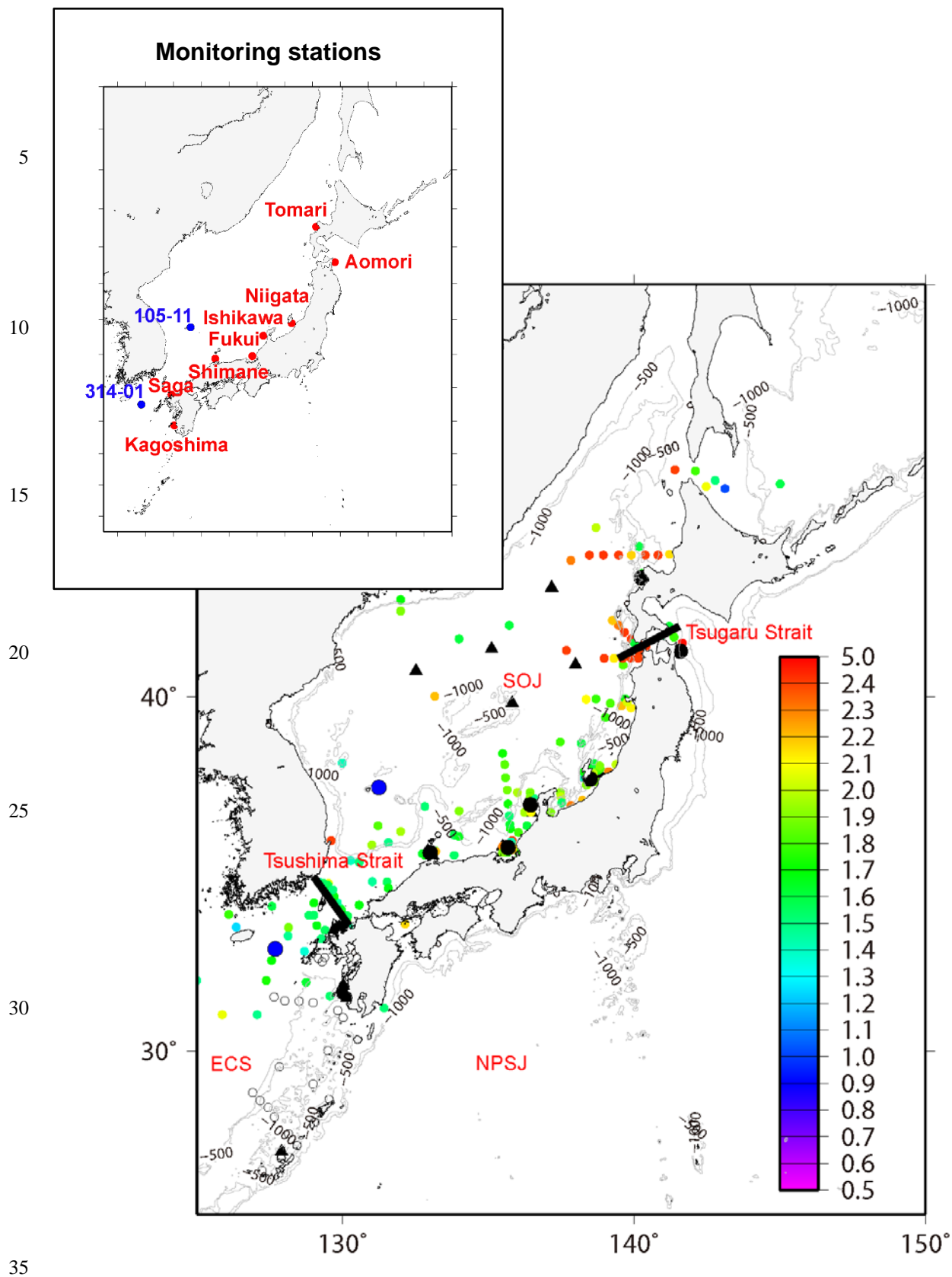


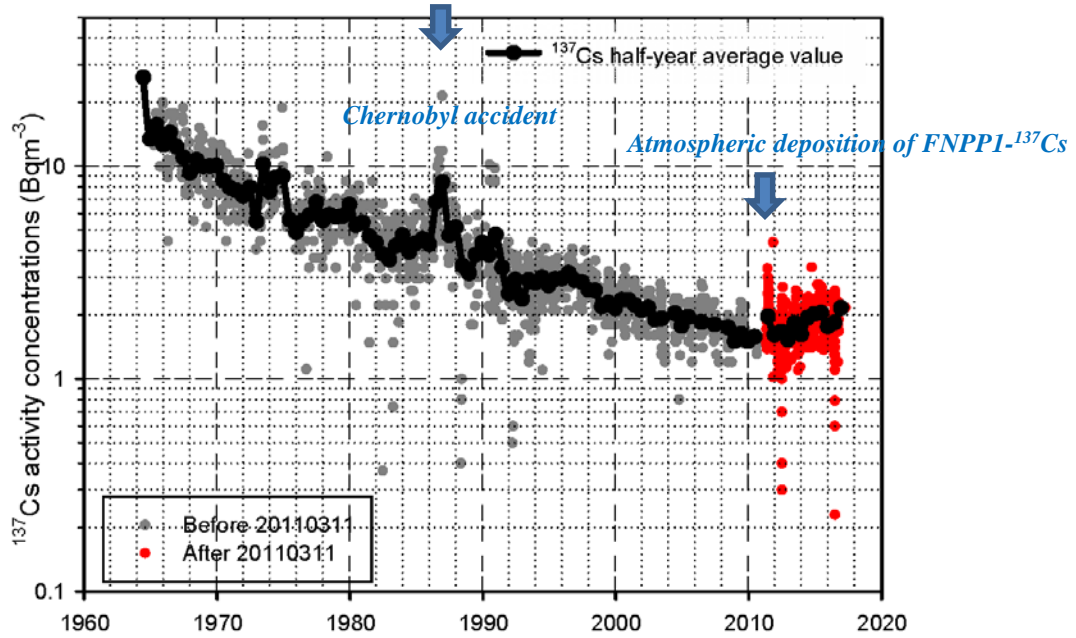
Figure 1: Location of the sampling points after the FNPP1 accident. Large black circles are sites monitored by the Japanese government. Blue circles are sites monitored by the Korean government. Black triangles are sites with measured vertical profiles. Circle colours correspond to the ^{137}Cs activity concentrations measured in 2011. Open circles are sites measured after the FNPP1 accident. The area around Japan was divided into 3 regions: the SOJ, ECS, and NPSJ ($<141.5^\circ\text{E}$). The locations of monitoring stations are also plotted in the upper Figure.

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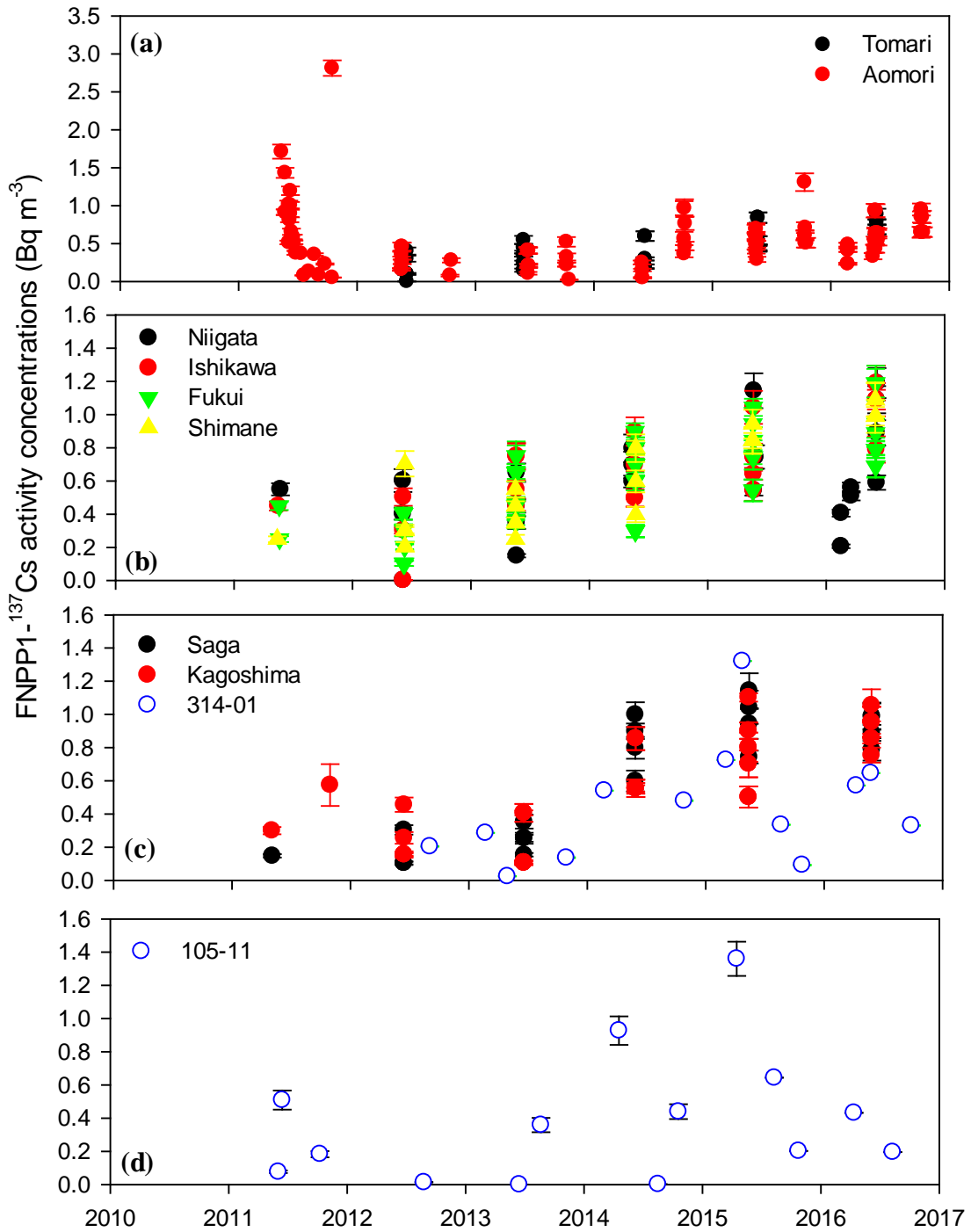
25 Figure 2: Temporal variations in the ^{137}Cs activity concentrations in surface seawater in the Sea of Japan during the period from 1960 to 2016. Circles indicate measured values, and red circles indicate data measured after the accident (2011 March 11). Black circles indicate the 0.5-year average value. Standard deviations of data were removed to clearly show the temporal variation.

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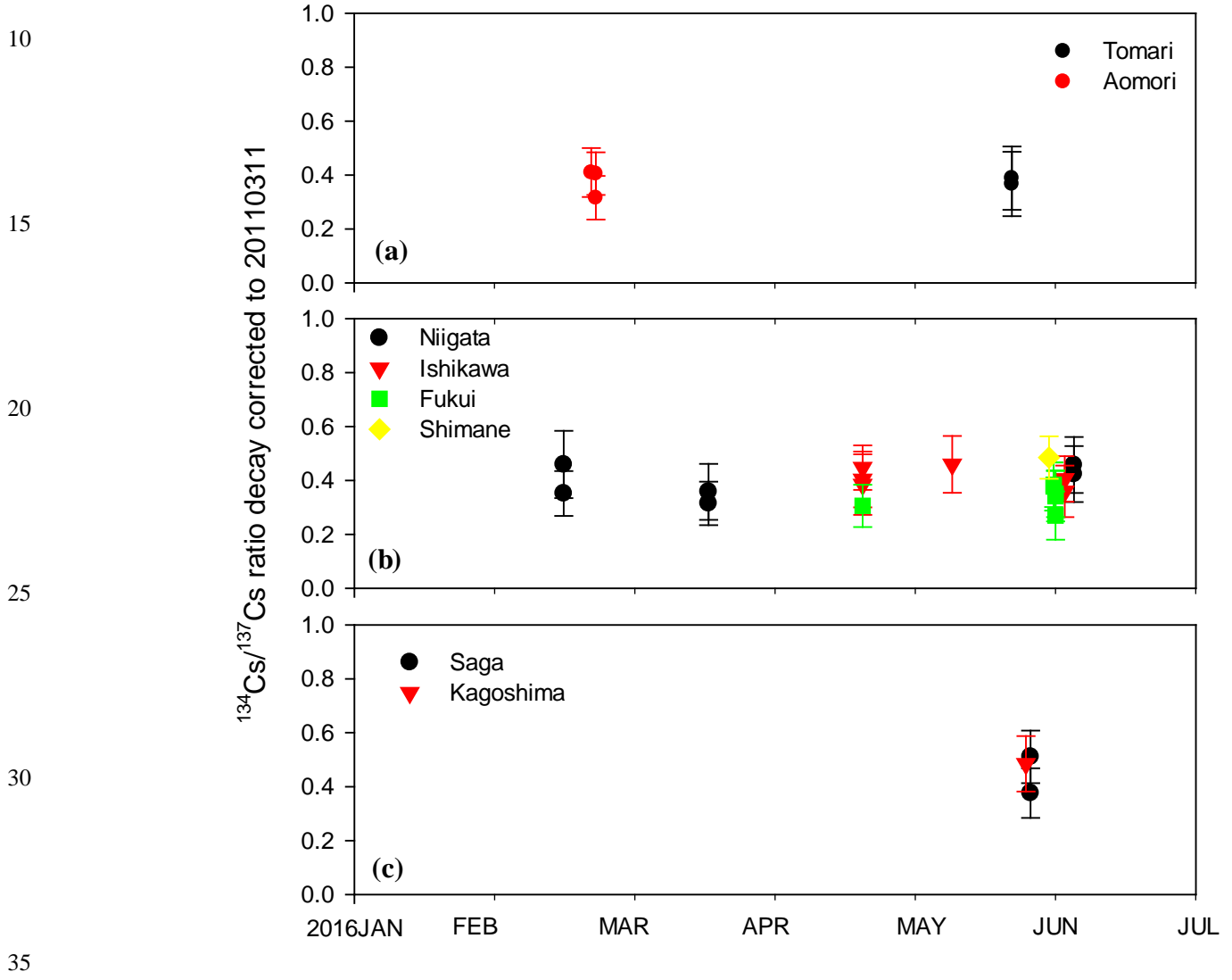
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45 Figure 3: Temporal variations in the FNPP1-¹³⁷Cs activity concentrations in the surface seawater at the monitoring stations in the SOJ after 2011. (a) Tomari, Aomori, (b) Niigata, Ishikawa, Fukui, Shimane, (c) Saga, Kagoshima, 314-01, and (d) 105-11.

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Figure 4: Temporal variations in the $^{134}\text{Cs}/^{137}\text{Cs}$ activity concentration ratio in the surface seawater at the monitoring sites in Japan from Jan 2016 to Jun 2016: (a) Tomari, Aomori, (b) Niigata, Ishikawa, Fukui, Shimane, and (c) Saga, Kagoshima.

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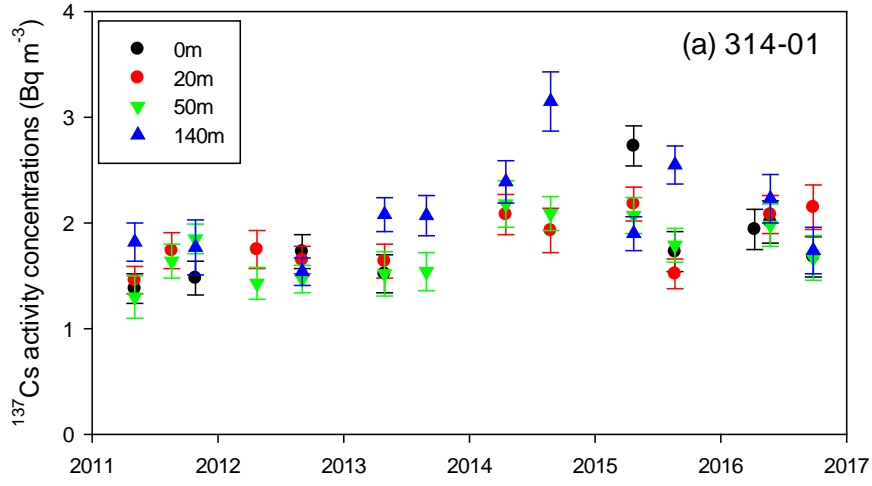
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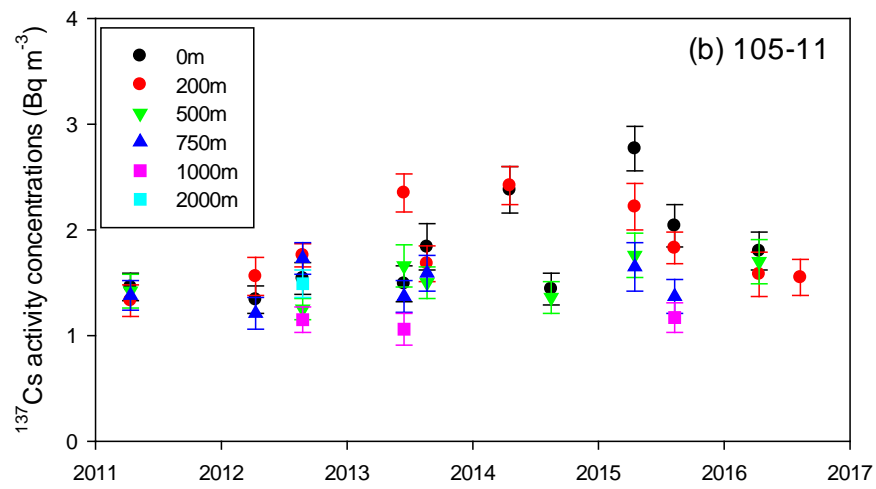
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Figure 5: Temporal variations in the ^{137}Cs activity concentrations at sites (a) 314-01 and (b) 105-11.

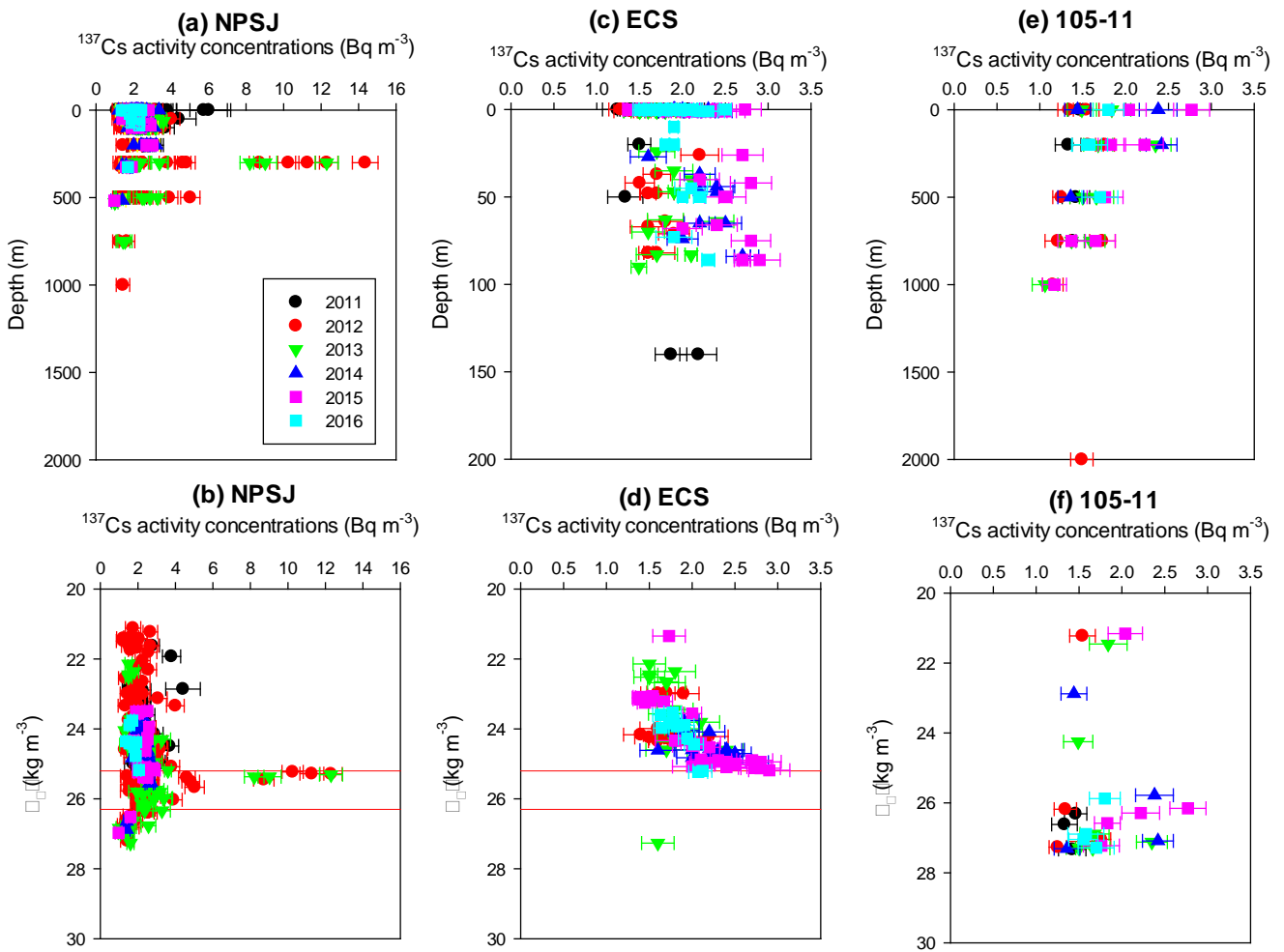


Figure 6: Vertical distributions of the ^{137}Cs activity concentrations over the (a) depth profile in the NPSJ, (b) potential density profile in the NPSJ, (c) depth profile in the ECS, (d) potential density profile in the ECS, (e) depth profile at 105-01 along the WTWC, and (f) potential density profile at 105-01 along the western TWC. Colour indicates the collection time (year). Horizontal red line shown in (b) and (d) means the boundary value of σ_θ of the STMW.

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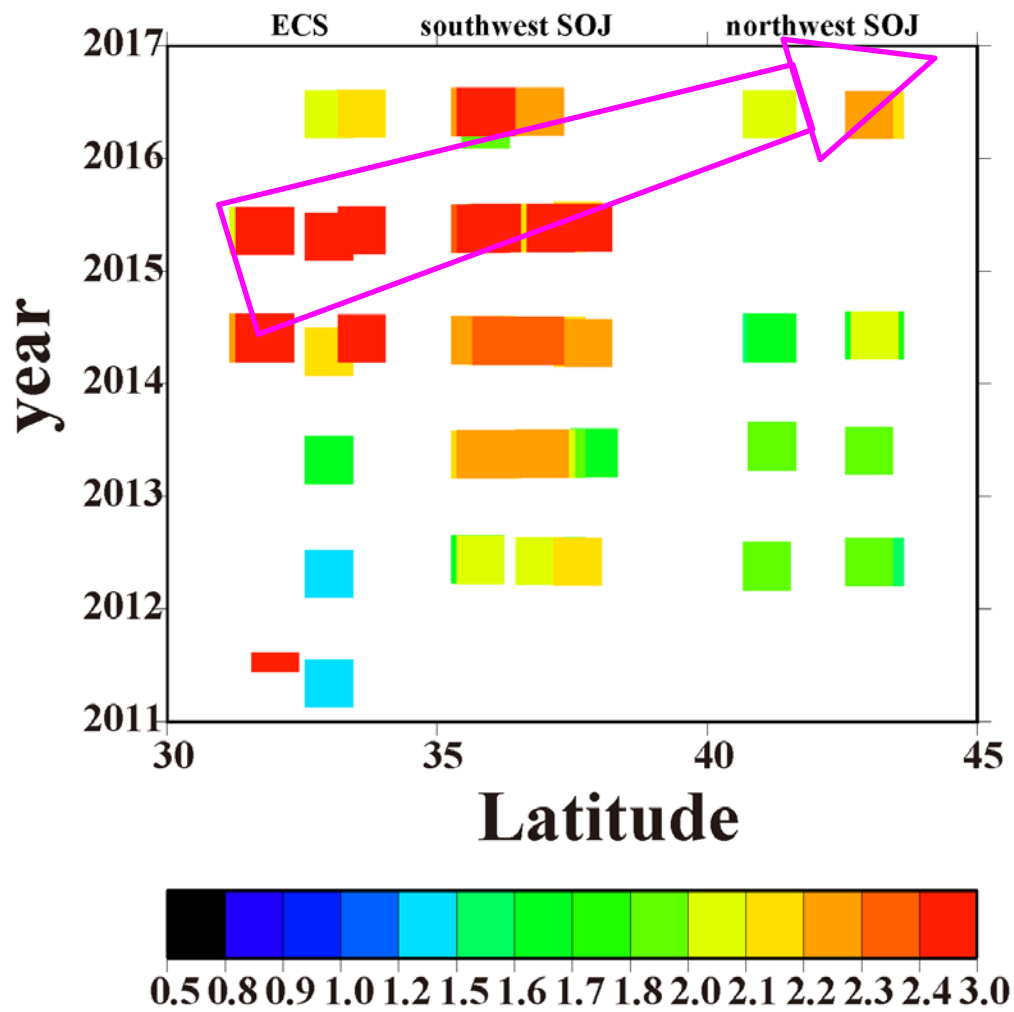


Figure 7: Hovmoller diagram of the ¹³⁷Cs activity concentrations at a potential density along with an eastern TWC at a potential density of $25.2 \pm 0.5 \text{ kg m}^{-3}$. The ECS stations described on the x-axis are Kagoshima and Saga stations. Southwestern SOJ includes the monitoring stations Shimane, Fukui, Ishikawa, and Niigata. The northeastern SOJ includes the monitoring stations Aomori and Tomari. Color indicates the ¹³⁷Cs activity concentrations (Bq m^{-3}). Arrow indicate the propagation of FNPP1-¹³⁷Cs transport from ECS to SOJ.

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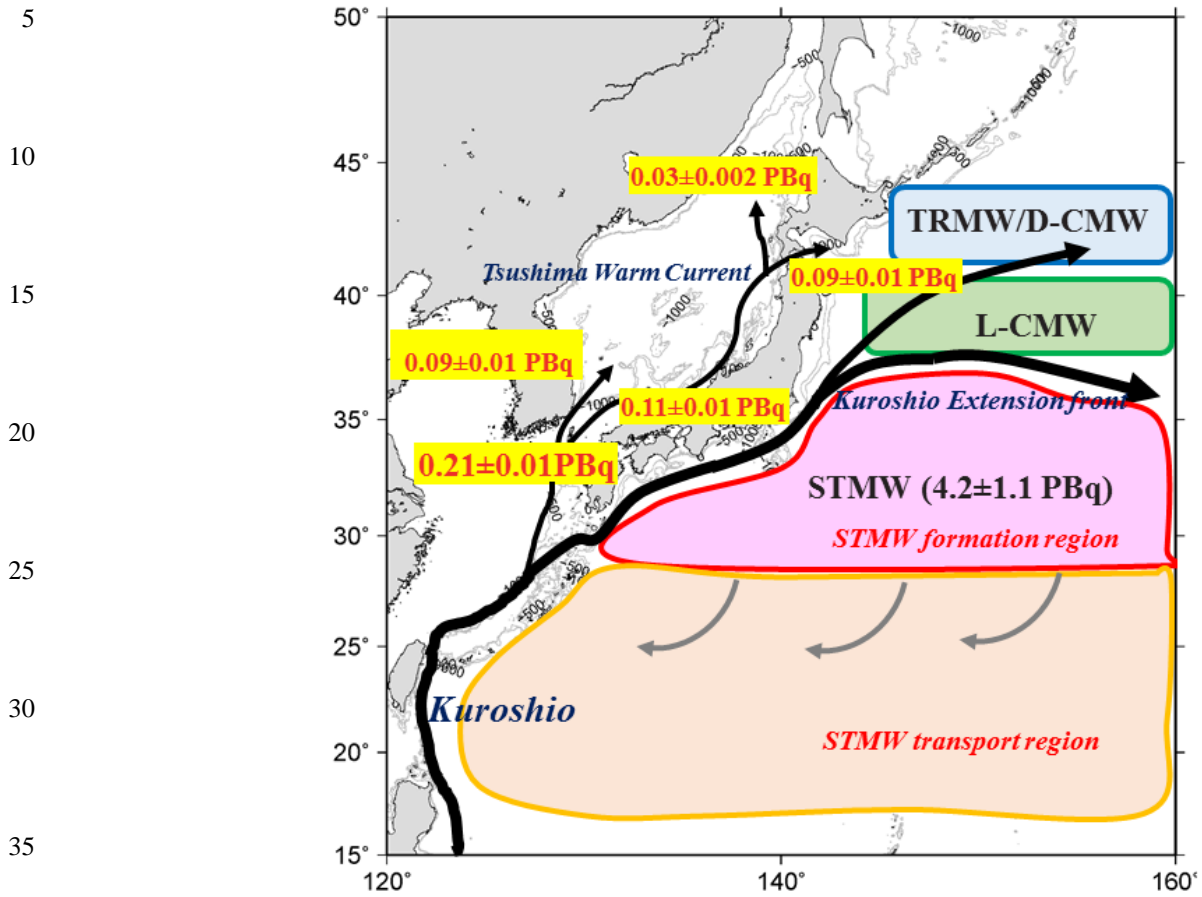


Figure 8: Schematic diagram of the FNPP1-¹³⁷Cs transport in the North Pacific Ocean. The bold line indicates the Kuroshio pathway. The thin black lines indicate the flow pathway of the Tsushima Warm Current. The pink region is STMW formation area. The green region is the L-CMW formation area, and the blue region indicates the TRMW and D-CMW formation area. The estimated accumulated flux during the period from 2012 to 2016 at each section is shown. The inventory in the STMW was deduced by Kaeriyama et al. (2016).

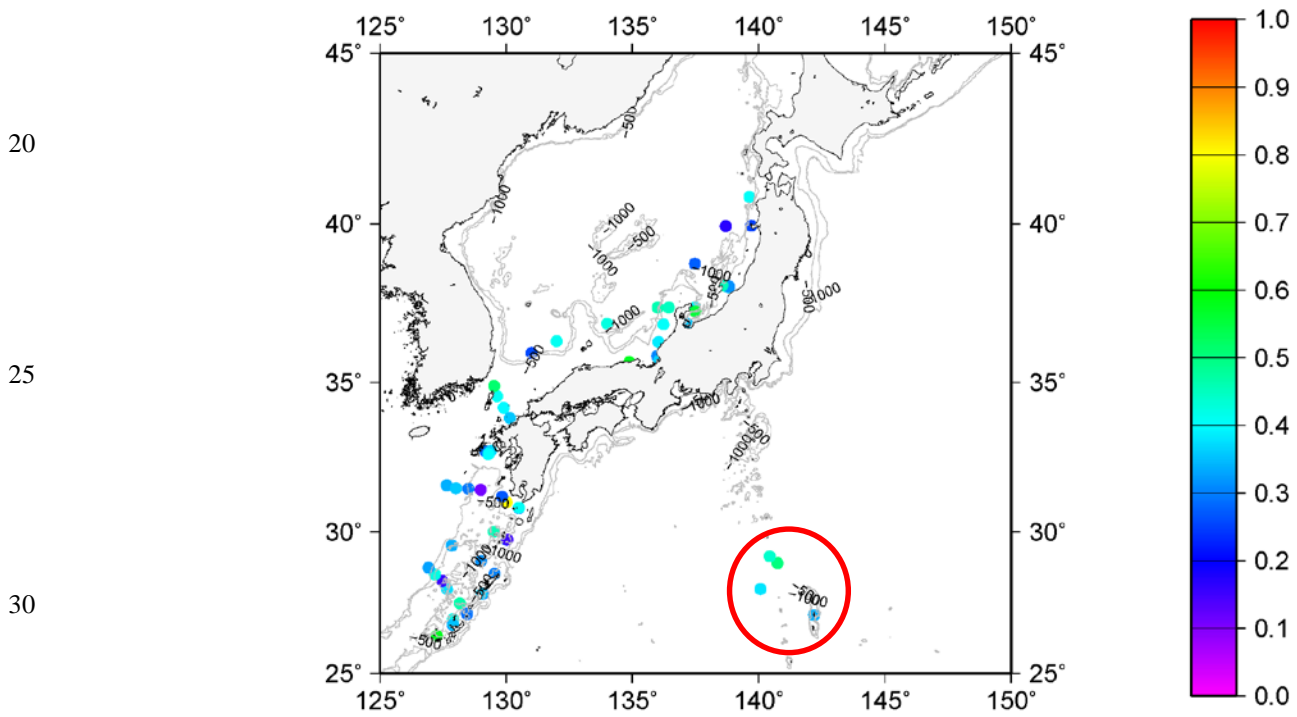
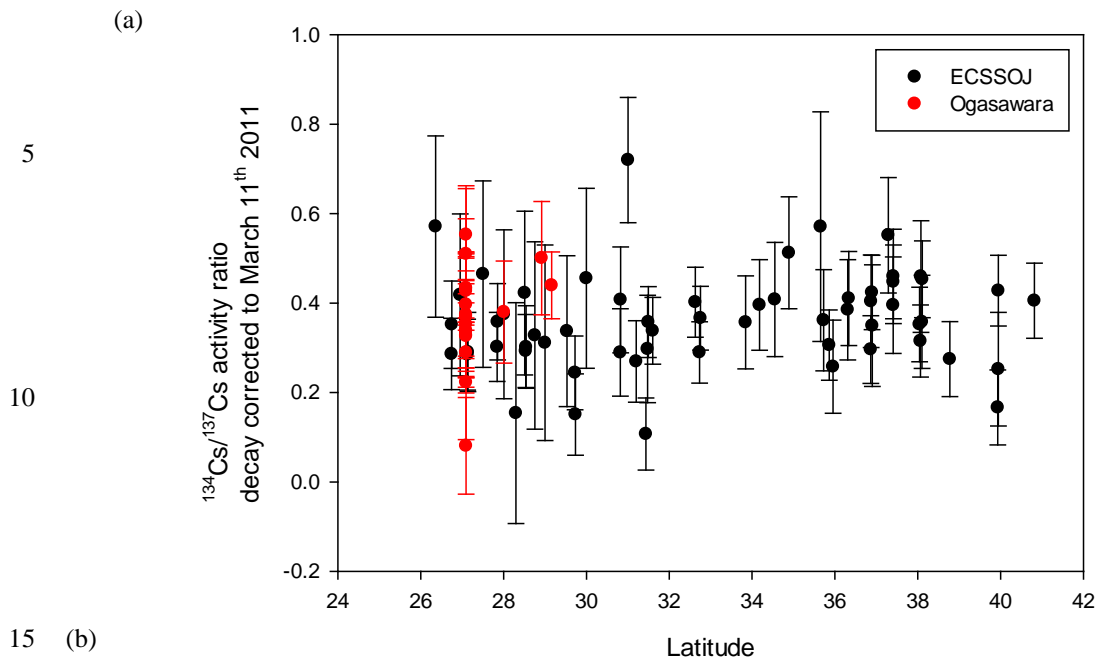


Figure 9: Latitudinal and horizontal distributions of the $^{134}\text{Cs}/^{137}\text{Cs}$ activity ratios measured at the coastal sites of the SOJ and ECS in 2015-2016. The values were radioactive decay corrected to 11th March 2011. The data measured in the Ogasawara area (red circles in (a)) were also added. (a) Latitudinal distribution, (b) horizontal distribution.

Table 1. Estimated transport amount of FNPP1-¹³⁷Cs in the monitoring site in the SOJ along with the Tsushima Warm Current.

| Flow | Station | Transport amount (PBq)* | Ratio against to STMW(%) | Ratio against to inflow into the SOJ (%) |
|---------------------------|-------------|------------------------------------|--------------------------|--|
| | | <i>Inflow to SOJ</i> | | |
| ECS to SOJ | Saga/304-01 | 0.21 ± 0.01 | 5.0 ± 2.3 | |
| | | <i>At Tsushima Strait</i> | | |
| Western TWC | 105-11 | 0.09 ± 0.01 | 2.1 ± 1.2 | 43 ± 10 |
| Eastern TWC | Shimane | 0.11 ± 0.01 | 2.6 ± 1.4 | 50 ± 7 |
| | | <i>Two branches of eastern TWC</i> | | |
| Outflow to Pacific Oceans | Aomori off | 0.09 ± 0.01 | 2.1 ± 1.2 | 43 ± 10 |
| Northward Transport | Tomari | 0.03 ± 0.002 | 0.7 ± 0.3 | 14 ± 2 |

5 *The value was decay corrected to 11th March 2011. ECS were estimated by using the average value of 314-01 and Saga monitoring sites. The transport amount (*) were estimated by sum of the duration from 2012 to 2016.

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