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Philippe Jean-Baptiste, Elise Fourré, Jean-Luc Charlou, Christopher R. German, Joël Radford-Knoery. Helium isotopes at the Rainbow hydrothermal site (Mid-Atlantic Ridge, 36°14'N). Earth and Planetary Science Letters, 2004, 221 (1-4), pp.325-335. 10.1016/S0012-821X(04)00094-9 . hal-03116204

HAL Id: hal-03116204 https://hal.science/hal-03116204v1

Submitted on 24 Jun 2021

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Helium isotopes at the Rainbow hydrothermal site (Mid-Atlantic Ridge, 36°14'N)

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Abstract: The 3He/4He ratio and helium concentration have been measured in the vent fluids and the dispersing plume of the Rainbow hydrothermal site, on the Mid-Atlantic Ridge (MAR). The 3He/4He ratio (7.51 Ra) and 3He end-member concentration (25 pmol/kg) are in the range of observed values elsewhere on mid-ocean ridges, pointing to the relative homogeneity of the upper mantle with respect to helium isotope geochemistry. 3He is linearly correlated with methane and manganese throughout the plume, with CH4/3He and Mn/3He ratios identical to those measured in the hot fluids. The bulk residence time of the plume in the rift valley estimated from the plume 3He budget is not, vert, similar20 days. The 3He flux transported by the plume, calculated from current-meter data, is 12.3±3 nmol/s, requiring a flux of 490 kg/s of high-temperature fluid. The scaling of the heat flux emitted by the Rainbow site to that of 3He, using the 3He/heat ratio measured in the hydrothermal fluids(9.3±2×10-18 mol/J), indicates a heat output of 1320 MW. With a regional spreading rate of not, vert, similar25 mm/year, we calculate that the annual 3He flux for this section of the MAR is of the order of 0.5±0.2 mmol per kilometre of ridge per millimetre of newly formed crust. This figure compares well with the flux calculated for the neighbouring Lucky Strike segment. Although the uncertainties remain large, both fluxes are not, vert, similar40-50% above the world average (0.33 mmol/km/mm), thus supporting earlier suggestions that the intensity of the hydrothermal activity south of the Azores is higher than expected from the regional spreading rate.

Keywords: helium; hydrothermal vents; isotopes; heat flux; Mid-Atlantic Ridge

1. Introduction

The Rainbow hydrothermal plume (Mid-Atlantic-Ridge, $36^{\circ}14^{\circ}N$) was first identified in 1994 from the detection of optical back-scatter anomalies [1]. In the year 1997, a detailed survey of the site was undertaken as part of the EU Amores project, including physical oceanography and geochemical study of the hydrothermal plume dispersal. Seafloor investigations where also carried out with the submersible *Nautile*, which led to the discovery and study of the active vents [2]. Here we report helium isotopes results for the high temperature fluids and the hydrothermal plume. In the following, the helium ${}^{3}\text{He}/{}^{4}\text{He}$ isotope ratio, R, is expressed either relative to the atmospheric ratio ($R_a = 1.38 \times 10^{-6}$) or using the delta notation, $\delta^{3}\text{He}$ (%) = (R/R_a -1)×100. With regards to hydrothermal processes, the helium isotopes represent a useful tracer in two different ways : first, owing to the clearly distinct ${}^{3}\text{He}/{}^{4}\text{He}$ signature of the different solid earth reservoirs, including the depleted upper mantle (${}^{3}\text{He}/{}^{4}\text{He}$ <0.1 R_a), helium isotope ratio is a good indicator of the source of the hydrothermal fluids. Second, due to its high concentration in vent fluids compared to seawater and to its chemical

2. Geological and hydrographic settings

The Rainbow site lies at $36^{\circ}13.8$ 'N – $33^{\circ}54.12$ 'W along the western flank of a topographic high (the Rainbow ridge) located at the western end of a non-transform discontinuity between the South AMAR and AMAR segments (fig.1). The hydrothermal site consists of a 100×200 m active vent field, between 2270 m and 2330 m depth, displaying extensive stockwork mineralization. Ten major groups of active black smokers emit hot temperature (T=365°C) acidic fluids. Unlike the previously discovered Menez Gwen and Lucky Strike hydrothermal sites south of the Azores, which are located in a basaltic environment, the Rainbow site is nested in serpentinized ultramafic rocks. This specific

inertness, ³He provides a faithful tracer of fluid dilution and plume dispersion.

geological environment strongly influences the chemistry [3] and gas content of the fluids, which show exceptionally high hydrogen concentration attributed to the serpentinization process [4].

Physical oceanography data, including CTD, LADCP and long-term current-meter moorings [5-6], as well as the spatial distribution of optical back-scatter anomalies, reveal that the hydrothermal plume is advected to the northeast along the rift valley of the AMAR segment as a single spatially and temporaly coherent structure, detectable over a distance of several tens of kilometers away from its source [6-7].

3. Sample collection and analytical methods

Water samples were taken in the rift valley on each side of the Rainbow ridge from a series of CTD-rosette vertical and "tow-yo" casts designed to obtain a 3-D picture of the hydrothermal plume. The water samples were stored in sealed copper tubes and returned to the laboratory for subsequent gas extraction and mass spectrometry measurement with a VG 3000 mass spectrometer using routine analysis procedures [8]. The accuracy in the δ^{3} He(%) excess is better than ±0.3, and the relative standard deviation of the absolute ⁴He concentration is 0.8%.

Helium isotope analyses were also performed on high temperature vent samples collected in 750-ml gas-tight titanium syringes. These sampling devices, developed at IFREMER for the french submersibles *Cyana* and *Nautile*, have been used routinely since the early 80's for hydrothermal gas studies. Leak tests carried out at various pressures in the high pressure facility at IFREMER have shown that these syringes, if properly maintained, are totally reliable to recover gassy samples such as those found in hot hydrothermal systems. The consistency of the Mn/³He and Mn/CH₄ ratios recorded in the plume and in the hot fluids (see fig.4) is a further evidence that the gas phase was adequately sampled. On deck, the fluid samples were transferred immediately in a specially designed evacuated degassing tank

ensuring 100% gas recovery. During the transfer procedure, the syringes were kept under water to avoid any contamination by atmospheric air. The degassing tank is a development of the prototype used in previous studies [9]. The main improvement consists in the addition of a membrane pump which ensures the complete transfer of the dissolved species from the degassing tank to a small stainless steel container through a drying column. The pressure in various parts of the equipment is monitored by miniature digital sensors. The measured pressure after the drying column makes it possible to determine the exact amount of dissolved gases [10-12]. In parallel to the stainless steel container, gas aliquots were also taken in small copper tubes tightly sealed with special clamps for helium isotopes analysis. In the laboratory, the copper tubes were directly connected to the inlet system of the mass spectrometer for isotopic analysis [8]. The uncertainty in the isotopic ratio is within $\pm 0.05 R_a$, with a relative error in the total helium concentration < 5%.

4. Results

4.1. Plume data

The vertical structure of the plume (fig. 2) shows pronounced δ^3 He anomalies in a near constant depth range between 2000 and 2300 meters, with increasing values up to 37% close to the vent site. This value is high compared to the background value of the deep Atlantic waters, around 5% at this latitude and depth [13-14]. The near-field plume exhibits two maxima at two different depths off-set by 40-50 meters (fig. 2), consistent with the nephelometry data [6]. This interval nearly corresponds to the depth interval spanned by the active vents on the Rainbow ridge so that this double plume is likely due to discrete contributions from the different vents. However, it has been observed in the TAG hydrothermal site [15] that a plume rise-height can vary by more than a hundred meters as a function of tidal cycling. Therefore, this double peak structure could also be due to a similar phenomenon. The stations southwest of the ridge show no δ^3 He anomaly (fig.3). This is in

agreement with the structure of the deep currents recorded by long term moorings (~ 400 days), which indicate a persistent northeastward advection of the plume [5]. The δ^3 He horizontal section at plume height (fig. 3) shows that the plume is detectable over a distance of about 30 km.

Throughout the dispersing plume, helium-3 is linearly correlated with manganese and methane (fig. 4), with slopes of $(110\pm10)\times10^6$ and $(131\pm16)\times10^6$ for Mn/³He and CH₄/³He ratios respectively.

4.2. Vent fluids data

The fluids are characterized by a rather uniform chemical composition indicating a single deep supply for the various vents [3-4]. The ${}^{3}\text{He}/{}^{4}\text{He}$ ratio, R, obtained from the slope of the ${}^{3}\text{He}-{}^{4}\text{He}$ diagram (Fig. 5a), is (7.51±0.02) R_a, which is within the 7-9 R_a range of MORB value. Helium concentrations (Table 1) are plotted in Fig. 5b against magnesium, used as an indicator of the dilution factor of the hydrothermal end-member with ambient seawater. The data define a helium (${}^{4}\text{He}$) end-member of 2.41±0.5 µmol/kg, corresponding to a ${}^{3}\text{He}$ concentration of 25±5 pmol/kg.

End-member hydrothermal tracer ratios in the hot fluids are $(94\pm20)\times10^6$ and $(108\pm37)\times10^6$ for Mn/³He and CH₄/³He ratio respectively. The ³He/heat ratio corresponding to the fluid output temperature of 365°C is $(9.3\pm2)\times10^{-18}$ mol/J.

5. Discussion

One of the main achievement of the Amores project was to deliver a coherent integrated data set of the physical and geochemical variables relevant to the study of an active venting site and its plume dispersal. In addition to presenting new data for hydrothermal tracers in a rather unusual (ultramafic) environment that permits comparison with other sites, this combined vent site/plume approach allows us 1) to compare plume tracer data to their

source composition, 2) to establish the ³He and thermal budget for a single vent site, and to put the results in perspective with respect to global hydrothermal fluxes.

5.1. Comparison of the vent fluid results with other hydrothermal sites

The ³He end-member (25±5 pmol/kg) lies in the range of previously studied hydrothermal sites on mid-ocean ridges (Table 2). Unlike major and trace elements andreactive gases such as CH₄, H₂, H₂S, whose concentrations in the hydrothermal fluids strongly depend on the geological nature of the substratum and local thermodynamical conditions, the mean ³He end-member concentration averaged over the MAR and EPR sites where helium isotopes data are available (Table 2) is relatively constant (17.1 pmol/kg, with a standard deviation of 5.4 pmol/kg, or 32% in relative terms). This appears thus to reflect primarily the relative homogeneity of the upper mantle source with respect to helium geochemistry. This is all the more true for the ³He/⁴He ratio in hot fluids which varies in a very narrow range of(8.0 ± 0.5) R_a (relative standard deviation $\approx 6\%$), in line with the range of most MORB ³He/⁴He data (7-9 R_a).

³He/heat ratios for mature hydrothermal vents - mean value = $(8.9 \pm 3.6) \times 10^{-18}$ mol/J (Table 2) - exhibit a higher variability than that of the helium end-member, presumably in connection with variations in the water-rock ratio which translate into a variable rate of extraction of heat and ³He from the hot basalt [16]. However, anomalously large shorttimescale variations of the ³He/heat ratio at a single vent site have been observed, related to magmatic events on the Juan de Fuca Ridge [16-19] or directly associated with fresh lava flows (see for instance fig. 2 in ref. [19]). With a value of $(9.3\pm2)\times10^{-18}$ mol/J, the Rainbow site exhibits a ³He/heat ratio typical of that observed for mature sites, characterized by a well established steady-state circulation.

Large variations in the $CH_4/{}^{3}He$ ratios are observed along mid-ocean ridges. Even for basaltic environments, the variability can be substantial : the $CH_4/{}^{3}He$ ratio at EPR-13°N and

EPR-21°N are $(1.7\pm0.4)\times10^6$ [20] and $(5.0\pm1.5)\times10^6$ [21] respectively while it is 65×10^6 at Lucky Strike and 85×10^6 at Menez Gwen on the MAR [11]. Mn/³He ratios also display a large variability: for example, the Mn/³He ratio of the EPR-13°N fluids is 13.7×10^6 [22] while it is more than 160×10^6 at the Galapagos Spreading Center [23]. Due to relative homogeneity of the ³He end-member in mid-ocean ridge hydrothermal systems, the variations in the CH₄/³He and Mn/³He ratios essentially reflect the variations in the end-member valueof these tracers. In the Rainbow fluids, CH₄ concentrations are indeed among the highest for abiogenic methane [4] and reflect the production of methane by the serpentinization reactions in the ultramafic substratum. Manganese and other transition metals concentrations are also unusually high [3]. A comprehensive discussion of the geochemistry of these tracers inrelation to the geological setting of the Rainbow site is available in [3] and [4].

5.2. ³*He and heat plume budgets*

Direct current observations, spanning the time when the plume samples were collected, were carried out with lowered ADCP. The results indicate that the local deep velocity field is strongly controlled by the topography [6] and show that the entire plume is advected across the Rainbow sill situated immediately NE of the vent site (see fig. 1 and fig. 3). LADCP measurements made in the saddle of the sill are in good agreement with those of the current meters moorings deployed during the cruise to monitor the water flow across valley section on the sill and recovered in 1998 (i.e., after ~ 400 days of operation). This long-term monitoring shows that below 2000 m, the direction and the intensity of the deep current advected over the sill are remarkably stable, corresponding to a mean water flux of

 $\Phi = (0.070\pm0.012) \times 10^6$ m³/sec [5]. This cross-valley section on the Rainbow sill is an ideal location for calculating the ³He flux advected with the plume. The best estimate of the mean δ^3 He integrated vertically across the sill between 2000 m and the sill depth is $\delta = (12.3 \pm 0.5) \%$ (fig. 6). Hence, the hydrothermal ³He flux Q_{3He} transported across the sill

is given by $Q_{3He} = \Phi \times [^{4}He] \times R_{a} \times (\delta - \delta_{BG})/100$, where $[^{4}He]$ is the helium concentration and δ_{BG} is the $\delta^{3}He$ background. With a mean measured helium concentration $[^{4}He] = 1.75 \pm 0.1 \ \mu mol/m^{3}$ and $\delta_{BG} = 5\%$, the ^{3}He flux transported by the plume is $Q_{3He} = (12.3 \pm 3) \times 10^{-9}$ mol/sec. Knowing the end-member vent fluid ^{3}He concentration, we calculate that a flux of 490 ± 220 kg/sec of high-temperature fluid is required to sustain the ^{3}He plume we observe.

The above ³He flux can also be combined with the ³He/heat ratio of the fluids to derive a thermal output of the Rainbow site of 1320 ± 600 MW. This calculation assumes that ³He/heat ratios in the focussed and diffuse vent fluids are identical. As a matter of fact, since both helium and heat are conservative tracers, they may locally occur in constant proportions. Thus we think that the above assumption is valid. This value is to compare with that for the Endeavour vent field on the Juan de Fuca Ridge, estimated around 1700 ± 1100 MW based on advective transport of heat [24].

5.3. Hydrothermal tracers dispersal

Most plume tracer data were obtained within a distance of ~ 20km from the vent site. This portion of the plume, which corresponds roughly to the isoline δ^3 He = 7.5%, contains 0.021 ³He moles in excess of the regional background (this figure was computed from the grid file used for the plot of figure 3, with a plume shape between 2000m and 2300m depth approximated by triangles). Dividing this plume ³He inventory by the ³He flux transported by the dispersing plume, $Q_{3He} = (12.3\pm 3)$ nmol/sec, leads to a bulk residence time of the plume in the rift valley of ~ 20 days. Over this period of time, the CH₄/³He and Mn/³He ratios calculated for the dispersing plume (fig. 4) and those measured in the vent fluids are identical, within the experimental uncertainties. These linear correlation among the tracers and the consistency of the ratios in the hot fluids and in the diluted plume show that, on this time scale, the different hydrothermal tracers behave conservatively with respect to one another in the plume and with respect to their source fluids. This is in agreement with available observations which show that the oxidation kinetics of Mn^{2+} and CH_4 are such that they behave conservatively close to the vents [25-26].

5.4. Comparison with global hydrothermal fluxes

The spatial pattern of ³He excess in the deep ocean strongly points to the prominent role played by the mid-ocean ridges with respect to ³He mantle degassing, with major water column δ^{3} He anomalies being centered above spreading centers in the East Pacific [27-28], in the Indian Ocean [29-30] and in the Atlantic [31]. To the first order, 3-D oceanic general circulation models make a good job at simulating the ³He distribution with the simple assumption that the ³He source intensity is linearly correlated to the spreading rate [32-33]. The slope of the correlation is ~330 ³He mol/km² of crustal spreading, translating to a world average ³He annual flux of 0.33 mmol per kilometer of ridge and per millimeter/year of full spreading rate (fig. 7). The section of the MAR to which the Rainbow site belongs has a spreading rate close to 25 mm/year [34]. In order to plot the Rainbow site in the diagram displayed in fig. 7, one has to define the length of ridge involved. While Lucky Strike and Menez Gwen venting sites, situated further north, are in segment centres, the Rainbow site occurs within a non-transform discontinuity between the South AMAR and the AMAR segment. Therefore, it cannot directly be affected a single segment length. On the other hand, surveys of hydrothermal plumes on the MAR 36-38°N have yielded evidence for an average spacing between adjacent vent-sites of 25-30 km along-axis [35]. Assigning this length scale to the Rainbow site leads to a ³He flux of 0.5 ± 0.2 mmol per kilometer of ridge per millimeter of spreading rate. This value is comparable to that for the Lucky Strike segment (0.45 mmol/km/mm) [36]. Both sites are $\sim 40-50\%$ above world average (0.33 mmol/km/mm), thus supporting earlier the suggestion [35] that hydrothermal activity along

this section of ridge axis close to the Azores Triple Junction is more intense than has been reported elsewhere on slow spreading ridges.

6. Conclusion

The main results of this helium isotopic study of the Rainbow hydrothermal site include the following :

1- The Rainbow site, although hosted in serpentinized diapir, i.e. a rather unusual geological environment compared to most already studied sites, displays a helium isotope geochemistry very similar to "classic" mid-ocean ridge sites.

2- From the compilation of mid-ocean ridge sites for which helium isotopes data are available, we come to the more general conclusion that, as far as helium geochemistry is concerned, the variability among hydrothermal sites is rather limited, with relative standard deviations of 34% and 6% for the ³He end-member concentration and ³He/⁴He ratio respectively. This result points to the relative homogeneity of the helium isotope composition of upper mantle, also noted in MORB studies.

3- The ³He budget leads to a plume residence time of ~ 20 days in the rift valley. On this time scale, ³He correlates linearly with methane and manganese hydrothermal tracers throughout the dispersing plume. Furthermore, $CH_4/^3$ He and $Mn/^3$ He ratios are identical in the

plume and in the hot fluids showing that the different hydrothermal tracers behave conservatively with respect to one another in the plume and with respect to their source fluids.

4- The ³He flux transported by the plume could be deduced from the measurement of the water transport across the sill which marks the entrance of the AMAR segment, allowing us to infer a thermal output of 1320 MW. Translated into a flux per km of ridge and per millimeter of spreading rate, this ³He flux (0.5 mmol/km/mm) is comparable to the flux calculated for the neighbouring Lucky Strike segment (0.45 mmol/km/mm). Although the

uncertainties are large, both fluxes are $\sim 40-50\%$ above the world average (0.33 mmol/km/mm), thus showing that the MAR south of the Azores discharges substantially more than its share of the global hydrothermal input to the ocean.

Acknowledgements

We would like to thank D. Pinti, M.D. Rudnicki, and one anonymous reviewer for their constructive comments on the manuscript. This research was undertaken as part of the Amores project supported by the European Commission (MAST program under contract MAS3-CT95-0040).

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Table captions

Table 1: Rainbow vents fluids helium isotopes data (the uncertainty in the ${}^{3}\text{He}/{}^{4}\text{He}$ ratio is within $\pm 0.05 \text{ R}_{a}$, with a relative error in the helium concentration <5%).

Table 2: Compilation of helium isotopes and heat data for hydrothermal vents from the Mid-Atlantic Ridge and East Pacific Rise.

Figure captions

Fig. 1: Bathymetric 3-D map of the Rainbow area.

Fig. 2: Selected δ^3 He vertical profiles along the dispersing plume (solid lines). The location of the stations (a to d) is shown in figure 3. δ^3 He vertical profiles at nearby stations outside the central rift (dashed lines) are also displayed to define the regional helium-3 background : NATS (1981) stations n°60 (36°25N/34°38W), n°61 (36°25N/33°67W),

 $n^{\circ}65$ (36°25N/30°25W) and $n^{\circ}66$ (36°25N/29°27W) [13], and TOPOGULF (1983) station $n^{\circ}83$ (40°00N/35°00W) [14]. (N.B. the δ^{3} He maximum in the upper 1000 meters reflects the transient signal produced by the radioactive decay of bomb tritium).

Fig. 3 : Horizontal distribution of the δ^{3} He (%) maximum associated with the Rainbow hydrothermal plume in the 2000-2300 m range (coordinates in decimal degree). Letters a to d indicate the location of the stations displayed in figure 2. The dashed line shows the position of the Rainbow sill, NE of the vent site. Darkly shaded regions are less than 2000 m.

Fig. 4 : Manganese (a) and methane (b) as a function of ${}^{3}\text{He}$ in the hydrothermal plume.

Fig. 5 : 3 He- 4 He relationship (a) and helium (4 He) concentration versus magnesium (b) in the vent fluids.

Fig. 6 : δ^{3} He (%) vertical section across the sill located NE of the vent site. Due to the lack of stations at the exact location of the sill, the section was reconstructed from the horizontal grid file used for plotting figure 3 (on the vertical, the shape of the profile in the range 2000-2300 m was approximated by a triangle).

Fig. 7 : ³He annual flux per kilometer of ridge as a function of spreading rate. The dashed line is the linear relationship used in GCM simulations [32-33]. Lucky Strike and Red Sea values

17

are calculated from reference [36] and [44]. The global average corresponds to the classic value of 4 ± 1 ³He atom/cm²/sec estimated by Harmon Craig and his colleagues thirty years ago [45].

Vent marker	Sample	Temperature (°C)	R/Ra	[⁴ He] (µmol/kg)	Mg (mmol/kg)
2	PL03-G2	-	6.80	0.22	45.5
3	PL04-D3	351	7.75	1.00	29.6
3	PL07-G2	351	7.18	2.04	11.2
4	PL03-D3	354	7.02	0.70	33.3
4	PL03-G3	354	5.71	0.23	48.3
4	PL05-EG	354	7.34	1.93	9.5
5	PL04-G2	362	6.73	0.12	46.1
5	PL09-D2	362	7.66	2.49	3.3
7	PL06-D2	-	7.61	1.55	3.7
9	PL10-D2	364	7.56	2.89	2.1
SW	deep water	3-4	1.05	0.0017	52.9

Table 1

Hydrothermal sites	³ He conc. (pmol/kg)	R/R _a	³ He/Heat (10 ⁻¹⁸ mol/J)	Reference
MAR				
Logatchev (14.45°N)	-	7.2	-	[37]
Snake Pit (23°N)	23.8 ± 1.6	8.4	12.4 ± 1.2	[9]
Snake Pit (23°N)	12.5 ± 1.5	7.9	6.5 ± 0.8	[38]
TAG (26°N)	20.1 ± 2.2	7.5	9.3 ± 1.1	[38]
TAG (26°N)	16.7 ± 1.4	8.2	7.7 ± 0.8	[39]
Broken Spur (29°N)	-	8.9	-	[37]
Rainbow (36°14N)	25.0 ± 5	7.51	9.3 ± 2	this work
Lucky Strike (37°17N)	10 ± 3.5	8.13	6.5 ± 1.2	[36]
Menez Gwen (37°50N)	20 ± 4	8.65	17.8 ± 3.5	[11]
EPR Galapagos EPR 21°N EPR 13°N EPR 11°N EPR 18°S Global average for mature sites standard deviation (σ)	$ \begin{array}{c} 12.4 \pm 2.3 \\ 21 \pm 7 \\ 17.9 \pm 8 \\ 9.2 \pm 3 \\ \begin{array}{c} 17.1 \\ (5.4) \end{array} $	7.8 7.8 7.6 8.32 8.3 8.0 (0.5)	5.2 ± 0.4 5.2 ± 0.9 11.0 ± 3.7 9.5 ± 4 6.2 ± 2 8.9 (3.6)	[40] [21] [41] [42] [43]
Juan de Fuca : - event Plume - associated deep plume - over fresh lava flows	- - -	7.9 7.9	3.2-4.3 14-47 22-75	[19] [19] [19]

Table 2





Figure 2



Figure 3



Figure 4



Figure 5



Figure 6



Figure 7