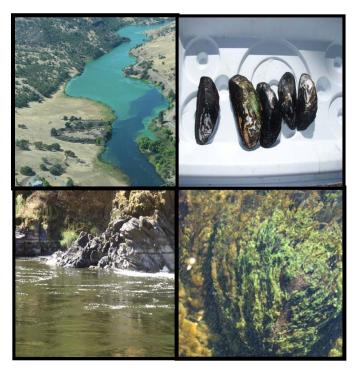
Technical Memorandum

Microcystin Bioaccumulation in Klamath River Freshwater Mussel Tissue: 2009 Results



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INTRODUCTION

Copco and Iron Gate Reservoirs (the lowermost projects of PacifiCorp's Klamath Hydropower Project-- KHP) experienced extensive blooms of toxigenic *Microcystis aeruginosa* (MSAE) from 2004-2009 (Kann and Corum 2009; 2010; Jacoby and Kann 2007). These blooms were associated with high levels of the cyanotoxin microcystin, a potent hepatotoxin capable of causing chronic liver damage and acting as a tumor promoter (Carmichael 1995; Chorus et al. 1999; Chorus 2001).

The results of the 2005-2009 sampling program demonstrated widespread and high abundance of toxigenic MSAE blooms in Copco and Iron Gate reservoirs and in the Klamath River downstream, exceeding World Health Organization Moderate Probability of Adverse Health Effect Levels for both cell density and toxin by 10 to over 1000 times.

In addition, bioaccumulation studies undertaken in 2007 and 2008 showed accumulation of microcystin toxin in muscle and/or liver tissues of yellow perch, hatchery salmon, and freshwater mussels (Mekebri et al. 2009; Kann 2008; Kanz 2008). Microcystin levels in biota exceeded public health threshold values for safe consumption (Kann 2008; OEHHA 2008). The following report summarizes the Karuk Tribe Department of Natural Resources and Yurok Tribe Environmental Program 2009 sampling program to evaluate microcystin bioaccumulation in freshwater mussels in the Klamath River.

METHODS

Station Location and Frequency

During the 2009 sampling season freshwater mussels were collected by the Karuk and Yurok Tribes at a variety of locations on the Klamath River (Figure 1 and Figure 2). Karuk Tribe stations included the Klamath River at the I5 Bridge (IB), Brown Bear river access (BB), Seiad Valley (SV), Happy Camp (HC), and Orleans (OR). The primary Yurok Tribe station was located on the Klamath River just upstream of Starwein Riffle (KA), as well as one sample collected on the Trinity River (TR) just upstream from the Klamath River confluence (Figure 2). The primary stations were generally sampled monthly from July through October, but TR was sampled only once in September, and BB was sampled one additional time in December (Figure 3).

Ambient concentrations for microcystin were collected in conjunction with the freshwater mussel samples, as well as at additional locations to evaluate bioaccumulation trends relative to ambient concentrations (Figure 3). Ambient data from the Karuk Tribe's 2009 public health monitoring program (Kann and Corum 2010) was also evaluated with respect to bioaccumulation trends. Ambient data for the Yurok Tribe station TG, which is close to the KA mussel collection station, were available on dates when mussels were collected and is also included in below analyses.



Figure 1. Location of Karuk Tribe freshwater mussel sampling locations, 2009.

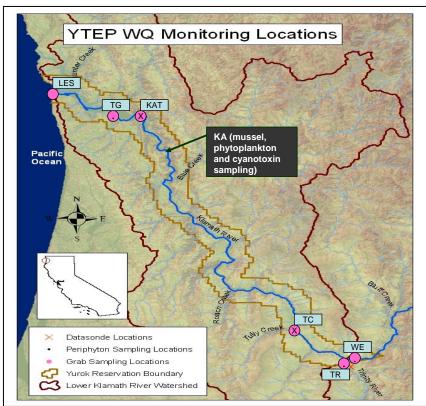


Figure 2. Location of Yurok Tribe freshwater mussel sampling locations, 2009. Note that mussel's were collected at stations KA and TR .

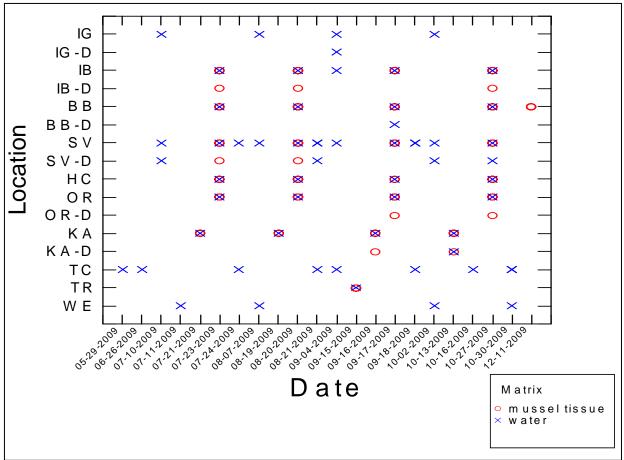


Figure 3. Freshwater mussel and ambient (water) samples collected from the Klamath River system in 2009. Stations ordered longitudinally top (upstream) to bottom (downstream). "D" denotes laboratory split.

Sample Collection and Lab Analysis

Klamath River freshwater mussel beds were sampled either via snorkeling or wading (if depth permitted) at the above locations (Figure 4). In general 5 to 6 individual mussels (whole *Gonidea angulata*) from each location were selected from the bed, wrapped in foil, and then frozen prior to shipment to the California Department of Fish and Wildlife Water Pollution Control Laboratory in Rancho Cordova, CA. Extraction of tissue samples included homogenization and sonification prior to using the LC-ESI-MS-MS method to determine the concentration of major microcystin congeners (Mekebri at al. 2009). Congeners analyzed included: MCY-RR, MCY-Desmethyl-RR (MCY-RRDM), MCY-LR, MCY-Desmethyl-LR (MCY-LRDM), MCY-YR, MCY-LA, MCY-LW, MCY-LF, and MCY-LY. In addition, the neurotoxins anatoxin-a, domoic acid, and okadaic acid were also analyzed.

Samples for ambient microcystin concentration were collected using the standard operating procedure (SOP) developed by the Klamath Blue-Green Algae Working Group (see Kann and Corum 2010 for details). These were analyzed as per the liquid sample extraction method in Mekebri et al. (2009). Methodology for additional ambient samples that were collected as part of the Karuk Tribe's public health monitoring program for cyanobacteria is outlined in Kann and Corum 2010.



Figure 4. Collection of Klamath River freshwater mussels, 2009.

Laboratory quality assurance consisting of split samples for both tissue and ambient microcystin toxin concentrations were also provided (Figure 3; denoted with a "D" after the station name). These data are contained in Appendix I, and generally showed good agreement between tissue duplicates (Figure 5).

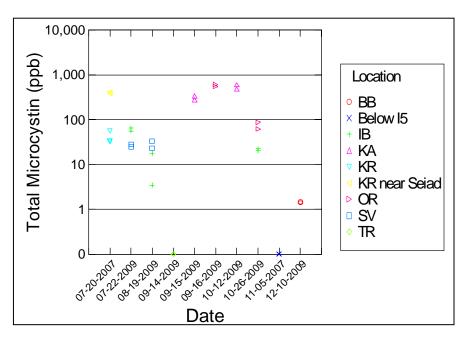


Figure 5. Comparison of duplicate mussel samples for stations in the Klamath River system collected in 2007 and 2009.

Comparison to Public Health Threshold Values

The following comparison of Klamath River microcystin tissue concentrations to public health guideline values is based on a recent comprehensive review of cyanobacterial toxin accumulation by Ibelings and Chorus (2007). Table 2 from Ibelings and Chorus (2007) entitled *"Tolerable doses to microcystin-LR in relation to frequency and duration of exposure"* is reproduced here:

Temporal pattern of exposure and ensuing Tolerable Intake (TI)	Assumptions	Tolerable Intake per kg	Tolerable Intake for a 10 kg child	Tolerable Intake for a 75 kg adult		ne value <u>l (μg kg⁻¹)</u> AF = 0.2
Acute TI	NOAEL ¹ of 250 μ g/kg and day, extrapolation factors of 100	2.5 µg per kg and single exposure	25 μg per single exposure	190 µg per single exposure	Adult: 1900, Child: 250	Adult: 380, Child: 50
Seasonal TDI	NOAEL of 0.4 µg/kg and day, extrapolation factors of 100 (Chorus and Bartram, 1999, adapted)	0.4 μg per kg and day	4 μg per day	30 µg per day	Adult: 300, Child: 40	Adult: 60, Child: 8
Lifetime TDI	NOAEL of 0.4 µg/kg and day, extrapolation factors of 100 and uncertainty factor of 10 (Chorus and Bartram, 1999)	0.04 μg per kg and day	0.4 μg per day	3 μg per day	Adult: 30, Child: 4.0 ²	Adult: 6, Child: 0.8 ²

Tolerable doses in seafood related to the frequency and duration of the exposure. A distinction is made between intake by small children and adults, and a further distinction between an Allocation Factor (AF) of 1 (toxins present in food only) and – following the derivation of the provisional WHO GV for Drinking-water – an AF of 0.2 (80% of the dose is taken im—mainly—via drinking water, only 20% via food). For calculating guideline values, following eq. (2) in Section $\underline{3}$ a consumption (C) of 100 g fish (per day) is assumed. Acute TI: single exposure event (e.g. week-end fishing trip). Seasonal TDI: ongoing, "daily" exposure for several weeks during the cyanobacterial season. Lifetime TDI: ongoing "daily" exposure for many months in settings where microcystin-producing cyanobacteria proliferate perennially. ¹NOAEL= no observed adverse effect level. ²Original values in Ibelings and Chorus contained a typo and were listed incorrectly as 0.4 and 0.08 µg/kg; correct values are as shown above.

Previous analyses (Kann 2008) evaluated three of the congeners (-LR, LR-DM, and –LA) with respect to the guideline values derived by Ibelings and Chorus (2007) that are based on toxicity work for microcystin-LR. However, because this approach likely underestimates toxicity due to the exclusion of several of the congeners, similar to the California Office of Environmental Health Hazard Assessment (OEHHA), the below comparisons utilize a more conservative approach with respect to public health that includes the sum of all microcystin congeners. For the following comparisons to Ibelings and Chorus (2007) it is assumed that the only exposure is through ingestion; therefore guideline values were evaluated for Allocation Factor=1 (see reproduced Table 2 above for AF description as well as assumptions regarding frequency and quantity of tissue consumed). In addition, OEHHA calculated the maximum number of 8-oz meals per month at varying microcystin levels in tissue (see Appendix II below); these values are equivalent to the Seasonal TDI guidance value as shown by Ibelings and Chorus in Table 2 above. For example the recommended OEHHA microcystin level above which a child should not consume even one meal per month is $40 \mu g/L$, and is the same level as the Ibelings and Chorus (2007) Seasonal TDI guideline value for a child as shown above.

RESULTS/DISCUSSION

Trends in Tissue and Ambient Toxin Concentration

The neurotoxins anatoxin-a, domoic acid, and okadaic acid were not detected in any of the samples (Appendix I) and will not be discussed further. Of all microcystin congeners measured, only four were detected: MCY-RR, MCY-LR, MCY-LA, and MCY-LRDM (Appendix I). These showed a distinct seasonal pattern in freshwater mussel tissue, with most stations in July dominated primarily by MCY-LR and MCY-LA, and secondarily by MCY-LRDM (Figure 6). Aside from the Yurok station KA, which increased in August from non-detects in July, concentration at the remaining upstream stations decreased in August, with MCY-LRDM not detected for the remainder of the season (Figure 6). The concentration of MCY-LA increased again in September, but MCY-LR remained proportionally lower, especially with respect to July. In addition, MCY-RR was first detected in September at several stations (Figure 6). The overall seasonal pattern of the 3 most dominant congeners over all stations shows July domination by both MCY-LR and MCY-LA and although MCY-LRDM was present in July it was not detected for the remainder of the season; both MCY-LR and MCY-LA declined in August, but only MCY-LA rebounded in September and October; only MCY-LA was detected in a final set of samples taken in December at station BB (Figure 7).

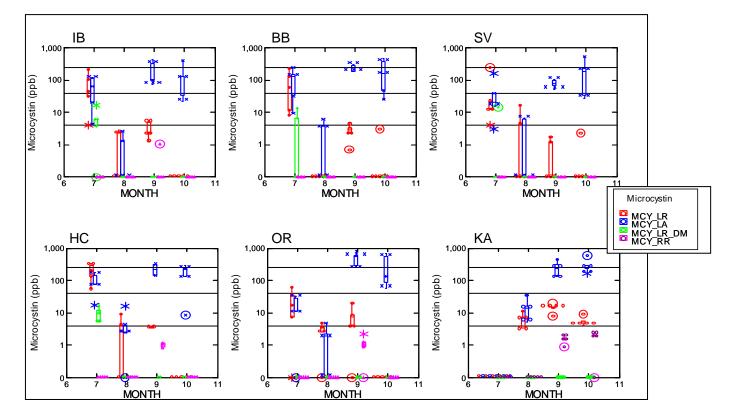


Figure 6. Concentration of microcystin-LR, LA, LR-DM, and RR (ng/g or ppb) in whole freshwater mussels collected from stations (IB, BB, SV, HC, OR, KA) in the Klamath River system in 2009. Horizontal lines denote the Lifetime, Seasonal, and Acute Public health threshold levels.

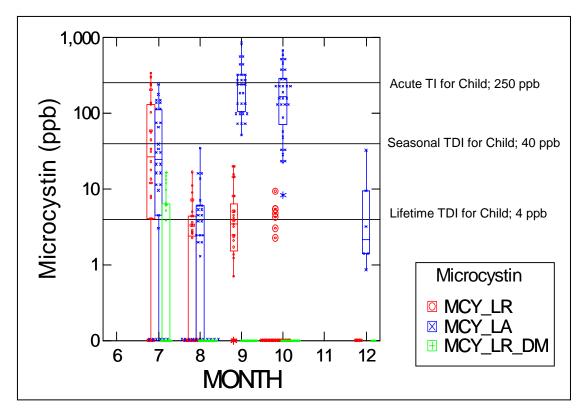


Figure 7. Concentration of microcystin-LR, LA, and LR-DM (ng/g or ppb) in whole freshwater mussels collected from stations (IB, BB, SV, HC, OR, KA) in the Klamath River system in 2009. TDI values are as described in Ibelings and Chorus (2007; Table 2 reproduced above).

A comparison of tissue concentration data relative to ambient concentrations reveals that both MCY-LR and MCY-LA were detected in both media, while MCY-RR and MCY-LRDM were detected in tissue data but they were not detected in water samples (Figure 8). In addition, the frequency of MCY-LR detection in tissue was greater than that for water. The lack of detection or low frequency of detection in water relative to tissue samples indicates that although these congeners may have been below detection in water samples, through bioaccumulation mechanisms were then found in proportionally greater frequency (as well as higher concentration) in freshwater mussel tissue.

Corresponding to the August decline in tissue bioaccumulation, the seasonal trend in ambient data also indicates a decline (most noticeable at station BB) in the total microcystin concentration during some of the August dates (Figure 9). A decline in August *Microcystis aeruginosa* cell density and microcystin concentration was also observed in the upstream reservoir system during August of 2009 (Kann and Corum 2010).

The monthly pattern in total microcystin concentration in freshwater mussel tissue indicates relatively high values in July, September, and October, with highest overall values occurring in September and October (Figure 10a). Microcystin was still detected in mussels from the December sample, with several values continuing to exceed the Lifetime TDI for a child (Figure 10a).

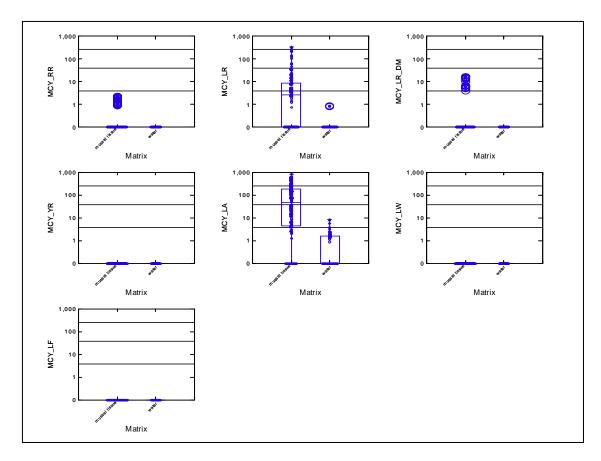


Figure 8. Concentration of microcystin congeners (ng/g or ppb) in freshwater mussel samples and ambient (water) samples from the Klamath River system in 2009.

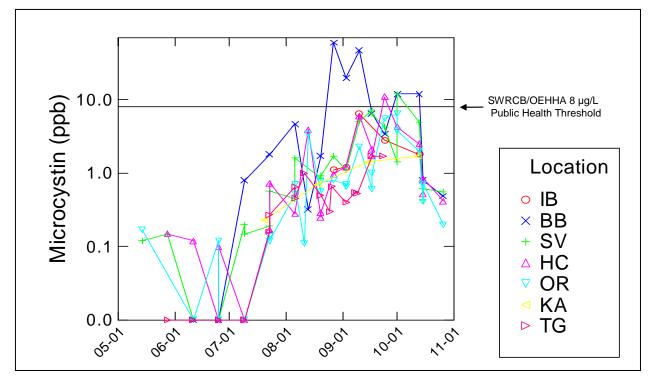


Figure 9. Ambient concentration of total microcystin samples collected from the Klamath River system in 2009.

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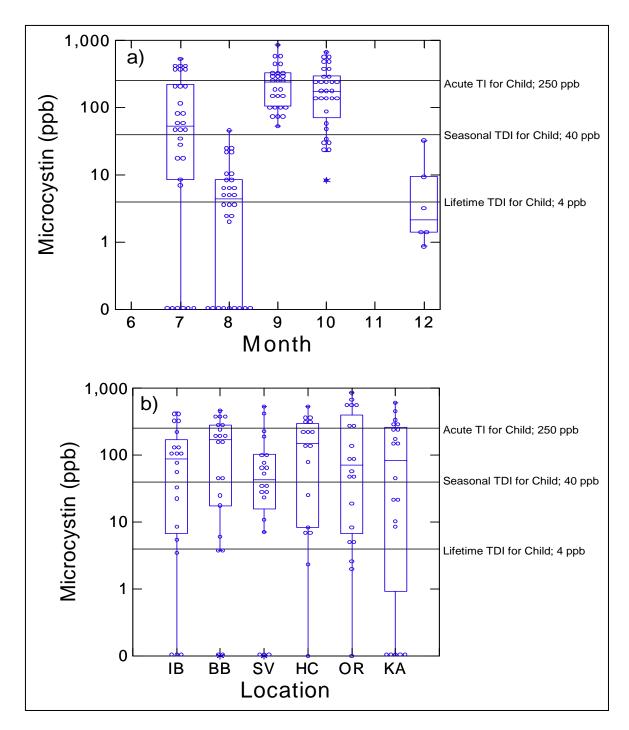


Figure 10. Concentration by month (a) and location (b) of the sum of microcystin-RR, RR-DM, LR, LR-DM, YR, LA, LW, LF and LY in whole freshwater mussels collected from the Klamath River system in 2009 July to October. Stations ordered longitudinally left (upstream) to right (downstream).

A comparison among stations reveals that all stations showed some level of microcystin bioaccumulation, with no clear longitudinal pattern in the median or upper quartile values (Figure 10b). However, there was some indication of a longitudinal pattern in the lower quartile values, which aside from the I5 Bridge (IB) decreased downstream. Seiad Valley (SV) tended to show a lower overall distribution, and the highest upper quartile value was observed at Orleans (OR). Although the farthest downstream station, KA, showed the lowest lower quartile value, there were still numerous occurrences of total microcystin that exceeded the Acute TDI value for a child (Figure 10b). No microcystin was detected in mussels sampled on September 14th from the Trinity River (Appendix I: station TR).

Total microcystin values in ambient water were above the public health guideline values for posting on several occasions at stations BB, SV, and HC, but in general the majority of samples were below the 8 μ g/L posting level (Figure 11). An overall comparison of ambient data to the tissue data reveals that despite relatively low ambient concentrations (Figure 11: median microcystin values were often less than 1 μ g/L), substantial microcystin bioaccumulation occurred at all stations (Figure 10b). This was especially true for the most downstream station, KA, where numerous non-detects and low microcystin values were observed (stations KA and TG in Figure 11), yet levels found in freshwater mussel tissue were above the various public health thresholds (Figure 10b).

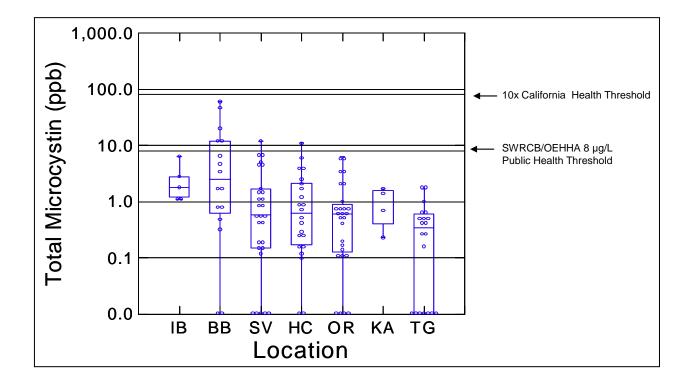


Figure 11. Ambient concentration of total microcystin samples collected from stations in the Klamath River system in 2009.

Comparison to Bioaccumulation in 2007 Freshwater Mussel Samples

Microcystin bioaccumulation in freshwater mussels was evaluated on two dates in 2007; one in July and one in November (Kann 2008). With the exception of MCY-RR which was present in July of 2007 but not 2009, other congeners were similar between the two years during July (the only time period for which samples overlapped in 2007 and 2009); with MC-LR, MCY-LA and MCY-LRDM all present in both years (Figure 12). It should be noted that although MCY-RR was not detected in July of 2009 it was detected in September of 2009. Comparison of the total microcystin concentration between the two years shows elevated bioaccumulation in July for both years, but by November of 2007 no microcystin was detected in the mussel tissue (indicating depuration), while microcystin continued to be detected in December of 2009 (Figure 13).

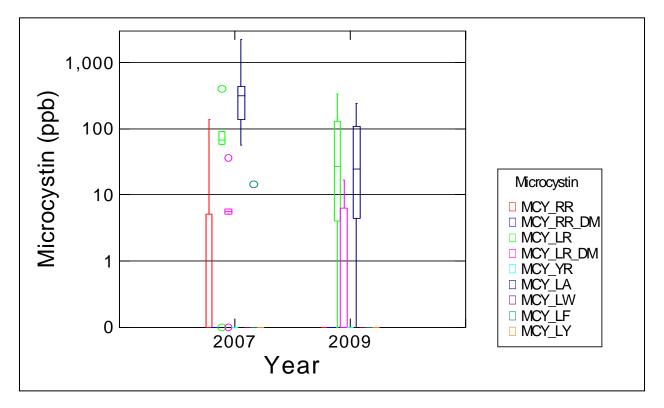


Figure 12. Concentration of Total Microcystin from samples collected in July of 2007 and 2009 from the Klamath River System. Note: Positive concentrations of Microcystin RR found beginning in September 2009.

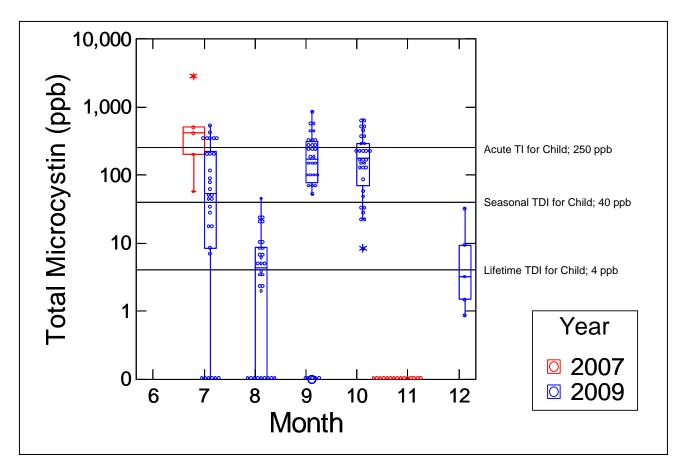


Figure 13. Box plot comparing concentration of Total Microcystin for 2007 and 2009 samples collected from the Klamath River System.

Comparison to Public Health Threshold Values

Concentration of the three most prevalent microcystin congeners found in Klamath River freshwater mussel tissue shows that MCY-LR and MCY-LA levels exceeded all three guideline TDI levels for children (Lifetime, Seasonal, and Acute) at varying times in July, September and October, and that the concentration of MCY-LRDM often exceeded the Lifetime TDI guideline level when it was present in July (Figure 7). In addition, MCY-RR remained below the Lifetime TDI when it was present in September (Figure 6).

Temporal and spatial summaries of the total tissue microcystin concentration (the sum of all congeners) show that ingestion of freshwater mussels in the Klamath River system would result in microcystin doses that exceed various public health thresholds for safe consumption (Figures 14 and 15). This is especially true for children, where in the months of July, September, and October, the Acute Tolerable Intake (TI) dose was exceeded by up to ~4 times (Figure 14a). This means that even one meal of freshwater mussels for a child would exceed what is considered to be a safe level (OEHHA 2008). Exceedance of the Acute TI was shown to occur

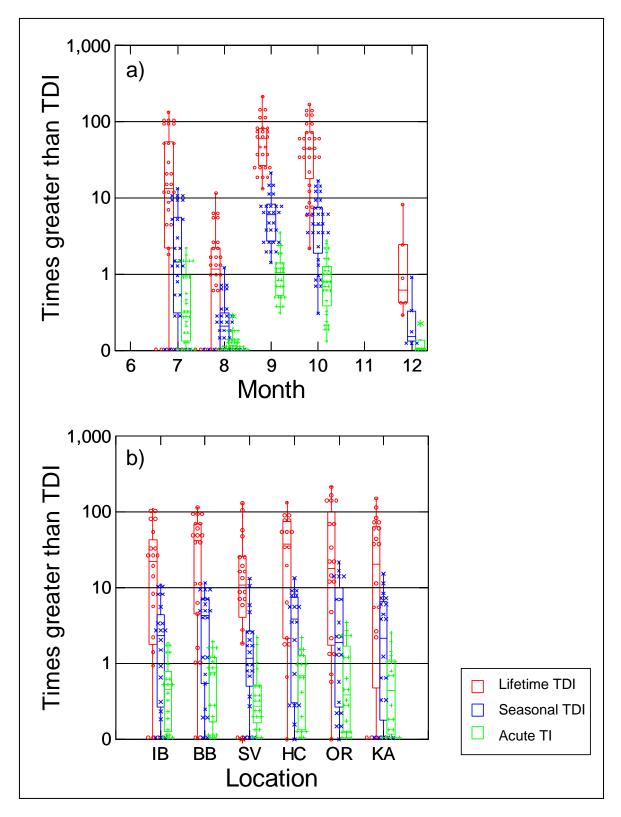


Figure 14.. Exceedance of Child Lifetime, Seasonal, and Acute TDI for the sum of microcystin-RR, RR-DM, LR, LR-DM, YR, LA, LW, LF and LY in whole freshwater mussels collected from the Klamath River system in 2009 July to October; by month (a) and location (b). TDI values are as described in Ibelings and Chorus (2007; Table 2 reproduced above).

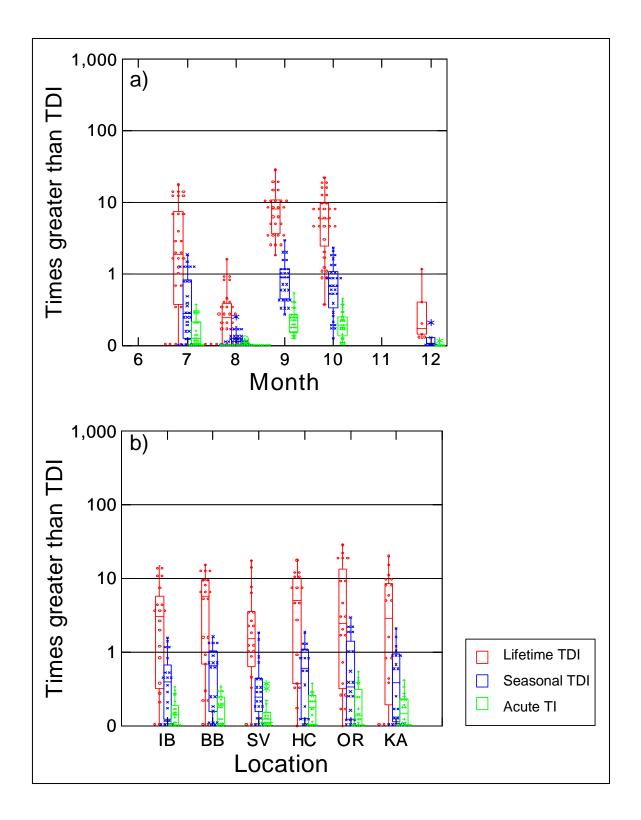


Figure 15. Exceedance of Adult Lifetime, Seasonal, and Acute TDI for the sum of microcystin-RR, RR-DM, LR, LR-DM, YR, LA, LW, LF and LY in whole freshwater mussels collected from the Klamath River system in 2009 July to October; by month (a) and location (b). TDI values are as described in Ibelings and Chorus (2007; Table 2 reproduced above).

at all stations, including the most downstream station, KA (Figure 14b). Total microcystin levels in freshwater mussel tissue consistently exceeded the Lifetime TDI guideline for children by 100's of times, and the Seasonal TDI by over 10 times (Figure 14). In addition, although no exceedances of Acute TI levels for adults were observed, numerous exceedances of the adult Seasonal and Lifetime TDI levels were observed during 2009 (Figure 15). Microcystin levels in the December tissue samples continued to exceed the Lifetime TDI level for both adults and children.

A review of literature pertaining to microcystin bioaccumulation in freshwater organisms reveals that the microcystin bioaccumulation patterns observed in the Klamath River system are commonly observed in other systems as well, and that World Health Organization guidelines for public health were commonly exceeded in a variety of organisms, including fish and freshwater mussels (Table1). A recent review by Smith et al. (2008) found that in 47% of the aquaculture studies they evaluated, hepatotoxin accumulation occurred in edible tissues that exceeded the WHO TDI (.04 μ g kg⁻¹ body weight d⁻¹; assuming 100-300 g fresh weight of tissue consumed). Further work by Smith et al. (2010) indicates that the majority of microcystins are likely covalently bound to target proteins in tissues and thus are not quantified or included in typical assessments. If, as Smith et al. (2010) indicate, these covalently bound microcystins may be made bioavailable in the digestive system of a consumer through the digestion of their attached protein phosphatase, then public health risk may be underestimated.

SUMMARY

The expanded sampling program in 2009 (relative to 2007) included broader spatial and temporal coverage, and clearly demonstrated bioaccumulation of various microcystin congeners in Klamath River freshwater mussels. Similar to 2007 results, evaluation of bioaccumulation in Klamath River freshwater mussels in 2009 with respect to public health guidelines indicates that, aside from a September sample from the Trinity River, all TDI guideline levels as defined by Ibelings and Chorus (2007) were exceeded to varying degrees, including observations of values exceeding Acute TDI thresholds.

A comparison of tissue concentration data relative to ambient concentration revealed that the congeners MCY-LR and MCY-LA were detected in both media, while MCY-RR and MCY-LRDM were detected in tissue data but they were not detected in water samples. In addition, the frequency of MCY-LR detection in tissue was greater than that for water. The lack of detection or low frequency of microcystin detection in water relative to tissue samples clearly indicates that bioaccumulation mechanisms then cause a proportionally greater frequency of detection (as well as higher concentration) in freshwater mussel tissue. The importance of this finding is that even when microcystin may be below detection in the ambient water, accumulation in freshwater mussel tissue can still occur.

A decline in tissue bioaccumulation was observed in August and coincided with a decline in ambient concentration, as well as a decline in *Microcystis aeruginosa* cell density and microcystin concentration in the upstream reservoir system during August of 2009. The

Author	Date	System	Toxin	Organism	Exceeds WHO TDI	Notes
Amorim et al.	1999	fresh water	MC-LR	mussels	yes	
Cazenave et al.	2005	reservoir	MC-RR	fish	yes	Presence of MC-RR in brain
Chen et al.	2005	lake	MC-LR, MC-RR, MC-YR	mussels	yes	Mean daily intakes were 8-23.5 times TDI value when mussels are eaten whole
Chen et al.	2006	lake	мс	silver carp	yes	Silver Carp should not be consumed during period of dense <i>Microcystis</i> blooms
Garcia et al.	2010	lake	МС	blue crab	yes	MC levels exceeded WHO drinking water guidelines.
Gkelis et al.	2006	fresh water	МС	various	yes	
Hang-jun et al.	2006	river	MC-LR, MC-RR	fish	no	Concentration close to WHO TDI warning of health risk
Kopp et al.	2005	pond	MC-LR, MC-RR, MC-YR	fish	NA	
Magalhaes et al.	2001	coastal lagoon	MC	fish	ves	
Magalhaes et al.	2003	bay	мс	fish, crustaceans	yes	19% of animal samples were above WHO TDI
Masango et al.	2008	fresh water	MC-LR		yes	MC-LR concentration was 1000 times more than the WHO provisional guideline for drinking water
Mohamed et al.	2003	fish farm	MC	fish	NA	Highest MC concentration found in guts
Osswald et al.	2008	laboratory aquarium	Anatoxin-a	mussels	no	One day after beginning depuration the toxin could not be detected
Valeria et al.	2010	lake	MC-LR, MC-RR, MC-YR, MC-LA	fish	no	MC-RR was dominant in water samples.
Van Buynder et al.	2001	lake	MC-LR	mussels	yes	
Vasconcelos	1995	estuary	MC-LR	mussels	yes	96% of toxin found in digestive gland and stomach
Vasconcelos	1999	fresh water	MC-LR	fish, crayfish, mussels	yes	
Wilson et al.	2008	lake	мс	fish	no	MC levels exceeded WHO drinking water guidelines.
Wood et al.	2006	lake	мс	fish, mussels	NA	MC detected in mussels but not in fish
Xie et al.	2005	lake	MC-LR, MC-RR	fish	yes	MC found in bile and blood
Zhao et al.	2006	flow-through	МС	fish	yes	

Table 1. Literature review of cyanotoxin bioaccumulation in freshwater organisms	Table 1.	Literature	review of	cvanotoxin	bioaccumulatio	n in	freshwater	organisms.
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monthly pattern in total microcystin concentration in freshwater mussel tissue indicated relatively high values in July, September, and October, with highest overall values occurring in September and October of 2009. Microcystin was still detected in mussels from the December sample, with several values continuing to exceed the Lifetime TDI for a child.

An overall comparison of ambient data to the tissue data revealed that despite relatively low ambient concentrations (median microcystin values were often less than $1\mu g/L$), substantial microcystin bioaccumulation occurred at all stations. This was especially true for the most downstream station, KA, where numerous non-detects and low microcystin values were observed (stations KA and TG), yet levels found in freshwater mussel tissue were substantially above the various public health thresholds for safe consumption.

These data show that ingestion of freshwater mussels in the Klamath River system would result in microcystin doses that exceed various public health thresholds for safe consumption throughout the summer and fall. This is true especially for children in the months of July, September, and October, when the Acute Tolerable Intake (TI) dose was exceeded by up to ~4 times. These are the months when traditional and subsistence use of fresh water mussels by Tribal members occurs, and at these times even one meal could exceed safe consumption levels. It should be realized that if the use of these organisms is curtailed during these months that coincide with harvest times, their use would be effectively eliminated both from a dietary and Tribal cultural standpoint. Regardless, given that ambient microcystin levels below public health guidelines for recreation can results in substantial microcystin bioaccumulation freshwater mussels, caution must be exercised when consuming these organisms in the mainstem Klamath River system.

Acknowledgements

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APPENDIX I

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri X	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-362-09-1	TC	TC052809	Yurok	28/May/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-463-09-1	тс	TC062509- OC	Yurok	25/Jun/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-431-09-2	IG	IG070909-IG	Karuk	09/Jul/2009	water	ND	ND	0.730	ND	ND	ND	ND	ND	ND	0.73	ND	ND	ND
L-431-09-1	SV	SV070909- SG	Karuk	09/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-431-09- 1Dup	SV-D	SV070909- SG	Karuk	09/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-463-09-2	WE	WE071009- OC	Yurok	10/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-2	KA	KASR072009 -01	Yurok	20/Jul/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-3	KA	KASR072009 -02	Yurok	20/Jul/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-4	KA	KASR072009 -03	Yurok	20/Jul/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-5	КА	KASR072009 -04	Yurok	20/Jul/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-6	KA	KASR072009 -05	Yurok	20/Jul/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-447-09-1	KA	KASR072009 -OC	Yurok	20/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-456-09- 21	BB	BB072209-1	Karuk	22/Jul/2009	muss el	ND	ND	59.3	ND	ND	146	ND	ND	ND	205.58	ND	ND	ND
L-456-09-		BB072203-1	παιώκ	22/30//2009	muss			53.5			-				17.54			
22 L-456-09-	BB	BB072209-2	Karuk	22/Jul/2009	el muss	ND	ND	8.05	ND	ND	9.49	ND	ND	ND	17.54	ND	ND	ND
23	BB	BB072209-3	Karuk	22/Jul/2009	el	ND	ND	129	6.52	ND	240	ND	ND	ND	375.59	ND	ND	ND
L-456-09- 24	BB	BB072209-4	Karuk	22/Jul/2009	muss el	ND	ND	11.8	ND	ND	32.7	ND	ND	ND	44.58	ND	ND	ND
L-456-09- 25	BB	BB072209-5	Karuk	22/Jul/2009	muss el	ND	ND	233	13.4	ND	131	ND	ND	ND	376.58	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-456-09- 16	НС	HC072209-1	Karuk	22/Jul/2009	muss el	ND	ND	56.0	5.12	ND	17.2	ND	ND	ND	78.32	ND	ND	ND
L-456-09-	пс	HC072209-1	Natuk	22/Jul/2009	muss	ND	ND	30.0	0.12	ND	17.2	ND	ND	ND		ND	ND	ND
17	HC	HC072209-2	Karuk	22/Jul/2009	el	ND	ND	299	12.0	ND	75.7	ND	ND	ND	386.52	ND	ND	ND
L-456-09-					muss										527.38			
18	HC	HC072209-3	Karuk	22/Jul/2009	el	ND	ND	335	16.5	ND	176	ND	ND	ND	021.00	ND	ND	ND
L-456-09- 19	НС	HC072209-4	Karuk	22/Jul/2009	muss el	ND	ND	202	9.62	ND	148	ND	ND	ND	359.72	ND	ND	ND
L-456-09-			riarun		muss				0.02						218.92			
20	HC	HC072209-5	Karuk	22/Jul/2009	el	ND	ND	138	5.91	ND	74.9	ND	ND	ND	210.92	ND	ND	ND
L-456-09- 26	IB	IB072209-1	Karuk	22/Jul/2009	muss el	ND	ND	103	6.23	ND	110	ND	ND	ND	219.45	ND	ND	ND
L-456-09-		10072209-1	Maluk	22/30/2009	muss	ND	ND	103	0.23	ND	110	ND	ND	ND		ND	ND	ND
27	IB	IB072209-2	Karuk	22/Jul/2009	el	ND	ND	31.3	3.80	ND	21.1	ND	ND	ND	56.19	ND	ND	ND
L-456-09-		1202000			muss				- · -						115.72			
28 L-456-09-	IB	IB072209-3	Karuk	22/Jul/2009	el	ND	ND	44.7	6.15	ND	64.9	ND	ND	ND		ND	ND	ND
L-458-09- 29	IB	IB072209-4	Karuk	22/Jul/2009	muss el	ND	ND	211	16.5	ND	124	ND	ND	ND	351.26	ND	ND	ND
L-456-09-					muss										8.40			
30	IB	IB072209-5	Karuk	22/Jul/2009	el	ND	ND	4.01	ND	ND	4.38	ND	ND	ND	0.40	ND	ND	ND
L-456-09- 27Dup	IB-D	IB072209-2	Karuk	22/Jul/2009	muss el	ND	ND	34.5	4.90	ND	25.4	ND	ND	ND	64.86	ND	ND	ND
27.Dup	0-0	10072209-2	Maluk	22/30/2009	muss	ND	ND	34.5	4.90	ND	23.4	ND	ND	ND		ND	ND	ND
L-456-09-6	OR	OR072209-1	Karuk	22/Jul/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-456-09-7	OR	OR072209-2	Karuk	22/Jul/2009	muss el	ND	ND	7.55	ND	ND	11.3	ND	ND	ND	18.85	ND	ND	ND
	011	0110122002	rtarun	22,000,2000	muss	110		1.00		110	11.0				50.94			
L-456-09-8	OR	OR072209-3	Karuk	22/Jul/2009	el	ND	ND	17.6	ND	ND	33.3	ND	ND	ND	50.94	ND	ND	ND
L-456-09-9	OR	OR072209-4	Karuk	22/Jul/2009	muss el	ND	ND	34.0	ND	ND	11.3	ND	ND	ND	45.31	ND	ND	ND
L-456-09-					muss										89.91			
10	OR	OR072209-5	Karuk	22/Jul/2009	el	ND	ND	60.9	ND	ND	29.0	ND	ND	ND	09.91	ND	ND	ND
L-456-09- 11	SV	SV072209-1	Karuk	22/Jul/2009	muss el	ND	ND	247	14.6	ND	158	ND	ND	ND	419.56	ND	ND	ND
L-456-09-					muss										61.67			
12	SV	SV072209-2	Karuk	22/Jul/2009	el	ND	ND	22.9	ND	ND	38.8	ND	ND	ND	01.07	ND	ND	ND
L-456-09- 13	SV	SV072209-3	Karuk	22/Jul/2009	muss el	ND	ND	3.97	ND	ND	2.95	ND	ND	ND	6.93	ND	ND	ND
L-456-09-	SV	SV/072200 A	Koruk	22/10/2000	muss	ND		10.0		ND	15 F		ND	ND	27.84	ND	ND	ND
14		SV072209-4	Karuk	22/Jul/2009	el		ND	12.3	ND		15.5	ND			34.60			
L-456-09-	SV	SV072209-5	Karuk	22/Jul/2009	muss	ND	ND	13.4	ND	ND	21.2	ND	ND	ND	34.00	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
15					el													
L-456-09-					muss										24.20			
14Dup	SV-D	SV072209-4 BB072209-	Karuk	22/Jul/2009	el	ND	ND	10.9	ND	ND	13.3	ND	ND	ND	220	ND	ND	ND
L-456-09-4	BB	OC	Karuk	22/Jul/2009	water	ND	ND	ND	ND	ND	1.67	ND	ND	ND	1.67	ND	ND	ND
		HC072209-													2.29			
L-456-09-3	HC	OC	Karuk	22/Jul/2009	water	ND	ND	ND	ND	ND	2.29	ND	ND	ND		ND	ND	ND
L-456-09-5	IB	IB072209-OC OR072209-	Karuk	22/Jul/2009	water	ND	ND	ND	ND	ND	1.90	ND	ND	ND	1.90	ND	ND	ND
L-456-09-1	OR	OR072209- OC	Karuk	22/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
		SV072209-													0.00			
L-456-09-2	SV	OC SV072309-	Karuk	22/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-464-09-1	SV	SG SV072309-	Karuk	23/Jul/2009	water	ND	ND	ND	ND	ND	1.62	ND	ND	ND	1.62	ND	ND	ND
		TC072309-	Yurok												0.00			
L-463-09-3	TC	OC		23/Jul/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND		ND	ND	ND
L-501-09-2	IG	IG080609-OC SV080609-	Karuk	06/Aug/2009	water	ND	ND	ND	ND	ND	1.87	ND	ND	ND	1.87	ND	ND	ND
L-501-09-1	SV	SV080809- SG	Karuk	06/Aug/2009	water	ND	ND	ND	ND	ND	2.54	ND	ND	ND	2.54	ND	ND	ND
		WE080609-		5											2.98			
L-527-09-2	WE	OC KASR081809	Yurok	06/Aug/2009	water muss	ND	ND	0.690	ND	ND	2.29	ND	ND	ND	2.00	ND	ND	ND
L-527-09-3	KA	-01	Yurok	18/Aug/2009	el	ND	ND	3.23	ND	ND	5.27	ND	ND	ND	8.51	ND	ND	ND
		KASR081809			muss										10.22			
L-527-09-4	KA	-02 KASR081809	Yurok	18/Aug/2009	el muss	ND	ND	3.29	ND	ND	6.93	ND	ND	ND	10.22	ND	ND	ND
L-527-09-5	KA	-03	Yurok	18/Aug/2009	el	ND	ND	11.1	ND	ND	34.3	ND	ND	ND	45.34	ND	ND	ND
		KASR081809			muss										20.65			
L-527-09-6	KA	-04 KASR081809	Yurok	18/Aug/2009	el muss	ND	ND	7.07	ND	ND	13.6	ND	ND	ND		ND	ND	ND
L-527-09-7	KA	-05	Yurok	18/Aug/2009	el	ND	ND	7.17	ND	ND	16.4	ND	ND	ND	23.55	ND	ND	ND
		KASR081809													1.05			
L-527-09-1 L-537-09-	KA	-OC	Yurok	18/Aug/2009	water muss	ND	ND	ND	ND	ND	1.05	ND	ND	ND		ND	ND	ND
23	BB	BB081909-1	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	3.67	ND	ND	ND	3.67	ND	ND	ND
L-537-09-			Kernit	10/1	muss	ND		ND	ND		2.00	ND			3.80	ND	ND	ND
24 L-537-09-	BB	BB081909-2	Karuk	19/Aug/2009	el muss	ND	ND	ND	ND	ND	3.80	ND	ND	ND		ND	ND	ND
25	BB	BB081909-3	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09-	BB	BB081909-4	Karuk	19/Aug/2009	muss	ND	ND	ND	ND	ND	5.99	ND	ND	ND	5.99	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
26					el													
L-537-09-					muss										0.00			
27	BB	BB081909-5	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09- 13	нс	HC081909-1	Karuk	19/Aug/2009	muss el	ND	ND	4.27	ND	ND	2.51	ND	ND	ND	6.78	ND	ND	ND
L-537-09-	110	1100010001	Karak	10// (09/2000	muss	THE .		7.21		TTD .	2.01	ND	ND	ND	04.04		ND	
14	HC	HC081909-2	Karuk	19/Aug/2009	el	ND	ND	8.95	ND	ND	16.0	ND	ND	ND	24.91	ND	ND	ND
L-537-09-		110004000 0	14 a mala	40/4	muss			0.50			4.05				6.85			ND
15 L-537-09-	HC	HC081909-3	Karuk	19/Aug/2009	el muss	ND	ND	2.59	ND	ND	4.25	ND	ND	ND		ND	ND	ND
16	HC	HC081909-4	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	2.25	ND	ND	ND	2.25	ND	ND	ND
L-537-09-				Ŭ	muss										0.00			
17	HC	HC081909-5	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09- 28	IB	IB081909-1	Karuk	19/Aug/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09-		120010001	Rarak	10// (09/2000	muss			ND	ND	TTD .	ND	ND	ne.		0.00		ND	
29	IB	IB081909-2	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09-	IB		1 and 1	40/4	muss	ND	ND	0.45			4.40	ND	ND	ND	3.34	ND		ND
30 L-537-09-	IB	IB081909-3	Karuk	19/Aug/2009	el muss	ND	ND	2.15	ND	ND	1.19	ND	ND	ND		ND	ND	ND
31	IB	IB081909-4	Karuk	19/Aug/2009	el	ND	ND	2.74	ND	ND	2.55	ND	ND	ND	5.29	ND	ND	ND
L-537-09-				-	muss										0.00			
32	IB	IB081909-5	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09- 30Dup	IB-D	IB081909-3	Karuk	19/Aug/2009	muss el	ND	ND	2.85	ND	ND	14.5	ND	ND	ND	17.38	ND	ND	ND
00D.up		120010000	Raran	10,7 (09,2000	muss			2.00			1 1.0				2.48		110	
L-537-09-8	OR	OR081909-1	Karuk	19/Aug/2009	el	ND	ND	2.48	ND	ND	ND	ND	ND	ND	2.40	ND	ND	ND
L-537-09-9	OR	OR081909-2	Karuk	10/400/2000	muss el	ND	ND	ND	ND	ND	1.91	ND	ND	ND	1.91	ND	ND	ND
L-537-09-9	UK	UR061909-2	Natuk	19/Aug/2009	muss	ND	ND	ND	IND	ND	1.91	ND	ND	ND		ND	IND	ND
10	OR	OR081909-3	Karuk	19/Aug/2009	el	ND	ND	3.56	ND	ND	4.72	ND	ND	ND	8.28	ND	ND	ND
L-537-09-					muss										4.89			
11 L-537-09-	OR	OR081909-4	Karuk	19/Aug/2009	el	ND	ND	4.89	ND	ND	ND	ND	ND	ND		ND	ND	ND
L-537-09- 12	OR	OR081909-5	Karuk	19/Aug/2009	muss el	ND	ND	3.26	ND	ND	2.08	ND	ND	ND	5.33	ND	ND	ND
L-537-09-					muss										0.00			
18	SV	SV081909-1	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09- 19	SV	SV081909-2	Karuk	19/Aug/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09-	30	31001909-2	raiuk	1 <i>9/A</i> ug/2009	muss						שא	שא	טא	טא		טא		
20	SV	SV081909-3	Karuk	19/Aug/2009	el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-537-09- 21	SV	SV081909-4	Karuk	19/Aug/2009	muss el	ND	ND	16.6	ND	ND	6.52	ND	ND	ND	23.14	ND	ND	ND
L-537-09- 22	SV	SV081909-5	Karuk	19/Aug/2009	muss el	ND	ND	4.40	ND	ND	6.30	ND	ND	ND	10.70	ND	ND	ND
L-537-09- 21Dup	SV-D	SV081909-4	Karuk	19/Aug/2009	muss el	ND	ND	20.9	ND	ND	12.5	ND	ND	ND	33.34	ND	ND	ND
L-537-09-4	BB	BB081909- OC	Karuk	19/Aug/2009	water	ND	ND	20.9 ND	ND	ND	1.34	ND	ND	ND	1.34	ND	ND	ND
	НС	HC081909- OC		5		ND	ND	ND	ND	ND	1.47	ND	ND	ND	1.47	ND	ND	ND
L-537-09-2 L-537-09-5	IB	IB081909-OC	Karuk Karuk	19/Aug/2009 19/Aug/2009	water water	ND	ND	ND	ND ND	ND	1.47	ND	ND	ND	1.23	ND	ND	ND ND
L-537-09-1	OR	OR081909- OC	Karuk	19/Aug/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-537-09-3	sv	SV081909- OC	Karuk	19/Aug/2009	water	ND	ND	ND	ND	ND	1.40	ND	ND	ND	1.40	ND	ND	ND
L-537-09-6	sv	SV082009- SG	Karuk	20/Aug/2009	water	ND	ND	ND	ND	ND	8.24	ND	ND	ND	8.24	ND	ND	ND
L-537-09-7	sv	SV082009- OC	Karuk	20/Aug/2009	water	ND	ND	ND	ND	ND	1.39	ND	ND	ND	1.39	ND	ND	ND
L-537-09- 7Dup	SV-D	SV082009- OC	Karuk	20/Aug/2009	water	ND	ND	ND	ND	ND	1.69	ND	ND	ND	1.69	ND	ND	ND
		TC082009-		Ŭ											0.00			
L-632-09-1 L-566-09-2	TC IB	OC IB090309-SG	Yurok Karuk	20/Aug/2009 03/Sep/2009	water water	ND ND	ND ND	ND ND	ND ND	ND ND	ND ND	ND ND	ND ND	ND ND	0.00	ND ND	ND ND	ND ND
L-566-09-3	IG	IG090309-OC	Karuk	03/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-566-09- 3Dup	IG-D	IG090309-OC	Karuk	03/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-566-09-1	SV	SV090309- SG	Karuk	03/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-632-09-2	тс	TC090309- OC	Yurok	03/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-2	TR	TR091409-01	Yurok	14/Sep/2009	muss	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-3	TR	TR091409-02	Yurok	14/Sep/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-4	TR	TR091409-03	Yurok	14/Sep/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-4 Dup	TR	TR091409-03	Yurok	14/Sep/2009	muss	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-5	TR	TR091409-03 TR091409-04	Yurok	14/Sep/2009 14/Sep/2009	el muss	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri X	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
					el													
L-584-09-6	TR	TR091409-05	Yurok	14/Sep/2009	muss el	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-584-09-1	TR	TR091409- OC	Yurok	14/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
		KASR091509	TUTOR	•	muss													
L-591-09-2 L-591-09-2	KA	-01 KASR091509	Yurok	15/Sep/2009	el muss	1.38	ND	14.3	ND	ND	317	ND	ND	ND	332.85	ND	ND	ND
Dup	KA	-01	Yurok	15/Sep/2009	el	1.51	ND	13.0	ND	ND	260	ND	ND	ND	274.43	ND	ND	ND
L-591-09-3	КА	KASR091509 -02	Yurok	15/Sep/2009	muss el	1.57	ND	15.4	ND	ND	439	ND	ND	ND	455.54	ND	ND	ND
L 001-00-0		KASR091509	TURK	10/000/2000	muss	1.07												
L-591-09-4	KA	-03 KASR091509	Yurok	15/Sep/2009	el muss	1.81	ND	19.9	ND	ND	134	ND	ND	ND	155.25	ND	ND	ND
L-591-09-5	KA	-04	Yurok	15/Sep/2009	el	0.787	ND	7.94	ND	ND	141	ND	ND	ND	149.37	ND	ND	ND
L-591-09-6	KA	KASR091509 -05	Yurok	15/Sep/2009	muss el	1.50	ND	14.5	ND	ND	243	ND	ND	ND	259.01	ND	ND	ND
		KASR091509		•														
L-591-09-1 L-601-09-	KA	-OC	Yurok	15/Sep/2009	water muss	ND	ND	ND	ND	ND	1.50	ND	ND	ND	1.50	ND	ND	ND
20	BB	BB091609-1	Karuk	16/Sep/2009	el	ND	ND	2.43	ND	ND	187	ND	ND	ND	189.38	ND	ND	ND
L-601-09- 21	BB	BB091609-2	Karuk	16/Sep/2009	muss el	ND	ND	3.10	ND	ND	275	ND	ND	ND	278.54	ND	ND	ND
L-601-09- 22	BB	BB091609-3	Karuk	16/Sep/2009	muss el	ND	ND	0.615	ND	ND	184	ND	ND	ND	185.03	ND	ND	ND
L-601-09- 23	BB	BB091609-4	Karuk	16/Sep/2009	muss el	ND	ND	2.25	ND	ND	237	ND	ND	ND	239.28	ND	ND	ND
L-601-09- 24	BB	BB091609-5	Karuk	16/Sep/2009	muss el	ND	ND	4.43	ND	ND	293	ND	ND	ND	297.82	ND	ND	ND
L-601-09- 13	HC	HC091609-1	Karuk	16/Sep/2009	muss el	0.768	ND	3.33	ND	ND	145	ND	ND	ND	149.12	ND	ND	ND
L-601-09- 14	HC	HC091609-2	Karuk	16/Sep/2009	muss el	1.10	ND	3.87	ND	ND	321	ND	ND	ND	326.30	ND	ND	ND
L-601-09- 25	IB	IB091609-1	Karuk	16/Sep/2009	muss el	ND	ND	1.26	ND	ND	106	ND	ND	ND	107.25	ND	ND	ND
L-601-09- 26	IB	IB091609-2	Karuk	16/Sep/2009	muss el	ND	ND	2.38	ND	ND	98.6	ND	ND	ND	101.02	ND	ND	ND
L-601-09- 27	IB	IB091609-3	Karuk	16/Sep/2009	muss	ND	ND	2.02	ND	ND	74.8	ND	ND	ND	76.84	ND	ND	ND
L-601-09- 28	IB	IB091609-4	Karuk	16/Sep/2009	muss el	0.982	ND	5.07	ND	ND	317	ND	ND	ND	323.41	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-601-09- 29	IB	IB091609-5	Karuk	16/Sep/2009	muss el	ND	ND	4.98	ND	ND	440	ND	ND	ND	444.84	ND	ND	ND
L-601-09-8	OR	OR091609-1	Karuk	16/Sep/2009	muss el	2.12	ND	19.6	ND	ND	555	ND	ND	ND	576.60	ND	ND	ND
L-601-09-9	OR	OR091609-2	Karuk	16/Sep/2009	muss el	1.14	ND	8.22	ND	ND	840	ND	ND	ND	849.74	ND	ND	ND
L-601-09- 10	OR	OR091609-3	Karuk	16/Sep/2009	muss el	0.960	ND	4.06	ND	ND	598	ND	ND	ND	603.38	ND	ND	ND
L-601-09- 11	OR	OR091609-4	Karuk	16/Sep/2009	muss el	ND	ND	ND	ND	ND	295	ND	ND	ND	294.56	ND	ND	ND
L-601-09- 12	OR	OR091609-5	Karuk	16/Sep/2009	muss el	0.820	ND	3.76	ND	ND	271	ND	ND	ND	275.46	ND	ND	ND
L-601-09- 10Dup	OR-D	OR091609-3	Karuk	16/Sep/2009	muss el	0.876	ND	3.57	ND	ND	544	ND	ND	ND	548.48	ND	ND	ND
L-601-09- 15	SV	SV091609-1	Karuk	16/Sep/2009	muss el	ND	ND	1.62	ND	ND	98.8	ND	ND	ND	100.41	ND	ND	ND
L-601-09- 16	SV	SV091609-2	Karuk	16/Sep/2009	muss el	ND	ND	ND	ND	ND	77.2	ND	ND	ND	77.24	ND	ND	ND
L-601-09- 17	SV	SV091609-3	Karuk	16/Sep/2009	muss el	ND	ND	ND	ND	ND	105	ND	ND	ND	105.20	ND	ND	ND
L-601-09- 18	SV	SV091609-4	Karuk	16/Sep/2009	muss el	ND	ND	ND	ND	ND	69.8	ND	ND	ND	69.79	ND	ND	ND
L-601-09- 19	SV	SV091609-5	Karuk	16/Sep/2009	muss el	ND	ND	1.13	ND	ND	51.5	ND	ND	ND	52.62	ND	ND	ND
L-601-09-4	BB	BB091609- OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	1.13	ND	ND	ND	1.13	ND	ND	ND
L-601-09- 4Dup	BB-D	BB091609- OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	2.22	ND	ND	ND	2.22	ND	ND	ND
L-601-09-2	HC	HC091609- OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	1.51	ND	ND	ND	1.51	ND	ND	ND
L-601-09-5	IB	IB091609-OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	3.73	ND	ND	ND	3.73	ND	ND	ND
L-601-09-1	OR	OR091609- OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	0.789	ND	ND	ND	0.79	ND	ND	ND
L-601-09-3	SV	SV091609- OC	Karuk	16/Sep/2009	water	ND	ND	ND	ND	ND	3.22	ND	ND	ND	3.22	ND	ND	ND
L-601-09-6	SV	SV091709- OC [†]	Karuk	17/Sep/2009	water	ND	ND	ND	ND	ND	7.70	ND	ND	ND	7.70	ND	ND	ND
L-601-09-7	SV	SV091709- SG	Karuk	17/Sep/2009	water	ND	ND	ND	ND	ND	5.68	ND	ND	ND	5.68	ND	ND	ND
L-632-09-3	тс	TC091709- OC	Yurok	17/Sep/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri X	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-626-09-2	IG	IG100109-OC	Yurok	01/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-626-09-1	SV	SV100109- SG	Karuk	01/Oct/2009	water	ND	ND	ND	ND	ND	1.50	ND	ND	ND	1.50	ND	ND	ND
L-626-09- 1Dup	SV-D	SV100109- SG	Karuk	01/Oct/2009	water	ND	ND	ND	ND	ND	1.68	ND	ND	ND	1.68	ND	ND	ND
L-632-09-4	WE	WE100109- OC	Yurok	01/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-653-09-2	КА	KASR101209 -1	Yurok	12/Oct/2009	muss el	1.80	ND	4.64	ND	ND	282	ND	ND	ND	288.86	ND	ND	ND
L-033-09-2	104	KASR101209	TUIOK	12/00/2009	muss	1.00	ND	4.04	ND		202	ND	IND	ND	200.00		ND	IND
L-653-09-3	KA	-2	Yurok	12/Oct/2009	el	1.97	ND	4.21	ND	ND	234	ND	ND	ND	239.99	ND	ND	ND
L-653-09-4	KA	KASR101209 -3	Yurok	12/Oct/2009	muss el	2.24	ND	9.24	ND	ND	589	ND	ND	ND	600.36	ND	ND	ND
L-653-09-	101	KASR101209	raiok	12/00/2003	muss	2.27		0.24			000	ND			000.00		TTD .	
4Dup	KA	-3	Yurok	12/Oct/2009	el	2.00	ND	7.66	ND	ND	474	ND	ND	ND	483.92	ND	ND	ND
L-653-09-5	KA	KASR101209 -4	Yurok	12/Oct/2009	muss el	ND	ND	5.32	ND	ND	247	ND	ND	ND	252.72	ND	ND	ND
	101	KASR101209	ruion		muss										LOL.IL			
L-653-09-6	KA	-5 KASR101209	Yurok	12/Oct/2009	el	1.94	ND	5.33	ND	ND	167	ND	ND	ND	174.27	ND	ND	ND
L-653-09-1	KA	-OC	Yurok	12/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-653-09-		KASR101209																
1Dup	KA	-OC TC101509-	Yurok	12/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-691-09-1	тс	OC	Yurok	15/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-685-09- 11	BB	BB102609-1	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	24.8	ND	ND	ND	24.80	ND	ND	ND
L-685-09- 12	BB	BB102609-2	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	459	ND	ND	ND	458.72	ND	ND	ND
L-685-09- 13	BB	BB102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	48.3	ND	ND	ND	48.29	ND	ND	ND
L-685-09- 14	BB	BB102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	374	ND	ND	ND	373.72	ND	ND	ND
L-685-09- 15	BB	BB102609-5	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	149	ND	ND	ND	149.41	ND	ND	ND
L-685-09- 16	BB	BB102609-6	Karuk	26/Oct/2009	muss el	ND	ND	2.93	ND	ND	166	ND	ND	ND	168.56	ND	ND	ND
L-685-09- 22	HC	HC102609-1	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	227	ND	ND	ND	226.56	ND	ND	ND
L-685-09- 23	HC	HC102609-2	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	8.25	ND	ND	ND	8.25	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri x	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
L-685-09- 24	HC	HC102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	296	ND	ND	ND	296.31	ND	ND	ND
L-685-09- 25	HC	HC102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	130	ND	ND	ND	130.48	ND	ND	ND
L-685-09- 26	HC	HC102609-5	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	238	ND	ND	ND	237.64	ND	ND	ND
L-685-09-6	IB	IB102609-1	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	393	ND	ND	ND	392.70	ND	ND	ND
L-685-09-7	IB	IB102609-2	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	130	ND	ND	ND	129.89	ND	ND	ND
L-685-09-8	IB	IB102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	22.3	ND	ND	ND	22.26	ND	ND	ND
L-685-09-9	IB	IB102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	32.8	ND	ND	ND	32.81	ND	ND	ND
L-685-09- 10	IB	IB102609-5	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	132	ND	ND	ND	131.83	ND	ND	ND
L-685-09- 8Dup	IB-D	IB102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	20.4	ND	ND	ND	20.37	ND	ND	ND
L-685-09- 27	OR	OR102609-1	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	666	ND	ND	ND	666.34	ND	ND	ND
L-685-09- 28	OR	OR102609-2	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	57.7	ND	ND	ND	57.73	ND	ND	ND
L-685-09- 29	OR	OR102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	534	ND	ND	ND	534.23	ND	ND	ND
L-685-09- 30	OR	OR102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	87.1	ND	ND	ND	87.14	ND	ND	ND
L-685-09- 31	OR	OR102609-5	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	136	ND	ND	ND	135.74	ND	ND	ND
L-685-09- 30Dup	OR-D	OR102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	62.1	ND	ND	ND	62.08	ND	ND	ND
L-685-09- 17	SV	SV102609-1	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	188	ND	ND	ND	188.35	ND	ND	ND
L-685-09- 18	SV	SV102609-2	Karuk	26/Oct/2009	muss el	ND	ND	2.17	ND	ND	521	ND	ND	ND	523.09	ND	ND	ND
L-685-09- 19	SV	SV102609-3	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	229	ND	ND	ND	229.10	ND	ND	ND
L-685-09- 20	SV	SV102609-4	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	34.4	ND	ND	ND	34.36	ND	ND	ND
L-685-09- 21	SV	SV102609-5	Karuk	26/Oct/2009	muss el	ND	ND	ND	ND	ND	28.3	ND	ND	ND	28.26	ND	ND	ND
L-685-09-2	BB	BB102609-	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND

WPCL Lab#	Station	Sample Identification	Collecto r	Date Collected	Matri X	MC- RR ppb (ng/g)	MC- Desm ethyl- RR* ppb (ng/g)	MC- LR ppb (ng/g)	MC- Desme thyl-LR ppb (ng/g)	MC- YR ppb (ng/g)	MC- LA ppb (ng/g)	MC- LW ppb (ng/g)	MC- LF ppb (ng/ g)	MC- LY ppb (ng/ g)	TOTA L_MC YST ppb (ng/g)	Anato xin A ppb (ng/g)	Domoi c acid ppb (ng/g)	Okadaic acid ppb (ng/g)
		OC																
L-685-09-4	HC	HC102609- OC	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-685-09-1	IB	IB102609-OC	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-685-09-5	OR	OR102609- OC	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-685-09-3	SV	SV102609- OC	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-685-09- 3Dup	SV-D	SV102609- OC	Karuk	26/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-691-09-2	тс	TC102909- OC	Yurok	29/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-691-09- 2Dup	тс	TC102909- OC	Yurok	29/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-691-09-3	WE	WE102909- OC	Yurok	29/Oct/2009	water	ND	ND	ND	ND	ND	ND	ND	ND	ND	0.00	ND	ND	ND
L-760-09-1	BB	BB121009-1	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	1.38	ND	ND	ND	1.38	ND	ND	ND
L-760-09- 1Dup	BB	BB121009-1	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	1.31	ND	ND	ND	1.31	ND	ND	ND
L-760-09-2	BB	BB121009-2	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	0.766	ND	ND	ND	0.766	ND	ND	ND
L-760-09-3	BB	BB121009-3	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	3.11	ND	ND	ND	3.11	ND	ND	ND
L-760-09-4	BB	BB121009-4	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	9.31	ND	ND	ND	9.31	ND	ND	ND
L-760-09-5	BB	BB121009-5	Karuk	10/Dec/2009	muss el	ND	ND	ND	ND	ND	32.3	ND	ND	ND	32.3	ND	ND	ND

APPENDIX II

MEMORANDUM

- TO: Elmer Dudik, Environmental Scientist
 North Coast Regional Water Quality Control Board
 5550 Skylane Boulevard, Suite A
 Santa Rosa, California 95403
- FROM: Regina Linville, Ph.D., Associate Toxicologist Integrated Risk Assessment Branch

DATE: June 10, 2008

SUBJECT: APPLYING RECOMMENDATIONS BY THE WORLD HEALTH ORGANIZATION TO THE ISSUE OF FISH CONSUMPTION FROM THE KLAMATH RIVER.

In response to a request by the North Coast Regional Water Board (Water Board), the Office of Environmental Health Hazard Assessment (OEHHA) is providing general guidance on the application of the provisional **tolerable daily intake (TDI)** of **microcystin-LR (MC-LR)** that has been recommended by the **World Health Organization (WHO)**. This guidance is based on WHO (1999) and is intended to assist the Water Board in addressing the 2008 bloom of cyanobacteria on the Klamath River. The provisional TDI provided by WHO (1999) is currently the best available guideline; no other major public health organization has published a TDI or comparable guideline for microcystin exposure at this time. The information presented here represents a general interpretation of WHO recommendations, but does not represent an official OEHHA recommendation.

The WHO has published a provisional TDI for MC-LR of 0.04 μ g/kg/day (WHO, 1999). The provisional TDI was derived following standard risk assessment protocol as described below.

- 1. The most appropriate study representing the relationship between dose and toxicological response was found to be a 13-week MC-LR oral exposure in mice (Fawell et al., 1994 as described in Fawell et al., 1999).
- The No Observed Adverse Effect Level (NOAEL) was identified. The NOAEL is determined as the highest dose of a chemical that does not induce the targeted toxic effect. In the Fawell et al. (1994) study, 40 μg MC-LR per kg of mouse body weight was the highest dose of MC-LR that did not cause liver injury.
- 3. An **uncertainty factor (UF)** of 1000 was applied to the NOAEL, as is typical when applying limited animal data to this type of human risk assessment. This UF incorporates three major areas of uncertainty: 1) humans could be more sensitive to MC-LR toxicity than mice, 2) some humans are potentially more sensitive to MC-LR than others (i.e., protection of the most sensitive people) and 3) additional studies could reveal adverse

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effects at lower dosages. The NOAEL was reduced by a factor of 1000 to safeguard against the three uncertainties described above, resulting in a TDI of $0.04 \mu g/kg$ -day.

The WHO TDI can be used to calculate the maximum recommended consumption rate of fish or shellfish from an affected water body. For a seasonal bloom such as that occurring on the Klamath, this represents a conservative approach since the TDI is based on a longer exposure period than is expected during the bloom season. The study used by WHO (1999) to derive the TDI (Fawell et al., 1994) is categorized as a chronic study because the exposure lasted more than ten percent of the organism's average lifespan (13 weeks corresponds to roughly 13 percent of the average life span in mice). Seasonal exposures to microcystin in humans are more likely to be categorized as subchronic (exposures between 1 month and ten percent of lifespan) or short-term (exposures up to 30 days). Another issue is the application of a TDI derived for one microcystin variant (MC-LR) to all of the microcystin variants present in the bloom. Although many variants have been shown to cause acute toxicity similar to MC-LR, WHO (1999) determined that the existing data on other variants was insufficient to derive TDI values. The TDI was used to derive a consumption rate as follows:

 $CR (kg/day) = \frac{TDI (\mu g/kg-day) * BW (kg)}{MC_{edible} (\mu g/kg)}$

where,

CR = consumption rate, TDI = tolerable daily intake, BW = body weight, $MC_{edible} = concentration of MC in edible portions of fish or shellfish (fresh weight).$

To calculate CR as oz/month use:

 $CR (oz/mo.) = \frac{TDI (\mu g/kg-day) * BW (kg) * 1,073 oz/mo^{\dagger}}{MC_{edible} (\mu g/kg)}$

[†] Calculated as follows: 1 kg/day * 30.4 days/mo * 35.3 oz/kg = 1073 oz/mo.

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And finally, to calculate CR as the number of 8-oz meals/month use:

 $CR (\# 8-oz meals/mo.) = \frac{TDI (\mu g/kg-day) * BW (kg) * 134 meals/mo^{*}}{MC_{edible} (\mu g/kg)}$ * Calculated as follows: 1 kg/day * 30.4 days/mo * 35.3 oz/kg * 1 meal/8oz = 134 8-oz meals/mo.

The above equation was used to calculate the maximum number of 8-oz meals per month from affected areas, with the results shown in the following table (exact values shown in parentheses):

Microcystin (µg/kg)	Adult [‡]	Child [§]						
10	> 30 (38)	5 (5.4)						
20	19 (19)	3 (2.7)						
40	9 (9.4)	1 (1.3)						
60	6 (6.3)	< 1 (0.9)						
80	5 (4.7)	< 1 (0.7)						
100	4 (3.8)	< 1 (0.5)						
200	2 (1.9)	0 (0.3)						
300	1 (1.3)	0 (0.2)						
400	1 (0.9)	0 (0.1)						
500	< 1 (0.8)	0 (0.1)						
[‡] Adult = 70 kg (~155 lbs)	[§] Child = 10 kg (~ 22 lbs)							

References

Fawell, J.K., James, C.P. and James, H.A. 1994 Toxins from Blue-Green Algae: Toxicological Assessment of Microcystin-LR and a Method for its Determination in Water, Water Research Centre, Medmenham, UK.

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- cc: James C. Carlisle, D.V.M., Chief
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