

TECHNICAL MEMORANDUM

Date: February 26, 2016

To: Brian Ross, USEPA Region 9, Dredging & Sediment Management Team

Beth Christian, San Francisco Bay Regional Water Quality Control Board

From: Don Yee and Phil Trowbridge, SFEI

Re: Updated ambient concentrations of toxic chemicals in San Francisco Bay area

sediments

Introduction

The purpose of this memo is to communicate the most recent results for the ambient concentrations of mercury, Total PAHs and Total PCBs in San Francisco Bay area sediments.

Regulation of dredging and dredged material disposal or reuse under the San Francisco Bay Long Term Management Strategy for Dredging (LTMS) involves, in part, evaluation of sediment chemistry characteristics. For dredged material in California, for most chemicals, there are no numeric sediment quality standards or objectives that define chemical suitability for disposal or reuse at particular placement sites. Consequently, Bay dredged sediments are compared to various published screening values and other guidelines, to reference site chemistry, and where available and appropriate, to ambient sediment contaminant concentrations for the Bay as a whole. Therefore, starting in 2011, the LTMS agencies have asked SFEI to calculate the ambient concentrations for a small subset of sediment contaminants that are either subject to TMDL limits for in-Bay dredged material disposal or are relevant for bioaccumulation testing for dredging (DMMO, n.d.).

It is important to note that even if contaminant concentrations in a dredged material project are below ambient values, this alone does not indicate that the dredged material is suitable for placement at any particular site. Ambient levels may be higher than environmentally desirable for some placement sites, while at other sites ambient concentrations may be far below levels of concern that need to be regulated. Ambient concentrations by themselves are not sediment screening values, and should not be used directly as guidelines for any particular regulatory use. However, understanding how potential regulatory goals or limits under consideration may relate

¹ Pursuant to the Water Quality Control Plan for Enclosed Bays and Estuaries - Part 1 Sediment Quality, California's Sediment Quality Objectives (SQOs) do not apply to dredged material suitability determinations. However, strict sediment chemistry limits may be established for individual compounds or locations under Bay-wide TMDLs, site-specific remediation decisions, etc.

to existing Bay ambient sediment concentrations can be important in informing regulatory decisions that are both effective and feasible to implement.

Methods

For this report, RMP staff calculated updated the ambient concentrations for mercury, polycyclic aromatic hydrocarbons (PAHs), and polychlorinated biphenyls (PCBs) using data through 2014. Ambient concentrations were calculated for individual PAH compounds and PCB congeners routinely reported by the RMP, as well as the Total PAHs and Total PCBs.

The previous report on ambient sediment concentrations (SFEI, 2015) was based on data through 2012. Another round of sediment data were collected in 2014, but not 2013. Therefore, to update the ambient concentrations, the 2014 dataset was added to the data considered for the last report and then the combined dataset was reanalyzed. Specifically, for PAHs and PCBs, the dataset included the years from 2002-2014 (excluding data from 2004-2006 that was censored due to QA issues). For mercury, the previous calculations were based on data from 2003-2012, so the current update used mercury data from 2003-2014.

The approach taken for this update was consistent with the recommendations from SFEI (2015) that calculations of ambient concentrations use data from the most recent 13-year period, instead of the 10-year period that had been used previously. The frequency of RMP sediment sampling has been changed to every 4 years. A 13-year time window is needed so that the calculations for each update would be based on at least 4 sediment sampling events. The 13-year trailing window from the most recent RMP data is 2002-2014. The data for PAHs and PCBs used for this update exactly matched this window. For mercury, data used for the update were from 2003-2014, which falls within this window.

The ambient values calculated are the 90% upper tolerance limit of the 90th percentile concentrations (U.S. Army Corps of Engineers and US Environmental Protection Agency, 2011). To avoid skewing the results, data from historical fixed sampling stations that are not part of the RMP randomized sampling design, including all pre-2002 data, have been excluded. Outliers were removed to exclude any highly contaminated samples, using an updated methodology previously described in the 2015 report (and summarized in Appendix A). The RMP visually selects for fine-grained sediments during field sampling; thus the ambient concentrations are representative of sediments with greater than 40% fines (which is typical of most sediments dredged in the Bay area).

The updated results for mercury, Total PAHs and Total PCBs are summarized in a simple table for the RMP website (Table 1). For comparison, the concentrations for these parameters from the 2015 report are shown in Table 2.

The updated concentrations for mercury, Total PAHs, and Total PCBs plus updated concentrations for individual PAH and PCB congeners are shown in a master table in Appendix B. Appendix B also includes the ambient concentrations for all other analytes updated in the last report (SFEI, 2015) but are unchanged here. Appendix C contains histograms of the latest results for all analytes.

Results

Table 1. Ambient contaminant concentrations in San Francisco Bay area sediments calculated from data between 2002 and 2014 for Mercury, Total PAHs (and HPAH/LPAH subtotals), and Total PCBs.

	1998 Ambient Sediment Values	90th Percentile 90% UTL (2015 Ambient Value)	99th Percentile 90% UTL ¹
MERCURY (2003-2014) (mg/kg dw)			
Mercury	0.43	0.332	0.47
PAH (2002-2003, 2007-2014) (ug/kg dw)			
Total PAH	3390	4520	9090
Total HPAH	3060	3640	5830
Total LPAH	434	568	746
PCB (2002-2003, 2007-2014) (ug/kg dw)			
Total PCB (sum of 40 congeners)	21.6	18.2	29.5

¹99th percentile required only for Mercury and PCBs. Indicates approximate (non-outlier) upper limit for other contaminants.

Table 2. Ambient contaminant concentrations in San Francisco Bay area sediments calculated from data between 2002 and 2012 for Mercury, Total PAHs (and HPAH/LPAH subtotals), and Total PCBs as reported in SFEI (2015).

	1998 Ambient Sediment Values	90th Percentile 90% UTL (2014 Ambient Value)	99th Percentile 90% UTL ¹
METALS (2003-2012) (mg/kg dw)			
Mercury	0.43	0.332	0.47
PAH (2002-2003, 2007-2012) (ug/kg dw)			
Total PAH	3390	4540	9100
Total HPAH	3060	3870	5830
Total LPAH	434	574	946
PCB (2002-2003, 2007-2012) (ug/kg dw)			
Total PCB (sum of 40 congeners)	21.6	18.3	29.6

¹99th percentile required only for Mercury and PCBs. Indicates approximate (non-outlier) upper limit for other contaminants.

As would be expected based on the calculation algorithm (which includes multiple years to provide some stability), only small changes to each of the ambient concentrations were observed from the last report.

After receiving your approval, RMP staff will add the updated concentrations in Table 1 to the RMP's Dredged Material Testing Thresholds website: http://www.sfei.org/projects/dmmo-ambient-sediment-conditions.

The next RMP sediment sampling event will occur in 2018. Once the data are finalized, the ambient concentrations for mercury, PAHs and PCBs will be updated using data from the 13-year window between 2006 and 2018.

References

- Dredged Material Management Office (n.d.). Retrieved from http://www.spn.usace.army.mil/Missions/DredgingWorkPermits/DredgedMaterialManagementOffice(DMMO).aspx
- San Francisco Bay Regional Water Quality Control Board (SFBRWQCB) staff. May 1998. Staff Report: Ambient Concentrations of Toxic Chemicals in San Francisco Bay Sediments. Oakland, CA.
- SFEI. 2015. Updated Ambient Concentrations of Toxic Chemicals in San Francisco Bay Sediments. San Francisco Estuary Institute, Richmond, CA. Contribution # 749. Published online: http://www.sfei.org/documents/updated-ambient-concentrations-toxic-chemicals-san-francisco-bay-sediments.
- U.S. Army Corps of Engineers, U.S. EPA Region 9. Memo to: Robert S. Hoffmann, National Marine Fisheries Service, Long Beach, CA. 9 June 2011.

Appendix A: Addendum on Auto-Outlier Detection Function

Using the original coding of the R script for identifying outliers, addition of RMP PCB data from 2011 to the previous data set (2002-2010) resulted in a statistical artifact that caused a large percentage of the data (~11%) to be regarded as outliers; previous years' analyses had most of these same points not as outliers (1% or less of the data as outliers). This arose due to a randomly occurring tight cluster of values in one portion of the distribution. The outlier function used a moderately small percentage of the total data set (2.5%, with a minimum of 12 values) to determine a local average difference expected between points; this very tight cluster thus resulted in a gap to the next higher value (between 15.9 and 16.7 ppb) previously considered unremarkable (<10x the local average difference) in previous years' data, being labeled as the gap to a set of outliers when the 2011 data were included.

To prevent future flagging of such chance statistical anomalies as outlier gaps, the following modifications to the outlier auto-detection function have been made:

- 1. A larger percentage of the nearest lower values (5% of the total data) are used to calculate a local average difference. Increasing from 2.5 % to 5% of the population decreases the influence of any small cluster of tightly spaced points (the anomaly seen in the 2002-2011 set). However, in the upper tail of the data distribution where we usually expect sparser spacing, using a larger % of the population decreases the apparent local difference by averaging in some more tightly spaced points in the main body of the distribution, potentially flagging more gaps. This change stabilizes the local average difference, decreasing the probability of finding gaps in the main body of the distribution, but increasing the likelihood in sparse areas of the upper tail.
- 2. Any outlier has to be at least 25% greater than the mean of its nearest lower neighbors (a running average 10% of the total data) to be considered "meaningful". In many laboratory chemical analyses, a 25% difference in replicate measurements is acceptable, so the difference between a possible outlier and its nearest lower neighbors should be at least as large to account for measurement uncertainty. The difference to a running average of nearest lower neighbors rather than a single nearest lower neighbor is used, as otherwise a few values between low and high concentration groups could bridge apparent gaps, causing no outliers to be found. The use of 10% of the data for the running average further ensures that a small set of sparse values cannot bridge the zone between low and high concentration groups.
- 3. Since our concern is about high outlier values, any outlier has to occur above the upper quartile value. This avoids a case where a minority of very low values (e.g., clean sites near Golden Gate) would create an apparent gap to the main population of results, causing all of the latter to be called outliers. Using the upper quartile as the threshold, a bimodal distribution where a slight majority of the data is in the lower group would still result the upper portion labeled as outliers.

Imposing all three of the modifications above makes the detection of outlier groups more stable and resistant to chance anomalies in the data distribution. The current data set now has 10 years of data, and future calculations will only include the most recent 10 sets of data. Existing outlier groups are therefore unlikely to change just by the filling in of gaps using new data with similar concentration distributions; only larger and real shifts in distribution (e.g., 5% or more of the total data bridging the gaps) will cause appreciable shifts in the subset of points being identified as outliers.

Appendix B Full List of Ambient Values (including previous updates)

Table B-1. Ambient contaminant concentrations in San Francisco Bay area sediments calculated using data through 2014 for mercury, PAHs, and PCBs.

	1998 Ambient Sediment Values	90th Percentile 90% UTL (2015 Ambient Value)	99th Percentile 90% UTL ²		
MERCURY (2003-2014) (mg/kg dw)					
Mercury	0.43	0.332	0.469		
	PAH (2002-2003, 2007-2014) (ug/kg dw)				
Total PAH	3390	4520	9090		
Total HPAH	3060	3640	5830		
Total LPAH	434	568	746		
Acenaphthene	26.6	13.6	20.3		
Acenaphthylene	31.7	34.0	52.2		
Anthracene	88	80.5	110		
Benz(a)anthracene	244	212	252		
Benzo(a)pyrene	412	395	746		
Benzo(b)fluoranthene	371	225	280		
Benzo(e)pyrene	294	244	467		
Benzo(g,h,i)perylene	310	408	654		
Benzo(k)fluoranthene	258	229	452		
Biphenyl	12.9	11.8	17.0		
Chrysene	289	243	481		
Dibenz(a,h)anthracene	32.7	47.8	106		
Dibenzothiophene	NA	16.3	28.4		
Dimethylnaphthalene, 2,6-	12.1	12.9	17.6		
Fluoranthene	514	562	791		
Fluorene	25.3	27.2	57.3		
Indeno(1,2,3-c,d)pyrene	382	336	490		
Methylnaphthalene, 1-	12.1	13.8	20.1		
Methylnaphthalene, 2-	19.4	20.8	29.8		
Methylphenanthrene, 1-	31.7	37.8	70.2		
Naphthalene	55.8	56.2	70.6		
Perylene	145	213	319		
Phenanthrene	237	173	253		
Pyrene	665	783	1115		
Trimethylnaphthalene, 2,3,5-	9.8	7.31	9.77		

Total PCB (sum of 40 congeners)	21.6	18.2	29.
PCB 8	NA	0.143	0.29
PCB 18	NA	0.0668	0.094
PCB 28	NA	0.284	0.43
PCB 31	NA	0.134	0.22
PCB 33	NA	0.0819	0.14
PCB 44	NA	0.333	0.46
PCB 49	NA	0.249	0.30
PCB 52	NA	0.39	0.71
PCB 56	NA	0.135	0.23
PCB 60	NA	0.0661	0.13
PCB 66	NA	0.467	0.80
PCB 70	NA	0.58	0.94
PCB 87	NA	0.462	0.60
PCB 95	NA	0.6	1.2
PCB 99	NA	0.648	1.2
PCB 101	NA	1.11	2.2
PCB 105	NA	0.364	0.48
PCB 110	NA	1.04	1.9
PCB 118	NA	0.979	1.3
PCB 128	NA	0.276	0.50
PCB 132	NA	0.37	0.44
PCB 138	NA	1.84	2.6
PCB 141	NA	0.207	0.30
PCB 149	NA	1.26	2.5
PCB 151	NA	0.562	1.2
PCB 153	NA	1.82	2.6
PCB 156	NA	0.164	0.22
PCB 158	NA	0.147	0.21
PCB 170	NA	0.459	0.56
PCB 174	NA	0.468	0.67
PCB 177	NA	0.359	0.53
PCB 180	NA	1.02	2.1
PCB 183	NA	0.356	0.47
PCB 187	NA	0.852	1.1
PCB 194	NA	0.328	0.40
PCB 195	NA	0.112	0.13
PCB 201	NA	0.0551	0.11
PCB 203	NA	0.181	0.39

Table B-2. Ambient contaminant concentrations in San Francisco Bay area sediments for other analytes calculated using data through 2012 and reported in SFEI (2015).

	1998 Ambient Sediment Values	90th Percentile 90% UTL (2014 Ambient Value)	99th Percentile 90% UTL ²
METALS (2003-2012) (mg/kg dw)			
Arsenic	15.3	13.9	19
Cadmium	0.33	0.329	0.483
Chromium ¹	112	NA	NA
Copper	68.1	53.9	71.2
Lead	43.2	25.1	33.2
Nickel	112	98.3	118
Selenium	0.64	0.359	0.493
Silver	0.58	0.315	0.496
Zinc	158	136	162
PESTICIDES (2002-2003, 2007-2012)	(ug/kg dw)		
Aldrin	NA	0.0335	0.0585
Total Chlordane	1.1	0.342	0.599
Dieldrin	0.44	0.157	0.203
Endrin	NA	0.00897	0.0177
Total DDT (total of 6 isomers)	7	4.68	6.29
DDD(o,p')	NA	0.509	0.742
DDD(p,p')	NA	1.98	2.74
DDE(o,p')	NA	0.109	0.221
DDE(p,p')	NA	1.98	2.92
DDT(o,p')	NA	0.0371	0.0593
DDT(p,p')	NA	0.265	0.599
Total HCH	0.78	0.0512	0.115
Hexachlorobenzene	0.48	0.196	0.416
PBDE (2002-2003, 2007-2012) (ug/kg dw)			
Total PBDE	NA	5.47	21.4
DIOXIN (2008-2010) (ug TEQ/kg dw)			
Sum of Dioxins TEQ	NA	0.00183	0.00269
Sum of Furans TEQ	NA	0.00107	0.00135

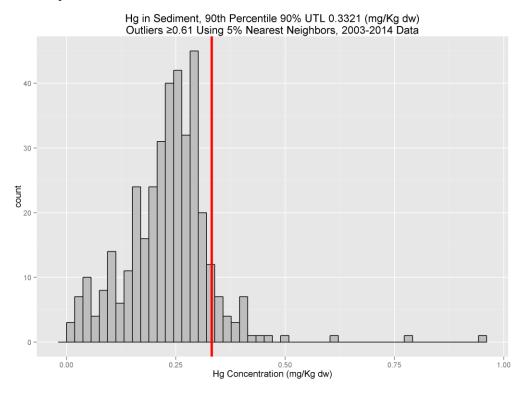
NA = Data Not Available

¹RMP stopped monitoring Cr in 2002 because it was not a concern for water quality in the Bay.

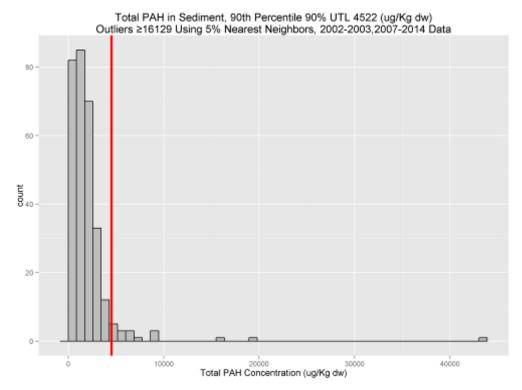
²99th percentile required only for Mercury and PCBs. Indicates approximate (non-outlier) upper limit for other contaminants.

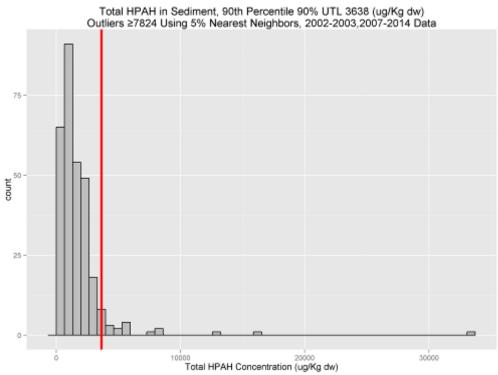
Appendix C Histograms for Individual Analytes

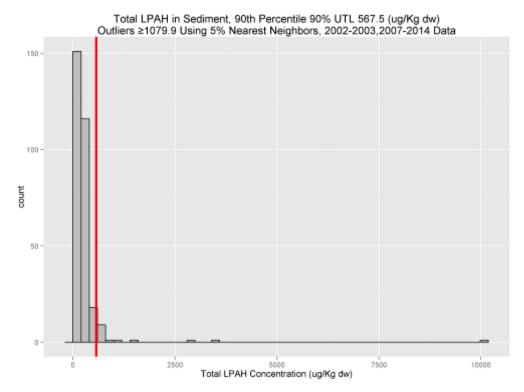
C-1 Short List Contaminants (changed since 2014 update) Mercury

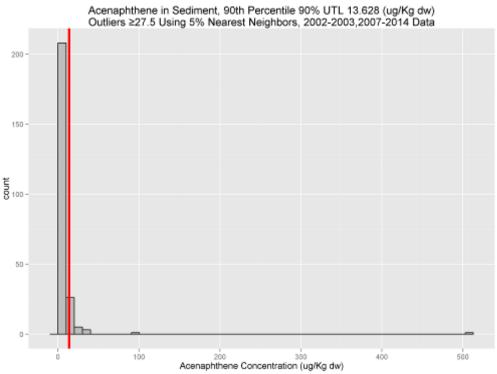


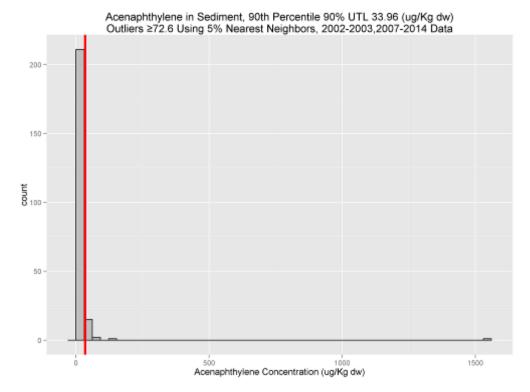
PAHs

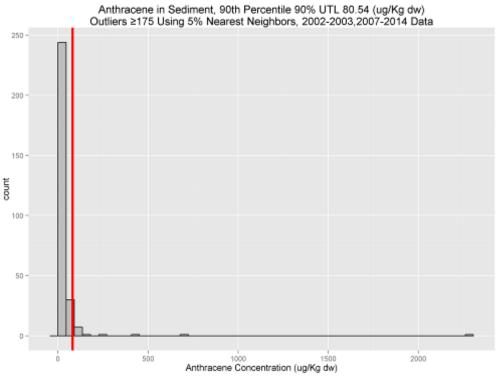


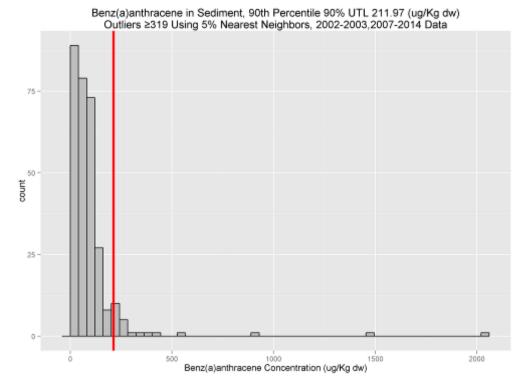


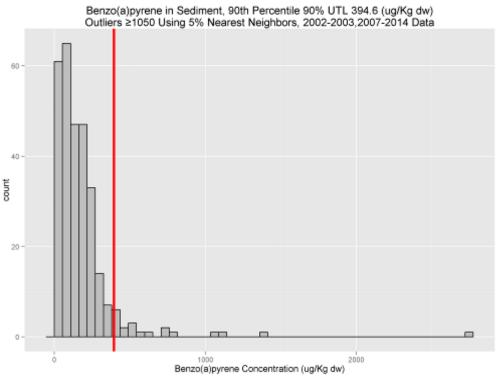


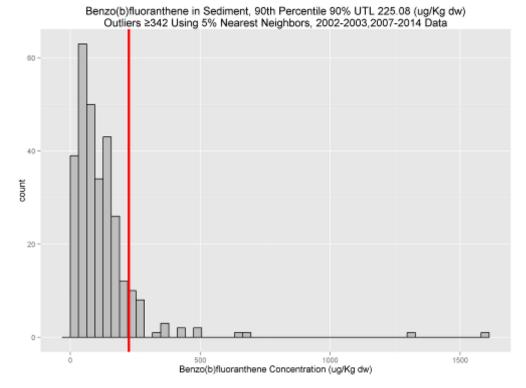


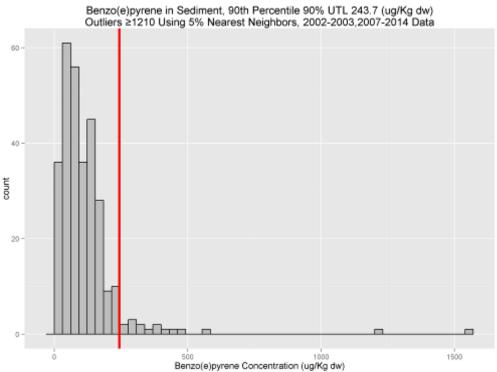


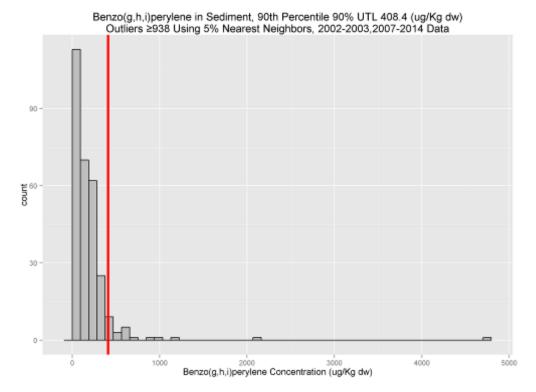


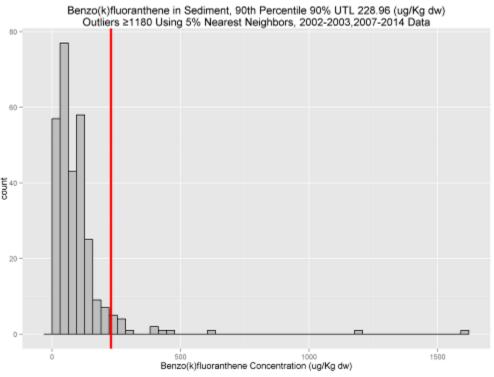


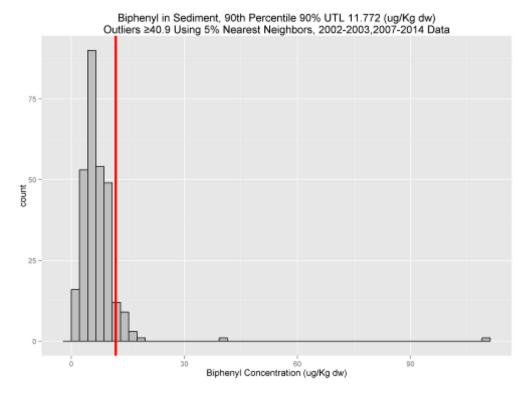


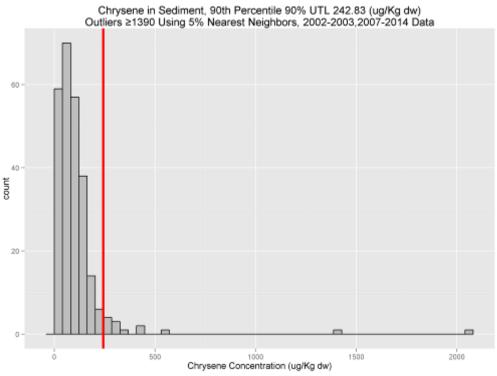


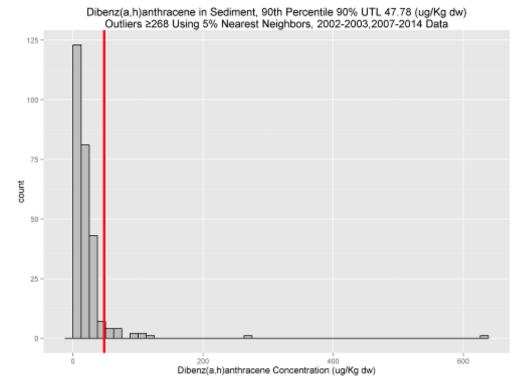


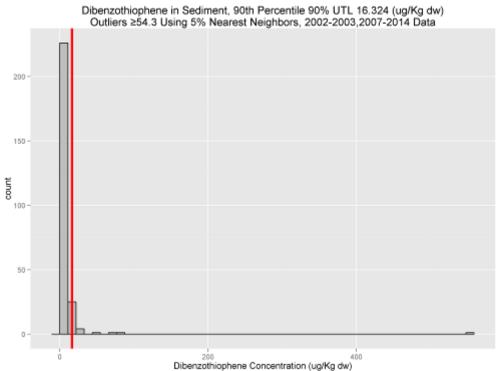


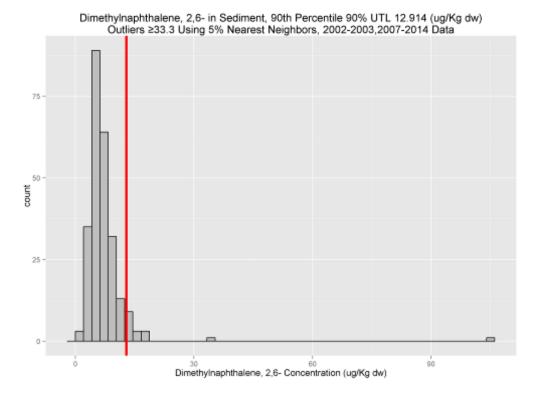


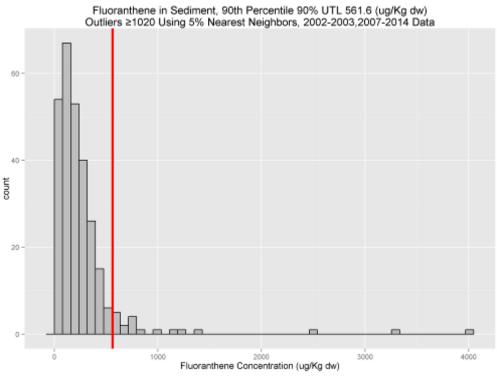


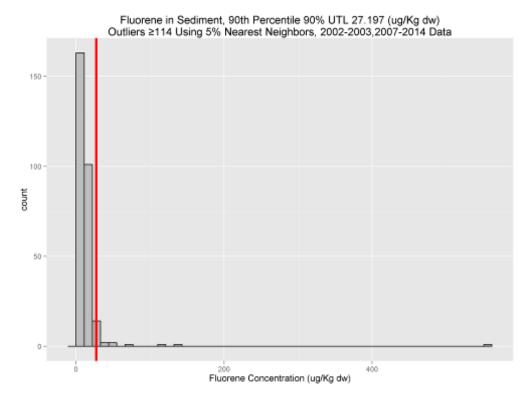


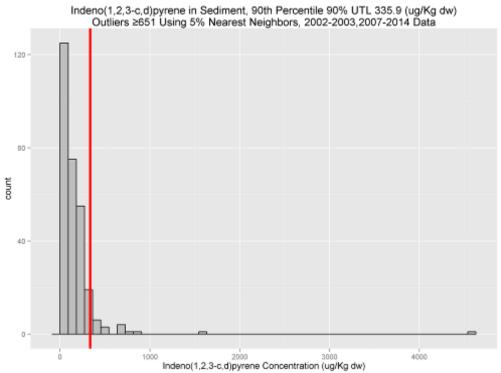


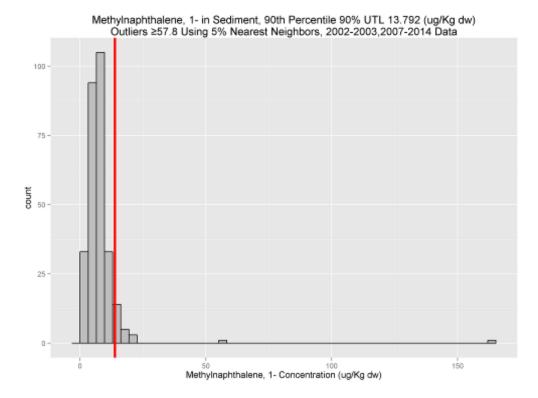


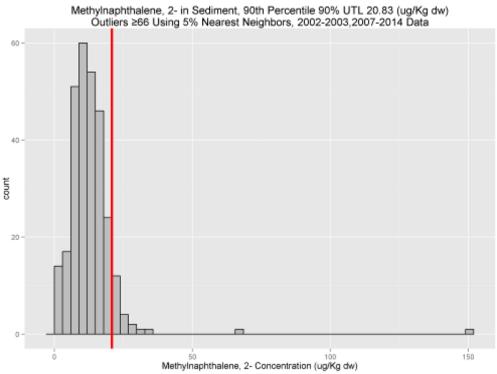


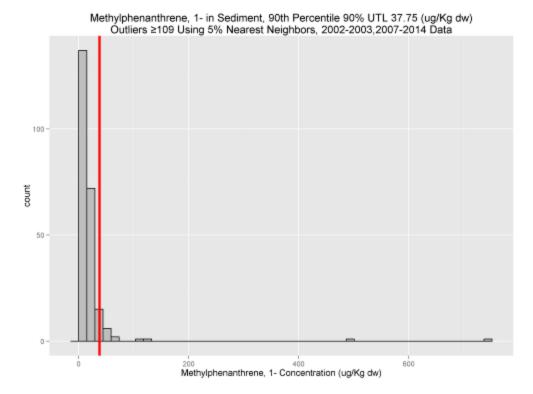


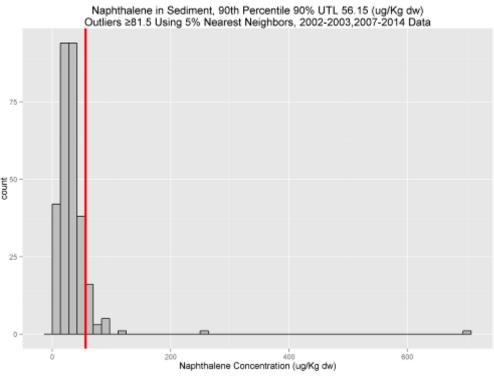


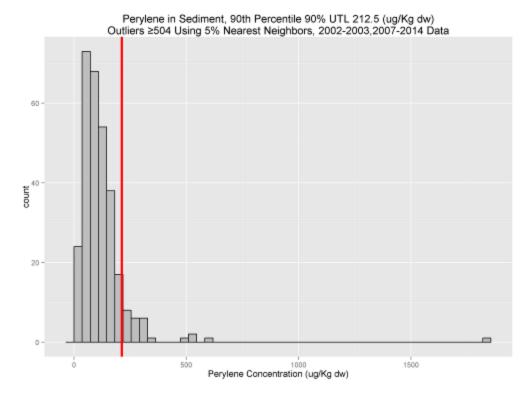


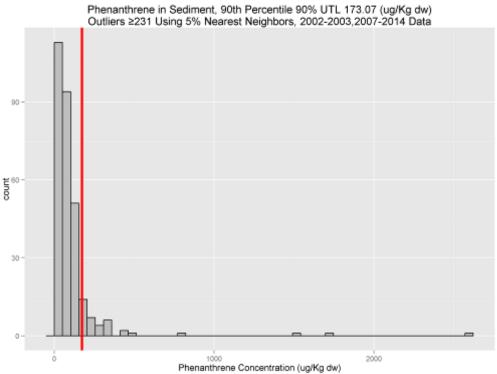


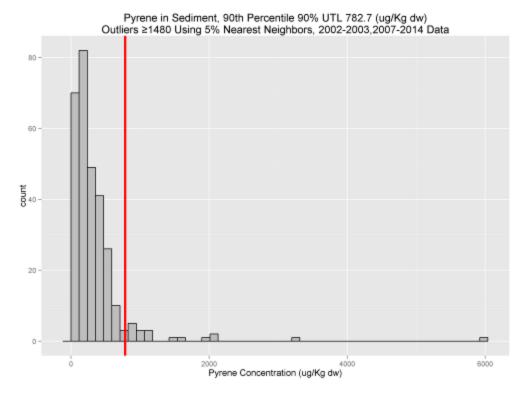


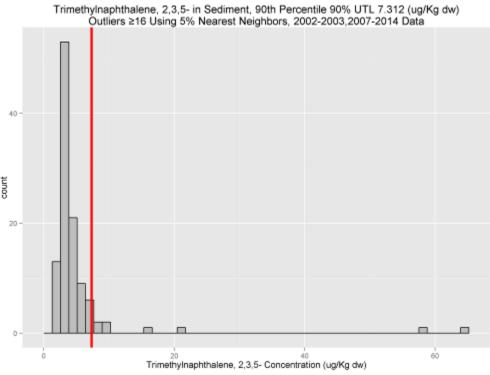




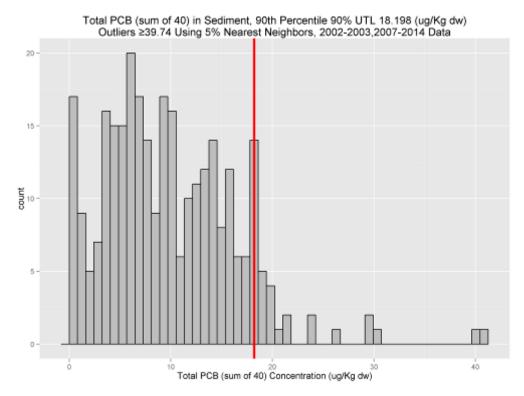


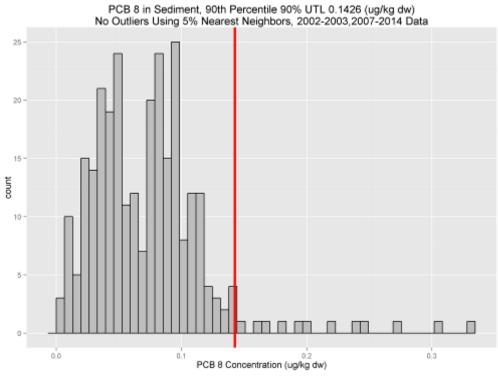


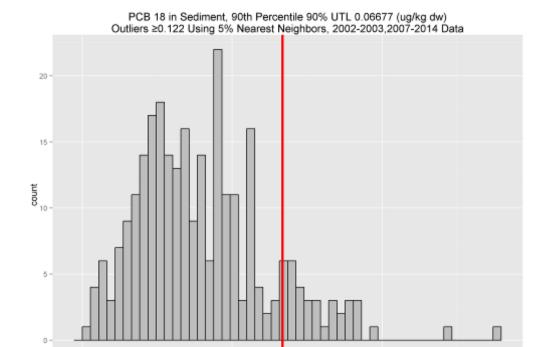




PCBs

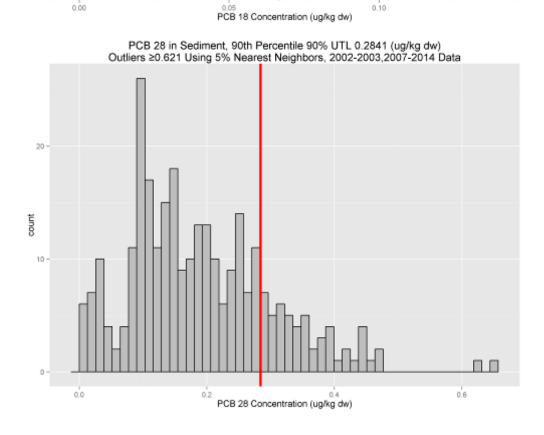


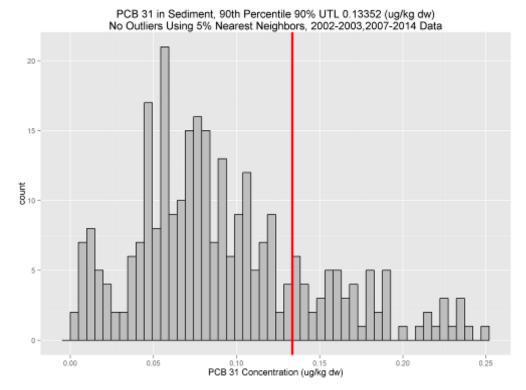


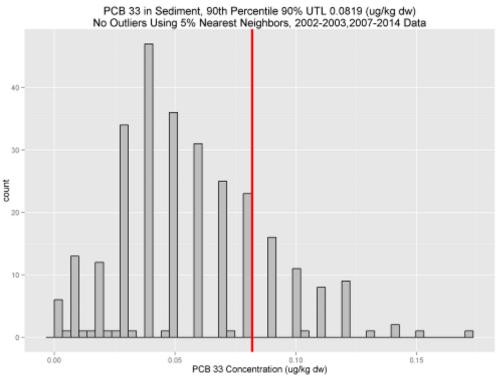


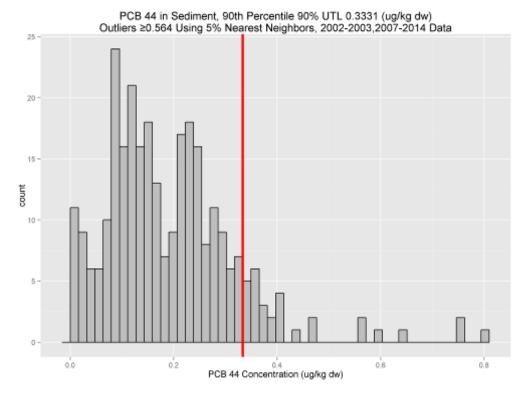
0.10

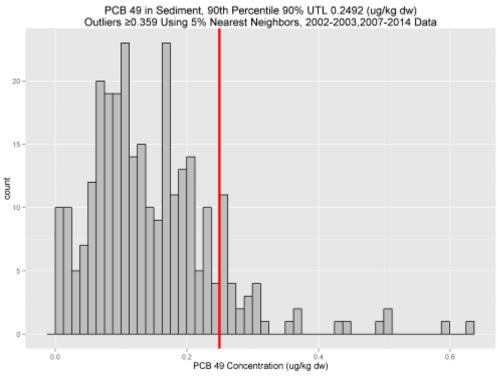
0.00

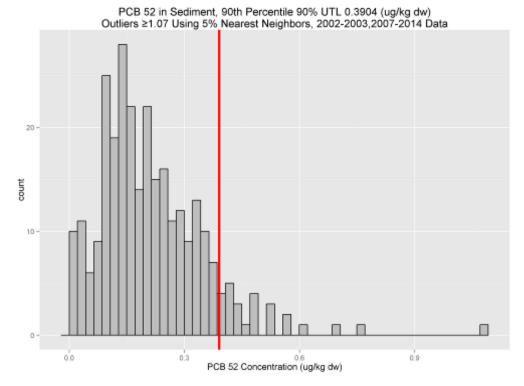


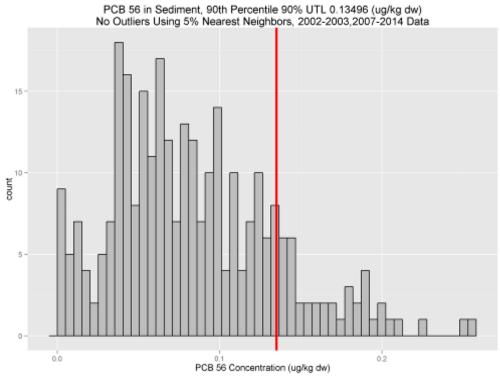


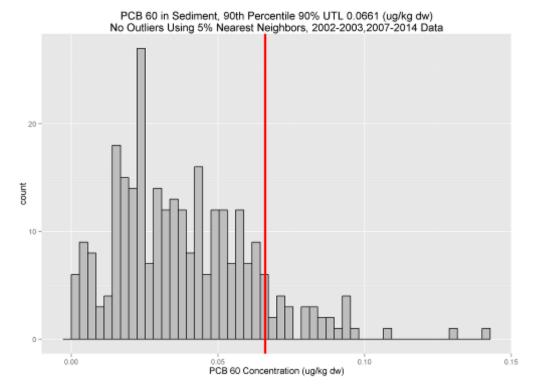


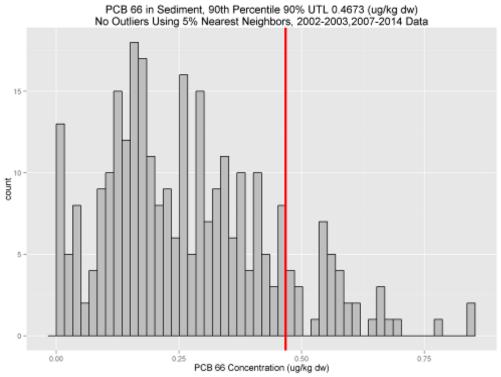


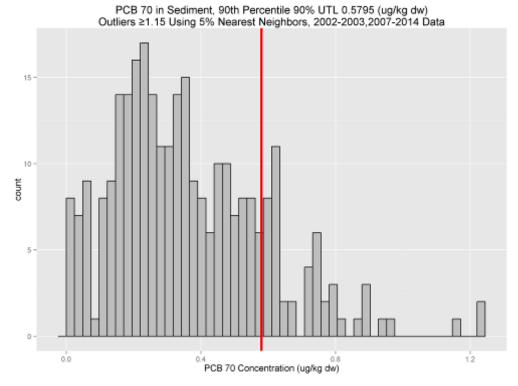


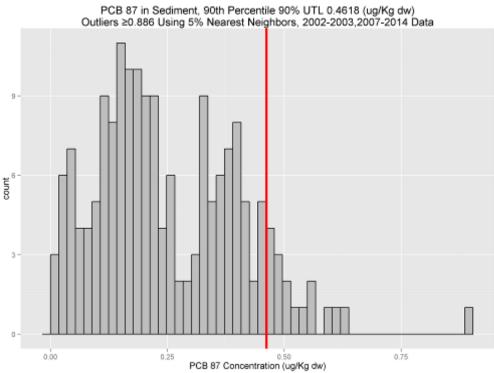


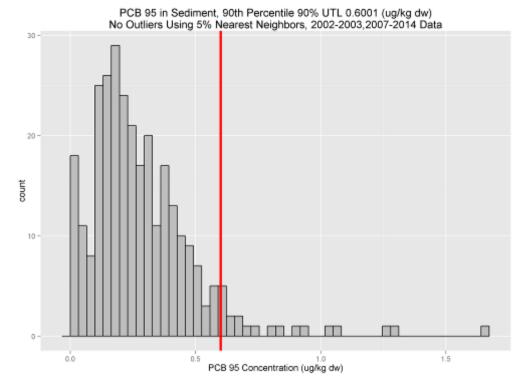


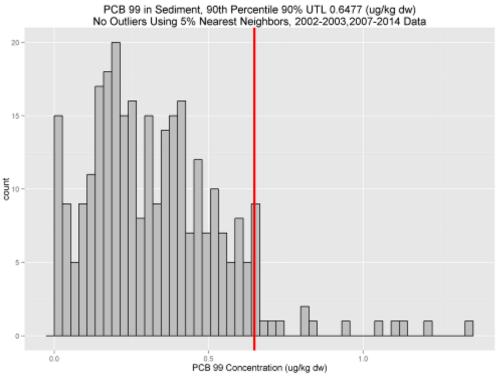


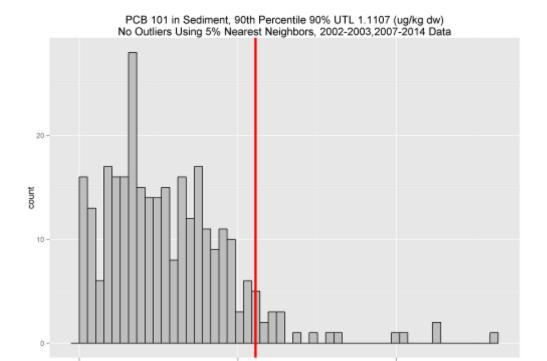


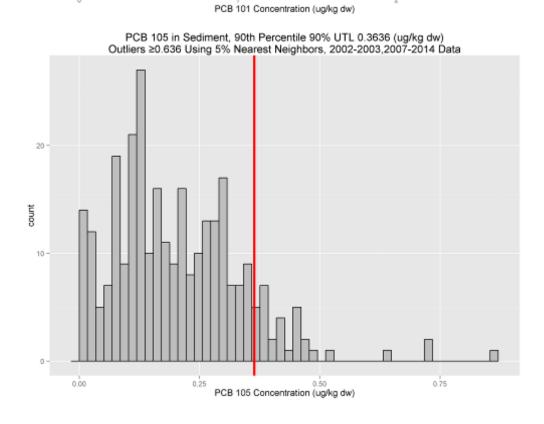


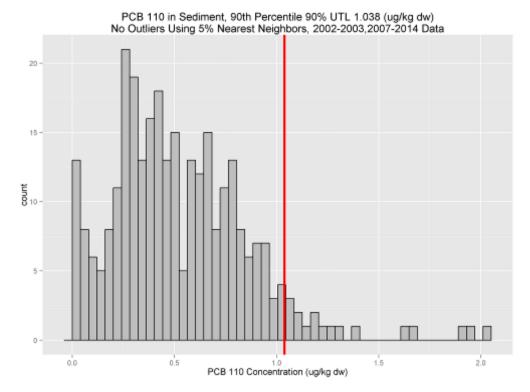


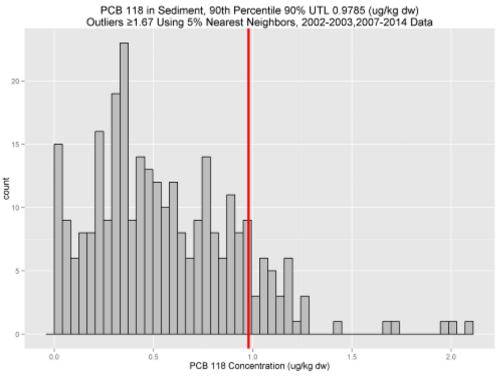


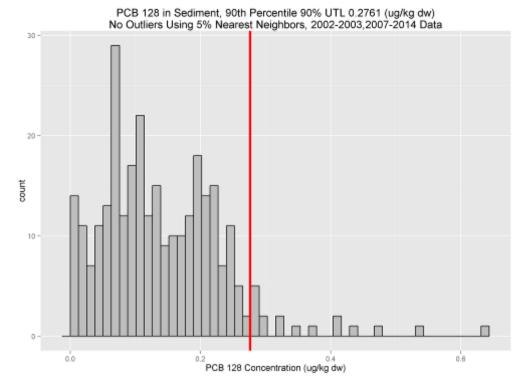


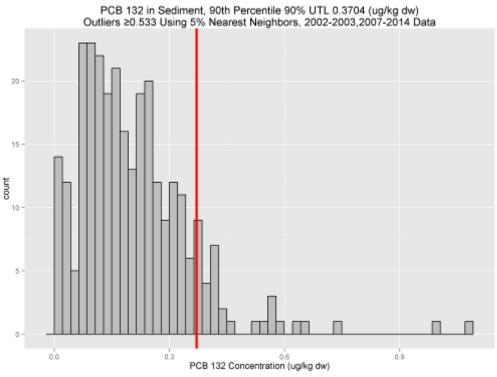


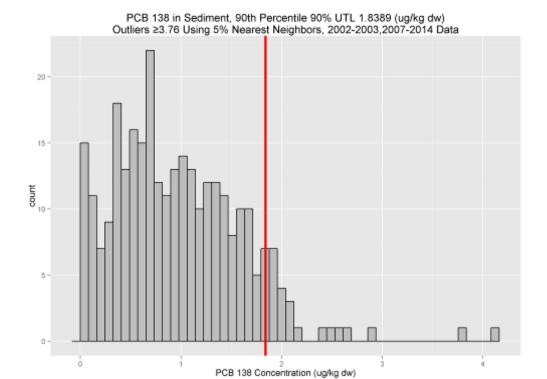


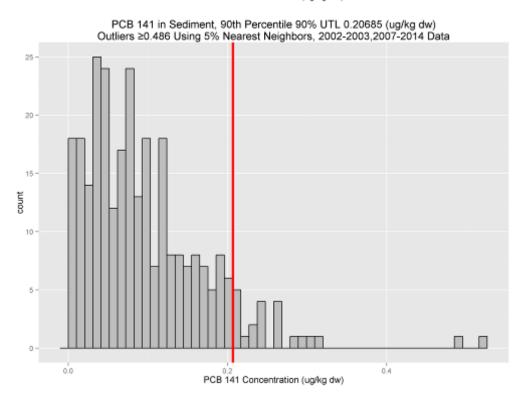


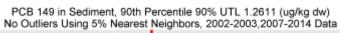


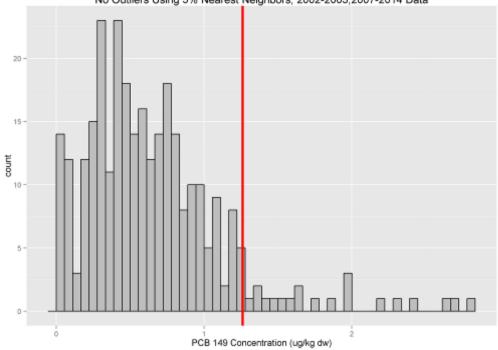


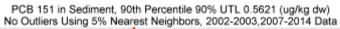


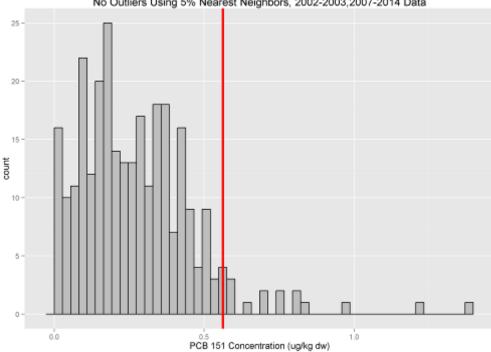


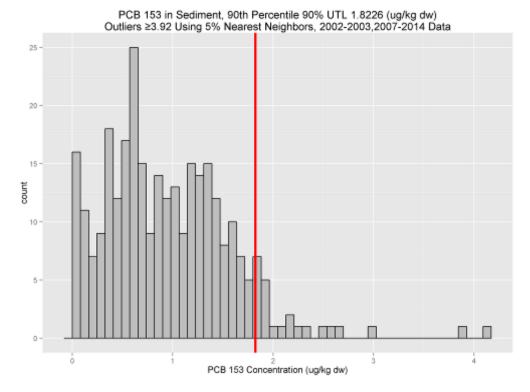


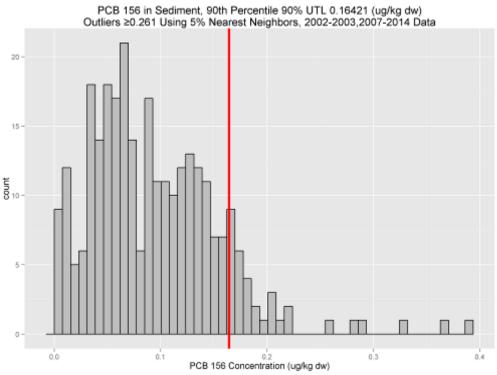


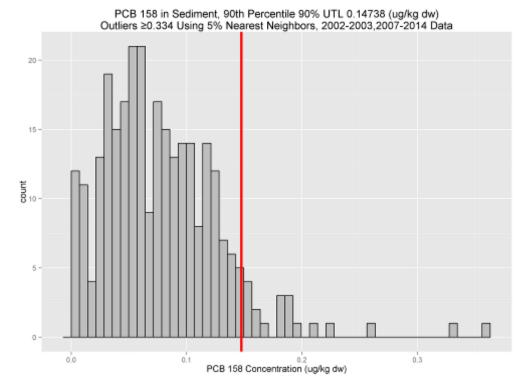


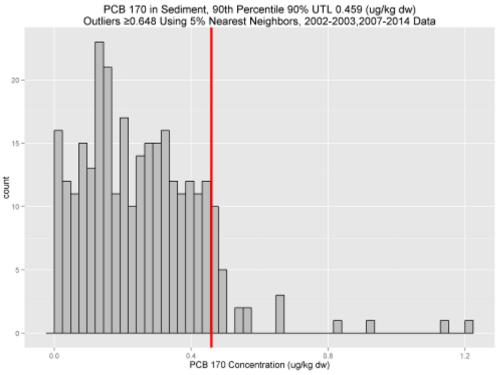


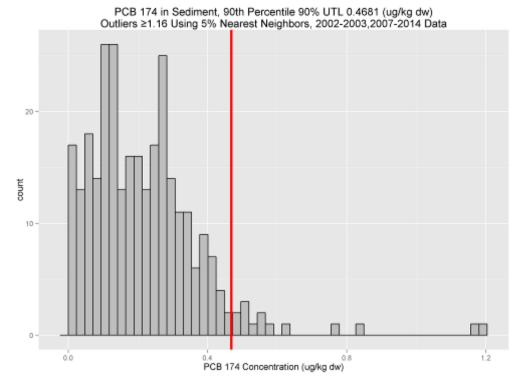


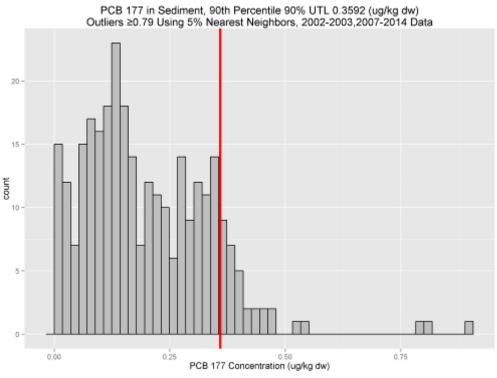


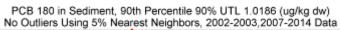


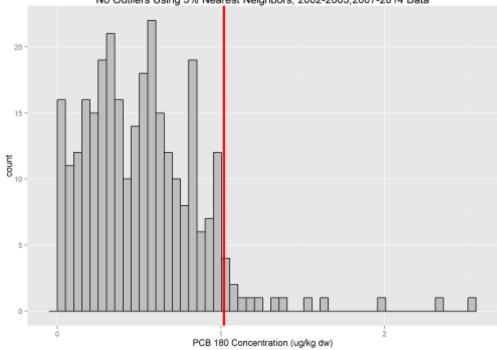


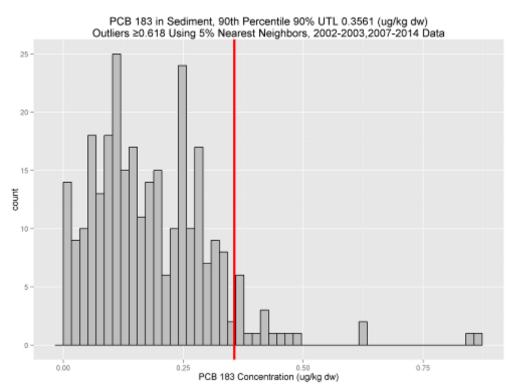


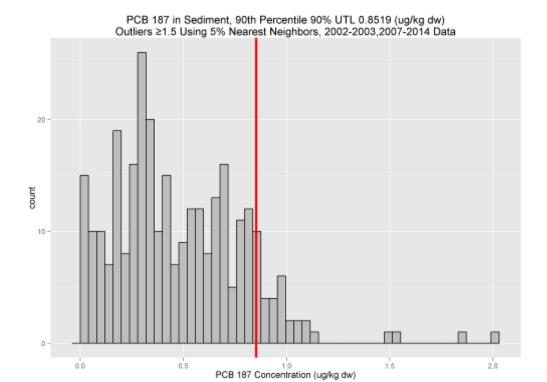


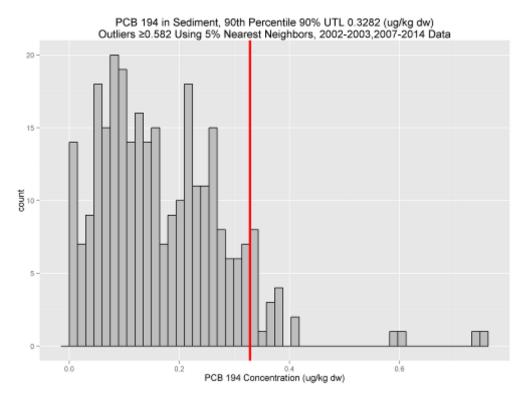


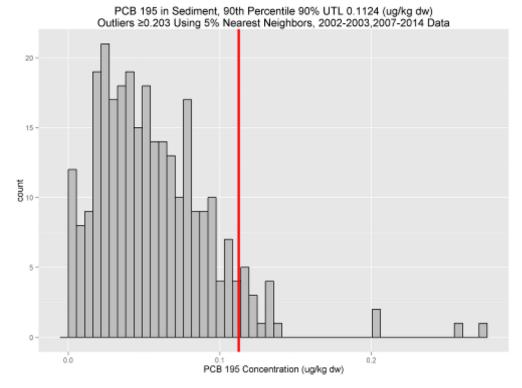


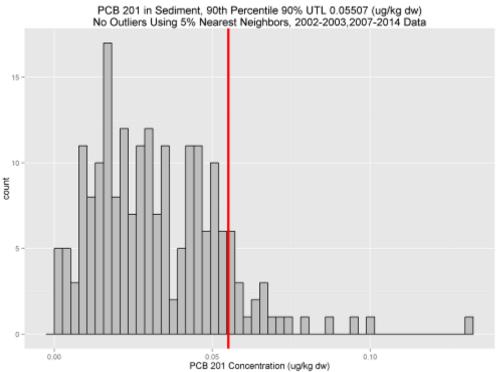


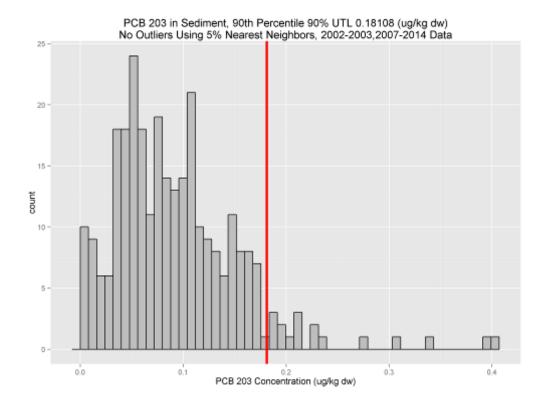




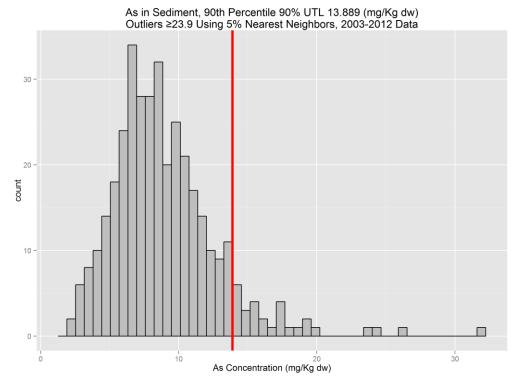


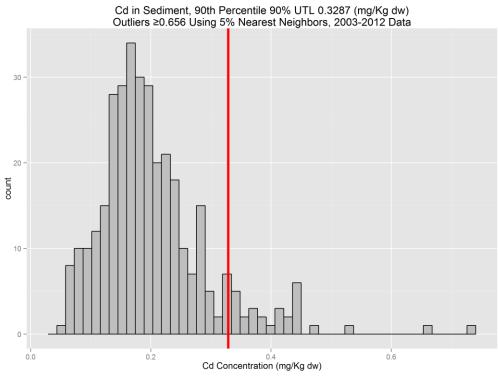


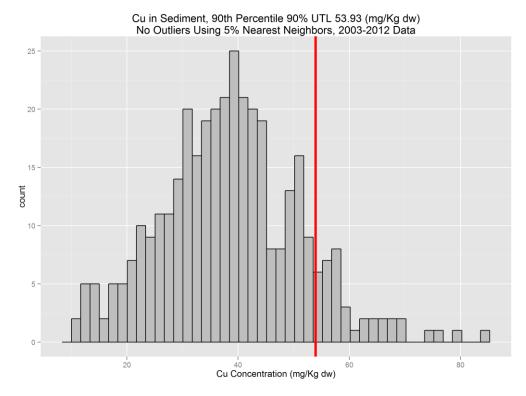


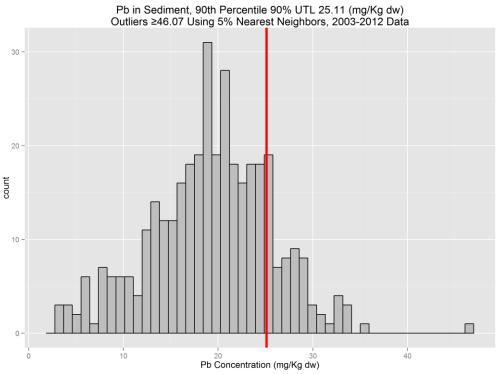


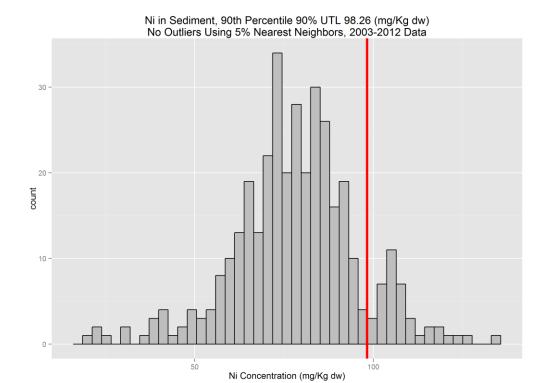
C-2 Remaining Full List Contaminants (unchanged from 2014 update) Trace Metals

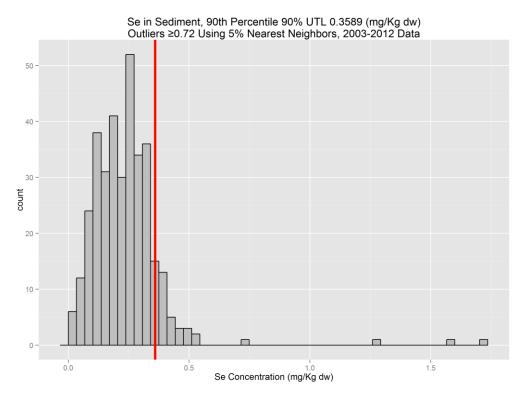


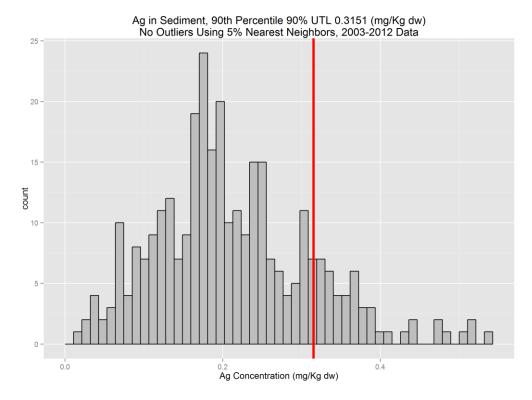


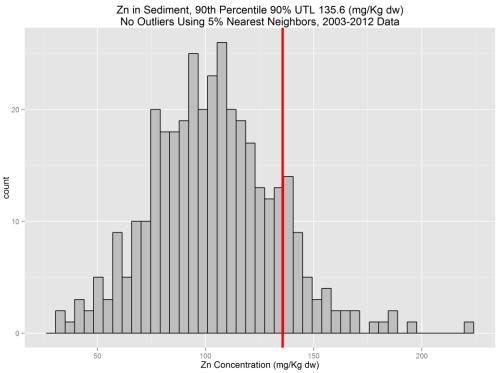




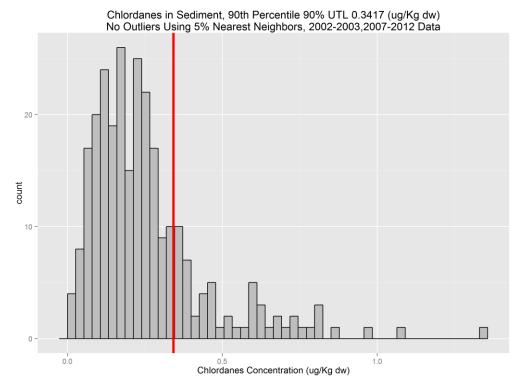


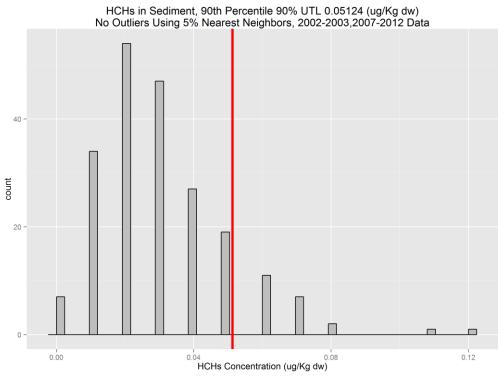


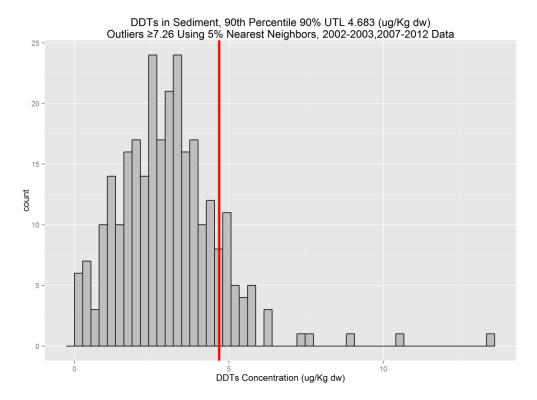


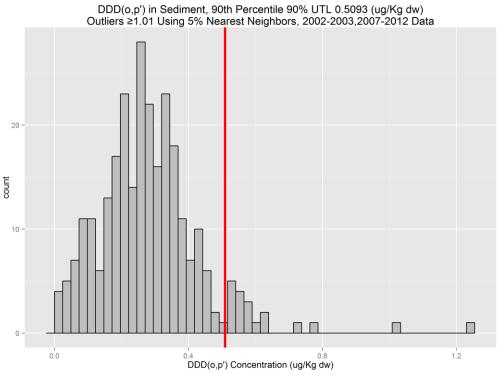


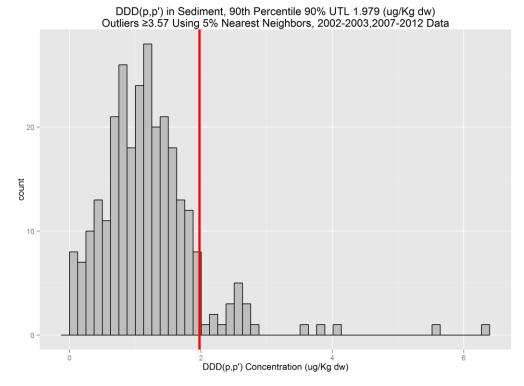
Pesticides

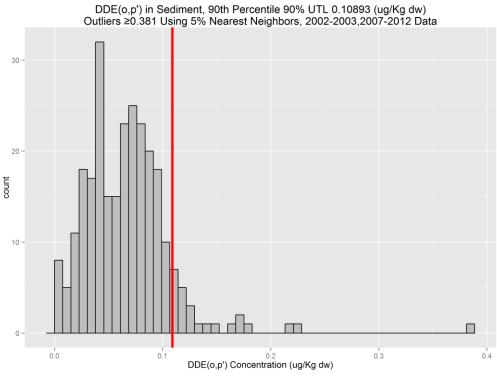


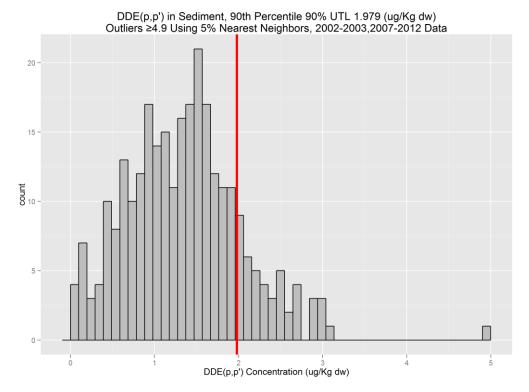


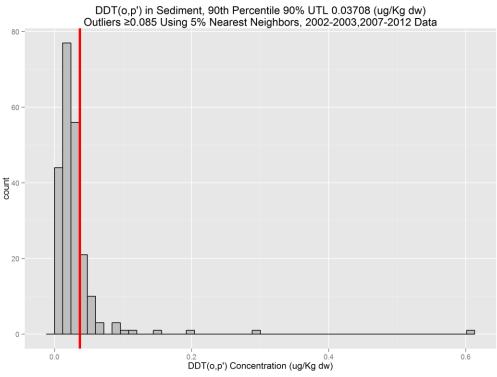


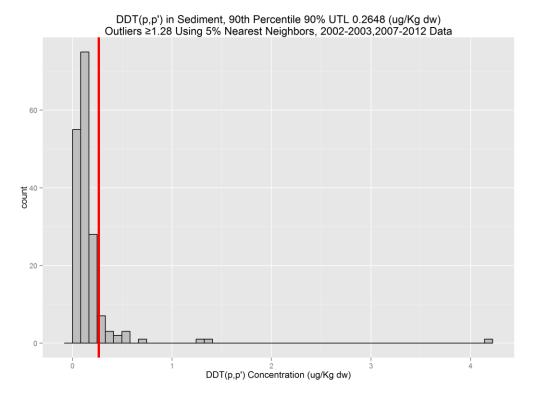


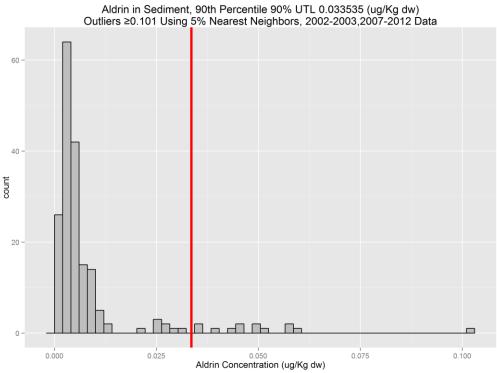


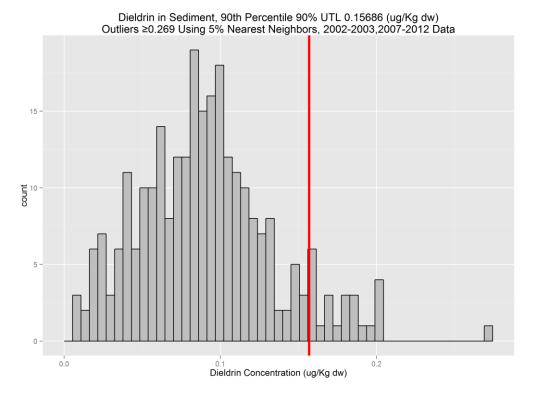


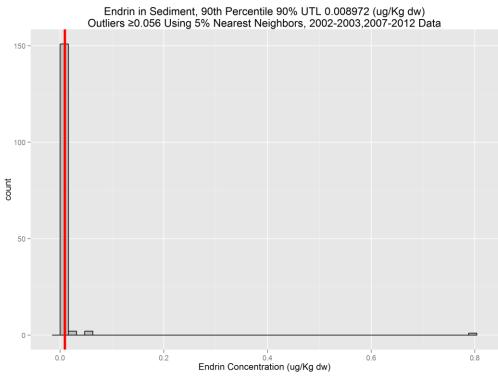


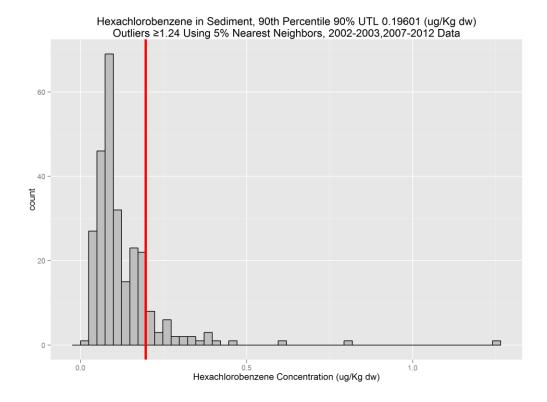




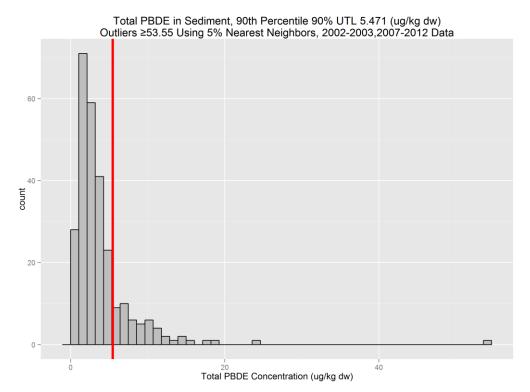








PBDEs



Dioxin TEQs

