

Assessment of contaminant body burdens and histopathology of fish and shellfish species frequently used for subsistence food by Alaskan Native communities

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Assessment of contaminant body burdens and histopathology of fish and shellfish species frequently used for subsistence food by Alaskan Native communities.

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NORTH PACIFIC RESEARCH BOARD PROJECT FINAL REPORT

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List of Acronyms

As	arsenic
ASTM	American Society of Testing and Materials
Cd	cadmium
CIRCAC	Cook Inlet Regional Citizens Advisory Council
Cr	chromium
Cu	copper
DDT	Dichlorodiphenyltrichloroethane
EMAP	Environmental Monitoring and Assessment Program
EPA	Environmental Protection Agency
EVOS	Exxon Valdez Oil Spill
GC/ECD	Gas Chromatography/Electron Capture Detector
GC/MS	Gas Chromatography/Mass Spectroscopy
g	gram
GOA	Gulf of Alaska
HCH	Hexachord-Cyclohexane
Hg	mercury
ICP	Inductively Coupled Plasma
KBNERR	Kachemak Bay National Estuarine Research Reserve
MDL	Method Detection Limit
mg	milligram
Mn	manganese
MS	Matrix Spike
MSD	Matrix Spike Duplicate
Ni	nickel
NIST	National Institute of Standards and Technology
NOAA	National Oceanic and Atmospheric Administration
NCCOS	National Centers for Coastal Ocean Science
NS&T	National Status and Trends
PAH	Polycyclic Aromatic Hydrocarbon
Pb	Lead
PCB	Polychlorinated Biphenyl
POP	Persistent Organic Pollutant
PWSRCAC	Prince William Sounds Regional Citizens Advisory Council
Se	selenium
SRM	Standard Reference Material
TBT	Tributyltin
µg	microgram
Zn	zinc

Abstract

Subsistence food items can be a health concern in rural Alaska because community members often rely on fish and wildlife resources not routinely monitored in these areas for contaminants and diseases. Subsistence activities are a large part of the traditional culture, as well as a means of providing protein in the diets for Tribal members. In response to the growing concerns among Native communities, contaminant concentrations in shellfish and indicators of disease in chum and sockeye salmon (*Oncorhynchus keta* and *Oncorhynchus nerka*) and the shellfish cockles and softshell clams (*Clinocardium nuttallii* and *Mya arenaria*) were assessed. In the Spring of 2010, the fish and shellfish were collected from traditional subsistence harvest areas in the vicinity of Nanwalek, Port Graham, and Seldovia, AK, and were analyzed for trace metals and residues of contaminants routinely monitored by the NOAA National Status & Trends Program (NS&T). Additionally, the fish and shellfish were analyzed for the presence, prevalence and severity of tissue abnormalities (histopathology), disease, and parasite infection. The fish and shellfish sampled showed low tissue contamination, and effects of the parasites and diseases were absent or minimal. Taken together, the results showed that the fish and shellfish were healthy and pose no safety concern for consumption. This study provides reliable information for local resource managers and Alaska Native people regarding subsistence fish and shellfish use and management needs.

1. INTRODUCTION

The Chugach and Cook Inlet Native communities of Nanwalek, Port Graham and Seldovia are located at the southwestern tip of the Kenai Peninsula near the entrance to Kachemak Bay, a bay off the lower Cook Inlet. In these villages, subsistence activities are a large part of the traditional culture, as well as a means of providing protein for Tribal members. As defined by the Division of Subsistence, Alaska Department of Fish and Game (ADF&G), and subsistence living is the customary and traditional use of wild food gathered through fishing and hunting (<http://www.adfg.alaska.gov>). In rural Alaska and particularly in Native villages, subsistence activities are a large part of the traditional culture and in many communities hunting and fishing provide the main source of protein (Wolfe, 1996). Based on the ADF&G most recent comprehensive assessment conducted in the 1990s, the Division of Subsistence estimated an annual per capita average consumption of about 375 pounds of food harvested in rural Alaska statewide. In contrast, for the average American in the contiguous U.S., the estimate is less than 255 pounds of meat, fish and poultry, primarily derived from commercial

grocery outlets (<http://www.adfg.alaska.gov>). Further estimates by Wolfe, (1996) indicated that in some rural Alaskan villages the average per capita harvest reaches well over 600 pounds per person. Drawing from population density, Wolfe estimated that close to two pounds of wild food is consumed per person per day, a figure that highlights the significance of subsistence food consumption in Alaska.

Although wild foods are traditionally considered more nutritious than commercially available food, they may not be any healthier because of potential exposure to environmental stressors. Pollution and other environmental factors, such as climate change, constitute stressors that are impacting the health of marine and coastal resources in Alaska. Remote Alaskan regions, which were once considered pristine, are now known to be subjected to exposure to contaminants (AMAP, 2005; Wolfe, 1996). Studies have found that a wide variety of pollutants, including synthetic organic chemicals and, polycyclic aromatic hydrocarbons (PAHs) from natural sources, industrial, and accidental spills, are finding their ways into food chains within ecosystems in Alaska (Short *et al.*, 2002; AMAP, 2011). While studying mercury accumulation in fish, MacFarlane, (2004) noted that possible sources of mercury in south-central Alaska include gold mining activities and volcanic eruptions. There are five active volcanoes on the western side of Cook Inlet. Intermittent eruptions from these volcanoes periodically contribute volcanic ash to the region. Thus, in addition to the weathering of mineral-rich soil, likely sources of natural inorganic contaminant inputs into area ecosystems could be linked to volcanic eruptions. With better understanding in recent years of global geochemical circulation in the Arctic region, there has been increasing concern about the grasshopper effect, by which metals and contaminants known as “persistent organic pollutants” (POPs) from warmer regions are being transported and deposited into Alaska’s ecosystems (UNEP, 2005). Thus, along with mercury, persistent organic pollutants, such as toxic chlorinated pesticides (e.g., DDTs) and industrial contaminants (e.g. polychlorinated biphenyls or PCBs) emitted as results of man-made activities in the Americas, Europe and Asia, could be transported and deposited in the Kachemak Bay ecosystem and stress vital coastal resources.

Most fish and shellfish species harbor a natural array of parasites that can affect their health, exposure to contaminants is known to impact their immune system and facilitate parasitism and occurrence of diseases (Weis *et al.*, 1995; Johnson *et al.*, 1992; and MacKenzie *et al.*, 1995). The Alaska Department of Fish and Game assessed the infection pattern of the unicellular parasite *Ichthyophonus hoferi*, which was said to be harmless to humans, but was blamed for devastating infections in salmon (Kocan *et al.*, 2004; Dehn, 2008). Recently, a number of biochemical alterations and emergence of diseases in marine and coastal environments have

been linked to climate change that is shifting the disease landscape globally (Harvell et al., 2002). Additionally, the presence of biological toxins such as paralytic shellfish poisoning (PSP) in shellfish related to harmful algal bloom events can pose a serious health risk. Recent outbreaks of PSP in Alaska have been linked to the consumption of shellfish (RaLonde, 1996).

Resources used for subsistence foods in Alaska could be potentially exposed to dangerous compounds and toxins but, there is no systematic wild food testing in Alaska (Wolfe, 1996). Native communities that rely on subsistence foods have minimal information about the safety of their harvest (Wolfe, 1996). In response to the growing concerns within Native communities, this project sampled commonly used subsistence foods (chum and sockeye salmon (*Oncorhynchus keta* and *Oncorhynchus nerka*) and cockles and soft-shell clams (*Clinocardium nuttallii* and *Mya arenaria*), to assess their overall health condition and level of contamination. The fish and shellfish were collected from traditional subsistence harvest areas in the vicinity of Nanwalek, Port Graham and Seldovia, AK, and were analyzed for metal contamination and organic contaminants routinely monitored by the NOAA National Status & Trends Program (NS&T). Additionally, the fish and shellfish were examined for diseases and parasitic infections.

To put results from this study into perspective, concentration levels in salmon and clams were compared to the Alaska Department of Environment Conservation, Fish Monitoring Program (DEC-FMP) data and, when possible, to the U.S. Food and Drug Administration (FDA) action levels for seafood safety and consumption levels shown by the Environmental Protection Agency to have no negative health effects. This study provides useful chemistry and disease information on salmon, cockles and clams for concerned native community members and coastal resource managers in Alaska. As the Nation's longest running coastal contaminant monitoring and assessment program, the NS&T program maintains a publically available national database of map based chemical, physical and biological information. The data from this study were incorporated into the NS&T data portal and are available to the public (<http://egisws02.nos.noaa.gov/nsandt/index.html#>).

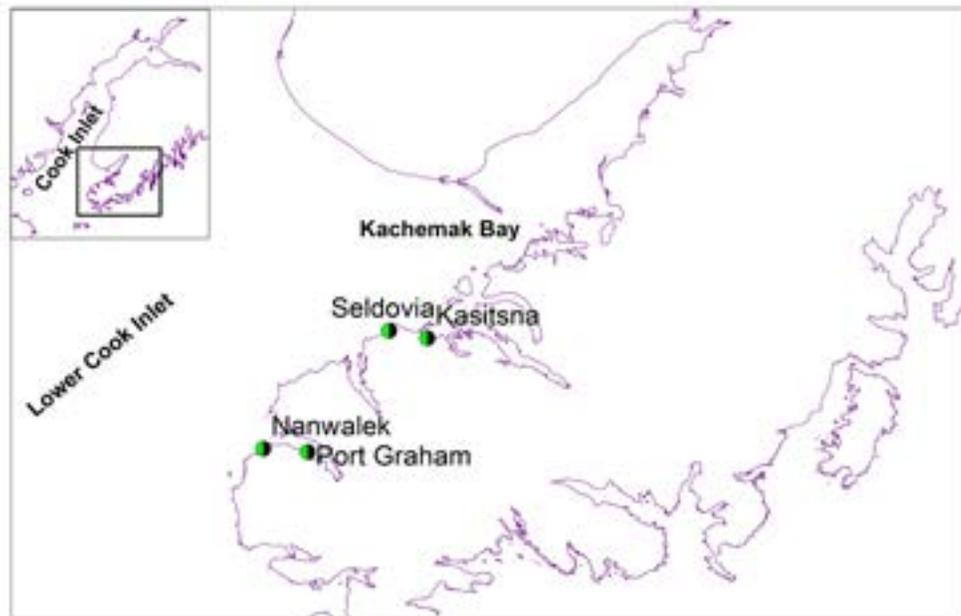


Figure 1. Map showing the geographic location of the villages of Nanwalek, Port Graham and Seldovia in Lower Cook Inlet, and the NOAA Kasitsna Bay lab. The inset depicts the general location of Cook Inlet and Kachemak Bay.

2. OBJECTIVES

The overall goal of the study was to assess the health risks associated with consumption of subsistence food items collected in the traditional harvest ground in Native communities of south-central Alaska. To achieve this goal the following objectives were accomplished:

1. Characterize the potential contamination of sessile (shellfish) vs. mobile (salmon) components of the subsistence fishery, using contaminant concentration in the organisms as well as incidence of disease
2. Evaluation of the health condition of the fish and shellfish
3. Assess the potential consumption risk of these fish and shellfish

3. METHODS

3.1 Sampling

Samples were collected by community members in each village. Sample collectors were trained and assisted by NOAA scientists to conduct quality assured sampling and sample handling. Sampling followed quality

controlled and quality assured procedures of the NS&T and national Marine Fisheries Service (NMFS) for sample collection (Apeti et al., 2012; NMFS, 1995). The target shellfish samples were collected in triplicate from each of three harvest areas in May, 2010. Fish were collected in July, 2010, during the salmon run.

3.1.1 Shellfish sampling

Three locations were identified in each of the traditional harvest areas of Nanwalek, Port Graham and Seldovia. At each location, edible sized and co-located cockles and clams were sought for hand collection at low tide. Clams were collected at all locations, while cockles could only be collected at Port Graham and Seldovia. Shellfish were identified in the field based on local traditional knowledge using common (colloquial) names and a scientific name (Foster, 1991). After collection, specimens from each location were kept separate, brushed clean with ambient water and sorted. Organisms of 6 to 7 cm in length were combined into a single sample (composite). For each species, three composite samples were selected for each location, one for each method (trace elements, organic contaminant and disease (histopathology)). For trace metal and organic contaminant analysis, sample composites consisted of 30 organisms, while only 12 organisms were collected for the histopathology analysis (Apeti et. al, 2012). Due to depleted stock of cockles in Port Graham, samples were only sufficient for organic contaminants and histopathology measurement. The selected samples were put into labeled double Ziploc bags and preserved on ice. The shellfish samples were air-shipped to government contracted analytical laboratories: TDI Brooks in Texas for organic analysis, Texas A&M University for trace element analysis, and Rutgers University, Haskin Shellfish Research Laboratory for histopathology analysis. Histological examination included summarizing the incidence of parasites and lesions in the gills and other organs and tissues.

3.1.2 Fish sampling and necropsy

The two species of salmon were collected from the traditional harvest grounds in each village using gillnets. Sample handling and preparation followed established protocols by Northwest Fisheries Science Center (NWFSC) a division of the NOAA National Marine Fisheries Service (NMFS, 1995; Lauenstein and Cantillo, 1993). For each species of salmon, five males and five females were collected from each village. Immediately after collection, the fish samples were delivered to the NOAA/University of Alaska Fairbanks, Kasitsna Bay laboratory for processing and necropsy. Muscle and liver samples were collected for chemical analyses. Liver, kidney, and gill samples were collected for histological assessment.

Fish necropsy occurred as soon after death as possible. The fish were sorted by species and sex, and measured to determine their weight and length. To prevent cross contamination of samples during necropsy, multiple separate sets of dissection tools were used for the removal of fish muscle, liver, kidney and gill tissues for the various analyses, depending on the type of analysis to be performed on the tissue. One set of “external only” dissection tools (that were used only for external procedures) was used to make the initial cuts through the epidermis for access to the fish muscle to be collected for chemical analyses. Using a pair of hemostats, a strip of skin was removed behind the head parallel to and about 5 to 10 mm dorsal to the lateral line, exposing the underlying muscle, and using “external only” dissection tools. Using separate sets of distilled water-rinsed Teflon knife and polyamide forceps (for the metals analyses) and isopropanol-rinsed scalpel and stainless steel scalpel and forceps (for the organic analyses), two separate blocks of muscle tissue were removed from the exposed area, from clearly inside of the margin of the original cut made through the skin, to prevent any contamination of the muscle samples by contact with the external skin or mucus, for the separate analysis of trace metals and organic contaminants. All samples of tissues for organic chemical analysis were collected using dissection tools that had been rinsed with isopropanol between fish; separate samples of tissues for metals analysis were rinsed with distilled water. Samples for chemical analyses were composited by site, species and sex; five fish of separate species and sex at each site consisted of a composite sample, with separate composite for the metals and organics analyses.

The “external only” set of dissection scissors, spawning knife and scalpels were then again used to open the abdominal cavity to access the internal organs, without contacting the liver. A separate set of dissection tools were used to collect the histological samples of gill, head and trunk kidney, and liver. When removing liver tissue, care was taken to not puncture the gall bladder, so that bile was not spilled on the liver sample. Liver tissue for the separate chemical samples (organics, metals) was collected using a separate set of dissection tools, consisting of separate distilled water-rinsed Teflon knife and polyamide forceps for the metals sample, and a separate set of stainless steel scalpel, scissors and forceps for the organics sample; these dissection tools were only used to collect samples for chemical analyses from the liver. A total of 90 individual kidney, gill and liver tissues were collected for the histopathology characterization. Liver, kidney and gill tissues were placed into tissue cassettes and immediately preserved in Davidson’s fixative (Fournie et al., 2000) until shipped to and analyzed at the Northwest Fisheries Science Center. A total of 24 composite samples of liver and muscle tissues were collected for fish contaminant body burden assessment. The tissue samples destined

for contaminants analysis were placed into labeled I-Chem jars and kept frozen until shipped to TDI Brooks and Texas A&M University. The project sought to collect and analyze stomach contents from the salmon to determine fish prey and to quantify the concentration of organic contaminants in prey organisms. However, all attempts to collect stomach contents were unsuccessful as they had stopped feeding in brackish waters during the fish run up the rivers prior to spawning (Pecquerie et. al, 2011). Unused tissues were discarded.

3.2 Contaminant analysis

Analyses of organic contaminants and metals in clam and cockles soft tissue and in fish liver and muscle followed the standard NS&T analytical protocols described in Kimbrough and Lauenstein (2006a and 2006b). The shellfish were shucked and the whole soft tissue of the 30 organisms composite were homogenized and freeze-dried. Fish liver and muscle composite samples were also subjected to the same blending and freeze drying processes before analysis.

During analysis, quality control samples were processed with every sample batch.

3.2.1 Major and trace metals analysis

Major and trace metals measured in the fish and shellfish tissue are listed in Table 1. All tissue types were subjected to the same digestion and analytical methods (Kimbrough and Lauenstein, 2006).

3.2.2 Organic contaminants analysis

Organic contaminants analyzed in fish and shellfish as part of this study are listed in Table 1. Polychlorinated Biphenyl ether (PCBs) and Polycyclic aromatic hydrocarbons (PAHs) were analyzed in shellfish tissue only, because fish effectively break down PAHs to compounds not detectable by routine analytical procedures. All analyses were conducted with methods described by Kimbrough and Lauenstein, 2006.

3.2.3 Method detection limits

For each metals and organic compound measured, an analytical method's limits of detection (MDL) were determined. Determination of MDL followed procedures described by the Environmental Protection Agency in 40 CFR Part 136, (EPA, 2005) and it was defined as the Student's t for 99% confidence level times the standard deviation of seven or more replicate measurement of the same low level spiked samples.

Table 1. List of organic pollutants and metals analyzed by the NS&T program.

Metals: Silver (Ag), Arsenic (As), Cadmium (Cd), Chromium (Cr), Copper (Cu), Iron (Fe), Lead (Pb), Mercury (Hg), Manganese (Mn), Nickel (Ni), Selenium (Se), Tin (Sn), Zinc (Zn)
Butyltins: monobutyltin, dibutyltin, tributyltin, tetrabutyltin
Chlordanes: <i>alpha</i> -chlordane, <i>gamma</i> -chlordane, oxychlordane, <i>cis</i> -nonachlor, <i>trans</i> -nonachlor, heptachlor, Heptachlor-Epoxyde
Chlorpyrifos
DDTs: <i>ortho</i> and <i>para</i> forms of parent 2,4' DDT and 4,4' DDT and metabolites 2,4' DDE; 4,4' DDE; 2,4' DDD; 4,4' DDD
Dieldrins: aldrin, dieldrin and endrin
Chlorobenzenes: 1,2,3,4-Tetrachlorobenzene, 1,2,4,5-Tetrachlorobenzene, Hexachlorobenzene, Pentachlorobenzene, Pentachloroanisole
Hexachlorocyclohexanes (HCHs): Alpha-Hexachlorocyclohexane, Beta-Hexachlorocyclohexane, Delta-Hexachlorocyclohexane, Gamma-Hexachlorocyclohexane
Endosulfans: Endosulfan I, Endosulfan II, Endosulfan sulfate
PAHs: Naphthalene, C1-Naphthalenes, C2-Naphthalenes, C3-Naphthalenes, C4-Naphthalenes, Benzothiophene, C1-Benzothiophenes, C2-Benzothiophenes, C3-Benzothiophenes, Biphenyl, Acenaphthylene, Acenaphthene, Dibenzofuran, Fluorene, C1-Fluorenes, C2-Fluorenes, C3-Fluorenes, Anthracene, Phenanthrene, C1-Phenanthrenes/Anthracenes, C2-Phenanthrenes/Anthracenes, C3-Phenanthrenes/Anthracenes, C4-Phenanthrenes/Anthracenes, Dibenzothiophene, C1-Dibenzothiophenes, C2-Dibenzothiophenes, C3-Dibenzothiophenes, Fluoranthene, Pyrene, C1-Fluoranthenes/Pyrenes, C2-Fluoranthenes/Pyrenes, C3-Fluoranthenes/Pyrenes, Naphthobenzothiophene, C1-Naphthobenzothiophenes, C2-Naphthobenzothiophenes, C3-Naphthobenzothiophenes, Benz(a)anthracene, Chrysene/Triphenylene, C1-Chrysenes, C2-Chrysenes, C3-Chrysenes, C4-Chrysenes, Benzo(b)fluoranthene, Benzo(k,j)fluoranthene, Benzo(e)pyrene, Benzo(a)pyrene, Perylene, Indeno(1,2,3-c,d)pyrene, Dibenzo(a,h)anthracene, Benzo(g,h,i)perylene, , Total PAHs,
Individual Alkyl Isomers, , 2-Methylnaphthalene, 1-Methylnaphthalene, 2,6-Dimethylnaphthalene, 1,6,7-Trimethylnaphthalene, 1-Methylphenanthrene, C29-Hopane, 18a-Oleanane, C30-Hopane,
PCBs: PCB8/5, PCB18, PCB28, PCB29, PCB31, PCB44, PCB45, PCB49, PCB52, PCB56/60, PCB66, PCB70, PCB74/61, PCB87/115, PCB95, PCB99, PCB101/90, PCB105, PCB110/77, PCB118, PCB128, PCB138/160, PCB146, PCB149/123, PCB151, PCB153/132, PCB156/171/202, PCB158, PCB170/190, PCB174, PCB180, PCB183, PCB187, PCB194, PCB195/208, PCB199, PCB201/157/173, PCB206, PCB209,
Mirex

Total is the arithmetic sum of the congeners or homologue groups of compounds

3.3 Histopathology analysis

The histopathology analyses are a set of measurements that determine the presence of parasites and degree of infection as well as the occurrence of disease in fish and shellfish tissues.

3.3.1 Shellfish histopathology

The histological analyses of shellfish were performed at Rutgers University's Haskin Shellfish Laboratory. A detailed account of the protocol is described in the NOAA's NOS/NCCOS technical memorandum number 27 (Kim *et al.* 2006). From each location, subsets of 5 individual organisms of legal harvestable size (38 mm) were randomly selected and prepared for the analysis.

The adductor muscles of organisms were cut with a sharp knife so that the valves remained open. The entire animal was placed in Davidson's fixative for 1 week and then transferred to 70% alcohol for storage. A sharp knife or scalpel was carefully run between the shell and the mantle to separate the meat from the shell. This procedure was repeated for the other shell to completely detach both sides of the mantle from the shell. Major tissue types examined included gill, mantle, gonad and gonoducts, digestive gland tubules, stomach/digestive gland, and connective tissue. The animals' gonadal stage evaluations include determination of sex and stage of gonadal development.

Quantitative Measures: Conditions scored quantitatively (Table 2) were evaluated by keeping a running count of occurrences of the condition as the slide is scanned to avoid re-examining each incident multiple times. Quantitative scores were used for parasites, pathologies, and selected morphological conditions that could be tallied individually (Kim *et al.*, 2006). Parasites counted quantitatively included prokaryotic inclusion bodies (rickettsia, chlamydia, etc.), various ciliates, gregarines, other protozoans, nematodes, encysted cestodes and metacercariae of trematodes, copepods and other unidentified organisms. Ciliates were quantified by tissue type (gill and digestive tract), as were the gregarines (body, gill, and mantle). Nematodes were also subjected to quantitative count based on their observed cross-sections. A number of tissue pathological conditions were also evaluated quantitatively, including the number of ceroid bodies, cases of hemocytic infiltration that were scored separately as focal and diffuse incidences of tissue inflammation, and tumors.

Semi-quantitative Measures: Some conditions are assigned to a semi-quantitative scale relative to the intensity or the extent of the affected area (Tables 2). Definitions of scale values can be found in Kim *et al.* (2006). A semiquantitative 0-to-4-point scale is used for invasive trematode sporocysts (*Felodistomidae* and *Bucephalidae*). For each specimen examined, the presence of neoplasia and unusual digestive tubules is recorded semi-quantitatively using the 0-to- 4-point scale. Abnormal gonadal development characterized by unusual development is given a semiquantitative 0-to-4-point score relative to the spatial coverage of

the condition (Kim *et al.*, 2006). For digestive gland atrophy, a condition known to be caused by a variety of stressors, most likely related to poor nutrition (Winstead, 1995), the average degree of thinning of the digestive tubule walls was assigned a numerical rating on a 0-to-4-point scale (Kim *et al.*, 2006). Semi-quantitative procedures for the assessment of the magnitude of parasitic infection and tissue diseases are exemplified in this document using scales for trematode sporocyst infection (Table 3) and histological condition of digestive gland atrophy (Table 4).

Table 2. List of parameters measured for the histopathological assessment of bivalves. Top: list of parasitic species. Bottom: list of diseases and tissue conditions. Parameters measured semi-quantitatively are in bold; all other parameters were measured quantitatively.

Parasite category	Parasites
Cestodes	Body cestode, Gill cestode, Mantle cestode, Cestode metacercaria
Copepods	Body copepod, Gill copepod, Gut copepod
Ciliates	Digestive tract ciliate, Large gill ciliate, Small gill ciliate, Gut ciliate
Protozoan	Digestive tubule protozoan, Gut protozoan
Nematode	Nematodes
Trematodes	<i>Trematode sporocyst gut</i> , <i>Bucephalid</i> trematode spore, Trematode sporocyst gill, Trematode metacercaria, Protoeces
Gregarines Nematopsis	Nematopsis body, Nematopsis gill, Nematopsis mantle
Rickettsia	Digestive tubule rickettsia, Gut rickettsia, <i>Chlamydia</i> , <i>Prokaryotic bodies</i>
Coccidian	<i>Pseudoklossia</i>
Hydra	Gill hydra
Nemertines	Gill nemertine
Pea crab	Pinnotherid crab
Unidentified organism	Unidentified gonoduct organism, Unidentified organism

Disease category	Diseases
Tissue Inflammation	Focal inflammation, Diffuse inflammation
Necrosis	Necrosis diffuse, Necrosis focal, Ceroid bodies
Digestive tubule conditions	Digestive tubule atrophy , Unusual digestive tubule
Edema	Edema
Gonads	Gonad abnormalities
Neoplasm	Neoplasm
Tumor	Tumor
Xenoma	Xenoma

Table 3. Semi-quantitative scale for trematode sporocyst infection.

Score	Description
0	Uninfected
1	Present in the gonads only (some gametic tissue still present)
2	Completely filling the gonads (no gametic tissue present); may be present in digestive gland or gills in very limited amount
3	Completely filling the gonads; extensive invasion of the digestive gland and/or the gills
4	Completely filling the gonad; substantially filling the digestive gland or gill; individuals appear to be a sac of sporocyst

Table 4. Semi-quantitative scale for digestive gland atrophy.

Score	Description
0	Normal wall thickness in most tubules (0% atrophy), lumen nearly occluded, few tubules even slightly atrophied
1	Average wall thickness less than normal, but greater than one-half normal thickness, most tubules showing some atrophy, some tubules still normal
2	Wall thickness averaging about one-half as thick as normal
3	Wall thickness less than one-half of normal, most tubules walls significantly atrophied, some walls extremely thin (fully atrophied)
4	Wall extremely thin (100% atrophied), nearly all tubules affected

3.3.2 Fish histopathology

The histopathological analysis of salmon was performed at the NOAA Northwest Fisheries Science Center, in Seattle, WA. Histopathologic diagnosis was performed on fish liver, head and trunk kidney, and gill tissues. Sections of liver, head and trunk kidney (1cm in thickness) and two gill arches collected from individual salmon were preserved in Davidson's fixative (Fournie et al., 2000) at a volume:tissue ratio of at least 10:1 for at least two full days, then transferred to 70% ethanol for storage and transfer to the histopathology laboratory in Seattle. In the laboratory, tissues were processed by an automated tissue processing center, embedded in paraffin, sectioned at a 5µm thickness, stained with hematoxylin and eosin and examined by light microscopy with the presence any lesions or detected parasites documented and scored as described for adult salmon in Fairgrieve et al., (2005). Lesions and parasites in tissue sections were identified and classified according to the

criteria specified in Meyers and Hendricks, (1985), Cotran et al. (1999) Chitwood and Lichtenfeld (1972) and Bruno et al. (2006).

3.4 Data analysis

3.4.1 Contaminant compounds data analysis

Laboratory concentration results were subject to regular NS&T performance-based quality assessment and quality control for data accuracy and precision. Concentration values for individual compound that were smaller than the minimum detectable level (MDL) were qualified as undetected and were assigned a value of zero. For organics, the “totals” were derived as the sum of all the individual congeners or homologues of the same group of compounds as listed in Table 1. Contaminant body burdens of toxic metals and organic compounds in salmon and clams were compared to FDA action levels, EPA chronic consumption limits, and to monitoring data from the Alaska DEC, Fish Monitoring Program. Alaska DEC and FDA both report concentration levels on wet weight basis. Assuming average percent moisture of 76% for the salmon and 86% for clam (values were derived from this study), factors of 4 and 7 respectively for salmon and clam were used to convert wet weight concentrations into dry weight concentrations.

3.4.2 Shellfish histopathology data analysis

The severity of parasitic infections and that of pathological conditions were assessed by calculating prevalence and intensity of the condition.

Prevalence describes the proportion of individuals in the population that are infected by a specific parasite or pathology and is calculated as:

$$\text{Prevalence} = \frac{\text{number of hosts with parasite or pathology}}{\text{number of hosts analyzed}}$$

Infection intensity is calculated as the average number of occurrences of the parasite or pathology in infected hosts. This is a measure of the intensity of infection in infected individuals.

$$\text{intensity} = \frac{\text{total number of occurrences of parasite or pathology}}{\text{number of hosts with parasites or pathology}}$$

For conditions measured semi-quantitatively, the scale rating replaced the number of occurrences in this computation. The protocol for the biological component of the NS&T Program stipulates analysis of five individuals per site.

For this study, parasites of the same taxa were pooled by class as indicated in Table 2 and the resulting prevalence and intensity values were determined as the sum of the prevalence and intensity values of the individual parasites. For instance, the class of Cestoda, or “tapeworms,” includes body cestodes, gill cestodes, mantle cestodes and cestode metacercariae. The class of Ciliates includes the digestive track ciliates, large gill ciliates, small gill ciliates and gut ciliates. The class of Gregarina includes the gregarines nematopsis in the body, gill and mantle. The class of Trematodes or flatworms includes *Bucephalid* trematodes spore, trematode sporocyst gill, trematode metacercariae and *Protoeces*.

3.4.3 Fish histopathology data analysis

Conditions were scored quantitatively by keeping a running count of occurrences of the condition as the slide is visually scanned to avoid re-examining each incident multiple times. Quantitative scores were used for parasites based on their observed cross-sections. Tissue pathological conditions (lesions) were also evaluated quantitatively. Parasite and pathologic conditions were tallied and scored in fish gill, liver and kidney for individual. For each site and each salmon species the severity of parasitic infections and that of pathological conditions were assessed by calculating prevalence values separately for the male and female fish. Prevalence describes the proportion of individuals in the population that are infected by a specific parasite or affected by a specific pathological lesion and was calculated as above.

3.4.4 Statistical analysis

SAS and JMP statistical packages were used for data processing and analysis. For the histopathology parameters, the severity of parasitic infections and occurrence of disease, or histologic conditions, were assessed by deriving the prevalence and intensity of each or group of parameters measured. Both prevalence and intensity were derived for parameters measured in clams and cockles. Due to the nature of the parameters measured in the salmon species, only prevalence were calculated for these parameters. For both contaminant concentrations and prevalence/intensity values, the Fisher's Exact Test was used to assess data comparability and degree of difference between values. Significance of statistical tests were reported at a probability level of 0.05.

4. RESULTS

Results are presented for the contaminant analyses and the histological assessments. Contaminant results are presented for shellfish and fish together because all tissues were analyzed for the same things.

The overall analytical results describing levels of the metal and organic contaminants measured in fish and shellfish tissues are presented in Tables 5, 6 (a,b, c). Location-specific assessment and data variation among the fish and shellfish species as well as comparison between different tissue types are discussed below for those chemicals with potential human health significance. Where possible, concentration levels of this study were put into context by comparison to reported safety threshold values from FDA and EPA. An FDA action level represents the limit at or above which FDA may take legal action to remove products from the marketplace. EPA-recommended values typically range from 2 to 120 times lower than the corresponding FDA action levels, as the EPA's values are derived from a risk-based approach to initiate local fish consumption advisories and are much more protective (EPA, 2000). For comparative purposes, the values presented here use the EPA chronic reference dose (RfD) and assume an average person's weight of 80 kg (176 lb) and a meal of fish to be 0.227 kg (1/2 lb). The chronic reference dose assumes consumption of a meal of fish every day. For comparative purposes, all values were converted into concentration based on dry weights.

$$\text{No Effect Chronic concentration} = (\text{RfD} * \text{weight}) / \text{meal weight}$$

Other assumptions for specific groups can be used (e.g. children). The tables also list concentration values derived for different bivalve species from the NOAA NS&T Mussel Watch and the AK DEC Fish Monitoring

Table 5a. Contaminant concentration in shellfish. Where available, results from this study were weighed against mean concentration values reported by the NOAA Mussel Watch for blue mussels and the Alaska DEC Fish Monitoring Program for razor clams. Results were also compared to the FDA action levels for shellfish consumption. (Metals $\mu\text{g}\cdot\text{g}^{-1}$, Organics $\text{ng}\cdot\text{g}^{-1}$ dry weight).

	Nanwalek		Port Graham		Seldovia		NOAA (2007)	FDA action level
Compound	Clam	Clam	Cockle	Clam	Cockle	Blue Mussel	shellfish	
Butyl Tin	0	0		0	0	1.18		
Chlordanes	0.27	1.48	1.09	0.96	0.45	1.46		
Chlorobenzenes	1.82	1.5	1.5	1.2	0.94	0.66		
Chlorpyrifos	0.44	0	0.41	0.23	0.37	0		
DDTs	0.31	0.17	0.17	0.73	0.41	0.47		
Dieldrins	0.13	0	0	0.38	0	0.44		
Endosulfans	0	0.26	0.22	0.04	0	0		
HCHs	3.85	5.7	4.32	2.93	3.53	0.71		
Mirex	0.22	0	0	0.5	0	0		
PAHs	143.8	302.0	235.1	262.9	175.4	167.9		
PCBs	5.41	6.45	0.92	10.1	6.94	2.47		
Compound	Clam	Clam		Clam	Cockle	Blue Mussel	shellfish	
Arsenic	8.14	7.14		12	6.57	9.22	602	
Cadmium	0.912	1.27		0.87	0.65	2.62	28	
Chromium	3.51	3.28		4.65	2.02	1.26	91	
Copper	18.6	23.1		33.9	38.4	7.96		
Iron	316	1170		2330	1100	670		
Lead	0.27	0.4		1.47	1.49	0.59	11.9	
Manganese	12.3	26.7		40.4	18.4	15.37		
Mercury	0.031	0.129		0.210	0.073	0.082		
Nickel	4.79	3.27		9.3	12.1	1.83	560	
Selenium	1.89	2.6		3.28	3.01	2.89		
Silver	0.31	0.27		0.50	0.03	0.06		
Tin	0.437	0.13		0.12	0.18	0.0		

Table 5b. Contaminant concentrations in shellfish reported by the Alaska DEC Fish Monitoring Program. Results were also compared to the FDA action levels for shellfish consumption. (Metals $\mu\text{g}\cdot\text{g}^{-1}$ dry weight).

	Butter clam	Littleneck clam	Maya sp.	Razor clam	Redneck clam	Cockle	FDA action level
Arsenic	21.18	22.75	5.60	5.18	3.50	4.90	602
Cadmium	0.74	2.18		1.67	1.09	0.00	32
Chromium	3.64	1.20	3.22	3.71	4.24	1.23	104
Lead	0.35	0.31	0.63	0.70	0.49	0.39	14
Nickel	0.35	0.31	0.63	0.70	0.49	0.39	640

Table 6a: Contaminant concentrations in liver and muscle tissue of salmon collected from Nanwalek. To put concentration values from this study into context, concentration levels reported by the Alaska DEC Fish Monitoring program are presented. Results were also compared to the FDA action levels and calculated EPA chronic consumption thresholds for fish consumption. (Metals $\mu\text{g}\cdot\text{g}^{-1}$, Organics $\text{ng}\cdot\text{g}^{-1}$ dry weight). m = male, f = female.

	Chum f		Chum m		Sockeye f		Sockeye m		AK FMP Sockeye			FDA	EPA
	liver	muscle	liver	muscle	liver	muscle	liver	muscle	Kodiak	Matanuska	fish	fish	fish
Chlordanes	0.77	1.6	1.97	1.04	3.73	2.64	1.04	2.38	11.32	6.88	1,200	704.8	
Chlorobenzene	2.27	2.78	2.72	2.05	2.99	2.84	3.55	2.22	5.00	4.00		1127.8*	
Chlorpyrifos	0	0	0	0	2.59	0	0	0				422.9	
DDT	0.46	2.43	15.45	2.88	21.84	9.74	1.58	5.99	38.12	19.04	20,000	704.8	
Dieldrins	0.4	0.62	0.56	0.24	0.42	0.45	2.15	0.61	1.76	0.88	1,200	70.5	
Endosulfan	0.19	0.12	0.8	0.23	2.05	1.32	0.3	0.17				8458	
HCHs	0.63	5.52	3.85	10.08	5.29	12.56	3.32	0.31	6.72	0.52		422.9#	
Mirex	0	0	0	0	0	0.6	0	0			400	281.9	
PCBs	3.86	6.79	17.78	9.29	15.35	14.7	7.45	11.11	23.60	11.85	8,000	28	
									Kenai R.				
Arsenic	1.44	0.83	1.42	0.73	1.31	0.90	1.66	1.06	1.20	1.20			
Cadmium	2.21	0.01	3.09	0.01	3.59	0.01	3.83	0.02	0.00	0.00		1.41	
Chromium	0	0	0	0	0	0	0	0	0.00	0.00			
Copper	73.5	2.01	329	2.11	365	1.81	1500	1.83		0.82			
Iron	449	17.6	563	15.4	558	13.9	452	12.4					
Lead	0.05	0	0.06	0	0	0	0	0	0.00	0.00			
Manganese	7.2	0.321	5.19	0.37	9.03	0.374	5.61	0.301					
Mercury	0.138	0.104	0.131	0.093	0.194	0.134	0.228	0.098	0.120	0.100	4000	0.14	
Nickel	0.07	0.13	0.09	0.06	0.20	0.00	0.19	0.05	0.00	0.00			
Selenium	7.21	1.14	18.6	1.3	26.1	0.99	79.4	0.89	0.88	0.84		7.05	
Silver	2.63	0	4.95	0	6.19	0	7.4	0					
Tin	0	0	0	0	0	0	0	0					
Zinc	98.9	13.6	108.0	13.9	128	12.2	173	14.3					

*Hexachlorobenzene

#gamma HCH

Table 6b: Contaminant concentrations in liver and muscle tissue of salmon collected from Port Graham. To put concentration values from this study into context, concentration levels reported by the Alaska DEC Fish Monitoring program are presented. Results were also compared to the FDA action levels and calculated EPA chronic consumption thresholds for fish consumption. (Metals $\mu\text{g.g}^{-1}$, Organics ng.g^{-1} dry weight). m = male, f = female.

	Chum f		Chum m		Sockeye f		Sockeye m		AK FMP Sockeye		FDA	EPA
	liver	muscle	liver	muscle	liver	muscle	liver	muscle	Kodiak	Matanuska	fish	fish
Chlordanes	0.35	1.3	0.64	1.13	1.69	3.12	2.81	3.2	11.32	6.88	1,200	704.8
Chlorobenzene	1.78	1.47	2.01	1.6	1.71	1.83	2.98	2.18	5.00	4.00		1127.8*
Chlorpyrifos	0	0	0	0	0	0	0	0				422.9
DDT	0.8	1.46	1.3	2.22	1.79	10.19	3.49	9.02	38.12	19.04	20,000	704.8
Dieldrins	0.42	0.31	1.82	0.33	0.89	0.6	0.47	0.49	1.76	0.88	1,200	70.5
Endosulfan	0	0	0.44	0	0.66	0.38	0	0.26				8458
HCHs	0.38	0.38	1.17	0.24	0.79	0.35	0.74	2.92	6.72	0.52		422.9#
Mirex	0	0	0	0	0	0	0	0			400	281.9
PCBs	3.07	3.98	1.76	4.92	2.35	14.33	1.35	11.97	23.6	11.85	8,000	28
									Kenai R.			
Arsenic	1.28	0.77	1.43	0.91	1.21	0.90	1.23	0.80	1.20	1.20		
Cadmium	2.67	0.02	4.13	0.02	3.10	0.02	5.45	0.02	0.00	0.00		1.41
Chromium	0	0	0	0	0	0	0	0	0.00	0.00		
Copper	151	2.1	423	1.97	415	1.94	1440	1.68		0.82		
Iron	627	16	1130	14.9	649	13.2	518	12.9				
Lead	0	0	0.10	0	0	0	0	0	0.00	0.00		
Manganese	6.93	0.35	5.21	0.39	7.57	0.39	4.28	0.37				
Mercury	0.133	0.121	0.118	0.091	0.231	0.127	0.347	0.141	0.120	0.100	4000	0.14
Nickel	0.08	0.06	0.12	0.06	0.22	0.08	0.33	0.09	0.00	0.00		
Selenium	8.29	1.23	21.10	1.14	27.30	0.92	51.60	0.95	0.88	0.84		7.05
Silver	3.84	0	6.46	0	5.56	0	7.84	0				
Tin	0	0	0	0	0	0	0	0				
Zinc	101.0	12.6	106.0	15.1	125.0	13.2	152.0	18.6				

*Hexachlorobenzene

#gamma HCH

Table 6c: Contaminant concentrations in liver and muscle tissue of salmon collected from Seldovia. To put concentration values from this study into context, concentration levels reported by the Alaska DEC Fish Monitoring program are presented. Results were also compared to the FDA action levels and calculated EPA chronic consumption thresholds for fish consumption. (Metals $\mu\text{g}\cdot\text{g}^{-1}$, Organics $\text{ng}\cdot\text{g}^{-1}$ dry weight). m = male, f = female.

	Chum f		Chum m		Sockeye f		Sockeye m		AK FMP Sockeye		FDA	EPA
	liver	muscle	liver	muscle	liver	muscle	liver	muscle	Kodiak	Matanuska		
Chlordanes	1.01	1.16	4.46	1.25	2.59	2.33	1.91	1.8	11.32	6.88	1,200	704.8
Chlorobenzene	3.01	1.72	1.94	3.65	1.49	1.96	2.1	1.56	5.00	4.00		1127.8*
Chlorpyrifos	0.79	0	0	0	0	0	3.48	0				422.9
DDT	3.67	2.14	7.24	3.47	15.54	6.99	33.67	8.79	38.12	19.04	20,000	704.8
Dieldrins	0.47	0.26	1.72	0.31	0.45	0.4	0.63	0.4	1.76	0.88	1,200	70.5
Endosulfan	0.61	0.57	0	0.42	2.16	1.1	1.67	0.98				8458
HCHs	5.39	10.58	3.68	15.45	5.84	16.72	6.17	14.35	6.72	0.52		422.9#
Mirex	0	0.18	0	0.24	0	0.52	0	0.46			400	281.9
PCBs	8.15	9.33	9.65	7.76	12.99	18.16	16.05	13.61	23.6	11.85	8,000	28
									Kenai R.			
Arsenic	1.32	0.79	1.38	0.88	1.34	1.29	1.46	1.18	1.20	1.20		
Cadmium	2.40	0.01	4.57	0.03	2.88	0.02	3.87	0.04	0.00	0.00		1.41
Chromium	0	0	0	0	0	0	0	0	0.00	0.00		
Copper	86.3	2.23	404	2	593	2.19	1570	1.86		0.82		
Iron	685	17.7	1060	15	249	12.1	427	13.9				
Lead	0.13	0	0.08	0	0	0	0	0	0.00	0.00		
Manganese	7.21	0.52	6.38	0.67	7.62	0.36	4.33	0.304				
Mercury	0.124	0.108	0.122	0.092	0.221	0.125	0.262	0.114	0.120	0.100	4000	0.14
Nickel	0	0	0.09	0.07	0.15	0.07	0.18	0	0.00	0.00		
Selenium	7.26	1.18	20.70	1.13	32.50	0.93	69.30	0.93	0.88	0.84		7.05
Silver	3.03	0	6.33	0	5.56	0	8.13	0				
Tin	0	0.04	0	0	0	0	0	0				
Zinc	98.3	13.8	119.0	16.2	134.0	12.8	135.0	12.7				

*Hexachlorobenzene

#gamma HCH

Programs (FMP) from the region. Bivalves were collected from various locations in and around Kachemak Bay. Salmon filet values from the FMP are from fish captured in the Mantanus River above the north end of Cook Inlet, the Kenai River in the middle of Cook Inlet, or at Kodiak. Note that metals concentrations are expressed in ppm and organic contaminants are expressed in ppb. All values are presented as dry weight (dw).

4.1 Metal Contaminant concentrations in fish and shellfish

The results of contaminant body burdens including concentrations of the major and trace metals measured in in clams and cockles from the three villages are presented in Table 5a. Although results of all metals measured are presented, graphical representations and discussion of the most toxic and/or carcinogenic heavy metals (arsenics, cadmium, chromium, lead, mercury, nickel, and selenium) are examined in detail.

Arsenic

The concentrations of arsenic in the shellfish from the different villages are shown in Figure 2. Concentrations varied from 6.57 to 12.0 $\mu\text{g}\cdot\text{g}^{-1}$ dw with the minimum and maximum concentration values found in cockles and clams, respectively collected from the Seldovia village harvest grounds (Table 5a). The result showed little variation between arsenic concentration in clams from Nanwalek and Port Graham and cockles from Seldovia.

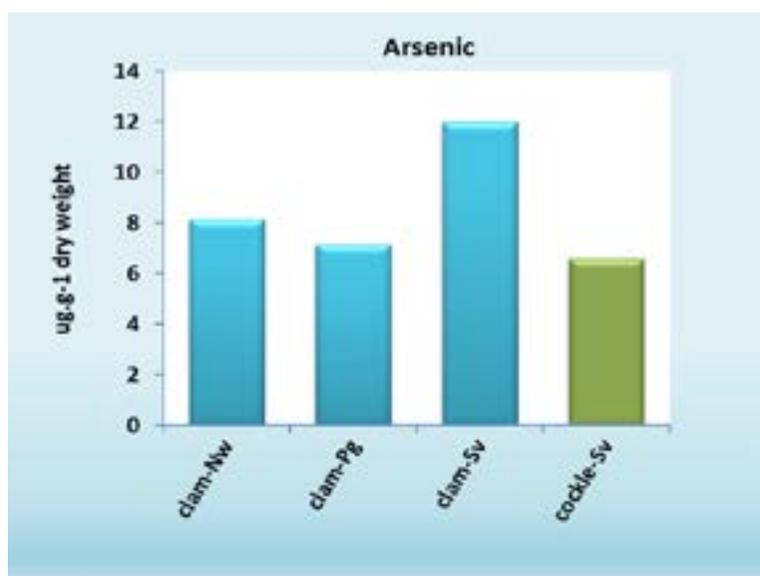


Figure 2. Concentration of arsenic in clam and cockle collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

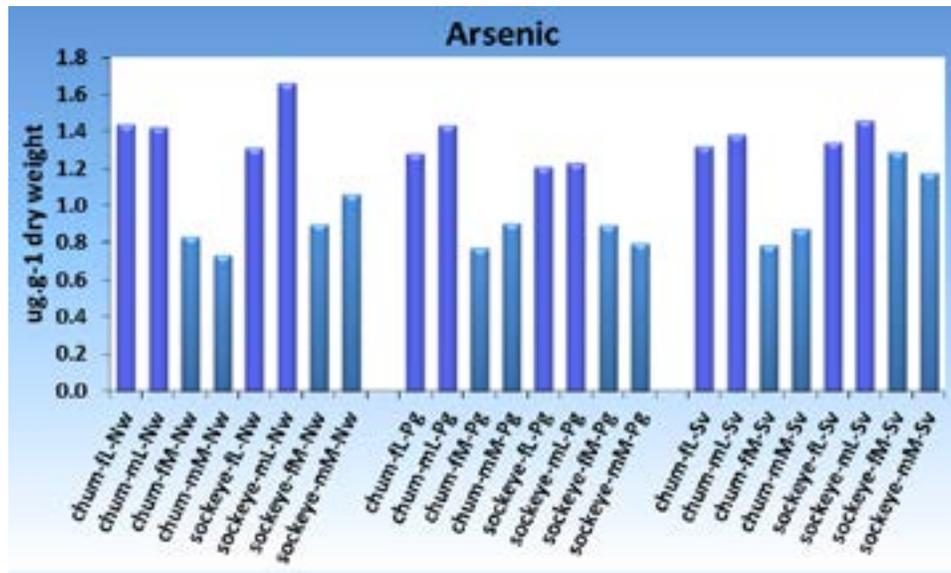


Figure 3. Concentration of arsenic in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

The average arsenic concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is 9.22 $\mu\text{g}\cdot\text{g}^{-1}$. The Alaska DEC-FMP (<http://dec.alaska.gov/eh/vet/fish.htm>) report tissue concentrations ranging from 3.5 to 22.75 $\mu\text{g}\cdot\text{g}^{-1}$ in bivalves from the Kachemak Bay area. These results indicate that arsenic concentration in the shellfish used as subsistence food is within the regional concentration range found in shellfish.

Arsenic body burdens in fish showed little variation between the two species or sex of salmon (Tables 6a, 6b and 6c). Body burdens varied from 0.73 to 1.66 $\mu\text{g}\cdot\text{g}^{-1}$ dw. The Wilcoxon/Kruskal Wallis test showed no differences among the three locations ($p > 0.05$). These values were an order of magnitude lower than concentrations found in the shellfish. Arsenic levels in the salmon were comparable to average value of 1.2 $\mu\text{g}\cdot\text{g}^{-1}$ dw derived from Alaska DEC-FMP respectively for sockeye from Kodiak and the Mantanuska River in 2012. Results of the Wilcoxon/Kruskal Wallis test indicated concentration differences between the two types of fish tissue analyzed ($p < 0.05$) with elevated concentration of arsenic in fish liver relative to fish muscle.

The FDA has set the maximum permissible action level of 76 and 86 $\mu\text{g}\cdot\text{g}^{-1}$ arsenic wet weight (ww) in crustaceans and molluscan shellfish respectively. Using the measured 86 % moisture content in shellfish we derived an equivalence value of 602 $\mu\text{g}\cdot\text{g}^{-1}$ arsenic dw in shellfish. The highest arsenic concentrations found in the clams, cockles and salmon from the villages were very low relative to the FDA criterion. EPA (2000) has calculated a reference dose for inorganic arsenic, whereas the data presented here are for total arsenic.

Cadmium

The concentrations of cadmium in the shellfish from the different villages are shown in Figure 4. The highest cadmium concentrations ($1.27 \mu\text{g}\cdot\text{g}^{-1}$ dw) were found in clams from Port Graham, while the lowest concentrations were measured in cockles from the Seldovia harvest grounds. In the 2011 survey, the Alaska DEC-FMP reported tissue values of 0.0 to $2.18 \mu\text{g}\cdot\text{g}^{-1}$ dw cadmium in bivalves from Kachemak Bay. The average cadmium concentration in blue mussels from Figure Figure 3. Concentration of arsenic in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek, Port Graham and Seldovia.

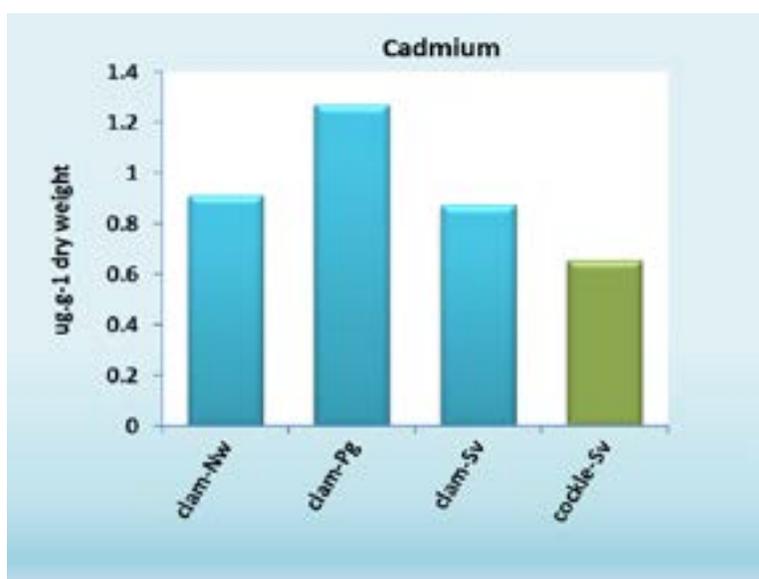


Figure 4. Concentration of cadmium in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

Kachemak Bay in the NOAA NS&T data base is $2.62 \mu\text{g}\cdot\text{g}^{-1}$ dw. As in the case of arsenic, cadmium concentrations in littleneck clams and cockles used for subsistence food were within the regional concentration range found in shellfish.

As illustrated in Tables 6 (a, b, and c), cadmium concentration varied from 0.010 to $5.45 \mu\text{g}\cdot\text{g}^{-1}$, with no obvious concentration differences among the two species of salmon. Wilcoxon/Kruskal Wallis test applied to the combined data indicated significant differences for cadmium concentration between fish gender and

tissue types ($p < 0.05$). For both salmon species, cadmium concentrations in liver were more than 500 times higher than concentrations found in muscle (Figure 5). Also, at all locations, liver tissue of male fish had higher cadmium content compared to liver tissue from female fish ($p < 0.05$). The low cadmium values found in fish muscle were consistent with those of the Alaska DEC-FMP, which reported cadmium concentrations that were at or below reporting limits.

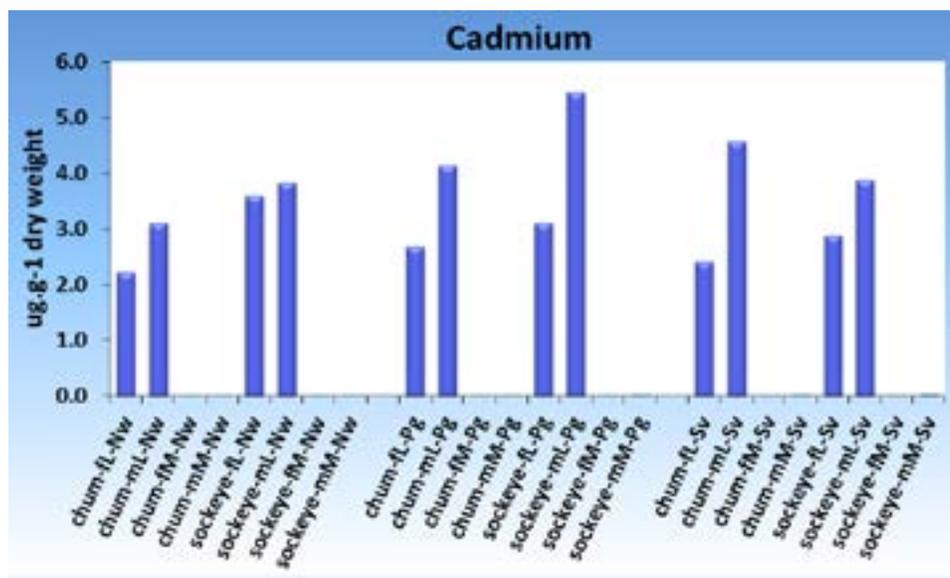


Figure 5. Concentration of cadmium in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

The FDA action level for cadmium in shellfish is $4 \mu\text{g}\cdot\text{g}^{-1}$ wet weight. Using the measured 86 % moisture content in shellfish, we derived an equivalence value of $28 \mu\text{g}\cdot\text{g}^{-1}$ cadmium dw in shellfish. There is no FDA action level for cadmium in fish tissue. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for cadmium in fish filets, the concentration which would be expected to cause no adverse effects for an average person consuming fish daily, is $1.41 \mu\text{g}\cdot\text{g}^{-1}$ dry weight. The average measured concentration in fish muscle was $0.012 \mu\text{g}\cdot\text{g}^{-1}$. There is no comparable reference for fish liver from any source. Thus, concentrations of cadmium in clams and fish tissue are one to two orders of magnitude below applicable safety thresholds.

Chromium

The concentrations of chromium in the shellfish from the different villages are shown in Figure 6. The highest chromium concentration ($4.65 \mu\text{g.g}^{-1} \text{ dw}$) was found in clams from Seldovia, while the lowest concentration ($2.02 \mu\text{g.g}^{-1} \text{ dw}$) was measured in cockles from the Seldovia harvest grounds. In the 2012 survey, the Alaska DEC-FMP reported tissue values from 1.23 to $4.24 \mu\text{g.g}^{-1} \text{ dw}$ chromium in bivalves from Kachemak Bay. The average chromium concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is $1.26 \mu\text{g.g}^{-1} \text{ dw}$. Chromium concentrations in littleneck clams and cockles used for subsistence food were within the regional concentration range found in shellfish.

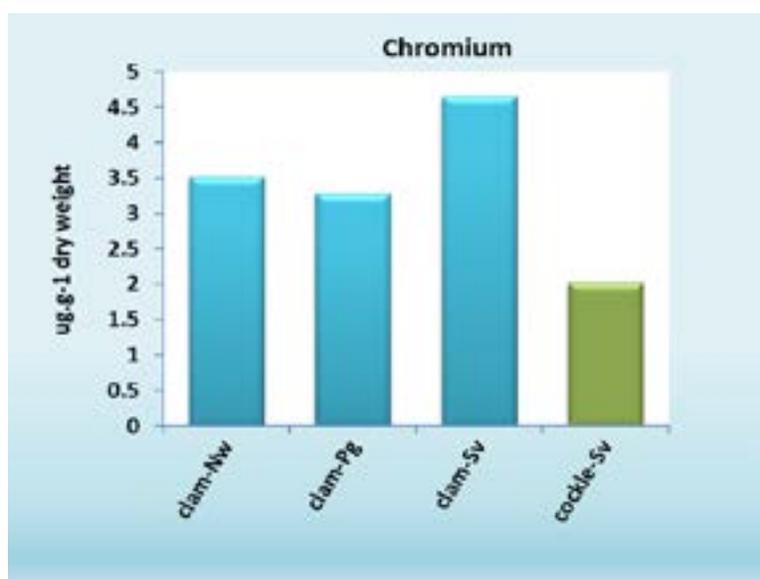


Figure 6. Concentration of chromium in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

Chromium was below detection limits in all fish samples. The low chromium values are consistent with those of the Alaska DEC-FMP, which reported chromium concentrations that were below reporting limits at all locations.

The FDA action level for chromium in shellfish is $13 \mu\text{g.g}^{-1}$. Using the measured 86 % moisture content in shellfish, we derived an equivalence value of $91 \mu\text{g.g}^{-1}$ chromium dw in shellfish. There is no FDA action level or EPA reference dose value for chromium in fish tissue.

Lead

The concentrations of lead in the shellfish from the different villages are shown in Figure 7. The highest lead concentration ($1.49 \mu\text{g}\cdot\text{g}^{-1}$ dw) was found in cockles from Seldovia, while the lowest concentration ($0.27 \mu\text{g}\cdot\text{g}^{-1}$ dw) was measured in clams from the Nanwalek harvest grounds. Both species had higher concentrations in the Seldovia samples. In the 2012 survey, the Alaska DEC-FMP reported tissue values from 0.31 to $0.70 \mu\text{g}\cdot\text{g}^{-1}$ dw lead in bivalves from Kachemak Bay. The average lead concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is $0.59 \mu\text{g}\cdot\text{g}^{-1}$ dw. Concentrations in littleneck clams and cockles used for subsistence food were slightly above the regional concentration range found in shellfish.



Figure 7. Concentration of lead in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

As illustrated in Tables 6 (a, b and c), lead concentrations in fish tissue were either very low or not detectable. Lead was found at detectable concentrations only in chum salmon (Figure 8). The low lead values found in fish muscle were consistent with those of the Alaska DEC-FMP, which reported lead concentrations below detection limits in 2012.

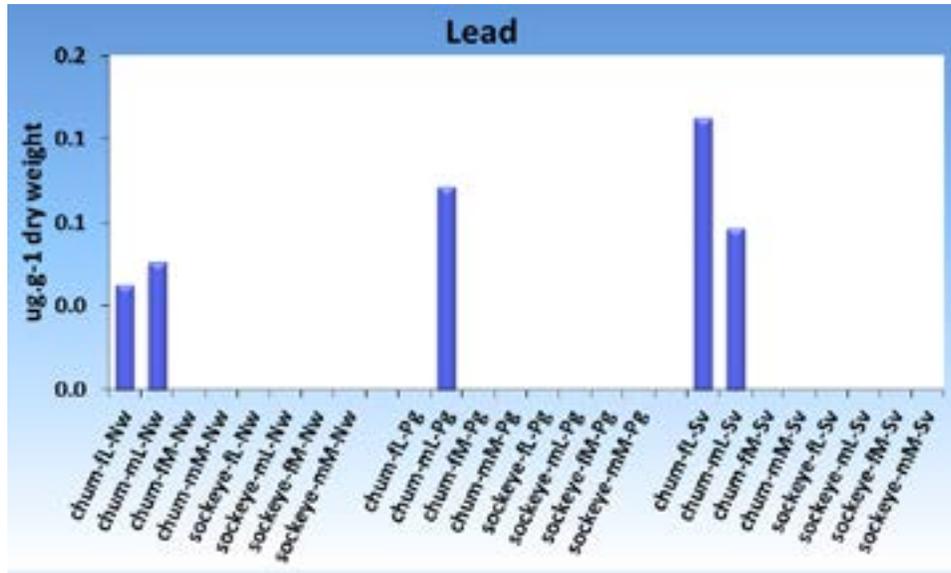


Figure 8. Concentration of lead in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

The FDA action level for lead in shellfish is 1.7 $\mu\text{g.g}^{-1}$. Using the measured 86 % moisture content in shellfish, we derived an equivalence value of 11.9 $\mu\text{g.g}^{-1}$ lead dw in shellfish. There is no FDA action level or EPA reference dose value for lead in fish tissue.

Mercury

The concentrations of mercury in the shellfish from the different villages are shown in Figure 9. The highest mercury concentration (0.21 $\mu\text{g.g}^{-1}$ dw) was found in clams from Seldovia, while the lowest concentration (0.03 $\mu\text{g.g}^{-1}$ dw) was measured in clams from the Nanwalek harvest grounds. In a 2011 survey, the Alaska DEC-FMP reported an average value of 0.1 $\mu\text{g.g}^{-1}$ dw total mercury in razor clams from the Lower Cook Inlet. In the 2012 survey, mercury was below detection limits in Razor clams from Redoubt Creek. The average mercury concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is 0.082 $\mu\text{g.g}^{-1}$ dw.

As illustrated in Tables 6a, 6b and 6c, mercury concentrations in fish tissue were similar to the shellfish levels. Mercury concentrations were significantly higher in liver than in muscle ($p < 0.01$), but only in the sockeye salmon (Figure 10). The mean mercury concentration in muscle was 0.11 $\mu\text{g.g}^{-1}$ dw. The mercury values found in fish muscle were consistent with those of the Alaska DEC-FMP, which reported mercury concentrations of 0.100 to 1.120 $\mu\text{g.g}^{-1}$ dw.

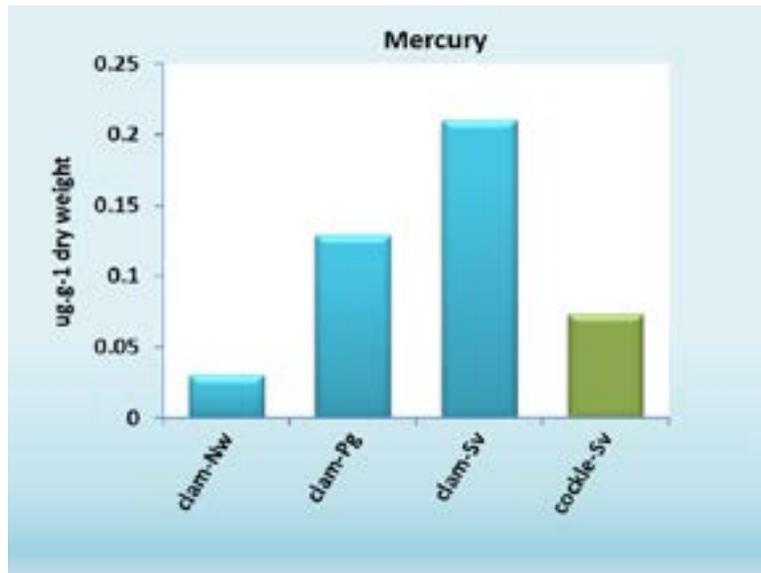


Figure 9. Concentration of mercury in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

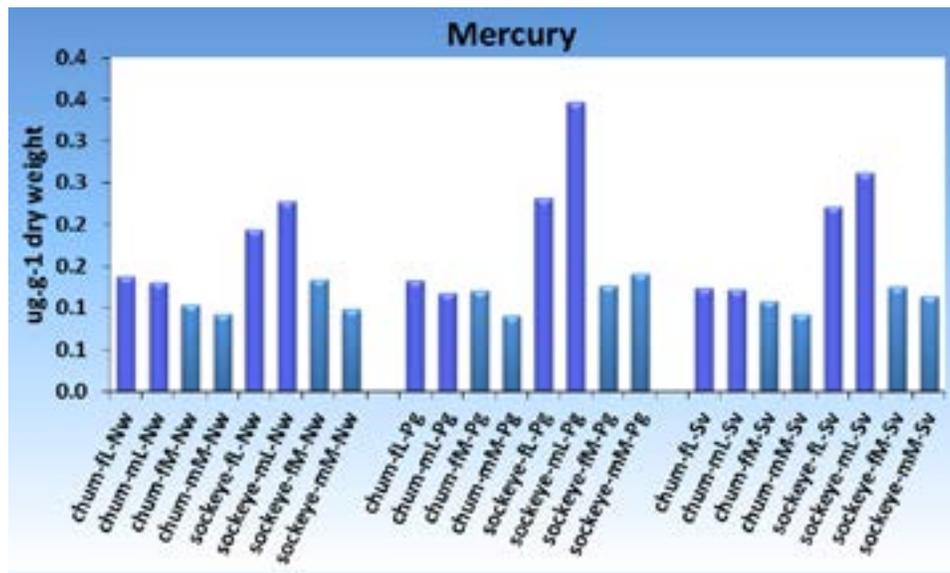


Figure 10. Concentration of mercury in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native village of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

There is no FDA action level for mercury in shellfish tissue. There is an FDA action level of 1 $\mu\text{g}\cdot\text{g}^{-1}$ for methylmercury in fish tissue. Using the measured 76% moisture content of the fish tissue, the FDA action

level is $4 \mu\text{g}\cdot\text{g}^{-1}$ dw. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for mercury in fish filets, the concentration which would be expected to cause no adverse effects for an average person consuming fish daily is $0.14 \mu\text{g}\cdot\text{g}$ dry weight. It should also be noted that the EPA reference dose is for methylmercury, whereas the data presented here are for total mercury. In fish tissue, the majority of mercury is methyl mercury (EPA, 2000). There is no comparable reference for fish liver from any source.

Nickel

The concentrations of nickel in the shellfish from the different villages are shown in Figure 11. The highest nickel concentration ($12.1 \mu\text{g}\cdot\text{g}^{-1}$ dw) was found in cockles from Seldovia, while the lowest concentration ($3.27 \mu\text{g}\cdot\text{g}^{-1}$ dw) was measured in clams from the Port Graham harvest grounds. Both species had higher concentrations in the Seldovia samples. In the 2012 survey, the Alaska DEC-FMP reported tissue values of 0.31 to $0.70 \mu\text{g}\cdot\text{g}^{-1}$ dw nickel in bivalves from Kachemak Bay. The average nickel concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is $1.83 \mu\text{g}\cdot\text{g}^{-1}$ dw. Concentrations in littleneck clams and cockles used for subsistence food were slightly above the regional concentration range found in shellfish.

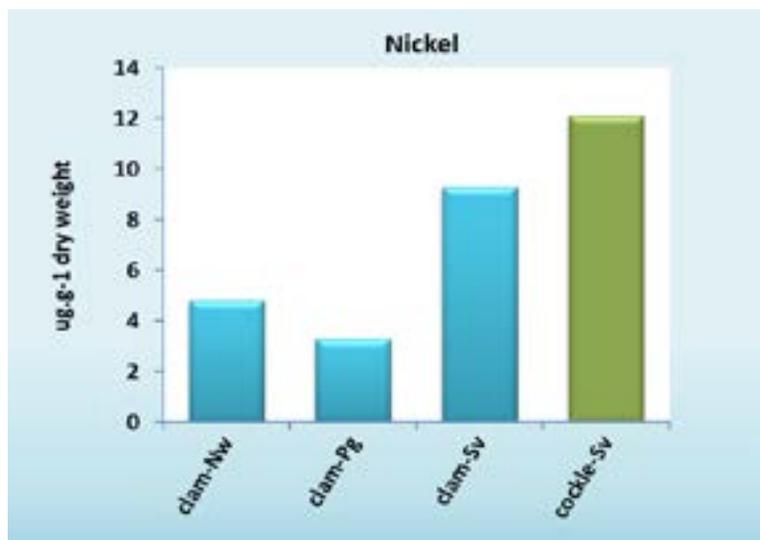


Figure 11. Concentration of nickel in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

As illustrated in Tables 6 (a, b and c), nickel concentrations in salmon vary by tissue, again, primarily in the sockeye salmon where liver concentrations were significantly higher than muscle ($p < 0.01$). Concentrations

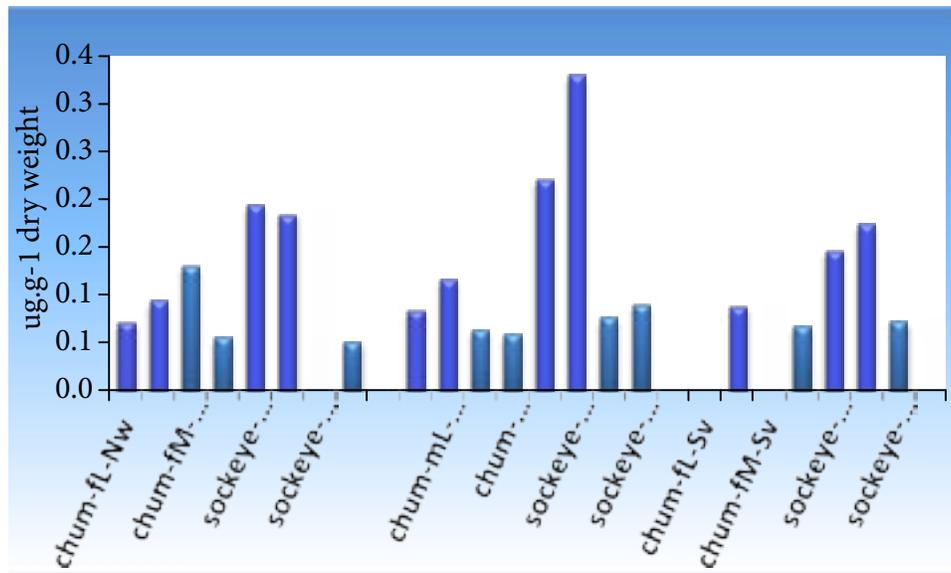


Figure 12. Concentration of nickel in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

were variable with some values at or below the detection limit, but with no obvious pattern (Figure 12). The low nickel values found in fish muscle were consistent with those of the Alaska DEC-FMP, which reported nickel concentrations below detection limits in sockeye salmon.

The FDA action level for nickel in shellfish is $80 \mu\text{g}\cdot\text{g}^{-1}$. Using the measured 86 % moisture content in shellfish, we derived an equivalence value of $560 \mu\text{g}\cdot\text{g}^{-1}$ nickel dw in shellfish. There is no FDA action level or EPA reference dose value for nickel in fish tissue.

Selenium

The concentrations of selenium in the shellfish from the different villages are shown in Figure 13. The highest selenium concentration ($3.28 \mu\text{g}\cdot\text{g}^{-1}$ dw) was found in clams from Seldovia, while the lowest concentration ($1.89 \mu\text{g}\cdot\text{g}^{-1}$ dw) was measured in clams from the Nanwalek harvest grounds. Both species had higher concentrations in the Seldovia samples. In the 2011 survey, the Alaska DEC-FMP reported an average value of $3.8 \mu\text{g}\cdot\text{g}^{-1}$ dw selenium in razor clams from the Lower Cook Inlet. In the 2012 survey, razor clams in Redoubt Creek had a mean concentration of $0.53 \mu\text{g}\cdot\text{g}^{-1}$. The average selenium concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is $2.89 \mu\text{g}\cdot\text{g}^{-1}$ dw. Concentrations in littleneck clams and cockles used for subsistence food were within the regional concentration range found in shellfish.

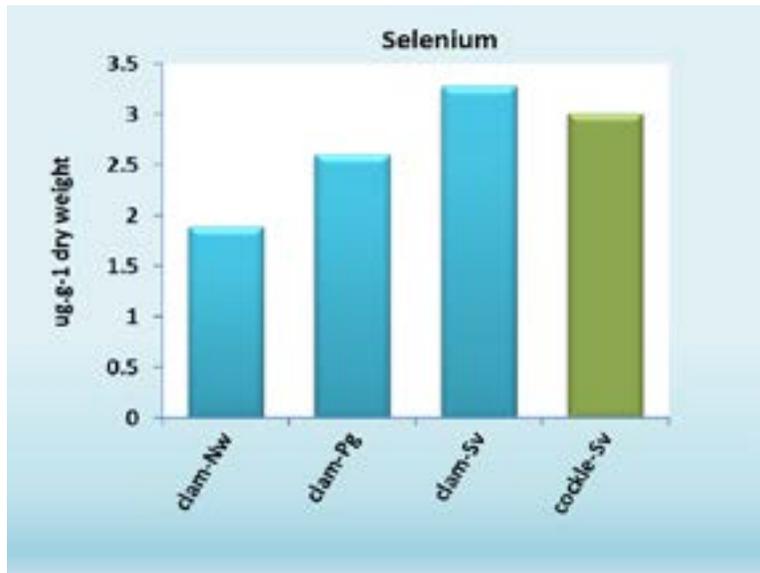


Figure 13. Concentration of selenium in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

As illustrated in Tables 6 (a, b and c), selenium concentrations vary by tissue, in both species (Figure 14). Liver concentrations were significantly higher than muscle ($p < 0.01$). The low selenium values found in fish muscle

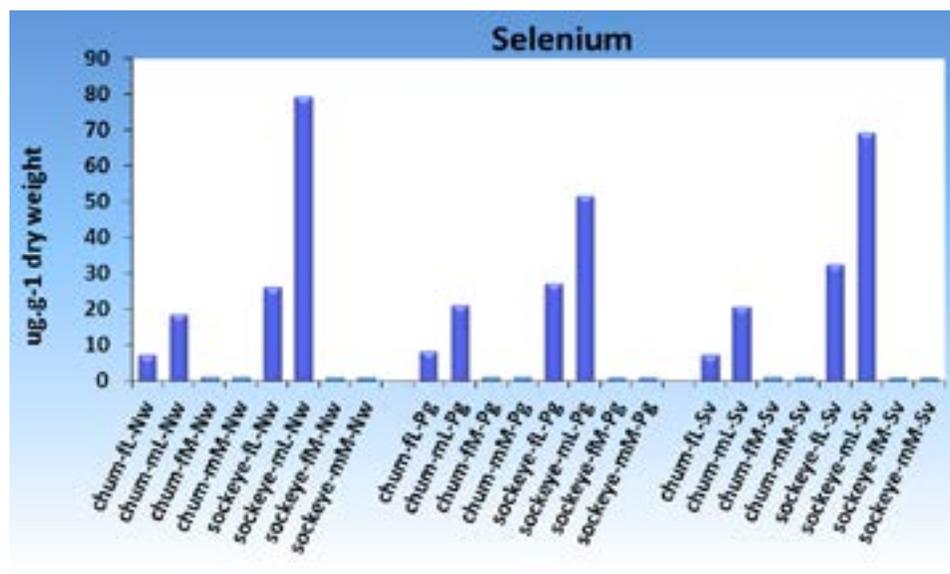


Figure 14. Concentration of selenium in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native village of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

were consistent with those of the Alaska DEC-FMP, which reported selenium concentrations of 0.88 and 0.84 $\mu\text{g}\cdot\text{g}^{-1}$ dw in sockeye salmon.

There is no FDA action level for selenium in shellfish tissue. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for selenium in fish filets, the concentration which would be expected to cause no adverse effects for an average person consuming fish daily is 7.1 $\mu\text{g}\cdot\text{g}^{-1}$ dry weight. The average measured concentration in fish muscle was 1.06 $\mu\text{g}\cdot\text{g}^{-1}$. There is no comparable reference for fish liver from any source.

4.2 Organic contaminant concentrations in fish and shellfish

Detailed descriptions of contaminant concentrations found in molluscs and salmon in this study are limited to well-known and well-studied compounds. Calculated EPA chronic threshold tissue concentrations and FDA action levels for all contaminants are shown in Tables 5 and 6a-c. All of the pesticide thresholds were one to two orders of magnitude greater than any tissue concentration seen in this study.

Chlordanes

The concentrations of chlordane in the shellfish from the different villages are shown in Figure 15. The highest chlordane concentration (1.48 $\text{ng}\cdot\text{g}^{-1}$ dw) was found in clams from Port Graham, while the lowest concentration (0.27 $\text{ng}\cdot\text{g}^{-1}$ dw) was measured in clams from the Nanwalek harvest grounds. Both species had higher concentrations in the Port Graham samples. The average chlordane concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is 1.46 $\text{ng}\cdot\text{g}^{-1}$ dw. Concentrations in littleneck clams and cockles used for subsistence food were within the regional concentration range found in shellfish.

As illustrated in Tables 6a, 6b and 6c, chlordane concentrations in salmon vary by tissue in both species (Figure 16). Sockeye salmon tended to have higher levels of chlordane, but this was not true in all cases. The low chlordane values found in fish muscle were not consistent with those of the Alaska DEC-FMP, which reported higher chlordane concentrations of 11.3 and 6.88 $\text{ng}\cdot\text{g}^{-1}$ dw in sockeye salmon. According to earlier reports, muscle tissue concentrations in Chinook, pink and Coho salmon were 21.52, 1.62, and 3.22 $\text{ng}\cdot\text{g}^{-1}$ dw, respectively.

There is no FDA action level for chlordane in shellfish tissue. The FDA action level for chlordane in fish tissue is 1.2 $\mu\text{g}\cdot\text{g}^{-1}$ dw. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for chlordane in fish filets, the concentration which would be expected to cause no adverse effects for

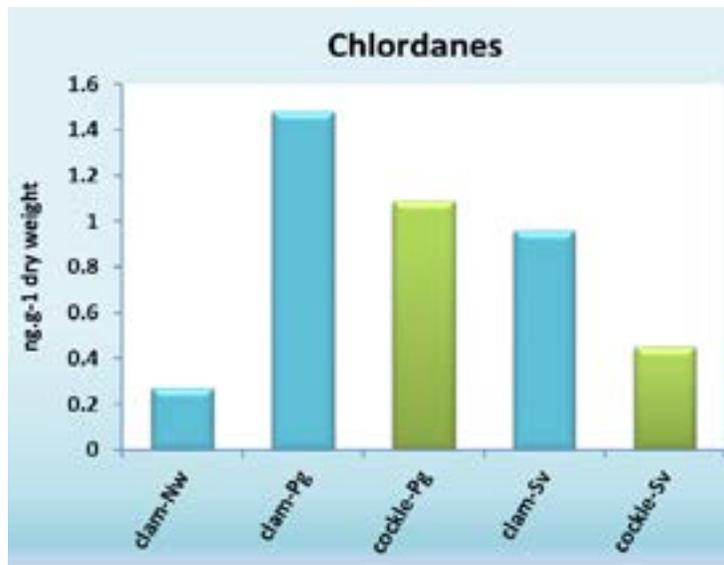


Figure 15. Concentration of total chlordanes in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

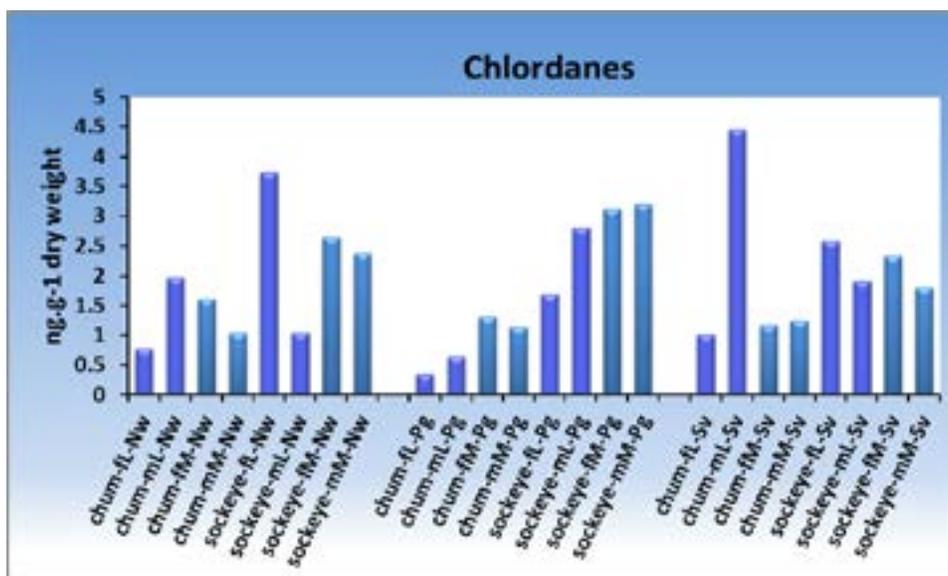


Figure 16. Concentration of total chlordanes in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

an average person consuming fish daily is 704.8 ng.g⁻¹ dw. The average measured concentration in fish muscle was 1.90 ng.g⁻¹. There is no comparable reference for fish liver from any source.

DDTs

The concentrations of DDT in the shellfish from the different villages are shown in Figure 17. The highest DDT concentration ($0.73 \text{ ng.g}^{-1} \text{ dw}$) was found in clams from Seldovia, while the lowest concentration ($0.17 \text{ ng.g}^{-1} \text{ dw}$) was measured in cockles from the Port Graham harvest grounds. Both species had higher concentrations in the Seldovia samples. The average DDT concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is $0.47 \text{ ng.g}^{-1} \text{ dw}$. Concentrations in littleneck clams and cockles used for subsistence food were within the regional concentration range found in shellfish.

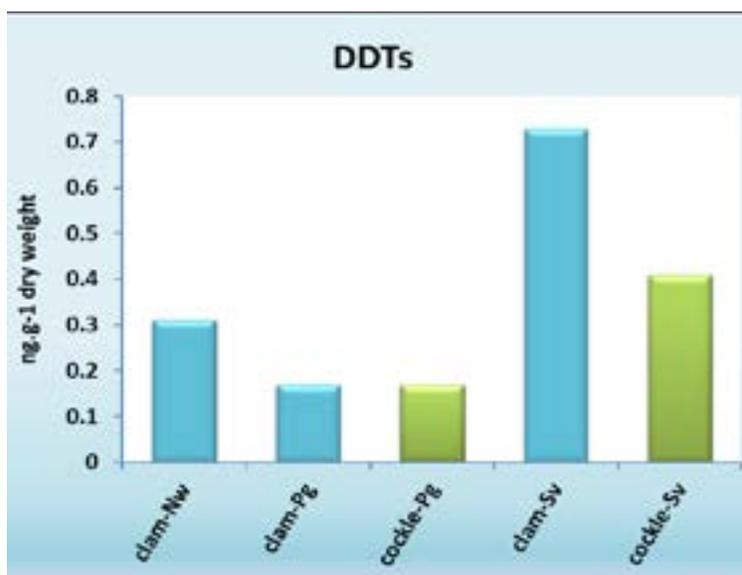


Figure 17. Concentration of total DDTs in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and

As illustrated in Tables 6a, 6b and 6c, DDT concentrations in salmon vary by tissue in both species (Figure 18). Sockeye salmon tended to have higher levels of DDT, but this was not true in all cases. In muscle tissue, however, sockeye had much higher concentrations (8.46 ng.g^{-1}) than the chum salmon (2.44 ng.g^{-1}). The differential DDT values found in the two species were consistent with those of the Alaska DEC-FMP, although they report higher values for both locations (19.04 and 38.12 ng.g^{-1}), as was seen with chlordane.

There is no FDA action level for DDT in shellfish tissue. Using the measured 76% moisture content of the fish muscle, the FDA action level for DDT in fish tissue is $20,000 \text{ ng.g}^{-1} \text{ dw}$. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for DDT in fish filets, the concentration which

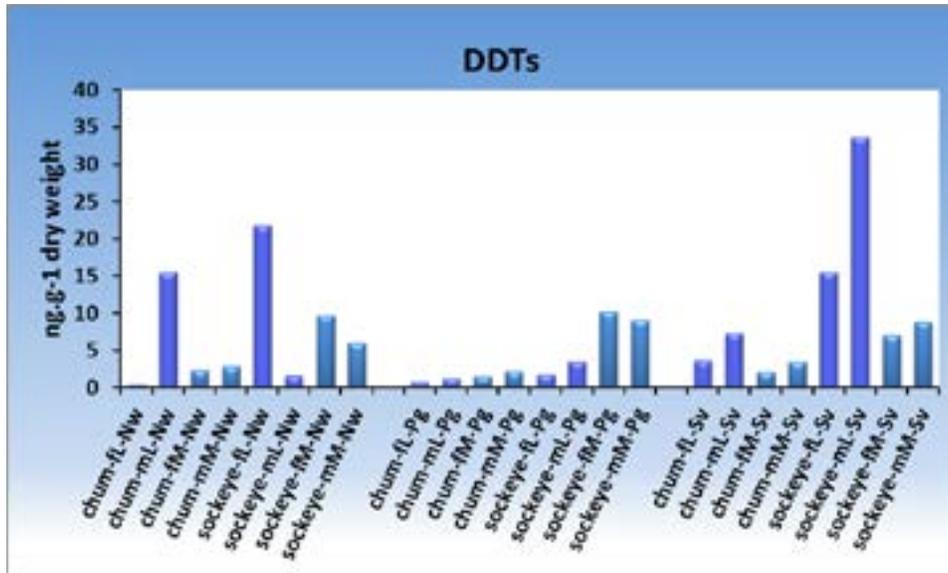


Figure 18. Concentration of total DDTs in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

would be expected to cause no adverse effects for an average person consuming fish daily is 704.8 ng.g⁻¹ dw dry weight. There is no comparable reference for fish liver from any source.

PAHs

The concentrations of PAHs in the shellfish from the different villages are shown in Figure 19. The highest PAH concentration (302 ng.g⁻¹ dw) was found in clams from Port Graham, while the lowest concentration (143.8 ng.g⁻¹ dw) was measured in clams from Nanwalek harvest grounds. The average PAH concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is 167.8 ng.g⁻¹ dw, but varied between 72.1 and 263.7 ng.g⁻¹ dw. Concentrations in littleneck clams and cockles used for subsistence food overlap the regional concentration range found in shellfish. There are no applicable FDA action levels or EPA reference doses for PAHs in shellfish. The average PAH concentration in blue mussels from the entire data set from southcentral and southeast Alaska (15 stations) in the NOAA NS&T data base is 304.84 ng.g⁻¹ dw, and varies between 28.13 and 1,026.23 ng.g⁻¹ dw. PAHs were not measured in fish tissue because vertebrates are able to break down PAHs to a much greater extent than mollusks.

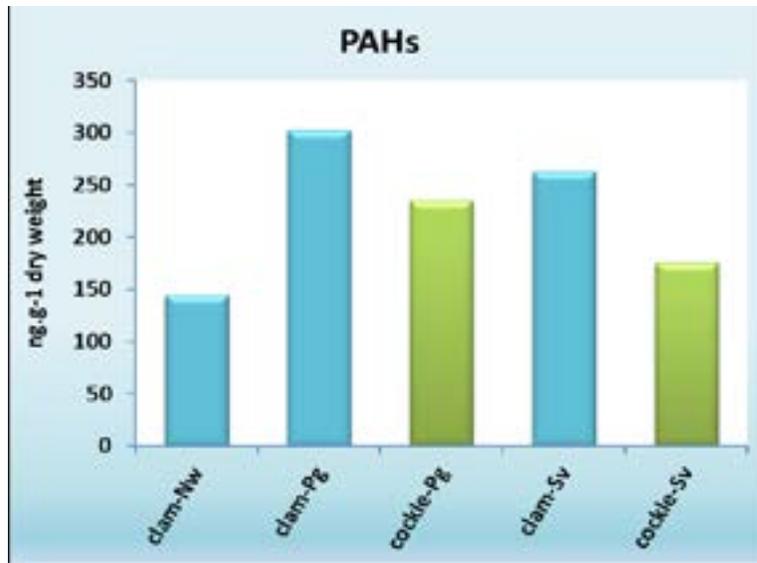


Figure 19. Concentration of total total PAHs in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

PCBs

The concentrations of PCBs in the shellfish from the different villages are shown in Figure 20. The highest PCBs concentration (10.1 ng.g⁻¹ dw) was found in clams from Seldovia, while the lowest concentration (0.92 ng.g⁻¹ dw) was measured in cockles from the Port Graham harvest grounds. The average PCBs concentration in blue mussels from Kachemak Bay in the NOAA NS&T data base is 2.47 ng.g⁻¹ dw. Concentrations in littleneck clams and cockles used for subsistence food overlap the regional concentration range found in shellfish.

As illustrated in Tables 6a, 6b and 6c, PCBs concentrations in salmon vary by tissue in both species and by location (Figure 21). Sockeye salmon tended to have higher levels of PCBs, but this was not true in all cases. Port Graham sockeye muscle tissue had much higher concentrations (13.15 ng.g⁻¹) than the other values, but the same was not seen at Nanwalek or Seldovia. The Alaska DEC-FMP reports values of 23.60 and 11.85 ng.g⁻¹ in sockeye salmon from Kodiak and the Matanuska River, respectively.

There is no FDA action level for PCBs in shellfish tissue. Using the measured 76% moisture content of the fish muscle, the FDA action level for PCBs in fish tissue is 8,000 ng.g⁻¹ dw. Using the measured 76% moisture content of the fish muscle, and the EPA reference dose value for total PCBs in fish filets, the concentration which would be expected to cause no adverse effects for an average person consuming fish daily is 28 ng.g⁻¹ dw dry weight. The EPA reference dose is based on analyses of a specific PCBs compound known as “aroclor.” Once PCBs are released into the environment, they spread out over and time and are slowly degraded. After a

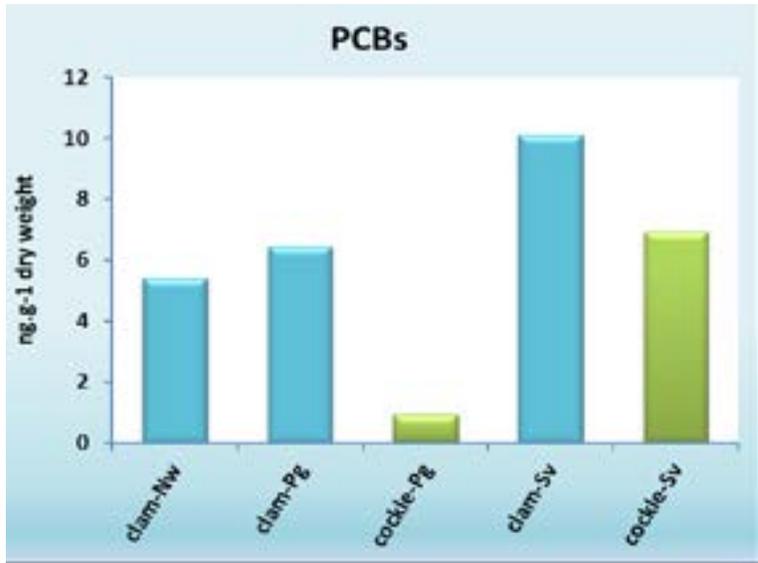


Figure 20. Concentration of total total PCBs in clams and cockles collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

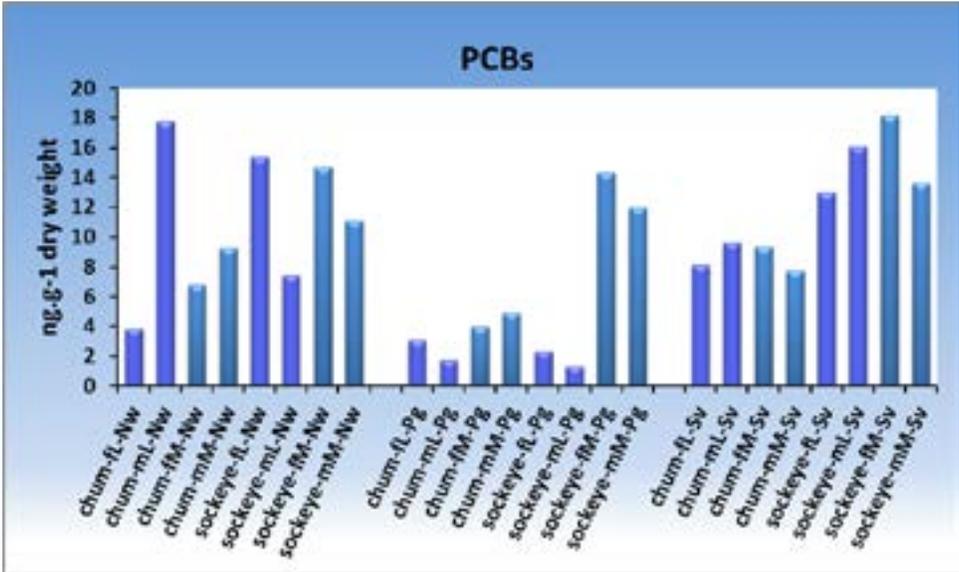


Figure 21. Concentration of total PCBs in liver and muscle tissues of chum and sockeye salmon collected from subsistence harvest grounds of the Alaskan Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

certain length of time, they no longer resemble their original structure. The data presented here are the sum of the PCBs actually measured. There is no clear relationship at this time between total Aroclors and total PCBs that have been degraded. There is no comparable reference for fish liver from any source.

4.3 Histopathology characterization in shellfish

Results of prevalences (% affected) and intensity of parasitic infections and pathological/disease in cockles and softshell clams sampled from subsistence harvest grounds in Nanwalek, Port Graham, and Seldovia, Alaska are summarized in Table 7, and presented graphically in Figures 22 and 23.

Parasitic infection

Parasitic copepods infection: Copepods are small, hard-shelled creatures related to crabs and shrimp that are frequently referred to as the “insects of the sea.” Copepod infections were observed in clams and cockles sampled from the harvest grounds of Seldovia (Figure 22). There are two groups of parasitic copepods which infect bivalves. They affect the digestive tract and the mantle and gills of bivalves (Heegaard, 1962; Darwin and Stefanich, 1966). However, in bivalves such as clams and cockles, parasitic copepods are typically found in the digestive tract and rarely in the gills (Johnson et al., 2004; Kim et al., 2006). In this study, they were found in about 40% of the samples (Figure 22), of softshell clams and cockles from the Seldovia harvest grounds. Intensity values ranged from 1 to 2.5 (Figure 23). These are not dangerous to humans.

Nematode (roundworms) infection: Nematodes, or roundworms, were detected in clams from the harvest grounds of Seldovia (Figure 22). With over a million species, roundworms are very diverse, with many thousands of species described as disease causing (Hugot et al., 2001). Roundworms have multiple development stages. According to Cheng (1978), roundworms that infect shellfish such clams and cockles are mainly in larval stages, while adults can be found in the predators of the mollusks. As illustrated in Figure 22, prevalence values for the occurrence of the roundworms in the softshell clams from this study were low (only 20% prevalence). Some roundworms can have negative health impacts on humans.

Gregarine infection: Also found in the softshell clams from the harvest grounds of Nanwalek and Seldovia were gregarines, which are single celled microscopic parasites (parasitic protozoans) (Figure 22). They have several life stages with larval stages frequently found in bivalves, while mud and stone crabs were found to be the most common final hosts (Kim et al., 2006). Gregarines are often found in the digestive region of their hosts, but may invade other tissues (Kim et al., 2006). The results indicated that gregarine infections were relatively more intense in clams from Seldovia than those from Nanwalek (Figure 23). However, with

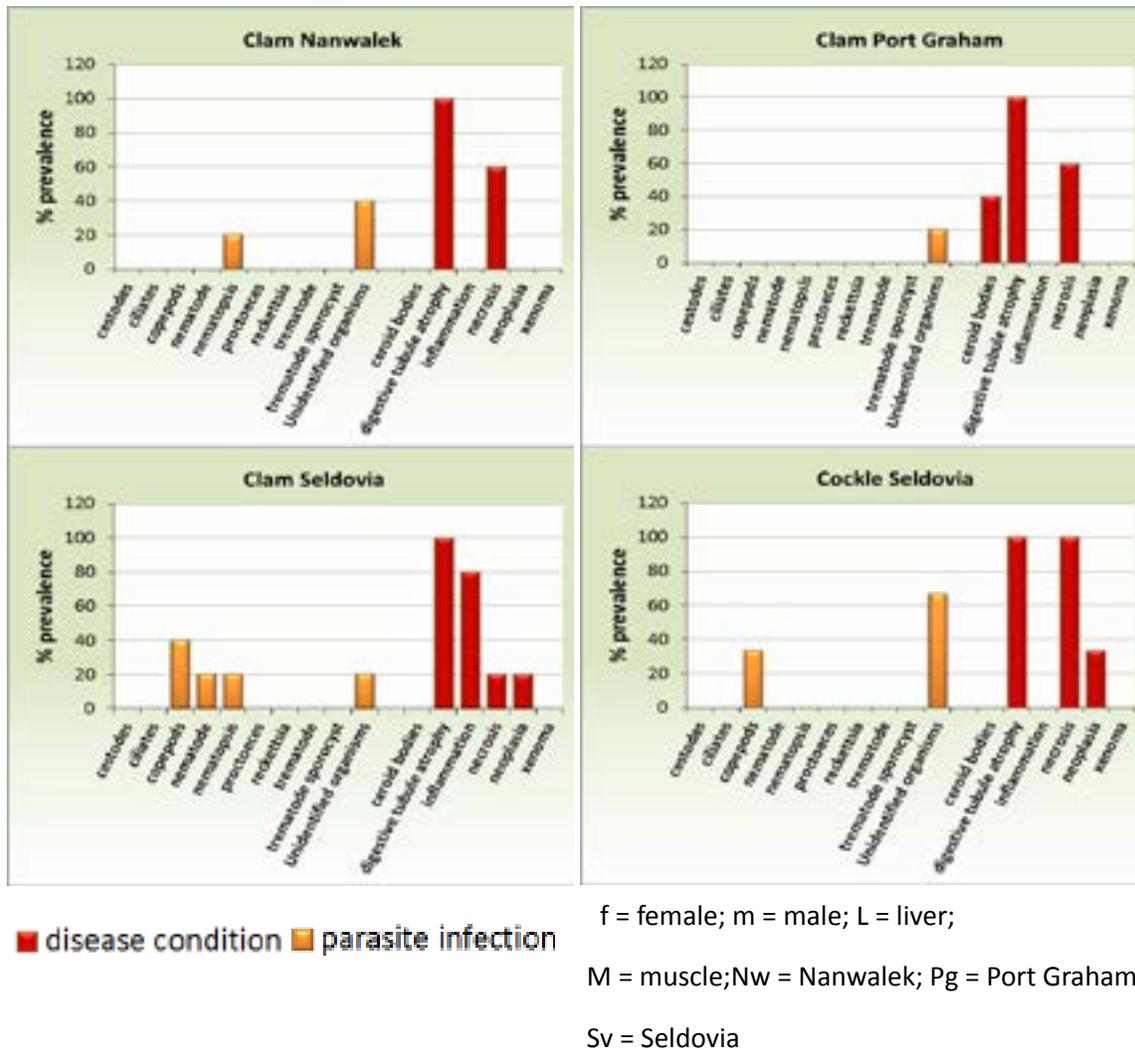
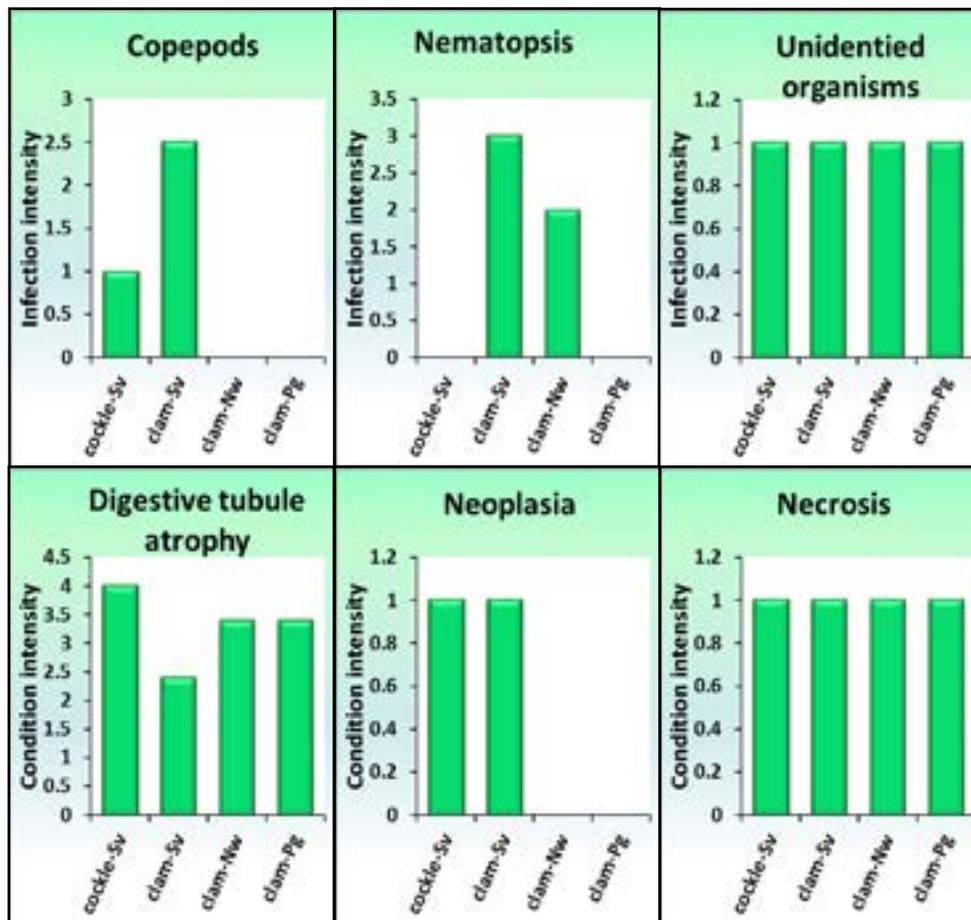


Figure 22. Prevalence (%) of parasite infections and histological lesions in cockles and softshell clams collected from subsistence harvest grounds of the Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

prevalence of occurrence of only 20% at both locations, gregarine infections appeared to be relatively low in the clams. Gregarines are not harmful to humans. Other harmful organisms that can infect shellfish, such as ciliates, cestodes, trematodes, Rickettsia, xenoma and MSX, were not observed in any specimen.

Disease conditions

Pathological and disease conditions were detected at various degrees in the softshell clams and cockles, depending on species and location (Table 7 and Figure 22 and 23). Xenoma, a condition which usually results from enlargement of tissue infected by parasites, were not observed, but other indications of disease, including inflammation, were detected in the shellfish.



f = female; m = male; L = liver; M = muscle; Nanwalek; Pg = Port Graham; Sv = Seldovia

Figure 23. Intensity of parasite infections and histological lesions in cockles and softshell clams collected from subsistence harvest grounds of the Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

Ceroid bodies: Ceroid bodies, a symptom of cellular disease, are primarily an accumulation of fats (lipofuscinosis) resulting from cellular damage and/ or metabolic unbalance. They were detected at about a 40 % prevalence in clams from Port Graham.

Digestive tubule atrophy: The condition of digestive tubule atrophy (wasting away) was the most common condition in the shellfish, occurring at a 100% prevalence, with intensity of occurrence reaching 4, particularly in cockles from the Seldovia harvest grounds (Figures 20 and 21).

Tissue inflammation: Cases of tissue inflammation were also observed in the softshell clams from the Seldovia harvest grounds. Tissue inflammation conditions were not detected shellfish from Nanwalek or Port Graham,

Table 7. Prevalences (% affected) and intensity of histological (disease) conditions and parasitic infections in cockles and softshell clams sampled in May 2010 from subsistence harvest grounds in Nanwalek, Port Graham, and Seldovia in Alaska.

Histopathology parameter	Nanwalek clams		Port Graham clams		Seldovia clams		Seldovia cockles	
	Prevalence (%)	Intensity	Prevalence (%)	Intensity	Prevalence (%)	Intensity	Prevalence (%)	Intensity
cestodes	0	0	0	0	0	0	0	0
ciliates	0	0	0	0	0	0	0	0
copepods	0	0	0	0	40	2.5	33.3	1
nematode	0	0	0	0	20	2	0	0
reckettsia	0	0	0	0	0	0	0	0
trematode	0	0	0	0	0	0	0	0
trematode sporocyst	0	0	0	0	0	0	0	0
proctoece	0	0	0	0	0	0	0	0
nematopsis	20	2	0	0	20	3	0	0
xenoma	0	0	0	0	0	0	0	0
Unidentified organisms	40	1	20	1	20	1	66.7	1
unidentified foll org	0	0	0	0	0	0	0	0
MSX	0	0	0	0	0	0	0	0
ceroid bodies	0	0	40	20	0	0	0	0
digestive tubule atrophy	100	3.4	100	3.4	100	2.4	100	4
inflammation	0	0	0	0	80	1	0	0
necrosis	60	1	60	1	20	1	100	1
neoplasia	0	0	0	0	20	1	33.3	1
unusual digestive tract	0	0	0	0	0	0	0	0
Replicates (N)	5		5		5		3	
wet weight (g) whole	10.5		17.66		17.02		13.43	
shell length (cm)	5.44		7.32		7.28		5.03	

however, the condition was frequent in softshell clams from Seldovia, with a prevalence value of 80% (Figure 22).

Tissue necrosis: Conditions of cell death in living tissue, or tissue necrosis, were detected at various degrees in shellfish from virtually all of the three harvest grounds (Table 7, Figures 20 and 21). While tissue necrosis was measured at 20% prevalence in clams from Seldovia and at 60% in clams from Nanwalek and Port Graham, the condition reached 100% prevalence in cockles from Seldovia (Figure 22).

Tissue neoplasia: Cases of tissue neoplasia (harmless tumors) were observed in both clam and cockle species collected from the Seldovia harvest grounds (Table 7, Figures 20 and 21). Tissue neoplasms were relatively

low from the Seldovia harvest grounds, at about 33% and 20% prevalence in cockles and softshell clams, respectively (Figure 22).

4.4 Histopathology characterization in fish

Parasitic infection

Nematodes (roundworms) infection: The histopathological examination of adult chum and sockeye salmon captured from traditional harvest grounds of Nanwalek, Port Graham and Seldovia were limited to parasitic infections/infestations of the gills, kidney and liver tissues, and a single noninfectious condition observed in the gills (Table 8 and Figure 24). Tissue inflammations were observed where roundworms infections occurred in the fish liver. These inflammations are a typical chronic host response to nematode infections. In female sockeye salmon, the prevalence of nematode infections ranged from 0% at Seldovia to 40% at Port Graham and Nanwalek. In male sockeye, prevalences ranged from 0% at Nanwalek to 20% at Seldovia and Port Graham. Overall prevalences of the nematode infections in sockeye were low, with 10% at Seldovia, 30% at Port Graham, and 20% at Nanwalek. These prevalences were not significantly different by the Fisher's Exact Test ($p < 0.05$). The vast majority of livers in both species were normal; in fact, in chum salmon, all livers were normal.

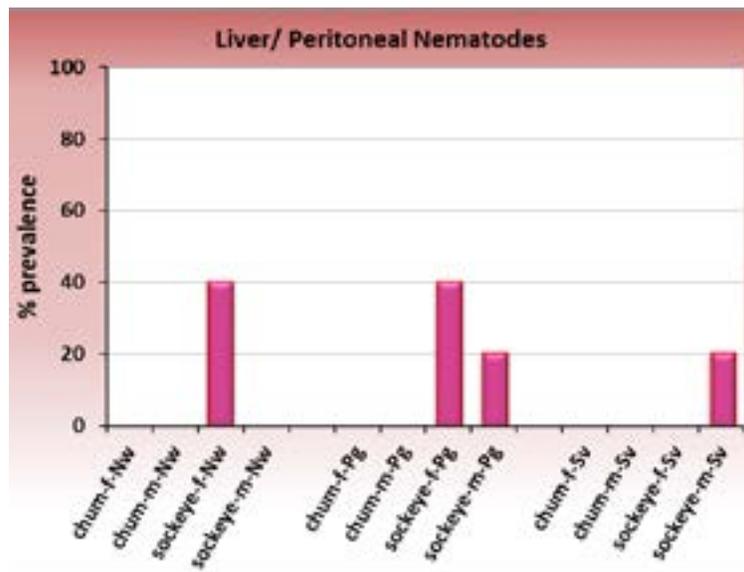
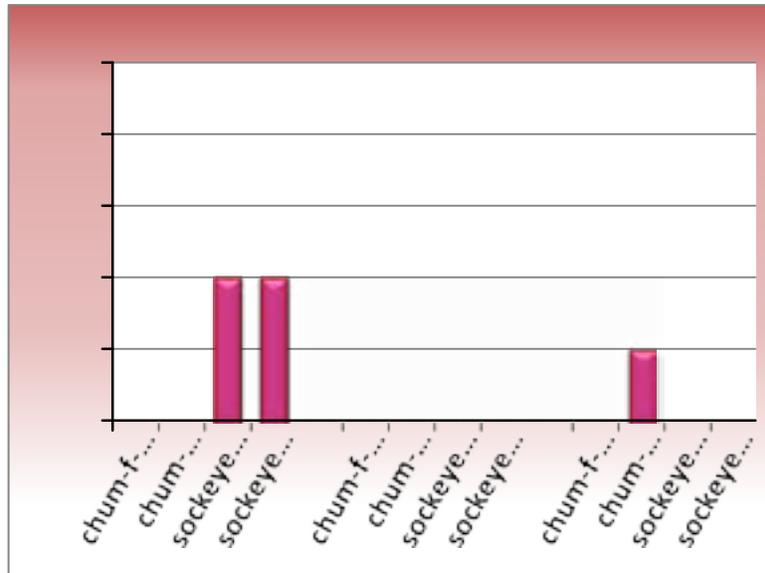


Figure 24. Prevalence of nematodes in liver peritoneal cavities of chum and sockeye salmon collected from subsistence harvest grounds of the Alaska Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

Table 8. Prevalences (% affected) of histological conditions and parasitic infections in adult chum and sockeye salmon (female, f and male m) sampled in July 2010 from subsistence fisheries in Nanwalek, Port Graham, and Seldovia in Alaska.

organ lesion/parasites	Nanwalek						Port Graham						Seldovia					
	Chum		sockeye		Chum		sockeye		Chum		sockeye		Chum		sockeye			
	f	m	f	m	f	m	f	m	f	m	f	m	f	m	f	m		
Sex	100	100	60	100	100	100	60	80	100	100	100	60	100	100	100	100	80	
Liver, Normal	0	0	40	0	0	0	40	20	0	0	0	40	20	0	0	0	20	
Liver, Peritoneal Nematodes	100	100	40	60	100	100	40	100	100	100	100	40	100	100	100	100	100	
Kidney, Normal	0	0	40	40	0	0	40	0	0	0	0	40	0	0	0	0	0	
Kidney, Tubular Myxosporidan	60	0	100	100	80	60	100	100	60	60	100	60	60	60	60	60	40	
Gill, Normal	20	100	0	0	20	20	0	0	20	20	0	20	20	20	20	20	20	
Gill, Loma salmonis (microsporidan)	20	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	40	
Gill, monogenetic trematodes (external)	20	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
Gill, Lamellar Microaneurysms	5	2	5	5	5	5	5	5	5	5	5	5	5	5	5	5	5	
Replicates (N)	60.7	64	56.8	55.3	61.5	62.5	55.3	59.7	60.7	60.8	51.9	54.3	54.3	54.3	54.3	54.3	54.3	
Average fish length (cm)	2946	3420	2132	2212	3140	3202	2165	2632	2950	2650	1700	1924	1924	1924	1924	1924	1924	
Weight (g) whole fish																		

Myxosporidians (*Myxidium sp.*) infection: In this study, myxosporidan (small, single celled organisms) parasites were detected in the chum and sockeye salmon (Figure 25) at various levels in the kidney tissue of the fish. Prevalence values were 20% in the kidney of male chum salmon from Seldovia, and 40% in both female and male sockeye from Nanwalek. The results indicated that the myxosporidan infections were minor in severity, and were not associated with any tissue pathology. Myxosporidan are not dangerous to humans.



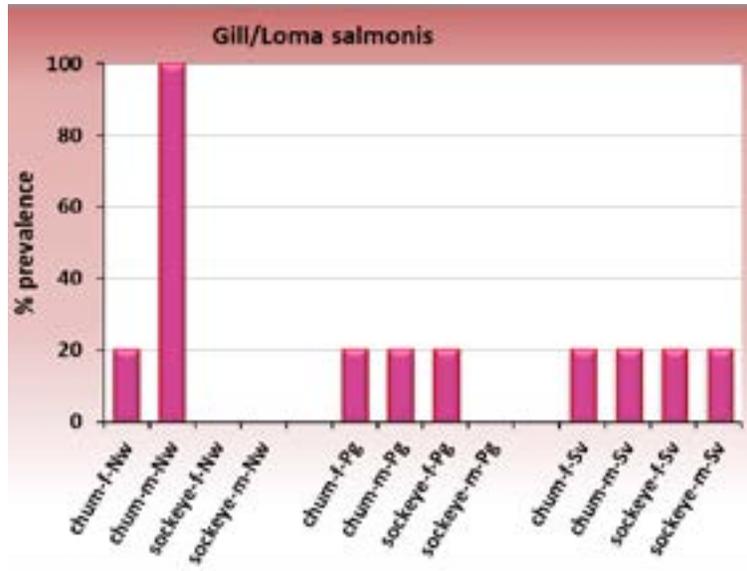
f = female; m = male; L = liver; M = muscle;

Nw = Nanwalek; Pg = Port Graham; Sv = Seldovia **Figure 25.**

Prevalence of myxosporidan parasites in kidney of chum and sockeye salmon collected from subsistence harvest grounds of the Alaska Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).

Microsporidan (*Loma salmonis*) infection: Infection by this microsporidan (spore-forming single celled parasites) which, unlike those mentioned above, specifically targets salmon, was detected in the gill tissue of fish of both species and sexes from virtually all of the areas sampled, with the exception of Nanwalek sockeye (Table 8 and Figure 26). In male chum salmon, prevalences of the microsporidan infection ranged from 20% at Seldovia and Port Graham to 100% at Nanwalek (N=2 male chum at Nanwalek). In female chum, prevalences at all three sites were 20%. The prevalences of the microsporidan infection in chum were 20% at Seldovia and Port Graham, and 43% at Nanwalek. In male sockeye, prevalences of the infection ranged from 0% at Port Graham and Nanwalek to 20% at Seldovia. In female sockeye, prevalences ranged from 0% at Nanwalek to 20% at both Seldovia and Port Graham. Overall prevalences were 0% at Nanwalek, 10% at Port Graham, and 20% at Seldovia Bay. Overall the differences among these prevalences were not statistically significant ($p < 0.05$). This parasite is not dangerous to humans.

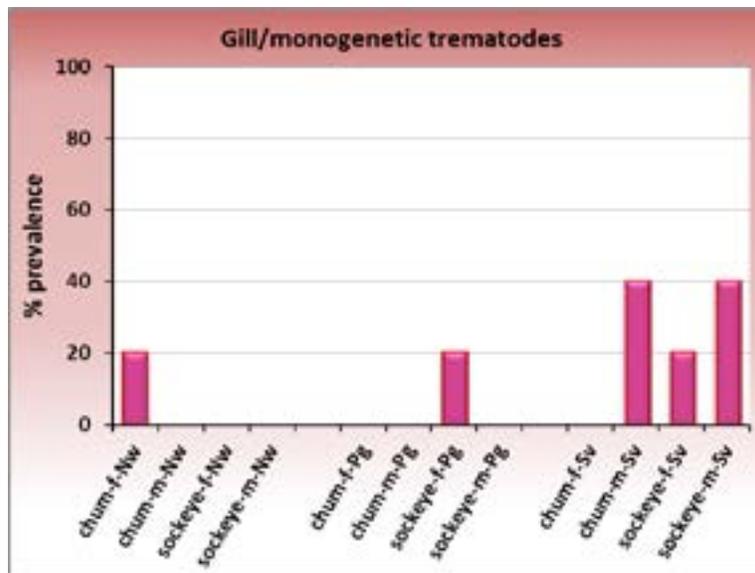
Trematodes (flukes or flatworms) infection: External infestations of the gill by trematode parasites, probably of the family Gyrodactylidae, were observed in both species of salmon (Table 8 and Figure 27). In male



f = female; m = male; L = liver; M = muscle;

Nw = Nanwalek; Pg = Port Graham; Sv = Seldovia

Figure 26. Prevalence of *Loma salmonis* in gill of chum and sockeye salmon collected from subsistence harvest grounds of the Alaska Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).



f = female; m = male; L = liver; M = muscle;

Nw = Nanwalek; Pg = Port Graham; Sv = Seldovia

Figure 27. Prevalence of trematodes in gill of chum and sockeye salmon collected from subsistence harvest grounds of the Alaska Native villages of Nanwalek (Nw), Port Graham (Pg) and Seldovia (Sv).



Figure 29. An unusual muscular lesion in a sockeye salmon.

Musculature lesions

The photographic image in Figure 29 illustrates a gross lesion in the flesh of a single male sockeye from Nanwalek. The lesion exhibited significant bleeding. The appearance of this lesion did not show any evidence of bacterial or other infections or any sign of disease, and there was no significant inflammation. It was concluded that this lesion was the result of capture-related trauma.

5. Discussion

Contaminant body burden

Trace and major elements are present everywhere because they are naturally occurring elements derived from surface soil and rock. Elevated concentrations may be the result of natural weathering of mineral-rich rock, volcanic eruptions, or man-made sources, such as industrial activity or mining. Many metals are essential nutrients at low levels, even some metals that are considered to be toxic at higher exposures. Organisms will absorb metallic elements and attain equilibrium concentrations in their tissues in proportion to their exposure and their release processes. Exposure may be by direct contact with sediment or water, ingestion of sediment, or via the food chain. The metals concentrations on the Kenai peninsula are reflective of metals eroded from rocks that have been subjected to a great variety of weathering forces including volcanic disturbances. Metals have also been deposited here after long range atmospheric transport from areas in lower latitudes. It is a highly variable environment in spatial terms, and local conditions may vary even between adjacent bays. Thus, the local geology influences the exposure of resident organisms in different locations to these metals,

and therefore the concentrations of these metals in their bodies. Variation in local sediment contaminant concentrations have been documented in Port Graham (Hartwell et al., 2009). In a cove near the village, sediment chromium and mercury concentrations were 50.9 and 0.353 ug.g^{-1} , respectively. In the adjacent cove at the head of the bay only 3 km away, the concentrations were 334 and 0.143 ug.g^{-1} , respectively.

The iron and zinc concentrations in clams from Seldovia are twice as high as in Port Graham, which in turn are two to three times as high as neighboring Nanwalek (Table 5). Iron and zinc are essentially non-toxic to humans so there is no issue with consumption, but it illustrates the importance of local geology. The pattern of elevated metals at Seldovia relative to the other locations is seen with virtually all of the trace metals. Port Graham concentrations exceed those of Nanwalek in half of the elements. Given the range of tissue concentrations in studies at wider spatial scales (NOAA NS&T and AK DEC FMP), the values on the Kenai are not greatly different from regional observations. The important point is that for those elements warranting FDA action levels (arsenic, cadmium, chromium, lead, nickel), the observed tissue concentrations are far below FDA thresholds (Table 5), some by more than an order of magnitude.

Chromium and coal mining activity has occurred historically on the Kenai Peninsula and the Kachemak Bay region. Chromite mines were located at Red Mountain in the interior of the peninsula, and at Chrome Bay at the lower tip of the Kenai. Ore from Red Mountain was transported to Kasitsna Bay where it was loaded onto transport vessels for shipping. Coal mining has been carried out at various times throughout Kachemak Bay. Coal mining typically exposes sulfide-bearing rock and releases ground water. Upon exposure to the atmosphere, the sulfides are oxidized into sulfuric acid, releasing metals, and the acid leaches more metals from the rock and soil through which it flows. Upon entry into marine water, the acid is neutralized, but many of the metals accumulate in local sediment deposits. These sediments can become a source of heavy metal contamination in resident organisms, and may be dispersed by tidal currents to settle elsewhere, or be diluted to insignificant concentrations. The data presented in this report does not indicate any accumulation of chromium, or other metal-laden sediments, in any of the three study areas, based on the observed tissue concentrations in the mollusks. Concentrations are within the ranges seen in other bivalves in Kachemak Bay, and the lower Cook Inlet in other monitoring data sets.

Mobile animals (such as salmon) which spend significant periods of their lives in other habitats will reflect the chemical makeup of those other habitats, influenced by their exposure to local conditions, only as they move

through them. It is well documented that Pacific salmon stop feeding on their spawning migration from the open ocean into coastal estuaries and rivers. None of the fish that were examined in this study had anything in their stomachs. Thus, their body contaminant concentrations are primarily a result of their diet in the open ocean. Within the range of overall variation in muscle tissue, there is very little difference in individual metal concentrations between locations or species. Muscle tissue levels are below calculated chronic no adverse effects levels for cadmium and selenium. Mercury levels are on the same order of magnitude, but still below the EPA chronic level. The reference dose for mercury is an order of magnitude below the other metals. Again, the standard is for methylmercury, and the measured value is total mercury, potentially providing an additional degree of safety relative to the standard. There is a large difference between muscle and liver tissue. Liver concentrations are considerably higher than muscle in all cases except arsenic and mercury. Unlike mammals, fish do not have the metabolic enzymes to regulate many metals levels. Consequently they accumulate in the liver and cannot be expelled.

There are no consumption standards for liver tissue, and the liver constitutes a much smaller proportion of the fish's mass compared to the muscle. Mercury does not tend to accumulate in the liver. Most mercury in tissues is converted into methylmercury and remains in the tissues. All of the organic contaminants assessed in this study are synthetic chlorinated compounds, except the PAHs. They are of interest due to their persistence, toxicity and tendency to accumulate in animals. PAHs may originate from natural seeps or human activities, while most of the compounds are synthetic chemicals banned or severely restricted in the U.S. With a few exceptions, the pesticides have likely never been used in the vicinity of the study area. Their presence indicates contamination from outside the region. The PCBs were used in a variety of industrial applications and may have local sources from previous uses and old machinery. The original mixtures, called Aroclors, contained specific mixtures of PCBs. Each Aroclor mixture had different uses. Mixtures of PCBs released to the environment will proceed through several transformations and behave differently in the environment. Thus, they will accumulate more or less strongly at different sediment depths due to things such as sediment size. Aroclors degrade and become structurally different over time but are still toxic. So this is why fish consumption reference doses based on analysis of Aroclors alone are not entirely reliable for environmental samples.

Some compounds degrade into distinct by-products that can aid in assessing the relative proximity to sources. For example, the banned pesticide DDT breaks down into DDE and DDD. The higher the relative proportion

of DDT in the mixture, the fresher the source. DDT is still used in Asia for example, which is up wind from Alaska. DDT was frequently below detection limits in our samples, and DDE and DDD were found at higher concentrations than DDT, indicating old sources. Tri-butyl tin, formerly used in anti-fouling paint, also goes through a degradation sequence of decreasing butyl content, which can be used to infer age of release. None of the concentrations were above detection limits in the mollusks.

PAHs are derived from natural and man-made sources. Natural sources include coal, decaying vegetation, and natural oil seeps. Man-made sources are spilled fuel and oil, and burning organic material, including fuel, wood or plastics. All of these substances are transported long distances by the atmosphere and on ocean currents. Different classes of organisms have differing abilities to break down PAHs. Vertebrates can break down or neutralize them and excrete them. It is the break down of certain compounds (e.g. benzo[a]pyrene) that generates carcinogenic by-products which renders them dangerous to health. Bivalve mollusks cannot effectively break down PAHs, which makes them good indicators of local PAH contamination.

Usually, high proportions of “low molecular weight” PAHs are associated with oil and petroleum releases (petrogenic source). A high proportion of “high weight” PAHs is often linked to combustion by-products and/or long-term weathering. With the exception of the PAH naphthalene, the low weight PAHs were found at concentrations comparable to the higher weight PAHs (Figure 30). The make up of the PAHs indicates a mixture of sources from spilled fuel and atmospheric drift of exhaust fumes from a variety of sources. Naphthalene itself is derived from a variety of sources, chiefly combustion and off-gassing from natural hydrocarbon sources and fuel, but is a commonly used chemical and is also emitted from a variety of substances, from building materials, to tobacco smoke. It is the primary ingredient in moth balls. One of the largest components of the suite of PAHs was perylene. This is a harmless natural by-product of the breakdown of terrestrial plant material (NRC 1985). This indicates that naturally occurring PAHs are as large or larger contributor to concentrations in clams than any man-made source. Considering that, and the overall very low concentrations, the contribution of man-made PAHs in the harvest areas appears to be extremely limited. This observation holds for PAHs concentrations in other shellfish such as blue mussels from locations in the Gulf of Alaska (Figure 31) and the Shelikof Strait (Figure 32). In mussels samples collected in 2007 by the National Park Service, Southeast Network as part of their coastal water monitoring program, concentrations of low weight PAHs (naphthalene and benzothiophene) were higher than those of high weight PAHs (chrysene and

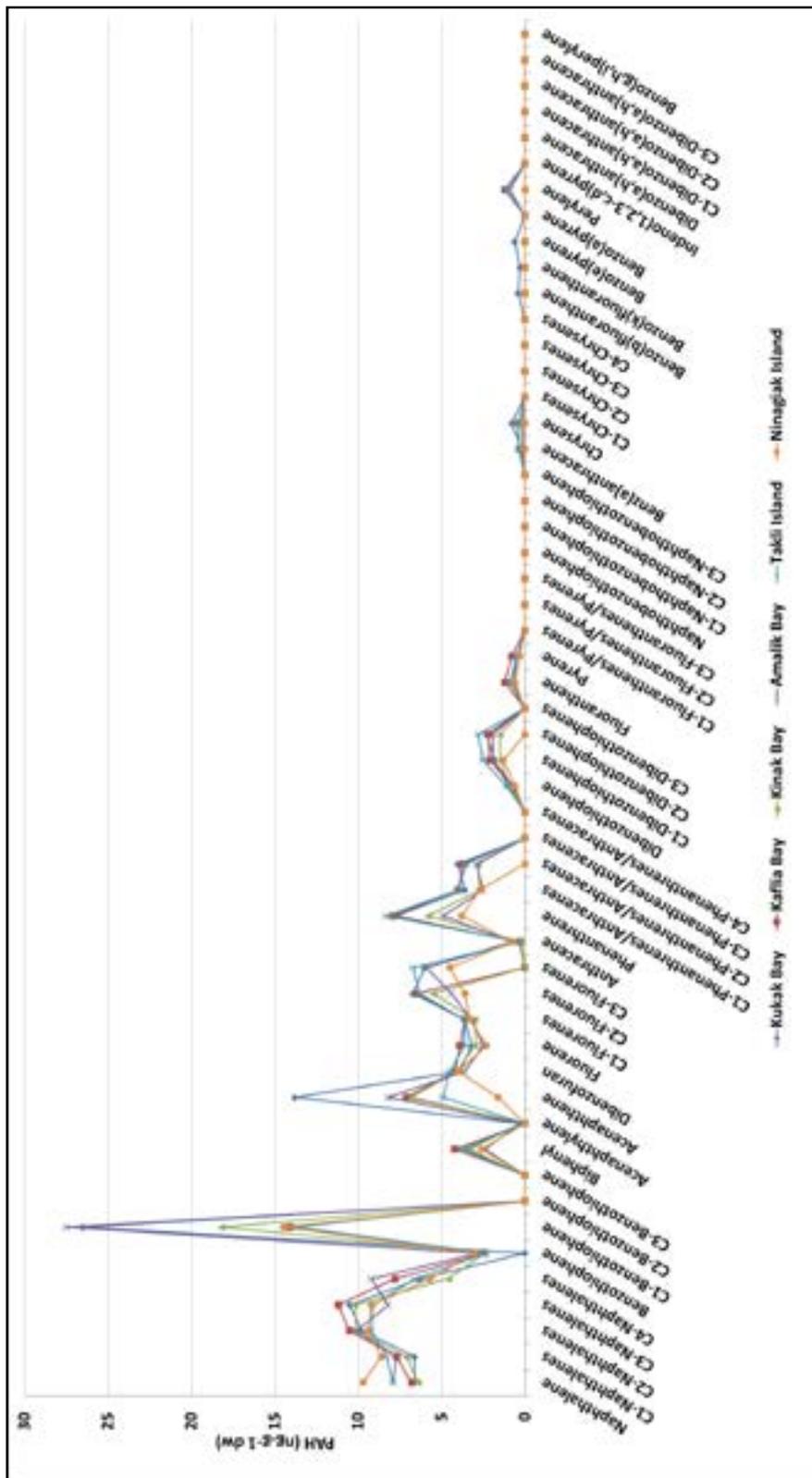


Figure 31. Individual PAH concentrations in blue mussels collected in 2007 from diverse NPS, Southeast Network monitoring locations along the northern shoreline of the Gulf of Alaska location. The PAHs are arranged from low molecular weight compounds (left) to high molecular compounds (right).

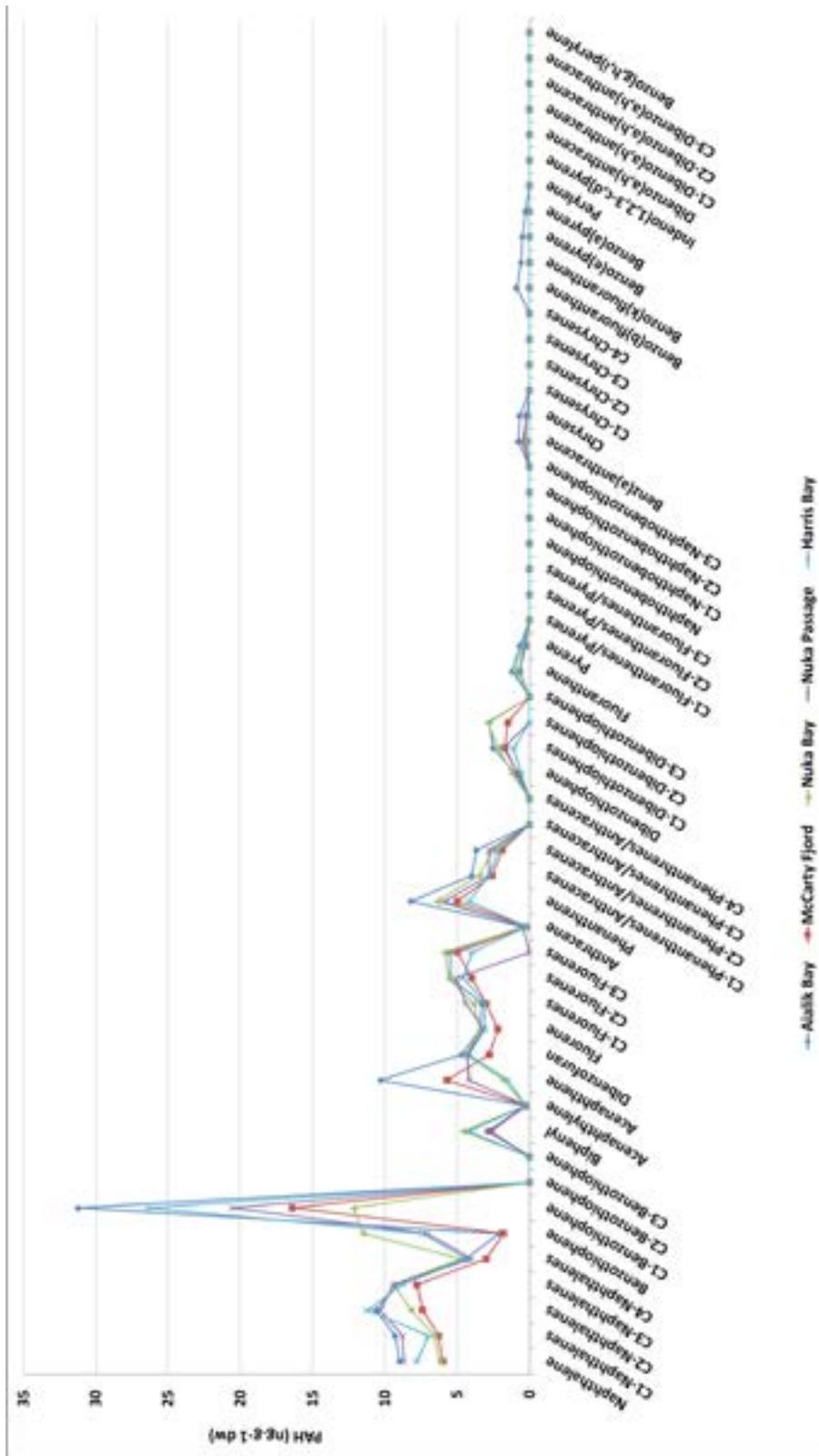


Figure 32. Individual PAH concentrations in blue mussels collected in 2007 from diverse NPS, Southeast Network monitoring locations along the western shoreline of the Shelikof Strait. The PAHs are arranged from low molecular weight compounds (left) to high molecular compounds (right).

chain. The chemistry data suggests there are between-species and between-site differences. The levels of DDT and PCBs in salmon livers are higher in the Nanwalek and Seldovia fish, relative to fish of the same species from Port Graham. This is not seen in the muscle tissue. There is no obvious reason for this and may simply be a consequence of small sample sizes and low absolute concentrations. Also, muscle tissue levels in the sockeye salmon were consistently higher than in the chum salmon. This is likely due to different feeding habits of the two species (Davis et al., (2005).

Histopathology

Every organism living today has some level of parasitic infection. In normal healthy organisms, this is of little consequence unless the parasites grow or proliferate to levels capable of producing significant pathology or disease (pathogenic) in the host. Among the parasites analyzed in the shellfish, only microcrustacean copepods, gregarines (nematopsis) and roundworms (nematodes) were detected. Although there are no human health concerns with the occurrence of parasitic copepods in shellfish, many of these parasitic microcrustaceans can have potential for ecological effects, such as the growth, fertility, and survival of their hosts (Johnson et al., 2004). Roundworm infections in mollusks can cause destruction of adjacent tissues and other problems (Kim et al. 2006). Additionally, many roundworms infect humans, with the most common ones including hookworms (ascaris) and trichina worms, which can cause trichinosis if raw or undercooked meat is ingested. Thus, roundworm infections of the shellfish could be harmful to the shellfish, but also to human consuming these resources. Gregarines were the other parasites detected in the shellfish. Heavy infections of gregarines has been suggested to have some harmful effects on the physiology of infested shellfish (Sindermann, 1990) however, Cheng (1967) concluded that, in general gregarine infections have “low pathogenicity in bivalve mollusks.” Although some of the parasites detected in the shellfish can cause ecological health and human health effects, the measurements indicated that the infections of virtually all parasites were relatively minor and may not have any significant health impacts on the shellfish or people.

The presence of diseases and other histopathologic conditions in the shellfish analyzed were limited to abnormal tissue inflammations, tissue death, digestive tube wastin and harmless tumors (Table 8). The conditions of digestive tubewasting detected in the shellfish are generally characterized by the thinning of the digestive tube walls and the conditions have been linked to a variety of stressors including exposure to contaminants and poor nutrition (Kim et al., 2006). Although digestive tube wasting is not necessarily

a pathologic condition, it can impact food uptake and potentially growth (Kim et al., 2006). Tissue inflammations, which were observed in some of the shellfish, are usually the result of intense infiltration of blood parasites. Tissue inflammation in shellfish can occur as wide-spread (diffuse) or very local inflammation. Diffuse inflammation is differentiated from local inflammation when the affected area does not appear to have a clear center or focal point of concentration and hemocytes are abundant and distributed broadly over a large section of tissue (Kim et al., 2006). Although tissue inflammation was not detected in shellfish from Nanwalek and Port Graham, the condition was very prominent in softshell clams from Seldovia (Figure 22). Although necrosis (tissue death) was seen at all locations, measurement of intensity indicated that occurrence of the condition was not pronounced (figure 23). In bivalve mollusks, most cases of tissue necrosis were observed in the connective tissue and is sometimes associated with the presence of parasites (Kim et al., 2006). Cases of tissue neoplasm conditions were observed in both clams and cockles collected from the Seldovia harvest ground (Figures 22 and 23). In bivalve mollusks, like clams and cockles, neoplastic sarcomas usually occur in vesicular connective tissues and could be harmful to the overall health of the bivalves.

The gender difference of the roundworm infections in fish did not appear to be universal since, in Seldovia, the parasites were detected in liver tissue of male sockeye, but not in liver tissue of female sockeye (Figure 24). Additionally, the Fisher's Exact Test results indicated that prevalence values were not significantly different between the genders of the fish ($p < 0.05$). Roundworm infections at high intensity in fish can impact fish reproduction, although in the majority of cases only larval stages are present in marine fish (Cheng, 1978). The final hosts for the parasitic roundworms are fish eating birds or mammals including humans. Certain roundworm varieties found in fish can infect humans and sometimes cause damage to stomach and intestinal tissue in human (Darwin and Stefanich, 1966). In this study, the vast majority of liver tissues in both species were normal; in fact, in chum salmon, all livers were normal. Thus the presence of the parasitic roundworms in the fish was not likely to be pathogenic.

Myxosporidia, a class of microscopic once celled parasites were also detected in the fish tissue (Figure 25). Myxosporidians are characterized by the presence of complex spores and having an infective life stage (Noble, 1944). Although they are primarily parasites of fish, some species of myxosporidians also infect amphibians and reptiles (Noble 1944). According to Jirk et al., (2006), most Myxidium usually infect the gallbladder,

urinary bladder, or urinary tubes in the kidneys of fish hosts. The myxosporidians, detected in the kidney tubes of the adult chum and sockeye salmon were likely to be of the *Myxidium* genus. Overall prevalences of myxosporidian infection in the kidney tubules of salmon showed variable infection frequencies (Figure 25), with sockeye salmon from Nanwalek being significantly more infested than either the Port Graham or Seldovia harvest grounds (Fisher's Exact Test, $p < 0.05$). *Myxidium* infection can be debilitating or even deadly to the fish (Alvarez-Pellitero and Sitja-Bobadilla, 1993). However, in this study, all infections were minor in severity and did not involve significant pathological change to the affected tubules or nephrons. Consequently, the higher prevalence of this condition at Nanwalek is unlikely to have significant physiological impacts on the affected fish.

The spore-forming microsporidian parasitic infections (*Loma salmonis*) can be pathogenic. They commonly infect fish gills and can cause serious xenomas (extremely large lesions in infected cells) in salmon species because they undergo division (sporogony) in host cells (Higgins et al., 1998). In this study, the *Loma salmonis* infections were found at relatively low prevalences in salmon of both species and sexes collected from Port Graham and Nanwalek's traditional harvest grounds (Figure 26). No infected sockeye were found at Nanwalek. Prevalence of the *Loma salmonis* infection was higher in male than female chum from Nanwalek, but this 100% prevalence was in a sample size of only two male fish. However, the statistical assessment indicated no significance difference in prevalences of infection among fish from different harvest grounds (Fisher's Exact Test, $p < 0.05$). Although infection in fish by *Loma salmonis* can be severely pathogenic, (Higgins et al., 1998), in the present study, infections were generally low (Figure 26), and the infections represented by the typically small xenomas were all minor in severity and nonpathogenic.

The monogenetic trematodes (roundworms) found in salmon from this study are likely of the family of Gyrodactylidae. Trematodes mainly infect mollusks, like clams and cockles, as a first host, but intermediate and final hosts may include animal ranging from invertebrates to mammals and even humans (Kumar, 1999). There are two types of parasitic trematodes; digenetic trematodes are internal parasites of mammals and humans, while monogenetic trematodes are external parasites in fish, mollusks and reptiles (Kumar, 1999; Darwin and Stefanich, 1966). Trematodes detected in the chum and sockeye salmon analyzed in this study were low in prevalence at nearly all three traditional harvest grounds (Figure 27). The infections

that were present did not appear to induce any significant host response or result in any significant lesions (nonpathogenic).

Microaneurysms are small distentions or swellings usually found in blood vessels, such as the capillaries of the gill as reported in this study. These lesions are quite commonly observed in histological preparations of fish gills sampled from the wild (Landolt and Busch, 1991). In fish, microaneurysm lesions are frequently associated with a variety of infectious diseases (Landolt and Busch, 1991). In this study, the microaneurysm lesions in the gill were observed only in chum salmon (Figure 28). The lesions were minor in severity and were probably an artefactual result of sampling trauma such as capture by gillnet.

A gross lesion in the muscular tissue (Figure 29) was observed in a single male sockeye salmon from Nanwalek. However, careful histopathologic assessment indicated that there was no significant inflammation associated with this lesion, and there was no evidence of bacterial or other infections or any systemic disease. It was concluded that the lesion was most probably caused by gill net capture and possibly trauma resulting from extraction from the gill net.

Histopathological conditions were characterized in the softshell clams and cockles, as well as in chum and sockeye salmon used for subsistence food in the Chugach communities of Nanwalek, Port Graham and Seldovia. Among the histologic parameters measured, only parasitic infections/infestations were detected with consistency in both the fish and shellfish specimens. Occurrences of noninfectious histologic conditions or diseases were limited and found mainly in the two shellfish species. In general, parasitic infections in the fish and shellfish were relatively few in type and minor in severity, resulting in a very low parasitic impact. Many parasitic infections are often associated with actual disease (pathogenic); however, in this study parasitic taxon richness and intensity/severity of infection were low and not adequate for assessing the parasite-disease linkages. The results indicated that parasitic infections and the rare noninfectious histologic conditions in the subsistence salmon and shellfish species were nonpathogenic, and no toxicopathic lesions (those likely to possess an etiology related to toxic chemical exposure) were detected in salmon. We conclude that none of the infections or noninfectious histologic conditions constitute a health hazard for the fish or shellfish analyzed, or to humans.

6. CONCLUSIONS AND RECOMMENDATIONS

Across the three sampling sites, the fish and shellfish sampled showed low tissue contamination. Pathological (disease) effects in shellfish and fish tissues for the parasites and diseases measured were absent or minimal. Taken together, our results showed that they were healthy and non-contaminated. These findings do not preclude the possibility of other factors impacting these coastal resources in the region. The mere presence of the synthetic contaminants at detectable levels in the tissues suggested some minimal exposure from remote sources.

Contaminant body data and information about histopathology (tissue disease) characterization in coastal and marine animals are important for resource managers. Chemistry and histopathology data from this study represent useful information for concerned native community members and coastal resource managers in Alaska. The data from this study were georeferenced and incorporated into the NS&T data portal and are available to the public.

Fish and shellfish have high nutritional value as they are excellent sources of essential protein, antioxidants, fatty acids (lipid), and vitamins. Of a particular importance for human health are omega-3 fatty acids, which provide many health benefits including protection from diabetes and cardiovascular disease. Omega-3 lipids also help improve maternal nutrition and neonatal/infant brain development. With low contamination and presence of few to no lesions (especially in salmon), this assessment indicated that the clams, cockle and salmon from the traditional harvest grounds in Nanwalek, Port Graham and Seldovia are safe for consumption by Native communities. However, we recommend the following:

- *Because fish and shellfish harbor potentially harmful pathogenic parasites, it should be a good practice to always keep harvests frozen until ready to be processed or cooked.*
- *To avoid the possibility of migration of intestinal worms into the edible parts, thoroughly clean the fish and shellfish as soon after catching as possible.*
- *During cleaning and processing of fish in particular, if lesions or parasites are observed, it is recommended to always remove the entire organ where the parasites were found (Darwing and Stefanich, 1966). Most parasitic worms die when heated. It recommended to refrain from consuming raw seafood of any kind*

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