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- 1 How significant is atmospheric metal contamination from mining activity adjacent to
- 2 the Tasmanian Wilderness World Heritage Area? A spatial analysis of metal
- 3 concentrations using air trajectories models
- 4
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23 Abstract:

This study investigated metal contamination from historical mining in lakes in the Tasmanian 24 Wilderness World Heritage Area (TWWHA) and surrounding region. The largest increase in 25 26 sedimentation and metal contamination occurred ca. 1930 when open-cut mining commenced and new mining technology was introduced into the region. The geochemical signal of lake 27 sediments changed from reflecting the underlying geology and lithology to that reflecting 28 mining activities. The HYSPLIT air particle trajectory model explains metal distribution in 29 the lakes, with those in the northwest region closest to the mines having the highest metal 30 31 contamination. Lake metal concentrations since mining activities commenced are in the order: Owen Tarn > Basin Lake > Perched Lake > Lake Dove > Lake Dobson > Lake 32 Cygnus, with Perched Lake and Lakes Dove, Dobson and Cygnus in the TWWHA. Metal 33 34 contamination affected sites up to 130 km down-wind of mining sites. Enrichment factors (EF) for Pb, Cu, As and Cd are > 1 for all lakes, with Owen Tarn and Basin Lake having very 35 high EFs for Cu and Pb (98 and 91, respectively). Pb, Cu, As and Cd concentrations are 36 above the Australia/New Zealand lower sediment guidelines, with Pb, Cu and As above the 37 upper guidelines in Owen Tarn and Basin Lake. This study demonstrated the legacy of metal 38 39 contamination in the TWWHA by mining activities and the consequences of a lack of execution of environmental regulations by past governments in Tasmania. 40

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47 **1. Introduction**

48 Mining has been a key factor in the economic development of Tasmania, Australia, with numerous abandoned mine sites that are still contaminating soils, rivers, lakes and estuaries 49 (Figure 1; Augustinus et al., 2010). This is of environmental concern in a wide geographical 50 area beyond the immediate vicinity mine sites, as particulate emissions released to the 51 atmosphere by mining operations can be transported over long distances by atmospheric 52 circulation (Suvarapu and Baek, 2017). Here, we use computer modelling of air particle 53 54 trajectories and lake sediment contamination measurements to develop and test a model of airborne contamination transport from historic mining activities in western Tasmania. 55

The west coast of Tasmania is characterised by folded and faulted geology containing several ore bodies that were exploited when the British arrived in Australia in the late 1700s. Principal among these are the major mining centres developed around the closely spaced Mt Lyell and Mt Read ore deposits, in Queenstown and Rosebery, respectively.

60 Analyses of sediment and water from Macquarie Harbour, downstream from the Queenstown 61 region, indicates a dramatic increase in metal and metalloid concentrations (hereafter collectively referred as metals) in the harbour resulting from contamination of the Queen and 62 King Rivers by the Mt Lyell mine (Augustinus et al., 2010; Carpenter et al., 1991; Eriksen et 63 al., 2001; Stauber et al., 2000; Teasdale et al., 2003). Further afield, an increase in metals in 64 isolated catchments downwind from both Queenstown and Rosebery reveal the same trends 65 66 in metal contaminants through the period of intensive mining and smelting operations, suggesting transportation of metal contaminants by wind from mining centres (Harle et al., 67 2002). 68

While the effects of mining on the environment around the Queenstown-Rosebery region are relatively well recognised, e.g. localised deforestation and downstream impacts on aquatic ecosystems (De Blas, 1994; Harle et al., 2002; Hodgson et al., 2000; Kozlov and Zvereva, 2006), there has been no attempt to understand the spatial distribution of airborne metal contaminants from Queenstown and Rosebery. This is important because the western boundary of the Tasmanian Wilderness World Heritage Area (TWWHA) lies just 11 and 12 km from both Queenstown and Rosebery, respectively, in the prevailing wind direction.

The atmospheric distribution of metal contaminants involves a complex interplay of environmental factors, climate and local meteorological characteristics. The principal environmental factors affecting atmospheric transport of metals include, but are not limited to, precipitation, temperature, air movement and pressure (Fang et al., 2005; Pacyna et al., 2009; Suvarapu and Baek, 2017). All of these must be taken into consideration when assessing atmospheric metal distribution and deposition into the environment.

A useful tool to understand the interplay of climate factors on metal distribution is The 82 Hybrid Single Particle Lagrangian Integrated Trajectory Model (HYSPLIT) model (NOAA, 83 2018). HYSPLIT produces forward trajectories that, when combined with satellite images 84 85 (from NASA's MODIS satellites), can calculate air particle trajectories over a set period of time and, thus, the direction atmospheric contamination has travelled (Kusumaningtyas and 86 Aldrian, 2016). Despite its apparent usefulness and value, this model has never been applied 87 88 to assess airborne contamination from historical mining sites in Tasmania to understand the potential effects of airborne contamination on the environment. 89

In this study, we assess the extent of metal contamination in the TWWHA and surrounding
areas using sediment cores from six freshwater lakes. In particular, we applied the HYSPLIT
model and statistical analyses to establish the main chemical and physical factors affecting

93 the airborne distribution of metals in these lakes. Furthermore, we compared lake sediment 94 metal concentrations with the Australia/New Zealand (ANZECC/ARMCANZ 2000) 95 sediment guidelines to assess the past and current health of the local environments. 96 Ultimately, the study was undertaken to inform the scientific community and the public about 97 the legacy of metal contamination within the TWWHA to support government initiatives in 98 establishing appropriate regulations and policies to protect the environmental values of this 99 wilderness area.

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101 **2.** Material and Methods

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103 **2.1.Regional description**

Western Tasmania is a mountainous area underlain predominantly by intensely folded and 104 faulted Cambrian and pre-Cambrian quartzite rocks and conglomerate units that are 105 intersected with highly mineralised volcanic belts (Corbett and Solomon, 1989). The area 106 includes more than 4,000 lakes and tarns, mostly of glacial origin, ranging from highly acid, 107 dystrophic lakes to ultra-oligotrophic clear water lakes (Hodgson et al., 2000). The west coast 108 receives high orographic rainfall produced by air masses rising over mountains (Gentilli et 109 al., 1972; Sturman and Tapper, 2006). The rainfall reaches a maximum of 3,400 mm and 110 there is an annual temperature range of 3 - 21 °C, with a mean annual temperature of 11 °C 111 at sea level, and 6 °C at 1000 m altitude (Langford, 1965). The climate is dominated by the 112 prevailing zonal westerlies that latitudinally migrate through the seasonal cycle, with west to 113 south-westerly airflow dominant in the austral winter and west to north westerly airflow 114 dominant in the austral summer. 115

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117 **2.2.Historical setting**

Tasmania has been occupied by humans for ca. 40,000 years (Cosgrove, 1999), with Tasmanian Aborigines responsible for maintaining an essentially open landscape through the use of fire (Fletcher and Thomas, 2010). Despite the arrival of the British in Tasmania in the late 1700s, it was not until the late 19th century with the arrival of mineral prospectors that western Tasmania was exposed to exploitation for deposits of gold, silver, lead, zinc and copper. Subsequently, major mining and smelting operations were established and concentrated at several centres around Queenstown and Rosebery (Figure 1).

Discharge of tailings, slag, toxic metals and acid drainage into the Queen River that runs through Queenstown, and downstream to the King River and ultimately into Macquarie Harbour, has eliminated all but the most robust forms of aquatic life in these waterways (Hodgson et al, 2000). Today, these towns are located along the boundary of the TWWHA, thus, there is a high probability that areas within the TWWHA have experienced some degree of long-range metal contamination and ecological change from historic mining activities (Harle et al., 2002; Hodgson et al., 2000).

Mineral exploration commenced in the 1880s however, it was not until the end of 1920s, with the advent of automation and changed work practices, that mining activities expanded from underground to open-cut. This was attributed to favourable copper prices and advances in transport (Rae, 1994).

The mining boom in Queenstown-Rosebery saw a downturn in the 1980s due to the low price of copper and activities were reduced to two mining companies: Copper Mines of Tasmania (CMT) in Queenstown which has now been active for 100 years, and MMG Rosebery, active in Rosebery since 1936.

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141 **2.3.** Site selection and sediment core collection

Sites were chosen to provide an adequate spatial coverage to characterize the aerial transport 142 of metals from mining sites and to document spatial differences in the deposition of metals 143 within the TWWHA. Given the aim of assessing atmospheric transport of particles and metal 144 deposition in lakes, we targeted lakes with small catchments to avoid major geochemical 145 influence from the drainage basins . A total of six lakes were identified as suitable for these 146 analyses, and a total of six cores (one per lake) were collected from the deepest point of the 147 lakes. A 25 m-resolution digital elevation model (DEM) was used to analyse lake catchment 148 morphologies from where the catchment area for each lake was derived using the suite of 149 Hydrology Tools (Arc Hydro) in ArcGIS 10.3 (ESRI, 2015). This approach allowed us to 150 map flow direction and stream paths based on aspect and slope for each cell of the DEM. The 151 catchment boundaries were delineated, and the surface was calculated using the same 152 program. 153

154 Sediment collection was conducted in two periods:

Collection 1: Sediment cores from lakes in the TWWHA (Dove Lake, Lake Cygnus, Lake
Dobson and Perched Lake) were collected in 2000 using a gravity corer and hammer driven
piston corer (Neale and Walker, 1996).

158 Collection 2: Sediment cores from lakes bordering the TWWHA (Lake Basin and Owen159 Tarn) were collected in 2011 and 2015, respectively, using a Universal Corer.

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161 **2.4. Sediment dating**

Lead-210 (²¹⁰Pb) samples were processed at the Australian Nuclear Science and Technology
Organisation (ANSTO) using alpha spectrometry and following methods described by

164 Harrison et al. (2003). Each dried sediment sample (2 g) was spiked with Polonium-209 (²⁰⁹Po) and Barium-133 (¹³³Ba) tracers. Each sample was then leached with hot nitric and 165 hydrochloric acids to release polonium and radium. Polonium was autoplated onto silver 166 disks after adding the reducing agent hydroxylammonium chloride. Radium and barium were 167 isolated by co-precipitation and collected as colloidal micro-precipitates of barium sulphate 168 on fine membrane filter papers. The activities of ²¹⁰Po on the silver disks and ²²⁶Ra on the 169 membrane filters were determined by alpha spectrometry. Each membrane filter was also 170 counted by gamma spectrometry to measure the ¹³³Ba tracer activity. Chemical yield 171 recoveries of ²¹⁰Po and ²²⁶Ra were calculated using the recoveries of ²⁰⁹Po and ¹³³Ba tracers, 172 respectively. Unsupported ²¹⁰Pb activity for each sample was calculated from the activity of 173 ²¹⁰Po (the proxy for total ²¹⁰Pb) minus the ²²⁶Ra activity (the proxy for supported ²¹⁰Pb). 174

The ²¹⁰Pb dating models Constant Initial Concentration (CIC) (Pennington et al., 1976; Robbins and Edgington, 1975) and Constant Rate of Supply (CRS) (Appleby and Oldfield, 1978) were used to determine sediment ages and mass accumulation rates for sediment cores with dry bulk density data available. A modified CIC ²¹⁰Pb dating model as described by Brugan (1978) was used to determine CIC ages and sedimentation rates for those sediment cores where dry bulk density data were not available.

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182 **2.5.Geochemical analyses**

All samples were transported from the field to the laboratory at ANSTO or at the University of Melbourne and kept stored at 4° C in a cool room. Samples were manually mixed and transferred to a clean glass vial, covered with parafilm and placed in a FreeZone Plus 6 freeze-drier (Labconco, Kansas City, MO) and lyophilized at -50 °C for 48 hours. Given the substantial gap between the analysis of samples from collection 1 and 2 (ca. 15 years), the methodologies differ slightly due to the technology and procedures in use at the time. Similar results for certified reference materials indicate the applicability and comparability of both procedures.

Samples from collection 1 (Lake Dove, Perched Lake, Lake Dobson and Lake Cygnus): 191 Approximately 0.5 g of dried sediment was weighed into a tetrafluormethaxil (TFM) closed 192 digestion vessel (Ethos Milestone) and 3 mL sub-boiled nitric acid, 1 mL of sub-boiled 193 hydrochloric acid, 0.1 mL of 50% w/v high purity hydrofluoric acid (Merck, Suprapur) and 3 194 195 mL of deonised water added. Each vessel was capped and placed in a Milestone MLS 1200 Mega microwave cavity, heated to 180°C for 25 mins, and then held at 180°C for 15 mins 196 before being cooled to room temperature and diluted with 30 mL of deonised water. One mL 197 198 of the digest was transferred to an 8 mL centrifuge tube and 4 mL of ICP-MS internal standard added (⁶Li, ⁴⁵Sc, ⁸⁹Y, ¹⁰³Rh, ¹¹⁵In, ¹⁸⁵Re, and ²⁰⁹Bi). 199

Metal concentrations in sediments were measured by inductively coupled plasma mass 200 spectrometer (ICP-MS) and inductively coupled plasma atomic emission spectrometer (ICP-201 AES). Mixed standard working solutions in the 500 to 0.001 µg/mL range and continuous 202 203 calibration verification solutions were measured at the same time as samples. Internal standard and suppression solutions (In, Rh, Rb) were prepared and added to the sample via 204 on-line addition. Certified reference materials, National Research Council of Canada (NRCC) 205 sediment SRMS MESS-3 and PACS-1 were also analysed and measured values were in 206 agreement with certified values (Supplementary Table 1). 207

Sediment samples from collection 2 (Owen Tarn and Basin Lake): approximately 1 g of sediment was weighed into a 60 mL polytetrafluoroacetate (PFA) closed digestion vessel (Mars Express), and 2 mL of concentrated nitric acid (Aristar, BDH, Australia) and 1 mL of 211 30% concentrated hydrochloric acid (Merck Suprapur, Germany) added (Telford et al., 2008). Each PFA vessel was then capped, placed into an 800W microwave oven (CEM 212 model MDS-81, Indian Trail, NC, USA), and samples heated at 120° C for 15 mins. The 213 digests were cooled to room temperature and diluted to 50 mL with deionised water 214 (Sartorius). The tubes were then centrifuged at 5000 rpm for 10 mins. One mL of the digest 215 was transferred into a 10-mL centrifuge tube and then diluted to 10 mL with ICP-MS internal 216 standard (Li⁶, Y¹⁹, Se⁴⁵, Rh¹⁰³, In¹¹⁶, Tb¹⁵⁹ and Ho¹⁶⁵). Digests were stored (0-5° C) until 217 analysis. Samples were analysed using an ICP-MS (PerkinElmer DRC-e) with an AS-90 218 autosampler (Maher et al., 2001). The certified reference NIST- 2710 Montana Soil was used 219 as controls to check the quality and traceability of metals. Measured concentrations were in 220 agreement with certified values (Supplementary Table 1). 221

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223 **2.6.**The Hybrid Single Particle Lagrangian Integrated Trajectory Model (HYSPLIT)

Wind trajectories from the sites were calculated using a map of the frequencies of HYSPLIT trajectories (Stein et al., 2015). A map with the average circulation of air masses over Tasmania during the period 1961-1990 for particles released at 42°S and 145.5°E (Queenstown, Tasmania) and 41.78°S and 145.5 °E (Rosebery, Tasmania) was created using ~1 million data points corresponding to the position of hourly-resolved HYSPLIT forwardtrajectories, overlayed with 10 x 10 km grid cells. The 1961-1990 period represents the full extent of the data available from the Australian Bureau of Meteorology.

Hourly-resolved meteorological data for calculating the HYSPLIT trajectories were derived from NOAA ARL NCEP/NCAR Reanalysis FTP (ftp://arlftp.arlhq.noaa.gov/pub/archives/reanalysis). The number of occurrences per grid cell was extracted in ArcMap 10.3 and relative frequencies calculated. Red indicates grid cells with a higher occurrence of air masses travelling from Queenstown/Rosebery. A directional
ellipse was derived using the 'Directional Distribution: Standard Deviational Ellipse'
function in ArcMap 10.3. This tool creates an elliptical polygon centred on the mean for all
features. The orientation of the ellipse indicates the average direction of flow during the
chosen time window and spatial scale. One standard deviation was chosen to cover
approximately 68% of all input feature centroids.

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242 **2.7. Enrichment factor (EF)**

The calculation of a normalized enrichment factor (EF) for metal concentrations above uncontaminated background levels enables an estimation of anthropogenic inputs of metals to sediments (Abrahim and Parker, 2007). The EF calculation seeks to reduce the variability of metal concentrations associated with fluctuations in clay/sand ratios and is a convenient tool for plotting geochemical trends across large geographic areas, which may have substantial variations in the sediment (i.e. clay rich) to sand ratios.

The EF method normalises the measured metal concentration with respect to a sample reference element such as iron (Fe) or aluminium (Al) (Cevik et al., 2009). In this approach the Fe or Al is considered to act as a "proxy" for the clay content. In this study, as Fe atmospheric deposition in lakes are known to have been altered by mining activities, we used Al as it was the element with least change through the profiles.

256 EF = (Mx/Alx)/(Mb/Alb)

The EF was calculated using the average contamination for the years comprising the peak in mining contamination (1930 to 1980), following the equation:

where Mx and Alx are the average metal and aluminium concentrations, respectively, for the mining period between 1930 to 1980. *Mb* and *Alb* are metal and aluminium background concentrations, respectively.

The lower metal concentrations in the bottom of the cores were interpreted as sediment deposited before the beginning of mining activities in 1880. From these results, natural background heavy metal values for the six lakes was proposed based on the average of premining trace element concentrations.

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265 **2.8. Statistical analyses**

All analyses were performed using the R Statistical Software (R Development Core Team, 267 2008) and the respective libraries used in particular analyses are cited.

268 To reveal differences in the metal concentrations among lakes and mining phases, we conducted a permutational multivariate analysis of variance (PERMANOVA) based on 269 Euclidean 270 distances (adonis, vegan package 2.5-1 https://cran.rproject.org/web/packages/vegan/index.html) using the function vegdist to find the 271 dissimilarities. Lakes and phases were included as fixed factors, and metal concentrations 272 were given as a matrix from where vegan calculated pairwise distance to find the 273 dissimilarities. 274

Principal Component Analysis (PCA) was used to explore the similarity of metal concentration in the lakes before and after mining activities in the region (dudi, ade4 package https://cran.r-project.org/web/packages/ade4/index.html). A multiple regression with backward-stepwise selection was performed to identify the main drivers of metal deposition in the sediments. Metal concentrations were log transformed to comply with the assumptions of linearity, normality and homoscedasticity. Before running the multiple regression,

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predictor correlations were checked to avoid problems for parameter estimation and potentially leading to the wrong identification of relevant predictors of the statistical model (Dorman et al., 2013). The predictors tested were: catchment size, precipitation, atmospheric temperature, distance from the mining site and frequency of particles passing through the lakes, calculated from wind directions and speed in HYSPLIT trajectories (section 2.5). If a correlation was > 0.7, then one of the predictors was removed.

287

288 **3. Results and Discussion**

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290 **3.1.** Sediment dating

²¹⁰Pb dating results are shown in Supplementary Table 2 and Supplementary Figure 1. The CIC and CRS ²¹⁰Pb dating model results were in close agreement in most lakes except for Lake Cygnus and Owen Tarn. The unsupported ²¹⁰Pb activities from these cores exhibited non-monotonic profiles, thus the use of the CRS model was more appropriate. (Supplementary Table 2).

The largest variations in sedimentation rates for sediment cores closer to the mines were recorded in the 1930s (Supplementary Table 3). At this time, the open-cut mine commenced in the region and new technology such as stamper and mills arrived in the region. This change in mining methods and technology increased sedimentation rates in these lakes as a result of increased atmospheric inputs. Details are discussed in a separate publication.

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302 3.2. Background metal concentrations and spatial distribution

Patterns of metal deposition in sediments have changed dramatically since the start of mining
activities (Table 1 and Supplementary Table 3). Sediment metal concentrations differed

significantly between lakes (PERMANOVA: *F* model = 104.1; r^2 = 0.729; *P* < 0.001; 999 permutations) and between mining periods (PERMANOVA: *F* model = 9.5; r^2 = 0.153; *P* < 0.001; 999 permutations). PCA of metal concentrations (Figure 2 A-B) illustrates the changes in metal concentrations (Axis 1) and their dramatic change since mining activities commenced.

Metal concentrations pre-mining were in the order of: Lake Dove > Perched Lake > Basin Lake > Lake Dobson > Owen Tarn > Lake Cygnus. Metal concentrations since mining activities commenced are in the order of: Owen Tarn > Basin Lake > Perched Lake> Lake Dove > Lake Dobson > Lake Cygnus (Table 1, Supplementary Table 2). These results demonstrate that mining activities have caused a shift in the geochemical signals of sediments in the lakes, from signals reflecting the specific geology and lithology to an association with mining activities.

The PCA performed using metal concentrations measured in sediments dated from before and after the 1880s clearly indicates that the most proximal sites to the mining centres (Owen Tarn and Basin Lake) have the highest metal concentrations since mining. Lake Dove and Perched Lakes, with the highest background concentrations, decreased in the rank of metal concentrations since mining activities commenced (Figure 2 A-B).

The majority of effort in determining the impact of mining contamination on aquatic 322 environments in western Tasmania has focussed on waterborne contamination down-stream 323 from mines (Augustinus et al., 2010; Carpenter et al., 1991; Dawson, 1996; De Blas, 199a; 324 Eriksen et al., 2001; McQuade et al., 1995; Stauber et al., 2000; Teasdale et al., 2003), with 325 airborne contamination receiving comparatively little attention (Harle et al., 200b). Our study 326 reveals that metal contamination can influence sites up to 130 km down-wind of mining sites, 327 with Lake Cygnus in the TWWHA displaying clear evidence of contamination. These results 328 329 indicate that most of the TWWHA area has potentially been impacted by airborne

contamination from the Queenstown-Rosebery mines (Figure 3). We thus urge a concerted
effort to understand the environmental and ecological consequences of this contamination in
the TWWHA.

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3.3. Drivers of metal spatial distribution

Predictors of metal atmospheric distribution in the TWWHA are given in Table 2 and Figures
3 and 4. Table 2 also summarises the main geographical and climatological information for
each lake that were considered to be the main factors influencing metal distribution in lakes
across the TWWHA.

Predictors (Table 2) were checked for between-predictors correlations to select the predictors to run the statistical model. The factors distance, precipitation and frequency had a correlation > 0.7 and were, therefore, removed from the model and the HYSPLIT-derived frequency of particles was used. This decision was based on knowledge that the HYSPLIT frequency model takes into consideration environmental variables and distance in its calculation. The final list of predictors for the model was therefore: catchment size, temperature and frequency of the particles.

The HYSPLIT frequency of particles model (Figure 5) successfully explained most of the metal atmospheric transport and metal deposition into the lakes (Table 3). The significance of the HYSPLIT model on metal distribution indicates that this model provides an effective predictive tool of the spread of airborne pollutants in the landscape. The decline in metal concentration over distance is indicative of atmospheric dispersion of the particles, resulting from mining activity.

The high precipitation rate within the TWWHA area suggests that wet deposition is an important factor in metal deposition into the environment. Although lakes with small catchment areas were only considered in this study, catchment size was a significant factor
only for the major elements Fe, Al and Zn. This indicates that metals deposited in these lakes
were mainly a result of atmospheric metal deposition rather than catchment leaching (Table
357 3).

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359 3.4. Enrichment Factors

To evaluate the extent of the historical metal contamination affecting the TWWHA lake sediments, EFs were calculated for the period 1930-1980, where an EF < 1 = no enrichment, EF 1- 3 = minor enrichment, EF 3 - 5 = moderate enrichment, EF 5 - 10 = moderately severe enrichment, EF 10 - 25 = severe enrichment, EF 25 - 50 = very severe enrichment, and EF > 50 = extremely severe enrichment (Cevik et al., 2009). All lakes had at least one of the metals with sediment concentrations showing moderate enrichment (EF > 3) since mining commenced (Table 4).

The EF values demonstrate significant metal contamination in the TWWHA. Mining contamination has reached distances as far as 130 km as demonstrated by the EF values > 1 for Lake Cygnus, the furthest lake from Queenstown and Rosebery in this study (Table 4, Figure 3). The effect of metal contamination distribution in the entire TWWHA and surrounding area can be visualised in Figure 6, which demonstrates the significant increase of metal inputs since mining started.

The EF values demonstrate that, from the metals measured in this study, As, Cd, Cu, Pb and Zn are the elements of most concern in the region. Owen Tarn and Basin Lake had the most significant metal enrichment in sediments. In Owen Tarn, specifically, Cu and Pb were 90 times higher relative to the background values (Table 4). This is of major concern as Pb and Cd bioaccumulate in the bodies of aquatic and soil organisms (Cresswell et al., 2015; Lanctôt et al., 2017; Storelli, 2008; Zheng et al., 2007). Even small concentrations of these metals can
affect body functions of aquatic organisms (Hodgson et al., 2000b).

The Pb and Cu EFs of 91 and 97.7, respectively, recorded in Owen Tarn are among the 380 highest reported in the scientific literature. These results are comparable to highly 381 contaminated places such as in the Kurang River in Pakistan, subjected to heavy metal 382 contamination from urbanisation and discharge of untreated domestic effluents (EF Pb= 4.46, 383 EF Cu = 12) (Zahra et al., 2014), the Shur River in Iran receiving inputs from copper mining 384 (EF Pb = 118.42, Cu = 264.1) (Karbassi et al., 2008), and the Lot River France receiving 385 inputs from mining and smelting activities since the late 19^{th} century (EF Pb = 10, EF Cu = 5) 386 (Audry et al., 2004). 387

EFs for As are of concern in Owen Tarn and Basin Lake. Cd in Owen Tarn and Basin Lake are also significantly higher, with Cd in Basin Lake yielding an increase of 25-fold from background concentrations (indicating severe enrichment) (Table 4).

Although Se concentrations increase in most lakes, only Owen Tarn has severe enrichment
while Perched Lake has moderate enrichment. Therefore, Se concentration increases in these
two lakes should be taken into consideration in further studies.

The extremely high enrichment of these elements in the TWWHA and surrounding area supports the need to investigate the effects of mining contamination in aquatic organisms in western Tasmania, given that 15,842 km² (one fifth of the island) is World Heritage Area.

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398 3.5. Comparison of metal concentrations and ANZECC/ARMCANZ (2000) sediment
399 quality guidelines

400 The Australian and New Zealand interim sediment quality guidelines (ISQGs)
401 (ANZECC/ARMCANZ, 2000) comprise two sediment guideline concentrations: (1) ISQG402 Low concentrations and (2) ISQG-High concentrations.

ISQG-Low concentrations are used as a threshold limit to appeal for checks on possible
adverse biological effects in aquatic organisms. The ISQG-High concentration is a threshold
limit above which adverse biological effects are expected to occur frequently in aquatic
organisms.

Table 5 shows the ratios of maximum concentration to sediment quality guideline values, being the concentration of a given metal in the sediment divided by that of the ISQG-low and ISQG high in the guidelines. The results show that Pb, Cu, As and Cd concentrations in all lakes are above the ANZECC/ARMCANZ ISQG-High threshold limit. Of concern are the Pb and As concentrations in Owen Tarn and Basin Lake, and Pb in Perched Lake sediments, which are above the ANZECC/ARMCANZ ISQG-High threshold limit.

In Australia, there are no selenium guidelines for sediments. The Screening QuickReference 413 Tables (SQuiRTs) developed by the National Oceanic and Atmospheric Administration 414 (Buchman, 2008) were, therefore, used to assess Se contamination in sediments. Although 415 SQuiRTs screening values are intended for preliminary screening purposes only, Owen Tarn 416 has shown a Se concentration 11 times higher than the SQuiRTs screening values, indicating 417 severe contamination and a likelihood of adverse biological affects in the area. The Se 418 concentrations in Owen Tarn (up to 16.8 mg/kg in the 1950s) is actually higher than 419 concentrations reported in Belews Lake, North Carolina, a lake heavily contaminated by Se 420 421 in wastewater released from a coal-fired electric generating facility during 1974-1985 (Lemly, 1997). In Belews Lake, Se concentrations of 4 to 12 mg/g in sediments were high 422

423 enough to cause severe reproductive failure and teratogenic deformities in fish. It is likely424 that Owen Tarn organisms might be been facing health issues due to Se contamination.

The EF and sediment guidelines indicate that the northwest side of the TWWHA has been severely contaminated (Table 4 and 5), and most likely have generated adverse biological effects in aquatic organisms. This is of great concern considering that contamination in organisms takes place through bioaccumulation from sediments to plants (Schneider et al., 2015) and its subsequent movement through trophic levels to animals and humans (Schneider et al., 2018). No study testing the health of aquatic organisms has been conducted in the area.

Studies in other areas of western Tasmania (De Blas, 1994b; Humphrey et al., 1997; Keele, 431 2003; Rae, 2005) have shown metal concentrations in food web organisms above guideline 432 limits proposed by the World Health Organisation (WHO, 1993). In Owen Tarn, a change in 433 434 diatom composition from oligotrophic to those more characteristic of dystrophic, acidic lake waters, and a decline in species richness occurred in response to mining activities (Hodgson 435 et al., 2000). It was also found that valve deformations in *Eunotia* species were a response to 436 chemical stress (Hodgson et al., 2000). A study of metal bioaccumulation and toxicity of 437 aquatic organisms within the TWWHA is highly recommended. 438

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440 **3.6-** Government Regulations and Inconsideration

This study demonstrates the atmospheric extent of deposition of metals in the TWWHA from past mining activities. Based on the ANZECC/ARMCANZ (2000) sediment quality guidelines, metal contamination is likely to be causing adverse health effects to aquatic organisms and humans feeding on them. During Tasmania's prosperous mining phase, mining companies were not subject to the same environmental regulations as the present day. Tasmania implemented and integrated environment protection legislation in 1973, when the *Environment Protection Act 1973* (comprising air, water and noise pollution, and waste management) was put in place. Even though Tasmania was one of the first states to have environment protection legislation in place in Australia (Bingham, 1992), mining companies were allowed to operate under exemptions which were granted by the government of the day. The argument supporting exemptions was that the cost of installing equipment to comply with emission standards would be such that the mine would have to close (De Blas, 1994).

Had regulations been strictly followed, it is possible the metal contamination in the TWWHA would be less severe and would have left a minor legacy of metal contamination. The high historical metal concentrations in lake sediments reported in this study leads to the question of how to and who should deal with the legacy of environmental problems arising from long running or discontinuing activities, which in earlier times had no environmental management protocols in place or lacked legal compliance to guidelines

459

460 **4.** Conclusions

This study demonstrates that historical metal concentrations in lake sediments can assist in interpreting the extent and severity of metal contamination in pristine areas. While independent studies and governmental reports have focused on the environmental effects of mining contamination in the King River and Macquarie Harbor, this study demonstrates that the atmospheric transport of metals has caused contamination to sites outside the mining catchment areas.

467 Atmospheric metal contaminates from mining activities in Queenstown-Rosebery in 468 Tasmania have contaminated most of the TWWHA area and have significantly altered the 469 natural geochemical signal of lakes. The precipitous increase in metal contamination from the 470 1930s, due to the start of open-cut mining and introduction of new technology, demonstrates471 the importance of considering historical records when interpreting metal contamination.

The HYSPLIT forward trajectories particle model has been demonstrated to be a useful tool 472 473 to track past metal contamination from airborne sources, explaining most of the metal atmospheric transport and metal deposition into the lakes of the TWWHA. Sediment EF 474 values > 50 (classified as extremely severe enrichment) and metal concentrations above 475 ISQG-High concentrations indicate that metal contamination might be posing health risks to 476 aquatic organisms and humans feeding on them. Further investigation of metal 477 478 bioaccumulation in ecosystems of the TWWHA are warranted starting in the northwest area where the metal contamination is highest. 479

480 Although mining activities have decreased significantly in the area, the metals deposited in 481 the sediment are constantly remobilised by redox reactions, wind, catchment leaching and 482 activities of microorganisms in the sediment. The environmental contamination in the 483 TWWHA, therefore, is not a past issue and justifies current attention.

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649	Figures and Legends
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651 652 653 654 655	Figure 1 – Map of Tasmania, Australia, with the Tasmanian World Wilderness Heritage Area (TWWHA) in grey. The yellow circles indicate the six lakes in this study. The red stars indicate the three mining centres in the area: Queenstown, Mount Lyell and Mount Read, Rosebery.
656 657 658 659 660	Figure 2 – Principal Component Analyses of metal concentrations (Cu, Se, Cd, Pb, As, Zn, Fe and Al) in sediments of the four lakes in the Tasmania Wilderness World Heritage Area (Perched Lake and Lakes Dove, Dobson and Cygnus) and closer to the mining centres (Basin Lake and Owen Tarn) A) before mining and B) since mining activities commenced.
661 662 663	Figure 3 – Distance between lakes studied for metal contamination and (A) Queenstown mining site and (B) Rosebery mining site. Maps developed in ArcMap 10.3.
664 665 666	Figure 4 – (A) Atmospheric temperature and (B) Precipitation in the Tasmanian Wilderness World Heritage Area lakes (1961 – 1990). Data from Australian Bureau of Meteorology, maps developed in ArcMap 10.3.
667 668 669 670 671 672	Figure 5 –The Hybrid Single Particle Lagrangian Integrated Trajectory Model (HYSPLIT) forward trajectories calculations demonstrating air parcel trajectories and directions of atmospheric particles and associated metals in the Tasmanian Wilderness World Heritage Area. This map represents the average circulation of air masses over Tasmania during the period 1961-1990 for particles released at Queenstown (42°S, 145.5°E) and Rosebery (41.78°S, 145.5 °E). 1sd is one standard deviation
673 674 675 676	Figure 6 – Metal concentrations (mg/kg) in the four lakes in the Tasmania Wilderness World Heritage Area (Perched Lake and Lakes Dove, Dobson and Cygnus) and two lakes closer to the mining centres (Basin Lake and Owen Tarn) pre-mining and during its peak.
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679 680 681	Table 1- Metal concentrations in lake sediments within the Tasmania Wilderness World Heritage Area. Metal concentrations are presented as mean concentrations per mining period.
682 683 684 685	Table 2: Tasmanian Wilderness World Heritage Area and surrounding lakes and their attributes: catchment size (km ²), geographic coordinates, annual precipitation (mm), annual temperature (°C), and distance from the mining sites in both Queenstown and Rosebery.

- 686Table 3 Linear model results (p value and R²) for environmental factors influencing metal atmospheric687transport and metal deposition in sediments of the four lakes in the Tasmania Wilderness World Heritage688Area (Perched Lake and Lakes Dove, Dobson and Cygnus) and two lakes closer to the mining centres (Basin689Lake and Owen Tarn).
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Table 4 - Enrichment factors of metals in the sediments of the four lakes in the Tasmania Wilderness World
 Heritage Area (Perched Lake and Lakes Dove, Dobson and Cygnus) and two lakes closer to the mining

- 693 centres (Basin Lake and Owen Tarn).
- 694 Footnote for Table 4: White = no enrichment, very light grey = minor enrichment, light grey = moderate
- 695 enrichment, mid grey = moderately severe enrichment, dark grey = severe enrichment, very dark grey = very
- 696 severe enrichment, and black = extremely severe enrichment.
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- **Table 5 Ratios of maximum concentration of sediments (average concentrations from 1930 to 1980)**
- 700 from lakes in this study to ANZECC/ARMCANZ (2000) sediment quality guideline values. Metal
- 701 concentrations highlighted in yellow indicate that the metal concentrations are above guidelines values.
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Lake	Mining	Al	Fe	Pb	Cu	Zn	As	Se	Cd	
ID	Phases				Concentration (mg/Kg)					
Owen Tarn	Before mine (1880)	1639	769	10	5	3	3	1.7	0.2	
Owen Tarn	Mining peak (1930 to 1980)	3336	1311	492	220	51	50	7.6	1.4	
Owen Tarn	Fold-increase	1.0	0.7	46	43	19	14	3.6	6.4	
Basin	Before mine (1880)	4185	19077	7	4	6	6	4.5	0.0	
Basin	Mining peak (1930 to 1980)	8680	26838	388	203	44	63	7.4	0.6	
Basin	Fold-increase	1.1	0.4	58	44	6	10	0.7	55.8	
Dove	Before mine (1880)	39337	15366	74	9	50	9	3.3	0.3	
Dove	Mining peak (1930 to 1980)	55179	16986	248	47	78	33	3.4	0.6	
Dove	Fold-increase	0.4	0.1	2.4	4.3	0.6	2.8	0.0	1.0	
Cygnus	Before mine (1880)	12292	2741	2	2	3	0.6	0.3	0.1	
Cygnus	Mining peak (1930 to 1980)	14662	4382	5	12	6	0.8	2.3	0.1	
Cygnus	Fold-increase	0.2	0.6	1.1	6.4	1.1	0.3	7.3	0.0	
Dobson	Before mine (1880)	15783	10567	1	9	19	0.5	BDL	BDL	
Dobson	Mining peak (1930 to 1980)	16214	12929	13	9	39	0.8	BDL	0.4	
Dobson	Fold-increase	1.0	1.2	11	1.1	2.1	1.6	N/A	N/A	
Perched	Before mine (1880)	10362	3505	8	18	27	2.5	4.3	1.3	
Perched	Mining peak (1930 to 1980)	18360	6760	118	71	37	6.4	8.5	2.3	
Perched	Fold-increase	0.8	0.9	14	2.9	0.3	1.5	1.0	0.7	

BDL= below detection limit.

Lake	Core depth analysed	Catchment size	Longitude	Latitude	Total Annual Precipitation (mm)	Mean Annual Temperature (°C)	Distance from (km)		
name	(cm)	(Km²)					Queenstown	Rosebery	
Owen Tarn	48	0.2	145.609434	-42.0997	2816	8.8	5	36	
Basin Lake	37	0.9	145.54829	-41.9808	3128	9.6	11	22	
Dove Lake	37	5.3	145.962222	-41.6575	2706	9	58	38	
Perched Lake	10	0.2	145.686163	-42.5648	2150	10.6	55	88	
Lake Dobson	10	1	146.617778	-42.6719	1443	8.1	109	133	
Lake Cygnus	10	0.3	146.241944	-43.1183	1974	8.9	128	160	

Mal		Predictors								
Nietai	Distance Precip	itation Si	ze Temp	perature Fr	equency	\mathbf{R}^2	Р			
Al	Removed Rem	oved **	** :	***	***	0.87	***			
Fe	Removed Rem	oved *	*	NS	NS	0.09	**			
Pb	Removed Rem	oved N	S	NS	***	0.51	***			
Cu	Removed Rem	oved N	S	NS	***	0.67	***			
Zn	Removed Rem	oved **	**	NS	**	0.16	***			
As	Removed Rem	oved N	S	NS	***	0.38	***			
Se	Removed Rem	oved N	S	NS	***	0.19	***			
Cd	Removed Rem	oved N	S ·	***	***	0.13	**			

NS= Not significant. (*) p < 0.05; (**) p < 0.01; (***) p < 0.001.

Lake	As	Cd	Cu	Fe	Pb	Se	Zn
Owen Tarn	30.2	14.8	97.7	3.4	91	8.7	26
Basin Lake	5.4	27.4	21.9	0.7	28.4	0.8	3.4
Lake Dove	2.4	0.3	1	0.4	7.5	0.2	0.7
Lake Cygnus	1.2	0.7	6.1	1.2	1.7	0.1	1.6
Lake Dobson	1.9	45	1.3	3.1	10.2	N/A	3.2
Perched Lake	2.2	1.6	2.8	1.5	11	3.3	1

Table 4 - Enrichment factor of six metals in sediments of six lakes in the Tasmanian Wilderness World Heritage Area.

	EF < 1	EF < 3	EF 3 - 5	EF 5 - 10	EF 10 - 25	EF 25 -50	EF > 50
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grey = severe enrichment, very dark grey = very severe enrichment, and black = extremely severe enrichment. N/A = not applicable, concentration below detection limit

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	Core [Depth	Age	Al	As	Cd	Cu	Fe	Pb	Se	Zn
name	(cm)	-	(yr AD)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)	(mg/kg)
Owen Tarr	1	0	2014.97	1523.2	20.07087	0.122256	119.299	4348.275	125.7053	2.132662	3.025703
Owen Tarr		6	1988.93	2047.3	16.44124	0.759985	328.1267	1059.174	245.1051	4.157364	31.42973
Owen Tarr		7	1982.42	1235.8	18.32454	0.97269	271.0552	940.3028	328.9093	4.895969	30.57227
Owen Tarr		8	1975.28	2001	22.51395	0.843436	254.8223	1206.159	334.481	5.406267	27.77645
Owen Tarr		9	1968.33	1751	26.3264	1.148413	151.1321	1190.886	439.1272	4.81381	37.37832
Owen Tarr		10	1962.39	1760.1	43.92913	1.27633	213.7414	1457.833	580.1228	8.471749	53.43515
Owen Tarr		13	1954.19	1122.8	161.8876	3.36744	228.1248	1830.494	888.4573	15.07879	105.9735
				1653	48.23714	1.394716	241.1671	1280.808	469.3671	7.137325	47.76091
Basin Lake		0	2011	<i>AA</i> 27	135 233	0 459566	71 69379	87635 3	160 6696	2 637912	25 16474
Dasin Lake		0	2011	4427	155.255	0.455500	/1.055/5	07055.5	100.0050	2.057512	23.10474
Basin Lake		2	1980	6155.7	65.07089	0.477691	172.7802	35460.04	252.6329	7.092378	29.06678
Basin Lake		3	1963	10200.5	56.06931	0.54898	211.2576	22878.33	373.9696	6.495491	51.57856
Basin Lake		4	1947	9682.4	66.67455	0.678804	224.8108	22174.23	538.1037	8.500396	52.31859
				8679.533	62.60492	0.568492	202.9495	26837.53	388.2354	7.362755	44.32131
Lake Dora		0	1999.9	52415.99	33.3	0.5	45.6	22500	173.9766	3.1	67.78297
Lake Dora		1	1990.5	60656.09	31.8	0.4	60.6	17600	230.0407	3.4	70.43287
Lake Dora		1.25	1985.5	56232.46	34.1	0.5	61.3	17800	245.9304	3.4	72.38321
Lake Dora		1.5	1980.4	63033.66	37.3	0.5	60.3	17200	281.6176	3.8	68.65178
Lake Dora		1.75	1973.8	62291.91	38	0.7	49.1	16500	305.0446	4	66.36334
Lake Dora		2	1967.3	62620.8	35.4	0.8	36.8	16300	274.0526	3.7	65.13988
Lake Dora		2.25	1961	20156.28	30.7	0.8	36.7	16600	215.4716	2.7	66.60153
Lake Dora		2.5	1954.8	61262.06	26.7	0.5	21.1	16900	154.375	2.9	65.80733
				55179.04	33.42857	0.6	46.55714	16985.71	243.7904	3.414286	67.91142
Lake Dove		0	1000	52/15 00	33.3	0.5	15.6	22500	170.6	3 1	57 5
Lake Dove		0	1999	52415.55	55.5	0.5	45.0	22500	170.0	5.1	57.5
Lake Dove		10	1990	60656.09	31.8	0.4	60.6	17600	233	3.4	52.1
Lake Dove		12.5	1984	56232.46	34.1	0.5	61.3	17800	248	3.4	51.5
Lake Dove		15	1978	63033.66	37.3	0.5	60.3	17200	288	3.8	48.6
Lake Dove		17.5	1972	62291.91	38	0.7	49.1	16500	308	4	45.7
Lake Dove		20	1967	62620.8	35.4	0.8	36.8	16300	287	3.7	42.8
Lake Dove		22.5	1961	20156.28	30.7	0.8	36.7	16600	214	2.7	62.6
Lake Dove		25	1955	61262.06	26.7	0.5	21.1	16900	157	2.9	48.2
Lake Dove		27.5	1949	52765.87	24.1	0.5	23.6	17200	135.3	2.6	67.4
				54877.39	32.2625	0.5875	43.6875	17012.5	233.7875	3.3125	52.3625
		~	4060 F	077 0010	-						_
Lake Cygni		0	1993.5	9/7.3913	1	0.2	9.7	5590	9.1	0.6	7
		0.05	4000 0	13/77.5	0.8	0.1	11.95	4440	5.45	0.325	6.4
Lake Cygni		0.25	1983.8	13100	0.8	0.1	11.9	4660	6.4	0.5	6.4
Lake Cygni		0.5	1974.1	19800	0.9	0.1	12.2	4270	5	0.1	6.2
Lake Cygni		0.75	1964.3	19200	0.8	0.1	13	4340	5.4	0.3	7.1
Lake Cygni		T	1954.6	3010	0.7	0.1	10.7	4490	5	0.4	5.9

Lake	Time	Al	As	Cd	Cu	Fe	Pb	Se	Zn
Owen	EF (2014)		13.7	1.4	52.5	12.4	26.4	2.8	1.8
Tarn	EF (1950-1990)		30.2	14.8	97.7	3.4	91.0	8.7	26.0
	Concentration 2014	1523.2	20.1	0.1	119.3	4348.3	125.7	2.1	3.0
	Average concentration (1950-1990)	1653	48.2	1.4	241.2	1280.8	469.4	7.1	47.8
	Background concentration (mg/kg)	3336	3.22	0.19	4.98	769.07	10.41	1.66	3.71
Basin	EF (2011)		22.9	43.4	15.2	4.3	23.1	0.6	3.8
Lake	EF (1950-1990)		5.4	27.4	21.9	0.7	28.4	0.8	3.4
	Concentration 2011	4427	135.2329814	0.459566	71.69379	87635.3	160.6696	2.637912	25.16474
	Average concentration (1950-1990)	8679.533	62.60491811	0.568492	202.9495	26837.53	388.2354	7.362755	44.32131
	Background concentration (mg/kg)	4185	5.58	0.01	4.47	19077	6.58	4.45	6.22
Lake	EF (2000)	-	2.6	0.3	1.1	0.6	5.9	0.2	0.9
Dora	EF (1950-1990)		2.5	0.3	1.1	0.4	7.8	0.2	0.9
	Concentration 2000	52415.99	33.3	0.5	45.6	22500	173.9766	3.1	67.78297
	Average concentration (1950-1990)	55179.04	33.42857143	0.6	46.55714	16985.71	243.7904	3.414286	67.91142
	Background concentration (mg/kg)	9914.6	2.422	0.325	7.916	7424.214	5.602	2.69	14.108
Lake	EF (2002)		2.6	0.3	1.1	0.6	5.8	0.2	0.8
Dove	EF (1950-1990)		2.4	0.3	1.0	0.4	7.5	0.2	0.7
	Concentration (2002)	52415.99	33.3	0.5	45.6	22500	170.6	3.1	57.5
	Average concentration (1950-1990)	54877.39	32.2625	0.5875	43.6875	17012.5	233.7875	3.3125	52.3625
	Background concentration (mg/kg)	9914.6	2.422	0.325	7.916	7424.214	5.602	2.69	14.108
Lake	EF (2001)		20.3	21.0	70.0	22.0	40.4	2.9	24.5
Cygnus	EF (1950-1990)		1.2	0.7	6.1	1.2	1.7	0.1	1.6
	Concentration (2001)	977.3913	1	0.2	9.7	5590	9.1	0.6	7
	Average concentration (1950-1990)	13777.5	0.8	0.1	11.95	4440	5.45	0.325	6.4
	Background concentration (mg/kg)	11292	0.57	0.11	1.6	2935	2.6	2.39	3.3
Lake	EF (2000)		2.2	56.2	1.4	3.3	12.4	N/A	2.8
Dobson	EF (1950-1990)		1.9	45.0	1.3	3.1	10.2	N/A	3.2
	Concentration (2000)	15300	0.9	0.5	11.4	31800	11	BDL	45
	Average concentration (1950-1990)	14520	0.72	0.38	9.94	28120	8.6	BDL	48.74
	Background concentration (mg/kg)	17200	0.45	0.01	9.35	10700	1	0.01	18.3
Perched	EF (2002)		3.4	1.5	3.5	2.1	7.5	4.7	1.3
Lake	EF (1950-1990)		2.2	1.6	2.8	1.5	11.0	3.3	1.0
	Concentration (2002)	12200	7.1	1.38	60.94	6800	50.2	9.5	46.19
	Average concentration (1950-1990)	18557.14	6.8	2.171429	72.72	7242.857	111.3714	10.31429	51.34
	Background concentration (mg/kg)	13560	2.29	0.99	19.18	3640	7.42	2.26	39.01

Lake Dobsc	0	1998.3	15300	0.9	0.5	11.4	31800	11	BDL	45	
			14520	0.72	0.38	9.94	28120	8.6	BDL	48.74	
Lake Dobso	0.75	1987.2	14000	0.7	0.3	10	41700	9	BDL	45.8	
Lake Dobso	1.25	1979	15000	0.6	0.4	10.3	28600	8	BDL	52.2	
Lake Dobso	1.75	1970.6	14400	0.8	0.5	9.9	24200	9	BDL	47	
Lake Dobso	2.25	1962.2	14500	0.6	0.3	9.5	23900	8	BDL	49.4	
Lake Dobso	2.75	1953.9	14700	0.9	0.4	10	22200	9	BDL	49.3	
			14520	0.72	0.38	9.94	28120	8.6	BDL	48.74	
Perched La	0	1995.8	12200	7.1	1.38	60.94	6800	50.2	9.5	46.19	
			18557.14	6.8	2.1/1429	/2./2	/242.85/	111.3/14	10.31429	51.34	
Perched La	0.25	1989.9	18500	6	1.84	73.4	8700	89.8	8.7	132.18	
Perched La	0.5	1984.1	19600	9.6	1.78	80.01	8200	98.3	20.9	43.29	
Perched La	0.75	1977.8	15500	4.6	3.19	63.1	5800	101.4	6.2	28.65	
Perched La	1	1971.5	23200	6.7	2.74	91.47	8600	142.9	8.3	38.81	
Perched La	1.25	1965	18000	5.6	2.07	74.06	6700	124.7	8.7	43.85	
Perched La	1.5	1958.5	18100	9.1	1.87	67.71	6700	112.6	11	38.42	
Perched La	1.75	1952.1	17000	6	1.71	59.29	6000	109.9	8.4	34.18	
			18557.14	6.8	2.171429	72.72	7242.857	111.3714	10.31429	51.34	

0.8 0.1 11.95

4440

5.45 0.325

6.4

13777.5

Lakos	Pb		Cu		Z	Zn		As		Cd	
Lakes	ISQG-Low	ISQG-High	SQuiRTs								
Owen Tarn	18.4	4.2	6.0	0.7	0.6	0.3	8.1	2.3	2.2	0.3	11.6
Basin Lake	10.8	2.4	3.5	0.0	0.3	0.1	6.8	1.9	0.5	0.1	0.4
Lake Dove	6.2	1.4	0.9	0.0	0.5	0.2	1.9	0.5	0.5	0.1	0.6
Lake Cygnus	0.2	0.0	0.2	0.0	0.0	0.0	0.1	0.0	0.1	0.0	0.1
Lake Dobson	0.4	0.1	0.2	0.0	0.3	0.1	0.1	0.0	0.5	0.1	0.6
Perched Lake	2.9	0.6	1.4	0.0	0.7	0.3	0.5	0.1	3.2	0.5	2.5

	Pb	Cu	Zn	As	Se	Cd	
ОТ		919.8	390.8	120.3	161.9	16.8	3.4
Basin		538.1	224.8	57.9	135.2	8.5	0.7
Dove		308.0	61.3	90.5	38.0	5.2	0.8
Cygnus		9.1	13.0	7.8	1.0	3.5	0.2
Dobson		19.0	11.4	54.8	1.4	0.0	0.7
Perched		142.9	91.5	132.2	9.6	20.9	4.8
Guidelines							

	Pb		Cu		Zn		As		Cd	
	ISQG-Low	ISQG-High								
OT	18.40	4.18	6.0	0.7	0.6	0.3	8.1	2.3	2.2	0.3
Basin	10.76	2.45	3.5	0.0	0.3	0.1	6.8	1.9	0.5	0.1
Dove	6.16	1.40	0.9	0.0	0.5	0.2	1.9	0.5	0.5	0.1
Cygnus	0.18	0.04	0.2	0.0	0.0	0.0	0.1	0.0	0.1	0.0
Dobson	0.38	0.09	0.2	0.0	0.3	0.1	0.1	0.0	0.5	0.1
Perched	2.86	0.65	1.4	0.0	0.7	0.3	0.5	0.1	3.2	0.5
Guidelines	50.0	220.0	65.0	270.0	200.0	410.0	20.0	70.0	1.5	10.0

