

Cooperative Research Centre for Landscape Environments and Mineral Exploration





# REGIONAL GEOCHEMICAL SURVEYS: RIVERINA PILOT PROJECT METHODOLOGY AND PRELIMINARY RESULTS

Patrice de Caritat, Subhash Jaireth, Megan Lech, John Pyke

CRC LEME OPEN FILE REPORT 160

August 2004

CRC LEME is an unincorporated joint venture between CSIRO-Exploration & Mining, and Land & Water, The Australian National University, Curtin University of Technology, University of Adelaide, Geoscience Australia, Primary Industries and Resources SA, NSW Department of Mineral Resources and Minerals Council of Australia, established and supported under the Australian Government's Cooperative Research Centres Program.







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This report presents outcomes of the 'Regional Geochemical Surveys-Murray Darling Basin Project', a collaborative research project between CRC LEME and Geoscience Australia that commenced in July 2003 and continued until June 2004.

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#### ABSTRACT

During financial year 2003-04, we carried out fieldwork and preliminary analyses for a collaborative project between CRC LEME and Geoscience Australia aimed at testing methodologies and strategies for regional geochemical surveys in Australia. The Riverina bioregion, in southern New South Wales and northern Victoria, was selected for this pilot project. Overbank sediments were chosen as sampling media, with a near-surface sample (TOP: O horizon, from 0-10 cm below the humus layer) and a bottom sample (BOT: B-C horizon, ~10 cm interval between approximately 70-90 cm below the humus layer) being collected. River Red Gum (*Eucalyptus camaldulensis*) leaves were also collected at a small number of locations. The sample sites were selected to be near outlets or spill points of large catchments, so that overbank sediments there could reasonably be assumed to represent well-mixed, fine-grained composite samples of all major rock and soil types present in the catchment. Three field trips visited 143 sites in the Riverina. This report documents the methodologies used in this survey and presents results to date.

Sediment samples were subjected to a detailed description and the determination of bulk parameters in the field (texture, moist and dry colour, field pH). In the laboratory, the samples were analysed for moisture content, EC 1:5, pH 1:5; some were also subjected to laser particle size analysis. Bulk (<180  $\mu$ m) composition was determined by X-Ray Fluorescence (XRF), Inductively Coupled Plasma Mass Spectrometry (ICP-MS) and Instrumental Neutron Activation Analysis (INAA), and the samples were also analysed by sequential digestion and prepared for heavy mineral separation. At the time of writing, bulk composition of the samples from the first fieldtrip (eastern Riverina) only was available. All together the concentrations of 62 elements were determined, and maps are presented showing the spatial and statistical distributions in the TOP and BOT samples and of the TOP/BOT ratios. These results will be the basis for further interpretative work and the backbone of a forthcoming geochemical atlas of the Riverina.

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### **1. INTRODUCTION**

Geochemical baselines provide information on the natural concentration of chemical elements and compounds in the regolith, which is the blanket of soils, sediments and weathered rocks common in Australian landscapes. These natural concentrations vary greatly in different areas, due to geology, biological processes and other factors (Reimann and Caritat, 1998). It is important to know the natural concentration and distribution of elements so that:

1. A baseline can be established against which future changes can be measured; and

2. Localised **anomalies** or **contaminations** can be identified and better understood. This knowledge will thus assist in environmental management and mineral exploration.

Regional geochemical surveys targeting 'normal' (background) sampling sites, as opposed to mineralised or polluted sites, are needed to generate baseline geochemical maps. These are fundamental data layers to develop environmentally appropriate and responsible land-use policies or, in mineral exploration, to aid discovery of ore bodies and promote the potential of certain regions.

Recent low-density geochemical surveys in China (Hangxin et al., 1997) suggest that wide-spaced geochemical sampling can provide good knowledge of broad geochemical patterns across a nation. To illustrate the potential of such studies. Xie and Ren (1993) reported "... the discovery of several hundreds of new mineral occurrences, including 400 new gold occurrences, many of which are being developed into workable mines". The Chinese study has prompted Geoscience Australia and CRC LEME to propose collaborative pilot studies which will determine whether low-density sampling can be applied with similar success in Australia. The present study in the Murray-Darling Basin is the first pilot project in what is hoped will become a widespread series of pilot studies and, eventually, a national geochemical baseline study. The rationale for low-density geochemical surveys has been discussed by Bølviken et al. (1992), among others, and proven to be effective (e.g., Fauth et al., 1985; Reimann et al., 1997, 2003; Li and Wu, 1999; Ruitenbeek, 2000; Gustavsson et al., 2001). This kind of work is likely to generate interest in a range of organisations in Australia (universities, government agencies concerned with land, water, agriculture, salinity, health, industry, research organisations, etc.) and add to the development of a global geochemical baseline (Darnley et al., 1995).

Given the potential environmental problems faced by present and future governments in this country, it is timely to provide them with a truly representative, nation-wide tool describing baseline conditions at the beginning of this decade. A regional geochemical survey at the national scale would provide this information. Australia is one of the last developed nations to *not* have regional geochemical information at the disposal of decision-makers. A geochemical survey and ensuing geochemical atlas of Australia would be a tremendously useful tool to address national priority issues that are important in terms of

- **Economy**: mineral discoveries through cover, new commodities, new provinces
- **Environment**: land-use appropriate for natural conditions, natural resources management

• Society: health and welfare of population, health of stock and safety of crops

The Murray-Darling Basin (MDB) (Figure 1) has been selected as the focus for this initial research proposal, being a vital region in terms of social, agricultural and mineral importance. The MDB

- Covers 1.06 M km<sup>2</sup> or 14% of the country's total area
- Contains 45% of the Australian crop area and 43% of the total number of farms
- Is Australia's most important agricultural region, accounting for 41% of the nation's gross value of agricultural production
- Is an important provider of resources such as wheat (34% of national production), cotton (96%), dairy, rice and viticulture
- Is home to nearly 2 M people, or 11% of the total Australian population (1996 census)

More specifically, we have undertaken a first geochemical survey in the Riverina bioregion (southern New South Wales and northern Victoria) (Figure 1).

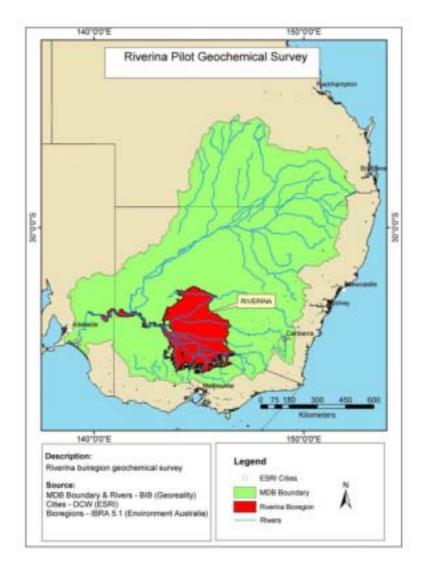


Figure 1: The Murray-Darling Basin with the Riverina pilot study area

Overbank sediment samples were collected for the pilot studies, as it is believed that this sampling medium can be used across the entire basin and beyond, providing comparable datasets between this and future pilot surveys. The merits of overbank sediments for low-density baseline geochemical surveys in Australia are borne out by previous studies (Ottesen et al., 1989; Murrell, 1991; Edén and Björklund, 1994, 1996; Ridgway et al., 1995; Volden et al., 1997; Xie and Cheng, 1997).

Top (0-10 cm) and bottom (~70-90 cm) samples have been taken at each site to detect the residence and mobility of chemical elements in natural environments and any impact of pollutants and land-use changes since European settlement. Data from the Riverina pilot study will be examined at a number of scales to ascertain suitability of low density sampling in the rest of the MDB.

### 2. AIMS

The aims of the project are (1) to provide internally consistent background geochemical data in the Riverina pilot region, and (2) to help develop a suitable geochemical sampling methodology for the whole MDB. This dataset will assist decisions in land-use management, geohealth and mineral exploration.

The aims of this report are to provide a detailed record of what was done during financial year 2003-04 in the Regional Geochemical Surveys project and, more importantly, to document the methods used for future reference.

### **3. ACHIEVEMENTS TO DATE**

This pilot project started on 1 July 2003, and had a duration of 12 months. In this period, the following work was carried out:

- overbank sediment sampling in the whole Riverina area (3 fieldtrips)
- vegetation sampling at selected sites in the western and northern Riverina area (2 of the above fieldtrips)
- groundwater sampling in the eastern Riverina area (1 separate fieldtrip) (not discussed here)
- sample preparation, field and laboratory measurements (field pH, texture, wet and dry colour, pH 1:5, EC 1:5, moisture content) of all eastern Riverina overbank sediments
- total (bulk) sediment (<180 μm) analysis by X-Ray Fluorescence (majors and traces) of all eastern Riverina overbank sediments
- total (bulk) sediment (<180 μm) analysis by Inductively Coupled Plasma Mass Spectrometry (traces) of all eastern Riverina overbank sediments
- total (bulk) sediment (<180 μm) analysis by Instrumental Neutron Activation Analysis (majors and traces) of all eastern Riverina overbank sediments
- laser grainsize analysis of 20 selected eastern Riverina overbank sediment samples
- sequential digestion of all eastern Riverina overbank sediments
- analysis of sequential digest by Inductively Coupled Plasma Mass Spectrometry (traces) of all eastern Riverina overbank sediments (not discussed here)
- presentations given at the 17<sup>th</sup> Australian Geological Congress in Hobart, February 2004 (Caritat et al., 2004) and at Geoscience Australia (15/9/03 by Patrice de Caritat, 4/2/04 by Patrice de Caritat, 7/6/04 by Megan Lech)
- preparation of a report detailing methods and showing results to date (this Report)

Outcomes (proposal - 1 July 2003)	Outcomes (achieved - 30 June 2004)
Geochemical database	Database populated with results to date
	- Sites for the eastern Riverina entered in RTMAP (to be migrated to DEVIANT)
	- Chemistry results sent from laboratory to Database Group (to be entered in OZCHEM)
Geochemical maps of sediments, plants and groundwater	Maps prepared for bulk (<180 μm) sediments (this Report). Geochemistry of plants and groundwater not yet available
Implications for land-use decisions, mineral exploration and geohealth	Interpretations started, but requires further work and availability of whole dataset
Dissemination of results to stakeholders	Presentations given locally and nationally, web release of this Report
Recommendations for geochemical survey of MDB	Started, but requires further work and availability of whole dataset

Outcomes achieved against the project proposal are given below.

#### 4. METHODS

#### 4.1. Site selection

The area, the Riverina bioregion, was chosen as the focus for this first pilot project because of the recent geophysical data acquisition carried out by the New South Wales Department of Mines here. Because of the nature of the main sampling medium selected, overbank sediments, the landscape units of interest here are hydrologic catchments. Thus, a coverage of Australian catchments was obtained from the Centre for Resource and Environmental Studies (CRES) at the Australian National University. The lowest points around each catchment's boundary were determined using ArcHydro, as detailed in Appendix 1. These points were the theoretical sampling sites.

All theoretical sampling sites were carefully checked with topographic, drainage and road coverages to see if the locations were reasonable and likely to generate the information sought. Where necessary, the sampling sites were moved from the software-generated ('theoretical') position to manually selected locations before going in the field. In the field, of course, the final site locations were decided by considering local landscape, likely anthropogenic interferences, and pragmatic considerations (e.g., access), and the actual site locations were recorded by GPS, after allowing for averaging, and recorded in the field log sheets (Appendix 2) and entered in the database.

#### 4.2. Field methods

The fieldwork for the Riverina geochemical survey took place in three phases:

- eastern Riverina: 20 September to 1 October 2003 (PdC, SJ, JP, ML)
- western Riverina: 14 to 22 April 2004 (PdC, Matthew Lenahan, Benjamin Ackerman)
- northern Riverina: 12 to 19 May 2004 (SJ, ML, Peter Taylor)

The main sampling medium targeted during this survey was overbank sediments. The rationale behind this sample selection is that the majority of Australia is covered by regolith, much of it transported. Thus by avoiding *in-situ* regolith or bedrock, the sampling medium would be most likely to be found in many environments around the country (but not all!). Further, overbank sediments, by their very nature, represent well-mixed composite samples of major rock and soil types present in the catchment, and are thus a particularly appropriate choice for low-density geochemical surveys, which aim to represent large tracts of land, as accurately as possible, with as few samples as necessary. An additional advantage of these sediments is that, by virtue of being deposited at times of flooding in low-energy environments, they are fine-grained and thus are more likely to be characterised by high signal-to-noise ratios for many chemical elements.

At each site, the sampling log sheet was filled to record, among others, date, GPS location, width and flow of the stream or river (if applicable), landscape position, vegetation, etc. The main sampling media targeted was overbank sediments, with two 10-cm intervals sampled: one in the O horizon from 0 to 10 cm below the humus layer (TOP), and another in the B-C horizon at depths generally ranging from 70 to 90 cm

(BOT). For each interval, two bags of ca 1-2 kg of material were collected, one being for all geochemical and mineral analyses, the other for heavy mineral separation. The samples were numbered according to the Geoscience Australia code:

**2003 751 X 601** to **2003 751 X 665**, for instance for the eastern Riverina, with '2003' being the year, '75' the project number, '1' the geologist or team, 'X' the layer/bag (with 'X' = '1' for TOP geochemistry, '2' for TOP heavy minerals, '3' for BOT geochemistry, '4' for BOT heavy minerals) and '601' the sample number (incrementally increasing for each different site). Samples for geochemistry and for heavy minerals were collected in exactly the same fashion, thus samples with 'X' = '1' and '2' are essentially field duplicates, as are those with 'X' = '3' and '4'.

Similarly, the western Riverina sample numbers are 2004 751 X 001 to 2004 751 X 043, and the northern Riverina sample numbers are 2004 751 X 100 to 2004 751 X 134. Thus, 65, 43 and 35 sites were visited and sampled during the eastern, western and northern Riverina fieldtrips, respectively, for a total 143 sites for the Riverina geochemical survey.

During the western and northern Riverina fieldtrip, River Red Gum (*Eucalyptus camaldulensis*) leaves were collected at several sites, and for these 'X' = '5'. The reasoning behind this is that we hope to gain some insight into the biological uptake of chemical elements in this widespread plant.

During the northern Riverina fieldtrip, two sites, 111 and 133, were selected for detailed soils profile sampling; for each of these sites eight 10-cm depth interval samples, labelled '0-10', '10-20', '20-30', '30-40', '40-50', '50-60', '60-70' and '70-80', were collected in addition to the four normal samples ('X' = '1' to '4'). These profiles were taken to test the representativity of the TOP and BOT samples in the context of continuous soil profiles.

The TOP sample was collected with a white plastic scoop after removal of any organic litter, humus and roots, over an area of ca 30 x 60 cm in size. For many sites, given the protracted drought during the last several months, the overbank sediments were extremely dry and hard. Thus, it was necessary at many sites to break up the material with a steel crowbar before scooping it up. It is possible that some samples have experienced minor contamination from (blue) paint flakes coming off the tools when first used, but this is not considered to be either a significant or a pervasive effect (the paint was analysed and shown to mostly consist of Ti). The BOT sample was collected after augering or digging down to the required depth, and collecting the disaggregated material either with the auger or with gloved hands. Where a trench had to be dug to reach the BOT sample, it was preferably collected using the white plastic scoop. Where augering was possible, three to six holes were generally drilled and the BOT sample was a composite of material from each hole; where digging was necessary, the BOT sample was collected from the bottom of the trench over a larger area. Up to four different hand augers (of different lengths and widths, 3 and 4 inch diameter) and a mechanical auger (posthole digger; 7 inch diameter) made of steel, a shovel and crowbar were used during fieldwork.

The TOP and BOT samples were collected in clear plastic bags (ca 30 x 40 cm), which were rolled and stapled closed, and placed in pairs in calico (cotton) bags with

string closing. The sample numbers were written on both the plastic and calico bags to minimise the risk of being rubbed off during transport (which did not occur).

For the western and northern Riverina fieldtrips, the texture, pH and Munsell<sup>TM</sup> colour were determined in the field according to standard protocols (Northcote, 1971; Munsell Color Company, 1975). Soil pH test kits from Inoculo<sup>TM</sup> Laboratories were used for pH determination. For the eastern Riverina samples, these parameters were determined in the laboratory within a few weeks of collection, together with other parameters (see below). The purpose of these field determinations is as follows.

Texture: provides context as to how and why elements may be concentrated or depleted in the soil, and can be a potential indicator of material origin. Clays can adsorb and exchange several elements. This is important where multiple material sources (which may influence element dispersion) need to be discriminated (e.g. alluvium vs aeolian material). It may also provide insights into the hydrology of the material, which in turn influences the geochemistry.

Colour: is important as it gives information as to whether it is an oxidising or reducing environment. Red soils are more oxidised and contain more iron oxides whereas the more yellow soils contain more goethite. Iron oxides may absorb or adsorb metals of interest including As and Au.

pH: provides information that can be related to element mobility and stability within the regolith materials. It can be correlated with various chemical and environmental factors that influence soils and plants. It must be recognised, however, that pH can vary markedly within a few metres of the sample site.

#### 4.3. Sample preparation methods

The various sample preparation methods used are described in detail below and summarised in Table 1.

Method	Analysis location	Sample 1 (TOP)	Sample 2 (TOP)	Sample 3 (BOT)	Sample 4 (BOT)	
Field pH	GA/field	Raw		Raw		
Texture	GA/field	Raw W		Raw W		
Colour (Dry)	GA/field	Raw		Raw		
Colour (Moist)	GA/field	Raw W		Raw W		
EC 1:5	GA		D45 U		D45 U	
рН 1:5	GA		D45 U		D45 U	
Moisture Content	GA		D105 U		D105 U	
XRF	GA	D40 S180		D40 S180		
ICP-MS	GA	XRF bead		XRF bead		
Neutron Activation		D40 S180		D40 S180		
Sequential leach Laser Particle	GA	D40 S180		D40 S180		
size analysis* Heavy Mineral	GA		Raw		Raw	
Fraction	GA		Raw		Raw	
Key:	Dx	Dried to x °C	(see text for o	luration)		
	W	Wet				
	Sx	Sieved to x µ	ım			
	U	Unsieved				
	Raw	As found under field conditions				
*The analysis was d	one on 2 and	1 hut thou are	Ishallad se 1 -	and 3 (pactorn	Divering only)	

 Table 1: Summary of sample preparation methods used for the Riverina geochemical survey

\*The analysis was done on 2 and 4 but they are labelled as 1 and 3 (eastern Riverina only)

#### 4.3.1. Drying and sieving for geochemical analysis

- For geochemical analysis, one TOP and one BOT samples ('X' = '1' and '3', respectively) were selected for each site (for instance 2003 751 1601 and 2003 751 3601 for the first sample site from the eastern Riverina)
- The last four digits of each sample number (e.g., 1601 and 3601) were engraved into the floor of a 250 x 380 x 50 mm aluminium baking dish
- Each sample was then spread evenly in its dish and dried in an oven at 40 °C for 48 hours
- The sample was disaggregated by placing it in a 250 x 350 mm sample bag, rolling the top closed and then pounding it with a large ceramic pestle. This process was repeated until sufficient sample was obtained, replacing the sample bag as required. The sample was then sieved using 180 µm nylon cloth until approx 80 g of <180 µm material was obtained</li>
- No further grinding of the sample was required. The samples were blended by shaking

#### 4.3.2. Fusion of glass beads for X-ray fluorescence (XRF) analysis

- Accurately weighing approx 0.27 g of sample and 1.73 g of 57:43 (lithium tetraborate:metaborate) flux into a Pt crucible
- Adding 0.5 mL of a 20% LiNO<sub>3</sub> solution to the crucible
- Sintering the mixture in a furnace set at 400 °C for 10 minutes to oxidise any sulphides present
- Transferring the crucible to the Bradway<sup>TM</sup> rocking furnace and heat at 1100 °C for a further 10 minutes

- Adding a few crystals of NH<sub>4</sub>I anti-wetting agent to the melt after 8 minutes
- Removing the melt after 10 minutes and quickly quench press on a preheated graphite disc using an aluminium press
- After pressing, annealing the disc on a hotplate at 180 °C for 20-30 minutes

#### 4.3.3. Heavy mineral separation

The sample bags collected for heavy mineral analysis ('X' = '2' and '4' for TOP and BOT samples, respectively) were emptied one by one in plastic basins and immersed in water overnight to disaggregate the sediments. The resulting slurry was subsequently poured into a black plastic panning pan with ridges on one side. The fines were washed off until only coarse and dense material was left over. This residue was then washed and dried in an oven at 40 °C overnight. It is intended to further separate heavy minerals using dense liquids and carry out X-ray diffraction (XRD), Scanning Electron Microscopy (SEM, including trials on the AutoGeoSEM at CSIRO, Perth) and/or optical determination of the dominating heavy mineral phases. Geochemical analysis of these is envisaged for selected samples, as is potentially SHRIMP analysis of zircons.

#### 4.4. Laboratory methods

#### 4.4.1. X-ray fluorescence spectrometry (XRF) of regolith and sediment samples

XRF analysis was carried out in the Minerals Division Laboratory at Geoscience Australia. The laboratory's XRF is a Philips PW2404 4kW sequential spectrometer using a Rh tube. It is used to determine the major elements present in samples presented to the laboratory and a range of the routinely more abundant trace elements to complement those analysed on the ICP-MS.

The instrument is calibrated using a range of USGS and SARM (South African Reference Material) and other international standards. Synthetically prepared standards and some Australian Soil and Plant Analysis Council (ASPAC) standards were also used. The 'Classic Model' as outlined in the 'Philips SuperQ/Quantitative Users Guide' was used to calculate alpha coefficients and line overlap factors.

Reference monitor standard Si3 is used.

The routinely determined major elements are SiO<sub>2</sub>, TiO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, Fe<sub>2</sub>O<sub>3</sub>, MnO, MgO, CaO, Na<sub>2</sub>O, K<sub>2</sub>O, P<sub>2</sub>O<sub>5</sub> and SO<sub>3</sub> (Table 2).

This program uses 57:43 X-Ray flux (57% Li tetraborate 43% Li metaborate) because of its ability to fuse a wide range of sample types. It is used for all regolith samples and any samples that may contain significant NaCl. The trace elements Sc, V, Cr, Ni, Cu, Zn, Ba, Cl and F are determined on these fusions because NaCl and other soluble salts migrate to the surface of powder pellets as the binder dries causing gross errors. Generally speaking, the longer the wavelength of the element being measured the worse the error. ICP-MS results are used for all other trace elements.

Appendix 3 shows the detection limits of the XRF method.

#### 4.4.2. Inductively Coupled Plasma Mass Spectrometry (ICP-MS)

ICP-MS analysis was carried out in the Minerals Division Laboratory at Geoscience Australia. The laboratory's ICP-MS is a Perkin Elmer Elan 6000. It is used to determine a range of trace elements to complement those analysed on the XRF.

The instrument is calibrated using synthetic standards. The calibrations obtained are verified against a range of USGS, SARM (South African Reference Material) and other international standards. Australian Soil and Plant Analysis Council (ASPAC) standards are used to verify calibrations for vegetable material.

The samples for ICP-MS analysis were prepared from the glass bead previously used for XRF analysis. After breaking up the bead, approximately 100 mg are weighted accurately and transferred to a Teflon<sup>TM</sup> container, to 1 mL distilled 40% HF, 5 mL distilled concentrated HNO<sub>3</sub> and 1 mL internal multi-element standard (see below) are added. The container is sealed and heated to 140 °C overnight. After cooling, the solution is transferred to a volumetric flask, made up to 200 mL with distilled water and run in the ICP-MS against prepared standards.

Samples are introduced to the ICP-MS as a mildly acidic aqueous solution containing approximately 100 ppm dissolved solids by way of a nebuliser that aspirates the sample with high velocity argon, forming a fine aerosol or mist. The aerosol then passes into a spray chamber where larger drops are removed via a drain. This process is necessary to produce droplets small enough to be vaporised in the plasma torch. Typically, only 2% of the original mist passes through the spray chamber.

Trace elements routinely determined by the ICP-MS are Be, V, Cr, Ni, Cu, Zn, Ga, Ge, As, Rb, Sr, Y, Zr, Nb, Mo, Ag, Cd, Sn, Sb, Cs, Ba, La, Ce, Pr, Nd, Sm, Eu, Tb, Gd, Dy, Ho, Er, Yb, Lu, Hf, Ta, Pb, Bi, Th and U (Table 2).

The analytical method used divides the analytes above into four groups. The count rate obtained for each analyte is ratioed against one of four isotopes introduced to the solution as internal standards, one for each of the analyte groups. Ni-61, Sm-147, Tm-169 and Tl-205 are the internal standard commonly used.

Inter-element interference factors are determined using a range of single element solutions. Likely inter-element interferences were selected from a Perkin Elmer compilation of theoretical interferences.

Appendix 3 shows the detection limits of the ICP-MS method.

#### 4.4.3. Instrumental neutron activation analysis (INAA)

Neutron activation analysis was performed at Becquerel Laboratories, Sydney. For the determination of most elements samples of about 1 to 30 g are weighed and heat sealed in polypropylene vials. A flux monitor is attached to each sample and up to 200 samples and their monitors are activated for 10-30 minutes in a thermal neutron flux of 2-4 x  $10^{12}$  n cm<sup>-2</sup> s<sup>-1</sup>. The gamma spectrum for each radioactive sample and monitor is measured simultaneously after seven days decay. Each sample is tested for the amount of activity to adjust sample-detector geometries so no dead time corrections

are necessary. Samples are then mounted on automatic sample changer tables and counted for 10-30 minutes. The gamma ray spectra are measured using hyperpure Ge coaxial detectors linked to multichannel analysers as part of an integrated counting control and data handling system. Spectral data are analysed using in-house programs developed by Becquerel Laboratories.

#### 4.4.4. Sequential digestions

The sequential digestion method of Gray et al. (1999) was used to specifically extract weakly bound chemical elements, which may be indicative of more recently migrated and adsorbed or incorporated species (e.g., Cameron et al., 2004). The extraction sequence is:

- 1. Ammonium acetate at pH 5, targeting carbonates and surface adsorbed metals
- 2. 0.1 M hydroxylamine hydrochloride, targeting Mn oxides
- 3. 0.25 M hydroxylamine hydrochloride, targeting Fe oxides

After the sequential digestions, the samples are presented to the ICP-MS as described elsewhere. Results of these analyses are not available at the time of writing.

#### 4.4.5. Laser particle size analysis

Twenty raw (bulk) overbank sediment samples from TOP and BOT layers were delivered to the Sedimentology Laboratory at Geoscience Australia to be analysed with the laser particle size analyser. The equipment used is a Malvern<sup>TM</sup> Mastersizer 2000 coupled to a Hydro 2000 MU pump unit, which allows dispersion of large volume samples using standard laboratory beakers as the sample container. The basic principles of particle size analysis are summarised by Rawle (undated).

#### 4.4.6. pH 1:5

The following method is from the Bureau of Rural Sciences (A. Plazinska, pers. comm., 2003). Transfer 30-40 g of unsieved (raw) sample into a labelled aluminium tray. Put sample in oven at 45 °C for a minimum of eight hours, then place in desiccators to allow to return to room temperature. Accurately weight 10 g ( $\pm$  0.02 g) of sample into a plastic vial. Add 50 mL of distilled water, close vial and shake by hand to ensure thorough mixing of sample. Place sample vial in plastic bowls and place bowls in New Brunswick shaker for one hour at 160 rpm and 25 °C. Measured pH using calibrated bench top pH meter with pH probe and temperature probe. Probes are rinsed with distilled water and dried with lab tissues between samples.

#### <u>4.4.7. EC 1:5</u>

Same method as above, except EC is measured with calibrated EC meter and probe.

#### 4.4.8. Moisture content

The following method is from the Bureau of Rural Sciences (A. Plazinska, pers. comm., 2003). Accurately weight 100-120 g of unsieved (raw) sample and transfer into a labelled, pre-weighted aluminium tray. Put the sample in oven at 105 °C for a minimum of eight hours, then place in desiccator to allow to return to room

temperature. Accurately record weight of dry sample, subtract weight of aluminium tray and calculate moisture content (%).

#### 4.5. Data analysis methods

Data generated by this project was delivered to Geoscience Australia's database group to be stored on Geoscience Australia's corporate databases Deviant and Ozchem. Data analysis is performed using ArcMap® (Minami, 2000) for basic spatial representation (e.g., Figures 2-11), and DAS® (Dutter et al., 1995) for more advanced statistical treatment (boxplots, cumulative frequency distributions, histograms) and geochemical mapping (Appendix 4). For statistical analysis, results below the detection limit (DL) were replaced by ½ DL values, to still enable mathematical treatment (such as ratios or log-transformation).

XRF and ICP-MS-GA Labs		INAA-Becquerel Labs			
ELEMENTS	UNITS	DL*	ELEMENTS	UNITS	DL*
Ag ICP-MS	ppm	0.01	Ag	ppm	5
AI2O3 XRF	%	0.001			
As ICP-MS	ppm	0.5	As	ppm	1
			Au	ppb	5
Ba ICP-MS	ppm	2	Ва	ppm	100
Be ICP-MS	ppm	0.05			
Bi ICP-MS	ppm	0.01			
			Br	ppm	1
CaO XRF	%	0.002	Ca	%	1
Cd ICP-MS	ppm	0.1	Cd	ppm	20
Ce ICP-MS	ppm	0.02	Ce	ppm	2
CI XRF	ppm	5			
			Со	ppm	1
Cr XRF	ppm	4	Cr	ppm	Ę
Cs ICP-MS	ppm	0.01	Cs	ppm	1
Cu XRF	ppm	2			
Dy ICP-MS	ppm	0.01			
Er ICP-MS	ppm	0.01			
Eu ICP-MS	ppb	0.5	Eu	ppm	0.5
F XRF	ppm	50			
Fe2O3T XRF	%	0.002	Fe	%	0.02
Ga ICP-MS	ppm	0.1			
Gd ICP-MS	ppm	0.01			
Ge ICP-MS	ppm	0.02			
Hf ICP-MS	ppm	0.01	Hf	ppm	0.5
Ho ICP-MS	ppm	0.01			
			Hg	ppm	5
			Ir	ppb	20
K2O XRF	%	0.002	К	%	0.2
La ICP-MS	ppm	0.02	La	ppm	0.5
Lu ICP-MS	ppm	0.01	Lu	ppm	0.2
MgO XRF	%	0.004			
ΜΙΟΙ					
Calculate	%	0.001			
MnO XRF	%	0.001			
Mo ICP-MS	ppm	0.01	Мо	ppm	Ę
Na2O XRF	%	0.004	Na	%	0.01
Nb ICP-MS	ppm	0.04			
Nd ICP-MS	ppm	0.01			
Ni XRF	ppm	2			
P2O5 XRF	%	0.001			
Pb ICP-MS	ppm	0.5			
Pr ICP-MS	ppm	0.01			
Rb ICP-MS	ppm	1	Rb	ppm	20
Sb ICP-MS	ppm	0.01	Sb	ppm	0.2
Sc XRF	ppm	10	Sc	ppm	0.1
			Se	ppm	Ę
SiO2 XRF	%	0.006			

Table 2: Summary of methods used, elements analysed and detection limits obtained for the bulk (<180  $\mu m$ ) overbank sediments of the eastern Riverina

Table 2. Continued						
XRF and ICP-MS-GA Labs			INAA-Becque	INAA-Becquerel Labs		
ELEMENTS	UNITS	DL*	ELEMENTS	UNITS	DL*	
Sm ICP-MS	ppm	0.01	Sm	ppm	0.2	
Sn ICP-MS	ppm	0.01	Sn	ppm	500	
SO3 XRF	%	0.001				
Sr ICP-MS	ppm	1				
Ta ICP-MS	ppm	0.06	Та	ppm	1	
Tb ICP-MS	ppm	0.01				
			Те	ppm	5	
Th ICP-MS	ppm	0.01	Th	ppm	0.5	
TiO2 XRF	%	0.002				
U ICP-MS	ppm	0.07	U	ppm	2	
V XRF	ppm	6				
			W	ppm	2	
Y ICP-MS	ppm	0.22				
Yb ICP-MS	ppm	0.01	Yb	ppm	0.5	
Zn XRF	ppm	2	Zn	ppm	100	
Zr ICP-MS	ppm	1	Zr	ppm	500	

\* Detection Limit as provided by lab, unless different value reported with results from lab (eg, -0.01 ppm)

#### **5. RESULTS**

The actual location of sampling sites in the Riverina is shown in Figure 2. Geochemical results of the bulk analyses for the eastern Riverina region are illustrated for selected elements as 'growing-dots' geochemical maps (Figures 3-11) and as a complete series of diagrams (for all elements with at least one result above detection limit [DL]) combining statistical plots and Exploratory Data Analysis (EDA)-based maps (Appendix 4). Table 3 summarises the results of the bulk geochemical analyses carried out on the eastern Riverina overbank sediment samples. Where elements were analysed by more than one method, the preferred method is indicated in the Table, and the resulting maps are shown in Appendix 4.

Generally, ICP-MS and XRF are favoured over INAA because of the lower detection limits attained, the in-house analytical capability, and the internal consistency with the sequential digestion work. Only for two elements analysed in both laboratories is INAA preferred: Sb (because of erratic contamination risk by the Pt crucibles used in XRF bead preparation) and Sc (because of detection limits and potential overlap with Ca on XRF). Other elements analysed only by INAA are: Au, Br, Co, *Hg*, *Ir*, *Se*, *Te*, W (elements in italics had all results <DL in this study to date).

All elements analysed for are represented as EDA-based geochemical maps and statistical diagrams in Appendix 4 (except Hg, Ir, Se and Te, which had all results below detection limits), whilst a selection of parameters are shown as 'growing-dots' maps in Figures 3-11. Those maps illustrate how the distribution of various soil properties can be derived, for instance:

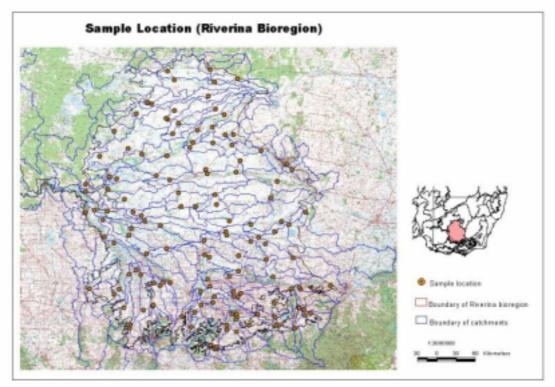


Figure 2: Location of factual sampling sites for the Riverina pilot geochemical survey (northern, eastern and western sub-regions).

- clay content (using Al<sub>2</sub>O<sub>3</sub> concentrations as a proxy, Figure 3)
- nutrient status (K<sub>2</sub>O concentrations, Figure 4)
- "sodicity risk" (Na<sub>2</sub>O concentrations, Figure 5)
- "acidity risk" (pH, Figure 6)
- "salinity risk" (EC, Figure 7; Cl concentrations, Figure 9)
- exploration pathfinders for Au mineralisation (As concentrations, Figure 8; Sb concentrations, Figure 11)
- exploration pathfinders for Cu or Cu-Au mineralisation (Cu concentrations, Figure 10)

The results of macroscopic properties are shown in Appendices 6 and 7. The quantitative results are summarised in Table 4.

Particle size analysis results are summarised in Figure 12 and detailed in Appendix 5. On average, 91% of the material has a grain size <180  $\mu$ m for these 20 samples. This suggests that the geochemical analyses performed on the <180  $\mu$ m fractions are likely to be representative of the bulk sample chemistry in these overwhelmingly fine-grained sediments. This is an important observation as many environmental geochemistry reports analyse the <2 mm fractions. Our results should thus be reasonably comparable with such studies.

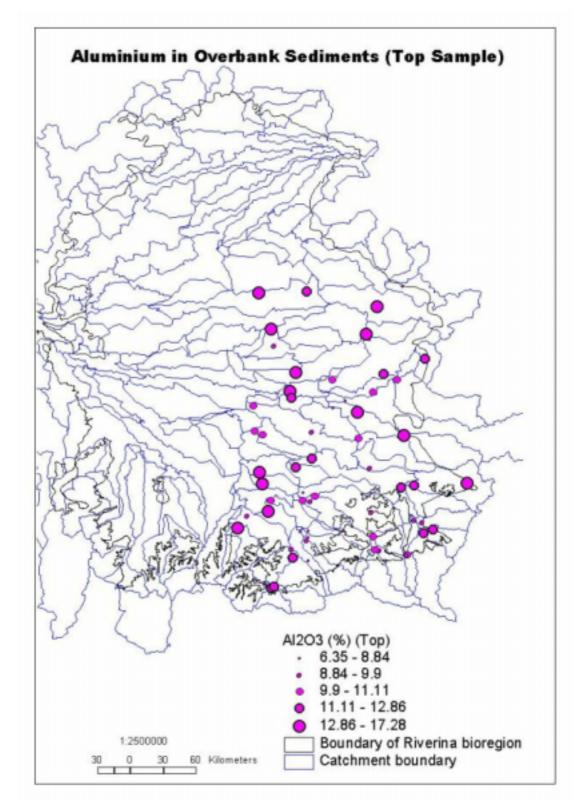


Figure 3a. Distribution of  $Al_2O_3$  (%, by XRF) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

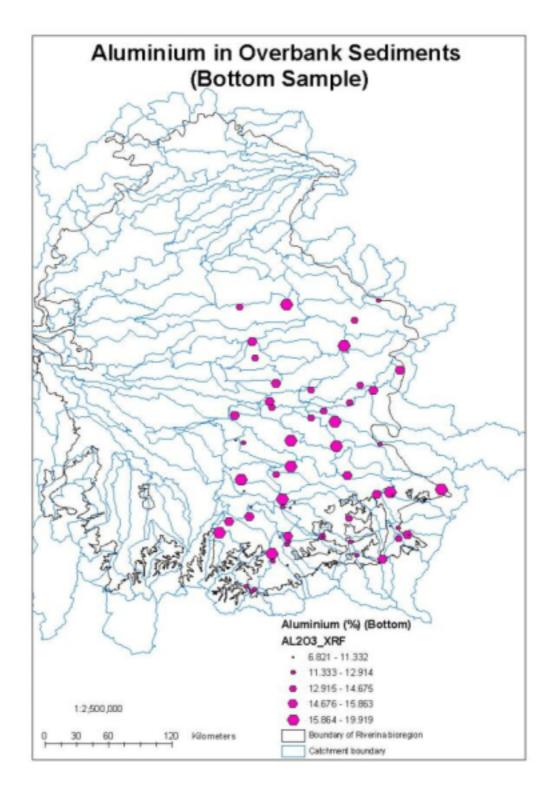


Figure 3b. Distribution of Al2O3 (%, by XRF) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

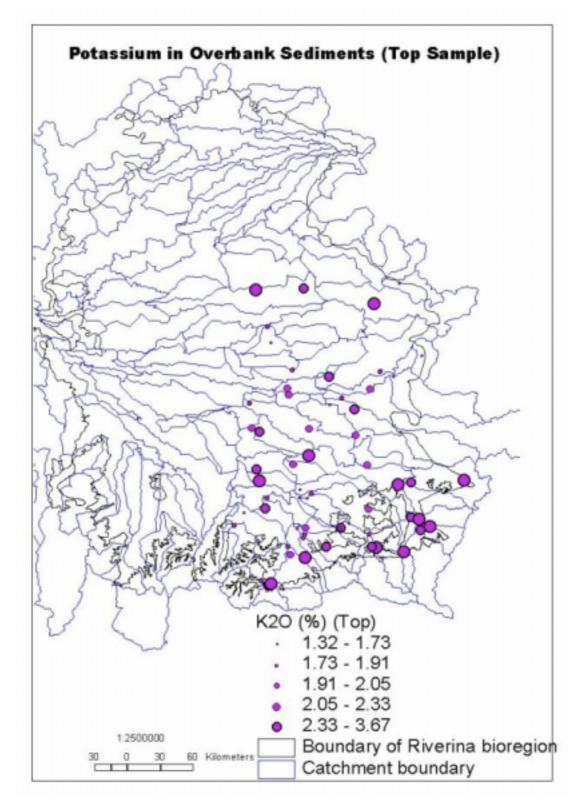


Figure 4a. Distribution of  $K_2O$  (%, by XRF) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

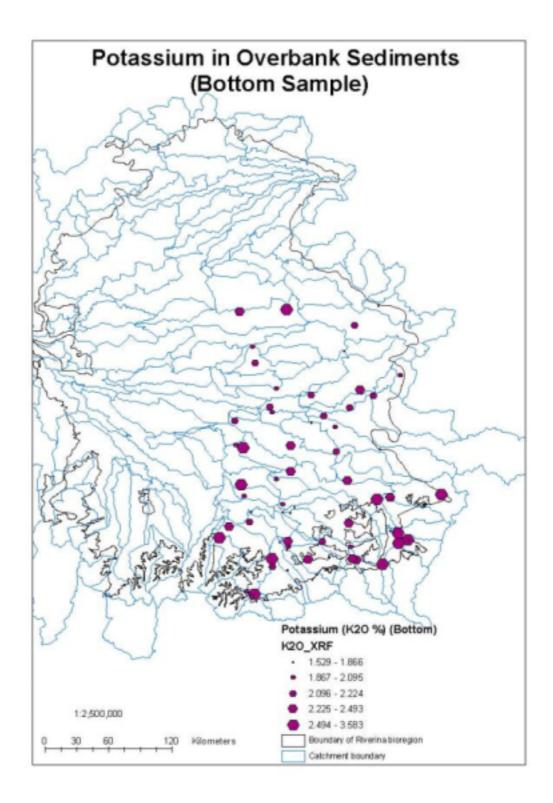


Figure 4b. Distribution of  $K_2O$  (%, by XRF) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

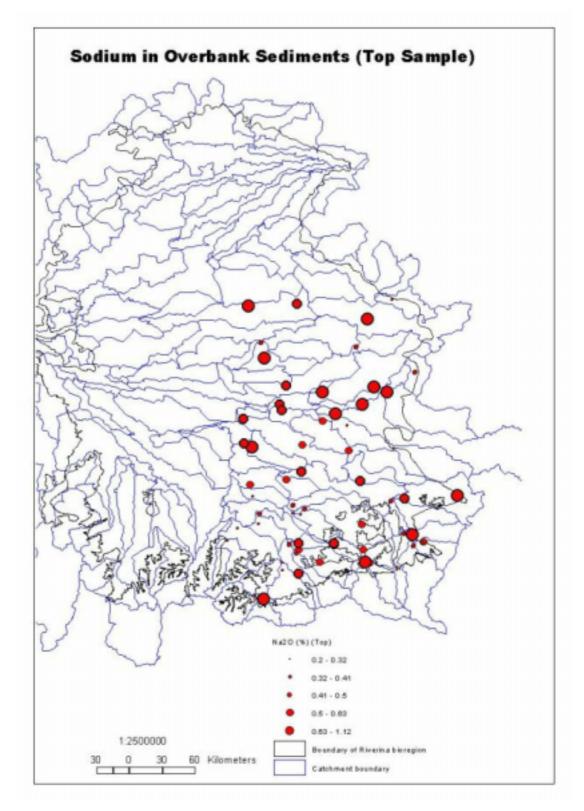


Figure 5a. Distribution of  $Na_2O$  (%, by XRF) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

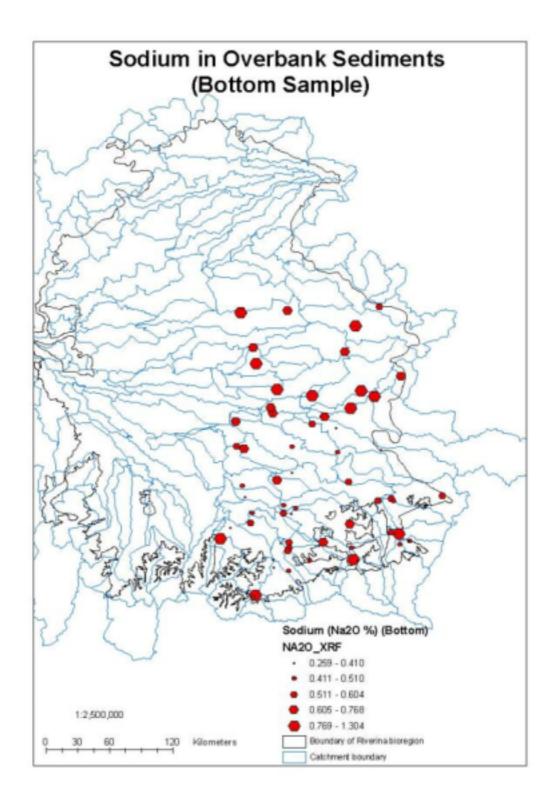


Figure 5b. Distribution of  $Na_2O$  (%, by XRF) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

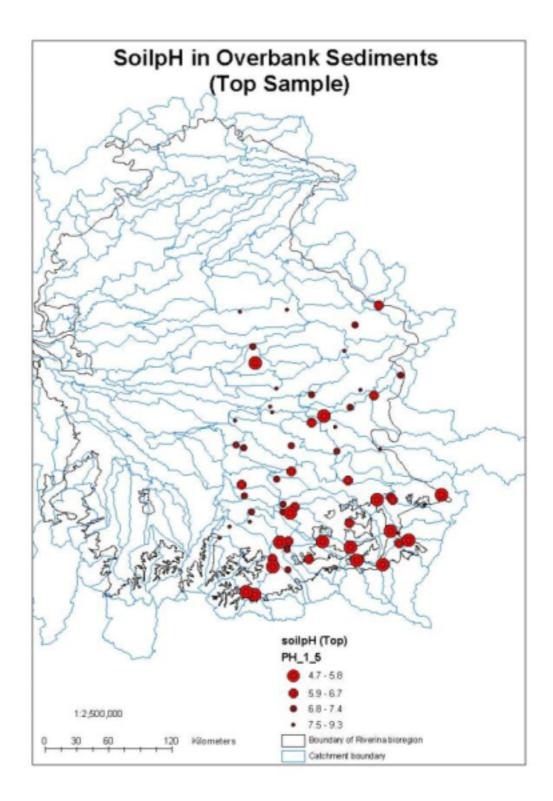


Figure 6a. Distribution of pH 1:5 in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

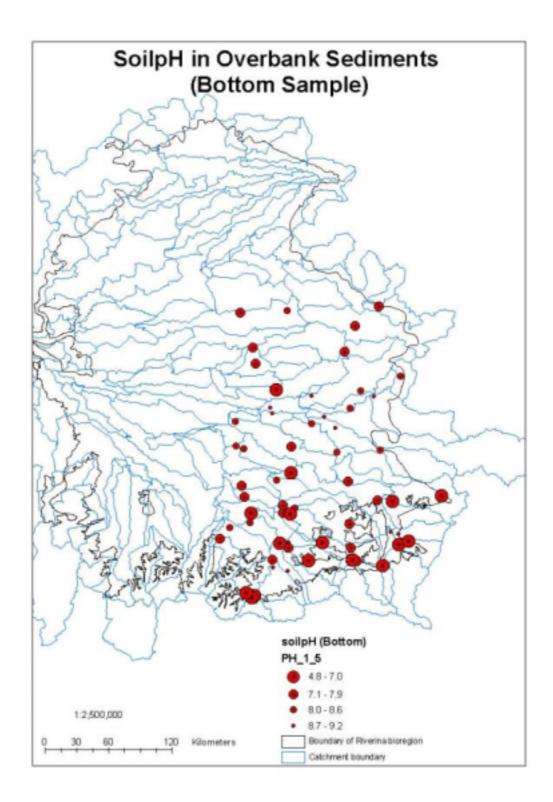


Figure 6b. Distribution of pH 1:5 in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

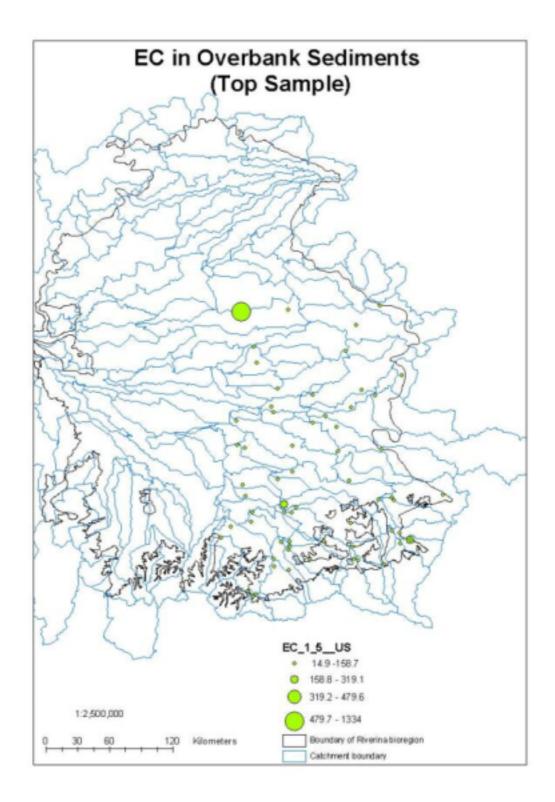


Figure 7a. Distribution of EC 1:5 in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

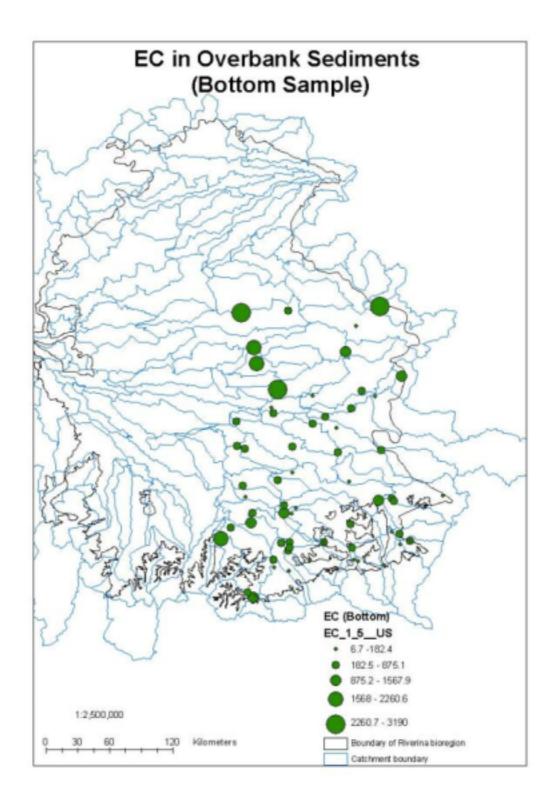


Figure 7b. Distribution of EC 1:5 in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

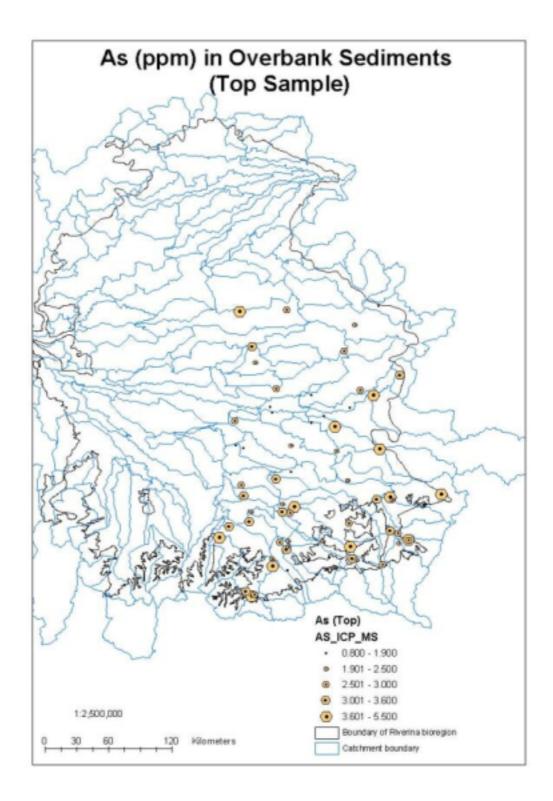


Figure 8a. Distribution of arsenic (ppm, by ICP-MS) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

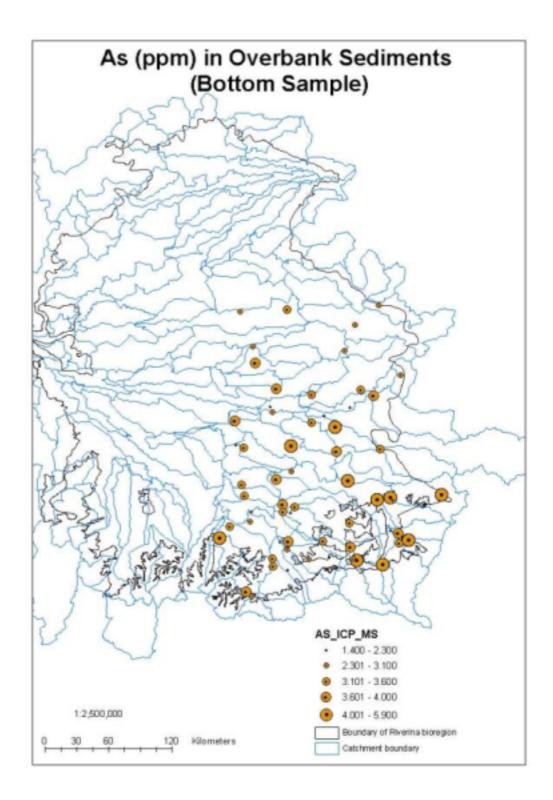


Figure 8b. Distribution of arsenic (ppm, by ICP-MS) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

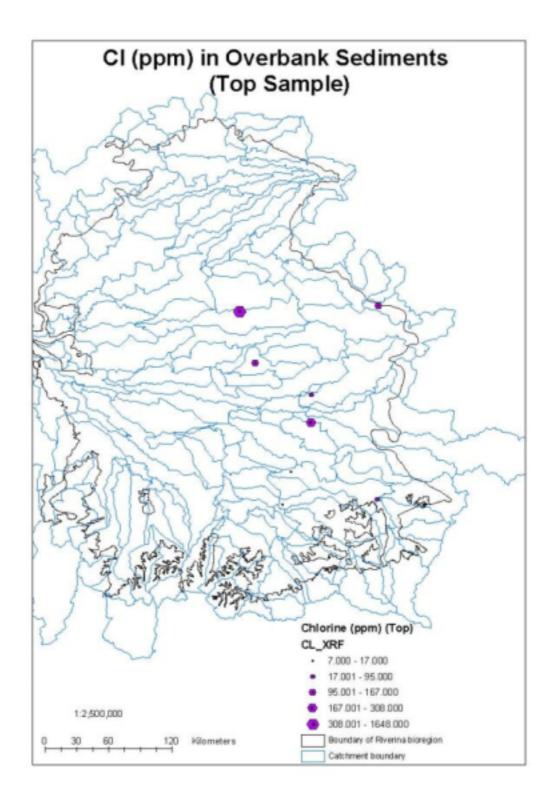


Figure 9a. Distribution of chloride (ppm, by XRF) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset. (Several values <DL not shown)

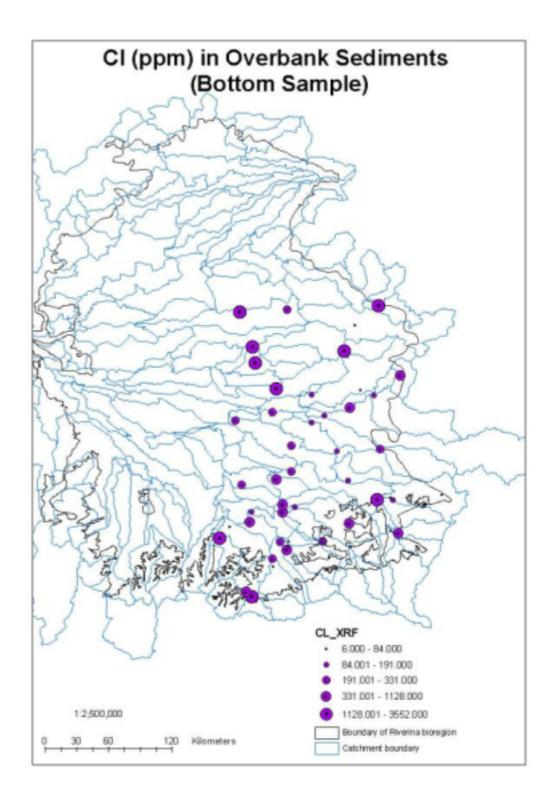


Figure 9b. Distribution of chloride (ppm, by XRF) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset. (Several values <DL not shown)

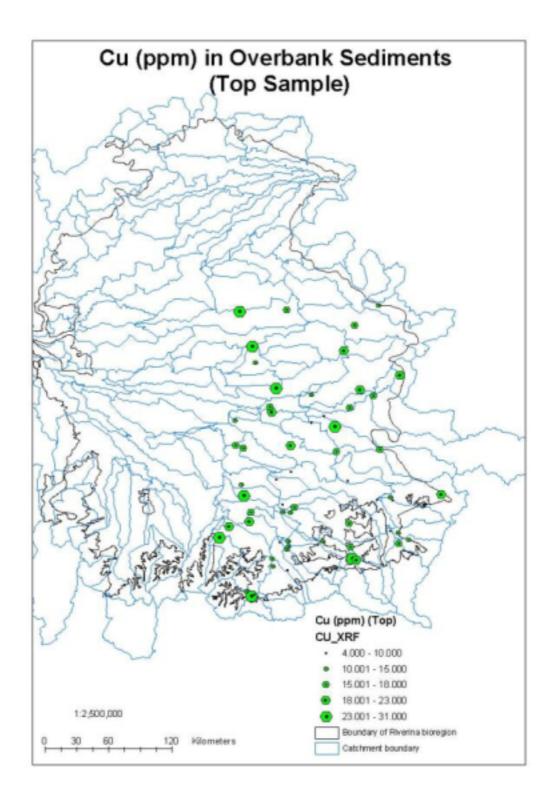


Figure 10a. Distribution of copper (ppm, by XRF) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

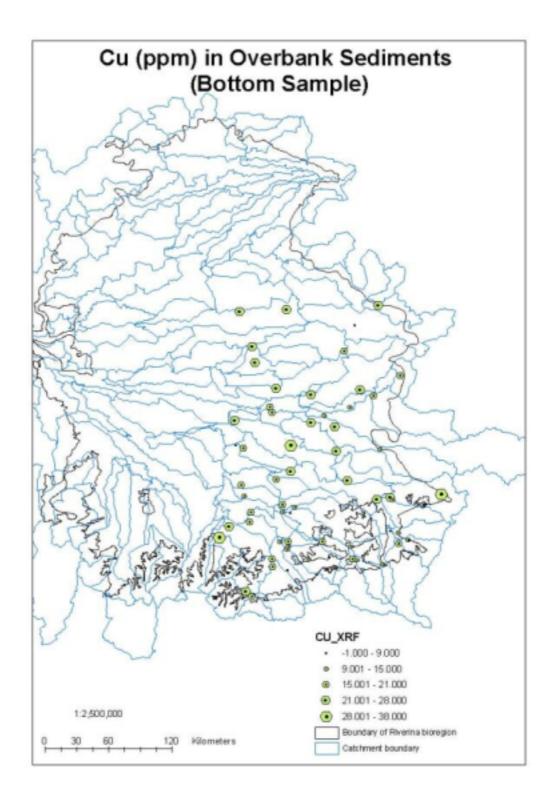


Figure 10b. Distribution of copper (ppm, by XRF) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

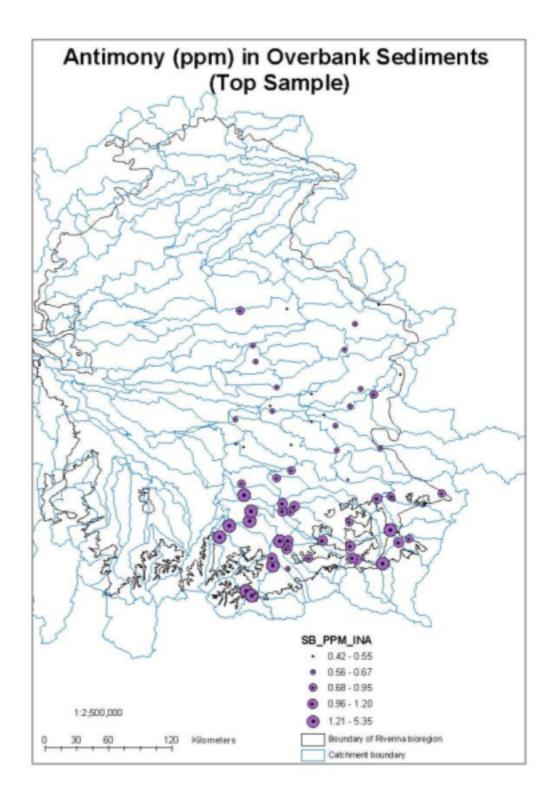


Figure 11a. Distribution of antimony (ppm, by INAA) in TOP overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

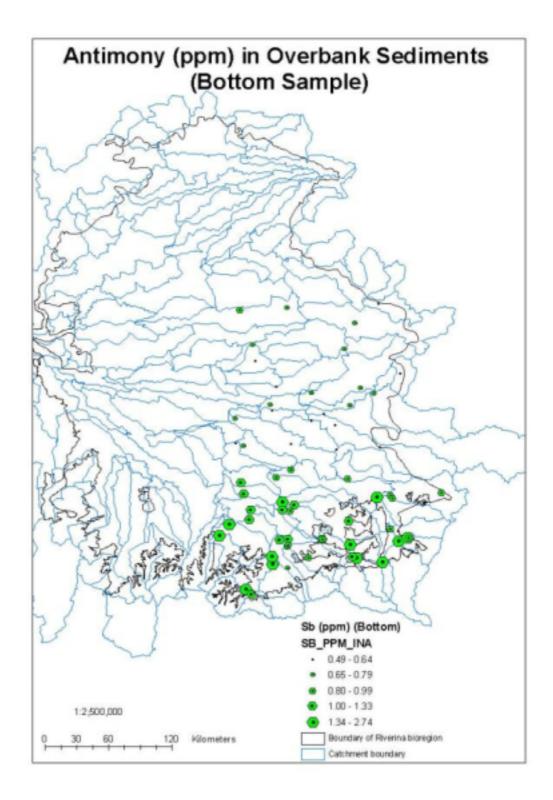


Figure 11b. Distribution of antimony (ppm, by INAA) in BOT overbank sediments of the eastern Riverina study area. Each of the five classes represents 20 % of the dataset.

El_unit_method	N	DL	Min	Med	Max	Units	% >DL	Preferred method?
Ag_ppm_ICPMS	129	0.01	0.01	0.07	0.13	ppm	100%	Y
Ag_ppm_INAA	129	5	<5	<5	<5	ppm	0%	
AI2O3_%_XRF	129	0.001	6.351	12.227	19.919	%	100%	Y
As_ppm_ICPMS	129	0.5	0.8	3	5.9	ppm	100%	Y
As_ppm_INAA	129	1	1.91	6.01	14.3	ppm	100%	
Au_ppb_INAA	129	5	<5	<5	50.7	ppb	9%	Y
Ba_ppm_ICPMS	129	2	249	476	1263	ppm	100%	Y
Ba_ppm_INAA	129	100	234	460	1220	ppm	100%	
Be_ppm_ICPMS	129	0.05	0.8	2.6	6.4	ppm	100%	Y
Bi_ppm_ICPMS	129	0.01	0.2	0.4	2.2	ppm	100%	Y
Br_ppm_INAA	129	1	<1	6.97	36	ppm	99%	Y
Ca_%_INAA	129	1	<1	<1	2.11	%	4%	
CaO_%_XRF	129	0.002	0.064	0.33	2.706	%	100%	Y
Cd_ppm_ICPMS	129	0.1	<0.1	<0.1	2.33	ppm	6%	Y
Cd_ppm_INAA	129	20	<20	<20	<20	ppm	0%	
Ce_ppm_ICPMS	129	0.02	56.86	86.12	170.4	ppm	100%	Y
Ce_ppm_INAA	129	2	58.8	88.4	163	ppm	100%	
Cl_ppm_XRF	129	5	<5	<5	3552	ppm	44%	Y
Co_ppm_INAA	129	1	2.96	11.1	34.2	ppm	100%	Y
Cr_ppm_INAA	129	5	27.1	69.3	182	ppm	100%	
Cr_ppm_XRF	129	4	14	56	150	ppm	100%	Y
Cs_ppm_ICPMS	129	0.01	2.57	6.96	16.08	ppm	100%	Y
Cs_ppm_INAA	129	1	2.21	7.32	17.7	ppm	100%	
Cu_ppm_XRF	129	2	<2	17	38	ppm	99%	Y
Dy_ppm_ICPMS	129	0.01	4.44	6.72	14.68	ppm	100%	Y
Er_ppm_ICPMS	129	0.01	2.64	3.94	8.39	ppm	100%	Y
Eu_ppb_ICPMS	129	0.5	734	1424	3704	ppb	100%	Y
Eu_ppm_INAA	129	0.5	0.71	1.39	3.65	ppm	100%	
F_ppm_XRF	129	50	<50	277	668	ppm	91%	Y
Fe_%_INAA	129	0.02	0.82	2.77	5.67	%	100%	
Fe2O3T_%_XRF	129	0.002	1.214	3.915	7.758	%	100%	Y
Ga_ppm_ICPMS	129	0.1	6.7	15.4	26.1	ppm	100%	Y
Gd_ppm_ICPMS	129	0.01	4.69	7.14	15.31	ppm	100%	Y
Ge_ppm_ICPMS	129	0.02	0.6	0.9	1.2	ppm	100%	Y
Hf_ppm_ICPMS	129	0.01	4.5	9.9	22.2	ppm	100%	Y
Hf_ppm_INAA	129	0.5	4.26	10.6	23.2	ppm	100%	X
Hg_ppm_INAA	129	5	<5	<5	<5	ppm	0%	Y
Ho_ppm_ICPMS	129	0.01	0.98	1.47	3.03	ppm	100%	Y
Ir_ppb_INAA	129	20	<20	<20	<20	ppb	0%	Y
K_%_INAA	129	0.2	0.81	1.69	3.28	%	100%	V
K2O_%_XRF	129 120	0.002	1.318	2.07	3.667	%	100%	Y
La_ppm_ICPMS	129 120	0.02 0.5	27.34	43.19 45 5	84.37 91.6	ppm	100%	Y
La_ppm_INAA	129 120		29.5	45.5	81.6	ppm	100%	V
Lu_ppm_ICPMS	129 120	0.01	0.49	0.62	1.32	ppm	100%	Y
Lu_ppm_INAA	129 120	0.2	0.39	0.51	1.12	ppm %	100%	V
MgO_%_XRF	129 120	0.004	0.144	0.7	1.761	% %	100%	Y Y
MLOI_%_Calculate MnO_%_XRF	129 120	0.001 0.001	1.01	6.618 0.063	12.199	% %	100% 100%	Y Y
₩IIIŲ_%_AK <b>F</b>	129	0.001	0.016	0.063	0.308	/0	100%	I

Table 3. Summary of geochemical composition of overbank sediments from the eastern Riverina

#### Table 3. Continued

Table 3. Continued								Preferred
El_unit_method	Ν	DL	Min	Med	Мах	Units	% >DL	method?
Mo_ppm_ICPMS	129	0.01	0.5	0.9	1.9	ppm	100%	Y
Mo_ppm_INAA	129	5	<5	<5	5.1	ppm	1%	
Na_%_INAA	129	0.01	0.169	0.421	0.996	%	100%	
Na2O_%_XRF	129	0.004	0.199	0.51	1.304	%	100%	Y
Nb_ppm_ICPMS	129	0.04	10.5	17	30	ppm	100%	Y
Nd_ppm_ICPMS	129	0.01	26.07	40.63	85.45	ppm	100%	Y
Ni_ppm_XRF	129	2	<2	20	81	ppm	98%	Y
P2O5_%_XRF	129	0.001	0.036	0.073	0.186	%	100%	Υ
Pb_ppm_ICPMS	129	0.5	14.1	21.8	33.8	ppm	100%	Y
Pr_ppm_ICPMS	129	0.01	7.26	11.16	23.19	ppm	100%	Y
Rb_ppm_ICPMS	129	1	65.3	123.3	216.8	ppm	100%	Y
Rb_ppm_INAA	129	20	58	136	236	ppm	100%	
Sb_ppm_ICPMS	129	0.01	0.5	2.9	11	ppm	100%	
Sb_ppm_INAA	129	0.2	0.42	0.82	5.35	ppm	100%	Y
Sc_ppm_INAA	129	0.1	5.22	12.2	22.9	ppm	100%	Y
Sc_ppm_XRF	129	10	<10	13	23	ppm	78%	
Se_ppm_INAA	129	5	<5	<5	<5	ppm	0%	Y
SiO2_%_XRF	129	0.006	54.524	72.593	87.273	%	100%	Y
Sm_ppm_ICPMS	129	0.01	5.62	8.27	16.6	ppm	100%	Y
Sm_ppm_INAA	129	0.2	5.1	7.79	15.7	ppm	100%	
Sn_ppm_ICPMS	129	0.01	2	4.2	15.6	ppm	100%	Y
Sn_ppm_INAA	129	500	<500	<500	<500	ppm	0%	
SO3_%_XRF	129	0.001	0.022	0.049	1.102	%	100%	Y
Sr_ppm_ICPMS	129	1	28.4	68.3	142.2	ppm	100%	Y
Ta_ppm_ICPMS	129	0.06	1	1.5	3.4	ppm	100%	Y
Ta_ppm_INAA	129	1	<1	1.37	2.91	ppm	91%	
Tb_ppm_ICPMS	129	0.01	0.79	1.2	2.59	ppm	100%	Y
Te_ppm_INAA	129	5	<5	<5	<5	ppm	0%	Y
Th_ppm_ICPMS	129	0.01	11.9	17.5	37.3	ppm	100%	Y
Th_ppm_INAA	129	0.5	12.2	18.4	34.7	ppm	100%	
TiO2_%_XRF	129	0.002	0.541	0.793	0.972	%	100%	Y
U_ppm_ICPMS	129	0.07	1.52	3.55	8.49	ppm	100%	Y
U_ppm_INAA	129	2	<2	3.23	7.3	ppm	88%	
V_ppm_XRF	129	6	31	78	145	ppm	100%	Y
W_ppm_INAA	129	2	<2	<2	5.13	ppm	47%	Y
Y_ppm_ICPMS	129	0.22	27.2	41.8	91	ppm	100%	Y
Yb_ppm_ICPMS	129	0.01	2.86	4.01	8.6	ppm	100%	Y
Yb_ppm_INAA	129	0.5	2.75	3.76	8.09	ppm	100%	
Zn_ppm_INAA	129	100	<100	<100	103	ppm	2%	V
Zn_ppm_XRF	129	2	14	54	104	ppm	100%	Y
Zr_ppm_ICPMS	129	1	156.3	386	868.3	ppm	100%	Y
Zr_ppm_INAA	129	500	<500	<500	840	ppm	40%	

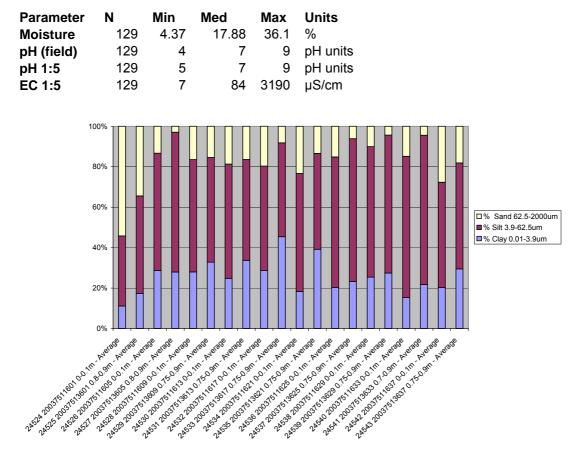


 Table 4. Summary of quantitative macroscopic properties of overbank sediments from the eastern Riverina (laboratory parameters)

Figure 12. Summary diagram of particle size analysis results.

### 6. FUTURE WORK

Future work in the Riverina will include:

- Completion of lab bulk parameters (moisture content, EC 1:5, pH 1:5) of TOP and BOT samples from western and northern Riverina
- Completion of sediment laser particle size analysis, as deemed necessary
- Bulk analyses (XRF, ICP-MS, ?INAA) of TOP and BOT samples from western and northern Riverina
- Selective digestion of all Riverina samples
- Heavy mineral separation and analysis (by X-Ray Diffraction, Scanning Electron Microscopy and/or AutoGeoSEM) for all Riverina samples
- Presentation of results at conferences, workshops and in publications/reports, including a web-based 'atlas' product

Beyond the Riverina pilot project, we are scoping a similar study in the Gawler region of South Australia, a collaboration between CRC LEME, PIRSA and Geoscience Australia. This study has started in 2004-05. Other possible areas being considered for further testing are in the wheat belt of Western Australia, the Tanami and perhaps the lower Balonne or Fitzroy-Burdekin catchments.

As an offshoot of the Riverina study, a collaborative PhD research project with University of Sydney began earlier this year in the southeastern Highlands and Australian Alps bioregions (New South Wales and Victoria). This area borders the Riverina, giving a greater coverage for the combined surveys, and will test the methodology and strategy in vastly different geomorphic environments.

### 7. CONCLUSIONS

A pilot low-density geochemical baseline survey was carried out in the Riverina bioregion during 2003-04, as a collaborative project between CRC LEME and Geoscience Australia. The pilot project aimed at testing methodologies and strategies for further regional geochemical surveys in Australia. The selected region was subdivided into large catchments whose drainage points were targeted as sampling locations. 143 locations were sampled in three field trips. At each sampling location, a near-surface (TOP: O horizon, 0-10 cm) and a deeper (BOT: B-C horizon, ~70-90 cm) overbank sediment sample was collected. Leaves of River Red Gum (*Eucalyptus camaldulensis*) trees were also collected at a small number of locations.

The methodologies used to collect, prepare and analyse the samples are detailed in this report, and preliminary results (pertaining to the first field trip in the eastern Riverina) are presented. The analyses carried out on the samples to date are:

- texture
- Munsell<sup>TM</sup> colour (moist and dry)
- field pH
- moisture content
- EC 1:5
- pH 1:5
- laser particle size analysis (selected samples)
- X-Ray Fluorescence (XRF) of bulk, sieved (<180  $\mu$ m) samples
- Inductively Coupled Plasma Mass Spectrometry (ICP-MS) of same
- Instrumental Neutron Activation Analysis (INAA) of same
- preparation for sequential digestion
- preparation for heavy mineral separation

Thus, the sediments are quantitatively characterised in terms of

- pH (field and 1:5), EC 1:5, moisture content
- grain size distribution (selected samples)
- Loss On Ignition (LOI)
- Bulk (<180 μm) composition for 62 elements/oxides (Ag, Al<sub>2</sub>O<sub>3</sub>, As, Au, Ba, Be, Bi, Br, CaO, Cd, Ce, Cl, Co, Cr, Cs, Cu, Dy, Er, Eu, F, Fe<sub>2</sub>O<sub>3</sub>, Ga, Gd, Ge, Hf, Hg, Ho, Ir, K<sub>2</sub>O, La, Lu, MgO, MnO, Mo, Na<sub>2</sub>O, Nb, Nd, Ni, P<sub>2</sub>O<sub>5</sub>, Pb, Pr, Rb, Sb, Sc, Se, SiO<sub>2</sub>, Sm, Sn, SO<sub>3</sub>, Sr, Ta, Tb, Te, Th, TiO<sub>2</sub>, U, V, W, Y, Yb, Zn, Zr)

Preliminary statistical treatment was carried out and geochemical maps prepared for all parameters (except Hg, Ir, Se and Te, which had all results below detection limits), and presented in Appendix 4. Those maps illustrate how the distribution of various soil properties can be derived and what the results can be used for, e.g., determine clay content, nutrient status, "sodicity risk", "acidity risk", "salinity risk", exploration pathfinders for Au, Cu or Cu-Au mineralisation. These results will be the basis for further interpretative work and the backbone of a forthcoming geochemical atlas of the Riverina.

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### APPENDICES

# Appendix 1: ArcHydro modelling

Methodology for selecting sample points using the Arc Hydro 1.1 Beta 2 extension in ArcGIS

<u>Datasets used</u>: Version 2 of the national 9 second Digital Elevation Model (DEM), Australian Nested Catchments and Sub-Catchments at 500 km<sup>2</sup> scale.

The national DEM was used to ensure that future geochemical surveys are fully comparable. NB: The DEM was not reconditioned as it is corrected using topographic information including drainage lines and point elevation data, and consequently is a robust dataset. This step may be useful for DEM's captured in airborne geophysical surveys. National Nested Catchments (500 km<sup>2</sup> scale) from the Centre for Resource and Environmental Studies (CRES) at the Australian National University were used instead of defining our own catchments with ArcHydro. This ensures that the dataset is fully comparable across the whole of Australia.

### Installation of the ArcHydro Extension

To be able to use the ArcHydro extension, the ArcInfo licence of the ArcGIS software is a requirement. The ArcHydro 1.1 Beta 2 extension was downloaded from <a href="http://www.crwr.utexas.edu/gis/gishydro03/ArcHydroTools/ArcHydroTools.html">http://www.crwr.utexas.edu/gis/gishydro03/ArcHydroTools/ArcHydroTools.html</a>

To install the toolbar:

- 1. Run the ArcHydro tools setup. After double-clicking the setup, browse to the desired installation location (use default location). The necessary files will be installed in the bin directory under the destination folder.
- 2. Open ArcMap. Right mouse click on the menu bar and scroll to "Customize."



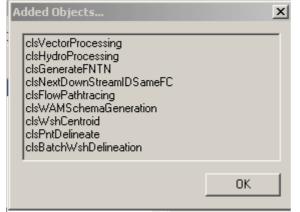
3. Click "add from file" when the Customize dialog box appears.

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- 4. Browse to the location of ArcHydroTools.dll. In this case, the .dll was located at: C:\Program Files\ESRI\ArcHydro
- 5. Select ArcHydroTools.dll and click Open to add to the Toolbar

Open		<u>? x</u>
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ArcHydroT	ools.dll	
	:Manager.dll	
SHPTools	s.dll	
File name:	ArcHydroTools.dll	Open
The name.		open
Files of type:	Type Libraries (*.olb,*.tlb,*.dll)	Cancel
		//.

6. Click OK to acknowledge the additions of the new objects



7. The ArcHydro toolbox should now appear as a toolbar

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✓Draw		
✓Layout		Delete
✓Editor		Reset
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Topology		
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8. The ArcHydro Toolbar should now appear.

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9. The Spatial Analyst Extension needs to be activated, by clicking Tools>Extensions..., and checking the box next to Spatial Analyst

### **Derivation of the Sampling Points**

To get the sample points, follow the steps below:

- 1. <u>Fill the sinks</u>: Terrain Processing>Fill Sinks. (rivdemv2\_fil). This function fills the holes in the grid. If a cell is surrounded by higher elevation cells water flows in but cannot flow out. By filling the sinks, the elevation is raised to prevent this problem from occurring.
- 2. <u>Flow direction</u>: Terrain Processing>Flow Direction. (fdir\_rivdem2) Insert the relevant DEM and name the output file. The values in the cells of the flow direction grid indicate the steepest decent from that cell – hence the flow direction.
- **3.** <u>Flow Accumulation:</u> Terrain Processing>Flow Accumulation. (fac-rivdem2). ArcHydro takes the flow direction grid and determines the flow accumulation grid.
- 4. <u>Drainage Point Processing</u>: Before this can be done, the nested catchments layer from CRES (CAT500) needs to be converted to a grid using the spatial analyst extension: Spatial Analyst>Convert>Features to Raster. The field needs to be ID, and the output cell size is 0.0025<sup>1</sup>. To obtain the drainage points: Terrain Processing>Drainage Point Processing. Insert the DEM and the catchment grid. The resultant points should be that the outlet points for each of the catchments. Check to ensure that there are not points in the middle of the catchments etc.

The first lot of drainage points were selected by following the whole process from start to finish. These points were then regenerated only redoing step 4. It was noted that the drainage points that were created were slightly different than the first lot of points generated. It is hypothesised that by creating the drainage points, the flow accumulation grid is modified slightly. This needs to be tested to ensure that when the process is repeated from start to finish that the same result can be achieved more than once.

<sup>&</sup>lt;sup>1</sup> This cell size was used as the 9 second grid spacing is in longitude and latitude approximates to 250 metres.

### Culling sample points for the Murray Riverina Low Density Survey

This methodology was used to cull points for the low density survey for the Riverina bioregion. Present at the meeting on 19<sup>th</sup> August were PdC, JP, SJ, ML. By documenting these procedures, it is hoped that there can be consistency in any other low density geochemical survey carried out within Australia.

As the survey is low density and has a define budget it was decided that approximately 120 points would be selected for the whole Riverina bioregion. This equates to around 60 points being selected for the area covering the east of the bioregion. These will be sampled in September 2003 field excursion.

The catchments (CRES National Nested Catchments with a threshold of 500km<sup>2</sup>) were sorted by perimeter<sup>2</sup>. These were classified into quantiles with 10 classes and were rounded to have no decimal places. The catchments were clipped to include those within the Riverina bioregion. It is now recognised that they should have been clipped to include those wholly or party within the bioregion. This was decided because the study boundary is arbitrary, and if the geochemical survey is to continue, picking up the samples close to those within the Riverina bioregion will save time in the long term. Those catchments that were party within the bioregion were added after the meeting and any sample points that were within these extra catchments were included into the sampling strategy.

Upon inspection, it was discovered that there were many errors<sup>3</sup> in the data which resulted in small "dangle" catchments in which sample points were assigned. It was essential to remove these erroneous points and this was done in a number of steps as described below:

- 1. Points within the bottom 10th percentile (i.e., 25 catchments with a perimeter of 947-965 m) were selected. These were randomly checked to determine that they were erroneous "dangle" catchments. The sample points minus the erroneous sample points (19 points) was named drnpts\_1sel.shp
- 2. This second pass cleanup looked at all the points from the 10<sup>th</sup> 20<sup>th</sup> percentiles (i.e., 53 points with a perimeter of 965-1013 m). This catchment selection was exported as a theme and converted to a shapefile called cat\_perlt1013.shp. This enabled a more thorough check to determine whether they were errors. There were 26 erroneous drainage sample points within the "dangle" catchments between 965-1013 m. The new file, minus the erroneous sample points, was named drnpts\_2sel. Catchments up to this size were determined to be erroneous with absolute certainty.

 $<sup>^{2}</sup>$  Alternatively the catchments could be sorted by area but were not in this case as the area statistics were not calculated at the time of the meeting.

<sup>&</sup>lt;sup>3</sup> These errors can result when the raster dataset is converted to a polygon. They can also be a result of small depressions in the DEM that don't drain into a stream e.g. clay pans. In this case they are not errors as such, but have such a small area that they are not worth worrying about.

- 3. The perimeters ≥1013 and ≤5000 m (9 catchment polygons) were selected<sup>4</sup>. These 201 "real" catchments were saved as cat\_pergt5000.shp. After 5000 m perimeter there looked to be a big jump in the perimeter length. Perimeters ≥5000 and ≤7000 m were selected and viewed. Some of these shapes looked more like catchments. These catchments were viewed to visually check their validity, but it was decided that they were still errors. All points between 1013 and 5000 were selected and deleted with the remaining points (158 drainage points) saved as drnpts\_3sel.shp.
- 4. From here on in, the smallest remaining catchments were scrutinised from the shapefile cat\_pergt5000.shp. The percentiles redefined for the perimeter with 10 quantile classes. The values were rounded to have no decimal places. These catchments had a mix of erroneous and "real" catchments. The points within the lower 10<sup>th</sup> percentile of the catchments were removed (140 drainage points left) and saved as drnpts\_4sel.shp.
- 5. Lastly, those points that were contained within catchments that were partly within the bioregion polygon were added to the coverage of drainage points.
- 6. The final step in selection of sampling sites consisted in a detailed inspection of each catchment and its computer-derived drainage point compared to topographic maps using GIS. This resulted in deleting still a few point and moving some others to be at more pragmatic locations (closer to tracks, outside swamps or reserves).

<sup>&</sup>lt;sup>4</sup> It was suggested at this stage that it may be worth cleaning up the nested catchments shapefile to that all those catchments with a perimeter of less than 5000m as these can be confidently assigned as erroneous catchments.

## Appendix 2: Field log sheet

						Method of	digging hole:	
			REGIO	NAL GEO	CHEMICAL S	URVEY		
			OVER	BANK SE	DIMENT SAM	PLING		
						Date	II	(dd/mm/yyyy)
SAMPLE ID	2004 75	51 <u>□</u> *_				Sampler/T	eam	
	REGION:				MAPSHE	ET		
	STREAM	NAME			NEARES PROPER	T TOWN		
	Zone			Latitu			Longitude	
	Datum			Way F	Pt		Altitude	m
DESCRIPTIO Approx si				(hectare	es/km <sup>2</sup> )	Catchmen	t ID	_ (ANU catch500
Predomin	ant bedrock	lithology	within catch	nment bas	sin		_	
SITE DESCR	IPTION							
	e/topograph	וy						
Land use								
	<ul> <li>Agriculture</li> <li>Pasture, g</li> </ul>							
					Plantation		Other	
	Wetland		-					
	Not-cultiva	ated						
	Urban	aif.						
	Other, spe	CITY						
River w	vidth		_m	depth	al c	m		
Channel	Characterist	ics		Natura	al 🚬 🛛	reinforced		
Time sinc	e last rainfa	Ш		□ Man-r	made (ditch, irrig	bours	el)	
Stream fl			□ Dry	uays	□ Flowing		Flooding	
Bedrock I	ithology				_		0	
					specify			
Grain size	2			□ No	% sand-silt	0/	clasts > 2 mm	
	, observed gr	oundwate	r table (cm)	)		/		
	interval from	n surface	□ 0-10	cm ( <b>1</b> )	_	Geochem	Sieved? < _	microns
	For heavy	minerals:	□ 0-10	• • •				microns
				0 cm ( <b>3</b> ) 0 cm ( <b>4</b> )				microns microns
				, specify: (		Tieavy Will		microns
Possible	sources of c	ontamina	tion, specify	/				-
OTHER	Field pH	Top			Munsell Colour	- Dry	Top	· · · · · · · · · · · · · · · · · · ·
		Bottom					Bottom	
	Texture	Top Bottom				- Moist	Top Bottom	
						BUGTOO		····
NUMBER OF	SAMPLE E	SAGS				PHOTOS		
COMMENTS								

Appendix 3: Practical detection limits for XRF and ICP-MS, Minerals Division Laboratory, Geoscience Australia

## ANALYTES WITH DETECTION LIMITS (as reported) FOR SAMPLES LIKELY TO CONTAIN NaCl

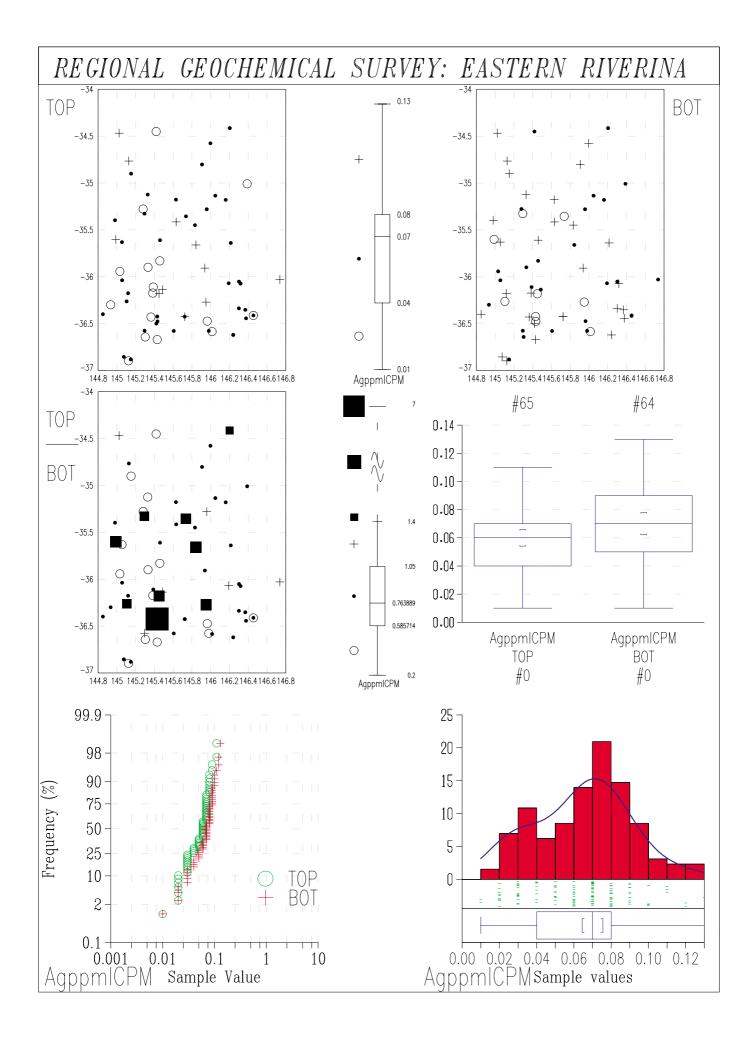
XRF MAJOR ELEMENTS								
Al <sub>2</sub> O <sub>3</sub>	0.001	%						
CaO	0.002	%						
Fe <sub>2</sub> O <sub>3</sub>	0.002	%						
K <sub>2</sub> O	0.002	%						
MgO	0.004	%						
MLOI	0.001	%						
MnO	0.001	%						
Na <sub>2</sub> O	0.004	%						
$P_2O_5$	0.001	%						
SiO <sub>2</sub>	0.006	%						
SO <sub>3</sub>	0.001	%						
TiO <sub>2</sub>	0.002	%						
XRF TRACE I	ELEMENTS							
(determined on	fused disc)	-						
Cl	5	ppm						
Cr	32	ppm						
Cu	2	ppm						
F	50	ppm						
Ni	2	ppm						
Sc	2	ppm						
V	4	ppm						
Zn	2	ppm						

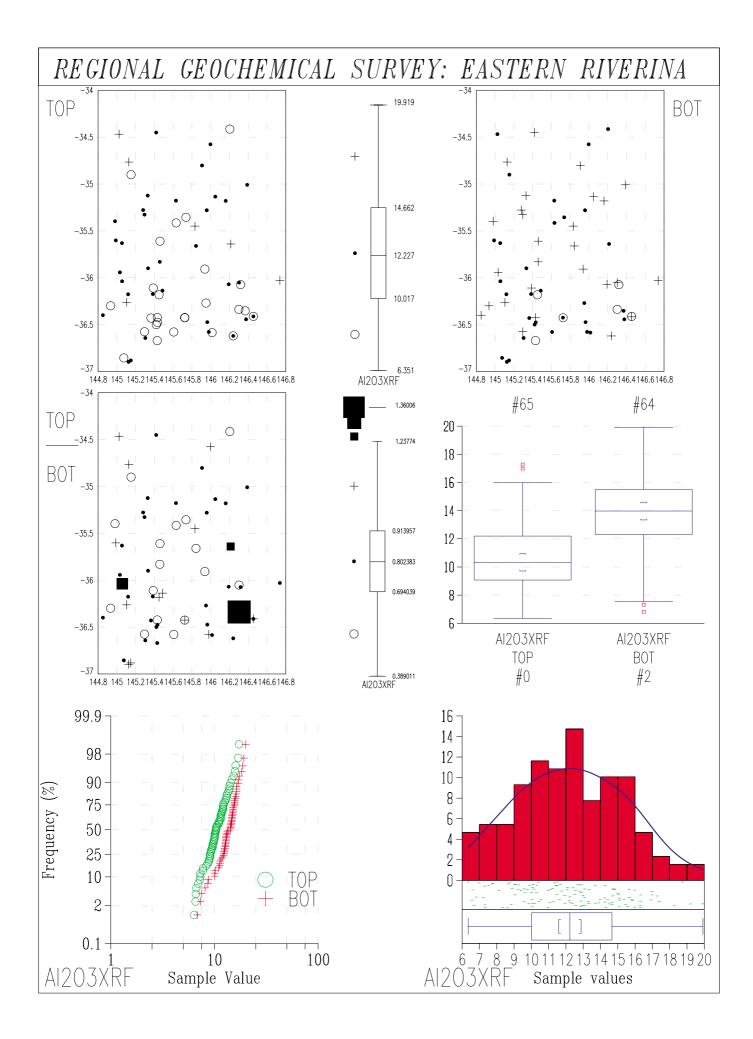
ICP-MS TRACE ELEMENTS							
Ag	0.01	ppm					
As	0.01						
Ba	2	ppm ppm					
Be	0.05						
Bi	0.03	ppm					
Cd		ppm					
	0.05	ppm					
Ce		ppm					
Cs	0.01	ppm					
Dy	0.01	ppm					
Er	0.01	ppm					
Eu	0.50	<u>ppb</u>					
Ga	0.1	ppm					
Gd	0.01	ppm					
Ge	0.02	ppm					
Hf	0.01	ppm					
Но	0.01	ppm					
La	0.02	ppm					
Lu	0.01	ppm					
Мо	0.01	ppm					
Nb	0.04	ppm					
Nd	0.01	ppm					
Pb	0.5	ppm					
Pr	0.01	ppm					
Rb	1.0	ppm					
Sb	0.01	ppm					
Sm	0.01	ppm					
Sn	0.01	ppm					
Sr	1.0	ppm					
Та	0.06	ppm					
Tb	0.01	ppm					
Th	0.01	ppm					
T1	0.01	ppm					
Tm	0.01	ppm					
U	0.07	ppm					
Y	0.22	ppm					
Yb	0.01	ppm					
Zr	1.0	ppm					
<i>L</i> 1	1.0	PPm					

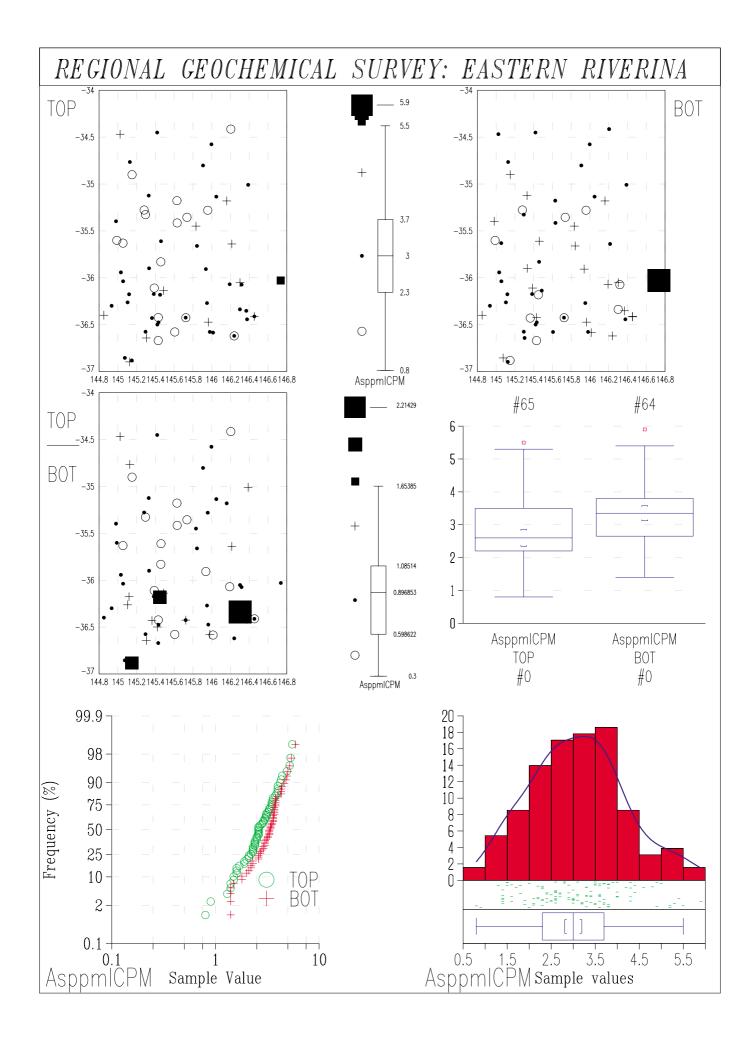
Appendix 4: Statistical and geochemical distributions of bulk (<180  $\mu$ m) overbank sediments from the eastern Riverina area. Where more than one method was used for a particular element, the preferred method results only are shown here. All values <DL (detection limit) were set to 1/2 DL before statistical manipulation.

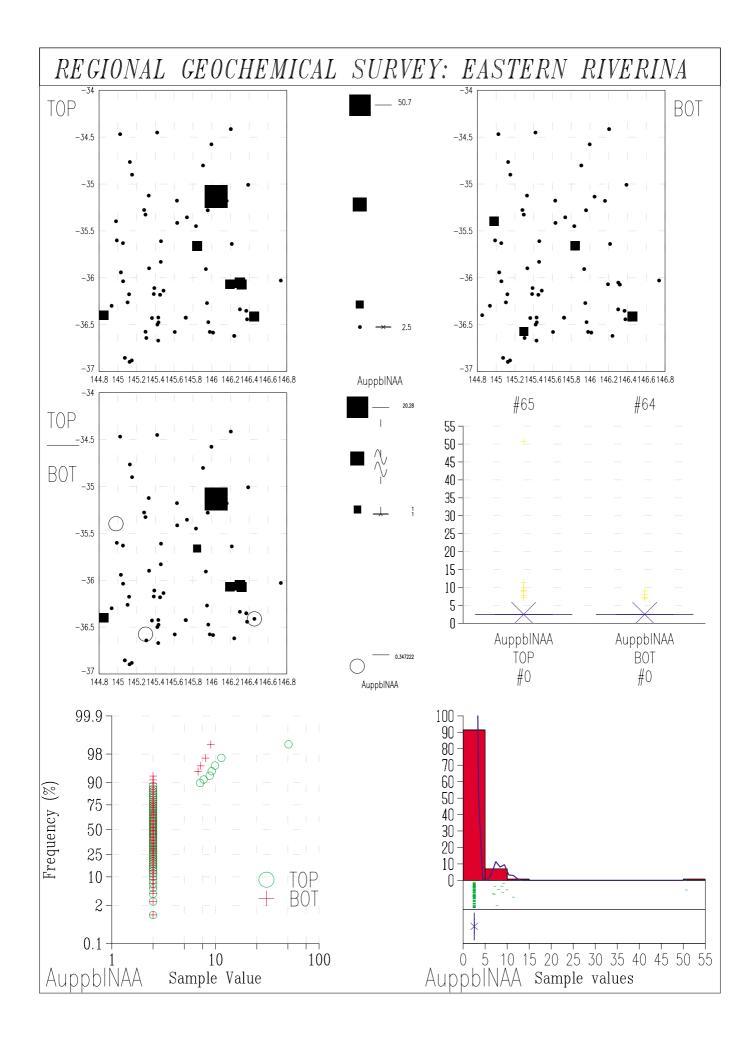
**Diagrams shown are:** 

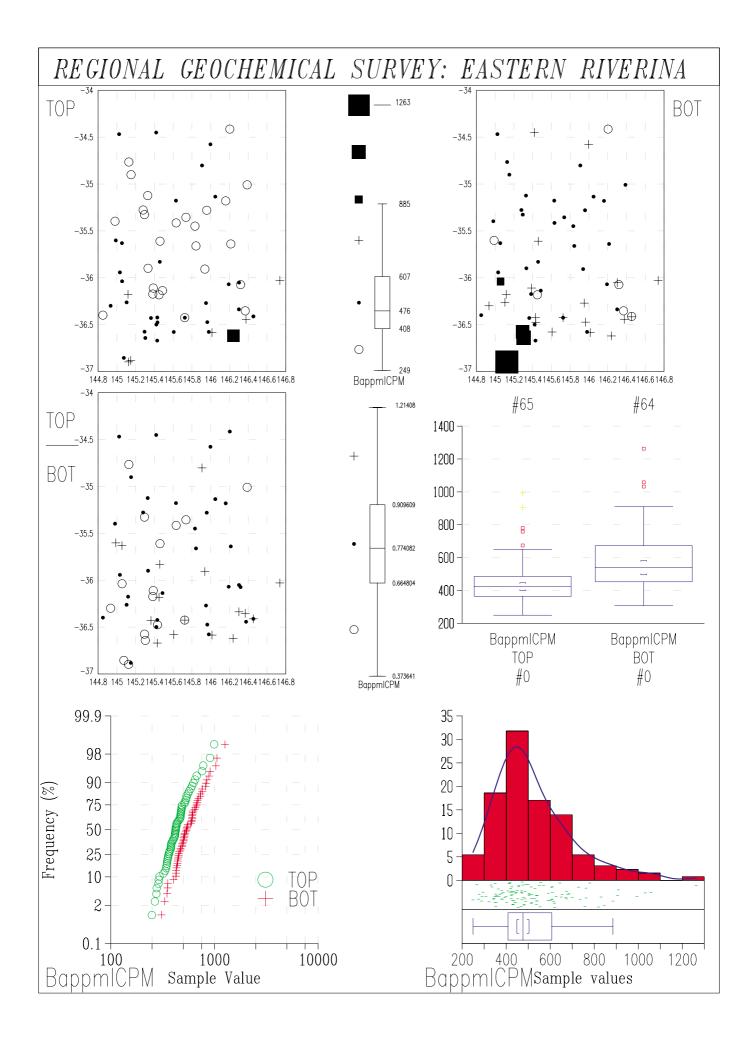
- **1.** Distribution map of element concentration in TOP overbank sediment, using boxplot classification of both TOP and BOT values
- 2. Distribution map of element concentration in BOT overbank sediment, using same boxplot classification
- 3. Distribution map of TOP/BOT ratio in overbank sediment
- 4. Boxplot distribution of TOP and BOT overbank sediments
- 5. Cumulative frequency diagrams of TOP and BOT overbank sediments
- 6. Histogram, density curve, scatter and boxplot of element concentration in TOP and BOT overbank sediment

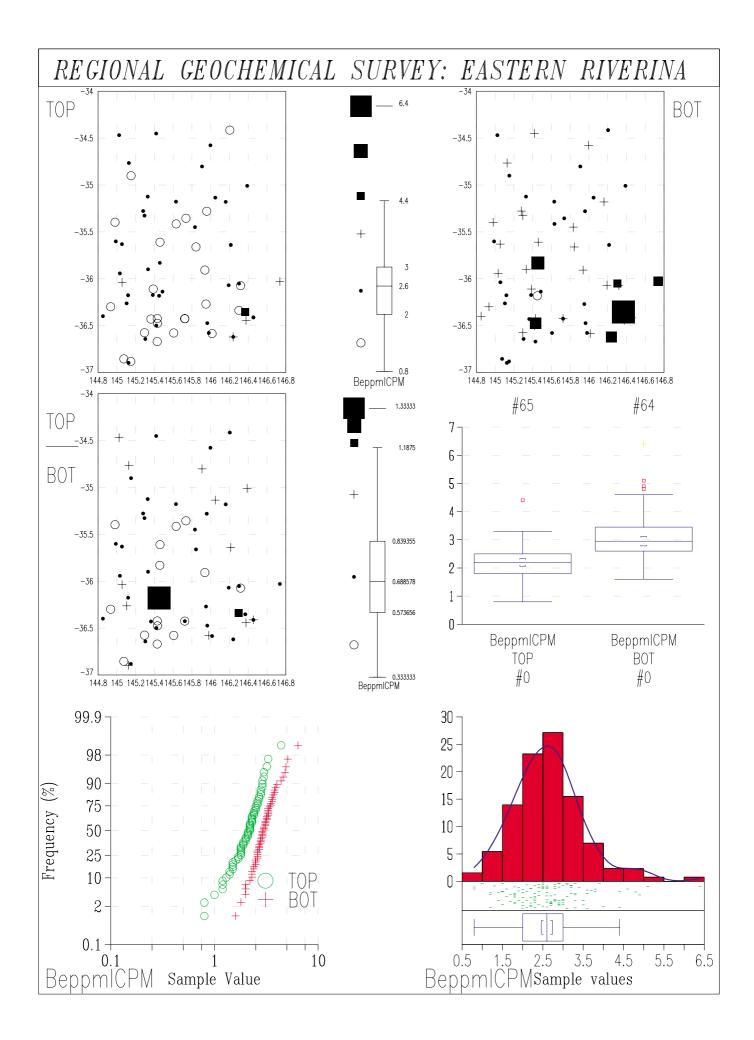


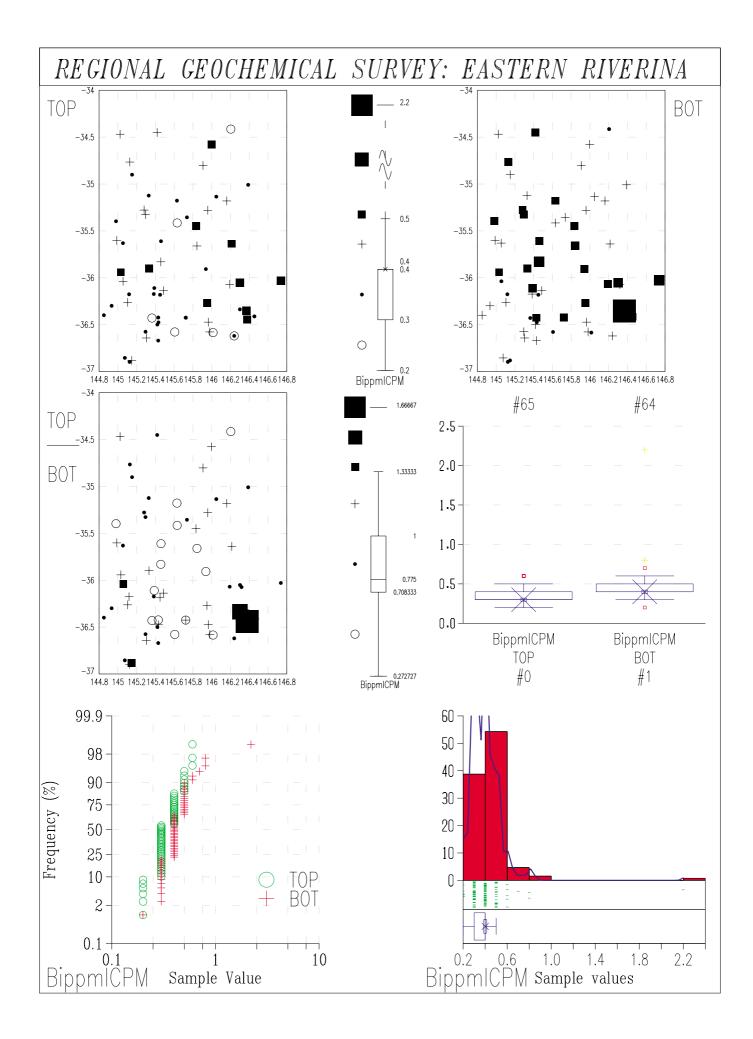


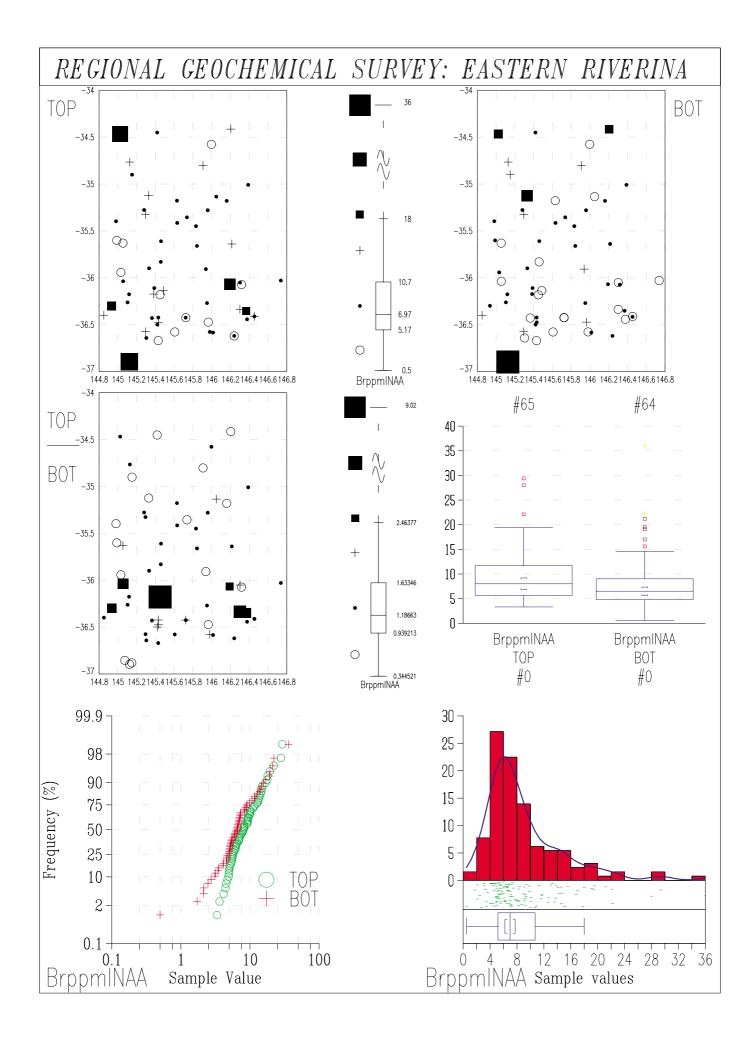


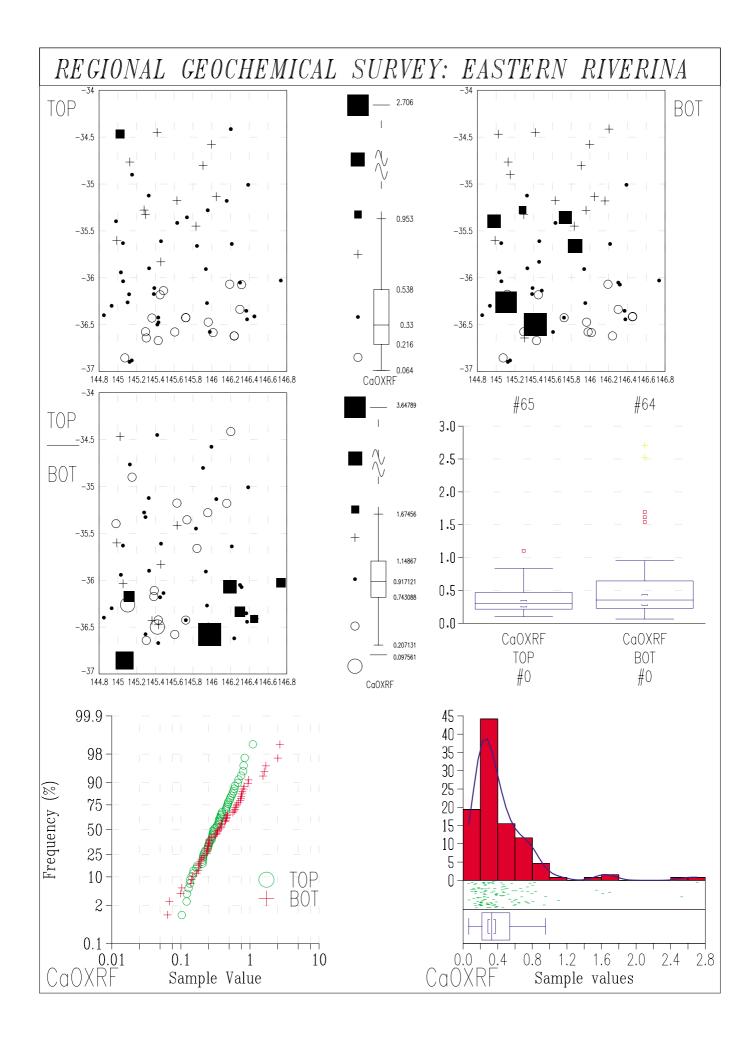


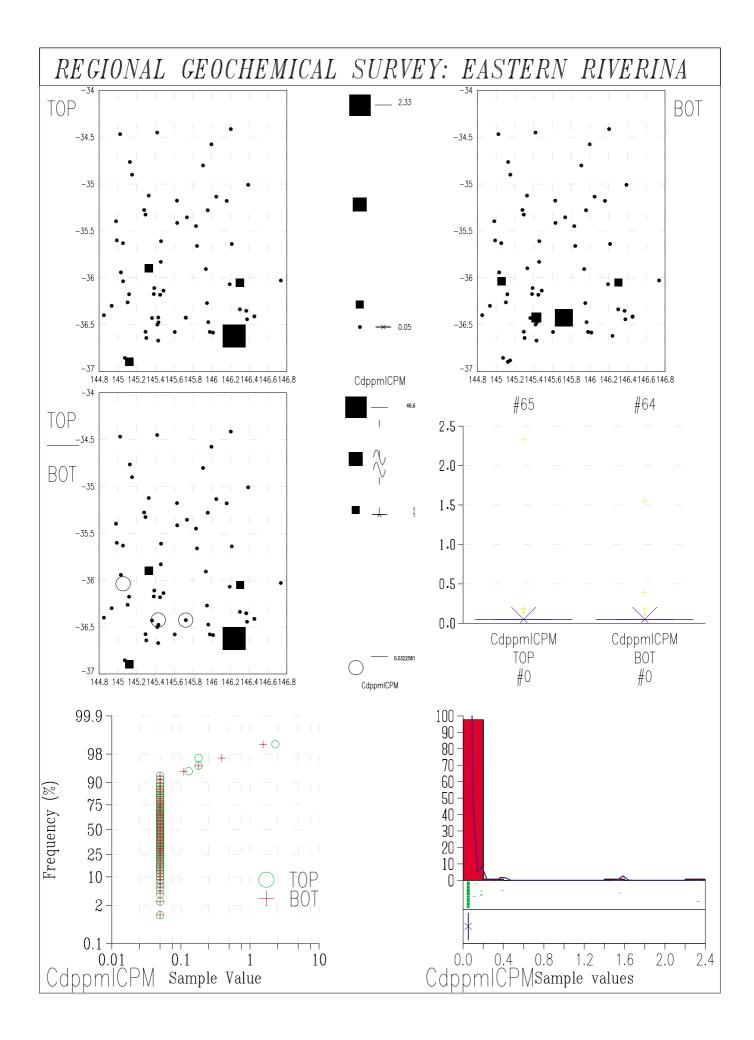


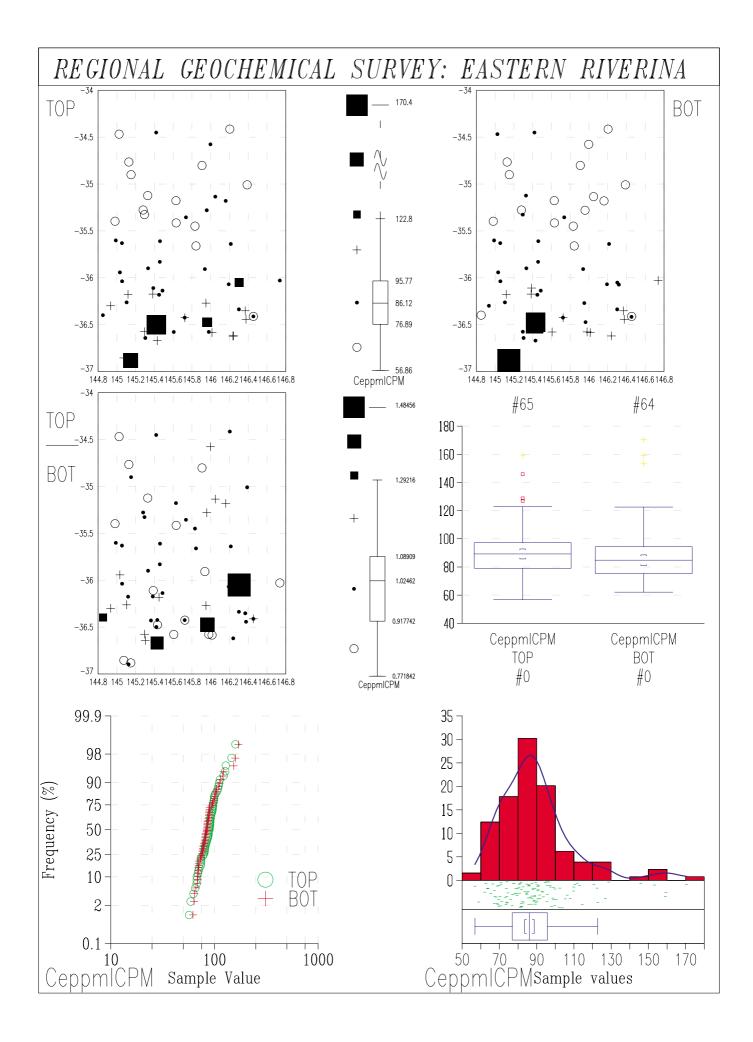


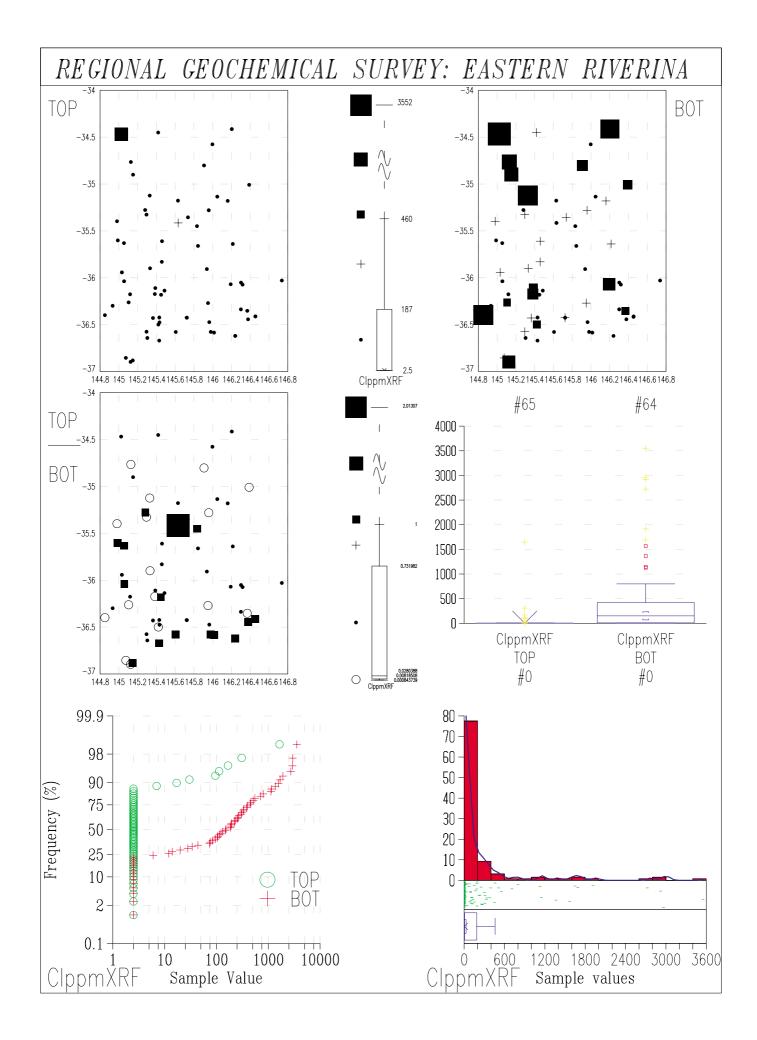


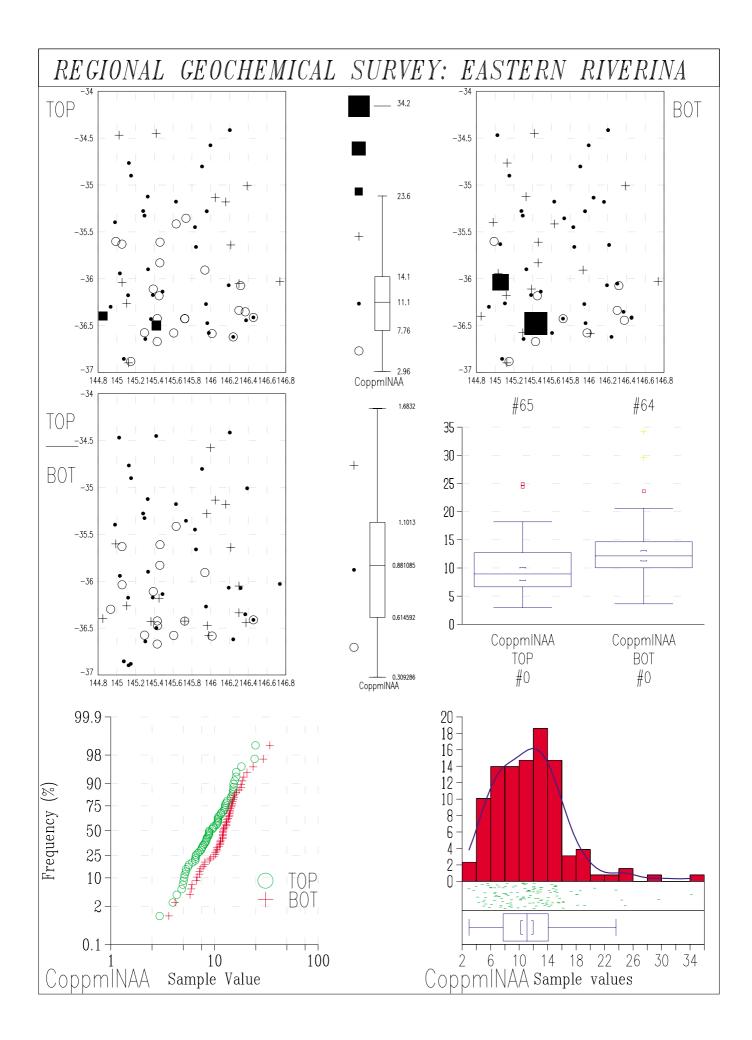


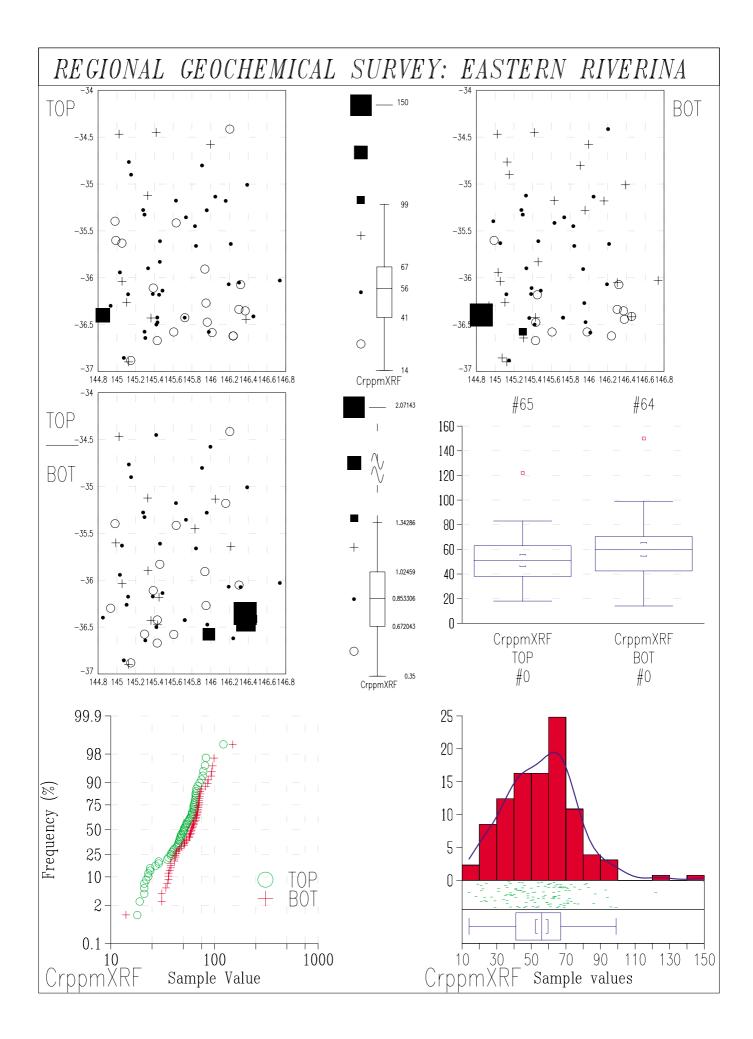


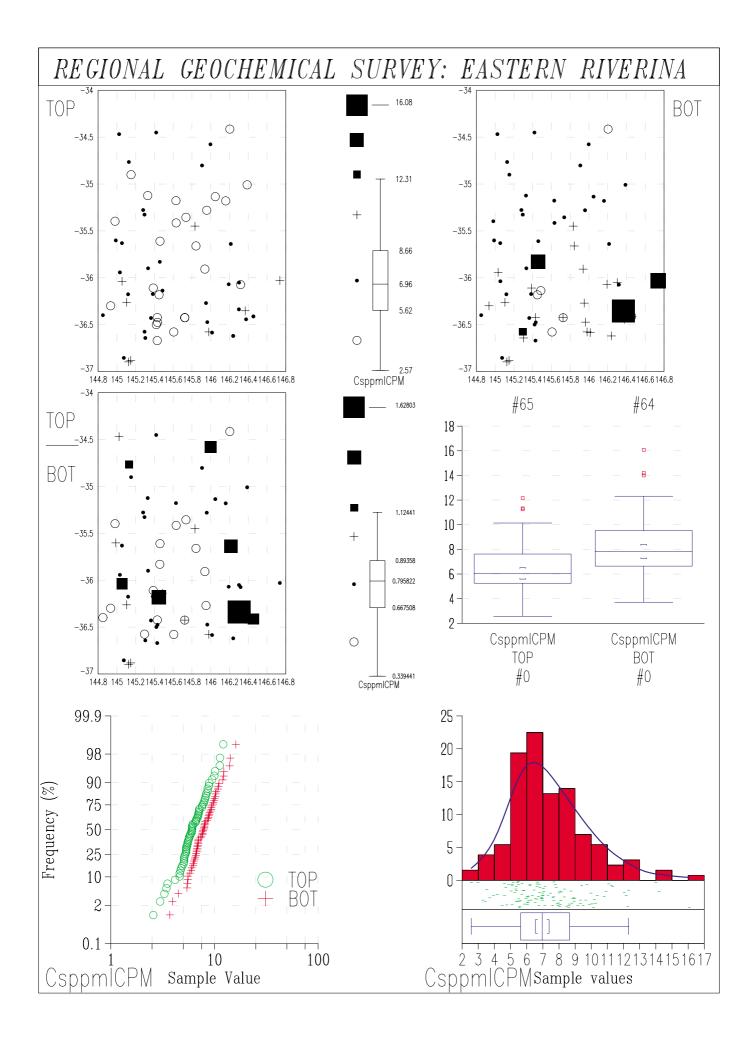


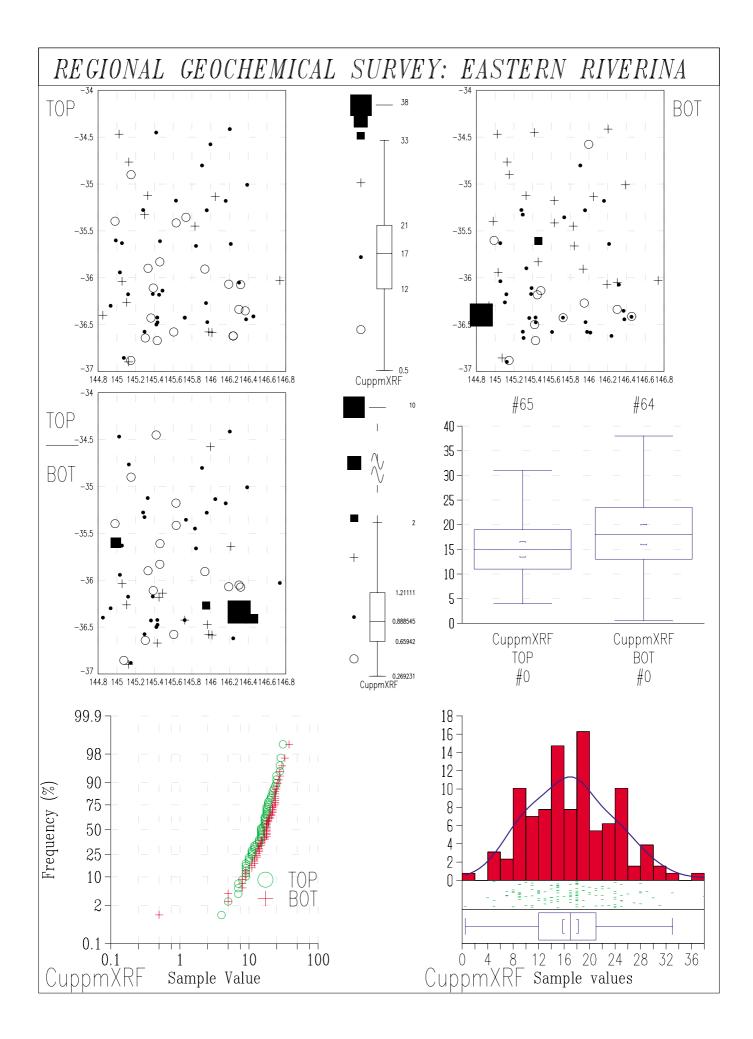


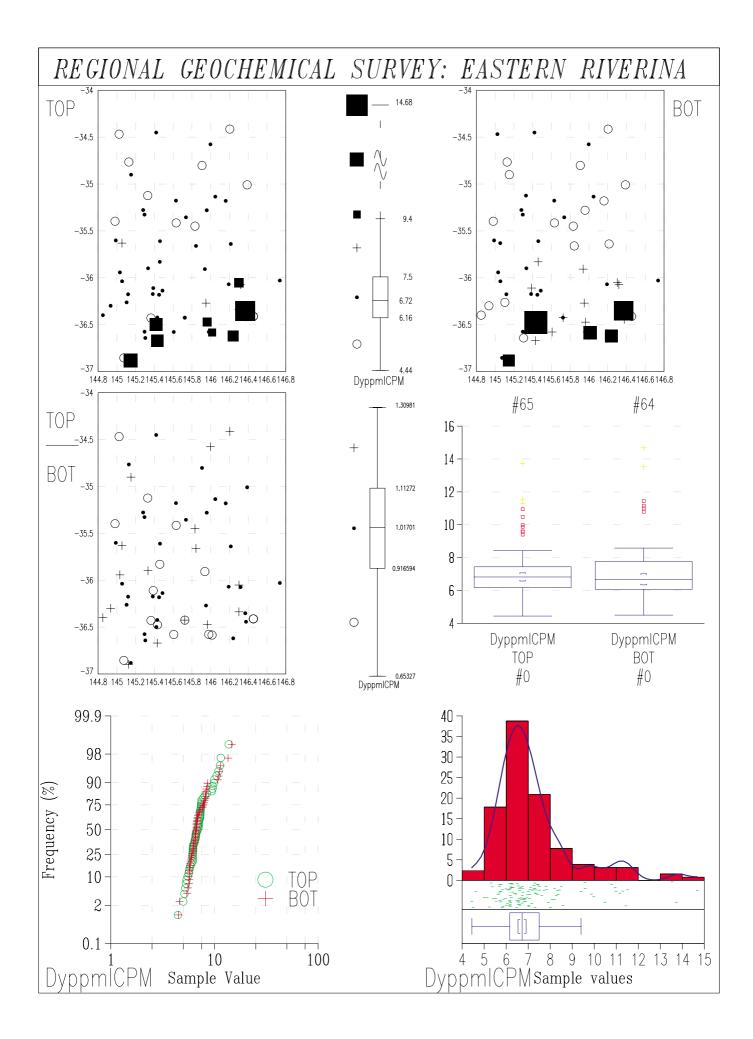


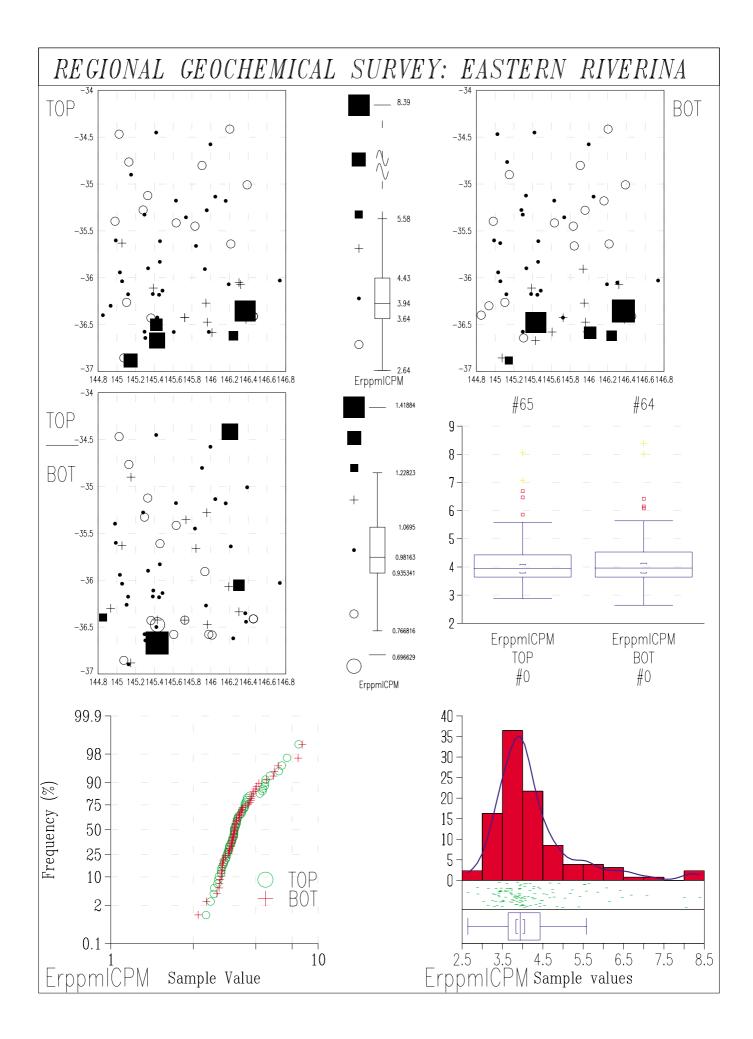


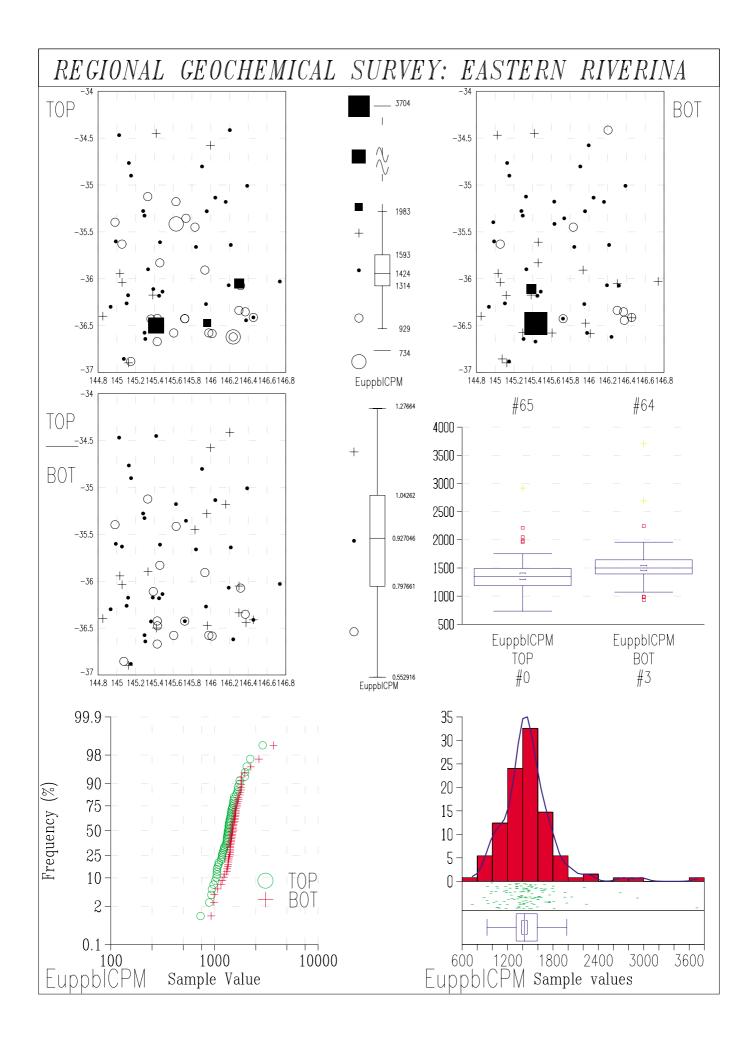


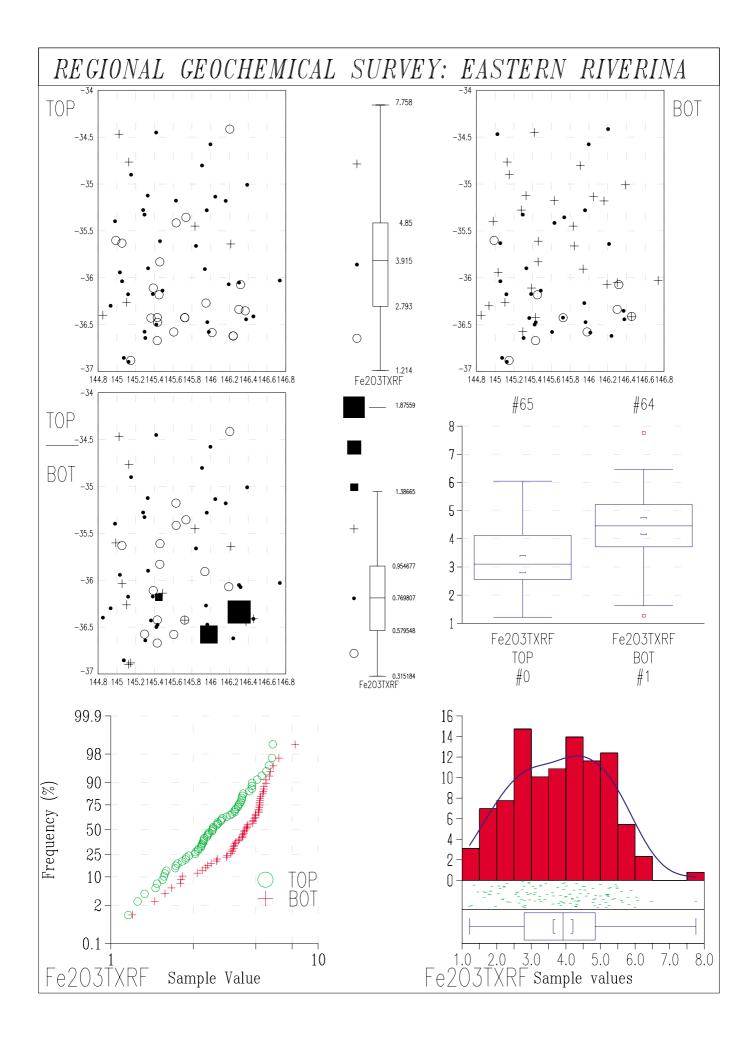


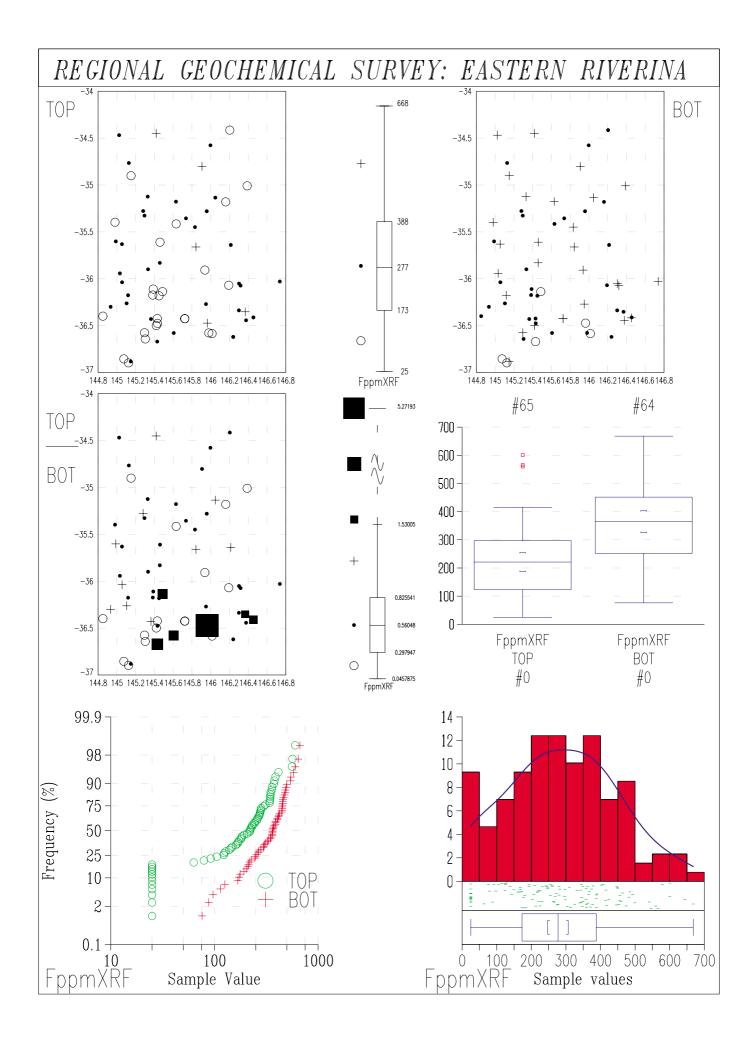


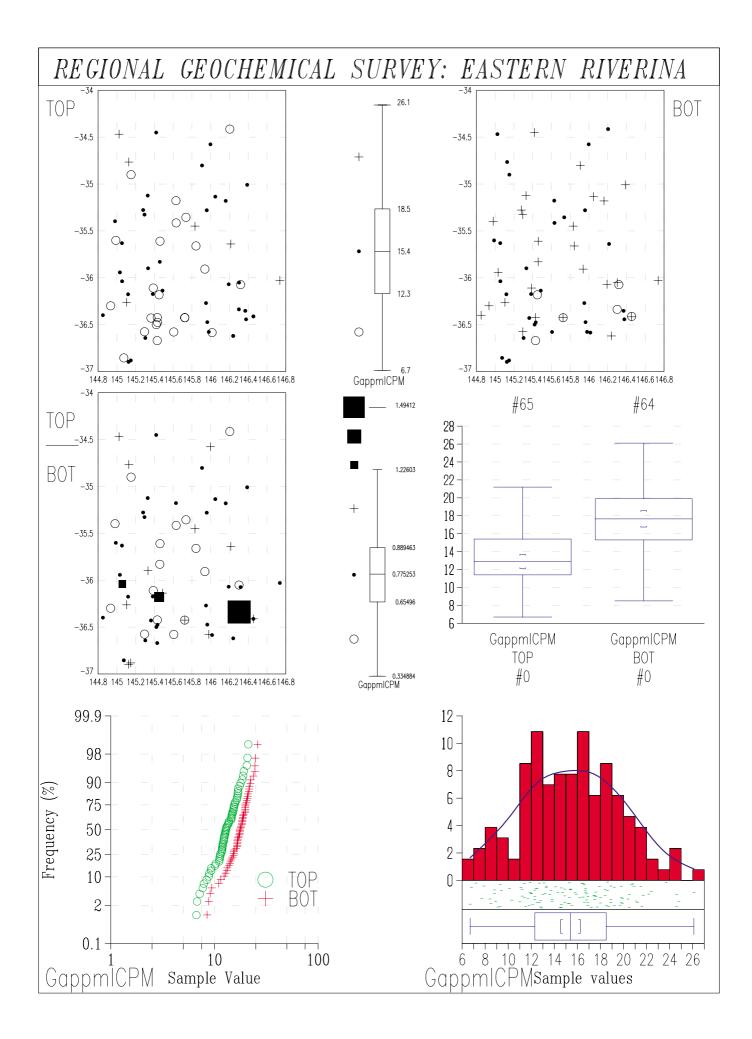


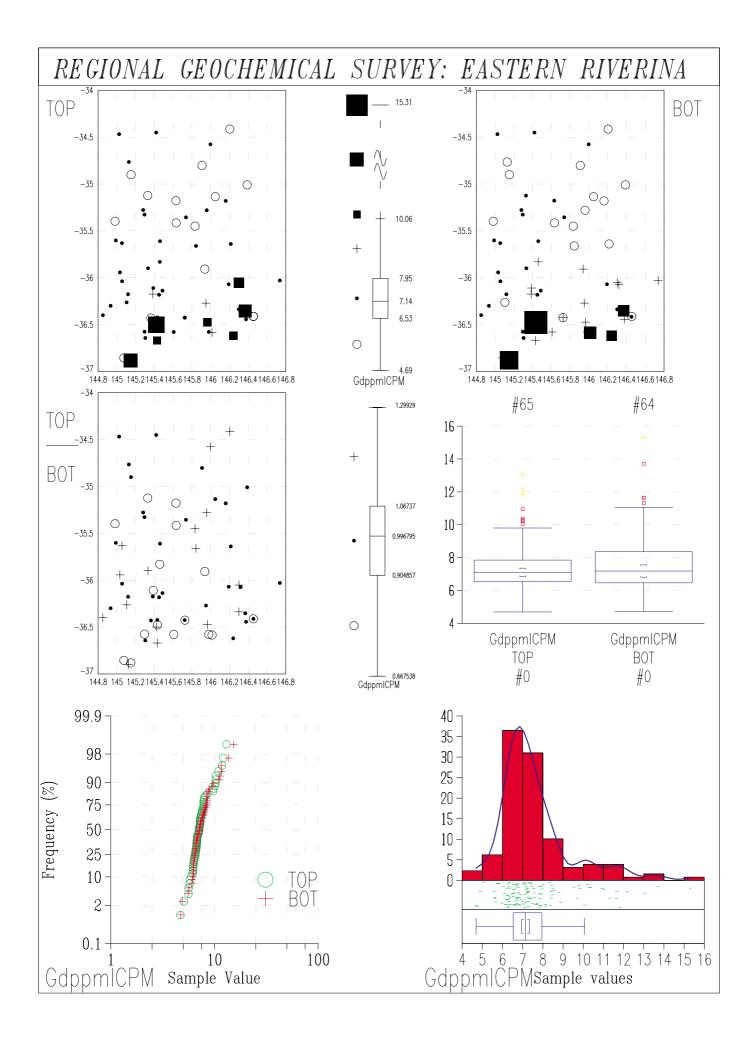


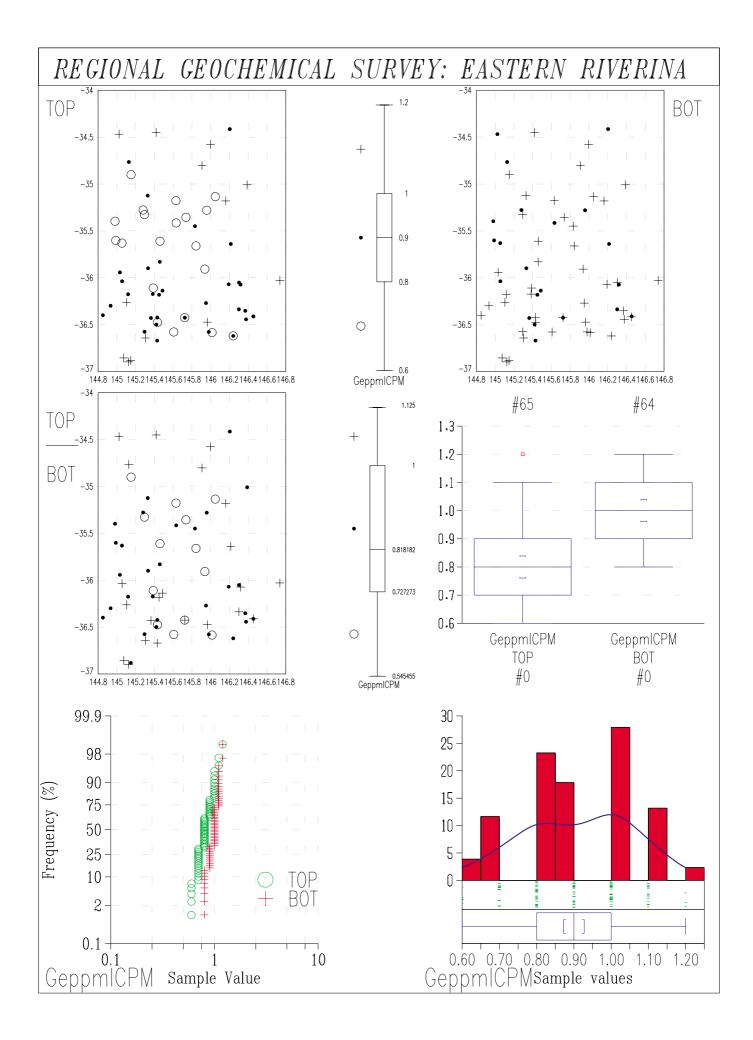


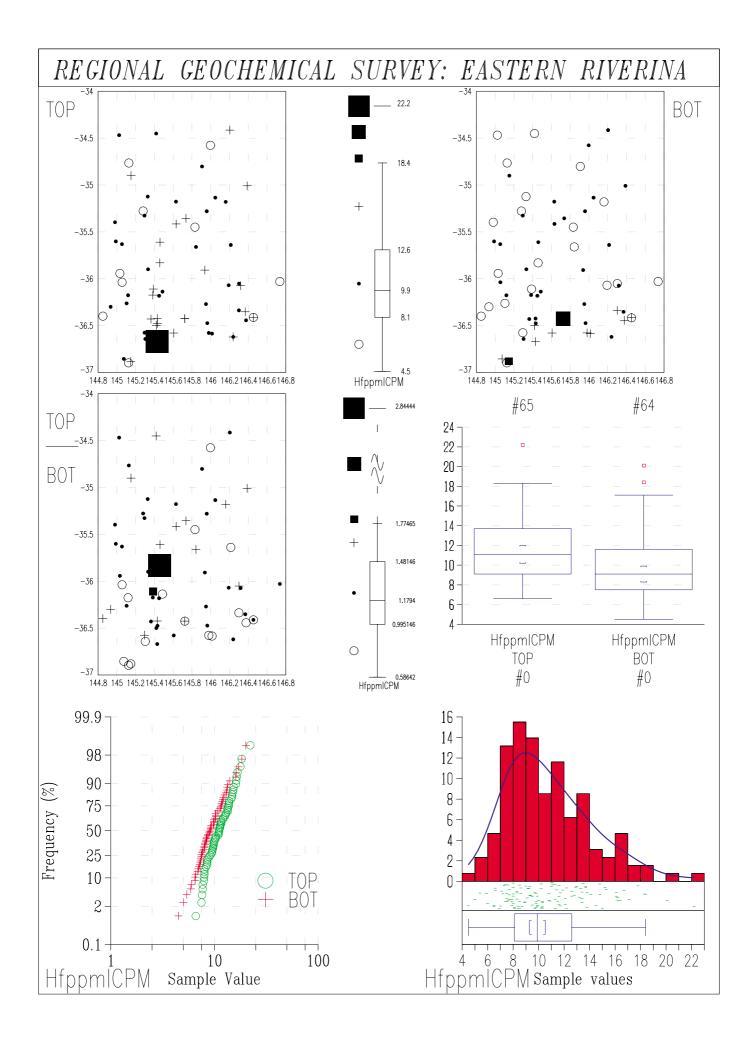


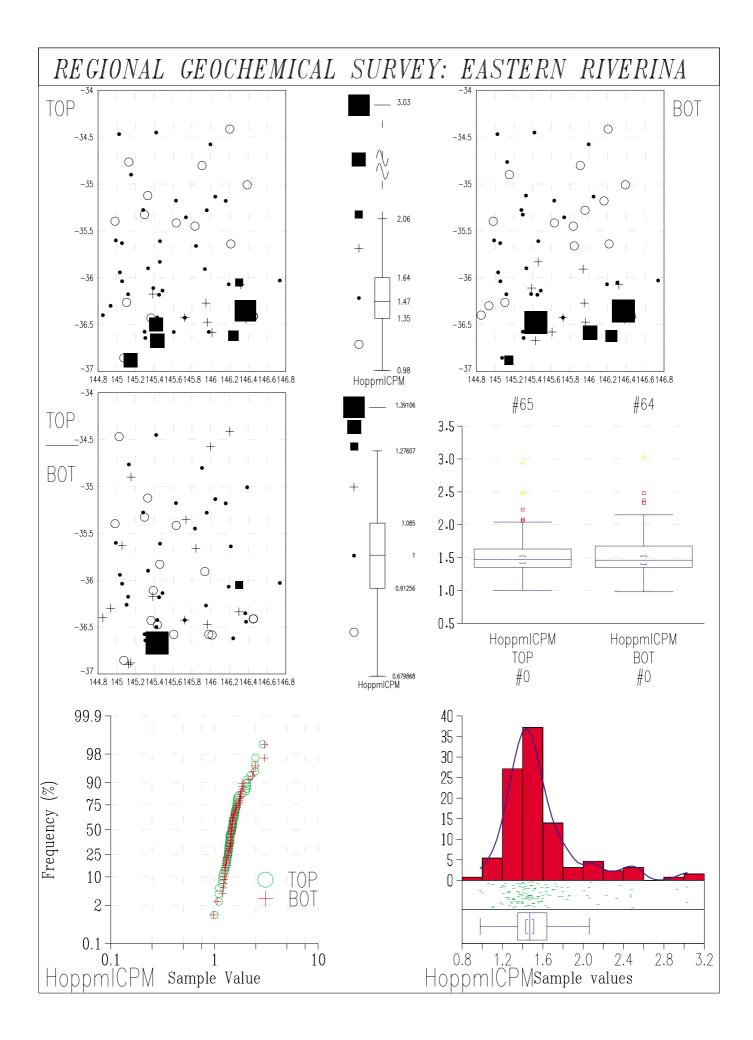


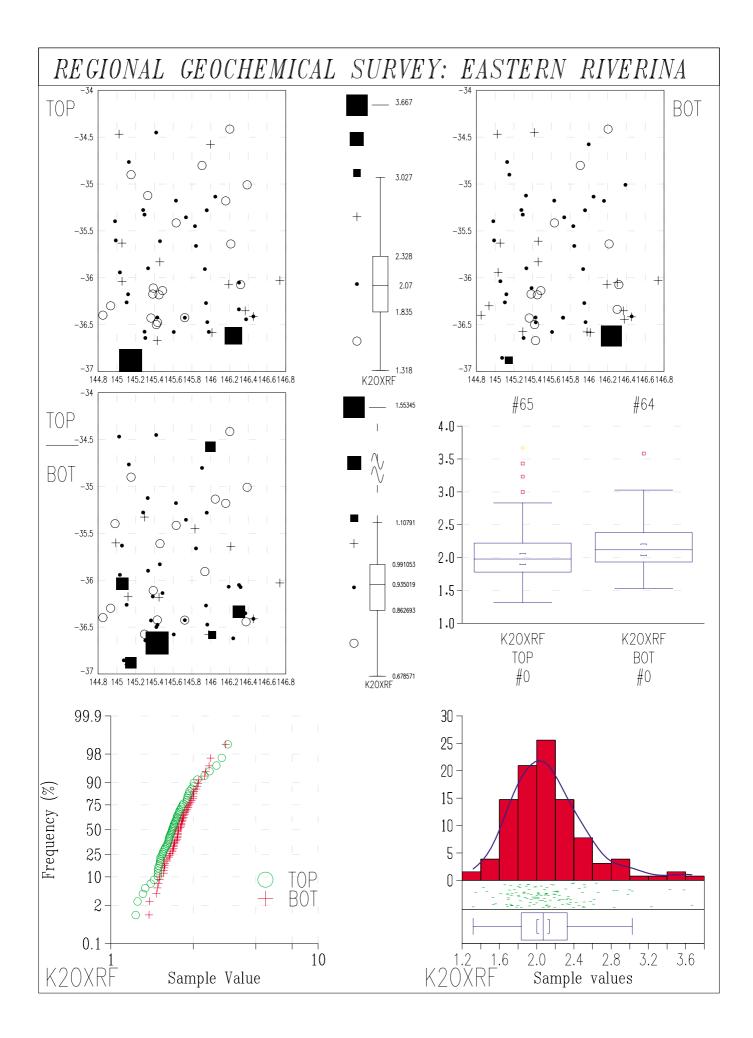


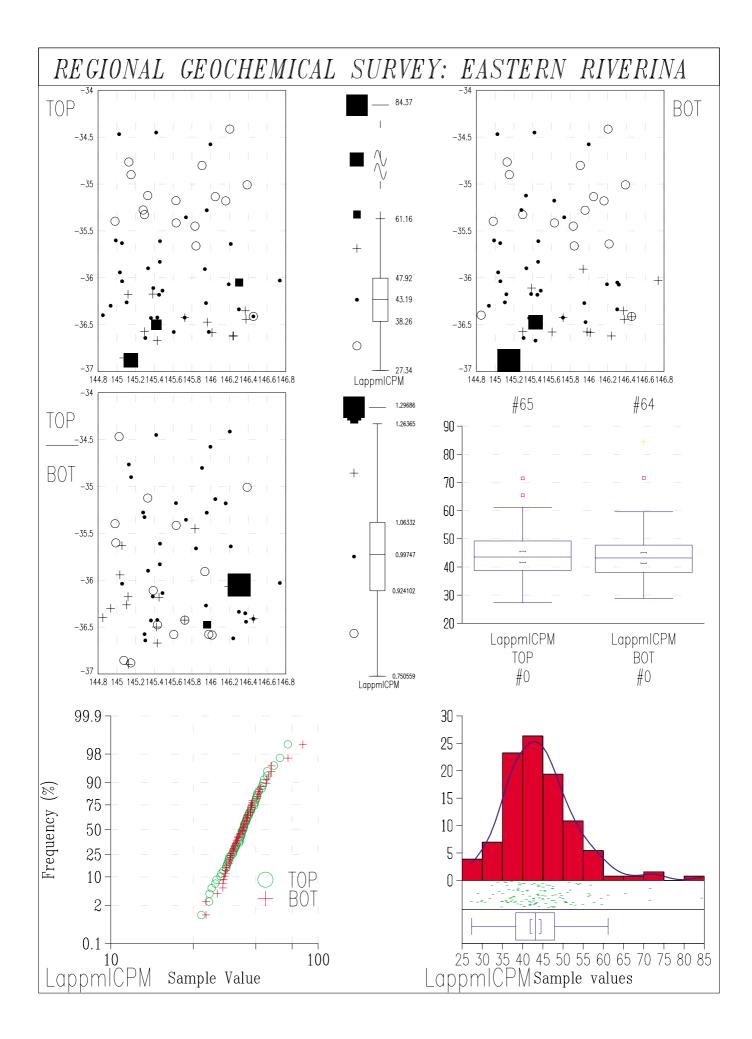


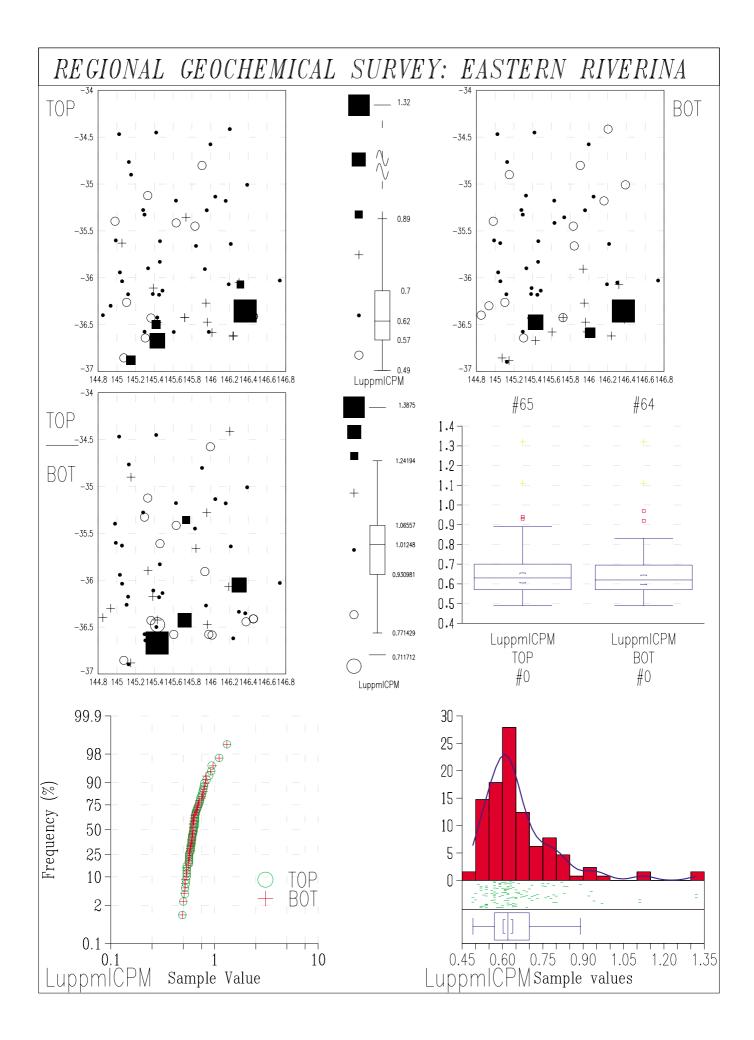


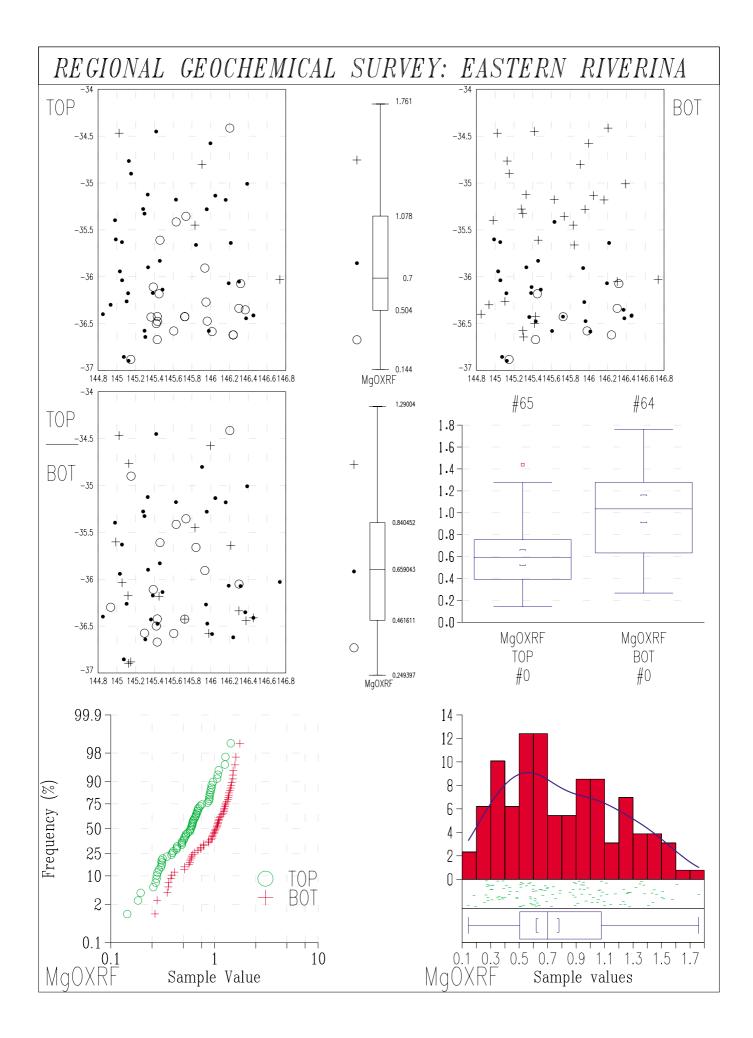


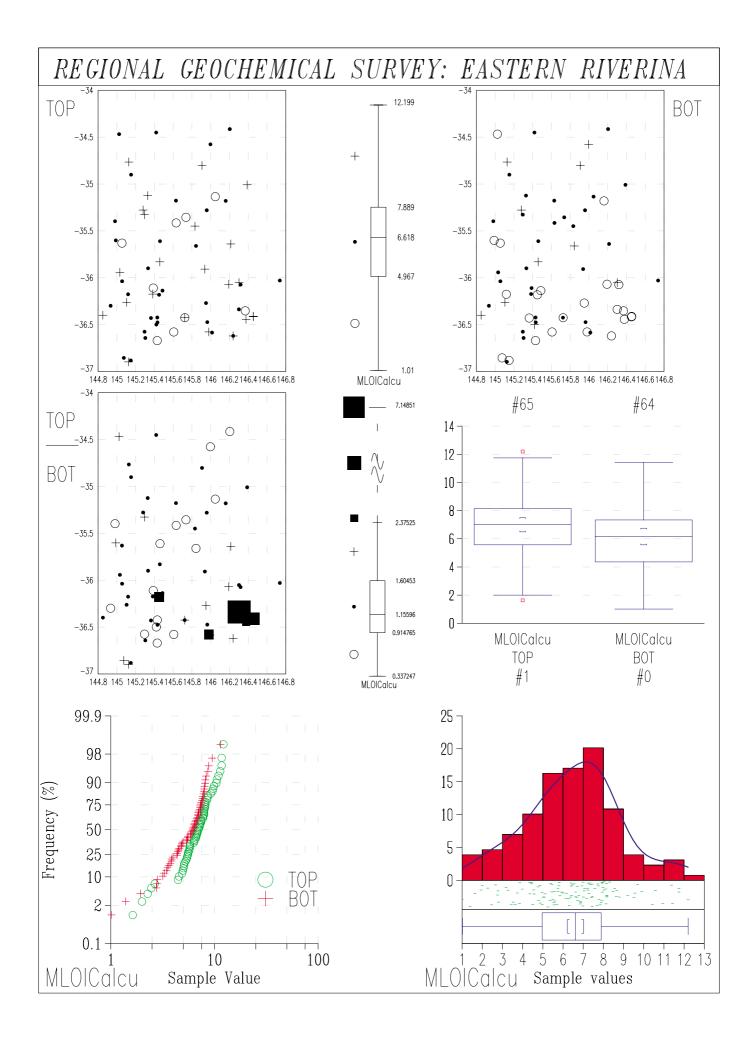


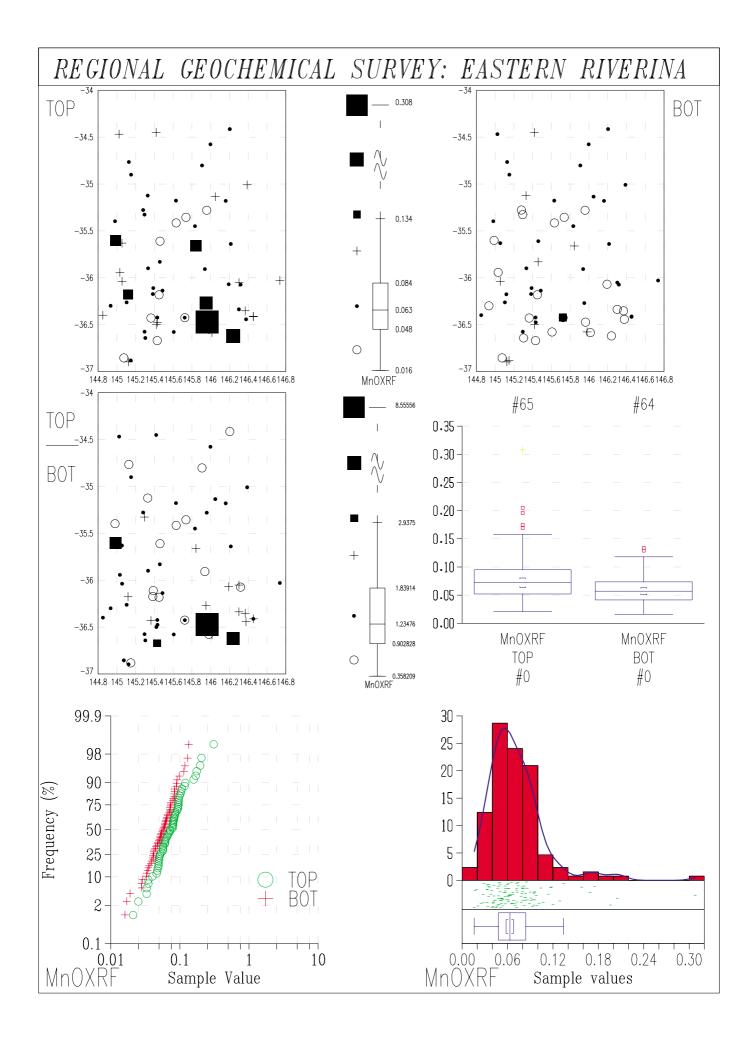


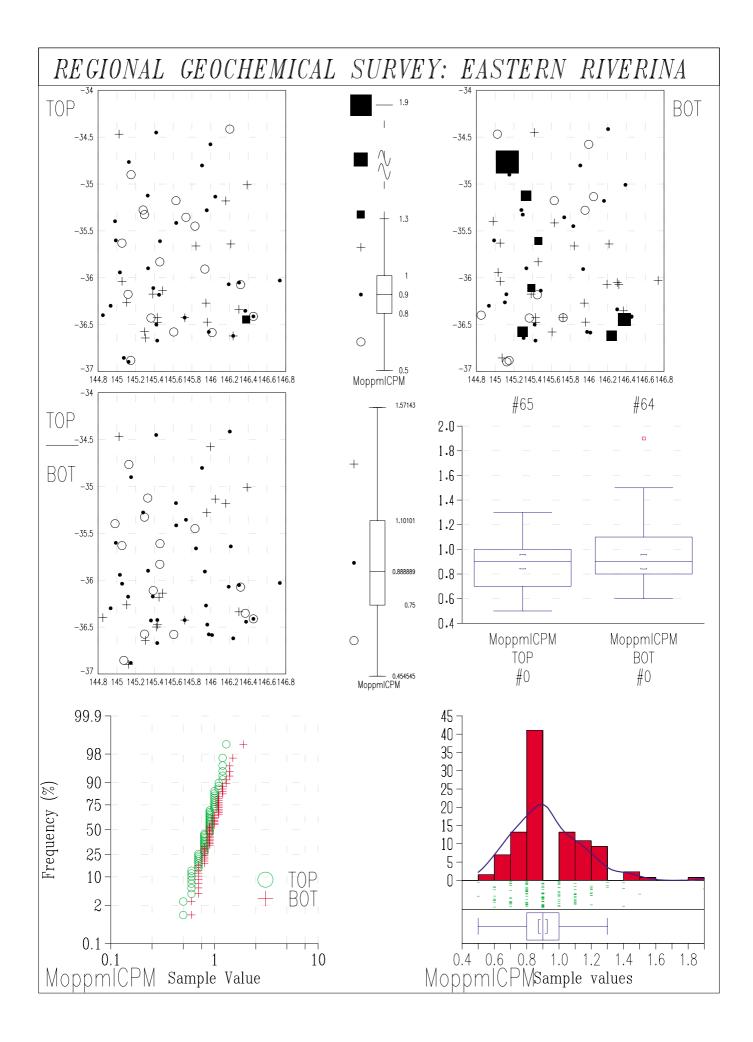


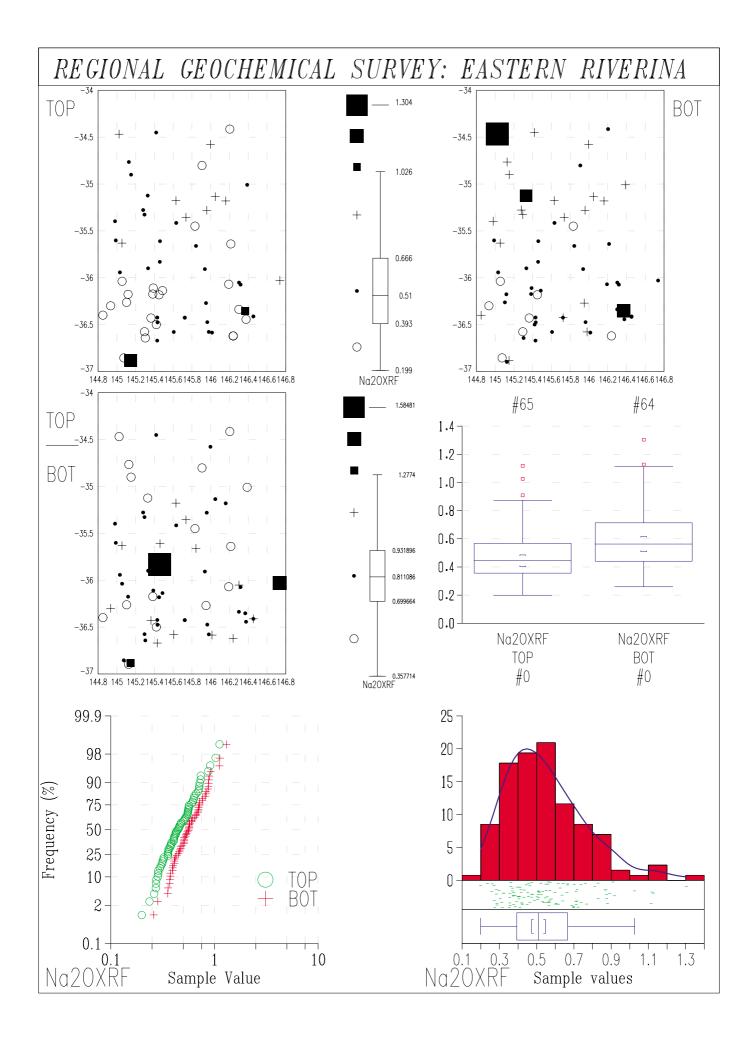


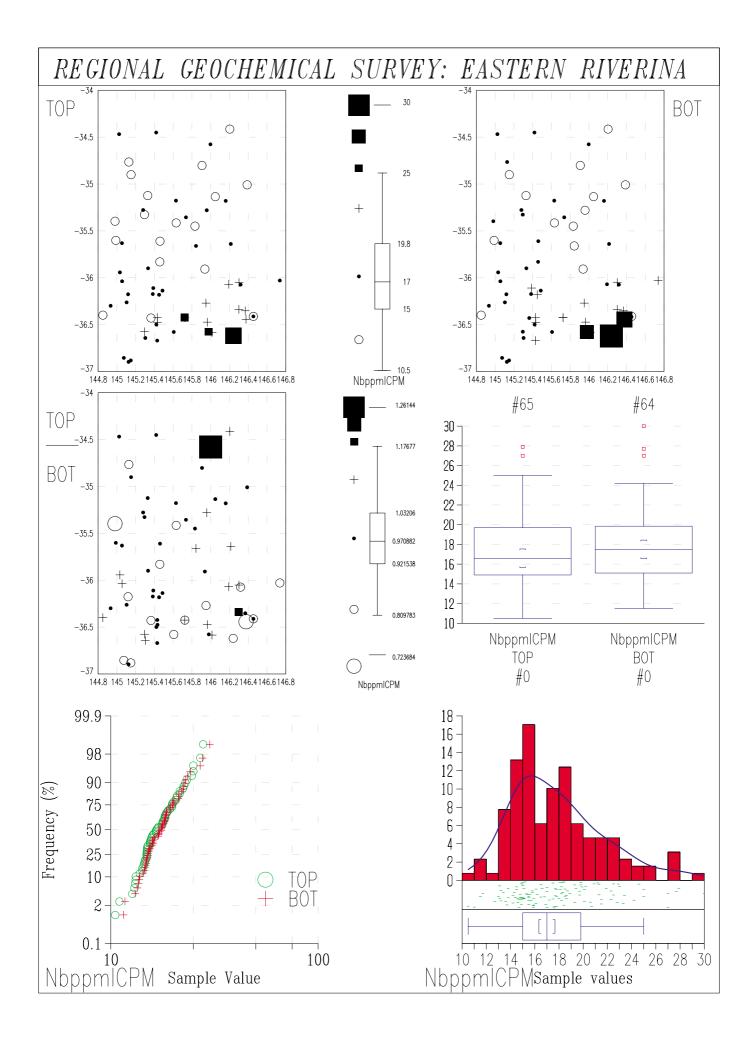


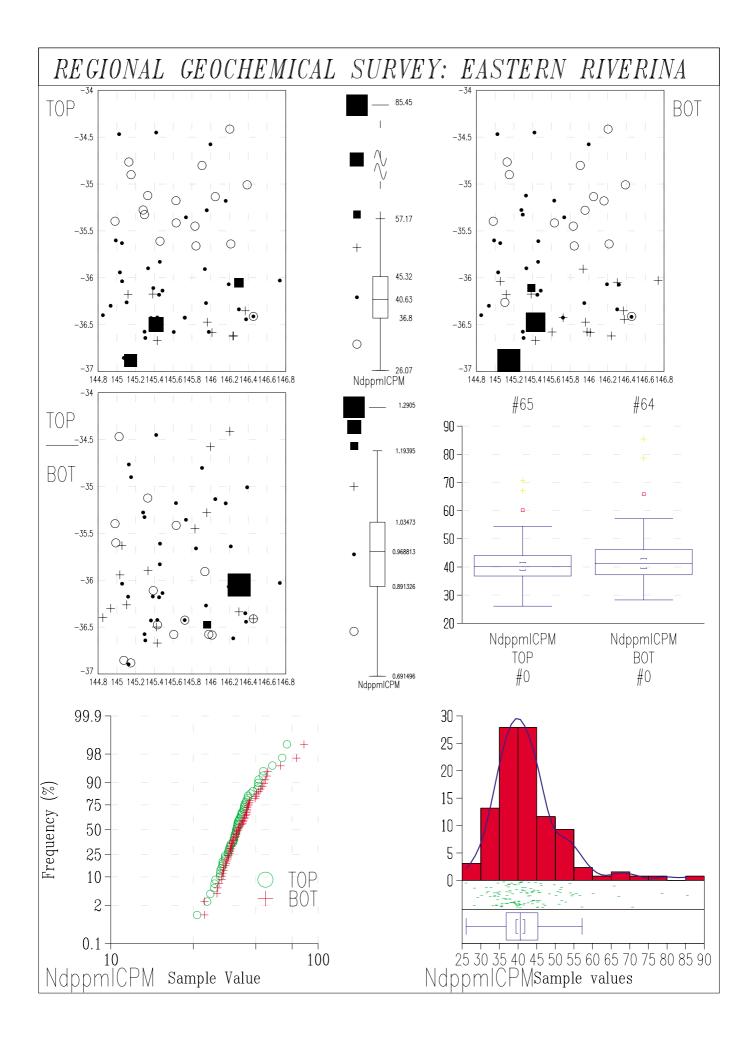


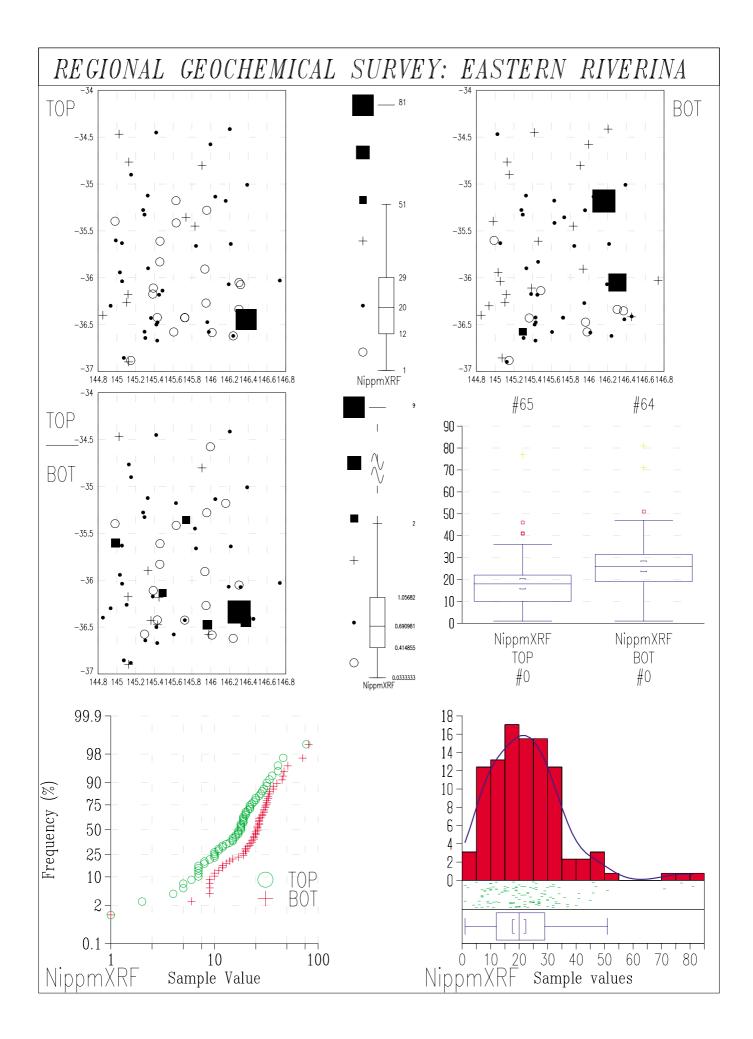


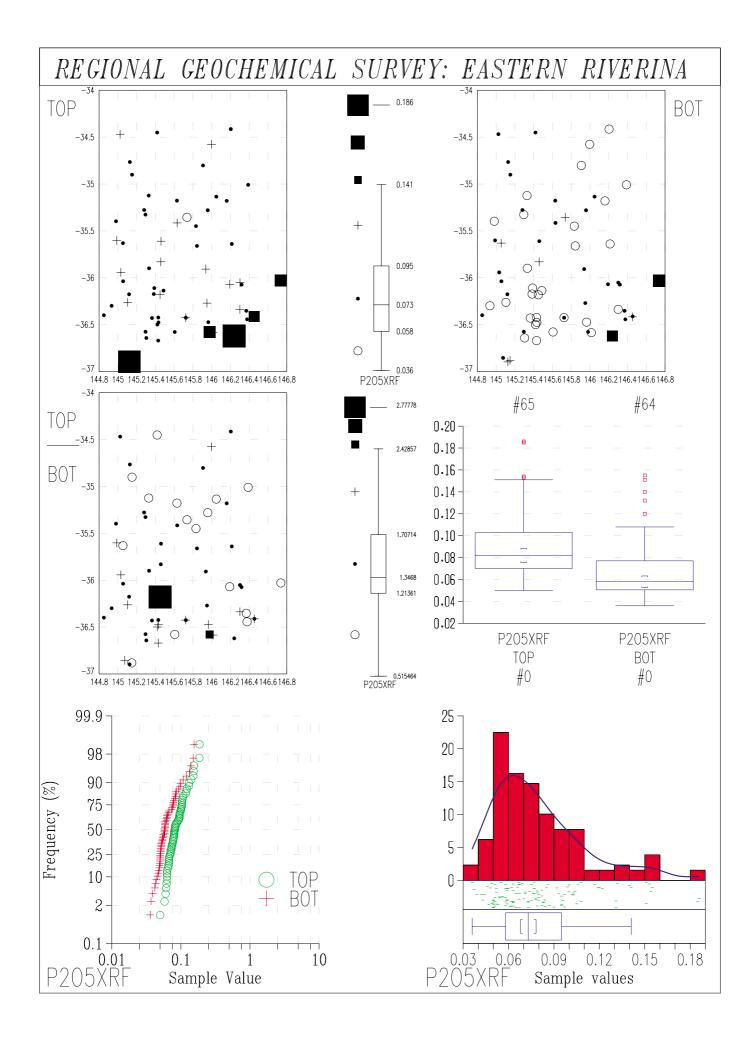


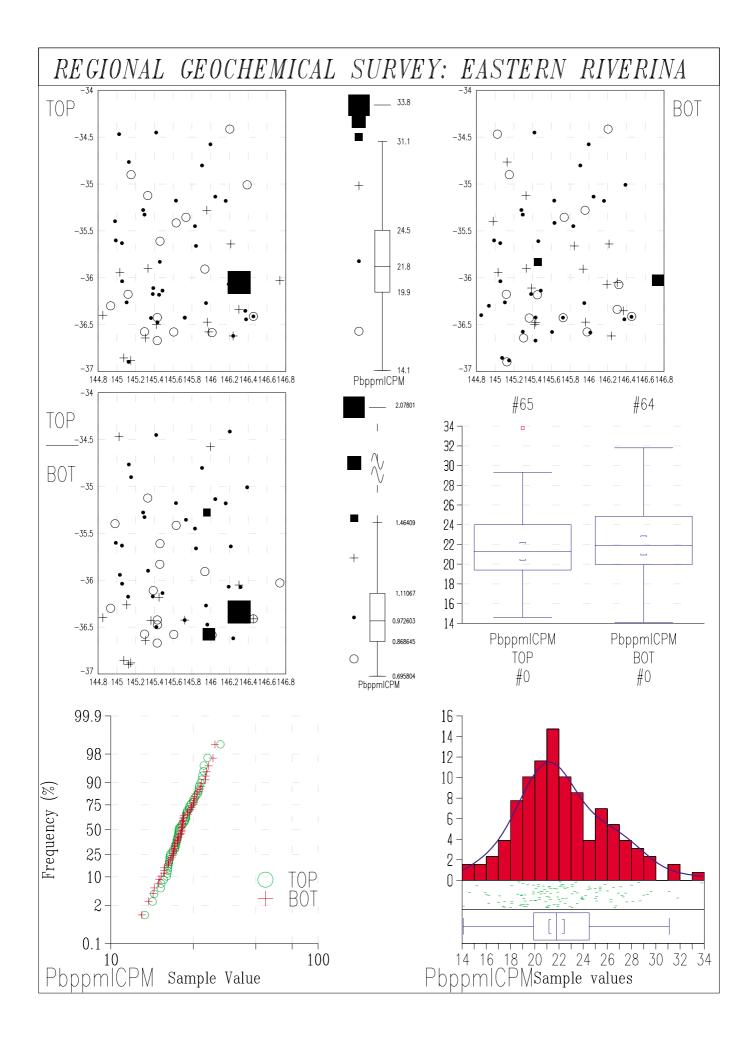


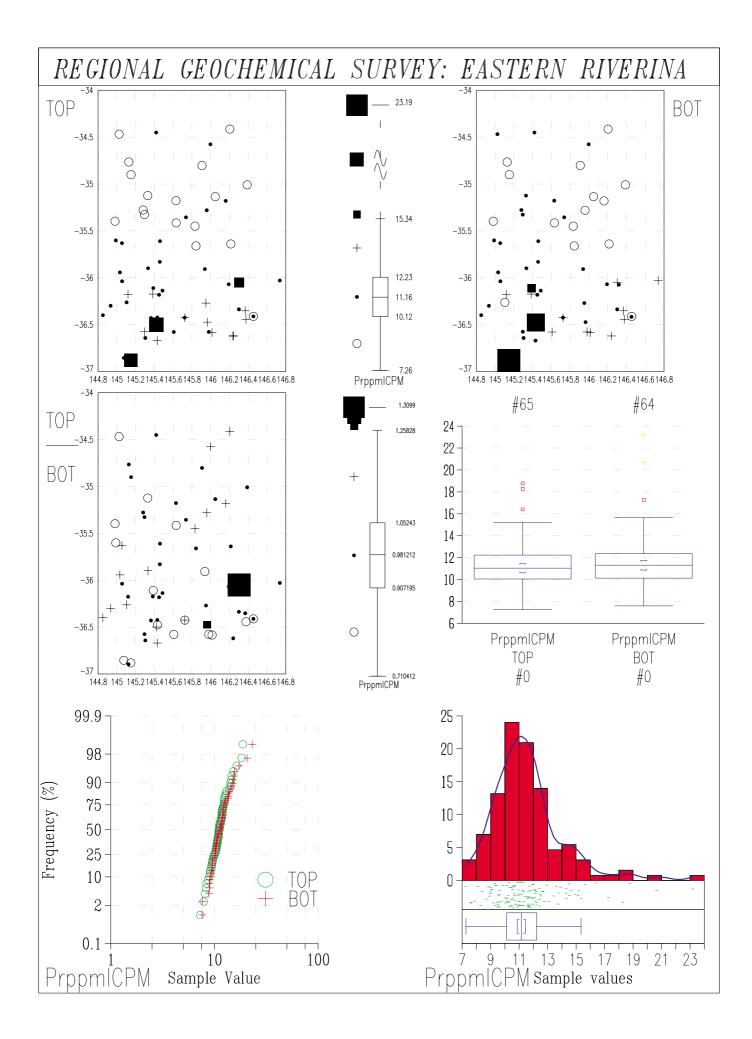


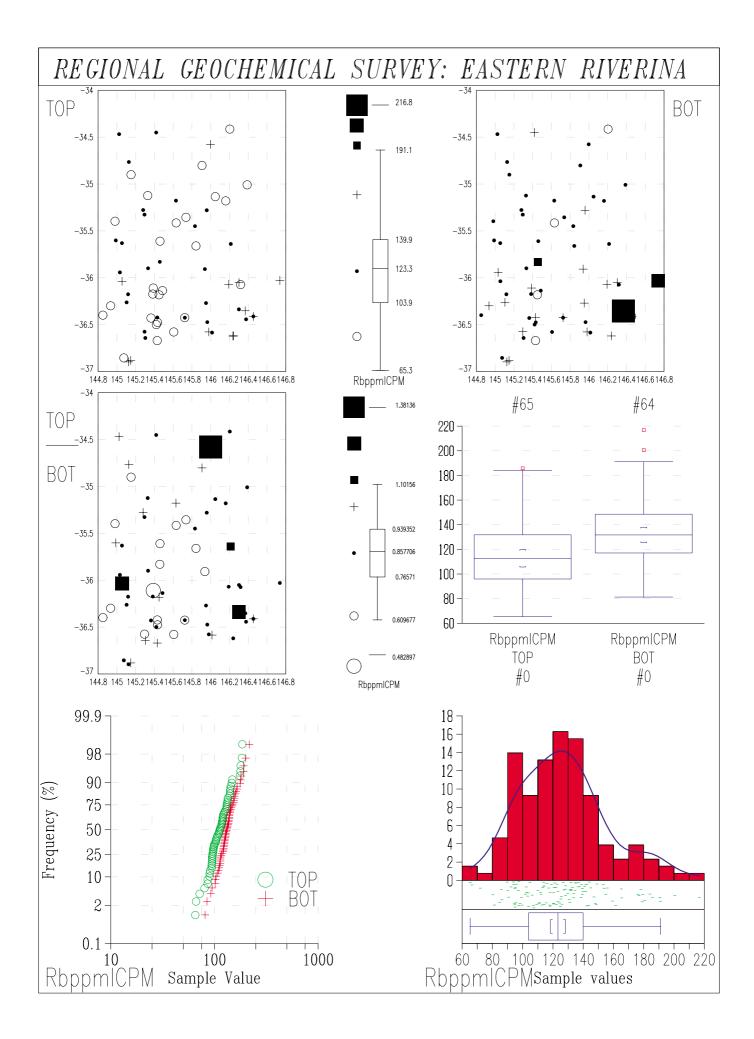


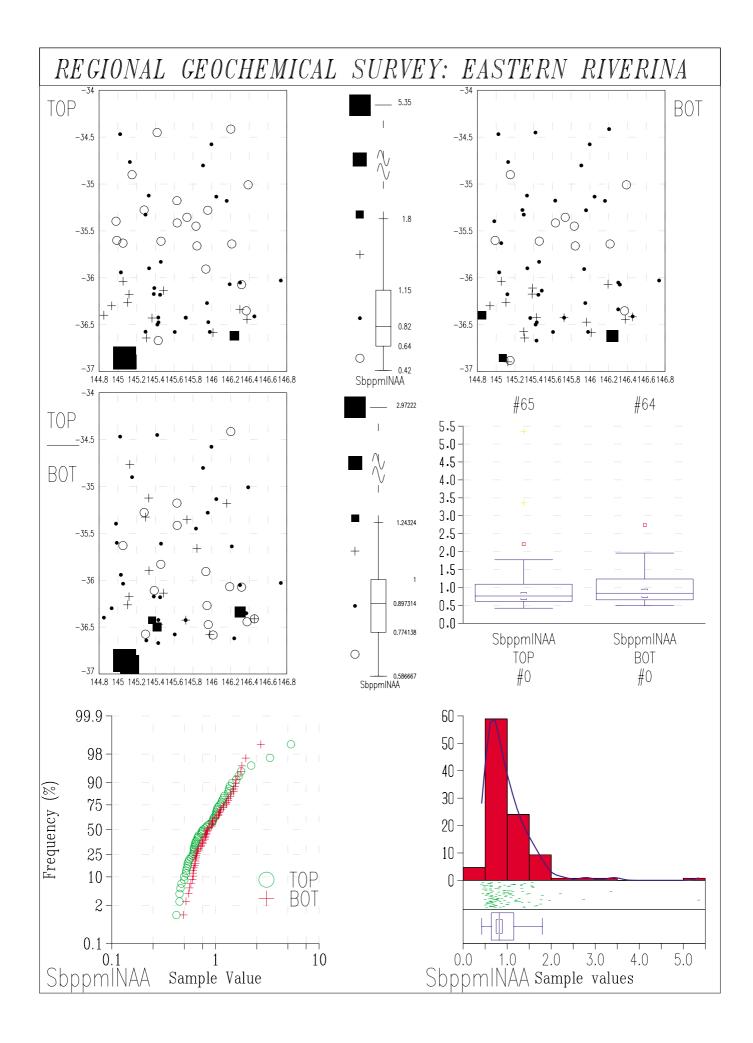


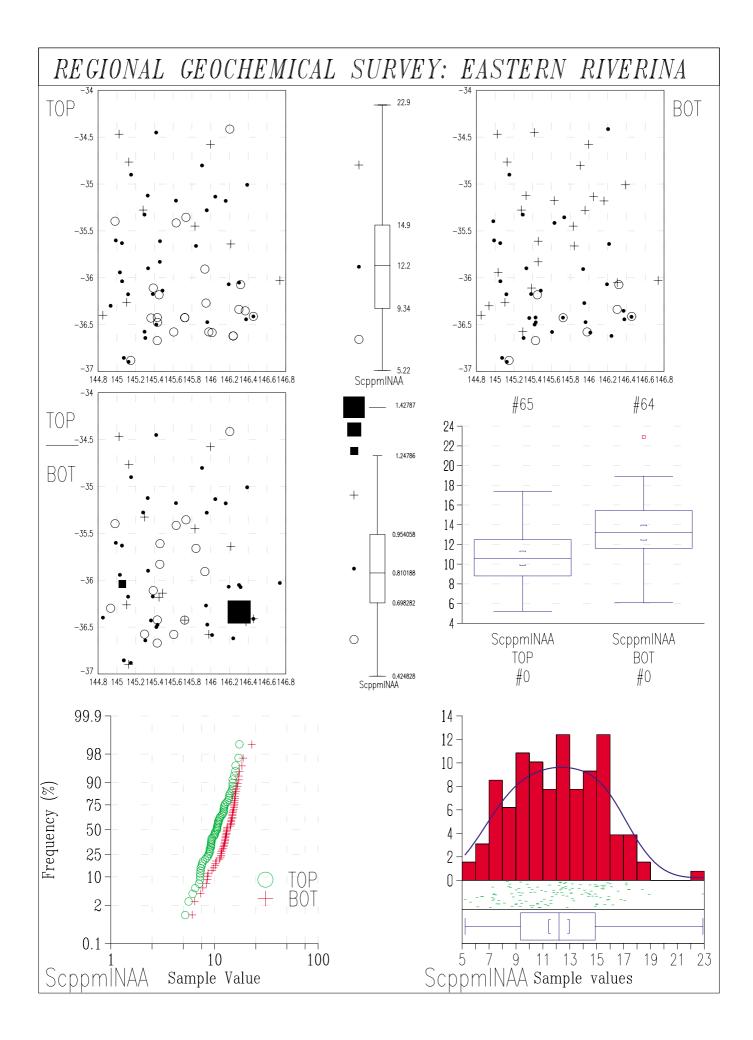


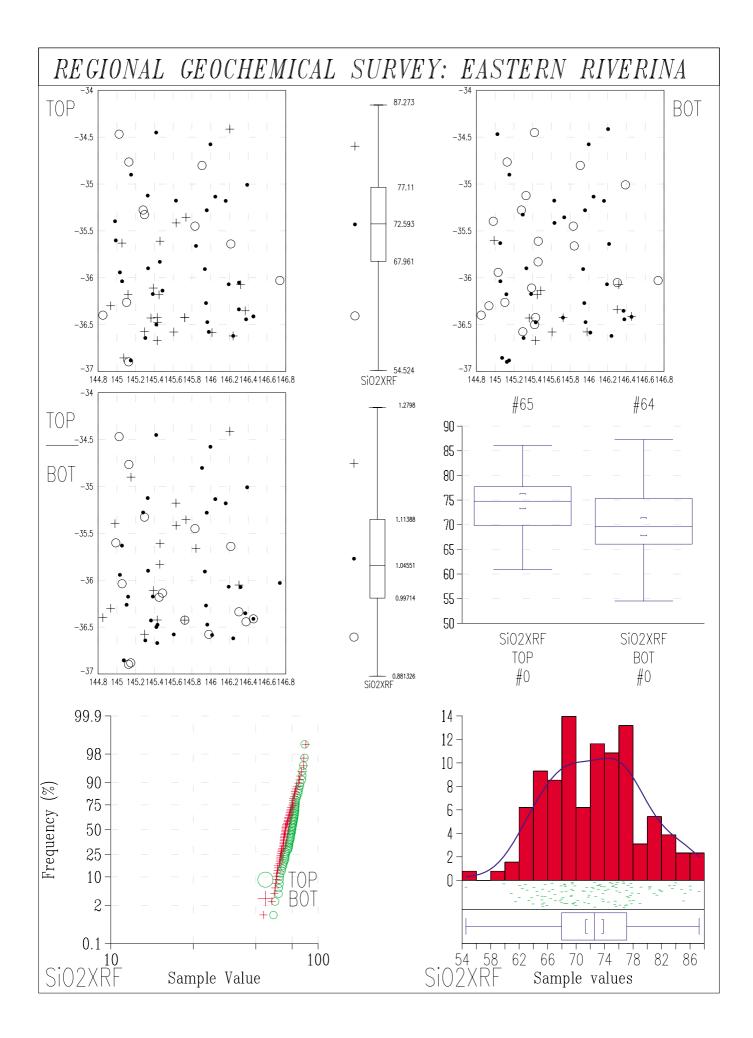


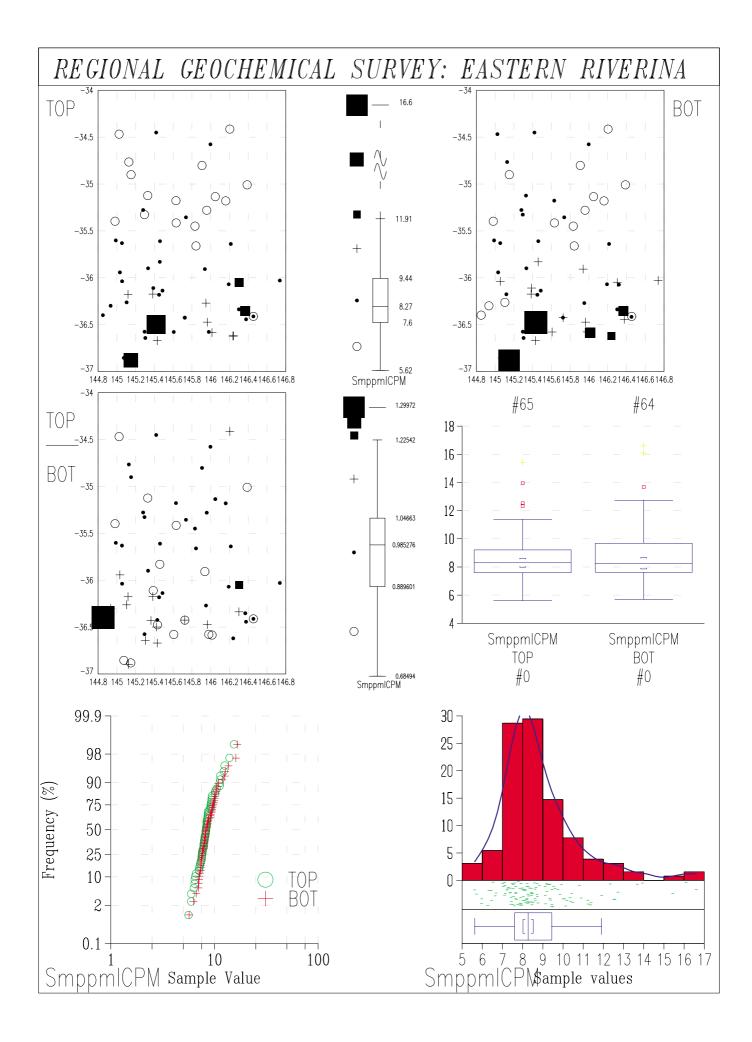


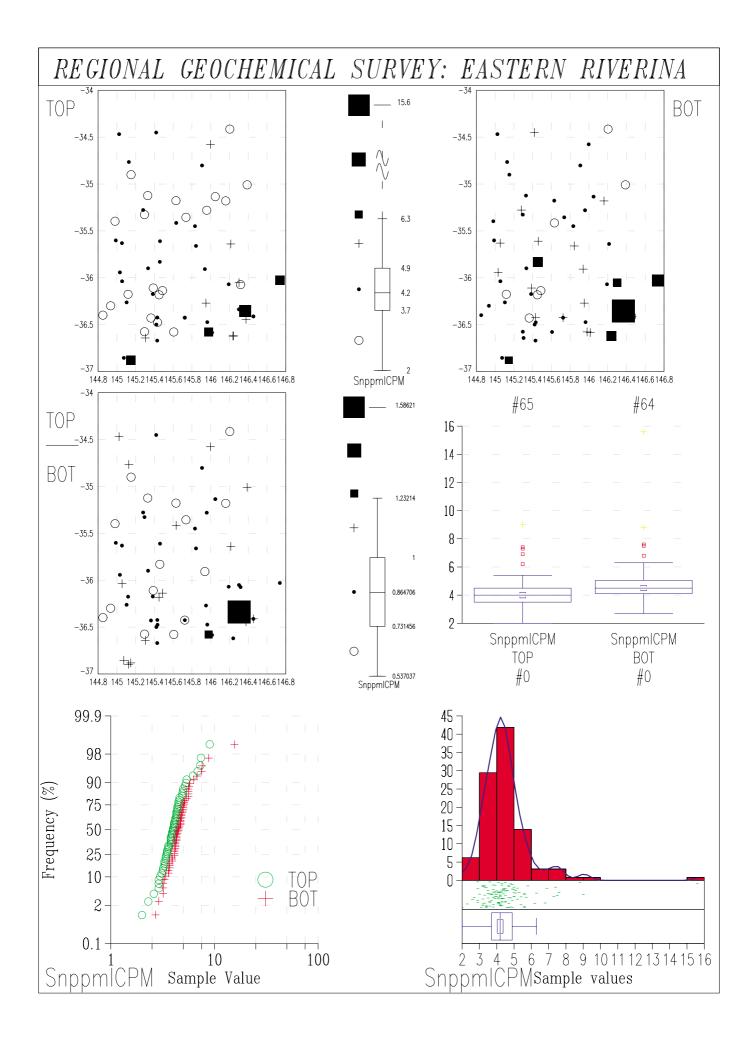


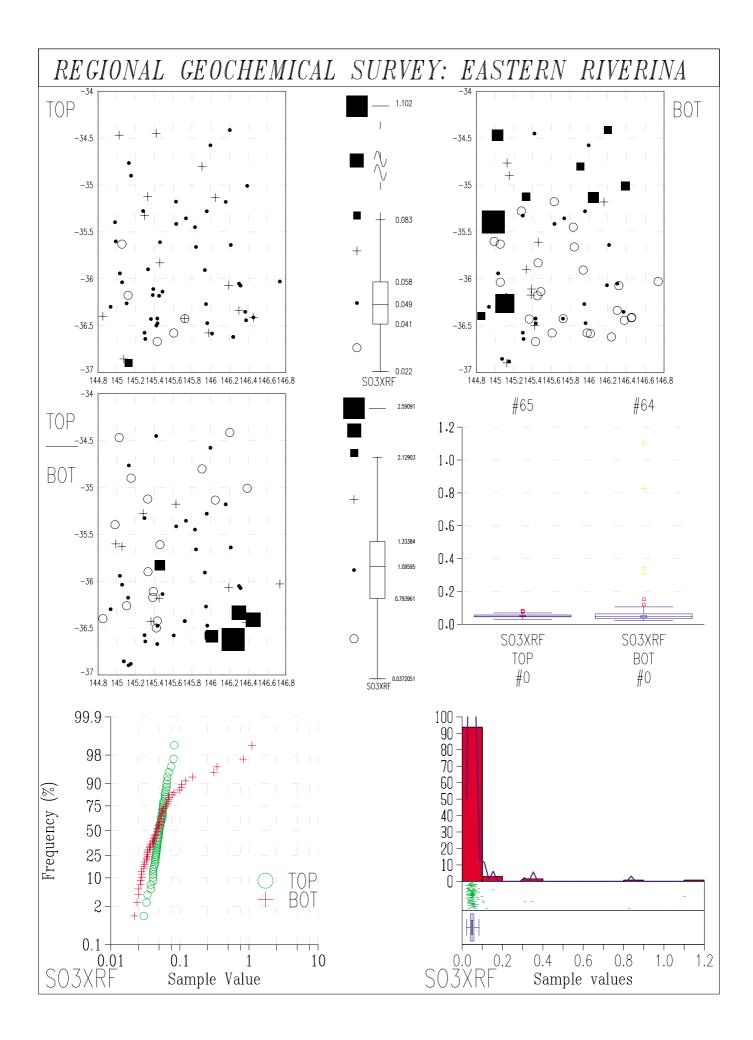


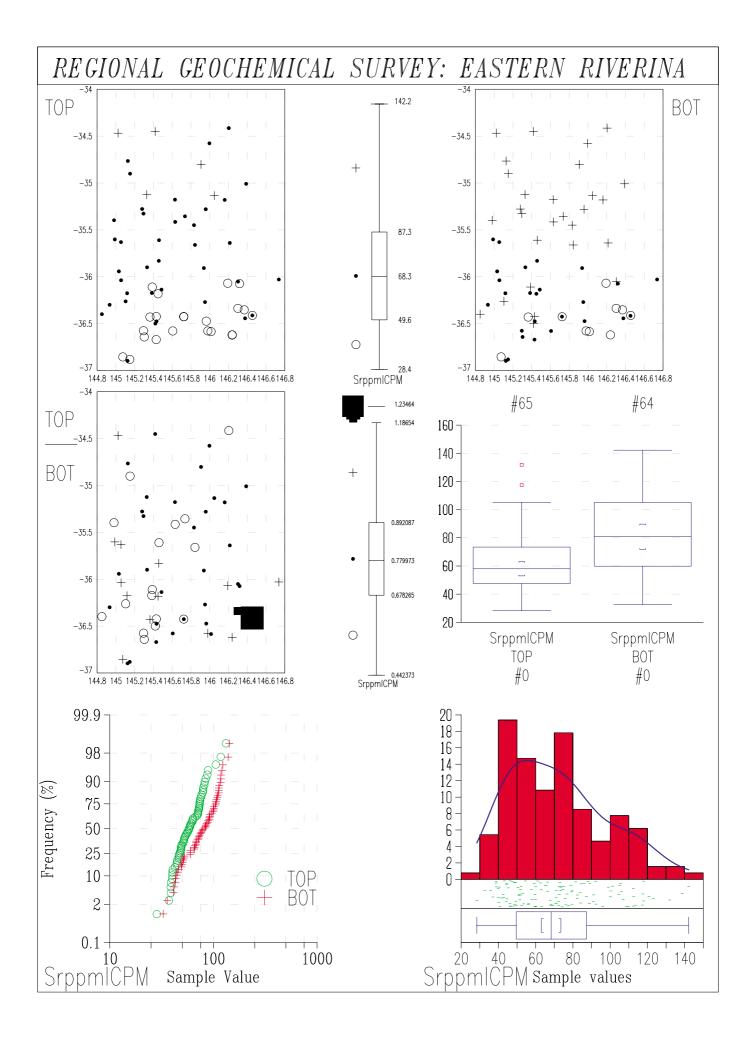


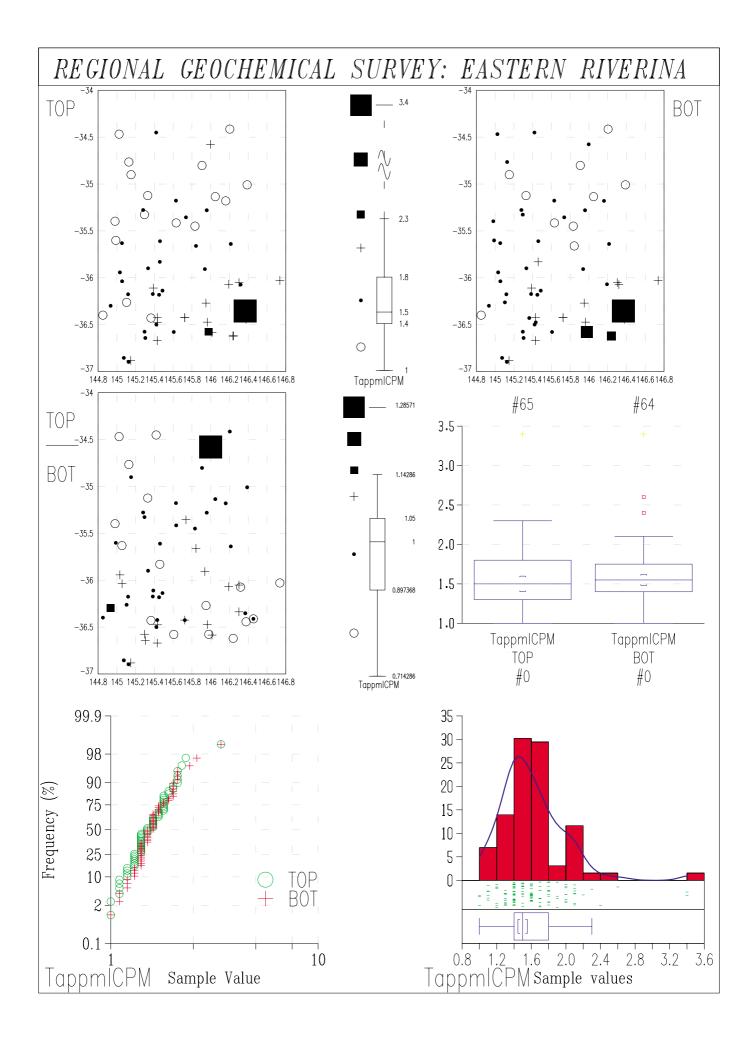


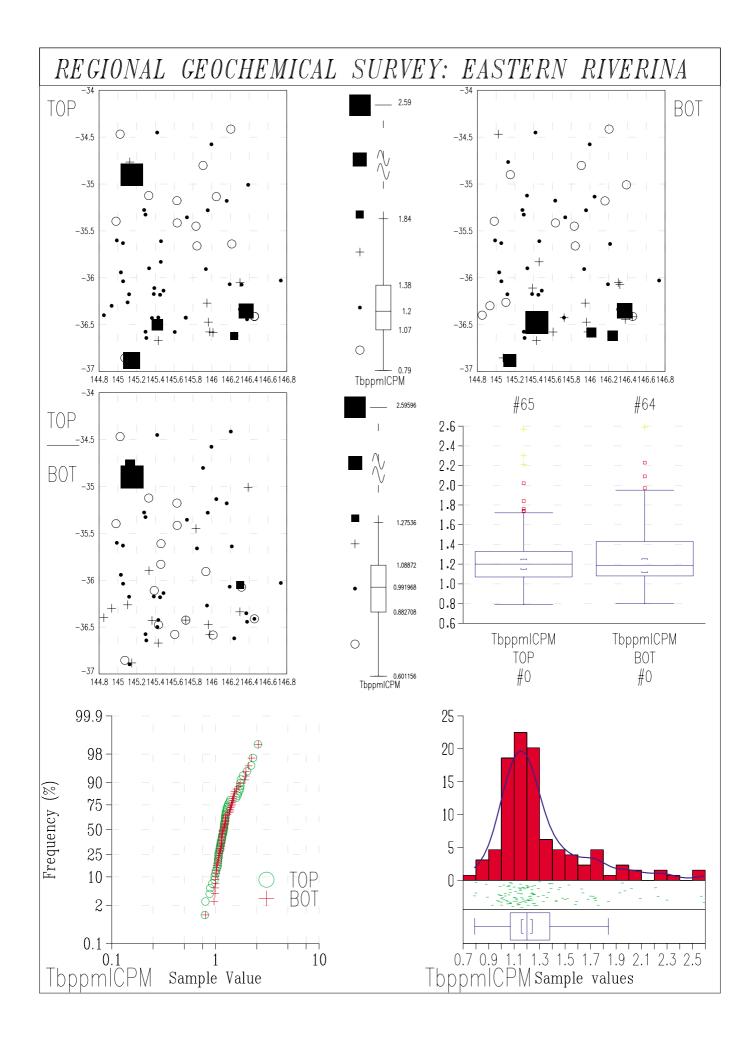


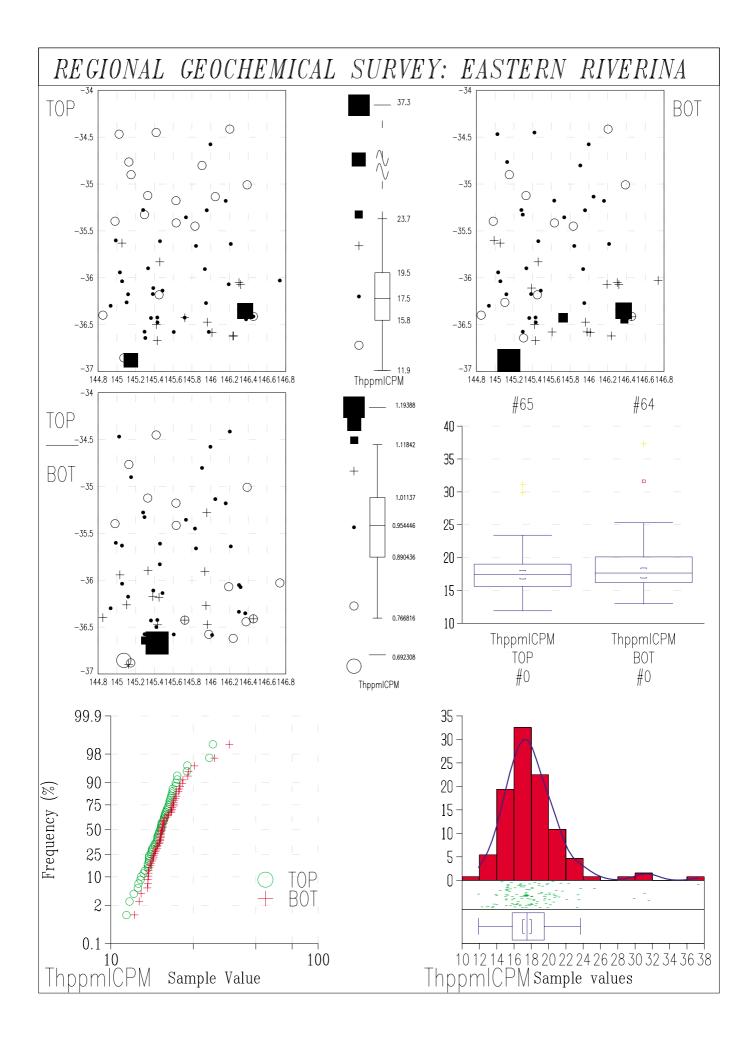


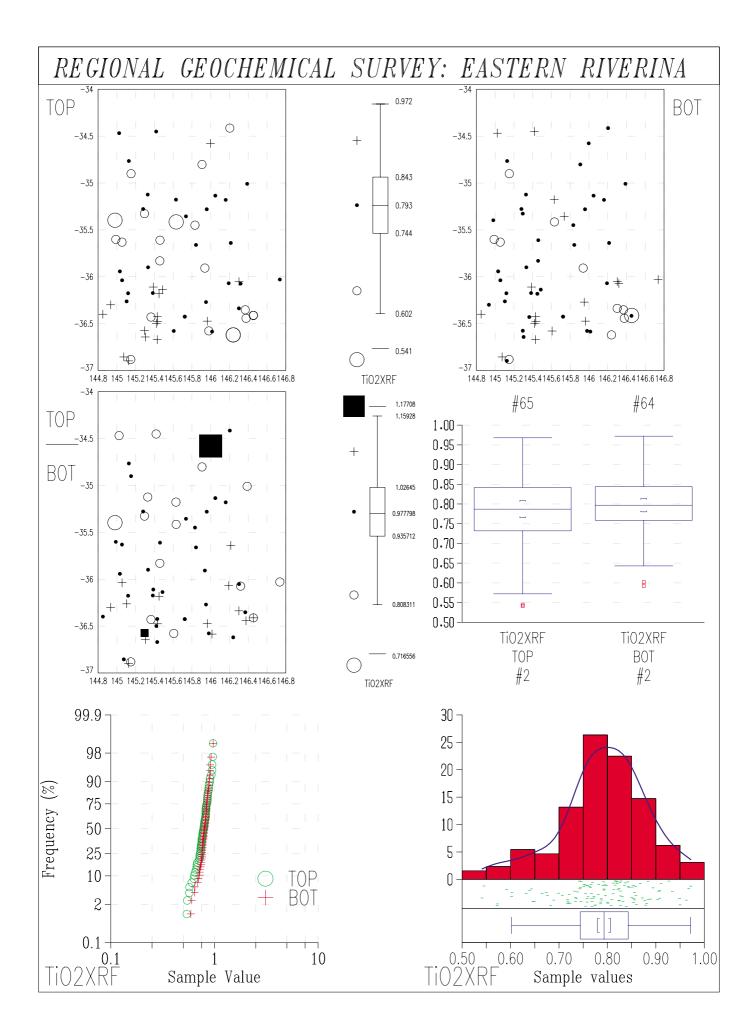


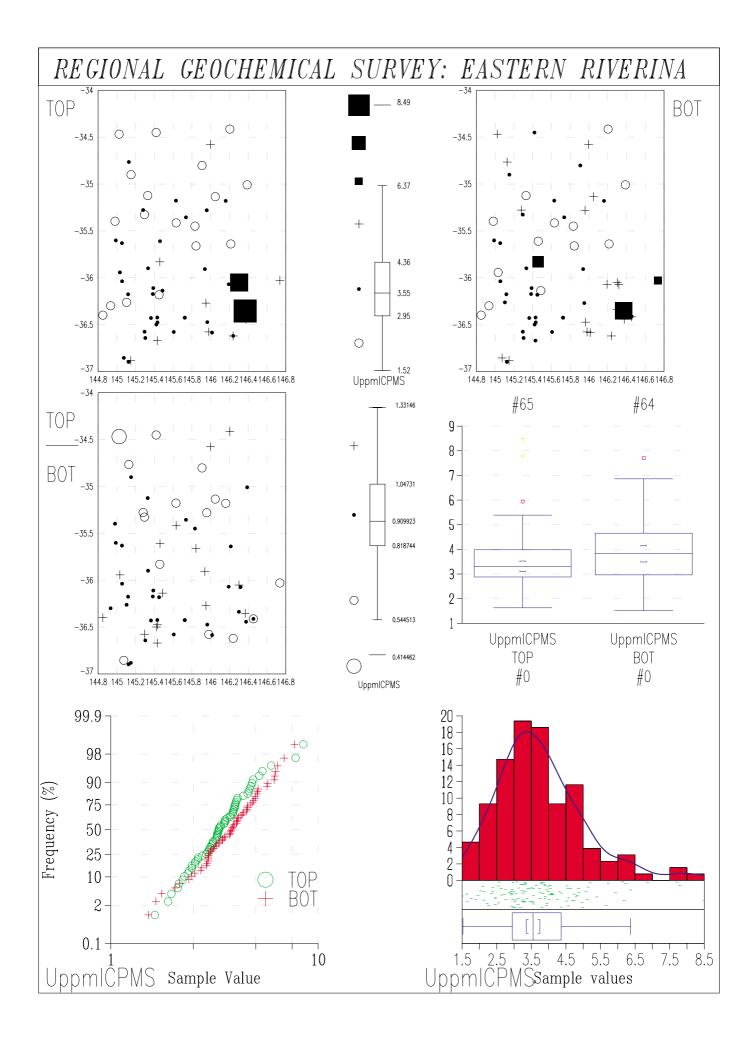


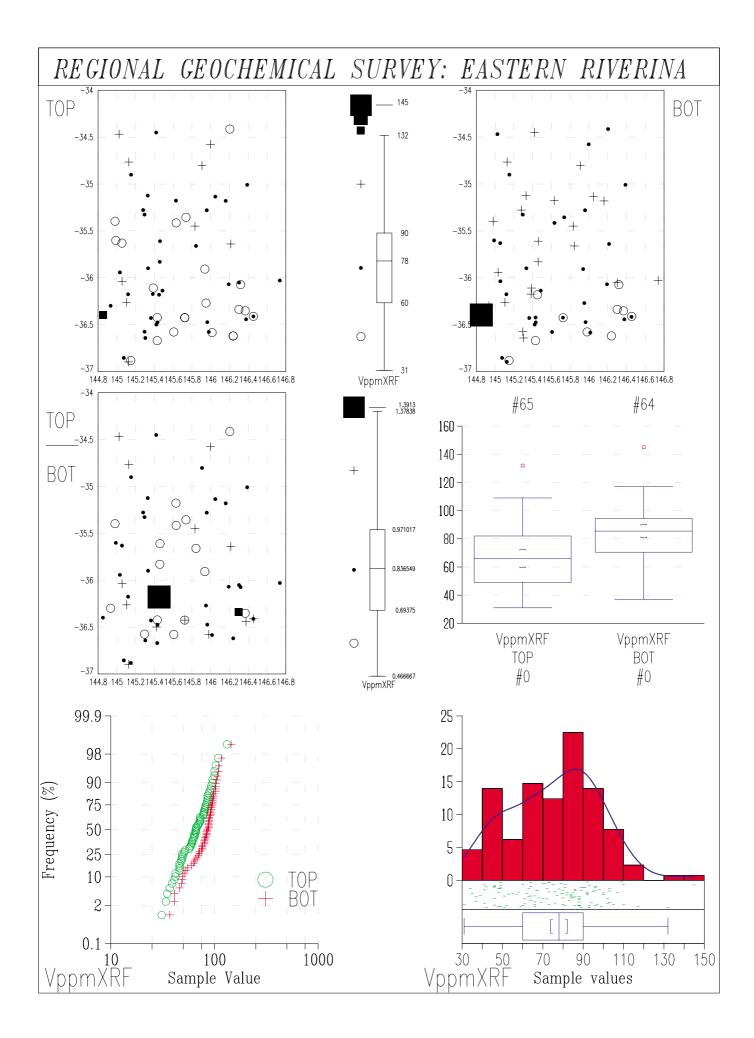


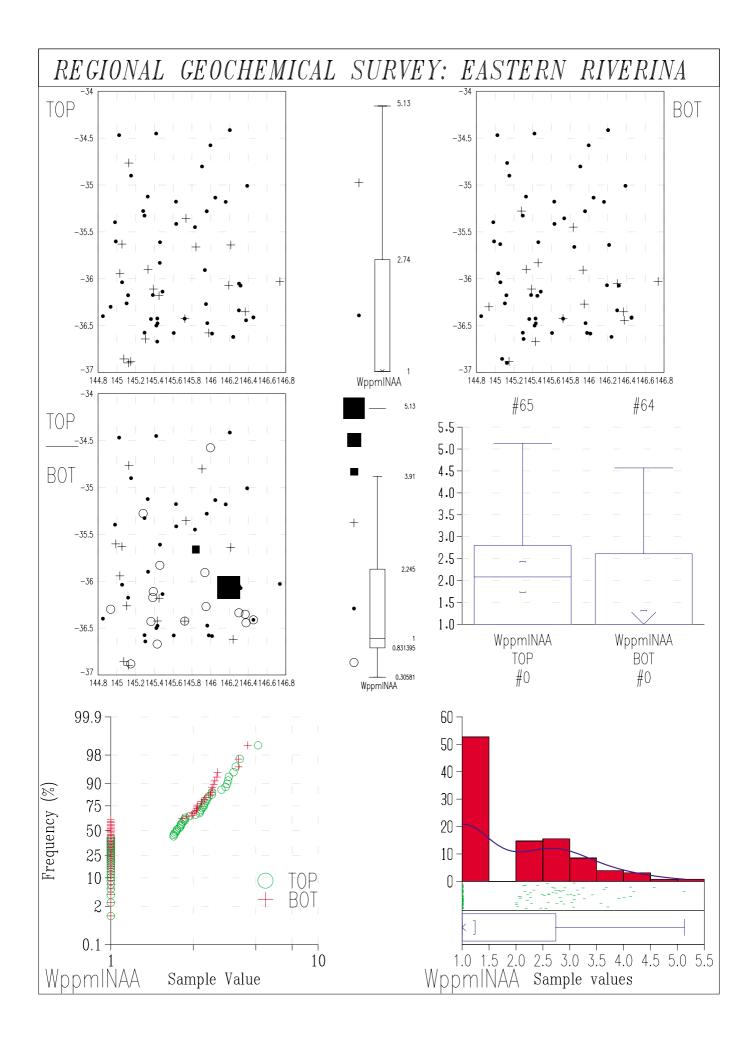


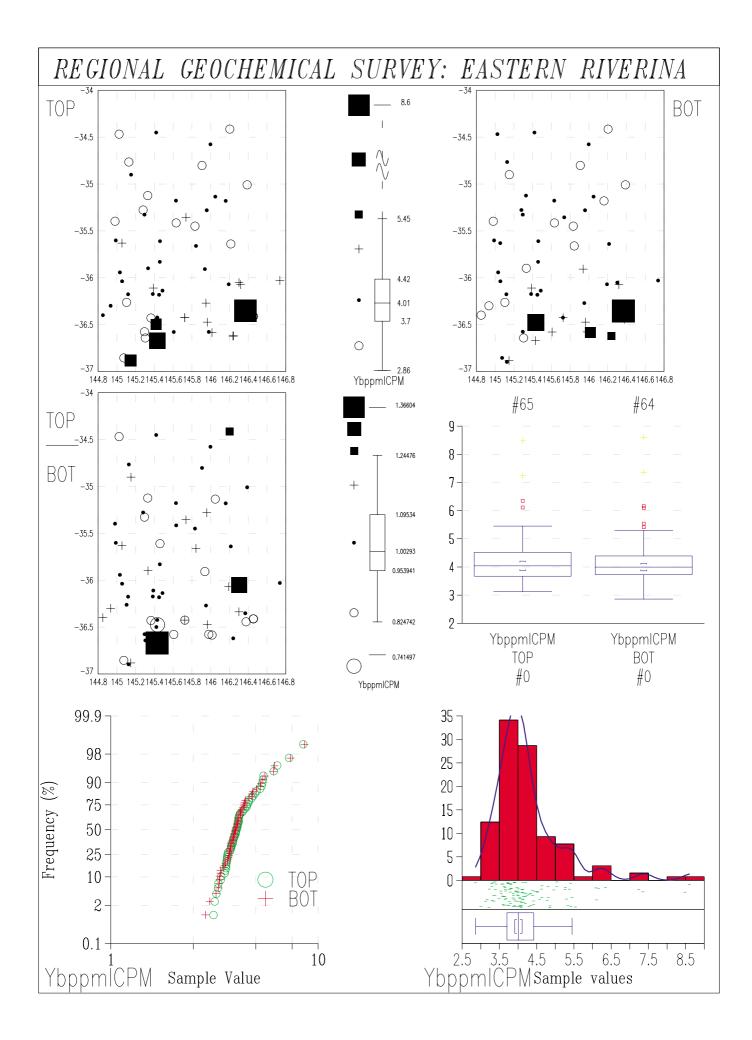


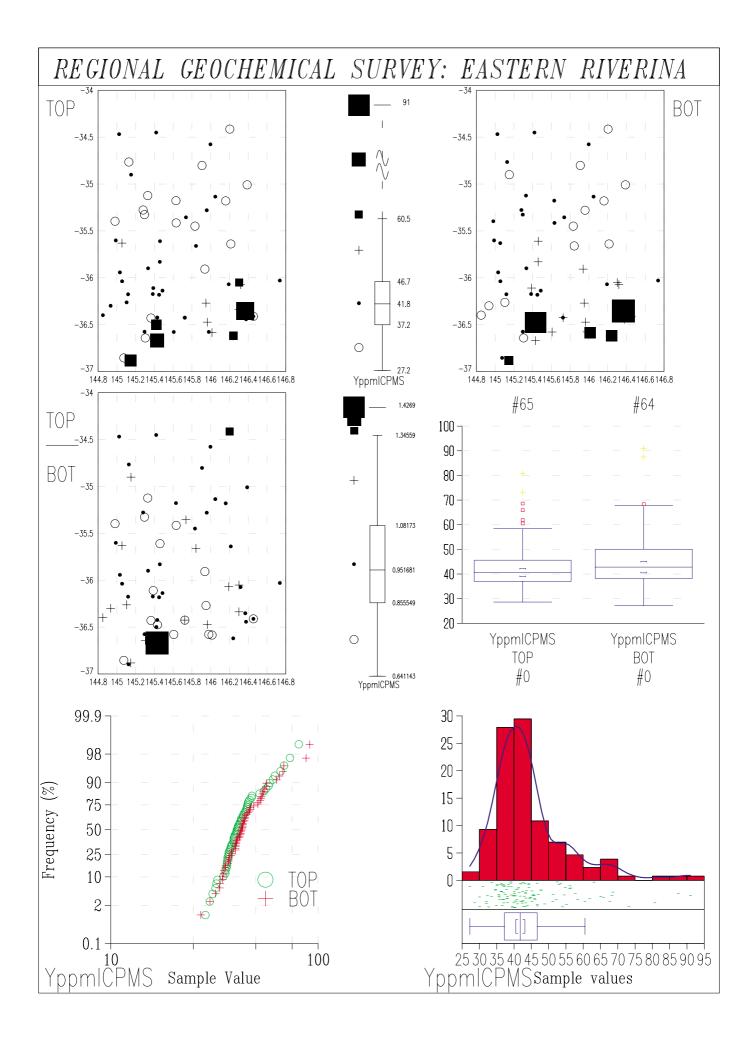


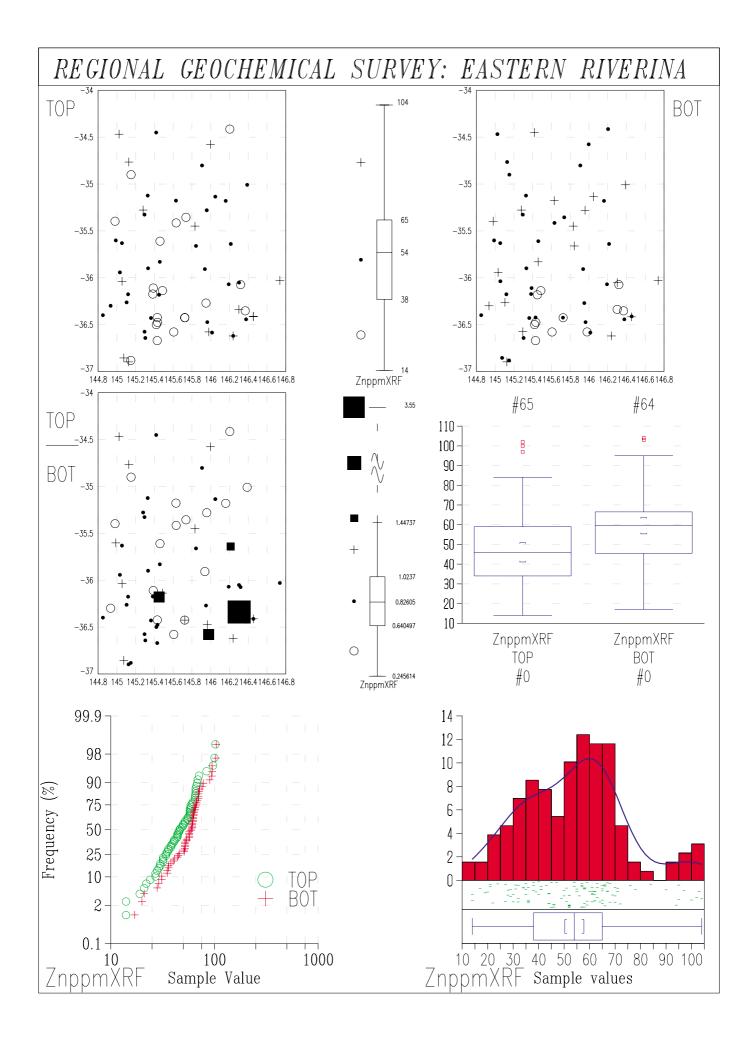


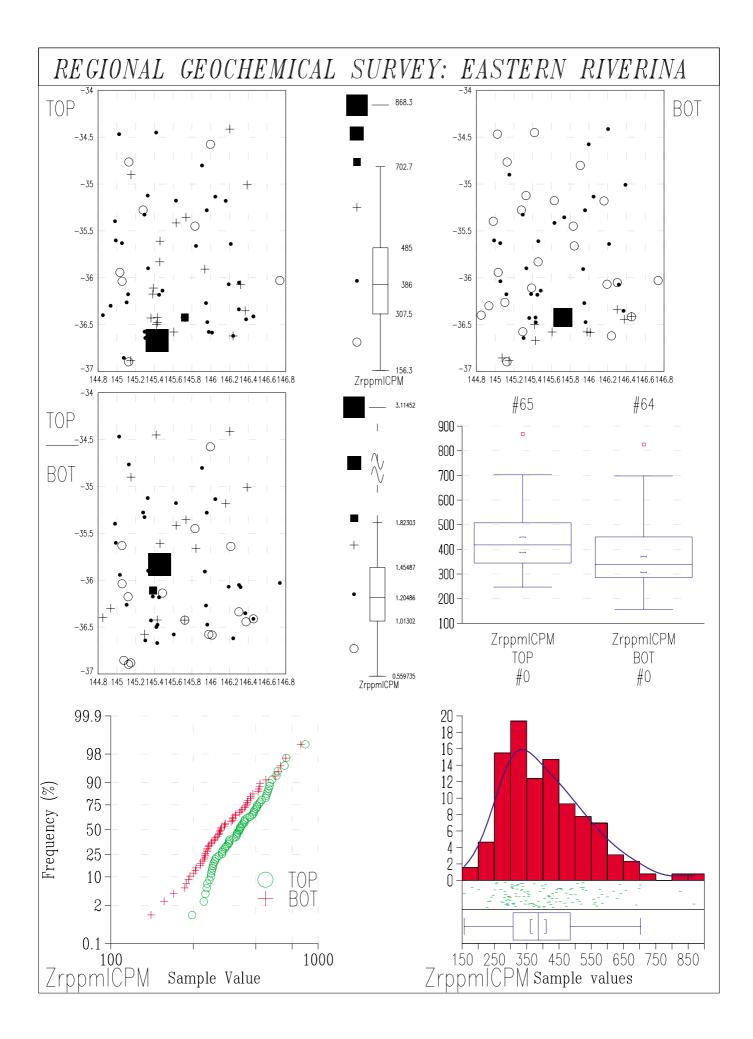




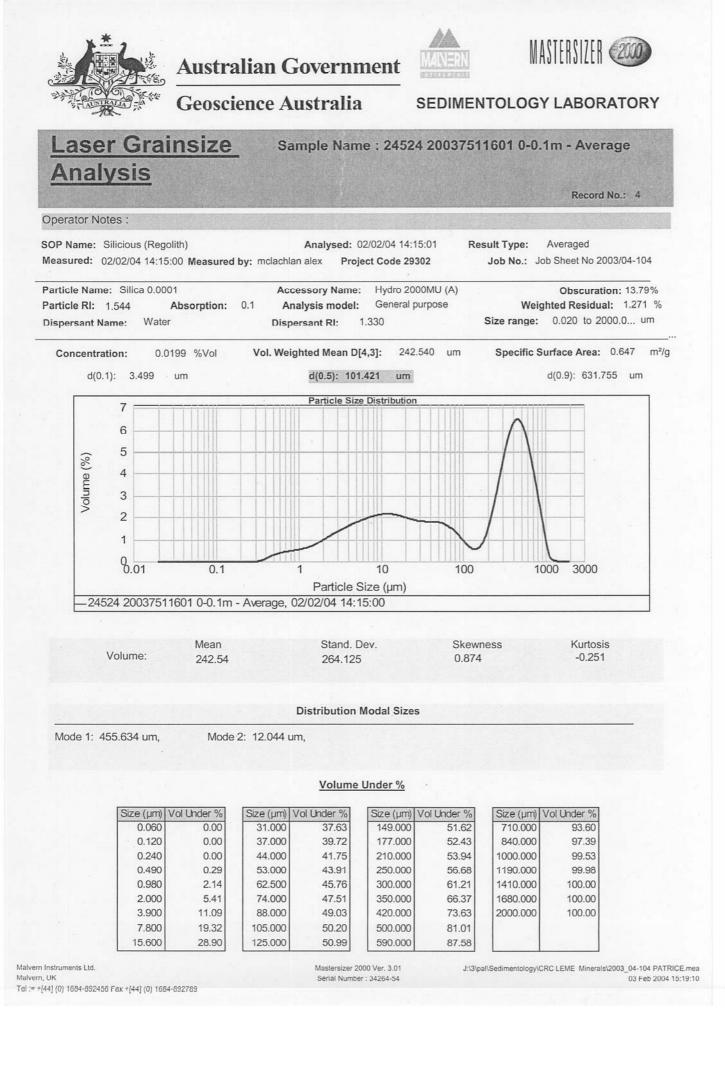








Appendix 5: Results of laser particle size analysis of 20 samples from the eastern Riverina region (bulk samples), Sedimentology Laboratory, Geoscience Australia





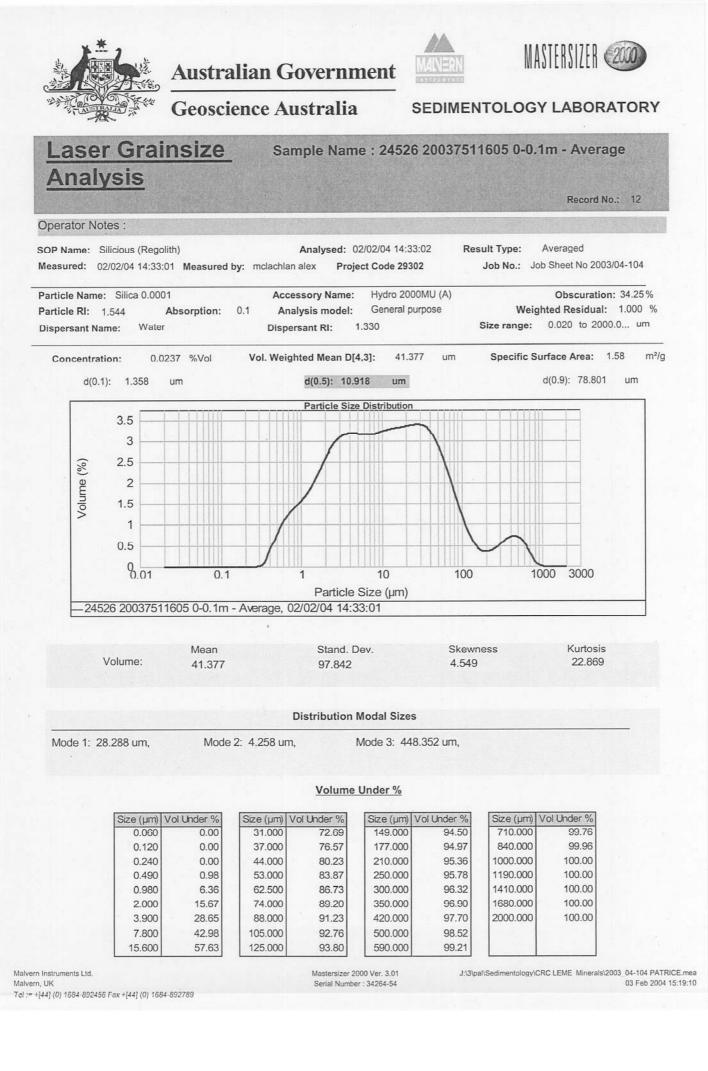




**Geoscience** Australia

SEDIMENTOLOGY LABORATORY

#### Laser Grainsize Sample Name : 24525 20037513601 0.8-0.9m - Average Analysis Record No.: 8 **Operator Notes :** Analysed: 02/02/04 14:24:01 Result Type: Averaged SOP Name: Silicious (Regolith) Job No.: Job Sheet No 2003/04-104 Measured: 02/02/04 14:24:00 Measured by: mclachlan alex Project Code 29302 Hydro 2000MU (A) Particle Name: Silica 0.0001 Obscuration: 28.48% Accessory Name: Weighted Residual: 1.147 % General purpose Particle RI: 1.544 Absorption: 0.1 Analysis model: Size range: 0.020 to 2000.0... um 1.330 Dispersant Name: Water Dispersant RI: Vol. Weighted Mean D[4,3]: 164.109 um Specific Surface Area: 1 m²/q 0.0291 %Vol Concentration: d(0.1): 2.358 d(0.5): 16.904 d(0.9): 557.199 um um um Particle Size Distribution 4.5 4 3.5 Volume (%) 3 2.5 2 1.5 1 0.5 8.01 100 0.1 1 10 1000 3000 Particle Size (µm) 24525 20037513601 0.8-0.9m - Average, 02/02/04 14:24:00 Kurtosis Mean Stand. Dev. Skewness Volume: 1.406 0.864 164.109 243.314 **Distribution Modal Sizes** Mode 1: 471.404 um, Mode 2: 10.409 um, Volume Under % Size (µm) Vol Under % 0.060 0.00 31.000 59.91 149.000 710.000 95.81 68.25 0.120 0.00 37.000 61.77 177.000 68.61 840.000 98.40 44.000 0.240 0.00 63.27 210.000 69.35 1000.000 99.78 0.490 0.46 53.000 64.60 250.000 70.85 1190.000 100.00 0.980 3.24 62.500 65.59 300.000 1410.000 100.00 73 56 8.20 2.000 74.000 66.44 350.000 76.86 1680.000 100.00 3.900 17.35 88.000 67.14 420.000 81.74 2000.000 100.00 7.800 31.67 105.000 67.66 500.000 86.88 15.600 48.26 125.000 67.99 590.000 91.55 Malvern Instruments Ltd. Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:10 Tel := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789









**Geoscience** Australia

SEDIMENTOLOGY LABORATORY

### Laser Grainsize Sample Name : 24527 20037513605 0.8-0.9m - Average Analysis Record No.: 16 **Operator Notes :** Analysed: 02/02/04 14:48:26 **Result Type:** Averaged SOP Name: Silicious (Regolith) Job No.: Job Sheet No 2003/04-104 Measured: 02/02/04 14:48:25 Measured by: mclachlan alex Project Code 29302 Obscuration: 26.84 % Hydro 2000MU (A) Particle Name: Silica 0.0001 Accessory Name: Weighted Residual: 1.096 % General purpose Particle RI: 1.544 Absorption: 0.1 Analysis model: Size range: 0.020 to 2000.0... um 1.330 Water **Dispersant RI:** Dispersant Name: Vol. Weighted Mean D[4,3]: 15.372 um Specific Surface Area: 1.54 m²/g 0.0176 %Vol Concentration: d(0.1): 1.528 d(0.5): 8.953 d(0.9): 38.016 um um um Particle Size Distribution 5 4 Volume (%) 3 2 1 0.01 1000 3000 0.1 1 10 100 Particle Size (µm) -24527 20037513605 0.8-0.9m - Average, 02/02/04 14:48:25 Kurtosis Stand. Dev. Skewness Mean Volume: 2.249 6.549 15.372 17.664 **Distribution Modal Sizes** Mode 1: 13.837 um, Volume Under % Size (µm) Vol Under % 100.00 0.060 31.000 85.65 149.000 100.00 710.000 0.120 0.00 37.000 89.47 177.000 100.00 840.000 100.00 1000.000 100.00 0.240 0.00 44.000 92.61 210.000 100.00 0.490 0.71 53.000 95.29 250.000 100.00 1190.000 100.00 5.34 62,500 97.07 300.000 100.00 1410.000 100.00 0.980 1680.000 100.00 2.000 13.96 74.000 98.37 350.000 100.00 88.000 99.22 420.000 100.00 2000.000 100.00 3.900 27.93 7.800 105.000 99.72 500.000 100.00 46.07 15.600 66.68 125.000 99.93 590.000 100.00

Malvern Instruments Ltd. Malvern, UK

Te) := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789

Mastersizer 2000 Ver. 3.01 Serial Number : 34264-54 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea 03 Feb 2004 15:19:10







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### Laser Grainsize Sample Name : 24528 20037511609 0-0.1m - Average Analysis Record No.: 20 **Operator Notes:** SOP Name: Silicious (Regolith) Analysed: 02/03/04 09:09:02 Result Type: Averaged Measured: 02/03/04 09:09:01 Measured by: mclachlan alex Job No.: Job Sheet No 2003/04-104 Project Code 29302 Particle Name: Silica 0.0001 Hydro 2000MU (A) Accessory Name: Obscuration: 23.91% Weighted Residual: 1.092 % Particle RI: 1.544 General purpose Analysis model: Absorption: 0.1 Size range: 0.020 to 2000.0... um Dispersant Name: Water Dispersant RI: 1.330 0.0159 %Vol Vol. Weighted Mean D[4,3]: 45.558 um Specific Surface Area: 1.5 m²/q Concentration: d(0.1): 1.425 d(0.5): 11.168 d(0.9): 114.963 um um um Particle Size Distribution 3.5 3 2.5 Volume (%) 2 1.5 1 0.5 8.01 0.1 10 100 1000 3000 1 Particle Size (µm) 24528 20037511609 0-0.1m - Average, 02/03/04 09:09:01 Kurtosis Mean Stand. Dev Skewness Volume: 94.023 3.558 13.862 45.558 **Distribution Modal Sizes** Mode 1: 11.918 um, Mode 2: 308.924 um, Volume Under % Size (µm) Vol Under % 0.060 0.00 31.000 71.41 149.000 91.74 710.000 99.98 0.120 0.00 37.000 74.78 177.000 92.77 840.000 100.00 0.240 0.00 44.000 77.93 1000.000 100.00 210.000 93.79 0.65 0.490 53.000 81.07 250.000 94.87 1190.000 100.00 0.980 5.65 62.500 83.56 300.000 96.05 1410.000 100.00 2.000 15.19 74.000 350.000 1680.000 85.77 97.06 100.00 3.900 27.97 88.000 87.69 420.000 98.18 2000.000 100.00 7.800 42.21 105.000 89.29 500.000 99.08 15.600 57.33 125.000 90.60 590.000 99.68 Malvern Instruments Ltd. Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:11 Tel := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789







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### Laser Grainsize Sample Name : 24529 20037513609 0.75-0.9m - Average Analysis Record No.: 24 **Operator Notes :** Analysed: 02/03/04 09:16:59 SOP Name: Silicious (Regolith) **Result Type:** Averaged Job No.: Job Sheet No 2003/04-104 Measured: 02/03/04 09:16:58 Measured by: mclachlan alex Project Code 29302 Hydro 2000MU (A) Obscuration: 24.30% Particle Name: Silica 0.0001 Accessory Name: Weighted Residual: 1.259 % General purpose 0.1 Analysis model: Particle RI: 1.544 Absorption: Size range: 0.020 to 2000.0... um Water 1.330 Dispersant RI: Dispersant Name: Vol. Weighted Mean D[4,3]: 47.452 um Specific Surface Area: 1.75 m²/a 0.0141 %Vol Concentration: d(0.9): 114.322 um d(0.1): 1.205 d(0.5): 8.768 um um Particle Size Distribution 3.5 3 2.5 Volume (%) 2 1.5 1 0.5 0.01 0.1 1 10 100 1000 3000 Particle Size (µm) 24529 20037513609 0.75-0.9m - Average, 02/03/04 09:16:58 Stand. Dev. Skewness Kurtosis Mean Volume: 3.601 13.72 107.213 47 452 **Distribution Modal Sizes** Mode 1: 3.529 um. Mode 2: 374.65 um, Volume Under % Size (µm) Vol Under % 0.060 0.00 31.000 73.88 710.000 149.000 91.40 0.00 37.000 76.85 177.000 92.20 840.000 100.00 0.120 0.240 0.00 44.000 79.63 210.000 93.02 1000.000 100.00 1190.000 100.00 0.490 0.95 53.000 82.39 250.000 93.95 0.980 7.26 62.500 84.57 300.000 95.06 1410.000 100.00 74.000 86.49 350.000 96.08 1680.000 100.00 2.000 18.57 2000.000 3.900 32.79 88.000 88.13 420.000 97.31 100.00 7.800 47.58 105.000 89.46 500.000 98.42 125.000 90.51 590.000 99.27 15.600 61.51 Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern Instruments Ltd. Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:11 To) := +[44] (0) 1694-892456 Fax +[44] (0) 1684-892789







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#### aser Grainsize Sample Name : 24530 20037511613 0-0.1m - Average Analysis Record No.: 28 **Operator Notes :** SOP Name: Silicious (Regolith) Analysed: 02/03/04 10:06:13 Result Type: Averaged Measured: 02/03/04 10:06:12 Measured by: mclachlan alex Job No.: Job Sheet No 2003/04-104 Project Code 29302 Particle Name: Silica 0.0001 Hydro 2000MU (A) Accessory Name: Obscuration: 23.01% Weighted Residual: 0.945 % Particle RI: 1.544 Absorption: 0.1 Analysis model: General purpose Water 1.330 Size range: 0.020 to 2000.0... um Dispersant Name: Dispersant RI: Vol. Weighted Mean D[4,3]: 50.215 Specific Surface Area: 1.36 Concentration: 0.0170 %Vol um m²/a d(0.1): 1.502 um d(0.5): 18.507 d(0.9): 102.574 um um Particle Size Distribution 4.5 4 3.5 Volume (%) 3 2.5 2 1.5 1 0.5 0.01 0.1 10 100 1000 3000 1 Particle Size (µm) -24530 20037511613 0-0.1m - Average, 02/03/04 10:06:12 Mean Stand, Dev. Skewness Kurtosis Volume: 17.247 3.915 50.215 98.593 **Distribution Modal Sizes** Mode 1: 41.515 um, Mode 2: 3.364 um, Mode 3: 388.524 um, Volume Under % Size (µm) Vol Under % 0.060 31.000 0.00 61.75 149.000 92.86 710.000 99.84 0.120 0.00 37.000 66.71 177.000 93.65 840.000 100.00 0.240 0.00 44.000 71.77 210.000 94.35 1000.000 100.00 0.490 0.69 53.000 77.06 250.000 95.08 1190.000 100.00 0.980 5.39 62.500 300.000 81.32 95.94 1410.000 100.00 74.000 2.000 13.94 85.02 350.000 96.74 1680.000 100.00 3.900 24.83 88.000 88.02 2000.000 420,000 97.74 100.00 7.800 35.93 105.000 90.25 500 000 98.65 99.35 15.600 46.92 125.000 91.79 590.000 Malvern Instruments Ltd. Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:11 Tel := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789





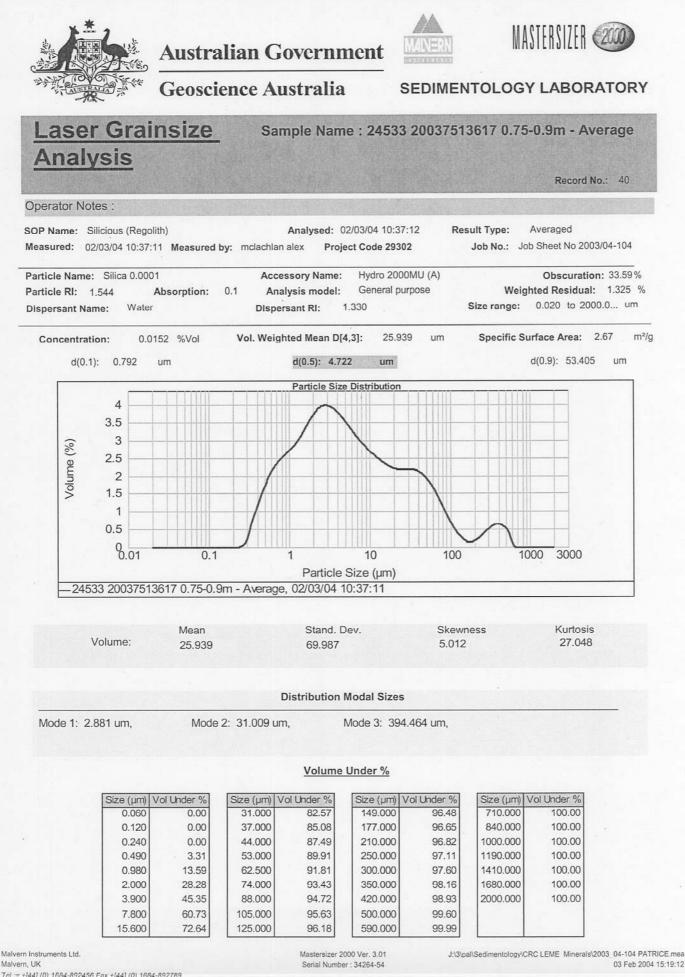


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### SEDIMENTOLOGY LABORATORY

#### Laser Grainsize Sample Name : 24531 20037513613 0.75-0.9m - Average Analysis Record No.: 32 **Operator Notes :** Analysed: 02/03/04 10:14:07 SOP Name: Silicious (Regolith) **Result Type:** Averaged Job No.: Job Sheet No 2003/04-104 Measured: 02/03/04 10:14:06 Measured by: mclachlan alex Project Code 29302 Hydro 2000MU (A) Particle Name: Silica 0.0001 Accessory Name: Obscuration: 23.34 % Analysis model: General purpose Weighted Residual: 1.130 % Particle RI: 1.544 Absorption: 0.1 Size range: 0.020 to 2000.0... um Water Dispersant RI: 1.330 Dispersant Name: Vol. Weighted Mean D[4,3]: 52.023 Specific Surface Area: 1.81 m²/a um Concentration: 0.0134 %Vol d(0.1): 1.151 um d(0.5): 9.732 d(0.9): 101.534 um um Particle Size Distribution 3.5 3 2.5 Volume (%) 2 1.5 1 0.5 0.01 0.1 10 100 1000 3000 1 Particle Size (um) 24531 20037513613 0.75-0.9m - Average, 02/03/04 10:14:06 Mean Stand. Dev. Skewness Kurtosis Volume: 17.503 3.993 52.023 121.234 **Distribution Modal Sizes** Mode 1: 41.047 um. Mode 3: 444.941 um. Mode 2: 2.943 um, Volume Under % Size (µm) Vol Under % 0.060 0.00 31.000 68.62 149.000 92.16 710.000 99.38 0.120 0.00 37.000 72.44 177.000 92.74 840.000 99.80 0.240 0.00 44.000 76.32 210.000 93.29 1000.000 99.96 0.490 1.33 53.000 80.36 250.000 93.91 1190.000 100.00 0.980 7.89 62.500 83.59 300.000 94.71 1410.000 100.00 2.000 19.34 74.000 86.38 350.000 95.52 1680.000 100.00 3.900 33.71 88.000 96.59 2000.000 100.00 88.62 420,000 7.800 46.59 105.000 90.27 500.000 97.65 98.58 15.600 56.82 125.000 91.39 590.000 Malvern Instruments Ltd. Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:11 Tel := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789

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Laser	CONSCRIPTION OF THE OWNER	nsize	San	nple Nam	ne: 24532	2003751	1617 0-0.1	m - Averag	e
Analy	<u>sis</u>							Record No.:	: 36
Operator Note	es :								
OP Name: Sil Measured: 02/	and the second second				02/03/04 10:28:2 ect Code 29302			Averaged Sheet No 2003/0	04-104
Particle Name: Particle RI: 1. Dispersant Nam	544 A	Absorption:	0.1 Anal	ssory Name: ysis model: sant RI:	Hydro 2000 General purp 1.330	oose		Obscuration: ated Residual: 0.020 to 2000.0	0.978 %
Concentratio		95 %Vol		ted Mean D[4	<b>4,3]:</b> 50.579	um	Specific Su	rface Area: 1.5	i3 m²/g
d(0.1)	1.397	um		d(0.5): 11.	740 um			d(0.9): 120.722	um
				Particle Size	e Distribution				
	3			$\bigcap$	$\searrow$				
(%)	2.5								
Volume (%)	1.5								
0N	1		/						
	0.5						$\sim$		
1.1	8.01	0.1		1	10	100	100	00 3000	
0450	00007544	617.0.0.1-	Augure 02		Size (µm)				
-2455	2 20037311	017 0-0. 111	- Average, 02	103/04 10.2	20.22	-			
Mean Volume: 50,570				Stand. Dev.		Skewness Kurtosis 3.819 16.539			
·	olume.	50.579		104.875	)	3.819		10.559	
			D	istribution	Modal Sizes				
Mode 1: 3.6	7 um,	Mode	2: 46.671 um		Mode 3: 395.5	539 um,			-
				Volume	Under %				
	Size (µm) V	/ol Under % 0.00	Size (µm) Vo	ol Under % 67.11	Size (µm) Vo	ol Under % 91.86	Size (µm) V	ol Under % 99.75	
	0.120	0.00	37.000	70.38	177.000	93.02	840.000	99.97	
	0.240	0.00 0.88	44.000 53.000	73.67 77.23	210.000 250.000	93.98 94.85	1000.000 1190.000	100.00	
	0.980	5.99 15.46	62.500 74.000	80.32 83.29	300.000 350.000	95.76 96.55	1410.000	100.00	
		10.40		86.04	420.000	97.52	2000.000	100.00	
	3.900	28.60 42.38	88.000	88.43	500.000	98.42			



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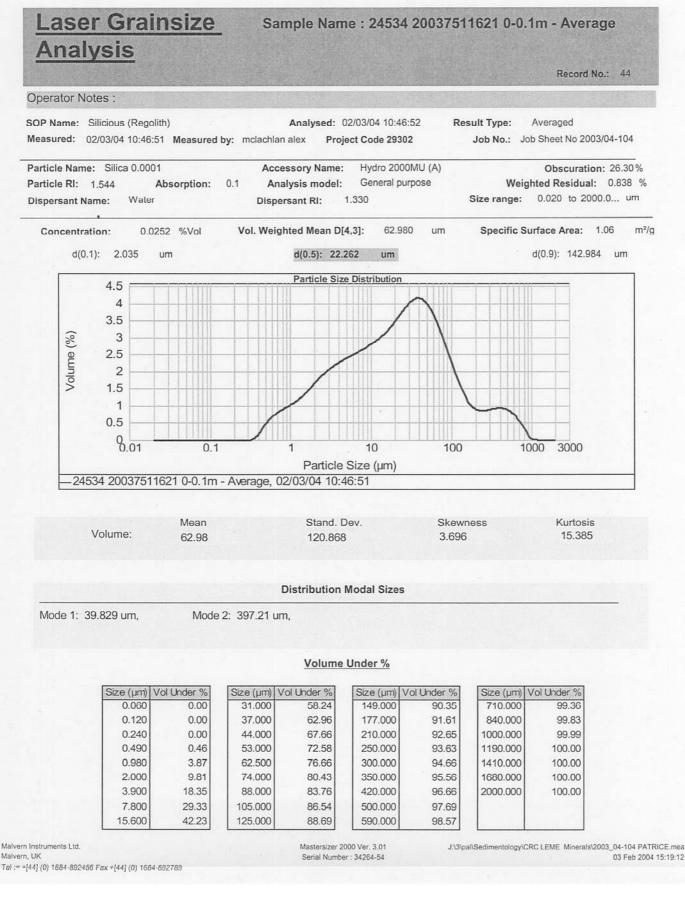






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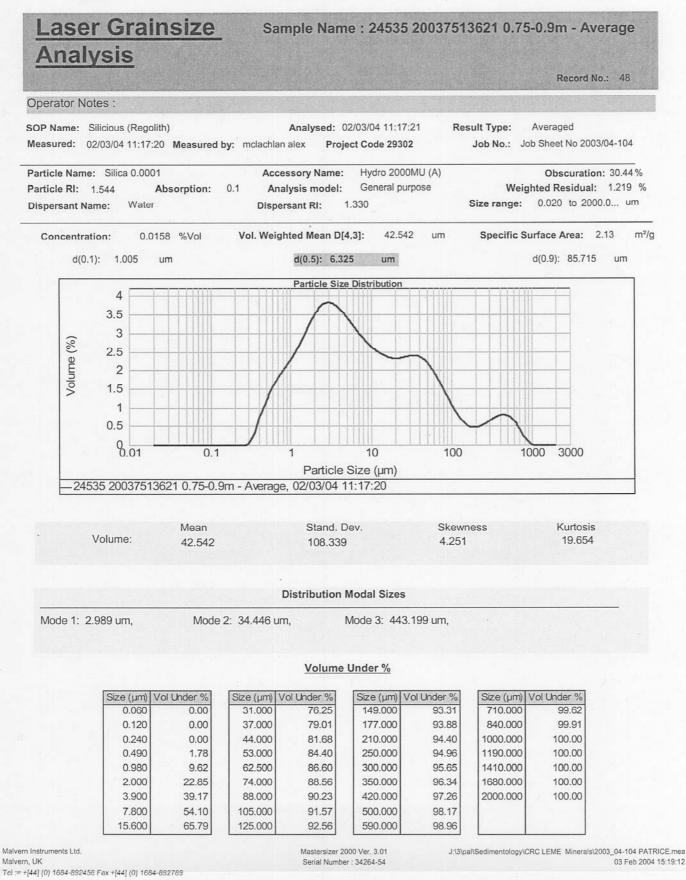


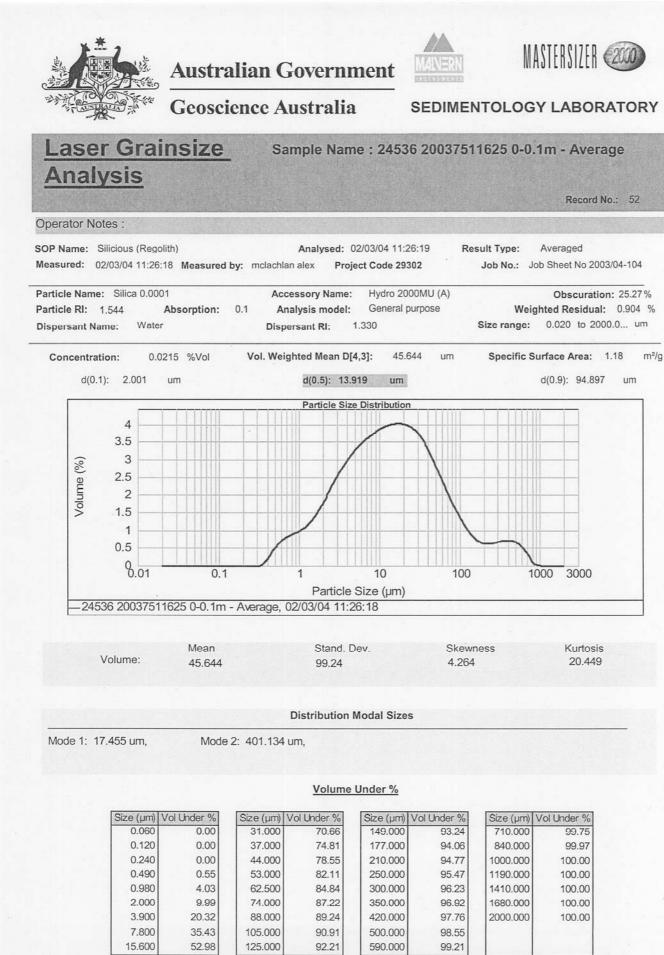




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## SEDIMENTOLOGY LABORATORY

#### Laser Grainsize Sample Name : 24537 20037513625 0.75-0.9m - Average Analysis Record No.: 56 **Operator Notes:** Analysed: 02/03/04 11:34:57 Averaged Result Type: SOP Name: Silicious (Regolith) Job No.: Job Sheet No 2003/04-104 Measured: 02/03/04 11:34:56 Measured by: mclachlan alex Project Code 29302 Obscuration: 23.35% Hydro 2000MU (A) Particle Name: Silica 0.0001 Accessory Name: Weighted Residual: 1.043 % General purpose Absorption: 0.1 Analysis model: Particle RI: 1.544 Size range: 0.020 to 2000.0... um Water 1.330 · Dispersant RI: Dispersant Name: Vol. Weighted Mean D[4,3]: 23.727 um Specific Surface Area: 1.41 m²/q 0.0170 %Vol Concentration: d(0.9): 46.961 d(0.1): 1.588 d(0.5): 12.117 um um um Particle Size Distribution 5 4 Volume (%) 3 2 1 0.01 100 1000 3000 0.1 1 10 Particle Size (µm) -24537 20037513625 0.75-0.9m - Average, 02/03/04 11:34:56 Kurtosis Stand. Dev. Skewness Mean Volume: 6.901 60.23 23.727 47.978 **Distribution Modal Sizes** Mode 1: 18.332 um. Volume Under % Size (µm) Vol Under % 98.22 710.000 100.00 0.00 31.000 0.060 80.65 149.000 0.00 37.000 85.19 177.000 98.51 840.000 100.00 0.120 100.00 0.240 0.00 44.000 88.84 210.000 98.73 1000.000 1190.000 100.00 0.490 0.90 53.000 91.86 250.000 98.92 1410.000 100.00 0.980 5.56 62.500 93.85 300.000 99.13 74.000 95.35 350.000 99.33 1680.000 100.00 2,000 12.73 99.58 2000.000 100.00 3.900 23.27 88.000 96.45 420.000 7.800 37.76 105.000 97.26 500.000 99.80 99.95 15.600 58.24 125.000 97.82 590.000

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### SEDIMENTOLOGY LABORATORY

#### Laser Grainsize Sample Name : 24538 20037511629 0-0.1m - Average Analysis Record No.: 60 **Operator Notes:** Analysed: 02/03/04 11:45:41 **Result Type:** Averaged SOP Name: Silicious (Regolith) Project Code 29302 Job No.: Job Sheet No 2003/04-104 Measured: 02/03/04 11:45:40 Measured by: mclachlan alex Hydro 2000MU (A) Particle Name: Silica 0.0001 Accessory Name: Obscuration: 29.82% General purpose Absorption: 0.1 Analysis model: Weighted Residual: 1.041 % Particle RI: 1.544 Size range: 0.020 to 2000.0... um 1.330 Water Dispersant RI: **Dispersant Name:** 32.602 Specific Surface Area: 1.4 m²/g Vol. Weighted Mean D[4,3]: um Concentration: 0.0219 %Vol d(0.9): 62.903 um d(0.1): 1.715 um d(0.5): 9.686 um Particle Size Distribution 4.5 4 3.5 Volume (%) 3 2.5 2 1.5 1 0.5 0.01 0.1 10 100 1000 3000 1 Particle Size (µm) 24538 20037511629 0-0.1m - Average, 02/03/04 11:45:40 Kurtosis Mean Stand. Dev. Skewness Volume: 32.326 5.374 32.602 81 449 **Distribution Modal Sizes** Mode 2: 464.971 um, Mode 1: 8.827 um. Volume Under % Size (µm) Vol Under % 0.00 31.000 99.88 0.060 78.72 149.000 96.19 710.000 0.00 37.000 840.000 100.00 0.120 82.03 177.000 96.60 0.240 0.00 44.000 84.97 210.000 96.89 1000.000 100.00 0.490 0.63 53.000 87.76 250.000 97.16 1190.000 100.00 0.980 4.66 62.500 300.000 97.50 1410.000 100.00 89.92 1680.000 2.000 12.12 74.000 91.81 350.000 97.87 100.00 3.900 25.39 88.000 93.39 420.000 98.41 2000.000 100.00 7.800 43.88 105.000 94.65 500.000 98.98 15.600 63.04 125.000 95.56 590.000 99.48 Mastersizer 2000 Ver. 3.01 J:\3\pal\Sedimentology\CRC LEME Minerals\2003\_04-104 PATRICE.mea Malvern Instruments Ltd. Malvern, UK Serial Number : 34264-54 03 Feb 2004 15:19:13 Tel := +[44] (0) 1684-892456 Fax +[44] (0) 1684-892789







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## SEDIMENTOLOGY LABORATORY

#### Laser Grainsize Sample Name : 24539 20037513629 0.75-0.9m - Average Analysis Record No.: 64 Operator Notes : Analysed: 02/03/04 13:50:53 **Result Type:** Averaged SOP Name: Silicious (Regolith) Measured: 02/03/04 13:50:52 Measured by: mclachlan alex Project Code 29302 Job No.: Job Sheet No 2003/04-104 Hydro 2000MU (A) Particle Name: Silica 0.0001 Accessory Name: Obscuration: 33.51% Weighted Residual: 1.150 % Analysis model: General purpose Particle RI: 1.544 Absorption: 0.1 Size range: 0.020 to 2000.0... um 1.330 Water Dispersant RI: **Dispersant Name:** 17.309 Specific Surface Area: 1.56 m²/g Vol. Weighted Mean D[4,3]: um Concentration: 0.0231 %Vol d(0.9): 38.634 d(0.1): 1.590 um d(0.5): 8.086 um um Particle Size Distribution 5 4 Volume (%) 3 2 1 8.01 0.1 1 10 100 1000 3000 Particle Size (µm) 24539 20037513629 0.75-0.9m - Average, 02/03/04 13:50:52 Kurtosis Mean Stand, Dev. Skewness Volume: 9.38 116.777 36.97 17 309 **Distribution Modal Sizes** Mode 1: 8.332 um, Volume Under % Size (µm) Vol Under % 100.00 0.00 31.000 710.000 0.060 86.43 149.000 99.35 37.000 89.35 840.000 100.00 0.120 0.00 177.000 99.43 0.240 0.00 44.000 91.82 210.000 99.44 1000.000 100.00 0.490 0.85 53.000 94.06 250.000 99.47 1190.000 100.00 0.980 5.44 62.500 95.68 300.000 99.52 1410.000 100.00 74.000 1680.000 100.00 2.000 13.11 97.01 350,000 99.60 3.900 27.40 88.000 98.03 420.000 99.73 2000.000 100.00

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98.73

99.15

500.000

590.000

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105.000

125.000

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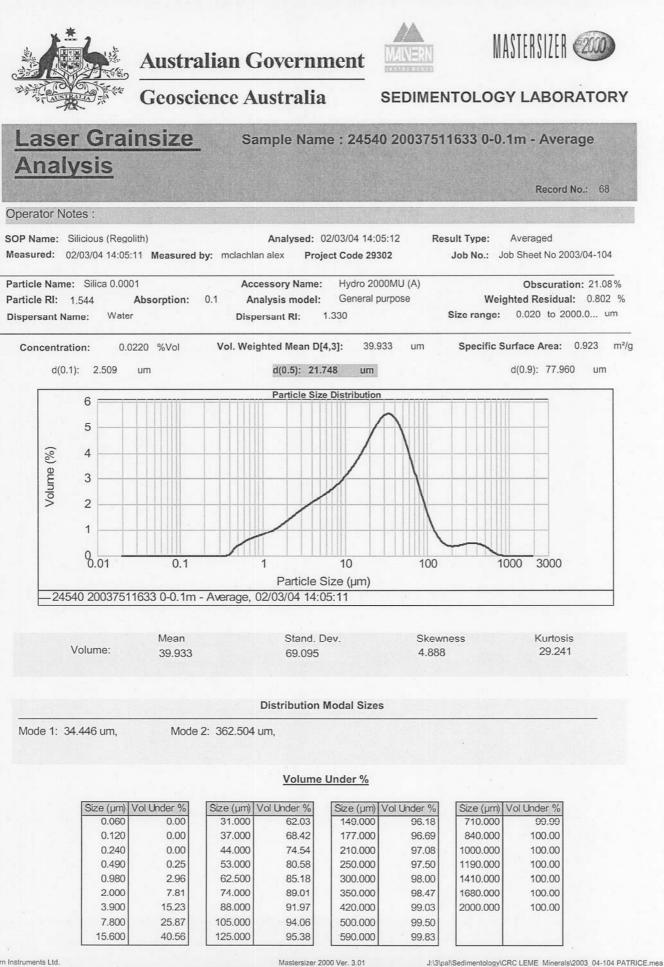
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7.800

15.600

48.80

70.76

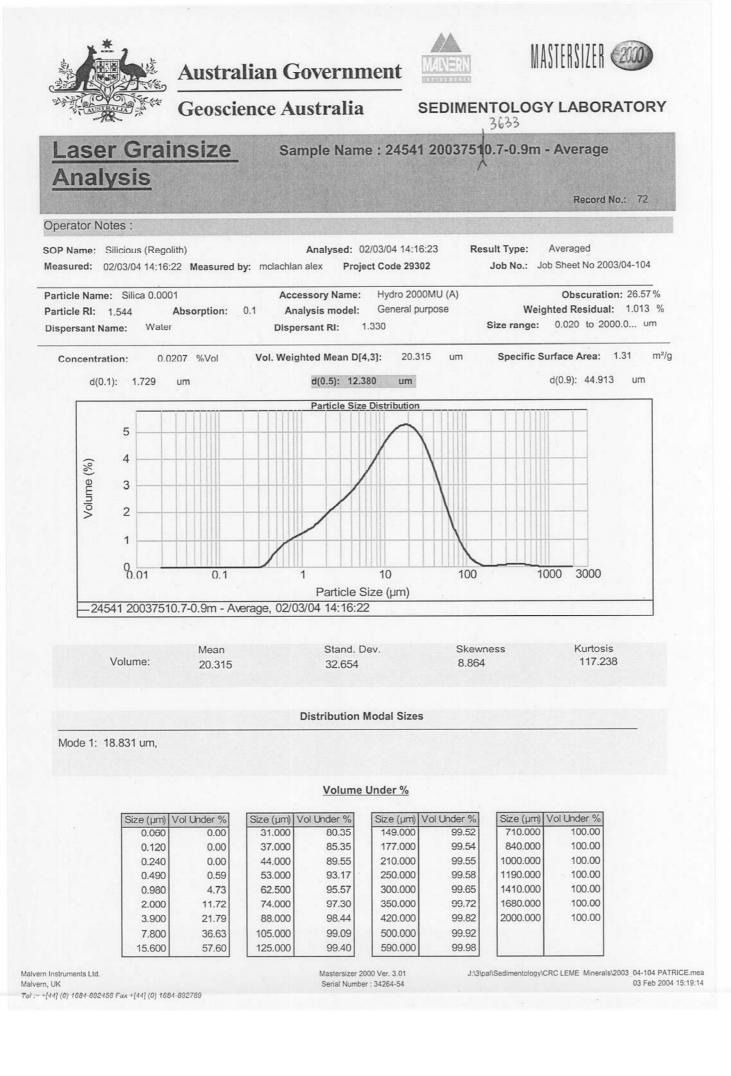


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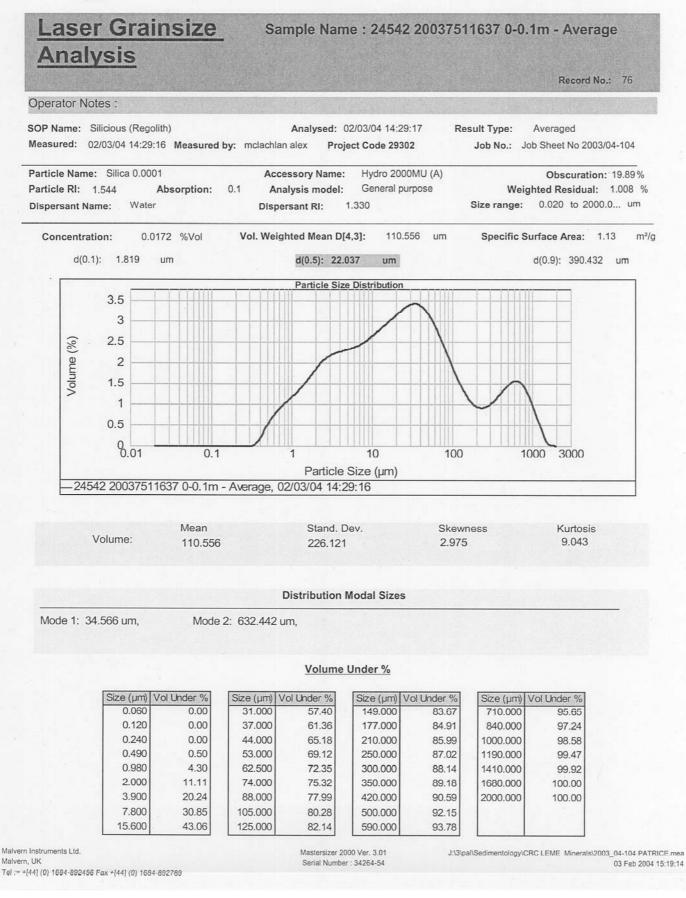






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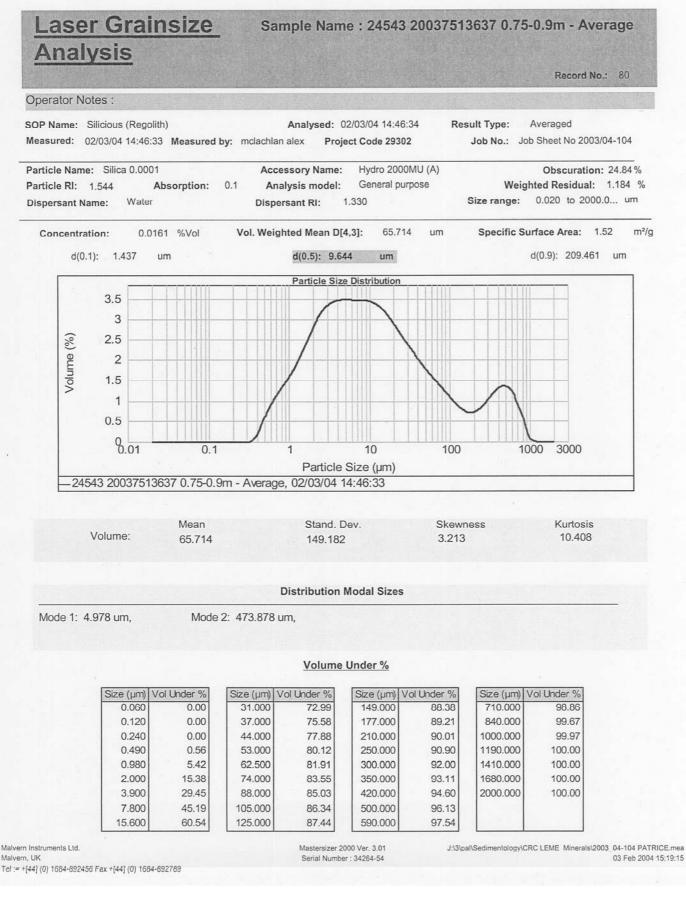






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Appendix 6: Macroscopic properties of overbank sediments form the eastern Riverina (field parameters)

SampleID	Texture	рH	Colour	Colour (dry)	Colour	Colour (wet)
Sampleid	TEXIULE	(field)	(dry)	Colour (ury)	(wet)	
20037512601	S	6	10YR6/6	Brownish Yellow	10YR4/3	Brown
20037514601	SCL	6.5	10YR7/6	Yellow	10YR4/4	Dark Yellowish
20037512602	SC	6	10YR5/3	Brown	10YR3/3	Brown Dark Brown
20037514602	LMC	6.5	10YR6/3	Pale Brown	7.5Y4/3	Brown
20037512603	CL	6	10YR5/2	Greyish Brown	10YR4/2	Dark Greyish
00007544000	0.01			2		Brown
20037514603	SiCL	5.5	10YR5/3	Brown	10YR3/2	Very Dark Greyish Brown
20037512604	LS	6	10YR5/3	Brown	10YR3/2	Very Dark Greyish
20027544004		C		Dark Crawish		Brown
20037514604	FSL	6	10YR4/2	Dark Greyish Brown	10YR2/2	Very Dark Brown
20037512605	FSCL	6	10YR7/6	Yellow	10YR5/4	Yellowish Brown
20037514605	LC	7.5	10YR6/4	Light Yellowish	10YR5/4	Yellowish Brown
20037512606	Lfsy	6.5	10YR6/4	Brown Light Yellowish	10YR4/3	Brown
2000/012000	Liby	0.0	1011(0/1	Brown	1011(1/0	Brown
20037514606	Μ	7	10YR7/4	Very Pale Brown	10YR4/4	Dark Yellowish
20037512607	Lfsy	6	10YR5/4	Yellowish Brown	10YR4/4	Brown Dark Yellowish
2000/01200/	LIGy	0	1011(0/4	T CHOWISH BIOWH		Brown
20037514607	CL	6.5	2.5Y7/4	Pale Yellow	2.5Y5/3	Light Olive Brown
20037512608	CLS	6	2.5Y6/2	Light Brownish	2.5Y4/2	Dark Greyish Brown
20037514608	НС	7	2.5Y7/2	Grey Light Grey	2.5Y5/2	Greyish Brown
20037512609	CL	6	10YR5/4	Yellowish Brown	7.5YR3/3	Dark Brown
20037514609	LC	7.5	2.5Y6/4	Light Yellowish	2.5Y5/3	Light Olive Brown
20037512610	CL	6.5	2.5Y7/2	Brown Light Grey	2.5Y6/2	Light Brownish
20037512010	0L	0.5	2.517/2	Light Grey	2.510/2	Grey
20037514610	MC	7	2.5Y5/2	Greyish Brown	7.5YR4/2	Brown
20037512611	LMC	6.5	10YR4/3	Brown	10YR3/3	Dark Brown
20037514611	LMC	8	2.5Y6/3	Light Yellowish Brown	2.5Y5/3	Light Olive Brown
20037512612	SiCL	6.5	10YR6/3	Pale Brown	10YR4/3	Brown
20037514612	MHC	7	10YR6/3	Pale Brown	10YR4/3	Brown
20037512613	CL	7	10YR5/3	Brown	10YR3/3	Dark Brown
20037514613	HC	9	10YR5/4	Yellowish Brown	10YR4/4	Dark Yellowish
20027542044	1.60.0	7		Light Cross		Brown
20037512614	Lfsy CL	7 8	10YR7/2 10YR7/3	Light Grey Very Pale Brown	10YR6/3	Pale Brown Pale Brown
20037514614 20037512615		6.5	101R7/3 10YR4/2	•	10YR6/3	
20037512015	LIVIC	0.5	101 K4/2	Dark Greyish Brown	10YR3/1	Very Dark Grey
20037514615	MC	8	2.5Y5/2	Greyish Brown	10YR4/2	Dark Greyish
20027512616	CL	7	10YR5/4	Yellowish Brown	10YR3/4	Brown Dark Yellowish
20037512616	CL	1	101 K3/4	reliowish brown	101 K3/4	Brown
20037514616	HC	8.5	2.5Y6/3	Light Yellowish	2.5Y5/3	Light Olive Brown
20027512617	МС	7	10YR5/4	Brown Yellowish Brown	10/04/2	Brown
20037512617 20037514617	HC	7 8.5	7.5YR5/4	Brown	10YR4/3 7.5YR4/4	Brown
20037512618	SiL	6.5	10YR5/3	Brown	10YR3/3	Dark Brown
20037512018	CL	5.5	10YR6/4	Light Yellowish	10YR5/4	Yellowish Brown
2000/014010	0L	5.5	101110/4	Brown	101110/4	
20037512619	L	5.5	10YR5/3	Brown	10YR3/3	Dark Brown
20037514619	Lfsy	8.5	10YR6/4	Light Yellowish	10YR5/4	Yellowish Brown
20037512620	Lfsy	5.5	10YR7/4	Brown Very Pale Brown	10YR5/4	Yellowish Brown
20037514620	CL	7	10YR7/3	Very Pale Brown	10YR5/4	Yellowish Brown
				-		

SampleID	Texture	pH (field)	Colour (dry)	Colour (dry)	Colour (wet)	Colour (wet)
20037512621	L	5	(01y) 10YR6/3	Pale Brown	10YR3/3	Dark Brown
20037514621	MC	5	10YR6/3	Pale Brown	10YR4/3	Brown
20037512622	Lfsy	7	10YR6/4	Light Yellowish Brown	10YR5/4	Yellowish Brown
20037514622	SC	7	10YR6/4	Light Yellowish Brown	10YR4/4	Dark Yellowish Brown
20037512623	L	4	10YR6/3	Pale Brown	10YR3/3	Dark Brown
20037514623	MC	7	2.5Y7/4	Pale Yellow	10YR4/6	Dark Yellowish Brown
20037512624	Lfsy	7	10YR5/3	Brown	10YR3/3	Dark Brown
20037514624	SCL	5.5	10YR7/2	Light Grey	10YR5/3	Brown
20037512625	SiL	5.5	10YR5/4	Yellowish Brown	10YR 3/3	Dark Brown
20037514625	SiCL	6.5	10YR6/6	Brownish Yellow	10YR4/6	Strong Brown
20037512626	SiL	5.5	10YR5/3	Brown	10YR3/3	Dark Brown
20037512627	LC	6.5	2.5Y7/4	Pale Yellow	2.5Y5/4	Light Olive Brown
20037514627	SC	6	2.5Y7/4	Pale Yellow	10YR5/4	Yellowish Brown
20037512628	SiL	5.5	10YR5/3	Brown	10YR3/2	Very Dark Greyish Brown
20037514628	S	5.5	10YR5/4	Yellowish Brown	10YR4/4	Dark Yellowish Brown
20037512629	SiCL	5.5	10YR6/3	Pale Brown	10YR4/2	Dark Greyish Brown
20037514629	SC	6	2.5Y7/4	Pale Yellow	10YR5/6	Yellowish Brown
20037512630	CL	5	10YR5/3	Brown	10YR3/2	Very Dark Greyish Brown
20037514630	SC	8	2.5Y7/2	Light Grey	2.5Y5/2	Greyish Brown
20037512631	L	5	10YR6/3	Pale Brown	10YR3/3	Dark Brown
20037514631	S	9	2.5Y7/3	Pale Yellow	2.5Y6/4	Light Yellowish Brown
20037512632	SL	9	10YR6/3	Pale Brown	10YR4/3	Brown
20037514632	LC	8.5	2.5Y7/4	Pale Yellow	2.5Y6/4	Light Yellowish Brown
20037512633	SiL	5.5	10YR6/3	Pale Brown	10YR4/3	Brown
20037514633	SiCL	6	2.5Y6/2	Light Brownish Grey	10YR3/2	Very Dark Greyish Brown
20037512634	LC	6	2.5Y6/3	Light Yellowish Brown	2.5Y4/2	Dark Greyish Brown
20037514634	SC	8.5	2.5Y6/3	Light Yellowish Brown	2.5Y5/3	Light Olive Brown
20037512635	L	5.5	10YR7/3	Very Pale Brown	10YR4/3	Brown
	LC	8	10YR6/6	Brownish Yellow	10YR4/6	Dark Yellowish Brown
20037512636	L	6.5	10YR4/4	Dark Yellowish Brown	7.5YR3/3	Dark Brown
20037514636	HC	7	7.5YR4/6	Strong Brown	7.5YR3/4	Dark Brown
20037512637	SiL	6	10YR5/3	Brown	10YR3/2	Very Dark Greyish Brown
20037514637	SC	8	10YR6/2	Light Brownish Grey	10YR5/3	Brown
20037512638	SiCL	7	10YR5/4	Yellowish Brown	10YR4/4	Dark Yellowish Brown
20037514638	LC	7.5	10YR6/4	Light Yellowish Brown	10YR5/4	Yellowish Brown
20037512639	SiC	6	10YR5/2	Greyish Brown	10YR3/2	Very Dark Greyish Brown
20037514639	LC	7	10YR5/2	Greyish Brown	10YR4/2	Dark Greyish Brown
20037512640	CL	6.5	10YR5/4	Yellowish Brown	10YR4/3	Brown
20037514640	LC	8.5	10YR5/4	Yellowish Brown	10YR4/4	Dark Yellowish Brown

SampleID	Texture	pH (field)	Colour (dry)	Colour (dry)	Colour (wet)	Colour (wet)
20037512641	SL	6.5	10YR6/2	Light Brownish Grey	10YR3/2	Very Dark Greyish Brown
20037514641	SC	6	10YR6/2	Light Brownish Grey	10YR4/2	Dark Greyish Brown
20037512642	LS	7	7.5YR4/6	Strong Brown	7.5YR3/3	Dark Brown
20037514642	MC	8	7.5YR4/6	Strong Brown	7.5YR3/4	Dark Brown
20037512643	FSL	6.5	10YR5/2	Greyish Brown	10YR3/2	Very Dark Greyish Brown
20037514643	SC	8.5	2.5Y6/4	Light Yellowish Brown	10YR5/4	Yellowish Brown
20037512644	SL	7.5	10YR5/2	Greyish Brown	10YR3/2	Very Dark Greyish Brown
20037514644	SC	8	10YR6/3	Pale Brown	10YR5/2	Greyish Brown
20037512645	FSL	7	10YR6/6	Dark Yellowish Brown	10YR3/4	Dark Yellowish Brown
20037514645	HC	8.5	10YR6/3	Pale Brown	2.5Y6/3	Light Yellowish Brown
20037512646	HC	7	2.5Y4/1	Dark Grey	2.5Y2.5/1	Black
20037514646	MC	8.5	2.5Y5/1	Grey	2.5Y4/1	Dark Grey
20037512647	CL	7.5	2.5Y5/1	Grey	2.5Y3/1	Very Dark Grey
20037514647	SiC	8	2.5Y6/2	Light Brownish Grey	2.5Y5/3	Light Olive Brown
20037512648	S	6	10YR4/3	Brown	7.5YR3/3	Dark Brown
20037514648	Lfsy	8.5	10YR4/4	Dark Yellowish Brown	10YR3/6	Dark Yellowish Brown
20037512649	SiCL	8	2.5Y5/1	Grey	2.5Y4/1	Dark Grey
20037514649	HC	9	2.5Y5/2	Greyish Brown	10YR4/2	Dark Greyish Brown
20037512650	L	7	10YR5/4	Yellowish Brown	10YR3/3	Dark Brown
20037514650	SiC	8.5	7.5YR5/6	Strong Brown	7.5YR4/6	Strong Brown
20037512651	MC	7	10YR5/2	Greyish Brown	10YR4/2	Dark Greyish Brown
20037514651	SCL	8	10YR5/4	Yellowish Brown	10YR4/3	Brown
20037512652	L	7	7.5YR4/4	Brown	10YR3/4	Dark Brown
20037514652	SiL	7.5	10YR6/4	Light Yellowish	10YR4/4	Dark Yellowish
20037512653	L	7	10YR5/3	Brown Brown	10YR5/3	Brown Brown
20037512653	LC	, 8.5	10YR6/4	Light Yellowish	10YR4/6	Dark Yellowish
				Brown		Brown
20037512654	CL	7.5	10YR5/3	Brown	10YR4/3	Brown
20037514654	LC	8	10YR6/4	Light Yellowish Brown	10YR5/4	Yellowish Brown
20037512655	L	6.5	10YR5/2	Greyish Brown	10YR3/1	Very Dark Grey
20037514655	CL	8.5	2.5Y6/3	Light Yellowish Brown	2.5Y5/3	Light Olive Brown
20037512656	LS	5	2.5Y6/2	Light Brownish Grey	2.5Y4/2	Dark Greyish Brown
20037514656	SiCL	8.5	2.5Y5/2	Greyish Brown	2.5Y4/2	Dark Greyish Brown
20037512657	CL	6.5	10YR6/2	Light Brownish Grey	10YR4/2	Dark Greyish Brown
20037514657	SiC	8.5	2.5Y6/2	Light Brownish Grey	2.5Y5/2	Greyish Brown
20037512658	LC	8.5	7.5YR4/4	Brown	7.5YR3/4	Dark Brown
20037514658	MC	8.5	10YR6/4	Light Yellowish Brown	10YR5/4	Yellowish Brown
20037512659	LC	6.5	10YR5/3	Brown	10YR3/3	Dark Brown
20037514659	CL	8.5	10YR5/4	Yellowish Brown	10YR4/3	Brown

SampleID	Texture	pH (field)	Colour (dry)	Colour (dry)	Colour (wet)	Colour (wet)
20037512660	L	(11010) 7	10YR5/3	Brown	10YR3/2	Very Dark Greyish Brown
20037514660	SiL	8.5	2.5Y6/3	Light Yellowish Brown	2.5Y5/3	Light Olive Brown
20037512661	MC	8	10YR4/2	Dark Greyish Brown	10YR3/2	Very Dark Greyish Brown
20037514661	CL	8	10YR5/3	Brown	10YR4/4	Dark Yellowish Brown
20037512662	L	6.5	2.5Y7/1	Light Grey	2.5Y4/2	Dark Greyish Brown
20037514662	LC	8	2.5Y6/3	Light Yellowish Brown	2.5Y5/3	Light Olive Brown
20037512663	Lfsy	6	2.5Y5/2	Greyish Brown	2.5Y3/2	Very Dark Greyish Brown
20037514663	CL	8.5	10YR6/3	Pale Brown	10YR5/3	Brown
20037512664	CL	7	2.5Y5/2	Greyish Brown	2.5Y3/2	Very Dark Greyish Brown
20037514664	LMC	8.5	10YR5/3	Brown	10YR4/3	Brown
20037512665	CL	8.5	7.5YR4/6	Strong Brown	7.5YR3/4	Dark Brown
20037514665	L	8.5	2.5Y6/4	Light Yellowish Brown	2.5Y4/4	Olive Brown

Appendix 7: Macroscopic properties of overbank sediments form the eastern Riverina (laboratory parameters)

SampleID	Moisture (%)	pH 1:5	EC 1:5 (µS/cm)	SampleID	Moisture (%)	pH 1:5	EC 1:5 (µS/cm)
20037512601	11.50	5.7	28.3	20037512628	18.22	5.4	159.5
20037514601	14.77	6.3	53	20037514628	20.09	5	187.8
20037512602	20.24	5.3	79.7	20037512629	32.31	5.7	80.9
20037514602	15.48	4.8	581	20037514629	24.20	6.9	6.7
20037512603	24.38	6	99.1	20037512630	21.20	5.8	26.7
20037514603	21.04	5.3	1528	20037514630	19.18	7.6	226
20037512604	15.01	5.6	61.1	20037512631	27.17	5.3	93.9
20037514604	22.48	6.2	39.8	20037514631	14.07	9.1	121.4
20037512605	23.30	5.5	76.7	20037512632	19.16	9.3	130.1
20037514605	14.22	8.9	169.8	20037514632	24.17	8.7	515
20037512606	22.35	6.5	22.2	20037512633	9.34	6.4	42
20037514606	22.10	7.8	403	20037514633	20.06	6	198.5
20037512607	15.99	5.6	23.7	20037512634	30.59	6.5	44.9
20037514607	12.91	5.1	252	20037514634	27.13	8.6	288
20037512608	24.47	5.3	60.3	20037512635	24.40	5.8	121.4
20037514608	15.54	7	29.6	20037514635	20.29	7.3	884
20037512609	15.13	6.7	29.9	20037512636	14.32	6.7	61.9
20037514609	14.89	8.3	170	20037514636	19.21	7.4	101.4
20037512610	8.14	6.8	164.5	20037512637	10.61	6.4	32.9
20037514610	20.18	7.1	575	20037514637	15.97	7.6	463
20037512611	22.35	7.2	47.7	20037512638	16.64	7	21.1
20037514611	19.20	7.8	1013	20037514638	15.23	7.9	25.6
20037512612	16.36	7.7	38.8	20037512639	16.16	6.6	34.5
20037514612	21.71	7.9	223	20037514639	20.85	7.1	291
20037512613	16.80	6.9	37.7	20037512640	14.68	7	37.2
20037514613	21.58	9.2	340	20037514640	14.34	8.2	668
20037512614	20.87	7.3	26.7	20037512641	16.73	6.2	94
20037514614	9.15	8.8	76	20037514641	18.59	6.4	104.9
20037512615	22.34	7.5	44.8	20037512642	5.52	7.2	83
20037514615	33.81	7.9	1628	20037514642	16.86	7.9	393
20037512616	15.57	7.6	45.5	20037512643	7.04	7	33
20037514616	23.46	8.6	216	20037514643	7.77	8.6	216
20037512617	19.62	7.6	52.5	20037512644	6.64	7.4	66.2
20037514617	22.99	8.3	1068	20037514644	11.79	8.4	245
20037512618	20.75	7.3	46	20037512645	10.68	7.6	53.3
20037514618	13.47	6.8	250	20037514645	20.92	8.4	733
20037512619	10.23	6.5	34	20037512646	18.75	7.6	25.4
20037514619	15.58	8.7	240	20037514646	18.28	8.8	123.7
20037512620	12.20	5.4	39.4	20037512647	17.56	7.9	78.6
20037514620	14.35	7.2	309	20037514647	17.89	8.7	317
20037512621	11.71	5.5	55.6	20037512648	4.37	6.7	36.5
20037514621	18.87	5	39.7	20037514648	18.09	8.6	382
20037512622	16.14	6.7	14.9	20037512649	36.10	8.1	83.7
20037514622	19.12	7	14.2	20037514649	31.41	9.2	136.4
20037512623	23.65	4.7	49.4	20037512650	7.45	7.4	48.3
20037514623	20.04	7.1	19.6	20037514650	21.13	8.5	463
20037512624	19.86	6.8	52.7	20037512651	20.83	7.5	39.8
20037514624	19.20	6.2	16.1	20037514651	13.34	8.1	240
20037512625	31.42	6	26.5	20037512652	16.52	7.4	100.7
20037514625	21.60	6.7	7.2	20037514652	17.88	7.1	2190
20037512626	22.78	5.8	23.1	20037512653	8.26	5.6	89.3
20037512627	20.94	6.2	22.1	20037514653	12.65	7.8	1734
20037514627	19.40	6.5	8.2				

SampleID	Moisture (%)	pH 1:5	EC 1:5 (µS/cm)	SampleID	Moisture (%)	pH 1:5	EC 1:5 (µS/cm)
20037512654	15.06	7.8	64.8	20037512660	11.10	7.4	63.4
20037514654	18.38	6.9	2890	20037514660	17.17	8.1	1251
20037512655	13.59	7.2	35.7	20037512661	19.81	8.3	97.7
20037514655	12.95	8.8	116.6	20037514661	21.52	7.8	1155
20037512656	4.99	4.7	59.6	20037512662	10.81	7.1	37.5
20037514656	16.69	8.9	391	20037514662	14.26	7.9	160.9
20037512657	11.65	7	47.2	20037512663	5.75	6.1	141.8
20037514657	18.76	8.4	320	20037514663	22.93	7.9	2520
20037512658	9.78	8.5	83.8	20037512664	10.95	7.5	76.9
20037514658	16.85	8.5	805	20037514664	15.76	8.6	349
20037512659	19.98	6.6	35.6	20037512665	11.79	8	1334
20037514659	19.98	8.7	169.9	20037514665	14.61	7.7	3190