INTRODUCTION

In stability but at the Estuarine areas where freshwater encounters seawater are characterized by a lateral variation in salinity, and can represent a transfer box for the sediments between land and the open ocean (Meade, 1972; Burton, 1988). They are very active areas where a huge amount of organic matter and trace metals are introduced into the ocean system through river runoff, in situ primary production and anthropogenic impacts. The behaviour of heavy metals as they pass through estuaries on their way to the ocean is of great interest and concern. Much evidence supports the key role estuaries play as biogeochemical barriers to toxic heavy metals. Although considerable information exists on some of the processes involved (Danielson et al., 1990; Windom et al., 1985), the need for additional data on the behaviour of pollutants in estuaries cannot be overstated. Concentrations of trace metals in estuaries can be elevated due to high inputs from natural and anthropogenic sources. In recent years, the Pahang River has been heavily impacted by discharges from
municipal and industrial outflows. This is due to the rapid development of the area via expansion of the industrialization area as well as the increase in population. Steel and petro-chemicals and automotives are the main industry activities in the area and are the catalyst for other supportive industries to develop around the same area. Very little work has been carried out on the determination of trace metal contents of rivers of Malaysia. The concentration of toxic metals are of interest as most of these rivers are used for the extraction of drinking water as well as their harmful effect to aquatic life at relatively high concentrations. The development of industrialisation and the greater utilisation of land for agriculture may undermine drinking water quality. As there is little published information on the levels of trace metals in riverine systems of Malaysia, the purpose of this study was to measure the background levels of Cd, Cu, Pb and Zn in a relatively clean river in a rural area.

**MATERIAL AND METHODS**

**Sampling Sites**

The study site area is located in the Pekan district which is about 50 km south of Kuantan, the capital state of Pahang (Fig. 1). This area has a humid tropical climate with Southwest and Northeast monsoons bringing an annual rainfall which varies between 1488 mm to 3071 mm. A total of 62 surface and bottom water samples were collected in this study. Stations 1 to 5 in the estuary represent the most seaward while station 52 to 63 inside the Pahang River, represent the landward most less saline water point. The samples were collected using the mercos water sampler and 1 liter of sample was collected at each station and was transferred into an acid cleaned polyethylene bottles. Sampling was done during high tides, twice during the dry season (July 2008 and May 2009) and twice during the rainy season (September 2008 and February 2009). Water samples were then acidified to pH 2, with HNO₃ (Suprapure), stored at 4°C until treatment and analysis were performed. In the laboratory, all samples were pressure filtered through a pre-weighted, acid rinsed 0.45 µm nucleopore polycarbonate filter, located in a clean air cabinet. Filtrates were transferred into a acid-rinsed polyethylene bottles and acidified to pH 2, using HNO₃, prior to trace element analysis. In this study seasonal changes in physico-chemical parameters of water were also monitored. The physical parameters (salinity, temperature and pH) for both the surface and bottom layers were obtained in situ by using the hydroLab. HydroLab were calibrated according to standard procedures before any data measurements were made in the field.

Analyses of metals were performed in a clean air cabinet (Class 100). Dissolved Cu, Pb, Zn and Cd were extracted using an ammonium pyrolidine dithiocarbamate/methyl isobuthyl ketone (APDC/MIBK) extraction into freon, and analysed by graphite furnace atomic absorption spectrophotometry (GFAAS) accordind to methods described by Danielsson et al (1982). The validity of the analytical techniques was assessed by analysis of blank and the standard reference materials SLRS-1 (Seawater and Riverine Water Reference Material for Trace Metals, supplied by the National Research Council of Canada). The contents of each element matched the certified values within 5% and the recoveries were obtained typically between 5 – 10%.

**RESULTS AND DISCUSSION**

Little information is available on the physico-chemical aspects of the Pahang River estuary. As for the tropical estuaries, their environmental characteristics are much influenced by tidal rhythm and monsoon cycle. The characteristic features of the Pahang River estuarine system are follow closely the pattern of river runoff and tidal variations and were mostly controlled by the monsoon seasons. According to the twenty years (1968 to 1987) accumulated data obtained from the Malaysian Meteorological Service, the monsoon seasons with strong winds with mean annual rainfall for 3064 mm occurred from November to January. Meanwhile the non-monsoon seasons with low rainfall occurred during April, May and June.

**Salinity**

Salinity varies from 0 – 32 ppt during non-monsoon seasons and 0 – 26 ppt during monsoon seasons. Results show that salinity at stn. 31 and upper stations are near the zero threshold throughout the year, signifying the fresh water region. Adopting the index of salt water penetration...
as 0 ppt, it is evident from Fig. 2a and 2b that the sea water intrusion is effective up to the maximum of 12 km (station 28, 29 and 30) in non-monsoon seasons and 5 km (station 17, 18 and 19) in the monsoon seasons (Fig. 2a and 2b). The relative lower salinity during monsoon seasons is due to dilution activity resulting from freshwater influx and land drainage into the riverine system. However, on many occasion the difference in salinity of surface and bottom layers is quite significant (P<0.05) but the bottom water is more saline than the surface water. The increase in salinity of both surface and bottom layers also indicates a lateral salinity gradient over most parts of the estuary.

**Temperature**

The temperature of surface and bottom waters are almost uniform or even sometimes they are identical (Fig. 3a and 3b). The magnitude of apparent variation of temperature in the water column is insignificant. There has been a gradual decrease in temperature from the mouth upstreams along the estuarine system reaching the riverine system in both seaasons. The mixed layer depth shoals and resulting detrainment inhibits the incoming surface heat flux from being transported downward and elevating the sea surface temperature (Wang et al, 2000; Wang et al., 2006a). These findings are in general agreement with that of Day (1981), where rivers are usually cooler than the coastal waters in the estuarine system.

**pH**

A small range of pH changes was encountered in the Pahang River River (Fig.4a and 4b). During non-monsoon seasons, pH values were relatively higher in the estuary (pH 7 – 8) due to the influence of seawater penetration and biological activity (Day, 1981), followed by lower pH in the monsoon seasons (pH 5 – 6), which may due to the lower primary productivity (Day, 1981). The lower pH during the monsoon seasons might also be due to the influence of freshwater influx and decomposition of organic matter (Day, 1981; Wang et al, 2000) carried by flood waters into the riverine system. This phenomenon was also reported by Law and Jong (2006), whereby more acidic pH was observed during pre-monsoon season.

**Trace metal concentrations**

The trace metals measured in the Pahang River are summarized in Table 2. In general, neither salinity nor the concentration of Cu, Pb, Zn and Cd in the Pahang River varied significantly during the seasonal changes. However, relatively lower concentrations during non-monsoon seasons were

<table>
<thead>
<tr>
<th>Metals</th>
<th>Monsoon seasons</th>
<th>Non-monsoon seasons</th>
</tr>
</thead>
<tbody>
<tr>
<td>Copper (Cu)</td>
<td>1.01 ± 0.13</td>
<td>0.57 ± 0.12</td>
</tr>
<tr>
<td>Lead (Pb)</td>
<td>1.34 ± 0.25</td>
<td>0.88 ± 0.45</td>
</tr>
<tr>
<td>Zinc (Zn)</td>
<td>12.8 ± 1.65</td>
<td>9.96 ± 0.78</td>
</tr>
<tr>
<td>Cadmium (Cd)</td>
<td>0.18 ± 0.04</td>
<td>0.09 ± 0.05</td>
</tr>
</tbody>
</table>

**Table 1: Average concentration of dissolved trace metals (µg/l^-1) along the Pahang River, showing a relatively lower metal concentrations during non-monsoon seasons**

<table>
<thead>
<tr>
<th>MetalLocation</th>
<th>Cu</th>
<th>Pb</th>
<th>Zn</th>
<th>Cd</th>
</tr>
</thead>
<tbody>
<tr>
<td>ZTerengganu River</td>
<td>0.11 – 1.49</td>
<td>1.67 – 12.9</td>
<td>1.32 – 9.65</td>
<td>0.17 – 0.65</td>
</tr>
<tr>
<td>Klang River</td>
<td>10</td>
<td>4.1</td>
<td>17.9</td>
<td>-</td>
</tr>
<tr>
<td>Bristol Channel</td>
<td>1.02 – 4.74</td>
<td>0.35 – 5.06</td>
<td>n.a</td>
<td>n.a</td>
</tr>
<tr>
<td>Pahang River(This Study)</td>
<td>0.57 – 1.01</td>
<td>0.88 – 1.34</td>
<td>9.96 – 12.8</td>
<td>0.09 – 0.18</td>
</tr>
</tbody>
</table>

**Table 2: The concentration of dissolved Cu, Pb, Zn and Cd in study area compared to other estuaries**
Fig. 1: Location of the study area showing sampling sites (62 stations) along Pahang River- Estuary, Pahang

Fig. 2a: Salinity gradient during pre-monsoon of Pahang River estuary

Fig. 2b: Salinity gradient during post-monsoon of Pahang River estuary
Fig. 3(a): Temperature gradient during pre-monsoon of Pahang River estuary

Fig. 3(b): Temperature gradient during post-monsoon of Pahang River estuary

Fig. 4(a): pH gradient during post-monsoon of Pahang River estuary
Fig. 4(b): pH gradient during pre-monsoon Pahang River estuary

Fig. 5: Dissolved metal concentrations (Zn & Cd) versus salinity in the Pahang River showing a conservative behaviour with statistical value of P<0.05.

a) sampling during non-monsoons and b) sampling during monsoon seasons
shown compared to the monsoon seasons. This could be result of successive dilution between two elemental sources of different origin, presumably identified as anthropogenic discharges and watershed flushing. The former would predominate during the non-monsoon seasons, and the latter during the monsoon seasons.

The lowest concentrations of dissolved metals measured in Pahang River estuary during the monsoon seasons were Cd (0.18 ± 0.04 µg l⁻¹) followed by Cu (1.01 ± 0.13 µg l⁻¹), Pb (1.34 ± 0.25 µg l⁻¹) and Zn (12.8 ± 1.65 µg l⁻¹). For all metals, as expected the lowest levels were found at stations away from the estuary. Furthermore, these stations are isolated from any human activities as it is within the Hulu Pahang Forest Reserve. For non-monsoon seasons, the concentrations of Cu (0.57 ± 0.12 µg l⁻¹) and Cd (0.09 ± 0.05 µg l⁻¹) showed relatively low concentrations, followed by Pb (0.88 ± 0.45 µg l⁻¹) and Zn (9.96 ± 0.78 µg l⁻¹). Some of the highest concentrations obtained in some stations are associated with water of low salinity and they may be attributed to anthropogenic inputs of the industrial area nearby. Meanwhile the lower metal concentrations near the lower estuary may be explained by the mixing of the contaminated runoff from land-based source and relative uncontaminated water from the sea in the mixing zone at the estuary itself. Variations between geographical areas were significant (P<0.05), when compared with some other estuaries. The concentration of dissolved Cu, Pb, Zn and Cd in the Pahang River were relatively higher than Terengganu estuary (Kong, 2001), and are generally lower than those reported by Law and Singh (1986) in the Klang river and Abdullah and Royle (1974) in the Bristol channel (Table 2).

In this study, concentration of dissolved metal of Cu, Pb, Zn and Cd were found to be decreased as the water salinity increased downstream. This trend is similar to many previous studies in other estuaries (Salomons et al. 1981; Zwolsman et al. 1997). In general, concentrations of Cu and Cd during non-monsoon seasons also show a good correlation (r² > 0.9) with salinity, meanwhile Cu during monsoon seasons, Pb and Zn during both seasons has a fairly correlation (r² = 0.7 – 0.8) with salinity (Figure 5). It is well known that these distributions are related primarily to physical mixing of fluvial and marine particulars. The decrease in trace metal content with the increasing salinity simply reflects the increase in the marine component in water (Zwolsman et al. 1997). It might be argued that the decrease in trace metals can also be due to desorption processes, which was demonstrated in the Scheldt estuary for Cd, Cu and Zn (Salomons et al., 1981). However, studies have stated that desorption for other trace metals in estuaries are insignificant (Zwolsman et al., 1997).

REFERENCES

7. Law A T and Singh A., Distribution of Mn,


