

Physico-Chemical Analysis in Surface Waters around the Closed Gaborone Sanitary Landfill in Botswana

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Abstract The aim of the study was to ascertain the pollution levels in water sources in the areas surrounding the closed Gaborone landfill site. The study focused on the physico – chemical analysis of surface water resources around the closed Gaborone landfill site. The specific objectives were to determine the properties of surface water around the landfill and compare them along a transect with the water sources upslope and downslope of the landfill. Furthermore, the probable impacts of the wastes on water resources are highlighted and the levels of heavy metal contamination in surface waters around the closed Gaborone landfill are shown, in addition to compare the findings in this study with set standards (WHO, USEPA, FAO, EU, USSR and BOBS) and other yardsticks from previous studies. Due to scarcity of water resources and poor drainage water levels, five (5) existing surface water samples were collected offsite along a spatial gradient transect while the sampling interval was based on the length of the slope below the landfill. Field sampling and laboratory analysis of surface water resources was done so as to ascertain physico – chemical and heavy metal pollution levels. The findings of the investigations show that physical parameters such as pH, Electrical Conductivity, turbidity, TDS and TCU levels at the closed Gaborone landfill are above the drinking water standards BOS 32: 2000, WHO (2004) and USEPA (1991) limits and there is a general decline in pH, EC, TSS, TVS, TSD and TCU with increasing distance from the landfill site. While the chemical properties show that alkalinity, nitrates, phosphates, sulphates, chlorides, calcium, magnesium, chromium, and ammonia levels for the closed Gaborone landfill are higher than BOS 32: 2000, WHO (2004) and USEPA (1991) limits indicating toxicity. The general trend patterns show that there is a general decline in alkalinity, BOD₅, sulphates, phosphates, nitrates, magnesium, calcium and chromium levels with increasing distance from the closed Gaborone landfill while chlorides levels are increasing. There was a significant decline in alkalinity, EC, sulphates, and calcium with increasing distance at $P = 0.05$. On the other hand, BOD₅ levels can be classified as

clean to moderately polluted. Overall, sanitary landfills have a far-reaching impact on the surrounding water resources and if left unmonitored increased pollution levels could lead to compromised drinking water quality, public health deterioration and descending environmental eminence.

Keywords Gaborone Landfill, Heavy Metals, Physico-Chemical, Surface Waters, Water Resources

1. Introduction

Initiatives in solid waste management are growing phenomenon in developing nations that have created employment (Areola, Segosebe and Gwisai, 2015; Strange, 2002; Abdelatif, 2001; Abd Malek *et al.*, 1996). On the other hand, the under – privileged have found opportunities to earn a living from landfill sites (Areola, Segosebe and Gwisai, 2015; Manyanhaire *et al.*, 2009; Masocha and Tevera, 2003). However, it has been observed that most employees in the sector have limited to no personal protective equipment. This poses serious risks on health hazards (Areola, Segosebe and Gwisai, 2015; Noel, 2010). Several studies note that limited attention has been granted towards investigations involving human health risks to which scavengers are exposed to (Areola, Segosebe and Gwisai, 2015; Noel, 2010; Chattopadhyay, Dutta and Ray, 2008; Mwiganga and Kansime, 2005; Chofqi, *et al.*, 2004) and to the human health concerns of the communities residing in the neighbourhoods of landfills (Gwebu, 2003). According to Elliot (2006), 80% of the population in developed nations live within a 2 – kilometre radius of a current or closed landfill site and have experienced serious health effects. In this regard, the closed Gaborone landfill site in Botswana, has been criticized poor waste operations and management (The Botswana Gazette, 2010; Ngole, 2000). In Botswana, generally there are fewer engineered landfill sites than dump sites (Gwebu, 2003); a situation

which makes the environmental health risks associated with waste disposal a major public concern.

Although sanitary landfill sites have been observed to constitute a potential hazard to the environment (Khan and Agarwal, 2006; Misra and Pandey, 2004; Christensen and Christensen, 1999; Moyo *et al.*, 1993), in many cases, the effects of the landfills are not easily discernible. The pertinent effects include ground and surface water pollution (Areola, Segosebe and Gwisai, 2015; Odukoya and Abimbola, 2010; Miller, 1996). Health risk related studies on employees in the handling, transporting, clean-up or maintenance of substances at landfill sites have been found to be very scarce. Yet, many chemicals present in landfill sites have been shown to have adverse effects on human health (Vrijheid, 2000). Hence, the focus of this study was to examine the environmental pollution challenges in surface water resources in and around municipal landfill sites as revealed in some studies (Areola, Segosebe and Gwisai, 2015). The concern for the plight of surface water resources near landfills is partly due to the lower levels of public awareness and respect for public opinion in decision making in developing countries as compared to what obtains in the developed world. Indeed, there are instances in some countries where governments and government officials have deliberately accepted the dumping of hazardous wastes at some locations in the developing world (Areola, Segosebe and Gwisai, 2015).

The aim of the study was to investigate and evaluate the surface water pollution levels of surface water resources nearby the closed Gaborone Municipal landfill. These investigations were based on the Gaborone landfill in Botswana in the southern African region of the continent. The specific objectives of the study were to determine the level of contamination, around the landfill sites by analyzing specifically the types and levels of physico – chemical elements pollution and heavy metal contaminants in surface water sources around the Gaborone landfill site. Furthermore to establish how physico – chemical properties and heavy metal concentrations in the closed Gaborone surface water sources compare with other yardsticks from previous studies and set standards (e.g. Areola, Segosebe and Gwisai, 2015; Matsa and Mutekwa, 2009; Odukoya and Abimbola, 2010).

Waste management has been one of the core values of the National Development Plan, NDP 10, in Botswana, which aimed at achieving the sustainable development goal. Thus, this study would contribute to knowledge specifically by helping to identify the primary chemical and heavy metals that constitute the major hazards. Such knowledge would help governments and international

organizations to develop appropriate mitigation or otherwise intervention measures as an aftermath of the decommissioning of landfills and dumpsites. Urbanization levels in Gaborone could have a significant influence on waste types and quantities and therefore pollution levels of water resources in the surrounding environment as shown in other studies (Areola, Segosebe and Gwisai, 2015). With the notion of almost two decades after the development of the landfill disposal ideology, legislation and implementation in 1998, it is necessary to carry out an audit on water contaminations related issues that will contribute to new knowledge on the impacts of landfills specifically on the surface water resources as recommended by other studies (Areola, Segosebe and Gwisai, 2015; Ngole, 2000).

2. Study Area

Botswana is located in the southern part of the African continent. It is bounded to the south and southeast by South Africa, to the west by Namibia, to the north by Zambia and to the northeast by Zimbabwe. The closed Gaborone landfill site was commissioned in 1993 close to the Gaborone dam and the Notwane river catchment area is one of the landfill sites that was commissioned before landfill legislature in 1998 (Government of Botswana, 1998). The landfill which served the capital city of the country was commissioned without a landfill leachate liner while management operations were poor and done haphazardly (Gwebu, 2003). Therefore the main focus of the study was to assess landfill site impacts, and the extent to which these pollutants have contaminated the environment and human health (Ngole, 2000). Gaborone landfill appeared to lack a lot of equipment that could have ensured a smooth running of operations, and this led to shortage of space in waste cells, possibilities of finding an admixture of items in the same zone of the landfill, and rare waste compaction (Ngole, 2000; Gwebu, 2003). The closed Gaborone landfill site is undergoing rehabilitation. Assessing the effects of landfill pollutants at this stage of the landfill site may act as a guide to future landfill rehabilitation programmes, which Local Authorities (LAs) and the central government may wish to take note of as this may reduce operational costs of landfill sites and help to develop sustainable economic budgets. The temperatures and rainfall have an influence on the biochemical reactions taking place in the landfill sites. This results in varying pollutant production rates affecting the environment and human health (Areola, Segosebe and Gwisai, 2015; Lobatse Town Council, 2002).

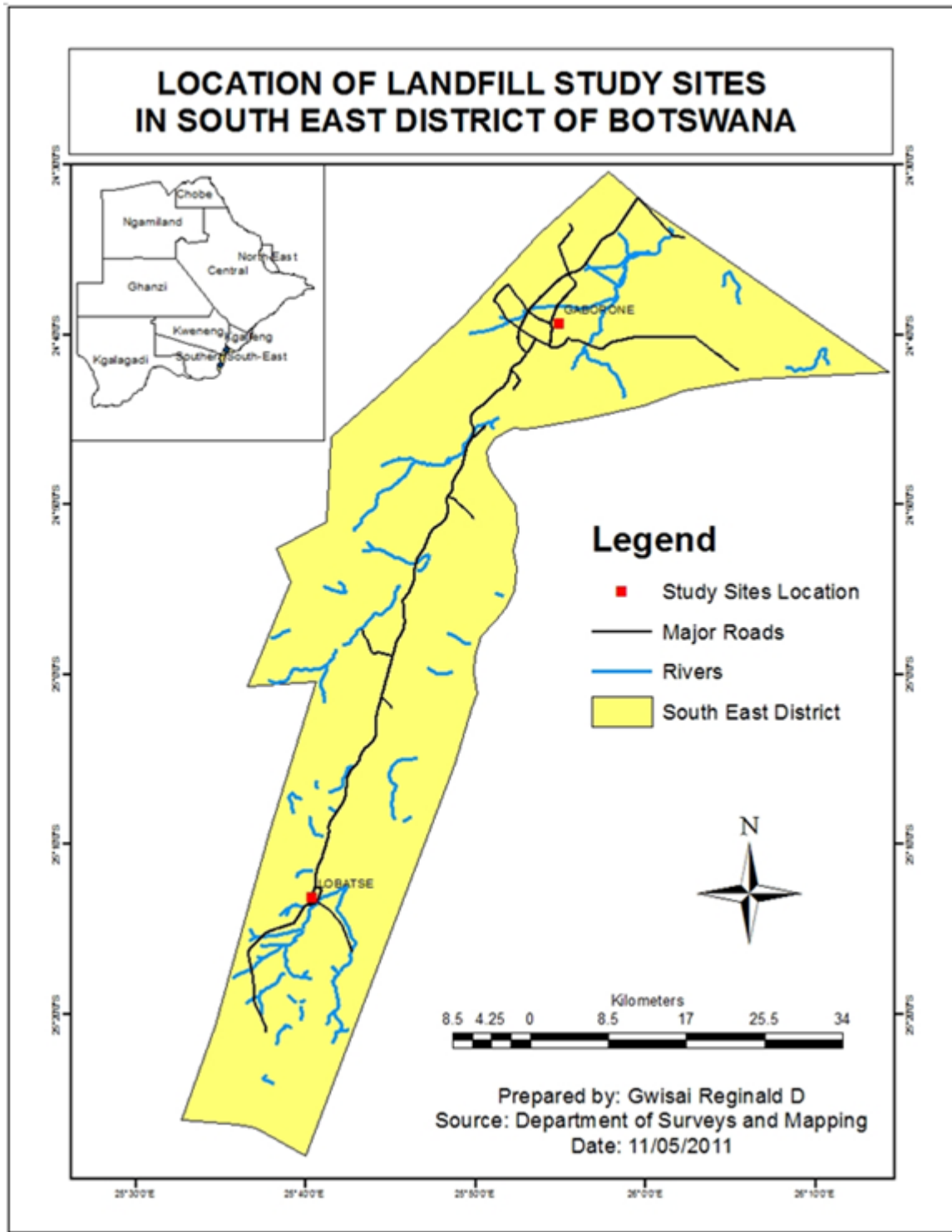


Figure 1. Map of Study Area

3. Materials and Methods

Due to scarcity of water resources and poor drainage water levels, five (5) existing surface water samples were collected offsite along a spatial gradient transect while the sampling interval was based on the length of the slope below the landfill. Field sampling and laboratory analysis of surface water resources was done so as to ascertain physico – chemical and heavy metal pollution levels. The basic principle followed was sampling from the landfill down to the foot of the slope below it. Surface water samples were collected (bulk sampling) from the existing water sources around the landfill sites A control sample from upslope was collected for comparative purposes. Figure 2 shows the sampling points at varying distances. The parameters measured were pH, colour (TCU), total solids (TS), total suspended solids (TSS), total dissolved solids (TDS), Biological Oxygen Demand (BOD_5), Chemical Oxygen Demand (COD), alkalinity, Biological Oxygen Demand(BOD_5), Chemical Oxygen Demand(COD), alkalinity, conductivity (EC), phosphates (PO_4^{3-}), sulphates (SO_4^{2-}), ammonia (NH_3), nitrates (NO_3^-) and chlorides (Cl^-), (APHA, 1995). Pollutant characterization may involve both quantitative and qualitative determination of heavy metal cations. The following metals were assessed; iron ($^{56}_{26}Fe^{2+}$), manganese ($^{55}_{25}Mn^{2+}$), copper ($^{64}_{29}Cu^{2+}$), zinc ($^{65}_{30}Zn^{2+}$), lead ($^{207}_{82}Pb^{2+}$), chromium ($^{52}_{24}Cr^{6+}$), nickel ($^{59}_{28}Ni^{2+}$), sodium ($^{23}_{11}Na^+$), magnesium ($^{24}_{12}Mg^{2+}$), potassium ($^{39}_{19}K^+$), calcium ($^{40}_{20}Ca^{2+}$) as these have been found to be common in other related landfill studies conducted earlier (Areola, Segosebe and

Gwisai, 2015; Nwachukwu *et al.*, 2010). The main analytical tool used was the Atomic Absorption Spectrophotometer (AAS). The Flame AAS was used to determine the concentration of the metals selected above. The AAS technique basically involves the principle of free atoms in elements that will absorb light at wavelength characteristics of that element which is determined by the outer electron structure (Areola, Segosebe and Gwisai, 2015; Ngole, 2000; Alloway, 1995). The amount of light is directly proportional to the concentration of the element in solution. The absorbance is normally measured and is used to determine the concentration of the specific element. Normally the solution is atomized in a flame in the Flame AAS.

4. Results and Discussions

Surface Water Physico-Chemical Pollution

Five (5) surface water samples were measured for pH, which ranged from 7.88 – 8.75 (Figure 2). All the samples recorded higher pH values than the standard pH values of 6.5 patently rendering them unsuitable for drinking purposes (Table 2), (BOS 32:2000; WHO, 2004; USEPA, 1991). The highest pH value recorded was 8.39 at a distance of 0.2 km downstream which is similar to other studies. However, there is complementary literature that shows surface water sources in other landfills having pH levels in the acidic to alkaline range which also signals unsuitability for drinking and pollution of the water sources (Areola, Segosebe and Gwisai, 2015; Longe and Balogun, 2010; Raman and Sathiya – Narayanan, 2008; Longe and Enekwechi, 2007; Haertling, 1989).

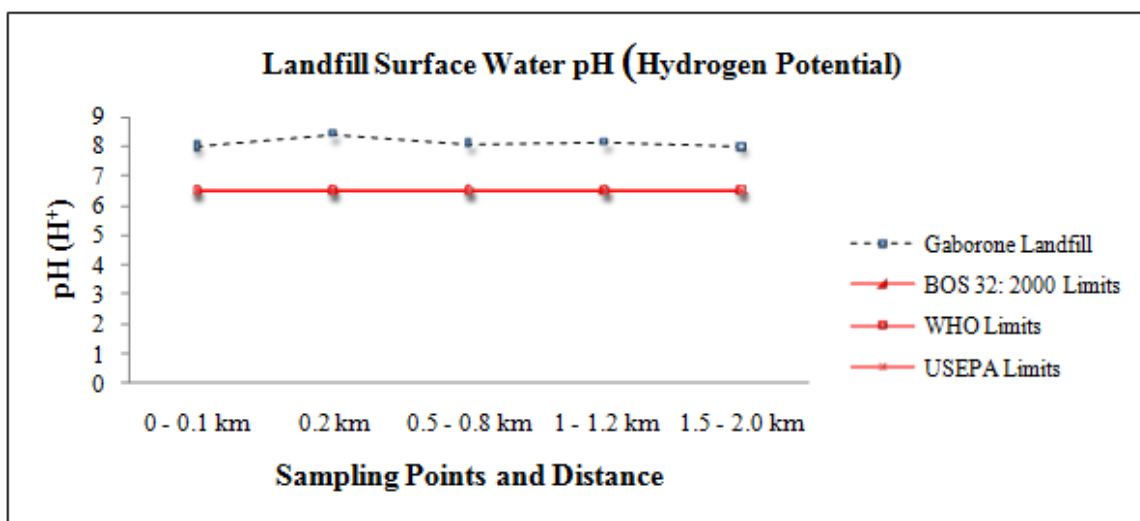


Figure 2. Landfill Surface Water pH

EC ($\mu\text{S cm}^{-1}$) Levels

Electrical Conductivity (EC) was measured at 25°C and all the surface water samples were in the range of 189.2 – 1577.70 $\mu\text{S cm}^{-1}$. All the samples were below the recommended drinking water standard BOS 32: 2000 value of 3100 $\mu\text{S cm}^{-1}$ (Figure 3) as observed by Areola, Segosebe and Gwisai (2015). This could have been because the Lobatse landfill is in the same geographical area and have similar rainfall and temperature patterns. On the other hand, this is contrary to findings by previous studies which found EC levels to be above the set EU limits (Table 2), (Alslaihi, Mogheir and Afifi, 2011; Al – Rawas and Valeo, 2011; Osei *et al.*, 2011; Al – Sabahi *et al.*, 2009; Salminen, 2005; Jeevan Rao and Shantaram, 2003).

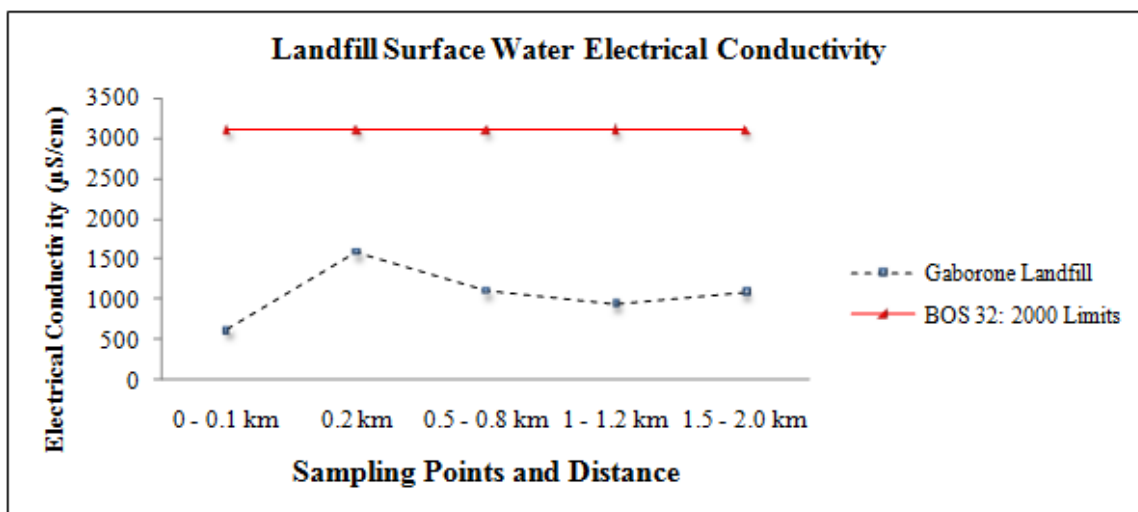


Figure 3. Landfill Surface Water Electrical Conductivity

Colour (TCU) Levels

The colour of surface water samples was measured using the True Colour Units (TCU), and all the samples were in the range 11 – 18 TCU with some samples surpassing the set drinking water standards of 15 TCU (BOS 32:2000; USEPA, 1991). At Gaborone only one sample (0.2 km) surpassed the standard (Figure 4). There is a general decline in colour units with increasing distance from the landfill signaling limited spatial influence of anthropogenic activities further away from the landfill sites. The trend is similar to that of other studies on wells around landfill sites (Areola, Segosebe and Gwisai, 2015; Akinbile, 2012; Raman and Sathiya – Narayan, 2008).

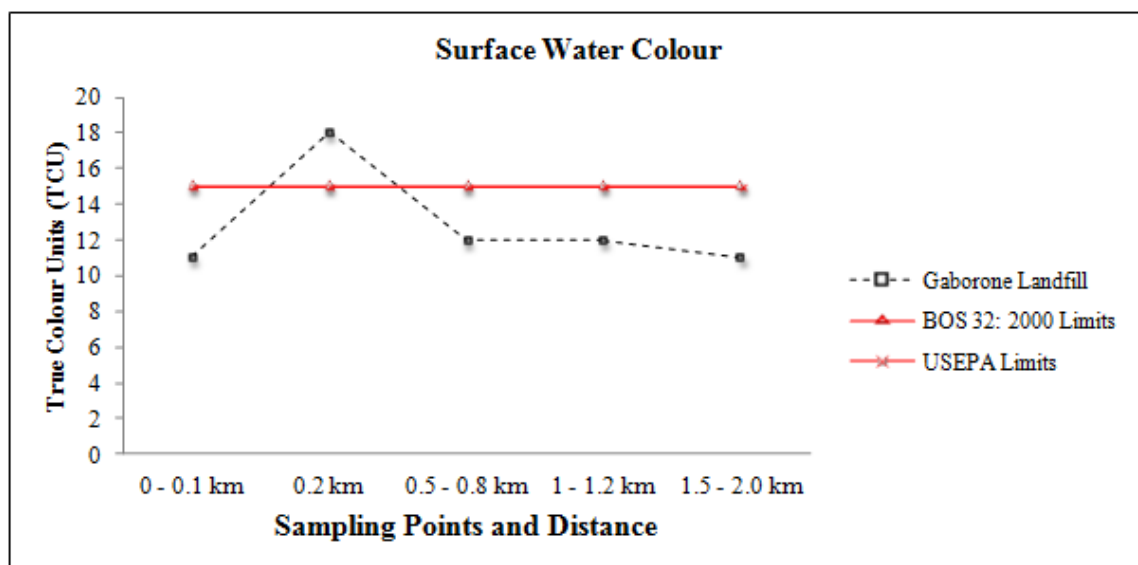


Figure 4. Surface Water Colour

Total Solids (TDS, TSS, TVS, TFS, TSD) Levels

The range of Total Dissolved Solids (TDS) is between 110 – 1015 mg which appears similar to those of previous studies (Akinbile, 2012; Longe and Enekwechi, 2007). All samples from Gaborone landfill have total solids more than the set drinking water standards (450 – 500 mg) (see Figure 5 and Table 5.7). The high amounts of TDS could decrease water palatability and cause gastro – intestinal irritation in humans (Al – Sabahi *et al.*, 2009; WHO, 2004). However, some studies showed TDS values lower than the set drinking water standards (Areola, Segosebe and Gwisai, 2015; Longe and Balogun, 2010; NSDQW, 2007; WHO, 2004).

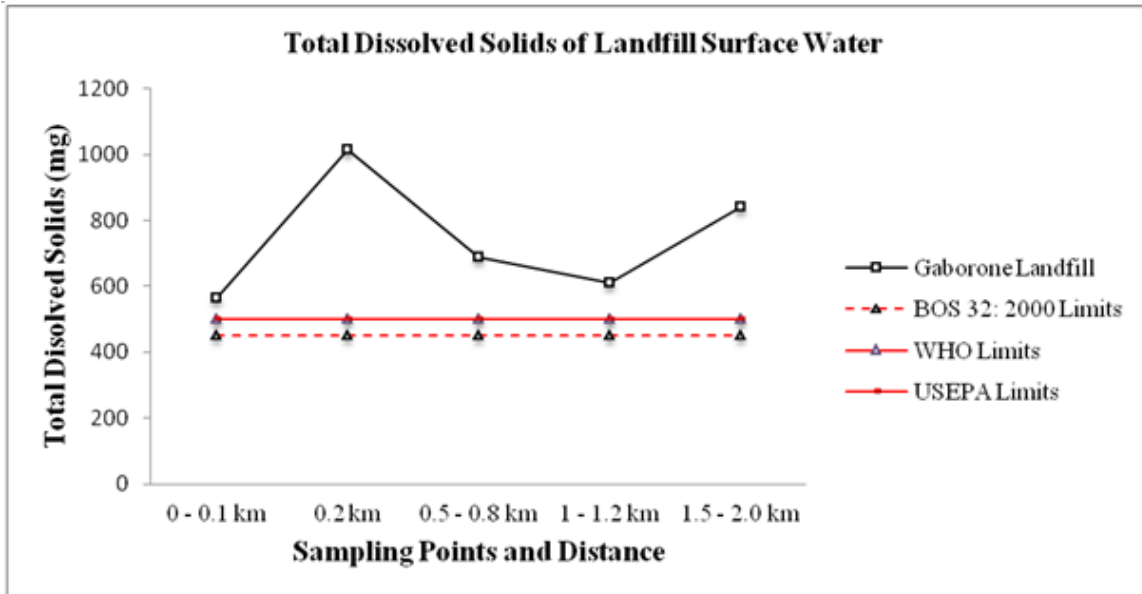


Figure 5. Total Dissolved Solids of Landfill Surface Water

Total Suspended Solids (TSS) in Gaborone (Figure 6) are below the findings in prior studies (Areola, Segosebe and Gwisai, 2015; Raman and Sathiya – Narayanan, 2008). However, there is a general decline in TSS with increasing distance from the landfill, perhaps as the influence of the waste dump and other anthropogenic factors wears off (Bada and Olarinre, 2012; Joel and Amajuoyi, 2009; Domska and Warechowska, 2008).

The Total Volatile Solids (TVS) are in the same range for all samples in Gaborone landfill. Gaborone samples show a significant gradual increase in TVS with increasing distance from the landfill ($r = 0.9244$, $p = \leq 0.05$ with a $R^2 = 0.8545$), (Figure 6). Again this could also be attributed to the amounts of wastes disposed of at the landfills and other independent human activities directly related to waste disposal at the Gaborone landfill site (Osei *et al.*, 2011; Edosomwan and Onwumah, 2008; Wong, 1988).

There is a limited amount of Total Fixed Solids (TFS) in all Gaborone samples (0.0032 – 0.0098 mg), (Figure 6). This could be due to limited spread of materials in semi – arid environments such as Botswana (Alslaibi, Mogheir and Afifi, 2011).

Also for Total Solids Dried (TSD) the range is 460 – 1040 mg and there is an increase in the amount of TSD with increasing distance at Gaborone landfill site (Figure 6). The influence of other media (other than the landfill) in transporting waste and soil materials into surface waters could be prevailing in increasing TSD load (Miguel *et al.*, 2012; Johnson and Zhang, 2012; Wuana and Okieimen, 2012; Alslaibi, Mogheir and Afifi, 2011).

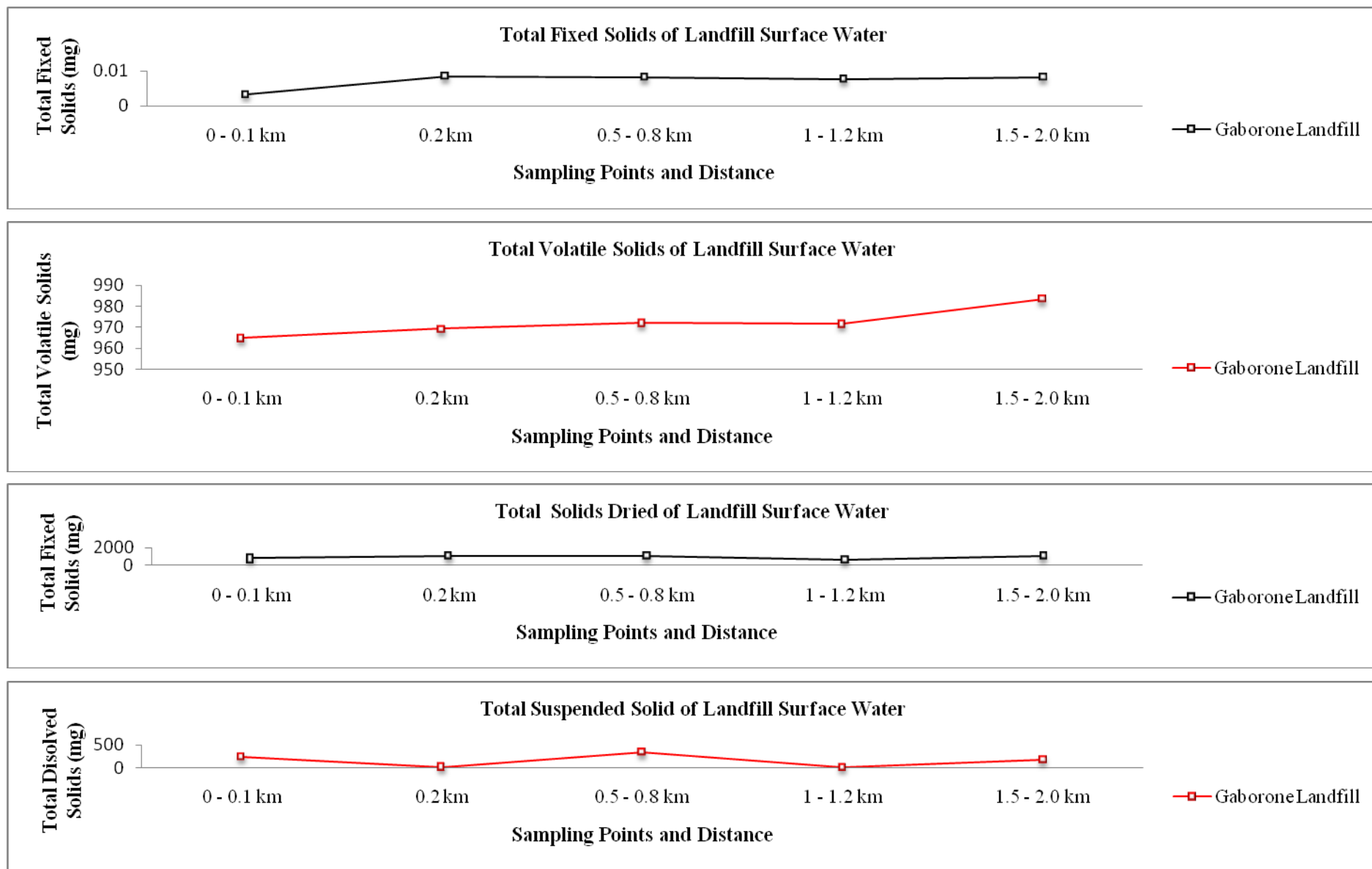


Figure 6. Gaborone Landfills Solids Levels

Turbidity (NTU)

Turbidity was measured using Nephelometric Turbidity Units (NTU) and the samples from Gaborone have a turbidity range of 1.65 – 11 NTU (Figure 7). Some studies show a higher turbidity than the present study signaling a lesser impact universally (Osei *et al.*, 2011). On the other hand studies in Lobatse show a lower NTU levels although in a similar range (Areola, Segosebe and Gwisai, 2015). Among the Gaborone samples the level of turbidity increased downstream with increasing distance from the landfill. The level of pollution in Gaborone is higher than that of other studies (Areola, Segosebe and Gwisai, 2015), while on the other hand it increases with distance from the landfill as opposed to other studies observed (Areola, Segosebe and Gwisai, 2015; Akinbile, 2012). This could be attributed to several factors chief being other sources around the surface water points independent of the landfill which could have produced more waste materials leading to high turbidity values (Loncnar *et al.*, 2010; Tamunobereton – ari, Omubo – Pepple and Tamunobereton – ari, 2010; Schenato, Schroder and Martins, 2008; Wuana and Okieimen, 2012; Osei *et al.*, 2011; Alslaibi, Mogheir and Afifi, 2011; Steinnes *et al.*, 1997). Also perhaps to a lesser extent this could be due to the influence of the landfill considering the volume of waste disposed of at the Gaborone landfill and a steep gradient from the landfill mounds to the surface water

points.

The majority of samples have an unobjectionable odour except one sample from Gaborone landfill at 0.2 km (Figure 7). This differed from the findings of other studies (Areola, Segosebe and Gwisai, 2015; Akinbile, 2012) where most of the water samples in wells around landfills had a mild odour or none at all. All turbidity readings are above the BOS 32: 2000 drinking water limits of Botswana signifying pollution of surface water around both landfills (Table 2). For Gaborone, some of the samples are above the USEPA, WHO and BOS 32: 2000 limits (Table 2), (Raman and Sathiya – Narayanan, 2008; WHO, 2004; Radojevic and Bashkin, 1999; USEPA, 1991). The high turbidity values could be an indication of pollution from other sources and the landfill as observed in other studies (Shyamala, Shanti and Lalitha, 2008) as both the landfill and surface water boundaries are not lined. Another observation is that waste and soil cover material could find its way to the surface water just as observed by prior studies hence increasing turbidity as these are not protected water reservoirs due to proximity to the landfill (Akinbile, 2012; Mohamed *et al.*, 2009; Jaji *et al.*, 2007). Generally Gaborone landfill have polluted the surface water around them and the water may need to be treated before any form of use is recommended as observed in other studies (Areola, Segosebe and Gwisai, 2015; Akinbile, 2012; Ogedengbe and Akinbile, 2004).

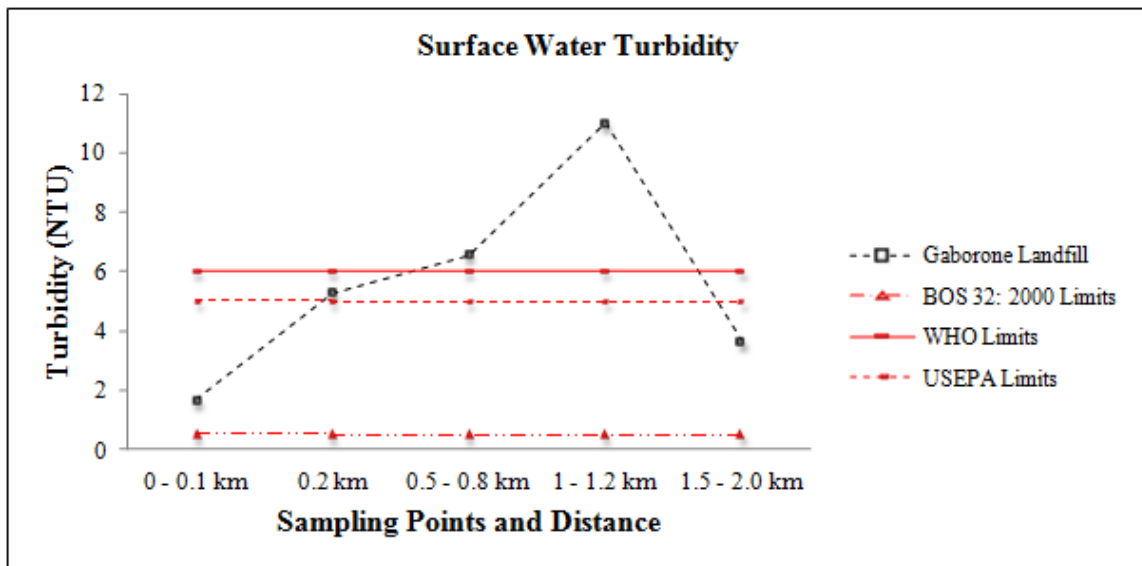


Figure 7. Surface Water Turbidity

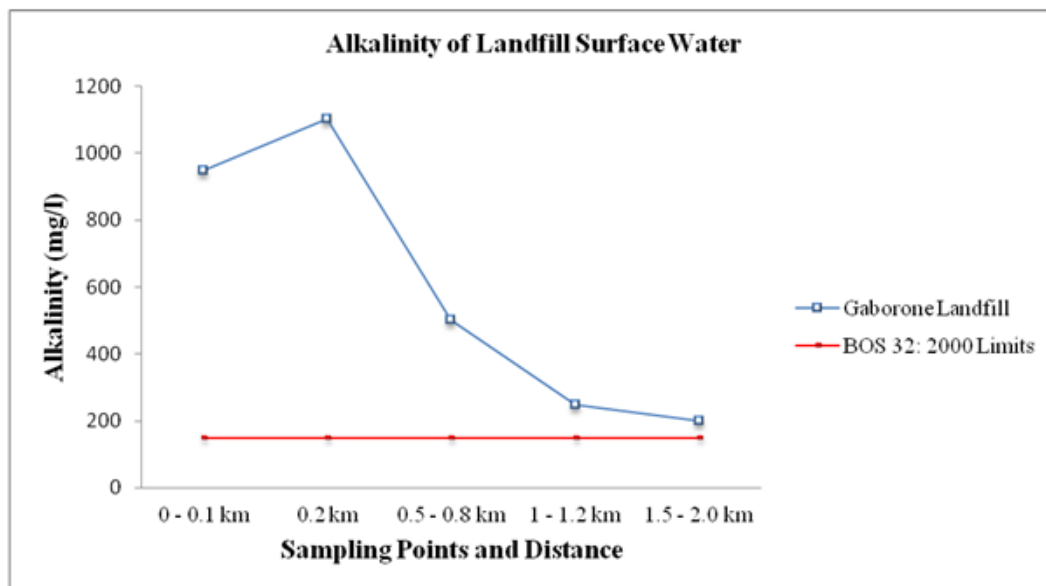
5. Chemical Pollution Levels of Surface Water Samples

Alkalinity

Alkalinity of the Gaborone samples is in the range, 200 – 1101 mg.l^{-1} . The majority (90%) of the samples are above the set drinking water alkalinity limits for Botswana (Figure 8 and Table 2) and also above other countries' thresholds (150 mg.l^{-1}), indicating toxicity in surface water around the landfill sites (Jeevan Rao and Shantaram, 2003; BOS 32: 2000; Le Seur Spencer and Drake, 1987; WHO, 1984; ISI, 1983). Gaborone samples have higher alkalinity levels than the previous studies (Areola, Segosebe and Gwisai, 2015), although they are similar trend to previous studies conducted (Areola, Segosebe and Gwisai, 2015; Raman and Sathiy – Narayanan, 2008). The alkalinity levels could be largely attributed to waste disposed of at the landfill sites (Bhambulkar, 2011). Furthermore, there is a significant general decline in alkalinity levels with increasing distance from the landfill sites ($r = -0.9382$, $p = \leq 0.05$ with a $R^2 = 0.8802$), as other studies observed (Miguel et al., 2012; Wong and Leung, 1989).

Biological Oxygen Demand (BOD_5)

The Biological Oxygen Demand (BOD_5) measured among the surface water samples is from 1.4 – 2.5 mg.l^{-1} for all the samples at the landfill site (Figure 9). The levels are below the set standard in Russia of 3.0 mg.l^{-1} although they have a similar range to other studies elsewhere (Areola, Segosebe and Gwisai, 2015). On the other hand Gaborone BOD_5 levels are also below the levels in other surface waters studied by others (Osei *et al.*, 2011; Sholichin, 2012; Alslaibi, Mogheir and Afifi, 2011; Radojevic and Bashkin, 1999). This shows that samples from the landfill are in the range of between clean and moderately polluted (Table 1). However, as to be expected the general trend for shows that there is a decline in BOD_5 downstream with increasing distance from the landfill sites. As observed by Husain, Hoda and Khan, (1989) there is an increase in the concentration of pollutants closer to the landfill downstream although the BOD_5 concentrations are higher as compared to the present study. On the other hand some studies concur with the findings of this study (Areola, Segosebe and Gwisai, 2015; Bhambulkar, 2011).



Source: Author's Findings (2012)

Figure 8. Alkalinity of Landfill Surface Water

Table 1. Classification of Surface Water Quality Based on (BOD_5) values

DEGREE	(BOD_5) ($\text{O}_2 \text{ mg l}^{-1}$)
Very Clean	< 1.0
Clean	1.1 – 1.9
Moderately Polluted	2.0 – 2.9
Polluted	3.0 – 3.9
Very Polluted	4.0 – 10.0
Extremely Polluted	>10.0

Source: Radojevic and Bashkin (1999)

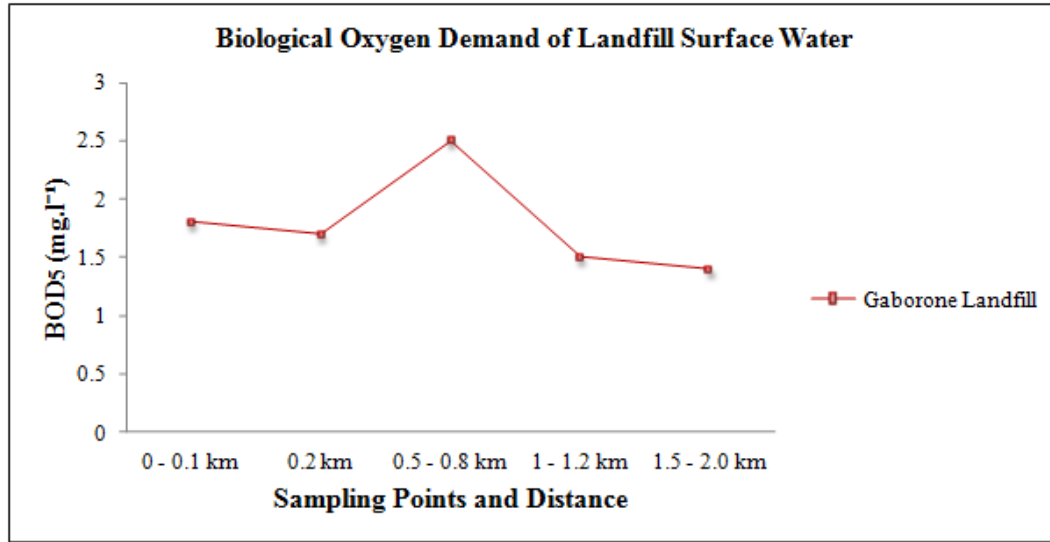


Figure 9. Biological Oxygen Demand of Landfill Surface Water

Phosphates (PO_4^{3-}), Nitrates (NO_3^-), Sulphates (SO_4^{2-}) and Ammonia (NH_3) Levels

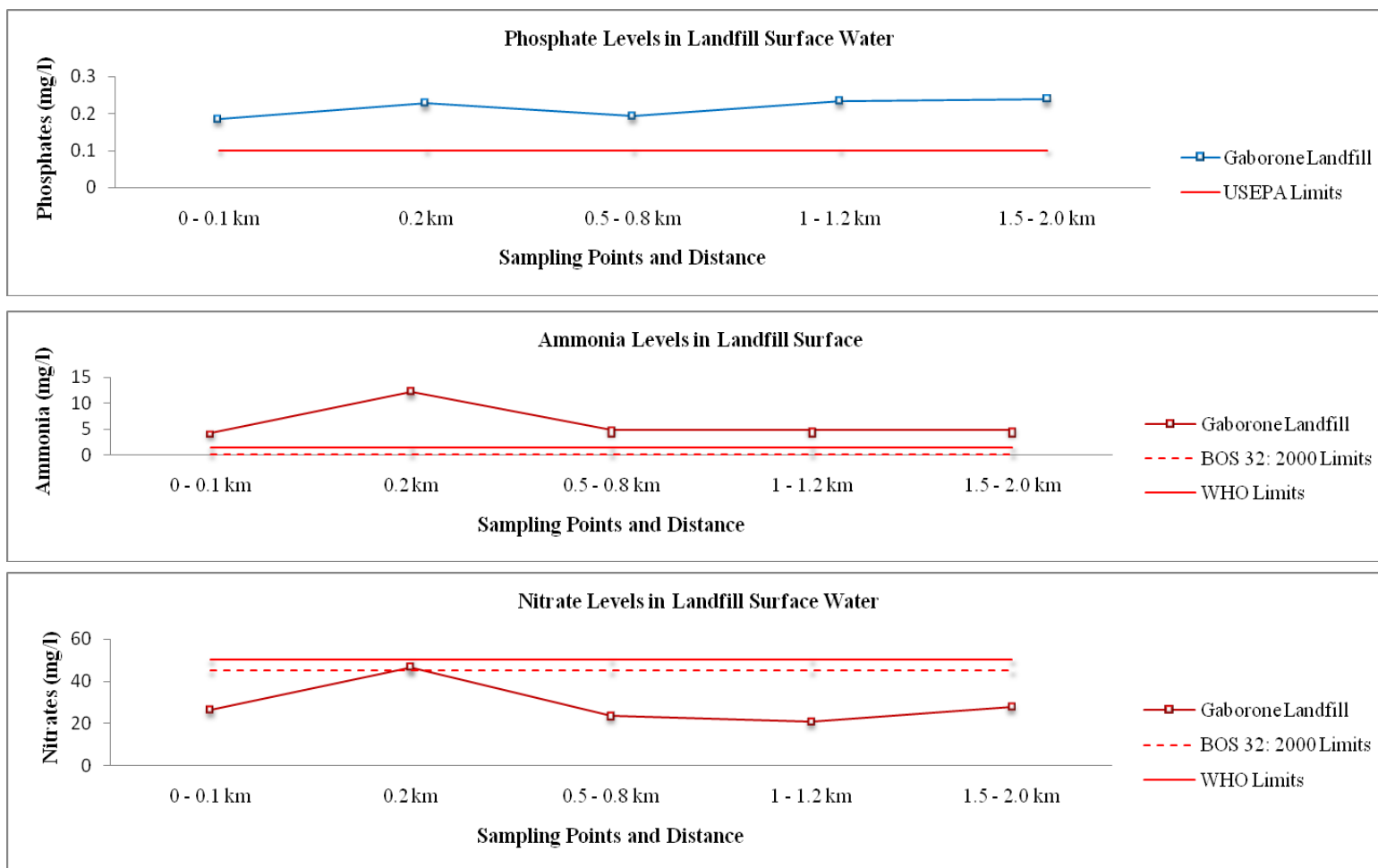
All the surface water samples have a mean phosphate (PO_4^{3-}) level of 0.22 $mg.l^{-1}$ above the USEPA drinking water set standard (0.1 $mg.l^{-1}$), (Table 2). The level is similar to that of other studies carried out earlier (Areola, Segosebe and Gwisai, 2015; Osei *et al.*, 2011; Raman and Sathiya – Narayanan, 2008; Longe and Enekwechi, 2007). The current levels could be due to waste materials from the landfill such as animal waste and fertilizers as revealed in previous studies (Longe and Balogun, 2010; Raman and Sathiya – Narayanan, 2008). There is a gradual increase in PO_4^{3-} levels with increasing distance (Figure 10). The Gaborone could be due to the amount of waste and other sources of waste deposition independent of the landfill that could have led to high PO_4^{3-} levels in Gaborone surface waters (Alslaibi, Mogheir and Afifi, 2011; Steinnes *et al.*, 1997).

Table 2. Mean values for physico – chemical parameters of surface water in Gaborone

Parameter	Gaborone Landfill Mean Values	WHO	USEPA	BOS 2000	EU Limits
pH	8.12	6.5	6.5	6.5	7.5
EC ($\mu S.cm^{-1}$)	1064.3	-	-	3100	44.6
Colour (TCU)	12.8	-	15	15	-
TDS (mg)	744	500	500	450	-
Turbidity (NTU)	5.62	6	5	0.5	-
Alkalinity ($mg.l^{-1}$)	600.20	-	-	150	-
BOD_5 ($mg.l^{-1}$)	1.78	-	-	-	-
PO_4^{3-} ($mg.l^{-1}$)	0.22	-	0.1	-	-
NO_3^- ($mg.l^{-1}$)	29.06	50	-	45	9.07
SO_4^{2-} ($mg.l^{-1}$)	447.3	250	250	200	52.1
NH_3 ($mg.l^{-1}$)	6.12	1.5	-	0.2	-
Cl^- ($mg.l^{-1}$)	2290.92	250	250	100	33.3

Nitrates (NO_3^-) in water samples of Gaborone landfills have a mean of 29.06 $mg.l^{-1}$ (Table 2). The range (20.93 – 46.51 $mg.l^{-1}$) is similar to that of other studies conducted earlier showing an appreciable presence of pollutants in all the water samples (Akinbile, 2012; Alslaibi, Mogheir and Afifi, 2011; Longe and Balogun, 2010); while other findings are far lower than for the current study and set thresholds for drinking water (Sholichin, 2012; Osei *et al.*, 2011; Al – Sabahi *et al.*, 2009; Raman and Sathiya – Narayanan, 2008; Longe and Enekwechi, 2007; Salminen, 2005). However the NO_3^- levels of the current study are below the set standards (NSDWQ, 2007; WHO, 2004; BOS 32: 2000). All samples except the 0.2 km [which is similar to previous studies (Gvozdic *et al.*, 2012)] are below the BOS 32:2000 drinking water standard (45 $mg.l^{-1}$), and the WHO drinking water standard (50 $mg.l^{-1}$) although above the EU limits (Table 2). There is a general decline in NO_3^- levels with increasing distance from the landfill.

Sulphate (SO_4^{2-}) levels are higher than the set standards for all the samples (Figure 10 and Table 2), (WHO, 2004; USEPA, 1991). The concentration range for all the samples is $353.7 - 651.6 \text{ mg.l}^{-1}$ which is statistically significantly higher than for other studies in other developing and developed countries indicating toxic pollution from the landfill waste (Osei *et al.*, 2011; Longe and Balogun, 2010; Raman and Sathiya – Narayanan, 2008; Longe and Enekwechi, 2007; Salminen, 2005). Indeed, some prior studies have shown that landfills may pollute surface waters with high SO_4^{2-} levels by as much as 784 mg.l^{-1} (Le Seur Spencer and Drake, 1987). The general trend shows that there is a decline in concentration of SO_4^{2-} downstream with an increase in distance from the landfill. However, this is contrary to the findings by Osei *et al.*, (2011), which indicate increased SO_4^{2-} levels with increasing distance.



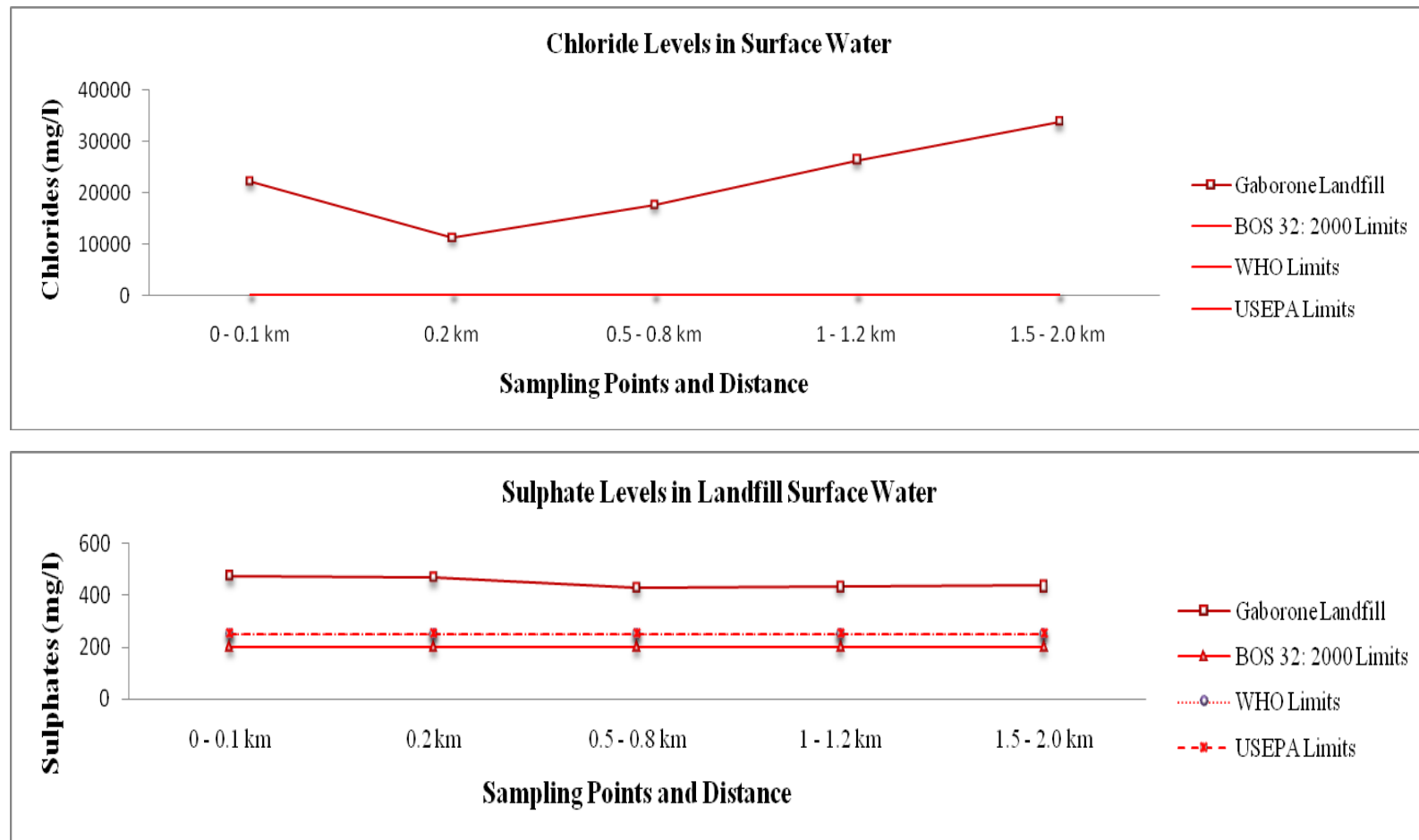


Figure 10. Gaborone Landfill Surface Water Levels PO_4^{3-} , NH_3 , NO_3^- , Cl^- and SO_4^{2-}

The concentration of Ammonia (NH_3) in Gaborone is 4.2 – 12.22 mg.l^{-1} (Figure 10). All samples from both landfills have NH_3 levels above the set Botswana and WHO drinking water limits, which are 0.2 mg.l^{-1} and 1.5 mg.l^{-1} respectively (Figure 10 and Table 2), (BOS 32: 2000; WHO, 2004). This is an indication of pollution from waste materials as the NH_3 levels are higher than in prior studies (Gvozdic *et al.*, 2011; Alslaibi, Mogheir and Afifi, 2011; Al – Sabahi *et al.*, 2009; Longe and Enekwechi, 2007). There is a general decline in NH_3 levels with increasing distance from the landfill (Longe and Balogun, 2010).

Chlorides ($^{36}_{17}\text{Cl}^-$) Concentration

Chlorides ($^{36}_{17}\text{Cl}^-$) concentrations for all the surface water samples range from 8224.4 – 37577 mg.l^{-1} (Figure 10). The $^{36}_{17}\text{Cl}^-$ levels of the current study are statistically significantly higher than those of prior measurements conducted in other developing countries (Akinbile, 2012; Alslaibi, Mogheir and Afifi, 2011; Longe and Balogun, 2010; Al – Sabahi *et al.*, 2009; Longe and Enekwechi, 2007; Salminen, 2005; Jeevan Rao and Shantaram, 2003). The high $^{36}_{17}\text{Cl}^-$ levels indicate pollution from landfill waste which requires treatment before use (Igbinsosa and Okoh, 2009). All samples are above the set Botswana, WHO and USEPA drinking water limits, which are 100, 250, and 250 mg.l^{-1} respectively (Figure 10 and Table 2), (Raman and Sathiya – Narayanan, 2008; WHO, 2004; BOS 32:2000; USEPA, 1991). There is a sharp increase in $^{36}_{17}\text{Cl}^-$ levels with increasing distance from the landfill. This could be due to livestock which utilize the surface water sources for drinking. However this is contrary to the findings of prior studies (Alslaibi, Mogheir and Afifi, 2011).

Hexavalent Chromium ($^{52}_{24}\text{Cr}^{6+}$) and Copper ($^{64}_{29}\text{Cu}^{2+}$) Concentration

All samples have $^{52}_{24}\text{Cr}^{6+}$ concentration in the range 0.004 – 0.02 mg.l^{-1} which is below the set limits (Figure 11). Though appreciable the $^{52}_{24}\text{Cr}^{6+}$ concentrations in the waters are below the set drinking standards of 0.05 mg.l^{-1} (Gvozdic *et al.*, 2011; Osei *et al.*, 2011; Raman and Sathiya – Narayanan, 2008). The results are similar to those of previous studies conducted in developing countries (Raman and Sathiya – Narayanan, 2008; Longe and Enekwechi, 2007). On the other hand some studies reveal that water samples have statistically significantly high $^{52}_{24}\text{Cr}^{6+}$ levels as compared to the current study (Longe and Balogun, 2010, Salminen, 2005). $^{52}_{24}\text{Cr}^{6+}$ concentration gradually increases with distance from the landfill. Similarly prior findings show the same pattern of $^{52}_{24}\text{Cr}^{6+}$ levels downstream (Al – Sabahi *et al.*, 2009).

$^{64}_{29}\text{Cu}^{2+}$ was detected in only one sample (0 – 0.1 km) from the Gaborone landfill at 0.054 mg.l^{-1} . The $^{64}_{29}\text{Cu}^{2+}$ level is lower or similar as compared to other research studies conducted earlier (Gvozdic *et al.*, 2011; Osei *et al.*, 2011; Tamunobereton – ari, Omubo – Pepple and Tamunobereton – ari, 2010; Al – Sabahi *et al.*, 2009; Longe and Enekwechi, 2007; Salminen, 2005; Haertling, 1989). The sample is the closest to the landfill indicating waste influence and pollution downstream. $^{64}_{29}\text{Cu}^{2+}$ is not detected in all the other samples suggesting that $^{64}_{29}\text{Cu}^{2+}$ could have been adsorbed by the soil strata or by the soil organic matter (Akinbile, 2012; Alslaibi, Mogheir and Afifi, 2011; Al – Sabahi *et al.*, 2009; Suman *et al.*, 2006).

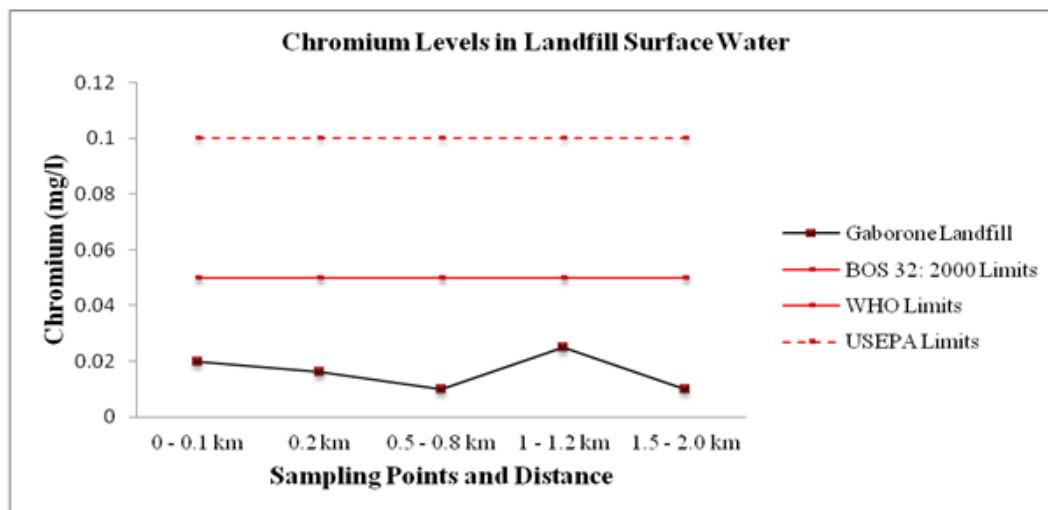


Figure 11. Chromium Levels in Landfill Surface Water

Sodium ($^{23}_{11}\text{Na}^+$) and Potassium ($^{39}_{19}\text{K}^+$) Concentration

The concentration range of $^{23}_{11}\text{Na}^+$ in the surface water samples for Gaborone landfill is 42.18 – 120.92 mg.l^{-1} . All the samples are below the WHO (200 mg.l^{-1}) drinking standards, while the majority of the samples are below the Botswana (100 mg.l^{-1}) drinking water threshold; on the other hand all samples are above the USEPA (10 mg.l^{-1}) and EU limits (23.1 mg.l^{-1}) of drinking water respectively (Figure 12 and Table 3). The $^{23}_{11}\text{Na}^+$ levels of the current study are statistically significantly higher than levels observed in other studies (Al – Sabahi *et al.*, 2009; Haertling, 1989). The impact of the landfill on the water samples appears to be limited based on the Botswana and WHO standards (BOS 32: 2000; WHO, 2004). The 0.5 – 0.8 km sample has the highest $^{23}_{11}\text{Na}^+$ level as compared to others, and a similar trend is observed among previous studies suggesting anthropogenic factors (Alslaibi, Mogheir and Affi, 2011; Steinnes *et al.*, 1997).

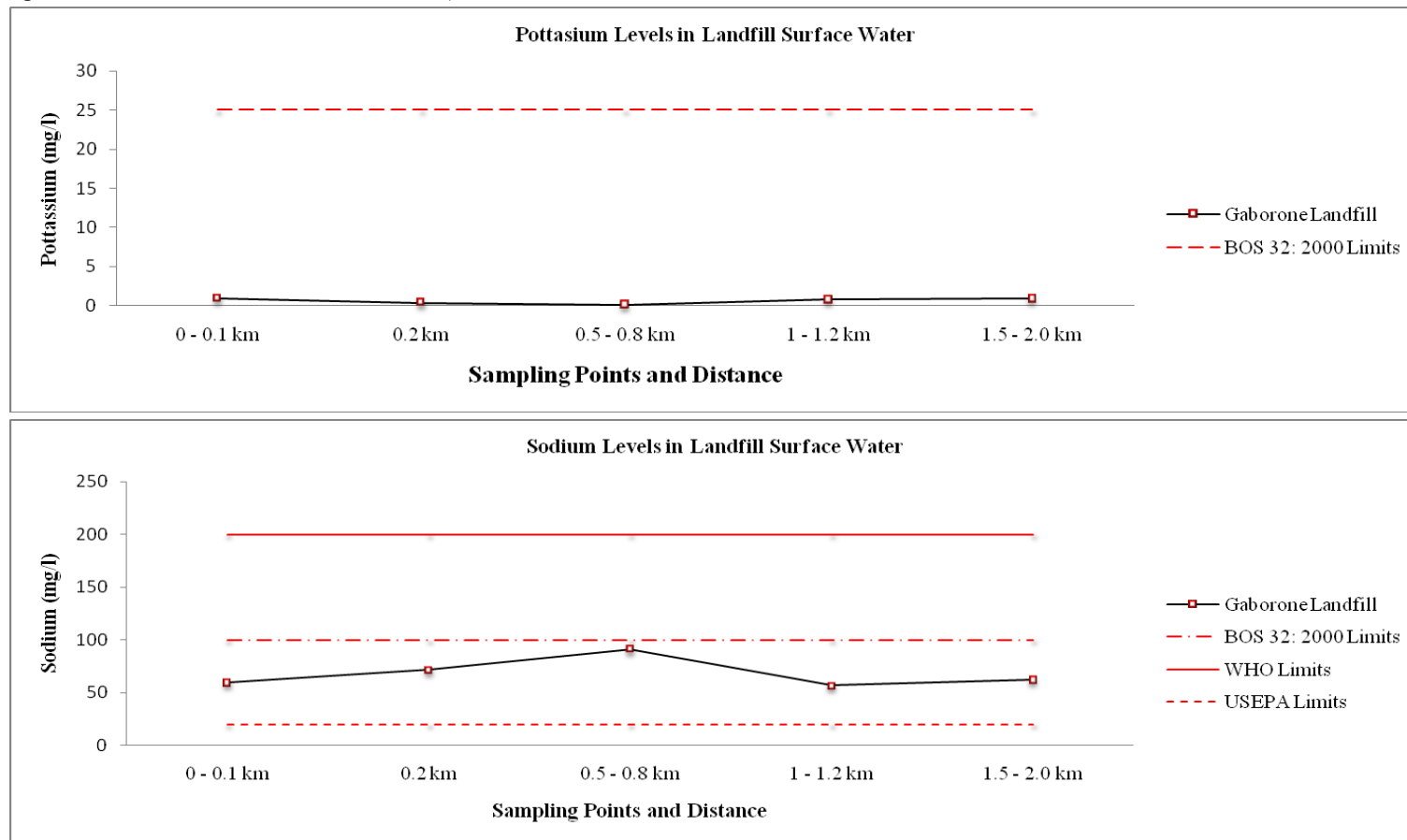


Figure 12. Sodium and Potassium Levels in Landfill Surface Water

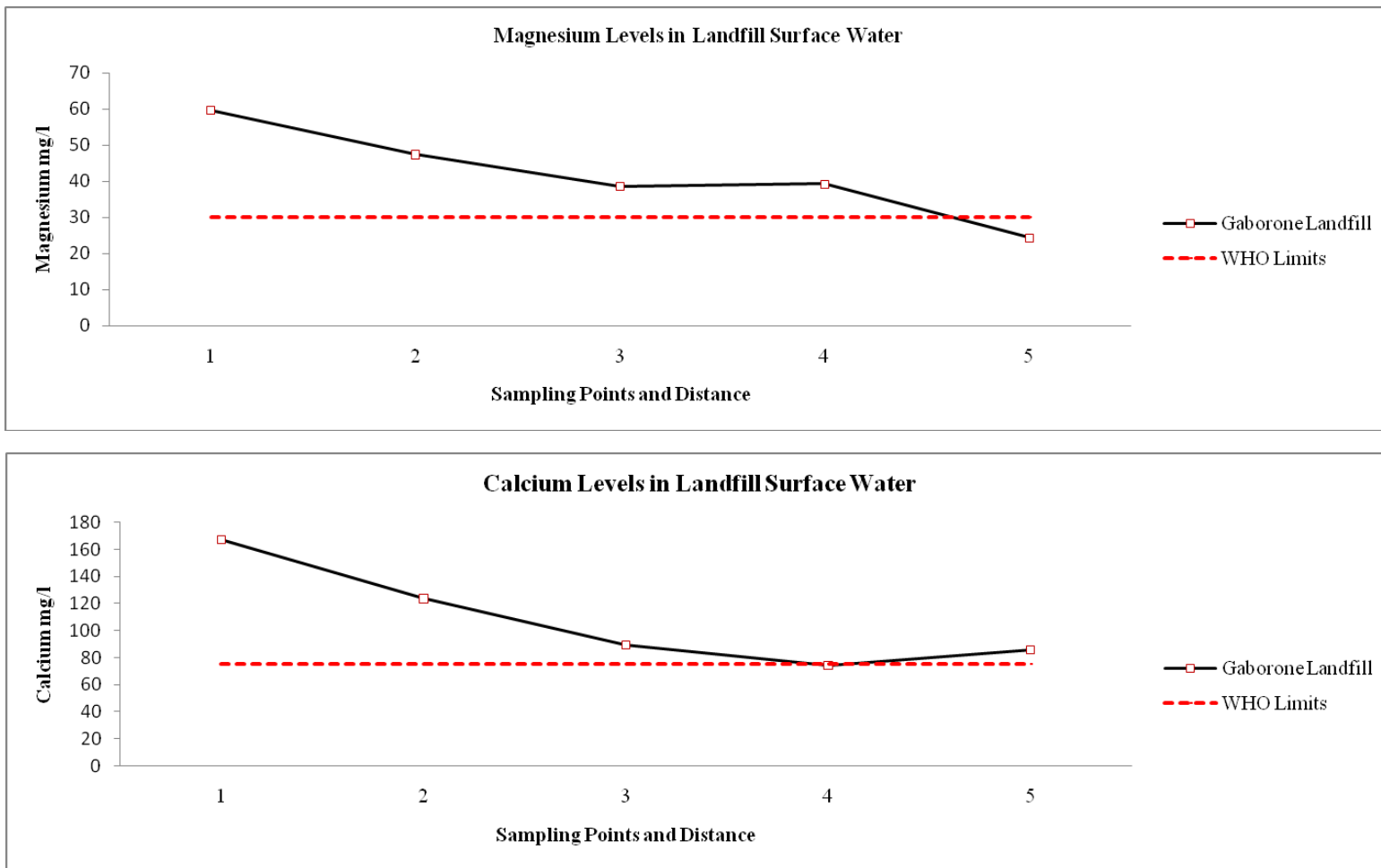


Figure 13. Calcium and Magnesium Levels in Landfill Surface Water

Table 3. Mean values for heavy metals of surface water in Gaborone.

Parameter	Gaborone Landfill Mean Values	WHO	USEPA	BOS 2000	EU Limits
$^{52}_{24}\text{Cr}^{6+}$ (mg.l ⁻¹)	0.02	0.05	0.1	0.05	0.792
$^{64}_{29}\text{Cu}^{2+}$ (mg.l ⁻¹)	0.05	1 – 2	1.3	-	1.23
$^{23}_{11}\text{Na}^{+}$ (mg.l ⁻¹)	68.17	200	20	100	23.1
$^{39}_{19}\text{K}^{+}$ (mg.l ⁻¹)	0.61	-	-	25	3.07
$^{40}_{20}\text{Ca}^{2+}$ (mg.l ⁻¹)	107.86	75	-	-	55.2
$^{24}_{12}\text{Mg}^{2+}$ (mg.l ⁻¹)	41.76	30	-	-	11.5

Source: Authors Findings (2012)

The range for $^{39}_{19}\text{K}^{+}$ concentrations in Gaborone is (0.145 – 0.8977 mg.l⁻¹), (Figure 12). However, all the samples are below the set drinking water limits in Botswana (25 mg.l⁻¹), EU countries (12 mg.l⁻¹) and Canada (10 mg.l⁻¹), (Salminen, 2005; Radojevic and Bashkin, 1999). Similarly other studies show that $^{39}_{19}\text{K}^{+}$ levels are lower than the set thresholds of international organizations and country regulations (Al – Sabahi *et al.*, 2009; YMWE, 1999). However, some studies have recorded very high $^{39}_{19}\text{K}^{+}$ levels beyond the set limits (Haertling, 1989). There is a general decline in $^{39}_{19}\text{K}^{+}$ levels with increasing distance from the landfill as observed in other literature (Omoniyi and Ogunsanwo, 2009).

Calcium ($^{40}_{20}\text{Ca}^{2+}$) and Magnesium ($^{24}_{12}\text{Mg}^{2+}$) Concentration

The range for $^{40}_{20}\text{Ca}^{2+}$ concentrations is 54.67 – 167.1 mg.l⁻¹ for both Gaborone samples (Figure 13). Most of the surface water samples are above the WHO and EU drinking water threshold as observed by prior studies (Longe and Enekwechi, 2007; Salminen, 2005; WHO, 2004; Le Seur Spencer and Drake, 1987). However, some studies have shown lower $^{40}_{20}\text{Ca}^{2+}$ levels than the current study (Haertling, 1989). There is a general decline in $^{40}_{20}\text{Ca}^{2+}$ levels with increasing distance from the landfill. The same spatial trend has been observed by studies in some developing nations (Akinbile, 2012; Al – Sabahi *et al.*, 2009). This indicates the influence of waste materials disposed of at the landfill site (Van Lynden, Mantel and Van Oostrum, 2004).

The range for $^{24}_{12}\text{Mg}^{2+}$ concentrations is (24.37 – 59.48 mg.l⁻¹) for Gaborone surface water samples (Figure 13). On the other hand some studies have recorded higher $^{24}_{12}\text{Mg}^{2+}$ levels than for the current study (Le Seur Spencer and Drake, 1987). There is a significant general decline in $^{24}_{12}\text{Mg}^{2+}$ levels with increasing distance from the landfill ($r = -0.9439$, $p = \leq 0.05$ with a $R^2 = 0.8909$), and the trend appears to be similar to that of previous

studies (Al – Sabahi *et al.*, 2009). Most of the samples at the landfill are above the set WHO and EU limits, (Salminen, 2005; WHO, 2004). On the other hand most studies have shown that $^{24}_{12}\text{Mg}^{2+}$ levels are higher downstream than upstream (Haertling, 1989; Le Seur Spencer and Drake, 1987; WHO, 1984).

6. Conclusions

All of the surface water samples were alkaline and there was a general decline in pH values with increasing distance from the landfill sites. On the other hand the pH was higher than the BOS 32: 2000, WHO and USEPA limits. Some water samples in Gaborone closer to the landfill surpassed the set colour limits.

Alkalinity levels were higher than BOS 32: 2000 limits. The BOD₅ test showed that the surface water samples were moderately polluted. All surface water samples had phosphate levels above the USEPA set limits, while some samples had surpassed the nitrates limits in Botswana. All samples from the landfill had sulphate, chloride and ammonia concentration levels which were above the set limits (USEPA, WHO and BOS 32:2000). The sodium concentrations were above the USEPA set limits. Calcium and Magnesium samples close to the landfill were above the WHO limits for Gaborone landfill.

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