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2016

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How to cite

KAPEMBO, Michel L. et al. Evaluation of water quality from suburban shallow wells under tropical conditions according to the seasonal variation, Bumbu, Kinshasa, Democratic Republic of Congo. In: Exposure and health, 2016, vol. 8, n° 4, p. 487–496. doi: 10.1007/s12403-016-0213-y

This publication URL: <https://archive-ouverte.unige.ch/unige:85288>

Publication DOI: [10.1007/s12403-016-0213-y](https://doi.org/10.1007/s12403-016-0213-y)

Evaluation of Water Quality from Suburban Shallow Wells Under Tropical Conditions According to the Seasonal Variation, Bumbu, Kinshasa, Democratic Republic of the Congo

Michel L. Kapembo¹ · Amandine Laffite² · Mathieu K. Bokolo¹ · Aimé L. Mbanga¹ · Marc M. Maya-Vangua¹ · Jean-Paul Otamonga³ · Crispin K. Mulaji¹ · Pius T. Mpiana¹ · Walter Wildi² · John Poté^{1,2,3}

Received: 12 February 2016 / Revised: 24 May 2016 / Accepted: 25 May 2016
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Abstract The contamination of freshwater resources with various organic and inorganic contaminants is still a major problem in many parts of the world, especially in developing countries in which the poor water quality continues to pose a serious threat to human health. In this study, the assessment of groundwater quality was performed in the municipality of Bumbu (Kinshasa, Democratic Republic of the Congo) according to the seasonal variation. Water physicochemical parameters [pH, electrical conductivity (EC), dissolved oxygen and soluble ions (Na^+ , K^+ , PO_4^{3-} , SO_4^{2-} , NO_3^- , NO_2^-)] and faecal indicator bacteria (FIB) including *Escherichia coli* (*E. coli*), *Enterococcus* (ENT) and Total coliforms (TC) were analysed. Except for EC and NO_3^- , the results revealed low concentration of other water physicochemical parameters, which are below the recommended limits, according to World Health Organization guideline for drinking water. Additionally, the result showed high concentration of FIB reaching the values of 1.6×10^4 , 1.5×10^4 and 9.0×10^5 CFU 100 mL^{-1} for *E. coli*, ENT and TC, respectively. The pollution was substantial in wet season compared to dry season. PCR amplification for human-*Bacteroides* indicated that more

than 90 % of bacteria were from human origin. Our results highlight the potential human risk associated with the exposure to water contamination from wells due to the high level of NO_3^- , EC, *E. coli* and ENT in both dry and wet seasons. The approach developed in this study helps provide a better understanding of the physicochemical and microbiological pollution of wells in large cities characterized with lack of wastewater and sanitation facilities.

Keywords Shallow wells · Water quality · Faecal indicator bacteria · Human-*Bacteroides* · Human health risk · Tropical conditions

Introduction

The contamination of freshwater resources with micropollutants and pathogenic organisms is still unsolved problem in many parts of the world, especially in developing countries, such as the Democratic Republic of Congo (DRC), in which there are lack of adequate sanitation facilities and where people practice open defecation (WHO 2004; Devarajan et al. 2015a, b; Mwanamoki et al. 2015). Data concerning the occurrence of biological contaminants such as faecal indicator bacteria (FIB) in aquatic environments in Sub-Saharan African countries is limited. Diarrhoeal diseases, mainly due to the consumption of microbiologically contaminated drinking water, cause about one billion illnesses and 2.2 million deaths per year (Montgomery and Elimelech 2007; WHO 2011). The majority of cases are in Sub-Saharan Africa and south Asia. In some of these regions, according to the economic situation and lack of effective infrastructure, a large proportion of the population uses highly contaminated surface water, shallow wells, boreholes, springs and streams for irrigation,

✉ John Poté
john.pote@unige.ch

¹ Department of Chemistry, Faculty of Science, University of Kinshasa, B.P. 190, Kinshasa XI, Democratic Republic of the Congo

² Faculty of Sciences, Earth and Environmental Sciences, Institute F. A. Forel and Institute of Environmental Sciences, University of Geneva, Bd Carl-Vogt 66, 1211, Geneva 4, Switzerland

³ Université Pédagogique Nationale (UPN), Croisement Route de Matadi et Avenue de la Libération. Quartier Binza/UPN, B.P. 8815, Kinshasa, Democratic Republic of the Congo

domestic and drinking purposes (Amanial 2015; Mwanamoki et al. 2015; Rochelle-Newall et al. 2015). Although most infections occur in developing countries, waterborne diseases are a worldwide problem that also concerns industrialized societies where highly populated centres draw their supply of drinking water (Haller et al. 2009; WHO 2011; Thevenon et al. 2012). In Democratic Republic of the Congo (DRC, with estimated population of 65.7 million inhabitants), despite the potential of its rich fresh water network, more than 75 % of the population have no access to safe water (UNEP 2011). The most common sources of domestic and drinking water for suburban and rural people are streams, shallow wells and springs. These water resources are in many cases polluted by micropollutants and pathogenic organisms which can represent significant potential impacts for human health.

At Kinshasa (capital of the DRC with more than 10 million inhabitants), waterborne diseases, including *Entamoeba* and *Shigella* and other diarrhoeal diseases, are great public health concerns. The prevalence of diarrhoeal in the province for the 15 last years affected more than 20.9 % of the population (EIES 2012). The city itself has experienced cholera outbreaks in years 1996, 2010 and 2011 and typhoid fever has affected nearly 1 % of the population (EIES 2012). To our knowledge, there have been no regular water quality monitoring and/or research programmes conducted to assess the microbial and physicochemical qualities of water resources. Our previous researches have been conducted to assess the quality of urban rivers surrounding the City of Kinshasa. The results of these research works indicated the deterioration of the quality of these water resources, especially rivers receiving urban and hospital effluent waters. The pollution can be explained with several aspects including domestic wastes, human open defecation and runoff of wastewater from septic tanks, and hospital wastewaters discharge (Mubedi et al. 2013; Mwanamoki et al. 2014). Our previous data showed also that sediments of urban river-reservoirs (such as Lake Ma Vallée and N'djili River) constitute important reservoirs of FIB (Tshibanda et al. 2014; Mwanamoki et al. 2015). To date, no programmes or researches have been conducted to assess the water quality from shallow wells according to the seasonal variation. Consequently, there is no quantitative information regarding the quantification of FIB in streams, rivers, shallow wells and other water sources in suburban municipalities of Kinshasa, other great cities as well as rural districts of DRC.

The main objective of the present study is to assess the seasonal variation of the water quality from shallow wells of the municipality of Bumbu, a suburban municipality located in south of the Capital City of Kinshasa. The assessment is based on (1) quantification of water physicochemical parameters including pH, electrical conductivity

(EC), dissolved oxygen (O_2) and soluble ions (Na^+ , K^+ , PO_4^{3-} , SO_4^{2-} , NO_3^- , NO_2^-), and (2) quantification of FIB including *E. coli*, ENT and TC and (3) to characterize the isolated FIB by molecular approach in order to identify the possible cause of water contamination. The analysis of water samples was performed during both the dry and wet seasons in order to identify the eventual changes in water quality with season. Molecular analysis was performed not only to confirm the FIB but also to identify the eventual human source of faecal pollution in wells. The approach used for this research is very important as it allows determining whether shallow wells can provide safe drinking water according to the seasonal variation or whether there is a need to develop ways to improve the water quality from shallow wells (Pritchard et al. 2008).

Materials and Methods

Study Site Description

The study was conducted in the commune of Bumbu, a suburban municipality located in the Funa district of Kinshasa (the capital city of the DRC), with total area of 5.3 km² and an estimated population of about 329,324 habitants. Seven shallow wells from four neighbourhoods (named Ubangi; Mfimi; Dipiya and Mongala) of the municipality were selected (Fig. 1; Table 1). The study area was chosen based on some criteria including, waterborne disease epidemiology, economic situation of local people as well as their access to safe water provided by national society (Regideso), the presence of several shallow wells which constitute the main water resource of more than 70 % of people for domestic use (drinking, cooking and washing) and sanitary risks such as the presence of uncontrolled landfills, pit latrines and septic tanks and how the wells are protected.

Sampling Procedure

Water samples from seven shallow wells were collected in June and July (dry season)/2015, and in October and November (wet season)/2015. The shallow wells (sampling sites) are labelled P1, P2, P3, P4, P5, P6 and P7 (Fig. 2). The frequency of water sampling from each shallow well was two times per month/season, that is, at the beginning (first week) and during the last week of each month. Water from wells was taken by a craft device made of one litre (L) clean polyethylene bottle attached to a rope. For physicochemical analysis, water samples (500 mL) were filled into clean plastic bottles. For bacteriological analysis, water samples (500 mL) were filled into sterile (autoclaved) plastic bottles.

Fig. 1 Sampling site, **a** Africa continental map, **b** Democratic Republic of Congo map, **c** maps location sampling shallow well water, Bumbu township, capital city of Kinshasa, **d** sampling sites location well water neighbourhood P1 (Ubangi); P2–P3 (Mfimi); P4–P5 (Dipiya) and P6–P7 (Mongala)



Table 1 Wells GPS location, depth and number of well users

Sampling site	Longitude	Latitude	Depth to water level		Colour	Number of users
			Dry season (m)	Wet season (m)		
P1	15°13'6.7"	4°22'4.0"	3	3	Clear	±800
P2	15°17'44.3"	4°22'49.7"	2	1	Soft	±700
P3	15°17'45.7"	4°22'50.2"	3	2	Clear	±900
P4	15°17'40.6"	4°22'43.1"	3	2	Clear	±250
P5	15°17'42.0"	4°22'41.1"	2.5	2	Clear	±200
P6	15°18'0.1"	4°22'29.0"	2	1	Soft	±150
P7	15°18'6.7"	4°22'1.4"	2.5	1.5	Clear	±450

After sampling, the samples were stored in an icebox at 4 °C and immediately transported to the laboratory for analysis within 24 h.

Water Physicochemical Parameter Analysis

Physicochemical parameters of water including temperature (T), pH, dissolved oxygen (O_2) and electrical conductivity (EC) were measured in situ using a Multi

parameter 350i (WTW, Germany). The concentration of dissolved major ions (Na^+ , K^+ , PO_4^{3-} , SO_4^{2-} , NO_3^- , NO_2^-) was measured using the spectrophotometer HACH LANGER (DR-3800, Germany) according to the manufacturer's recommendations. The reference material (certified water CRM, Ontario-99) from the National Water Research Institute, Canada was used to calibrate and verify the accuracy of the instrument. All CRM results were within the acceptance range on the CRM certificate.

Fig. 2 Selected shallow wells (P1–P7) of the municipality of Bumbu (photos taken by M. Kapembo in June 2015)



Faecal Indicator Bacteria (FIB) Analysis in Water

The FIB (including *E. coli*, ENT and TC) were quantified in water samples according to the international standard methods for water quality determination using the membrane filtration method (APHA 2005). Briefly, for each sample, triplicates of 100 mL of water was passed through a 0.45- μm filter (Sartorius stedim, biotech, Germany), and placed on different selective culture media (Biolife, Italian), using the following incubation conditions: *E. coli*: Tryptone Soy Agar (TSA) medium, incubated at 37 °C for 4 h and transferred to Tryptone Bile X-Gluc Agar (TBX) medium at 44 °C for 24 h; ENT: Slanetz Bartley Agar (SBA) medium, incubated at 44 °C for 48 h and transferred into Bile Aesculin Agar (BAA) medium at 44 °C for 4 h; and TC: Endo agar medium, incubated at 35 °C for 24 h. The results are expressed as colony forming units per 100 mL of water (CFU 100 mL⁻¹). The reproducibility of the whole experimental procedure was tested by means of triplicates.

Characterization of FIB Strains

The PCR amplification was performed directly on the colonies picked from selective-media plates (resuspended in 20 μL of sterile water) using human-specific *Bacteroides* primers *HF183F*- (5'-ATCATGAGTTCACATGTCCG-3') and *Bac708R*- (5'-CAATCGGAGTTCCTTCGTG-3') (Bernhard and Field 2000; Ahmed et al. 2007), with the PCR conditions as described by Thevenon et al. (2012). PCR reactions of 25 μL total volume consisted of 0.025 U/ μL Takara Ex Taq HS DNA polymerase (Takara Bio Europe, Saint-Germain-en-Laye, France), 1 \times PCR buffer (Takara, containing 2 mM of Mg²⁺), 800 μM of dNTP, 0.2 μM of each forward and reverse primers (Invitrogen), 50 ng/ μL of BSA. PCR was performed in a Biometra thermocycler (BIOLABO) using the following conditions: 15 min for

95 °C initial denaturation; 35 cycles of denaturation (94 °C for 30 s), of annealing (60 °C for 1 min) and of extension (72 °C for 1 min), followed by a final extension at 72 °C for 10 min. PCR products were visualized after electrophoresis on 0.8 % agarose gels containing 1 \times SYBR Safe DNA gel stain (Invitrogen) in 1 \times TAE buffer. The experiment was conducted in triplicate in each set of conditions. The negative (without DNA) and positive controls (e.g. the expected 520 bp length) (for HF183/Bac708) from sewage (Poté et al. 2009) were used for each PCR essays.

Data Analysis

All analyses were conducted in triplicate for each set of conditions. Statistical processing of data was performed using SigmaStat 11.0 (Systat Software, Inc., USA). The data were subjected to a Spearman Rank Correlation test to investigate possible relationship using R statistical software, version 3.2.2 (R Core Team 2015). Data were subsequently subjected to a principle component analysis (PCA) using a correlation matrix and the ade4 package, to investigate potential effect of season and sampling site.

Results and Discussion

Physicochemical Quality of Water

The water physicochemical parameters including *r*, pH, EC, O₂ are presented in Table 2. Due to the absence of DRC's regulation, water quality was compared with the recommendations of the World Health Organization (WHO 2011). The monthly average temperature values varied from 25.0 to 26.8 °C in dry season and from 26.5 to 30.7 °C during the wet season. The highest value was observed in November and the lowest during June/July. These results are comparable with other published data obtained under tropical

Table 2 Averages of physicochemical parameters of water samples from selected wells during the dry season (dry) and wet season (wet)

Wells	Temp (°C)				pH				EC ($\mu\text{S cm}^{-1}$)				O ₂ (mg L ⁻¹)			
	Dry		Wet		Dry		Wet		Dry		Wet		Dry		Wet	
	June	July	Oct.	Nov.	June	July	Oct.	Nov.	June	July	Oct.	Nov.	June	July	Oct.	Nov.
P1	26.0	26.4	27.4	29.3	5.7	5.4	6.1	6.6	649	618	605	678	1.3	0.9	1.3	1.0
P2	26.2	26.5	26.5	28.7	7.2	6.8	7.2	7.2	976	1051	915	1060	1.8	1.8	2.0	2.9
P3	25.8	26.8	28.8	27.5	5.8	5.4	6.2	6.1	913	719	806	765	2.9	3.5	2.5	3.5
P4	26.8	26.4	30.5	28.1	5.1	4.9	6.8	6.1	1080	1199	1087	1058	1.2	1.3	2.1	2.5
P5	26.2	26.1	29.6	28.2	6.7	6.2	6.7	6.9	989	631	991	602	1.3	1.0	1.4	1.5
P6	25.0	25.3	28.1	30.7	7.4	6.8	7.2	7.1	1151	865	805	1088	1.2	1.6	2.5	2.3
P7	25.2	25.0	28.4	29.2	6.6	7.0	6.9	7.0	1190	1547	1035	1121	2.0	1.7	2.0	1.8

conditions (Pritchard et al. 2008; Nola et al. 2013). The pH values varied significantly according to the seasonal variation ($p < 0.05$) for some wells such as P1 (5.4–6.6), P3 (5.4–6.2) and P4 (4.9–6.8). The water from these wells can be considered as slightly acid water especially during dry season. There was no significant difference in pH levels according to the seasonal variation ($p > 0.05$) for other wells: P2 (6.2–7.2), P5 (6.2–6.9), P6 (6.8–7.4) and P7 (6.9–7.0). Except for P1, P3 and P4 during dry season, the values of pH from investigated wells comply with WHO recommendation for drinking water. All wells show high values of conductivity, ranging from 618 to 1547 and from 605 to 1121 $\mu\text{S cm}^{-1}$ during the dry and wet season, respectively. According to our discussion with people in charge of P7, the commercial NaCl (put directly into the well) was used to eventually kill present pathogen. The concentration of the dissolved oxygen for each well was not significantly varied with the season variation, ranging the values from 0.9 to 3.5 mg L⁻¹.

The concentrations of soluble ions (Na⁺, K⁺, PO₄³⁻, SO₄²⁻, NO₃⁻, NO₂⁻) in water samples are reported in Table 3. The maximum average values were 94.5, 54.3, 1.9, 33.9 and 0.32 mg L⁻¹, for Na⁺, K⁺, PO₄³⁻, SO₄²⁻ and NO₂⁻, respectively. High values were obtained during the

dry period. The lowest level of these dissolved ions can be explained by the dilution effect during the wet season. These results indicate that these soluble ions are within the WHO guideline values for domestically users in both dry and wet seasons. In contrary, NO₃⁻ presents the highest values (which did not meet the WHO guideline of 50 mg L⁻¹) ranging between 393.5–775.6 mg L⁻¹ (dry season) and 189.2–302.0 mg L⁻¹ (wet season). As observed for other soluble ions, the values of NO₃⁻ were higher during dry season than wet season. This seasonal variation tendency of nitrate concentration in wells is in agreement with other studies performed under tropical conditions (Boutin 1987; Bricha et al. 2007; Mkandawire 2008; Pritchard et al. 2008).

It is very difficult to explain well the fluctuation of these parameters according to the wells and season. Some recent studies performed in the similar environment demonstrated that the various attributes of watersheds, such as morphological and geological factors, as well as the local socio-economic conditions, lack of adequate sanitation and uncontrolled landfills can influence the status of surface and ground water bodies through physical, chemical, biological or bacteriological parameters (e.g. Tshibanda et al. 2014; Devarajan et al. 2015a; Tallar and Suen 2015). The presence of high concentration of soluble ions such NO₃⁻

Table 3 Average concentration of soluble ions of water samples from selected wells during the dry season (dry) and wet season (wet)

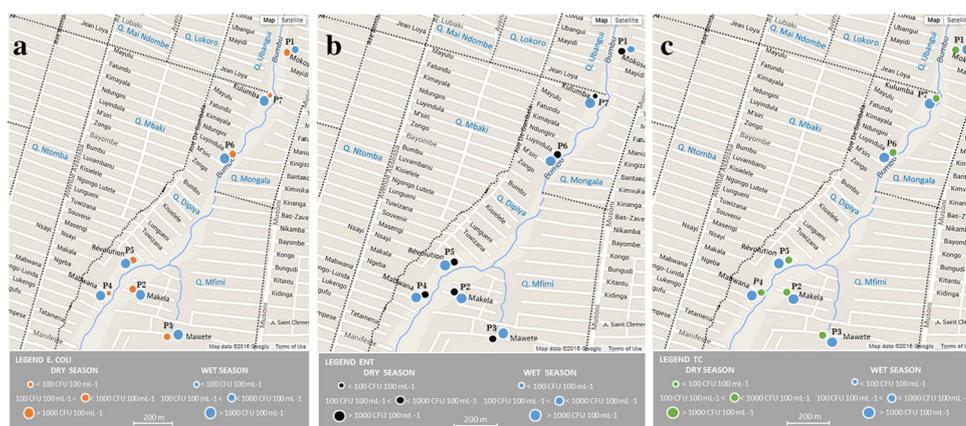
Wells	Na ⁺ (mg L ⁻¹)		K ⁺ (mg L ⁻¹)		PO ₄ ³⁻ (mg L ⁻¹)		SO ₄ ²⁻ (mg L ⁻¹)		NO ₃ ⁻ (mg L ⁻¹)		NO ₂ ⁻ (mg L ⁻¹)	
	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet	Dry	Wet
P1	83.69	31.15	51.11	32.75	0.32	0.13	24.75	12.13	546.02	200.06	0.08	0.06
P2	82.55	26.22	54.29	42.10	1.02	0.09	33.94	11.02	525.57	218.48	0.32	0.13
P3	61.83	19.76	38.53	18.69	1.85	0.10	13.52	5.89	693.62	272.12	0.07	0.05
P4	94.46	48.10	39.86	15.00	1.75	0.55	7.26	2.15	775.55	302.02	0.08	0.06
P5	43.39	18.42	29.78	10.13	0.06	0.01	25.59	17.20	500.55	198.08	0.08	0.09
P6	16.60	8.97	5.76	1.62	0.42	0.07	3.23	0.08	501.04	250.13	0.04	0.06
P7	9.50	0.69	2.37	1.01	0.04	0.01	3.66	1.01	393.50	189.20	0.03	0.01

in study area can probably be explained by water infiltration from crops (urban agriculture) which use fertilizers near wells (Ndembo Longo 2009; Ngelinkoto et al. 2014), mineralization of organic matter of septic tanks and uncontrolled landfills. It has been demonstrated that the concentration of NO_3^- and K^+ can be considered as the indicators of pollution from a range of organic sources, including latrines, landfills, leaking sewers, excrement from livestock or fertilizers or manures used on agricultural land (Banks et al. 2002).

Microbiological Quality of Water

Microbiological analysis of water showed that 100 % of analysed wells are substantially polluted with faecal matters in both dry and wet seasons (Fig. 3) and did not meet the WHO guideline for drinking/domestic using water. During the dry season, the FIB values ranged from $(0.9\text{--}3.5) \times 10^2$, $(1.1\text{--}5.8) \times 10^2$ and $(1.9\text{--}4.2) \times 10^3$ CFU 100 mL^{-1} for *E. coli*, ENT and TC, respectively. FIB concentrations for the wet season increased about 2 to 3 orders of magnitude than those observed for the dry season, with the values ranged from 8.6×10^2 to 1.6×10^4 , 0.8×10^2 to 1.5×10^4 and 2.5×10^2 to 9.0×10^5 100 mL^{-1} for *E. coli*, ENT and TC, respectively. These results indicate that water samples from tested shallow wells are heavily polluted in FIB. The contamination of wells in FIB can be attributed by several sources and mechanisms, including urban runoff, percolation from surface water, the absence of toilet facilities, septic systems' leaky sewer lines, direct injection of wastewater effluent and direct contamination by users (John and Rose 2005; Kelly et al. 2009; Nwachukwu et al. 2010; Nola et al. 2013). In our studied site, the deterioration of the quality of water from wells can probably explained by several aspects including the lack of adequate sanitation, protection of wells (there are many open wells), the presence of pit latrines located in the proximity of wells, open defecation and uncontrolled landfills (Fig. 2).

Fig. 3 Average **a** *Escherichia coli*, **b** *Enterococcus*, **c** total coliform quantification in shallow wells samples during the dry season (dry) and wet season (wet)



The surface recreation water as well as groundwater generally contains indigenous microorganisms, pathogenic and non-pathogenic microbes. The choice of bacterial indicators is thus very important for the management of aquatic environmental quality (Haller et al. 2009). The U.S. Environmental Protection Agency and the European Union recommend the use of *Escherichia coli* (*E. coli*), a subset of the faecal coliform group, and members of the genus *Enterococcus*, the *Enterococci* (ENT), to assess the hygienic safety of recreational waters (US EPA 2000; EU 2006). In this study, *E. coli*, ENT and TC were chosen to monitor the microbiological quality for the studied site under tropical conditions (Mwanamoki et al. 2014; Rochelle-Newall et al. 2015). Exposure to waters with high concentrations of *E. coli* and ENT is documented in epidemiological studies as being associated with an increased risk of contracting gastrointestinal and respiratory illnesses. The occurrence of these indicators in the studied site can therefore indicate water contamination in faecal material, and consequently, the possible presence of pathogenic organisms responsible for water-related diseases such as gastrointestinal, typhoid, cholera and other diarrhoeal diseases (US EPA 1984; Kay et al. 1994; Haile et al. 1999; An et al. 2002; Noble et al. 2004).

Characterization of FIB

Qualitative PCR assays were applied for large-scale screening of human-specific *Bacteroides*, in order to estimate as potential alternative indicators of human faecal material. Human material in water is generally considered to be a greater risk to human health risk (Scott et al. 2005; Converse et al. 2009). A total of 106 isolated colonies of *E. coli* from each wells were selected for PCR amplification to detect the human-specific *Bacteroides*. The PCR amplification performed for faecal human pollution showed high specific-human strains with the values ranging from 70 to 95 % (average for all test wells). Surprisingly, during both dry and wet seasons, PCR amplification

Table 4 Average *Escherichia coli*, *Enterococcus* and Total coliform quantification in shallow wells samples during the dry season (dry) and wet season (wet)

Samples	<i>E. coli</i> (CFU 100 mL ⁻¹)				ENT (CFU 100 mL ⁻¹)				T.C (CFU 100 mL ⁻¹)			
	Dry		Wet		Dry		Wet		Dry		Wet	
	June	July	Oct.	Nov	June	July	Oct.	Nov	June	July	Oct.	Nov
P1	1.6 × 10 ²	1.1 × 10 ²	1.9 × 10 ²	6.3 × 10 ²	2.9 × 10 ²	1.8 × 10 ²	8.2 × 10 ¹	7.9 × 10 ²	6.4 × 10 ²	3.5 × 10 ²	2.5 × 10 ²	1.9 × 10 ³
P2	3.5 × 10 ²	2.7 × 10 ²	4.4 × 10 ³	5.9 × 10 ³	5.8 × 10 ²	4.3 × 10 ²	2.8 × 10 ³	1.2 × 10 ³	1.3 × 10 ³	6.9 × 10 ²	1.3 × 10 ⁴	1.8 × 10 ⁴
P3	1.4 × 10 ²	1.5 × 10 ²	2.8 × 10 ³	1.1 × 10 ³	3.1 × 10 ²	3.5 × 10 ²	3.1 × 10 ²	2.4 × 10 ³	4.2 × 10 ²	4.6 × 10 ²	9.0 × 10 ⁴	3.4 × 10 ³
P4	5.9 × 10 ¹	7.8 × 10 ¹	4.7 × 10 ²	1.9 × 10 ³	1.3 × 10 ²	1.1 × 10 ²	1.7 × 10 ³	1.5 × 10 ³	1.9 × 10 ²	3.2 × 10 ²	1.4 × 10 ³	5.7 × 10 ³
P5	2.3 × 10 ²	2.1 × 10 ²	3.1 × 10 ³	8.6 × 10 ²	2.7 × 10 ²	2.9 × 10 ²	1.2 × 10 ³	9.0 × 10 ²	7.5 × 10 ²	7.1 × 10 ²	8.5 × 10 ³	2.7 × 10 ³
P6	1.6 × 10 ²	1.0 × 10 ²	1.6 × 10 ⁴	1.4 × 10 ⁴	3.1 × 10 ²	1.4 × 10 ²	1.1 × 10 ⁴	1.5 × 10 ⁴	4.2 × 10 ²	3.1 × 10 ²	4.7 × 10 ⁴	4.5 × 10 ⁴
P7	6.8 × 10 ¹	9.0 × 10 ¹	1.5 × 10 ⁴	2.1 × 10 ⁴	1.1 × 10 ²	7.9 × 10 ¹	1.4 × 10 ⁴	1.0 × 10 ⁴	3.3 × 10 ²	2.7 × 10 ²	4.4 × 10 ⁴	3.2 × 10 ⁴

E. coli *Escherichia coli*, ENT *Enterococcus*, T.C total coliform

demonstrated that 100 % of bacteria were from human origin for P2, P5, P6 and P7. These wells are characterized by highly values of physicochemical parameters during both dry and wet seasons (Table 4).

Statistical Correlation

FIB and physicochemical parameters were, for the most part, poorly correlated in sampled wells whether during dry season than during the wet season. In June, few correlations were observed between (Table 5); *E. coli* and TC ($r = 0.97, p = 0.001$); K⁺, Na⁺ and NO₂⁻ ($0.86 < r < 0.89, p < 0.01$); Na⁺ and NO₃⁻ ($0.76 < r < 0.86, p < 0.01$); NO₂⁻ and SO₄²⁻ ($0.82 < r < 0.85, p < 0.02$); NO₃⁻ and PO₄³⁻ ($r = 0.86, p < 0.01$); and SO₄⁻ and TC ($r = 0.81, p < 0.05$). Likewise, in October (Table 6) some sporadic

correlations were observed between *E. coli* and ENT ($0.85 < r < 0.93, p < 0.05$), Na⁺ and PO₄³⁻ ($r = 0.90, p < 0.01$); and O₂ and TC ($r = 0.72, p < 0.05$). These results indicate that, in general, FIB as well as soluble ion contamination wells could be considered to originate from different sources as explained above.

Inspection of ACP (Fig. 4) clearly shows the impact of seasonal variation on the wells contamination in FIB and dissolved ions, mainly NO₃⁻. Data show that the months of June and October are oppositely and separately clustered on separate axes of the first-principle component. During the rainy season, urban runoff, which is charged in contaminant, is continuously drained to wells leading to a large increase of bacterial contamination. On the contrary, during the wet season, there is no urban runoff to the well. FIB (mainly *E. coli*), which are indicator of recent

Table 5 Spearman’s Rank-Order Correlation of selected parameters^a analysed in water from shallow well in June 2015

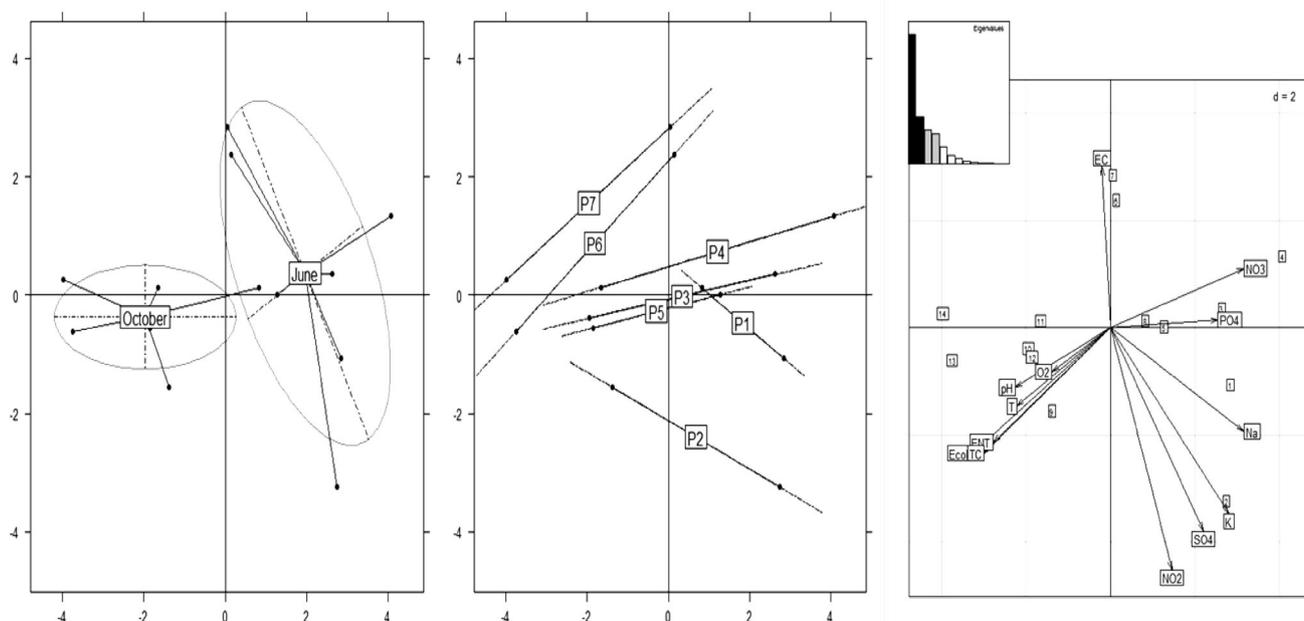
	EC	<i>E. coli</i>	ENT	K ⁺	Na ⁺	NO ₂ ⁻	NO ₃ ⁻	O ₂	pH	PO ₄ ³⁻	SO ₄ ²⁻	T	TC
EC		-0.38	-0.50	-0.75	-0.61	-0.59	-0.54	-0.25	0.32	-0.39	-0.68	-0.31	-0.54
<i>E. coli</i>			0.66	0.36	-0.07	0.54	-0.32	0.04	0.67	-0.16	0.68	0.07	0.97
ENT				0.50	0.14	0.39	0.23	0.12	0.50	0.52	0.38	-0.11	0.64
K ⁺					0.86	0.89	0.64	-0.04	-0.29	0.50	0.71	0.65	0.47
Na ⁺						0.74	0.86	-0.29	-0.64	0.57	0.43	0.76	0.04
NO ₂ ⁻							0.41	-0.17	-0.11	0.30	0.85	0.82	0.62
NO ₃ ⁻								-0.16	-0.64	0.86	0.11	0.47	-0.23
O ₂									-0.02	0.02	0.27	-0.23	0.17
pH										-0.25	0.00	-0.45	0.50
PO ₄ ³⁻											0.07	0.27	-0.13
SO ₄ ²⁻												0.63	0.81
T													0.16
TC													

^a Parameters include Physicochemical parameters (pH, temperature (T), electrical conductivity (EC), dissolved oxygen (O₂) and soluble ions (Na⁺, K⁺, PO₄³⁻, SO₄²⁻, NO₃⁻, NO₂⁻) and faecal indicator bacteria (FIB): *Escherichia coli* (*E. coli*), *Enterococcus* (ENT) and Total coliforms (TC). Significant coefficients ($p < 0.05$) are in bold

Table 6 Spearman's Rank-Order Correlation of selected parameters^a analysed in water from shallow well in October 2015

	EC	<i>E. coli</i>	ENT	K ⁺	Na ⁺	NO ₂ ⁻	NO ₃ ⁻	O ₂	pH	PO ₄ ³⁻	SO ₄ ²⁻	<i>T</i>	TC
EC		0.39	0.43	-0.36	0.00	-0.07	0.00	0.05	0.25	-0.13	-0.14	0.64	-0.11
<i>E. coli</i>			0.93	-0.68	-0.50	-0.15	0.04	0.49	0.85	-0.38	-0.82	0.07	0.36
ENT				-0.61	-0.61	-0.15	-0.21	0.33	0.88	-0.54	-0.71	-0.11	0.39
K ⁺					0.75	0.52	0.32	-0.24	-0.27	0.61	0.57	-0.43	-0.32
Na ⁺						0.37	0.57	-0.18	-0.38	0.90	0.39	0.07	-0.64
NO ₂ ⁻							0.04	-0.34	0.24	0.02	0.56	-0.26	-0.41
NO ₃ ⁻								0.69	0.02	0.74	-0.29	0.29	0.14
O ₂									0.39	0.12	-0.73	0.24	0.76
pH										-0.38	-0.58	-0.29	0.36
PO ₄ ³⁻											0.05	0.07	-0.40
SO ₄ ²⁻												-0.04	-0.54
<i>T</i>													-0.04
TC													

^a Parameters include Physicochemical parameters (pH, temperature (*T*), electrical conductivity (EC), dissolved oxygen (O₂) and soluble ions (Na⁺, K⁺, PO₄³⁻, SO₄²⁻, NO₃⁻, NO₂⁻) and faecal indicator bacteria (FIB): *Escherichia coli* (*E. coli*), *Enterococcus* (ENT) and Total coliforms (TC). Significant coefficients ($p < 0.05$) are in bold

**Fig. 4** PCA analysis of water physicochemical parameter and FIB, according to the sampling point and seasonal variation

contamination, showed a decrease and probably the evaporation of water can lead to a concentration of soluble ions in the wells. These hypotheses can explain higher concentrations of FIB during the wet season (less during the dry season) and higher concentrations of dissolved ions during the dry season (less during the wet season) (Mkandawire 2008; Pritchard et al. 2008; Msilimba and Wanda 2013). Additionally, domestic livestock and household activities as well as waste disposal could also be

major contributory factors of wells contamination during both dry and wet seasons (Banks et al. 2002).

Conclusion

The results of this study revealed that water from shallow wells of the selected neighbourhoods of the municipality of Bumbu is heavily polluted in faecal material, in both dry

and wet seasons. The situation is more alarming during the wet season. Physicochemical parameters, especially NO_3^- and EC in water samples, are unacceptable for the human consumption for both dry and wet seasons. Consequently, water from the studied zone is not appropriate for drinking or other domestic uses.

To our knowledge, this is the first study, which assessed the FIB concentration in the shallow wells of a suburban municipality of Kinshasa. Their presence in water suggests that viable human pathogens may also be present. Shallow wells represent the majority of domestic water supply in suburban municipalities of Kinshasa. More than 70 % of domestic water supply in this study site originated from shallow wells. Our results highlight on the human risk associated with exposure to water contamination from wells according to the high level of NO_3^- , EC, *E. coli* and ENT in both dry and wet seasons. The results of this study will help a better understanding of the microbiological pollution problematic in shallow wells under tropical conditions and will guide future municipality decisions on improving the water quality. Therefore, we recommend (1) the monitoring programme of water quality in shallow wells of other municipalities of Kinshasa, (2) implementation of wells protective measure (or construction of appropriate wells) and methods of water supply from shallow wells against to the risks of pollution, (3) education programme of population for sanitation tools and (4) the urgent development of appropriate and local treatment for purification of water such as the use of *Moringa oleifera* (Poumaye et al. 2012). The methods, scenarios and recommendations from this research can be performed in similar environment under tropical conditions. Nevertheless, further investigation is recommended for the assessment of emerging contaminants such as antibiotics, antibiotic resistant bacteria and antibiotic resistance genes in wells.

Acknowledgments We are grateful to the financial sources: the Swiss National Science Foundation (Grant No. 31003A_150163/1) and Swiss embassy in DRC via M. Christian Gobet. This result presents the collaboration between the University of Geneva (Institute F. A. Forel), University of Kinshasa and Pedagogic National University of Congo (Democratic Republic of Congo).

Compliance with Ethical Standards

Conflict of interest The authors declare no conflict of interest.

Ethical Standards We confirm that the field studies and sampling did not involve misunderstanding. The funder had no role in study design, data collection and analysis, decision to publish or preparation of the manuscript.

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