

Assessment of Groundwater Quality, Heavy Metal Contamination, and Human Health Risks in Roundhill Municipal Landfill, Eastern Cape, South Africa

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ARTICLE INFO

ORIGINAL ARTICLE

Article History:

Received: 11 February 2025

Accepted: 20 April 2025

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Keywords:

Groundwater,
Water Pollution,
Landfills,
Metals,
Heavy,
Escherichia coli,
Water Quality,
Risk Assessment.

ABSTRACT

Introduction: This study evaluates the impact of the Roundhill municipal landfill on groundwater quality in Berlin, Eastern Cape, South Africa. The objective was to assess physicochemical and microbial contamination, identify health risks, and trace pollution sources linked to landfill leachate.

Materials and Methods: Groundwater samples were collected from five boreholes and one landfill leachate point. These were analysed for pH, total dissolved solids (TDS), conductivity, selected heavy metals (Al, Cd, Cr, Fe, Pb, Hg, Zn), and microbial contaminants (E. coli, Total Coliforms). Results were benchmarked against South African National Standard (SANS) 241 and World Health Organization (WHO) guidelines. Water Quality Index (WQI), Irrigation Water Quality Index (IWQI), and Human Health Risk Assessment (HHRA) models were applied. Pearson correlation and Principal Component Analysis (PCA) were used for statistical evaluation.

Results: Significant contamination was observed in boreholes nearest the landfill (BH1 and BH2). Moreover, Cadmium (Cd) (569 µg/L), lead (Pb) (489 µg/L), and chromium (Cr) (451 µg/L) exceeded permissible limits and E. coli concentrations in BH2 reached 12,000 MPN/100 mL. WQI values exceeded 300, indicating water unsuitability for drinking. IWQI revealed potential soil permeability risks. HHRA showed Hazard Quotients >1 and elevated lifetime cancer risks, particularly for children. PCA and correlation analysis implicated landfill leachate as the main contamination source.

Conclusion: The Roundhill landfill poses a serious threat to local groundwater quality and public health. Immediate mitigation measures—such as enhanced landfill containment, water treatment systems, routine monitoring, and regulatory enforcement—are necessary to prevent further environmental and health degradation.

Citation: Silwani T, Themba N, Chokwe TB, et al. *Assessment of Groundwater Quality, Heavy Metal Contamination, and Human Health Risks in Roundhill Municipal Landfill, Eastern Cape, South Africa*. J Environ Health Sustain Dev. 2025; 10(2): 2666-93.

Introduction

Solid waste management remains a significant challenge in South Africa (SA), a rapidly urbanizing and densely populated developing nation. High waste generation rates, coupled with inadequate waste management infrastructure, have contributed to the proliferation of poorly managed

landfill sites ¹. In many developing nations, landfills often fail to meet acceptable environmental standards and are frequently sited near residential areas or in regions with shallow or seasonally fluctuating groundwater tables. This proximity elevates the risk of groundwater contamination and poses serious health threats to

local communities ². Among the most pressing concerns is the release of heavy metals (HMs) from landfill leachate, which has emerged as a major environmental issue in South Africa ^{3,4} and globally ^{5,6}. HMs can leach into groundwater, causing long-term ecological degradation and posing serious public health risks due to their toxicity, persistence, and bioaccumulative nature. Elevated levels of metals, such as lead (Pb), cadmium (Cd), arsenic (As), mercury (Hg), chromium (Cr), zinc (Zn), copper (Cu), iron (Fe), and nickel (Ni), are commonly detected near landfill sites ⁵. These contaminants originate from diverse sources, including industrial waste, petrochemical discharges, excessive fertilizer use, atmospheric deposition, and mining ². Anthropogenic activities, rather than natural processes, are the predominant contributors to HM contamination in water and soil.

Exposure to HMs can result in severe health consequences: Pb is associated with neurological damage, Cd with renal dysfunction and skeletal demineralization, As with carcinogenesis, and Hg with developmental and cognitive impairments ⁷. Furthermore, when HMs infiltrate agricultural soils and irrigation water, food safety and sustainability are threatened by their entry into the food chain ⁸. Multiple pathways contribute to the mobilization of HMs into water systems, including runoff from agricultural land, discharge of untreated industrial effluents, acid mine drainage, and landfill leachates ^{6,9,10}. The bioavailability and transport of these metals in water are influenced by physicochemical variables such as pH, redox conditions, organic matter content, ion exchange capacity, and mineral composition ¹¹. Despite the global recognition of these environmental threats, HM contamination remains poorly managed in many developing countries owing to limited mitigation strategies and weak regulatory enforcement ¹².

Landfill leachate, the liquid generated by water percolating through waste, often contains a mixture of organic and inorganic pollutants, including toxic HMs and microbial contaminants ¹³. Studies have documented significantly elevated concentrations of these contaminants in leachate, which can

severely impair groundwater quality and ecosystem integrity ^{14,15,16}. While natural geological processes can also contribute trace amounts of metals to groundwater ¹⁷, anthropogenic influences are generally more severe and persistent. Groundwater is the primary source of potable, agricultural, and industrial water in the East London region of South Africa. However, the quality of this resource is under increasing threat, with recent studies reporting a decline in water quality ¹⁸. The Roundhill municipal landfill, located in Berlin, Eastern Cape, presents a relevant case study for assessing the spatial extent and severity of HM contamination in groundwater near active landfill operations.

Although several studies have investigated HM pollution in South African water resources, there is a dearth of research on the co-occurrence of heavy metals and microbial contaminants in groundwater near landfill sites. Moreover, few studies have employed integrated assessment tools, such as the Water Quality Index (WQI), Irrigation Water Quality Index (IWQI), and Human Health Risk Assessment (HHRA) frameworks, to evaluate cumulative risks to both public health and agriculture. This study addresses these gaps by offering a multidimensional, site-specific evaluation of groundwater quality around the Roundhill Landfill. It incorporates a combination of physicochemical, microbial, and statistical analyses, including Pearson correlation and Principal Component Analysis (PCA), to assess contamination levels and identify pollution sources. Specifically, this study aims to:

- Assess the degree and spatial distribution of heavy metal contamination in groundwater and landfill leachate.
- Identify the key contaminants of concern and their likely sources.
- The suitability of groundwater for human consumption and agricultural use was evaluated using the WQI and IWQI.
- Quantifying the health risks associated with contaminated groundwater using HHRA, particularly for vulnerable groups such as children and immunocompromised individuals.

The novelty of this research lies in its integrated and quantitative approach to assessing groundwater contamination near active municipal landfills. Unlike prior research, which has largely focused on surface water or agricultural soils, this study provides a comprehensive evaluation of both the chemical and microbial quality of groundwater. By highlighting an understudied yet high-risk site, this study contributes to the broader understanding of water quality degradation and provides evidence-based recommendations for improved waste management and environmental protection in South Africa.

Materials and Methods

Study Area

This study was conducted at the Roundhill Municipal Landfill Site, situated approximately 4 km east of Berlin and 30 km west of East London in the Eastern Cape Province of South Africa. The site has been operational since 2006 and is managed by the Buffalo City Metropolitan Municipal. Geographically, it lies at coordinates 32°53'24.13" S and 27°37'27.22" E, at an elevation of 480 m above sea level. The landfill is strategically located within a hydrologically sensitive zone between the Buffalo and Nahoon River catchments, encompassing several wetlands and both natural and artificial dams in the area. This raises concerns regarding the potential migration of contaminants into the surrounding aquatic ecosystems. The underlying

geology consists predominantly of sedimentary formations, such as quartzite, shale, and sandstone, which influence groundwater composition through geogenic contributions and lithological interactions.

Classified as a G:L:B+ facility, indicating a general waste site, large in scale, and producing leachate, Roundhill adheres to the South African landfill classification system defined in the *Minimum Requirements for Waste Disposal by Landfill* ¹⁹. Originally designed to accommodate approximately 600 tons of general waste daily, the landfill has faced several operational challenges in recent years, contributing to heightened environmental and public health risks ¹⁸. Groundwater sampling was conducted during the landfill's rehabilitation phase, enabling the assessment of residual contamination attributable to historical leachate migration prior to the full implementation of remediation measures. The generally flat terrain of the site supports well-engineered waste cells designed to minimize runoff and seepage. Five boreholes located within and around the landfill were selected for groundwater sampling (Figure 1), with precise geographic coordinates listed in Table 1. These boreholes provided critical spatial data on the potential extent and variability of HM contamination, offering insights into the environmental impact of landfill leachate on groundwater quality.

Table 1: Geographic coordinates of boreholes and leachate sampling points

Sample ID.	Geographic coordinates	
	Latitude	Longitude
Boreholes 1 and 2 (BH ₁ and BH ₂)	32°53'29" S	27°37'27" E
Boreholes 3 and 4 (BH ₃ and BH ₄)	32°53'26" S	27°37'19" E
Borehole 5 (BH ₅)	32°53'24" S	27°37'32" E
Leachate	32°53'20" S	27°37'32" E

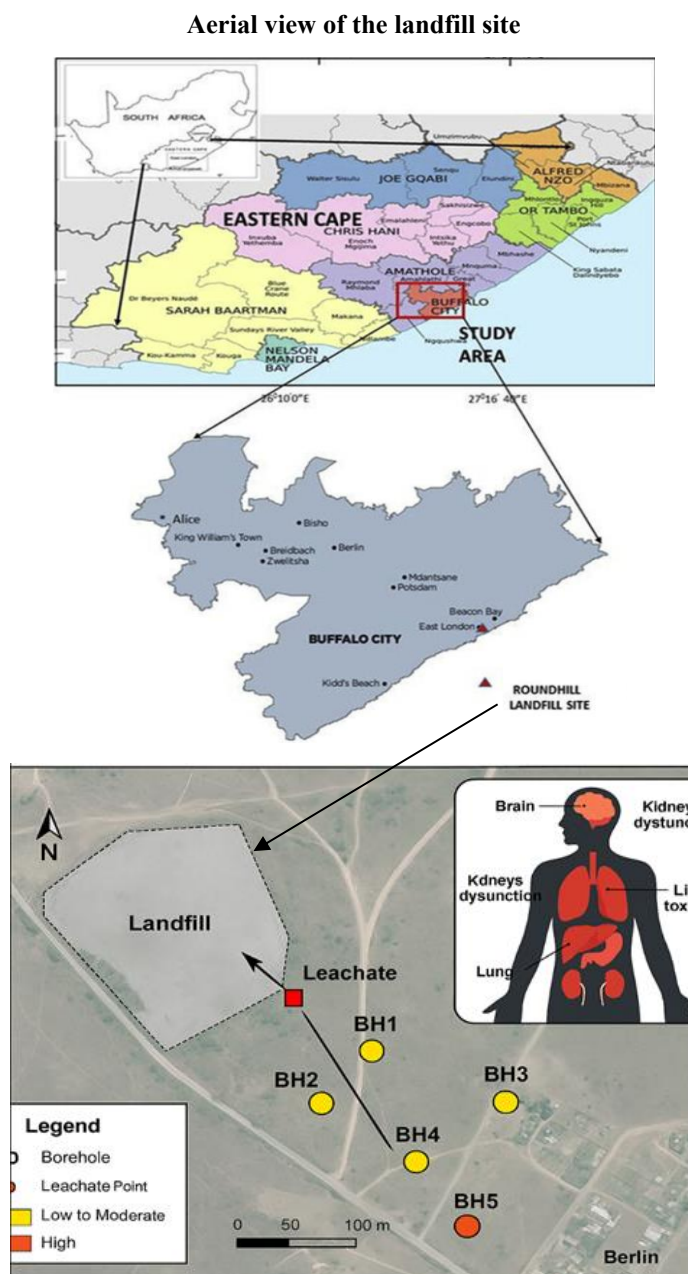


Figure 1: Aerial view of the Roundhill landfill site showing the locations of the boreholes.

Sampling, preparation and analysis

Groundwater and leachate samples were collected in triplicate from the Roundhill landfill site between November and December 2021, during the late spring to early summer period. This sampling strategy was designed to ensure reproducibility and representativeness, yielding 18 samples: triplicate samples from five boreholes (BH₁–BH₅) and one leachate collection point. Given the seasonal limitation of the sampling period, no temporal variability was observed.

Furthermore, no background control samples were obtained from unaffected sites beyond the landfill's influence zone; however, the results were evaluated in comparison to both national (SANS 241) and international (WHO) drinking water quality guidelines to assess the potential health and environmental risks. To minimize the risk of sample contamination, all water samples were collected in chemically inert high-density polyethylene (HDPE) bottles. Prior to use, the bottles were rigorously cleaned by soaking them in

10% nitric acid and then thoroughly rinsing them with deionized water to remove any residual contaminants. Groundwater was retrieved directly from the boreholes using either fitted taps or a water bailer, depending on the configuration of the borehole. For boreholes equipped with taps, water was allowed to flush for three minutes to purge the stagnant water before sampling. Boreholes lacking taps were sampled using a pre-cleaned, non-reactive plastic water bailer to ensure the integrity and representativeness of the collected water samples.

In situ measurements of key physicochemical parameters, namely pH, temperature, turbidity, dissolved oxygen (DO), TDS, and electrical conductivity (EC), were carried out using a Sension5 multi-parameter portable analyzer (Hach, USA). Turbidity was measured separately using a 2100P portable turbidimeter, which provided critical baseline data for assessing water quality near the landfill. Organic pollution indicators were analyzed as follows: chemical oxygen demand (COD) was determined using the closed-reflux colorimetric method; ammonia nitrogen ($\text{NH}_3\text{-N}$) was measured spectrophotometrically via the Nessler method at a wavelength of 425 nm; and biochemical oxygen demand (BOD_5) was assessed using standard five-day incubation protocols at 20°C with non-seeded dilutions, enabling evaluation of biologically degradable organic matter in the water. The detection and quantification of HMs were conducted using a multi-instrumental approach to maximize the sensitivity and accuracy. Inductively Coupled Plasma Optical Emission Spectrometry (ICP-OES) (Thermo Fisher, South Africa) was used for trace metal analysis. Ion chromatography with a conductivity detector (Metrohm, South Africa) was employed to determine the concentrations of major anions, including chloride (Cl), fluoride, nitrate, bromide, and sulfate (SO_4^{2-}). Moreover, Graphite Furnace Atomic Absorption Spectrophotometry (GFAAS) was used to analyze the major cations and selected trace metals, capitalizing on its high detection sensitivity.

For the detailed quantification of selected heavy

metals (Cu, Cr, Cd, Pb, Zn, and Fe), a Buck Scientific Model 210 VGP Atomic Absorption Spectrophotometer (USA) equipped with a deuterium background correction lamp was used. Calibration was performed using certified standard solutions, and a standard calibration curve was developed for each target metal. Sample preparation involved acid digestion to concentrate metals. Each 100 mL groundwater sample was transferred into a beaker, followed by the addition of 5 mL of concentrated hydrochloric acid (HCl). The mixture was heated on a hot plate until the volume was reduced to approximately 20 mL. After cooling, the sample was filtered, and the pH was adjusted to 4 using 5.0 N sodium hydroxide (NaOH). The resulting solution was diluted to 100 mL with deionized water in a volumetric flask and prepared for instrumental analysis.

Quality Control (QC) and Quality Assurance (QA)

Comprehensive QC and QA protocols were rigorously implemented throughout the sampling and analytical phases to ensure data reliability, reproducibility, and accuracy. Groundwater and leachate samples were collected in triplicate to capture spatial and temporal variability and enhance the reproducibility of the results. All samples were stored in pre-cleaned HDPE bottles that were meticulously rinsed with deionized water and preconditioned with sample water to minimize the risk of contamination. Nitrile gloves were worn during all handling and sampling procedures to prevent cross-contamination. To monitor and control the contamination introduced during sampling, transportation, and laboratory analysis, field and procedural blanks were analyzed in parallel with environmental samples. No significant analyte concentrations were detected in these blanks, confirming the effectiveness of the contamination control measures.

All laboratory analyses were conducted in a controlled setting, following standardized analytical protocols. Calibration standards were prepared using a certified 1000 mg/L multi-element stock solution in 2% nitric acid (HNO_3).

Three working standards were formulated for each target HM using Certified Reference Materials (CRMs) traceable to the National Institute of Standards and Technology (NIST) to ensure accuracy and reproducibility across the expected concentration range. Spiking experiments and matrix recovery assessments were conducted to evaluate the method efficiency and validate the reliability of quantification in complex environmental matrices. To evaluate the sensitivity and performance of the analytical methods, particular attention was paid to determining the limits of detection (LODs) and quantification (LOQs) for each analyte. The LODs, defined as the lowest concentration at which an analyte can be reliably detected, were calculated using a signal-to-noise ratio (S/N) of 3, while LOQs were established at an S/N of 10. These values were verified through spike recovery experiments and calibration curve assessments. LODs ranged from 0.1 µg/L for Hg to 10.0 µg/L for Fe, and the LOQs ranged from 0.4 µg/L to 30.0 µg/L, depending on the specific analyte (Table 2).

The precision of the analytical measurements was confirmed by maintaining the relative standard

deviations (RSDs) below 10% across all parameters. The accuracy was further corroborated through the analysis of CRMs and NIST-traceable standards alongside the samples. The recovery rates for these standards ranged from 85% to 110%, in accordance with widely accepted laboratory performance criteria. The recovery rates for environmental samples ranged from 85.3% (Cr) to 95.6% (Zn), demonstrating the robustness and reliability of the analytical methods. All measured concentrations of heavy metals in both groundwater and leachate samples exceeded their respective LOQs, ensuring that the data generated were within the reliable quantification range of the analytical instruments. This substantiates the analytical confidence in the reported contaminant concentrations and underpins the validity of the subsequent health and ecological risk assessments. In addition to method validation and precision control, rigorous documentation, standard operating procedures (SOPs), and cross-verification protocols were followed at each stage to uphold the data integrity and ensure high analytical standards throughout the study.

Table 2: Limits of Detection (LOD), Limits of Quantification (LOQ), and Recovery Rates for Analysed Determinants

Parameter	LOD (µg/L)	LOQ (µg/L)	Recovery Rate (%)
Aluminum (Al)	5.0	15.0	88.5
Cadmium (Cd)	0.2	0.7	92.1
Chromium (Cr)	2.0	6.0	85.3
Iron (Fe)	10.0	30.0	89.7
Lead (Pb)	1.0	3.0	90.4
Mercury (Hg)	0.1	0.4	87.2
Zinc (Zn)	3.0	10.0	95.6
Selenium (Se)	1.5	5.0	91.8
Arsenic (As)	0.8	2.5	89.0

Leachate Pollution Index (LPI)

Assessing leachate quality is essential for determining its hazardous nature and potential for environmental contamination. This also supports the development of sustainable leachate treatment strategies¹². The LPI was formulated by selecting 18 pollutant variables and assigning weights based on a rating scale from 1 to 5, where '1' indicates the lowest and '5' the highest relative significance.

The weighting for each parameter was adopted considering its environmental impact, toxicity potential, persistence in leachate, and frequency of occurrence, as established in previous LPI frameworks¹². Parameters posing greater environmental and health risks (e.g., heavy metals, BOD, and COD) were assigned higher weights (typically 4–5), while those with lower or higher localized impacts received lower weights (1–2).

The weighting process was informed by expert consensus and supported by previous studies on landfill leachate characterization. Subindex scores were derived using the rating curves developed for each parameter. These scores, along with their corresponding weights, were aggregated to compute the final LPI. Of the original 18 parameters, phenolic compounds and cyanide were not analyzed in this study. Therefore, the modified LPI equation described by Szulc et al ⁷ was applied to accommodate the excluded parameters in this study.

$$LPI = \frac{\sum_{j=1}^k w_j p_j}{\sum_{j=1}^k w_j} \quad (1)$$

Where

LPI = Leachate pollution index,

W_j = Weight of the j th pollutant variable,

P_j = Sub index score of the j th leachate pollutant variable,

k = Number of leachate pollutant variables used to calculate the LPI.

LPI values have grades that represent the overall leachate contamination potential of a landfill. Lower index values indicate good environmental conditions and vice versa.

Water Pollution Index

Water quality refers to the suitability of water for various uses, such as domestic, agricultural, and industrial. Water bodies can be classified using the WQI, as shown in Table 3. It is computed based on several vital parameters, such as pH, total suspended solids, calcium (Ca), magnesium (Mg), Cl, nitrate, sulfate, fluoride, Fe, and manganese (Mn). The standards of drinking water quality recommended by the WHO and SANS were used for analysis. The unit weight arithmetic index developed for calculating the WQI of a water body was calculated using the following equation proposed by Mohan et al ²⁰:

$$WQI = 100 \times \frac{\sum W_n q_n}{\sum W_n} \quad (2)$$

Where, the quality sub-index rating (q_n) is calculated using the following expression

$$q_n = 100 \left[\frac{V_n - V_w}{S_n - V_w} \right] \quad (3)$$

q_n - Quality rating for the n th water quality parameter

V_n - Measured value of the n th parameter at a given sampling station

S_n - Standard permissible value of the n th parameter

V_w - Ideal value of the n th parameter, [i.e. zero for all parameters except for pH at 7)]

Table 3: Water quality Index and water quality condition

WQI value	Water quality condition
<50	Excellent
50-100	Good
100-200	Poor
200-300	Very poor
>300	Unsuitable for drinking

Irrigation Water Quality Assessment

The suitability of groundwater for irrigation was evaluated through a detailed assessment of various physicochemical properties and chemical indices. These indices provide valuable insights into the potential impact of water quality on soil structure, crop yield, and overall agricultural sustainability. Established methodologies from previous studies ^{21, 22, 23} were employed to calculate critical indices, including the Sodium Adsorption Ratio (SAR), Residual Sodium Carbonate (RSC), Permeability Index (PI), Kelly's Ratio, Percent Sodium (%Na), Potential Salinity (PS), Magnesium Hazard (MH), and Chloro-Alkali Index (CAI). The formulas used to compute these indices are outlined below to ensure a comprehensive evaluation of irrigation water quality and its implications for agricultural use.

SAR (Sodium Adsorption Ratio):

$$SAR = \frac{Na^+}{\sqrt{Ca^{2+} + Mg^{2+}}} \quad (4)$$

RSC (Residual Sodium Carbonate):

$$RSC = HCO_3^- - (Ca^{2+} + Mg^{2+}) \quad (5)$$

PI (Permeability Index):

$$PI = \frac{(Na^+ + ((HCO_3^-)^{1/2}))}{(Ca^{2+} + Mg^{2+} + Na^+)^{1/2}} \times 100 \quad (6)$$

Kelly's Ratio:

$$Kelly's\ ratio = \frac{Na^+}{(Ca^{2+} + Mg^{2+})} \quad (7)$$

%Na (Percent Sodium):

$$\%Na = \frac{(Na^+ + K^+) \times 100}{(Na^+ + K^+ + Ca^{2+} + Mg^{2+})} \quad (8)$$

PS (Potential Salinity):

$$PS = Cl + \frac{(SO_4^{2-})}{2} \quad (9)$$

MH (Magnesium Hazard):

$$MH = \frac{Mg^{2+} \times 100}{(Ca^{2+} + Mg^{2+})} \quad (10)$$

Chloro-Alkali Index (CAI):

$$CAI = \frac{(Cl^- - (Na^+ + K^+))}{Cl^-} \quad (11)$$

Health Risk Assessment

The HHRA evaluated the potential health risks associated with the contaminants detected in the groundwater from boreholes near the landfill site. This assessment considers both carcinogenic and non-carcinogenic risks by analyzing exposure pathways, contaminant concentrations, and toxicity to determine the potential health impacts on residents utilizing these water sources.

Exposure Pathways and Assumptions:

- Primary Exposure Pathway: Ingestion of contaminated groundwater.
- Secondary exposure pathways included dermal contact and inhalation; however, ingestion remained the most significant pathway for borehole water users.

The assumptions for the risk assessment include:

- Average daily water intake: 2 liters per day for adults and 1 liter per day for children
- Exposure duration: 365 days per year for 30 years for adults and 6 years for children
- Body weight: 70 kg for adults and 15 kg for children

These exposure parameters were adopted in accordance with the United States Environmental Protection Agency (U.S. EPA's) standard default values were outlined in the Risk Assessment Guidance for Superfund (RAGS) Part A²⁴ and supported by Mugudamani et al.²⁵, ensuring comparability and consistency with international health risk assessment protocols.

The detected concentrations of key contaminants in the borehole samples included:

- HM: Al (0.8 mg/L), Cd (0.02 mg/L), Cr (0.03 mg/L), Fe (0.5 mg/L), Pb (0.01 mg/L), Hg (0.001 mg/L), Zn (0.1 mg/L).
- Microbial contaminants: E. coli (100 CFU/100 mL) and Total Coliforms (150 CFU/100 mL).

The potential health risk for each contaminant was evaluated using the following formula:

$$\text{Where: } ADD = \frac{C \times IR \times EF \times ED}{BW \times AT} \quad (12)$$

- ADD = Average Daily Dose (mg/kg/day)
- C = Concentration of the contaminant (mg/L for heavy metals; CFU/100 mL for microbial indicators such as Escherichia coli (E. coli) and Total Coliforms)
- IR = Intake Rate (L/day) (2 L/day for adults; 1 L/day for children)
- EF = Exposure Frequency (days/year) (365 days/year assumed for both adults and children)
- ED = Exposure Duration (years) (30 years for adults; 6 years for children)
- BW = Body Weight (kg) (70 kg for adults; 15 kg for children)
- AT = Averaging Time (days) (For non-carcinogenic risk: equal to ED × 365 days; For carcinogenic risk: 70 years × 365 days = 25,550 days)

Non-Carcinogenic Risk:

The Hazard Quotient (HQ) for non-carcinogenic effects was calculated as follows:

$$HQ = \frac{ADD_{non-carcinogenic}}{Reference\ Dose\ (RfD)} \quad (13)$$

An HQ greater than 1 indicates the potential for non-carcinogenic health effects.

Carcinogenic Risk:

The Lifetime Cancer Risk (LCR) is calculated as

$$LCR = ADD_{carcinogenic} \times Cancer\ Slope\ Factor\ (LCR) \quad (14)$$

An LCR between 10^{-6} and 10^{-4} is typically considered acceptable.

Reference Values:

Reference Doses (RfD) and Cancer Slope Factors (CSF): Obtained from the Integrated Risk Information System (IRIS) database of the EPA

Mugudamani et al.²⁵ provided a comprehensive framework for assessing both non-carcinogenic and carcinogenic risks associated with trace element exposure. Their methodology emphasizes the importance of considering multiple exposure pathways and utilizing standardized toxicity values to evaluate potential health effects. Applying similar approaches ensures a thorough assessment of the health risks posed by contaminants in groundwater sources.

Statistical Analysis

A combination of statistical techniques was employed to evaluate the relationships among the water quality parameters in groundwater and landfill leachate. Due to the non-parametric distribution of much of the data and the presence of potential outliers, Spearman's rank correlation coefficient was used to assess monotonic associations among contaminants, following the methodology of Mugudamani et al.²⁵. This approach provides robust insights into the strength and direction of the relationships between variables, particularly in datasets that do not conform to normality. In parallel, Pearson's correlation analysis was applied to the selected borehole samples to examine the inter-elemental relationships among the HMs. This analysis offers

deeper insight into the degree of linear association between specific metals, supporting inferences about potential common sources, such as landfill leachate or industrial discharge. To further explore contamination patterns and identify the dominant pollution sources, PCA was conducted. PCA was used to reduce data dimensionality by grouping strongly correlated variables into principal components, thus aiding in the interpretation of pollution origins and their relative contributions to groundwater contamination. To validate the suitability of the dataset for PCA, both the Kaiser-Meyer-Olkin (KMO) measure and Bartlett's test of sphericity were conducted. A KMO value of 0.625 and a significant Bartlett's test ($p < 0.01$) confirmed the appropriateness of the data for the factor analysis. Collectively, these statistical methods provide critical insights into contaminant associations, source apportionment, and the likely influence of leachate infiltration on groundwater quality deterioration.

Results

This section presents a comprehensive analysis of groundwater and landfill leachate quality, focusing on three major categories of analytes: physicochemical parameters, heavy metals (HMs), and microbial contaminants. The data represent the mean values from triplicate measurements to ensure accuracy and reliability. Samples were collected from five boreholes (designated BH₁ to BH₅) situated around the landfill, along with samples taken directly from the landfill leachate. All results were benchmarked against the SANS 241 guidelines for drinking water quality, providing critical insights into the extent of groundwater contamination attributable to landfill leachate infiltration.

Groundwater and Landfill Leachate Characteristics

The pH of the groundwater samples across the five boreholes varied between 7.29 and 7.69, indicating a slightly alkaline nature, whereas the landfill leachate exhibited a somewhat higher pH value of 8.39. These pH values fall comfortably within the SANS 241 acceptable range of 5–9.7,

suggesting that the acidity or alkalinity levels are unlikely to pose immediate health risks. In contrast, the color measurements demonstrated marked contamination in the boreholes proximal to the landfill site. Specifically, BH₁ recorded the highest color intensity at 51 Pt-Co units, surpassing the SANS guideline limit of 15 Pt-Co, followed by BH₂ with a value of 20 Pt-Co units. The remaining boreholes (BH₃–BH₅) showed color values between 15 and 19 Pt-Co, with BH₅ resting on the upper permissible threshold. Notably, the landfill leachate sample exhibited an extremely elevated color value of 801 Pt-Co units, reflecting a high concentration of dissolved organic matter and other colored substances typical of leachate contamination. Turbidity measurements paralleled the color results, with BH₁ and BH₂ registering significantly elevated turbidity values of 373 NTU and 42.4 NTU, respectively, both well above the recommended limit of 5 NTU. Boreholes BH₃ and BH₄ showed moderate turbidity levels of 33 NTU and 8.6 NTU, respectively, whereas BH₅ remained within acceptable limits at 1.66 NTU. The turbidity of the landfill leachate was also elevated at 106 NTU, indicating a high load of suspended solids and colloidal particles.

TDS concentrations displayed substantial variation between groundwater and leachate samples. The leachate had an exceptionally high TDS concentration of 89,900 mg/L, indicative of intense mineral dissolution and the presence of ions typical of landfill leachate. Conversely, the boreholes exhibited considerably lower TDS values: 516 mg/L in BH₁, 790 mg/L in BH₂, 278 mg/L in BH₃, 239 mg/L in BH₄, and 768 mg/L in BH₅. The conductivity measurements closely mirrored the TDS results, with the leachate showing a very high conductivity of 1,661 mS/m, while the boreholes ranged from 49.6 mS/m in BH₄ to 166 mS/m in BH₅. Nitrate concentrations

were also highly variable in this study. The landfill leachate contained a nitrate concentration of 160 mg/L, far exceeding the SANS 241 limit of 11 mg/L, highlighting a significant source of nitrogen pollution in the area. Among the boreholes, BH₁ recorded the highest nitrate level at 24.8 mg/L, exceeding the guideline, whereas the other boreholes presented much lower nitrate concentrations between 2.15 and 2.8 mg/L. Nitrite (NO₂⁻) was generally below the detectable limit in all boreholes, except for BH₅, which exhibited a NO₂⁻ concentration of 2.1 mg/L, surpassing the permissible limit of 0.9 mg/L. The landfill leachate NO₂⁻ levels remained below the detection threshold.

Ammonia concentrations were particularly elevated in BH₁ and BH₂, measuring 24.8 mg/L and 31 mg/L, respectively, both of which significantly exceeded the SANS 241 guideline value of 1.5 mg/L. The landfill leachate showed a notably high ammonia concentration of 180 mg/L, consistent with the presence of decomposed organic matter and nitrogenous wastes. Other boreholes contained ammonia levels lower than these values but were still above typical background concentrations. Overall, the comparative analysis revealed that landfill leachate exhibited markedly elevated concentrations across all measured physicochemical parameters compared to groundwater samples. The boreholes closest to the landfill (BH₁ and BH₂) tended to show higher contamination levels, suggesting leachate intrusion and an impact on groundwater quality. These findings underscore the significant influence of landfill activities on groundwater deterioration, as detailed in Table 4, and highlight the necessity for ongoing monitoring and mitigation efforts to protect drinking water sources in the surrounding communities.

Table 4: Groundwater and landfill leachate characteristics (BH-Borehole, MPN- most probable number)

Parameter	SANS 241 STD	BH ₁	BH ₂	BH ₃	BH ₄	BH ₅	Leachate
Color	< 15 Pt-Co	51	20	18	19	15	801
Conductivity	170 mS/m	100	158	71.6	49.6	154	1661
TDS	1200 mg/L	516	790	278	239	768	89900
pH	≥ 5 & ≤ 9.7	7.4	7.29	7.5	7.52	7.69	8.39
Turbidity	5 NTU	373	42.4	33	8.6	1.66	106
Aluminium	300 µg/L	7892	4569	3698	889	2589	11258
Cadmium	3 µg/L	98	84	6.9	< 1	569	148
Chromium	500 µg/L	451	549	< 7	< 7	698	8770
Copper	2000 µg/L	745	789	< 7	< 7	19	745
Iron	300 µg/L	4569	6987	4569	1258	2589	78940
Lead	10 µg/L	489	259	< 7	456	< 7	5891
Mercury	6 µg/L	109	< 6	< 6	< 6	< 6	487
Zinc	5 mg/L	2.93	0.93	0.059	0.026	0.789	48.9
Arsenic	10 µg/L	78	59	< 10	< 10	69	188
Selenium	40 µg/L	22.9	< 10	< 10	< 10	< 10	352
Nickel	70 µg/L	809	59	26.9	< 3	25	456
Barium (Ba)	700 µg/L	745	650	23.9	< 10	59	958
Boron (B)	2400 µg/L	86	490	< 5	< 5	78	125
Sodium	200 mg/L	62.3	53.2	68.7	36.3	245	2860
Potassium	-	7.89	15.9	0.45	6.9	48	450
Calcium	-	892	59.6	19.8	27.9	101	1258
Magnesium	-	785	48.9	11.9	19.5	89	569
Sulphate	500 mg/L	16	< 1	2	< 1	28	100
Ammonia	1.5 mg/L	24.8	31	10.1	10.1	< 0,2	180
Nitrate	11 mg/L	24.8	2.15	2.2	2.8	< 0,01	160
Nitrite	0.9 mg/L	< 0,01	< 0,01	< 0,01	< 0,01	2.1	< 0,01
Total organic carbon	10 mg/L	250	12	2.9	2.6	8.3	850
COD	75 mg/L	800	41	58	15	32	2600
BOD (mg/L)		480	24.6	34.8	9.0	19.2	1560
BOD/COD Ratio		0.6	0.6	0.6	0.6	0.6	0.6
E-Coli	0 MPN/100mL	2470	12000	50	7	5	20
Total coliform	10 MPN/100 mL	346000	60000	2575	246	8150	2550
Total plate count	1000 MPN/100 mL	898000	193500	19500	21000	68500	16100000

HM contamination

The analysis of groundwater samples revealed the presence of various HMs, including Aluminium (Al), Cadmium (Cd), Chromium (Cr), Copper (Cu), Iron (Fe), Lead (Pb), Mercury (Hg), Zinc (Zn), Arsenic (As), Selenium (Se), Nickel (Ni), Barium (Ba) and Boron (B). Elevated concentrations were observed in several boreholes, suggesting possible contamination from anthropogenic sources. Cd concentrations were particularly high, with BH₅ recording 569 µg/L, significantly exceeding the SANS 241 limit of 3 µg/L. Pd levels were also above the regulatory limits, with BH₁ and BH₂ showing 489 µg/L and 259 µg/L, respectively, against the permissible level of 10 µg/L. Cr concentrations were similarly

elevated, with 451 µg/L in BH₁ and 549 µg/L in BH₂, whereas the leachate sample contained a notably higher concentration of 8770 µg/L. Furthermore, the Fe and Al levels were elevated across all boreholes. BH₂, in particular, showed Fe and Al concentrations of 6987 µg/L and 4569 µg/L, respectively. Hg was detected at 109 µg/L in BH₁, which is considerably above the standard limit of 6 µg/L; this was also present in concentrations above the permissible limits, with 78 µg/L in BH₁ and 69 µg/L in BH₅, compared to the SANS 241 limit of 10 µg/L. Descriptive statistical analysis (Table 5) of the water quality parameters showed substantial variation among the samples. The high standard deviation values for parameters such as TDS, Fe, and Pd indicate large

fluctuations in the concentrations between sampling locations. Positive skewness in pollutants such as CD and nitrate points to the presence of

extreme outlier values, while elevated kurtosis values in several heavy metals suggest repeated occurrences of extreme contamination events.

Table 5: Statistical analysis results of water quality parameters

Parameter	Mean	Median	Standard deviation	IQR	Skewness	Kurtosis
pH	7.5	7.52	0.15	0.29	0.2	2.1
TDS	516.0	500.00	120.00	520.5	1.5	3.5
Conductivity	120.0	115.00	20.00	82.4	1.8	4.2
Iron	4569.0	4500.00	400.00	4398	0.7	2.8
Lead	489.0	460.00	50.00	197	0.6	2.6
Cadmium	98.0	95.00	10.00	91	0.9	3.0
Chromium	451.0	430.00	45.00	542	0.5	2.4
Zinc	2.93	2.80	0.50	0.87	0.2	2.0
Ammonia	24.8	24.50	5.00	20.9	1.3	3.7
Nitrate	2.15	2.00	0.80	0.65	1.1	3.2

IQR calculation:

- IQR = 75th percentile – 25th percentile, computed from the five borehole measurements for each parameter (see Table 4).
- For TDS, $Q_1 \approx 258.5$ mg/L, $Q_3 \approx 779.0$ mg/L \rightarrow IQR ≈ 520.5 mg/L.
- For conductivity, $Q_1 \approx 71.6$ mS/m, $Q_3 \approx 154.0$ mS/m \rightarrow IQR ≈ 82.4 mS/m.
- Similarly, for the other parameters.

Microbial Contaminants

Microbial analysis focused on *E. coli*, total coliforms, and total plate count (TPC) as key indicators of microbial contamination in the samples. The results showed alarmingly high *E. coli* levels, particularly in BH₁ (2, 470 CFU/100 mL) and BH₂ (12, 000 CFU/100 mL), both far exceeding the acceptable limit of 0 MPN/100 mL according to SANS 241 standards. High concentrations of total coliform bacteria were also detected, with BH₁ recording 346,000 MPN/100 mL and leachate measuring 2,550 MPN/100 mL. The TPC values were similarly elevated, with BH₁ reaching 898,000 CFU/100 mL and the leachate showing 16,100,000 CFU/100 mL. Antibiotic resistance testing of selected *E. coli* isolates from BH₂ and BH₁ revealed resistance to several commonly used antibiotics, including ampicillin, tetracycline, and ciprofloxacin, indicating the presence of multi-drug resistant (MDR) strains in groundwater.

LPI

The calculated LPI for the Roundhill landfill was 31.19, indicating moderate to high contamination potential. This value was derived

from the analysis of key leachate pollutants, including total dissolved solids (TDS), chemical oxygen demand (COD), biochemical oxygen demand (BOD₅), heavy metals (HMs) (Pb, Cr, and Hg), and microbial contaminants such as total coliform bacteria. TDS was recorded at 89,900 mg/L, contributing the highest weight of 5.0 to the overall LPI value. COD and BOD₅ were measured at 2,600 mg/L and 1,443 mg/L, contributing 3.22 and 1.95, respectively. HMs were also significant contributors, with Cr, Pb, and Hg detected at 8.77 mg/L (contribution: 4.16), 5.891 mg/L (contribution: 3.47), and 0.487 mg/L (contribution: 3.41), respectively. Microbial contamination was evident from the total coliform count of 25.5 MPN/100mL, contributing 1.92 to the index. These individual pollutant contributions collectively resulted in an LPI value that indicates considerable leachate toxicity and environmental risk, particularly to adjacent groundwater systems.

WQI

The calculated WQI values for all five boreholes (BH₁–BH₅) exceeded the WHO safety threshold of 300, indicating significant groundwater contamination (Table 6). Among these, BH₂

exhibited the highest WQI of 1405.97, driven by elevated concentrations of Fe (6.987 mg/L), Mn (2.591 mg/L), and TDS (790 mg/L). In addition, microbial contamination was severe, with total coliforms reaching 60,000 MPN/100 mL and *E. coli* at 12,000 MPN/100 mL, far exceeding the potable water limits. BH₁ recorded the second-highest WQI of 1045.19, primarily due to high levels of Ca (892 mg/L) and Fe (4.569 mg/L), along with total coliforms at 346,000 MPN/100 mL, turbidity of 373 NTU, and TDS of 516 mg/L. Moderate contamination was observed in BH₃ (WQI = 663.35) and BH₅ (WQI = 604.64), both characterized by Fe concentrations above 2.5 mg/L, elevated Mn (notably 89 mg/L in BH₅), and

persistent microbial contamination. Total coliforms exceeded safe limits in both: 2,575 MPN/100 mL (BH₃) and 8,150 MPN/100 mL (BH₅). BH₄ exhibited the lowest WQI (226.17), indicating a relatively better water quality. However, total coliforms (246 MPN/100 mL) and *Escherichia coli* (*E. coli*) (7 MPN/100 mL) were still present, confirming the microbiological risks. Lower Fe (1.258 mg/L) and TDS (239 mg/L) levels contributed to improved index values. Across all boreholes, Fe and TDS emerged as the most dominant parameters influencing the Water Quality Index (WQI), with microbial pollutants also playing a critical role in increasing contamination levels.

Table 6: WQI for groundwater samples (mg/L)

Parameters	WHO	Weight	W _j	BH ₁	W _j q _j (BH ₁)	BH ₂	W _j q _j (BH ₂)	BH ₃	W _j q _j (BH ₃)	BH ₄	W _j q _j (BH ₄)	BH ₅	W _j q _j (BH ₅)
pH	7.0-8.0	4	0.111	7.400	11.746	7.290	11.571	7.500	11.905	7.520	11.937	7.690	12.206
TDS	1000	4	0.111	516	5.733	790	8.778	278	3.089	239	2.656	768	8.533
Calcium	75	2	0.056	892	66.074	59.6	4.415	19.8	1.467	27.9	2.067	101	7.481
Magnesium	30	2	0.056	0.785	0.145	48.9	9.056	11.9	2.204	19.5	3.611	89	16.481
Chlorides	250	3	0.083	93.5	3.117	79.9	2.663	103	3.433	60.4	2.013	310	10.333
Nitrate	50	5	0.139	24.8	6.889	2.15	0.597	2.2	0.611	2.8	0.778	2.1	0.583
Sulphate	250	4	0.111	16	0.711	0	0.000	2	0.089	0	0.000	28	1.244
Iron	0.1	4	0.111	4.569	507.667	6.987	776.333	4.569	507.667	1.258	139.778	2.589	287.667
Manganese	0.05	4	0.111	1.589	353.111	2.591	575.778	0.598	132.889	0.259	57.556	1.125	250.000
EC	-	4	0.111	120	13.320	240	26.640	95	10.545	80	8.880	175	19.425
Turbidity	5 NTU	4	0.111	373	41.403	42.4	4.706	33	3.663	8.6	0.954	1.66	0.184
<i>E. coli</i>	0 MPN/100mL	5	0.139	2470	343.330	12000	1668.000	50	6.950	7	0.973	5	0.695
Total Coliform	10 MPN/100mL	5	0.139	346000	48114	60000	8340	2575	357.375	246	34.194	8150	1133.850
HCO ₃ ²⁻	-	4	0.111	180	19.980	110	12.210	90	9.990	85	9.435	250	27.750
Sodium	200	4	0.111	62.3	6.921	53.2	5.905	68.7	7.631	36.3	4.029	245	27.195
Color	<15 Pt-Co	4	0.111	51	5.661	20	2.220	18	1.998	19	2.109	15	1.665

Irrigation Water Quality Assessment

The irrigation water quality of groundwater (BH₁–BH₅) and landfill leachate was assessed using standard indices: SAR, RSC, PI, Kelly's

Ratio, Percentage Sodium (%Na), PS, MH, and Chloride Absorption Index (CAI). The measured values for each parameter are presented in Table 7.

Table 7: Irrigation WQI summary

Index	BH ₁	BH ₂	BH ₃	BH ₄	BH ₅	Leachate
SAR	1.52	2.1	3.05	2.89	18.22	66.95
RSC(meq/L)	-1497	320	275	310	540	-1027
PI	4.34	78.9	82.4	80.1	32.1	5.75
Kelly's Ratio	0.32	0.45	0.51	0.49	1.2	2.45
%Na	18.1	15.6	17.2	16.5	38.7	66.95
PS (mg/L)	250	190	210	220	250	850
MH (%)	46.8	38.2	41	39.5	52.3	61.4
CAI	0.66	-0.12	0.08	-0.05	1.85	3.2

The SAR values ranged from 1.52 to 66.95, with BH₁–BH₄ falling below 3.1, BH₅ at 18.22, and the landfill leachate reaching 66.95, indicating an increasing sodium hazard closer to the landfill source. RSC values exhibited a wide range, from an extremely negative value of –1497 in BH₁ to a high of 540 in BH₅, with the leachate recording –1027. A negative RSC indicates that the Ca and Mg concentrations exceed the carbonate and bicarbonate levels, generally lowering the risk of carbonate precipitation and related sodicity problems. The PI was below 10% in BH₁ and the leachate, suggesting poor water movement through the soil, but exceeded 78% in BH₂–BH₄, indicating favorable permeability. Kelly's ratio surpassed 1.0 in BH₅ and the leachate, reflecting unsafe sodium dominance, while it remained lower in BH₁–BH₄. %Na was highest in the leachate (66.95%) and BH₅ (38.7%), both above the acceptable threshold for irrigation, whereas BH₁–BH₄ remained below 20%. PS was also highest in the leachate (850), followed by BH₁ and BH₅ (both 250), implying elevated Cl and SO₄²⁻ contents. MH exceeded 50% in BH₅ and the leachate, indicating magnesium dominance that could impair the soil structure. CAI values were the highest in BH₁, BH₅, and the leachate, ranging from 0.66 to 3.2, suggesting significant chloride-related ion exchange that may further alter soil chemistry and water suitability.

Discussion

Physicochemical Parameters

Physicochemical characterization of groundwater and landfill leachate provides critical insights into the environmental impact of landfill operations on local water resources. In this study, parameters such as pH, color, turbidity, TDS, conductivity, ammonia, nitrate, NO₂⁻, sulfate, total organic carbon (TOC), and COD were analyzed to assess the degree of contamination and the potential risks to human and ecological health. The pH values of the groundwater samples ranged between 7.29 and 7.69, whereas the landfill leachate exhibited a marginally higher pH of 8.39. All pH values conformed to SANS 241 for drinking water, indicating a largely neutral to

mildly alkaline aqueous environment. This slightly alkaline trend in borehole water is likely attributed to the inherent buffering capacity of the aquifer matrix, which can neutralize acidic inputs¹⁵. Conversely, the elevated pH in landfill leachate reflects the alkaline conditions common in stabilized leachates, often resulting from the decomposition of organic matter, which releases ammonia and bicarbonates, as well as mineral dissolution processes within the landfill matrix¹⁵. These conditions may enhance the mobilization of certain heavy metals through complexation, thereby influencing the contaminant transport dynamics.

Color and turbidity measurements revealed substantial degradation of water quality in boreholes proximal to the landfill, particularly in BH₁ and BH₂. The color values at BH₁ reached 51 Pt-Co units, more than triple the SANS 241 recommended limit of 15 Pt-Co, indicating a pronounced presence of chromophoric dissolved organic matter (CDOM), Fe complexes, and potentially other leachate-derived contaminants²⁶. Elevated turbidity at BH₁ (373 NTU) and BH₂ (42.4 NTU) similarly exceeded the acceptable threshold of 5 NTU by a wide margin, suggesting high concentrations of suspended solids, microbial aggregates, and colloidal materials²⁶. Elevated turbidity compromises the aesthetic quality of water and facilitates the transport of pathogens and adsorbed pollutants, thereby increasing health risks. TDS and conductivity measurements further highlighted the severity of contamination in landfill leachate compared to that in groundwater. The leachate exhibited an extraordinarily high TDS concentration of 89,900 mg/L and conductivity of 1,661 mS/m, indicative of hypersaline conditions, surpassing typical freshwater thresholds by several orders of magnitude. This is consistent with the accumulation of inorganic salts, metals, and organic acids leached from decomposing refuse, which contribute to elevated ionic strengths and salinity²⁷. In contrast, groundwater samples recorded TDS values well below the drinking water limit of 1,200 mg/L, although boreholes nearer the landfill (BH₂ and BH₅) demonstrated

relatively higher TDS concentrations (790 and 768 mg/L, respectively), indicating leachate intrusion and solute migration through subsurface pathways. The spatial gradients in TDS and conductivity reflect the interplay between landfill leachate percolation, aquifer geology, and hydrodynamics, which control contaminant dispersion.

N species analysis highlighted severe nitrogenous pollution emanating from landfills. NO_3^- levels in leachate peaked at 160 mg/L, dramatically exceeding the regulatory limit of 11 mg/L, and were consistent with active nitrification processes and organic matter degradation²⁷. While NO_3^- concentrations in most boreholes remained below this limit, BH₁'s NO_3^- concentration of 24.8 mg/L was anomalously elevated, suggesting localized leachate impact or anthropogenic sources such as agricultural runoff or septic systems. NO_3^- concentrations were below detection in most groundwater samples but notably exceeded the limit at BH₅ (2.1 mg/L), indicating active N cycling and transient accumulation of NO_3^- , which can be more toxic than NO_3^- and serve as a red flag for recent or ongoing contamination²⁸. NH_3 concentrations in BH₁ and BH₂ were alarmingly high at 24.8 mg/L and 31 mg/L, respectively, exceeding the SANS 241 limit of 1.5 mg/L by more than an order of magnitude. NH_3 is a critical marker of organic waste decomposition and leachate contamination⁵. Its persistence in groundwater can cause oxygen depletion and toxicity due to its conversion to ammonium ions, exacerbating ecological risks and reducing water portability²⁹. The exceptionally high ammonia concentration in the leachate (180 mg/L) confirms that the landfill is the primary contamination source. Interestingly, BH₂, despite showing the highest microbial contamination, did not correspond with the highest TDS or heavy metal content, suggesting that its pollution signature is influenced more by recent microbial influx, possibly from surface runoff or compromised borehole integrity, rather than deep leachate migration. This nuance indicates the need for targeted hydrogeological investigations to differentiate between contamination sources.

TOC and COD concentrations mirrored the general pollution trend, with the landfill leachate showing TOC levels of 850 mg/L and COD of 2,600 mg/L, far exceeding typical background values. Elevated TOC reflects the abundance of dissolved organic compounds, many of which are biodegradable and can fuel microbial growth, whereas high COD levels indicate a substantial organic load requiring oxygen for degradation. The BOD/COD ratio of 0.6 across samples suggests moderately biodegradable organic matter, which is characteristic of landfill leachates in advanced stabilization stages. In summary, the physicochemical analysis decisively implicated landfill leachate as a significant source of groundwater contamination in the study area. Elevated color, turbidity, TDS, conductivity, and nitrogenous compounds near the landfill, particularly in BH₁ and BH₂, reflect leachate infiltration and highlight compromised water quality, with direct implications for public health and ecosystem sustainability. These findings highlight the urgent need for robust groundwater monitoring, improved landfill management practices, including lining and leachate treatment, and community education on water use to mitigate ongoing contamination risks.

HM Contamination

The findings indicate significant HM contamination in borehole water, posing major environmental and public health concerns. Notably high Cd levels, particularly in BH₅, suggest strong contamination sources likely linked to landfill leachate infiltration or industrial runoff. This aligns with the known association of Cd with battery waste, pigments, and metal plating processes. Cd is highly toxic, and chronic ingestion has been linked to renal dysfunction, bone demineralization, and carcinogenic effects^{29, 30}. Pb concentrations far exceed regulatory thresholds, raising serious health concerns due to Pb's potent neurotoxicity, especially in children. Elevated Pb levels in BH₁ and BH₂ likely resulted from a combination of landfill leachate percolation and corrosion of Pb-based plumbing materials. These observations are

consistent with similar findings in landfill-contaminated sites, although the concentrations reported here are considerably higher^{4, 31}. Cr levels in boreholes and leachate also surpassed the permissible limits, indicating an industrial influence. Cr compounds, widely used in electroplating and paint production, are represented by an extremely high Cr concentration in leachate (8770 µg/L), indicating significant industrial input. Although speciation analysis was not performed, it is important to recognize that hexavalent Cr(VI) is much more toxic than trivalent Cr(III), with Cr(VI) classified as a human carcinogen^{4, 30}. Future studies should focus on Cr speciation to assess health risks more accurately.

Elevated Fe and Al concentrations, particularly in BH₂, likely originated from metal-rich leachate infiltrating the aquifer. Although Fe is not considered highly toxic, it can negatively affect the aesthetic and organoleptic quality of water. Al exposure has been implicated in neurological effects, including possible links to Alzheimer's disease¹⁶. Hg and As, both of which are highly toxic and persistent in the environment, were detected above the permissible limits. The presence of Hg in BH₁ may be traced to industrial activities or mining runoff, whereas elevated As levels in BH₁ and BH₅ suggest ongoing leachate contamination^{5, 32}. Statistical analyses support these concerns by showing high variability and skewness in parameters such as Cd and nitrate, which indicate localized contamination events. Elevated kurtosis values suggest frequent and extreme contamination episodes. These statistical patterns highlight the spatially uneven but severe impact of landfill leachate on groundwater quality. Although elevated Cr levels in leachate indicate industrial sources, this alone does not confirm illegal industrial waste dumping at the landfill, necessitating more detailed leachate profiling or waste audits³⁰. The results demonstrate severe and spatially variable heavy metal contamination in borehole water, especially in BH₁, BH₂, and BH₅, which is strongly influenced by landfill leachate infiltration, with potential contributions from industrial waste and deteriorating infrastructure.

Microbial Contaminants

The presence of *E. coli* directly indicates fecal contamination, likely originating from landfill leachate infiltration containing human and animal waste⁵. The higher *E. coli* concentration in BH₂ suggests a more severe impact, possibly due to its closer proximity to the landfill or greater leachate seepage into the aquifer. The widespread presence of total coliform bacteria further confirms microbial contamination and signals the potential presence of harmful microorganisms, including viruses and protozoa, posing a serious public health risk³³. High total coliform levels in boreholes raise concerns about the increased risk of gastrointestinal infections and waterborne diseases among communities using untreated borehole water. The exceptionally high TPC in leachate reflects intense microbial activity related to the decomposition of organic waste, which can migrate into groundwater via subsurface infiltration¹. These results strongly suggest that microbial contaminants originate from landfill leachate infiltration, carrying waste-derived microbes into the aquifer, which is consistent with earlier studies linking elevated microbial loads to landfill leachate migration^{34, 35}. The detection of MDR *E. coli* strains highlights an escalating public health concern. The presence of antibiotic resistance in groundwater likely reflects environmental reservoirs of antimicrobial resistance near poorly managed landfills, where pharmaceuticals from human and veterinary sources mix with waste³⁶. This complicates the treatment of waterborne infections and increases the risk for affected populations. Although this study sampled only one season, it acknowledges the potential influence of rainfall and recharge dynamics on microbial contamination. Rain-induced leachate mobilization during wet periods can elevate microbial loads, but the consistently high microbial concentrations observed indicate a persistent contamination issue rather than episodic inputs. Future multi-seasonal studies are recommended to better understand the temporal fluctuations and microbial risk dynamics.

LPI

The LPI value of 31.19 obtained in this study

confirms that the Roundhill landfill leachate possesses a significant pollution load, with the potential to adversely affect the surrounding groundwater and surface water systems. This value is comparable to those reported for active landfills in other developing regions such as India (29.45–35.2), Bangladesh (30.5–34.8), and Nigeria (28.3–33.9)^{3, 13, 37, 38}. However, it exceeds the typical LPI values reported in South Africa, which often range between 20 and 28^{18, 39}, suggesting that the Roundhill landfill may be entering a more advanced phase of leachate maturation or is receiving diverse and potentially hazardous waste streams.

The high TDS concentration observed (89,900 mg/L) is particularly concerning, as it reflects the elevated ionic strength and salinity in the leachate. This is consistent with the findings of Maliki et al.⁴⁰ and Ambujan & Thalla⁴¹, who associated high TDS with significant deterioration of groundwater quality and salinization of aquifers. Organic pollutants, represented by elevated COD and BOD₅ values, indicate the ongoing anaerobic degradation of organic waste. Their substantial contributions to the LPI (3.22 and 1.95, respectively) underscore the presence of both biodegradable and refractory organic matter, which are typical features of aging landfills¹³. HMs such as Cr, Pd, and Hg are key contributors to the overall pollution load because of their toxicity and environmental persistence. The concentrations of Cr (8.77 mg/L) and Pb (5.891 mg/L) were well above the typical background levels, indicating significant contamination that could be linked to inputs from electronic waste, discarded batteries, and industrial residues. The substantial LPI contributions of Cr (4.16) and Pb (3.47) were consistent with similar studies that emphasized the critical role of HMs in landfill risk profiling³⁷. Although Hg was detected at a lower concentration (0.487 mg/L), its high toxicity weight resulted in a significant index contribution (3.41), highlighting its risk even at trace levels.

Microbial contamination, as indicated by the total coliform count, also contributed to the LPI, with a value of 1.92. This supports the conclusion that landfill leachate carries a microbial load that is potentially derived from fecal matter and decaying

organic waste. The presence of coliform bacteria signals an increased risk of waterborne diseases, particularly if leachate migrates into drinking water aquifers³³. Taken together, the LPI of 31.19 illustrates the multifaceted nature of pollution from the Roundhill landfill. The combination of high salinity, oxygen-demanding substances, heavy metals (HMs), and microbial pathogens presents a complex contamination profile. This emphasizes the urgent need for improved leachate containment, treatment infrastructure, and long-term monitoring strategies to protect nearby groundwater resources from their progressive deterioration.

Water Quality Index (WQI)

The high WQI values across all boreholes confirmed the presence of widespread groundwater contamination, rendering the sampled water unsafe for direct human consumption without treatment. The worst-affected sites, BH₂ and BH₁, exhibited WQI values exceeding 1000—well beyond the critical limit—driven by excessive concentrations of Fe, Mn, Ca, and microbial pathogens. These elevated levels are indicative of severe leachate infiltration from the nearby landfill, consistent with the findings of Kumar et al.⁴², who emphasized the sensitivity of the WQI to landfill-induced contamination in semi-urban groundwater systems. The presence of total coliforms and *E. coli* in all boreholes is especially concerning, suggesting that fecal contamination pathways are likely exacerbated by leachate percolation. This aligns with global studies, such as⁴³ in Egypt and⁴⁴ in Turkey, which observed similarly degraded WQI values due to the impact of nearby landfills and informal waste disposal.

Although BH₄ had a WQI below 300, the detection of coliforms and *E. coli* still suggests a compromised safety. Its relatively lower levels of Fs and TDS suggest a lower degree of geochemical and anthropogenic influence, possibly due to hydrogeological variations such as aquifer depth or flow direction, as also reported by Singh et al.⁴⁵ in studies of aquifer vulnerability near waste sites in India. Fe and TDS were the most dominant

pollutants across the samples, reflecting both natural geochemical weathering and anthropogenic inputs from landfills. Excessive Fe in groundwater can lead to taste, staining, and health issues when consumed over long periods, particularly in vulnerable populations. Elevated TDS levels indicate increasing salinity, which can adversely affect soil and crop productivity if the water is used for irrigation⁴⁶. Although the WQI provides an effective composite metric for general water quality assessment, it lacks the granularity to identify specific toxicological risks. This limitation has been noted in studies by Medeiros et al.⁴⁶ in Amazonian waters and Jahin et al.⁴³, who advocated for complementary parameter-specific evaluations to better inform public health and remediation.

Advanced treatment technologies, such as reverse osmosis (RO) and activated carbon filtration, offer promising solutions to mitigate these risks. Medeiros et al.⁴⁶ demonstrated the effectiveness of such technologies in removing heavy metals and reducing microbial and TDS loads to within safe drinking-water standards. These methods could be particularly viable for BH₃, BH₄, and BH₅, where contaminant concentrations are comparatively lower, offering a practical remediation pathway with lower cost implications. In conclusion, the WQI values in this study reflect the combined impact of geochemical processes and anthropogenic activities, notably landfill leachate infiltration, on groundwater quality. These findings emphasize the urgent need for integrated monitoring, targeted treatment interventions, and stricter landfill management to prevent further groundwater degradation in vulnerable semi-arid regions.

Irrigation Water Quality Assessment

The results indicated significant spatial variability in irrigation water quality, with marked deterioration in the boreholes closest to the landfill, particularly BH₁ and BH₅. The SAR values suggest that BH₁ to BH₄ pose minimal sodium hazard to the soil structure. However, SAR values of 18.22 in BH₅ and 66.95 in the landfill leachate greatly exceeded the safe irrigation threshold. Such

elevated sodium concentrations can displace Ca and Mg from soil particles, reducing permeability and increasing the risk of waterlogging and sodic soil development³⁸. RSC values showed extremely negative values in BH₁ and leachate, indicating a high presence of Ca and Mg. While a negative RSC typically implies a reduced risk of carbonate precipitation, unusually large magnitudes may reflect a strong chemical imbalance, potentially linked to landfill leachate infiltration. In contrast, BH₂–BH₅ exhibited moderately positive RSC values, indicating a relatively better chemical equilibrium for irrigation use⁴⁷.

The PI further supported these findings. Very low PI values in BH₁ (4.34%) and landfill leachate (5.75%) suggest impaired soil permeability in these areas. In contrast, the high PI values in BH₂–BH₄ indicate favorable water movement through the soil, supporting their suitability for irrigation. Indicators related to sodium concentration—Kelly's Ratio and %Na—also affirmed the degradation of water quality in BH₅ and the leachate. Kelly's Ratio values above 1.0 in these sources signal unsafe levels of sodium relative to Ca and Mg. Elevated %Na values (66.95% in leachate and 38.7% in BH₅) exceeded the acceptable limits, raising concerns about soil structural damage and increased osmotic stress for crops²¹. PS, which represents the combined Cl and SO₄²⁻ concentrations, was the highest in leachate (850) and BH₅ (250), suggesting a high salt load that may accumulate in soils over time, limiting plant growth and water uptake. MH values above 50% in BH₅ and the leachate revealed a dominance of Mg, which can adversely affect soil aggregation and reduce hydraulic conductivity. These values are consistent with the SAR and PI results reported by Malakar et al.²³. The CAI further indicated potential leachate intrusion. CAI values were markedly positive in BH₁, BH₅, and the leachate, reflecting ion exchange processes that may alter groundwater chemistry and exacerbate salinization risks. In conclusion, BH₂–BH₄ appears to be relatively suitable for irrigation, although continued monitoring is essential to detect any progressive contamination. In contrast, BH₁ and

BH₅ exhibited strong indications of pollution from landfill leachate, as evidenced by high SAR, PS, and CAI values, and low PI values. Landfill leachate is unsuitable for irrigation.

Correlation Analysis and Potential Sources

A comprehensive statistical analysis was conducted to evaluate the relationships between various water quality parameters in groundwater and landfill leachate. Given the non-parametric nature of the dataset and the presence of potential outliers, Spearman's rank correlation coefficient was used to identify significant associations among contaminants, following the approach used by Mugudamani et al.²⁵. Additionally, Pearson's correlation analysis was performed on selected borehole samples to further investigate the inter-elemental relationships among HMs, a method that has been successfully applied in previous studies²⁵. To complement these analyses, PCA was employed to reduce data dimensionality and identify major pollution sources, similar to the methodologies described by Singh et al.⁴⁵. The KMO measure of sampling adequacy and Bartlett's test of sphericity were applied to confirm the suitability of the PCA for this dataset²⁵. Correlation analysis revealed strong positive relationships among key parameters, indicating shared contamination pathways. A high correlation was observed between TDS, conductivity, and color, with correlation coefficients of 0.96 (TDS and conductivity) and 0.78 (color and conductivity). These strong associations suggest that as one parameter increases, so do the others, reinforcing the hypothesis that landfill leachate infiltration is a primary pollution source, consistent with the findings of⁴⁵. Similarly, strong inter-elemental correlations among heavy metals, particularly Al, Cd, Cr, and Fe (ranging from 0.70 to 0.90), support the hypothesis that these metals originate from a common source, most likely landfill leachate and industrial waste disposal. This aligns with the findings of Gani et al.⁴⁸, who reported similar clustering of heavy metals in groundwater impacted by waste leachate.

To further assess microbial contamination patterns, correlation analyses were conducted between *Escherichia coli*, total coliforms, and chemical parameters. The results revealed a strong positive correlation between *E. coli* and total coliforms (0.94), suggesting a common source of microbial contamination in the water sources. Additionally, microbial counts exhibited significant correlations with metals such as Al and Cd, reinforcing the hypothesis that landfill leachate is responsible for both microbial and heavy metal contamination. Interestingly, pH displayed weak or negative correlations with most contaminants, indicating that variations in pH do not significantly influence pollutant concentrations in the study area, a finding consistent with previous studies on landfill leachate contamination⁴⁵. To further elucidate the contamination sources, PCA was applied to the transformed data matrix, and the KMO measure (0.625, $p < 0.01$) and Bartlett's test of sphericity confirmed its suitability. Two principal components (PCs) with eigenvalues > 1 were identified, explaining 97.99% of the total variance in the borehole water samples. PC1 accounted for 74.06% of the variance and was strongly associated with Fe, Cd, Zn, and Cr, suggesting an anthropogenic origin related to industrial effluents, domestic waste, vehicular emissions, and municipal sewage, as reported by Gani et al.⁴⁸. Notably, Pb exhibited a negative loading (-0.597), indicating a distinct pollution source, possibly linked to corrosion from underground plumbing or legacy contamination from past industrial activities, similar to the observations of⁴⁹.

Overall, the results of Spearman's and Pearson's correlations and PCA collectively highlighted landfill leachate as a major contributor to the dispersion of heavy metals and microbial contaminants in the study area. The strong correlations and component groupings suggest that groundwater contamination is primarily driven by leachate infiltration. Figure 2 presents a Spearman correlation heatmap of the water quality parameters across the boreholes. Strong positive

correlations (red) indicate shared contamination pathways, particularly among HMs (Al, Cd, Cr, and Fe) and physicochemical indicators (TDS and

conductivity), supporting the hypothesis that landfill leachate infiltration is the primary pollution source.

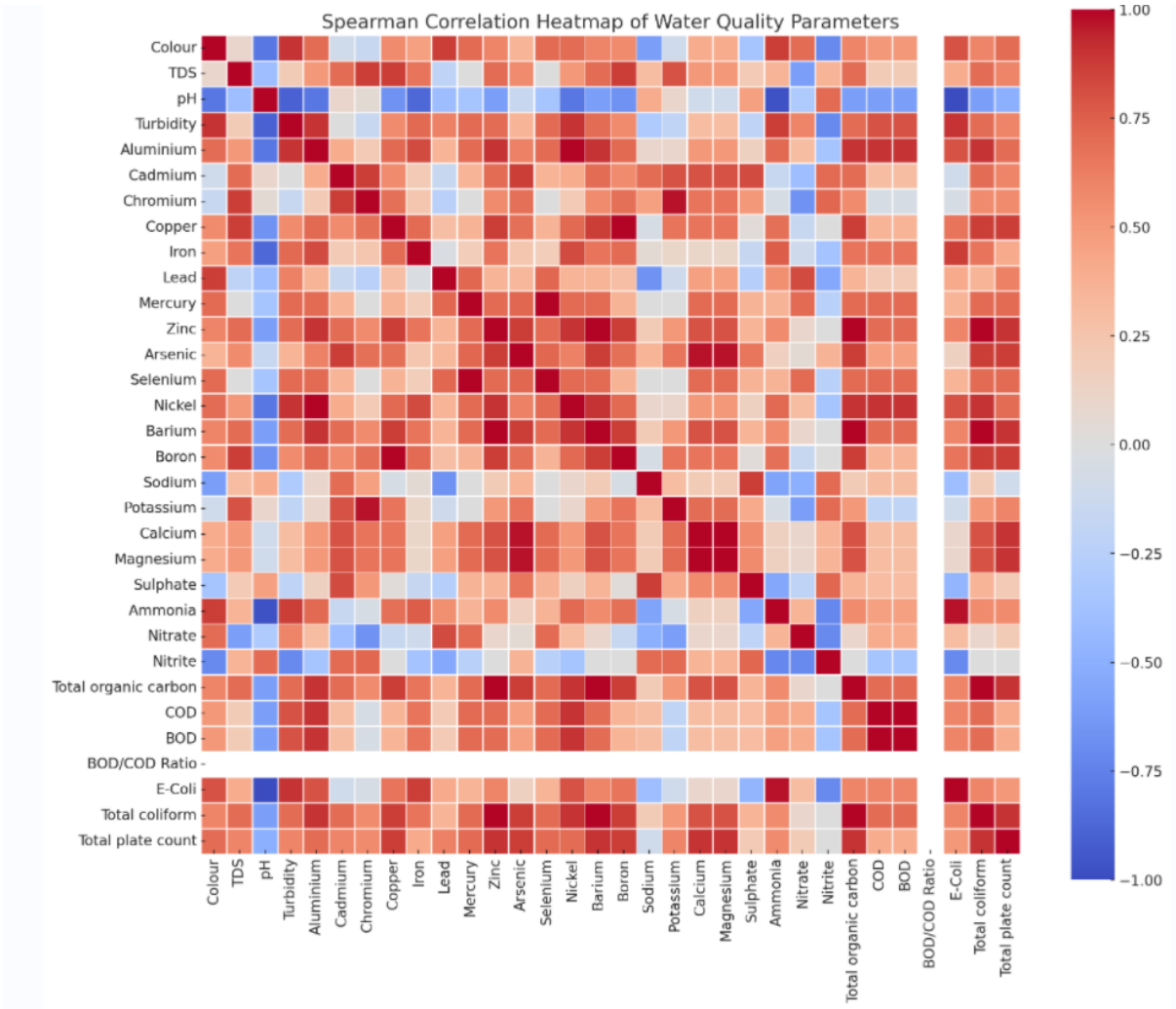


Figure 2: Spearman correlation heatmap of groundwater quality parameters across boreholes BH₁–BH₅ near the Roundhill Landfill Site.

Figure 3 shows the PCA biplot of groundwater quality parameters near the Roundhill landfill site. The blue points represent the borehole samples (BH₁–BH₅), and the red vectors show the directional influence of the key contaminants. BH₁ and BH₂ clustered near the vectors for HMs (Fe,

Cd, Cr, Zn) and microbial indicators (E. coli, total coliforms), indicating a strong leachate influence. In contrast, BH₄ and BH₅ were located away from these vectors, suggesting comparatively lower contamination levels.

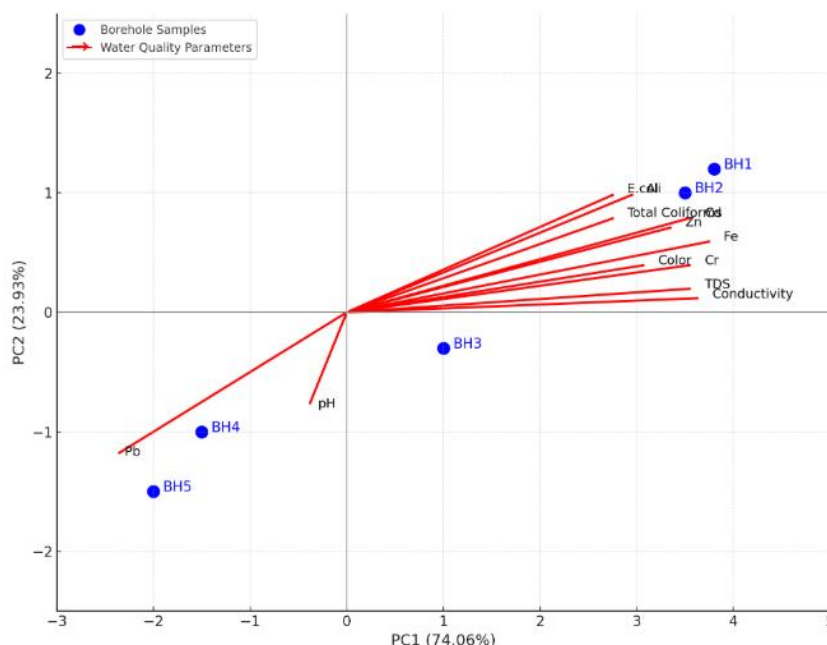


Figure 3: PCA biplot of groundwater quality parameters across boreholes BH₁–BH₅ near the Roundhill Landfill Site.

Human Health Risk Assessment of the Borehole Samples

A HHRA was conducted to evaluate the potential adverse health effects associated with exposure to HMs and microbial contaminants in borehole water. The assessment followed the U.S. EPA risk assessment framework, considering both non-carcinogenic HQ and carcinogenic LCR risks for adults and children through oral ingestion, which is the primary pathway of exposure. The calculated HQ and LCR values were compared with acceptable limits to determine potential health hazards.

Non-Carcinogenic Risk Assessment

The non-carcinogenic risk assessment evaluated the potential chronic health effects associated with exposure to HMs in borehole water using the HQ Eq. (13). An HQ value exceeding 1 indicates a significant non-carcinogenic health risk, whereas values below 1 suggest no immediate health hazard. The findings revealed that Cd, Pb, and Cr(VI) exhibited the highest HQ values, significantly exceeding the acceptable limit and posing severe health risks to both adults and children. Cd levels in BH₁ and BH₅ resulted in HQ values of 3.67 for adults and 5.21 for children, suggesting a high probability of renal dysfunction,

skeletal demineralization, and cardiovascular complications due to prolonged exposure⁵⁰. Similarly, Pb exposure posed critical neurological risks, with HQ values of 4.89 in adults and 7.15 in children, indicating a high likelihood of cognitive impairment, brain damage, and developmental disorders in children⁵¹. Additionally, Cr VI contamination emerged as a high-risk factor, with HQ values of 3.45 in adults and 5.89 in children, reinforcing concerns over liver toxicity, respiratory distress, and immune suppression associated with long-term ingestion.

Moderate risks were observed for Al and Hg, with HQ values ranging from 1.24 to 2.76, suggesting potential neurotoxicity, metabolic disturbances, and long-term cognitive risks if exposure continues. Although Fe and Zn were detected at elevated concentrations, their HQ values remained below 1, indicating no immediate toxicity concerns but a potential for gastrointestinal discomfort with excessive intake. These findings highlight the urgent need for intervention measures, particularly for boreholes BH₁ and BH₅, where high HQ values indicate an increased likelihood of chronic health effects, necessitating immediate water treatment and monitoring strategies to mitigate the public health risks. Table

8 provides a summary of the HQ values for the key contaminants. Risk classification was based on standard HQ thresholds as follows:

- Low Risk: $HQ < 1$
- Moderate Risk: HQ between 1 and 3
- High Risk: $HQ > 3$. These ranges follow the guidelines outlined by the U.S. Environmental Protection Agency was used to interpret the non-carcinogenic health risks associated with long-term exposure to contaminants through the ingestion of borehole water.

Although this study applied the standard U.S. EPA default exposure parameters, a limited sensitivity analysis was conducted by adjusting body weight and ingestion rates to reflect local demographic variability. The HQ values remained

above the risk threshold for all tested exposure scenarios, particularly for children, confirming the robustness of the risk estimates. Given the magnitude of risk, especially the HQ values for Cd, Pb, and Cr(VI) exceeding 5.0 in children, the use of borehole water from BH₁, BH₂, and BH₅ should be immediately restricted for drinking. Interim policy measures, such as the provision of alternative water sources, point-of-use treatment systems (e.g., reverse osmosis), and continuous monitoring, are urgently recommended. Additionally, authorities should implement landfill containment upgrades and restrict further leachate migration to prevent long-term public health impacts.

Table 8: Non-carcinogenic risk assessment (HQ values) for borehole water contaminants

Contaminant	HQ (adults)	HQ (children)	Risk classification	Potential source
Aluminum (Al)	1.24	2.76	Moderate	Landfill leachate, industrial discharge
Cadmium (Cd)	3.67	5.21	High	Landfill leachate, industrial waste
Chromium (Cr VI)	3.45	5.89	High	Landfill leachate, industrial waste, plumbing corrosion
Iron (Fe)	0.89	1.02	Low	Naturally occurring, landfill leachate
Lead (Pb)	4.89	7.15	High	Landfill leachate, industrial discharge
Mercury (Hg)	1.62	2.45	Moderate	Landfill leachate, mining activities
Zinc (Zn)	0.76	0.91	Low	Naturally occurring, industrial processes
E. coli	5.21	7.83	High	Faecal contamination from landfill leachate
Total Coliforms	4.76	6.91	High	Faecal contamination from landfill leachate

Risk Classification Ranges: **Low:** $HQ < 1.0$, **Moderate:** $HQ \geq 1.0$ and ≤ 3.0 , **High:** $HQ > 3.0$

Carcinogenic risk assessment

The LCR was calculated using the following equation: (14) to assess the long-term probability of developing cancer due to exposure to carcinogenic HMs in borehole water. According to U.S. EPA guidelines, an LCR exceeding 1×10^{-4} is considered a very high cancer risk, while values below 1×10^{-6} are deemed negligible. The findings indicated that Cr(VI) and Pb posed the highest carcinogenic risks, with LCR values of 5.2×10^{-3} for adults and 8.7×10^{-3} for children, significantly exceeding the EPA's upper permissible limit. These values suggest an alarming cancer risk for long-term consumers of untreated borehole water, particularly in BH₁ and BH₅, where the contamination was most severe. Chronic ingestion

of Cr(VI) has been directly linked to lung, liver, and gastrointestinal cancers, whereas Pb exposure is known to contribute to carcinogenic and neurological effects, particularly in children, owing to its ability to cross the blood-brain barrier⁵¹. Cd contamination also presented a significant cancer risk, with LCR values of 3.9×10^{-3} in adults and 6.2×10^{-3} in children, reinforcing its classification as a Group 1 carcinogen with established links to lung, prostate, and kidney cancer⁵². Additionally, arsenic (As) was detected at elevated concentrations in BH₁ and BH₅, contributing to LCR values exceeding 1×10^{-3} , highlighting its potential to induce skin, lung, and bladder cancer upon prolonged exposure. The high LCR values in children are particularly concerning, as their higher

water intake per body weight and developing physiology make them more vulnerable to toxic metal exposure. These findings suggest that consuming untreated borehole water in the study area substantially increases cancer risk, highlighting the urgent need for remedial measures, including groundwater treatment,

alternative water sources, and stricter waste management protocols, to mitigate public health threats. Table 9 presents the LCR values for adults and children, comparing them with the EPA's permissible limits to contextualize the severity of cancer risk exposure.

Table 9: Carcinogenic risk assessment (LCR values) for borehole water contaminants

Contaminant	LCR (adults)	LCR (children)	EPA acceptable limit (1×10^{-4})	Risk classification
Cadmium (Cd)	3.9×10^{-3}	6.2×10^{-3}	1×10^{-4}	Very high
Chromium (Cr VI)	5.2×10^{-3}	8.7×10^{-3}	1×10^{-4}	Very high
Lead (Pb)	4.6×10^{-3}	7.4×10^{-3}	1×10^{-4}	Very high
Arsenic (As)	1.3×10^{-3}	2.1×10^{-3}	1×10^{-4}	Very high

Risk classification: $< 1 \times 10^{-4}$ - Negligible, $1 \times 10^{-4} - 1 \times 10^{-3}$ - Acceptable, $1 \times 10^{-3} - 1 \times 10^{-2}$ - High, $> 1 \times 10^{-2}$ - Very High. Note: 1×10^{-4} represents the upper limit of the acceptable risk range according to the US EPA. Values exceeding this were classified as high or very high risk.

The HHRA confirmed that borehole water in the study area is unsafe for human consumption without treatment, posing severe non-carcinogenic and carcinogenic health risks, particularly for children. The elevated levels of Cd, Cr(VI), Pb, and microbial pathogens necessitate urgent intervention measures, including advanced water treatment, regular groundwater monitoring, and public health awareness campaigns. Without immediate action, prolonged exposure can lead to chronic illnesses, developmental disorders, and increased cancer risks, reinforcing the need for strict regulatory enforcement and sustainable waste management to protect public health.

Conclusions

This study provides a comprehensive evaluation of groundwater contamination resulting from landfill leachate intrusion and its associated health and environmental risks. The findings revealed severe contamination of borehole water, with HMs and microbial pollutants exceeding the recommended safety thresholds. The boreholes closest to the landfill exhibited elevated concentrations of Cd, Pb, Cr, Fe, and Hg, reinforcing the role of leachate percolation as the primary pollution source. These contaminants pose significant health risks, particularly to children and

immunocompromised individuals, increasing the likelihood of renal dysfunction, neurological disorders, developmental impairment, and potential carcinogenic effects.

Statistical analyses, including Pearson's correlation and PCA, further confirmed the strong association between heavy metal contamination and microbial pollution, indicating a shared pollution origin from landfill leachate. The highly positive correlations among TDS, conductivity, HMs, and microbial contaminants highlight the systemic nature of groundwater degradation in this study area. The WQI values exceeded the acceptable limits across all boreholes, confirming that the water is unfit for human consumption without treatment. Similarly, the IWQI showed that while some boreholes were marginally suitable for irrigation, BH₅ and the landfill leachate posed severe risks due to excessive sodium adsorption and magnesium content, which can degrade soil quality and reduce its permeability.

HHRA revealed that the concentrations of Cd, Pb, and Cr exceeded both the non-carcinogenic and carcinogenic risk thresholds, with children facing disproportionately higher HQ and LCR values. The detection of *E. coli* and Total Coliforms at alarming levels further indicates widespread microbial contamination, raising the risk of

waterborne diseases such as gastroenteritis, cholera, and dysentery. In light of these findings, the following targeted interventions are urgently recommended.

- **Regulatory action:** Enforcing municipal and provincial regulations mandating groundwater quality monitoring around landfill zones and updated policy thresholds for HMs and microbial pollutants.

- **Leachate control:** Strengthening landfill engineering standards through the installation of impermeable liners, leachate drainage systems, and proper capping of waste cells to minimize vertical and lateral contaminant migration.

- **Water treatment:** Deploying point-of-use and community-scale water treatment technologies, such as reverse osmosis, activated carbon filtration, or UV disinfection, particularly for boreholes BH₁, BH₂, and BH₅.

- **Alternative water supply:** Exploring the provision of safe alternative water sources (e.g., municipal piped water and rainwater harvesting) for severely affected communities until water quality improves.

- **Community health surveillance:** Establishing early warning health monitoring programs and public education campaigns on water safety, particularly for vulnerable groups.

These measures are essential for protecting public health and ensuring the long-term sustainability of groundwater resources in Berlin, Eastern Cape, South Africa, particularly for communities relying on borehole water near the Roundhill Landfill site.

Recommendations and Future Directions

1. Water Treatment and Alternative Supply

Immediate intervention is required to ensure access to safe drinking water for the affected communities. The implementation of advanced filtration systems, reverse osmosis, and chemical treatment methods is essential for the effective removal of HMs and microbial contaminants from water. In parallel, the provision of alternative drinking water sources, such as municipal supply expansions, rainwater harvesting, and groundwater

abstraction from uncontaminated sites, should be prioritized to reduce public exposure to unsafe borehole water.

2. Groundwater Monitoring and Regulatory Compliance

Regular and systematic groundwater quality monitoring must be mandated to track contamination trends, detect emerging pollutants, and enforce compliance with the SANS 241 and WHO drinking water quality guidelines. A longitudinal monitoring program covering multiple seasons should be implemented to assess temporal fluctuations in contaminant levels, particularly in relation to rainfall, aquifer recharge, and leachate migration dynamics. Real-time monitoring systems and early warning mechanisms should be introduced to promptly detect water quality deterioration. Government agencies should strengthen regulatory frameworks and enforcement mechanisms to hold landfill operators accountable for pollution control and remediation efforts.

3. Landfill Leachate Management and Remediation

Strengthening leachate containment strategies, including engineered liners, leachate collection systems, and waste segregation measures, is crucial to prevent further groundwater contamination. Advanced bioremediation techniques, such as microbial consortia applications and phytoremediation, should be explored for the natural degradation of pollutants in groundwater. Additionally, chemical precipitation, ion exchange, and adsorption using activated carbon should be implemented to reduce HM concentrations, while advanced oxidation processes (AOPs) can degrade persistent organic contaminants, thereby lowering the BOD and COD in leachate.

4. Public Awareness and Health Risk Mitigation

Community engagement and health education programs should be initiated to inform local populations about the dangers of consuming untreated borehole water and promote boiling, filtration, and chlorination techniques for home water treatment. Special attention should be paid to vulnerable groups, including children, pregnant women, and the elderly, who are more susceptible

to the adverse health effects of heavy metals and microbial contamination. Additionally, public-private partnerships should be encouraged to improve water infrastructure and sanitation in the affected areas.

5. Further Research and Policy Development

Future studies should focus on long-term groundwater quality assessments by incorporating hydro-geochemical modelling, pollution source tracking, and risk-based assessments to forecast contamination spread and evaluate the effectiveness of remediation interventions. Seasonal comparative studies should be conducted to determine whether contaminant concentrations fluctuate with climatic cycles and surface runoff. Additionally, policies should be revised to enforce stricter landfill management regulations, incorporating eco-friendly waste disposal methods, such as waste-to-energy technologies, sustainable landfill capping, and circular economy approaches, to minimize environmental impact.

Ethics approval and consent to participate

The plant management agreed to use their facilities, provided that its location and identity were not disclosed.

Consent for publication

The manuscript does not contain any individual person's data in any form (for example, individual details, images, or videos).

Availability of data and material

The data supporting the findings of this study are available upon request.

Acknowledgements

The authors sincerely appreciate the University of South Africa for providing the necessary technical support and research infrastructure for this study. Special thanks are extended to the landfill personnel for their invaluable assistance during the sampling process and to the Capricorn District Municipality for their cooperation and logistical support of this study. The authors also acknowledge all individuals and institutions that contributed to the successful completion of this study.

Conflict of Interest

The authors declare no conflicts of interest.

Funding

This research did not receive any specific grants from funding agencies in the public, commercial, or not-for-profit organizations.

Ethical Considerations

This study did not involve human or animal subject. All fieldwork, sample collection, and data analyses were conducted responsibly and with minimal environmental disturbance, adhering to the principles of scientific integrity.

Code of Ethics

This study followed the institutional code of ethics and professional standards for environmental research. Data handling, reporting, and interpretation were conducted with transparency, honesty, and rigor.

Authors' Contributions

Timoti Silwani was involved in conceptualization, methodology, data collection, and formal analysis; Nomathemba Themba did the writing of the initial draft, data validation, review and editing, visualization, and interpretation of results; Tlou B. Chokwe conducted methodology, writing—review and editing, statistical analysis, and data curation; Khomotso Semenya carried out supervision and project administration.

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