Associations between fluoride exposure in drinking water and cognitive deficits in children: A pilot study

Tewodros Rango Godebo, Marc Jeuland, Redda Tekle-Haimanot, Biniyam Alemayehu, Arti Shankar, Amy Wolfe, Nati Phan

| PII: | \$0892-0362(23)00143-5 |
|----------------|---|
| DOI: | https://doi.org/10.1016/j.ntt.2023.107293 |
| Reference: | NTT 107293 |
| To appear in: | Neurotoxicology and Teratology |
| Received date: | 9 April 2023 |
| Revised date: | 4 September 2023 |
| Accepted date: | 7 September 2023 |
| | |

Please cite this article as: T.R. Godebo, M. Jeuland, R. Tekle-Haimanot, et al., Associations between fluoride exposure in drinking water and cognitive deficits in children: A pilot study, *Neurotoxicology and Teratology* (2023), https://doi.org/10.1016/j.ntt.2023.107293

This is a PDF file of an article that has undergone enhancements after acceptance, such as the addition of a cover page and metadata, and formatting for readability, but it is not yet the definitive version of record. This version will undergo additional copyediting, typesetting and review before it is published in its final form, but we are providing this version to give early visibility of the article. Please note that, during the production process, errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

© 2023 Published by Elsevier Inc.





Associations between fluoride exposure in drinking water and cognitive deficits in children:

A pilot study.

Tewodros Rango Godebo^{a,*}, Marc Jeuland^b, Redda Tekle-Haimanot^c, Biniyam Alemayehu^c, Arti

Shankar^d, Amy Wolfe^e, Nati Phan^a

^a Department of Environmental Health Sciences, School of Public Health and Tropical Medicine, Tulane University, New Orleans, LA 70112 USA

^b Sanford School of Public Policy, Duke University, Durham, NC 27708 USA

^c Addis Ababa University, School of Medicine, Department of Neuroiosy, Addis Ababa, Ethiopia

^d Department of Biostatistics and Data Science, School of Public I leal h and Tropical Medicine,

Tulane University, New Orleans, LA 70112 USA

* Kentucky Geological Survey, University of Kentucky, KY. UCA.

*Corresponding author: Tewodros Godebo (tgodebo@tuic.ne edu)

Abstract

Fluoride (F⁻) exposure in drinking water may ead to reduced cognitive function among children; however, findings largely remain inconclusive. 'n this pilot study, we examined associations between a range of chronic F⁻exposures (low to 1.3h: 0.4 to 15.5mg/L) in drinking water and cognition in school-aged children (5-14 years) in rule. Euriopia. Fluoride exposure was determined from samples of community-based drinking water velue and urine. Cognitive performance was measured using: 1) assessments of ability to draw familiar objects (donkey, house, and person), and 2) a validated Cambridge Neuropsychologic. 'Test Automated Battery's (CANTAB) Paired Associate Learning (PAL), which examines vorking memory and new learning and is closely associated with hippocampus function of the brain. Associations between F⁻ and cognitive outcomes were evaluated using regression analysis, adjusting for demographic, health status, and other covariates. The median (range) of water and urine F^- levels was 7.6 (0.4-15.5mg/L) and 5.3 (0.5-15.7mg/L), respectively; these measures were strongly correlated (r=0.74), indicating that water is the primary source of F⁻ exposure. Fluoride in drinking water was negatively associated with cognitive function, measured by both drawing and CANTAB test performance. Inverse relationships were also found between F^- and drawing objects task scores, after adjusting for covariates (p<0.05). Further analysis using CANTAB PAL tasks in the children confirmed that F⁻level in drinking water was positively associated with the number of errors made by children (p<0.01), also after adjusting for covariates (p<0.05). This association between water F⁻ and total errors made became markedly stronger as

PAL task difficulty increased. Fluoride exposure was also inversely associated with other PAL tasks—the number of patterns reached, first attempt memory score and mean errors to success. These findings provide supportive evidence that high F^- exposures may be associated with cognitive deficits in children. Additional well-designed studies are critically needed to establish the neurotoxicity of F^- in children and adults exposed to both low levels known to protect dental decay, as well as excess F^- levels in drinking water.

Key words: Fluoride in drinking water, drawing performance, CANTAB cognitive tests, Children, Ethiopian Rift Valley

1. Introduction

Worldwide, millions of people are affected by fluorosis due to the consumption of drinking water containing levels of fluoride (F^-) that exceed the WHO recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). An optimal amount of F^- (0.7-1mg/L) is well-recommended level of 1.5 mg/L (WHO 2006). In text caries (O'Mullane et al. 2016; Medjedovic et al. 2/J15 U.3. DHHS, 2015; US CDC 1999), but excessive intake of F^- from sources such as wate: food and F^- -containing dental products can lead to dental and skeletal fluorosis (Rango et al., $2C \ge 2$, 2017; Ayoob and Gupta, 2006). In recent years, F^- exposure has received additional soluting due to findings linking F^- exposure with potential cognitive effects, such as a reduced intelligence quotient (IQ) in children (Goodman et al., 2022; Choi et al. 2012; Grandjean, 2019; Grandjean and Landrigan 2014; Tang et al. 2008; U.S. NRC, 2006); however, other studies have not found similar associations (Aggeborn and Oehman, 2021; Barberio et al., 2017; Broadbent et al., 2015).

Experimental studies on rodents provide one source of evidence for potential effects of F^- on the brain. These studies have shown that F^- crosses the placenta and blood brain barriers (Sharma et al. 2017; ATSDR 2003; Ron et al., 1986) and F^- related histopathological changes in various brain regions, including the hippocampus (critical for learning and memory), the prefrontal cortex (executive function), and the cerebellum (motor control) (Liu et al., 2018, Ge et al., 2018, Lee et al., 2016, Jiang et al., 2014, Whitford et al., 2009, Shivarajashankara et al., 2002; Mullenix et al., 1995).

Fluoride can also impair the function of myelin and neurotransmitters, increase lipid peroxidation, and inhibit several key neuronal enzymes, suggesting potential direct impairments to brain function (Niu et al., 2018; Shivarajashankara et al., 2002). The neurotoxic effect of high levels of F^- exposure is frequently referenced to the hippocampus, which is involved in learning, memory, and attention (Bittencourt et al., 2023; Grandjean, 2019 and reference therein; Mullenix et al. 1995; Pereira et al., 2011; Valdez-Jiménez et al., 2011, Bhatnagar et al., 2002). Given that F^- readily crosses the placenta (Sharma et al. 2017; ATSDR 2003), high exposures have been shown to damage the developing brain in utero, leading to permanent long-term effects (rian et al. 2016; Grandjean and Landrigan 2006). This is particularly relevant in F^- endemic ru al regions that have limited alternative water sources and where exposures are potentially chronic and long term, spanning from conception to adulthood. Several epidemiological studies from Asian countries, for example, have shown an association between higher drinking water exposure is for an et al., 2014; Choi et al. 2012; Ayoob and Gupta, 2006), as well as other cognitive effects on children such as inattentive behaviors (Bashash et al., 2007, 2018).

The Main Ethiopian Rift (MFR) '/alley is one of the best-known endemic regions for chronic exposures to low to elecated F^- in community-based groundwater sources. Extensive studies in the region have documented the adverse health effects—dental and skeletal fluorosis—of chronic exposures to elevated F^- (e.g., Rango et al., 2020, 2013, 2012). Most children in the region were born and raised in a single location, in- and out-migration in the region is low, and F^- exposures therefore typically occur over a lifetime starting at conception, but varying considerably in concentration, depending on the water source being used. Prior research in the Rift Valley has not examined effects of chronic F^- exposures on children's cognition.

To address this knowledge gap, we investigated the relationship between exposures to F⁻ from drinking water and children's performance in several cognitive tasks. The first such task involved

drawing of familiar objects and animals, which are considered valid and reliable instruments that correlate with general intelligence in children (Yong 2015; Imuta et al. 2013; Ebersbach and Hagedorn 2011; Brooks 2009; Van Niekerk, 1999; Reynolds 1978). For example, research using the Wechsler Intelligence Scale for Children has shown a moderate correlation between drawing ability and intelligence (Imuta et al., 2013; Reynolds and Hickman 2004; Abell et al., 1996; 2001). Ebersbach and Hagedorn (2011) showed that spatial drawing ability was positively related to cognitive flexibility in 7 to 11 years—old children (r=0.35). Successful drawing requires cognitive flexibility, which develops only gradually with age (Jolley 260c). Second, we examined the association of F⁻ exposure in drinking water with scores on the s'andardized cognitive tasks of the Cambridge Neuropsychological Test Automated Battery (CANTAB®, Cambridge Cognition Ltd., UK) of the Paired Associates Learning (PAL). In doing so, the stindy also provides an assessment of the feasibility of administering the CANTAB neurod strain drawing, for which the hippocampus of the brain region is critical.

In this study, we enrolled children e mosed to naturally occurring and wide-ranging drinking water concentrations of F^- (0.41 to 15.5 mg/L) since conception. We tested the hypothesis that chronic F^- exposure in drinking where is associated with a child's ability to draw familiar objects of varying difficulty (a donkey, a house and a person) and cognitive performance using a standardized CANTAB task. The children residing in sample communities come from a homogeneous rural population that generally engages in farming for its livelihood and has common living conditions, culture, and diet. Such similarity within a study population is rare in epidemiological studies, and dissimilar to the situation in studies in industrialized countries, where socioeconomic conditions vary substantially, and populations face much lower and less-variable F^- exposures. Thus, this location presents an ideal setting to study the health effects of wide range of chronic F^- exposures.

2. Materials and methods

2.1. Study population

We enrolled 74 MER Valley children, aged 5–14 years, in this cross-sectional study in two sampling periods in 2020 and 2021. In previous work, we mapped the distribution of F⁻ levels in several drinking water sources, including community-based wells in use in the study area. This work revealed that elevated levels of F⁻ due to deposition of volcanic *equile* sediments enriched with this element are widespread (Rango et al., 2010, 2012, 2013). Working from the prior mapping, 8 communities were selected to cover a wide range of F⁻ levels in drinking water sources that spanned from 0.4 to 15.5 mg/L (Rango et al. 2012, 2013, 2019). All chrolled children were confirmed to have been born and raised in sample villages that used these respective wells, and thus were known to have been chronically exposed to relatively stable F⁻ concentrations since conception. We assume that current concentrations are a reliable proxy for "historical" concentrations within the study area, which is consistent with our understancing of the hydrogeochemistry of the region and the consistent use of these sources (which have been in place for 15-65 years across the sample villages) in each study community.

Notably, the population c. these rural farming communities is characterized by similar genetic origin (i.e., ethnicity), education, dietary patterns, cultural and social values, livelihoods activities, and many behaviors (e.g., low rates of smoking, and rare if any use of toothpaste), but vary by F⁻ exposure. The relative uniformity of these parameters helps to minimize the risk of confounding of F⁻ exposures by other factors that influence health. Study participants live in households that are engaged in cereal-based agriculture as their primary livelihood activity and are generally low in income and wealth. We recruited children at community centers that are typically located near each community well. Study inclusion criteria were: Consent to participate (obtained from both parents

and children), permanent residence in the community, age between 5 and 14 years, and duration of residency that was at least as long as the age of the sampled well from which drinking water was being consumed. In each community, though we did not explicitly stratify by age and sex, we did attempt to select individuals to obtain a relatively good distribution of different ages and sex.

Survey data were then collected with each enrolled respondent, to record sex, age, place of birth, exposure to smoking, toothpaste usage, approximate daily water intake, and anemic appearance. For the latter, a trained neurologist looked for clinical signs c^{+} anemia, specifically pallor on conjunctiva, brittle or spoon-shaped fingernails, tongue rednets and swelling, and presence of headache symptoms. The recorded estimate of daily water intake, (in liters, L), weight (kg) and the water F⁻ concentrations (mg/L) were used to calculate the daily F⁻ intake dose (mg/kg/day), as the product of water F⁻ concentration (mg/L) and water intake (L)), divided by the body weight of each child. We also recorded weight and height using in electronic scale for weight and measuring tapes for height. These measures allowed calculation of the body mass index (BMI) for each study participant (BMI: weight(kg)/height(m²)) as c oroxy for nutrition status.

The study received ethical approve from the Institutional Review Board (IRB) at Tulane University (Protocol No. 2018-043) and locally from the National Research Ethics Review Committee (NRERC; reference no. MoSHE/144/1003/19). All participants provided consent, and parents/guardians gave permission for children to participate in addition to children giving their own assent.

2.2. Measurements of exposure

2.2.1. Sampling of drinking water and urine: We collected a total of 68 urine samples (24-hour urine (n=46) and spot urine (n=22) samples) from children residing in 8 community-based wells (**Fig S1**). These 8 wells had F^- concentrations ranging between 0.44 and 15.5 mg/L. All sample collection materials for water and urine were pre-cleaned with 1N HNO₃ and 1N HCl and rinsed

three times with deionized water. Water samples were then filtered in the field directly into 60 mL polyethylene bottles using luer lock syringes and 0.45µm Mixed Cellulose Ester membrane filters. Urine samples were collected in disposable plastic urine collection containers with a closing cap (each with a capacity of 1 gallon). The volume of each collected urine sample was registered, and the sample was immediately transferred into a 60 mL polyethylene bottle. Participants were shown how to avoid contamination and instructed to carefully collect urine samples. Water and urine samples were then kept in a zero-degree freezer, and properly pricked, stored, and transported to the lab at Tulane University, USA.

2.2.2. Fluoride, arsenic, and lead analysis in drinking many and urine

Fluoride content in water and urine was determined using the lon Selective Electrode (ISE), buffering the standards and samples using or al volume ratios with a total ionic strength adjustment buffer (TISAB II). The water and urine sample concentrations of As and Pb, which are known to also affect cognition, were also measured using an Inductively Coupled Plasma–Mass Spectrometer (Agilent 7900 ICP-MS) at furane lab. The recovery for F⁻, As, and Pb in samples with respect to the NIST SRM 2668 low standard was between 90 and 110%. In a recent and related biomonitoring study (Rango et al. 2019), we observed a significant positive correlation between F⁻ and As in drinking water and urine, highlighting the role of drinking water as the main route of exposure to both of these elements. Importantly, two (5.2 and 15.5mg/L of F⁻) of the eight community wells have only been used intermittently in the most recent period, because of well pumping malfunctions that were also observed during sampling. Given that urinary levels of these elements only reflect recent exposures that are not from these sources, we exclude them from the analysis of associations between urinary F⁻ and cognition.

2.2.3. Urine correction for urine dilution

Analyses of F⁻, As, and Pb concentrations in urine samples with a volume \leq 300mL (n=22), which can be considered spot samples rather than 24-hour samples, were adjusted for specific gravity (SG) in order to account for variations in urine dilution. This enhances comparability of results from these two types of urine samples. Strong correlations (*r*=0.78, p<0.0001) have been reported between SG-adjusted spot urine sample and fluoride in a 24-h urine sample (e.g., Zohouri et al., 2006), indicating that adjustment for urinary dilution approximates a 24-h biomarker. To apply this adjustment, specific gravity was measured using a handheld refractometer (National Instrument Company, Inc., Baltimore, MD) that was calibrated with deionized worter before each measurement. The refractometer prism head was rinsed in deionized water after each reading. Urinary F⁻ levels were normalized for dilution by SG adjustment using the follow ing formula (MacPherson et al., 2018; Hauser et al., 2004).

 $F_{SG} = F \times [(SG_M - 1)/(SG - 1)]$

Where: F_{SG} is the SG-corrected F⁻concentration (mg/L)

F is the observed F⁻ concentration (.mg/L), and

SG_M is the median specific $c_{1,CVI}$, or the study cohort (SG_M=1.012)

2.3. Measurements of out conces

2.3.1. Children's drawing asks: A total of 68 (37 males and 31 females) from the 74 children were enrolled and asked to participate in three drawing tasks of common objects that children readily encounter or experience in the study area, though their reproduction varies in complexity: a house, a person, and a donkey. They were provided with a pencil, rubber eraser, drawing pad, table, chair and allowed as much time to draw as they needed to complete their drawings, but no instruction or support was provided other than the name of the items to be produced. All children submitted drawings of each object, such that a total of 204 drawings were collected and scored. Most children took approximately 20-30 minutes to finish all three drawings. We developed scoring criteria based

on the completeness of each object such that a point was given for each part correctly drawn (**Table 1**) Each object also received an additional score for overall appearance ranging from 0 to 4 (bad (0), poor (1), fair (2), good (3), very good (4)). Similar figure drawing criteria were used in other studies to assess child cognition (Williams et al., 2006, Imuta et al., 2013; Panesi and Morra 2016). All drawings were independently scored by two examiners who were blinded to the F⁻ concentrations in well water; the inter-rater reliability was assessed and showed a strong correlation (r > 0.94).

2.3.2. Cambridge neuropsychological test automated battery (cr NrAB)

To further examine the association between F^- in water and cognitive outcomes, additional standardized tests were conducted using the Cambridge Neuropsychological Test Automated Battery (CANTAB®, Cambridge Cognition Ltd., UK). For this study, we selected one of the CANTAB tests, Paired Associate Leaning (PAL), which is new to ensitive to spatial memory and learning and linked to the medial temporal lobe (e.g., hi, oor ampus). This is the brain region most thought to be affected by F⁻ toxicity (Mullenix et al. 1995). The test administrator was trained in CANTAB, described the CANTAB instructions and demonstrated to each child how the iPad touch screen works using local language (Amharic or Oromegna). CANTAB is language and culture neutral and is a computerized test administered on a touch-screen interface that requires very little language comprehension, making it suitable for use with children (Fray and Robbins, 1996; Luciana and Nelson, 2002). Prior to in: dating the main tests, a prescreening test with the Motor Screening Task (MOT) was administered to introduce participants to CANTAB and provide a general assessment of sensorimotor, vision, movement, or comprehension difficulties. For the MOT task, a series of crosses that appeared in random locations on the screen was presented to the child. The examiner first demonstrated the correct touching procedure using the forefinger of the dominant hand to touch the cross, and the child then completed the test. The CANTAB PAL task also begins with two rounds of practice sessions. The examiner performed the first practice round, and the participant performed

the second. The task then moved to the assessment phase, which had four levels: 2, 4, 6 and 8 stimuli over the trials. No enrolled children had difficulties completing the tasks.

The PAL test assesses visual memory and new learning that depends on spatial planning ability. In this test, boxes are opened in random order on the screen to reveal their contents. The patterns inside the boxes are then displayed in the middle of the screen one at a time, and the participants must touch the box in which the pattern was originally located. The level of difficulty increases during the test, with 2-, 4-, 6-, and 8-pattern stages. The primary outcome, include the *total errors adjusted* (accounting for the number of trials completed), where a lower number is better, and the *number of patterns reached*, where a higher score is better (**Table S1**).

2.3.3. Statistical Analysis: Demographic, anthropomeulic. F⁻ concentrations, drawing scores and PAL measures (errors/stages completed) were frst described by their quartiles and means \pm standard deviation (**Table 2 and 3**). The F⁻ level in vater was analyzed as a continuous variable and categorized into three F⁻ exposure groups (in n g/L) (Group 1; reference low F⁻ group): <3, Group 2: >3-8, and Group 3: >8-15.5mg/L). Completions of means in the different F⁻ exposure groups, and by sex were carried out with one-weiv A, 'OVA. We then utilized a linear regression model to examine the associations between F⁻ exposure and the drawing scores for the three objects (a donkey, a person, a house), and CA'(1)' B PAL tasks, adjusting for As and Pb in drinking water, sex, grade levels, BMI, and anemic a, pearance, and tested water F⁻ and PAL difficulty interactions on PAL outcomes. Results are presented as regression coefficients β with their 95% confidence intervals (95% C/s), allowing for clustering of standard errors at the community level owing to the within-community correlation of exposures as well as other independent variables. Study hypotheses were evaluated at the 5% level of significance. All analyses, summaries, and graphs were performed using the Statistical Analysis System (SAS 9.4, SAS Institute, Cary, NC, and GraphPad Prism 9.3.1).

3. Results

3.1. General characteristics: The average age of children in the sample was 10 years; approximately half (54.4%; n = 37) of these were male (Table 2). Based on the WHO (2000) classification of BMI, most children would be categorized as underweight with the 75th percentile falling below 16.2 kg/m², and a mean BMI of 15.4±1.57kg/m². The F⁻ concentrations in the groundwater and urine samples ranged between 0.41 and 15.5mg/L and 0.54 and 15.7mg/L, respectively. All the water samples except that with 0.44mg/L exceeded the 1.5mg/L standard limit in drinking water (WHO 2017). The interguartile ranges of estimated daily F⁻ intake per day, F⁻ intake per body weight per day, and urinary F⁻ concentration were 3.22-3.59 mg/day, 0.124-0.35mg/kg bw/day and 4.02–4.65mg/L, respectively. Most of the children (8.2%; n=60) therefore ingested an estimated daily amount of F⁻ that exceeds the U.S. EPA: No-Observed-Adverse-Effects-Level (NOAEL) value of 0.06 mg/kg/day value for F⁻ (US EPA 20.2). Educational and health services, and dietary patterns were generally homogeneous collisis sample areas. Young children are usually breast fed for up to 2 years, then weaned on's household foods, and the diet is primarily cerealbased (maize, wheat, teff), with mean consumption being rare (Rango et al. 2012). Arsenic concentrations in groundwater rangeo ketween 0.92 and 22µg/L, and 2 of the 8 wells (11.8 and 22µg/L) exceeded the established subnards of 10µg/L for As concentration in drinking water (WHO 2017). The Pb concentrations in the groundwater were all below 1µg/L, and none of the water samples exceeded the 10 pg/L limit established for Pb in drinking water (US EPA 2001, WHO 2017).

3.1. Association of F^- As, and Pb in drinking water and urine: We observed strong correlation between F^- in drinking water and children's urine (r=0.74, *p*<0.001). This strongly suggests that drinking water is the main source of F^- for the children. Moderate correlations were observed between As and Pb in drinking water and children's urine—(r=0.33, *p*=0.023) and (r=0.28, *p*=0.054), respectively. A positive and moderate association was also found between F^- and As in drinking water (r=0.23, *p*=0.057). Conversely, a negative association was found between F^- and Pb in the drinking water samples (r=-0.61, *p*<0.001). The unique geochemistry of groundwater in the region

(i.e., the alkaline pH>8, and oxidizing aquifer conditions) limits the occurrence of Pb and other toxic heavy metals such as Cd and Ni, and results in a very low concentration of these elements in groundwater (Rango et al., 2019, 2013). Other than As, the concentrations of heavy metals are generally low in the Rift groundwater and expected to have minimal role on cognition.

3.2. Association of F⁻ in drinking water with drawing performance: The distribution of the scores for figure drawing (based on completed parts and overall appearance as shown in Table 1), in the three F⁻ exposure groups were as follows (Table 3): 1) <3mg L communities: 6.2 (donkey), 8.6 (person) and 6.1 (house), 2) >3-8mg/L communities: 4.6 (donkey), c 9 (person), 4.9 (house), and 3) >8-15.5mg/L communities: 4.0 (donkey), 7.3 (person), and 4.9 (house).

Regression analyses confirm the inverse associations be ween F^- concentrations in water and children's drawing scores (**Fig 1**). In continuous analysis (**Table 4**), the correlation between F^- in drinking water and the scores for drawings of a donkey is statistically significant with a β of -0.2 points (95% *Cl*: -0.37, -0.03) that explains *i* 1.% of the total variance in scores (**Table 4A**, **Fig 1A**). After adjusting for other factors (sex, chickren's grade levels, BMI, As and Pb in drinking water, anemic appearance), this association meakens slightly but remains significant. Representative children's drawings of a donkey for children in different F^- exposure groups are shown in **Fig 2**. Similarly, inverse relationships were found between water concentrations of F^- and drawing scores for a house (**Table 4B**, *i*: **q : B**, **Fig S2**), and person (**Table 4C**), but these relationships were not statistically significant. Fluoride in drinking water accounted for a relatively smaller percentage of the variance in scores for these two tasks-2.4% and 1.5%, respectively.

Adding scores for all three objects, a significant inverse relationship was found between F^- in water and drawing scores (**Fig 1C**). In a sensitivity analysis, with grade level replaced by age among the covariates, the association follows the same trend with a mildly lower effect size (**Table S3**).

In unadjusted categorical analysis (**Table 5**), the F⁻ water concentration category indicators explain 7.0 and 5% of the variability in donkey and house drawing scores. While all object drawing scores

decrease with higher F⁻ concentrations (**Table 3**), the drawing scores for the donkey task were significantly lower in the highest exposure Group 3 (>8-15.5 mg/L), compared to the lowest reference F⁻ exposure Group 1 (<3 mg/L). The adjusted β for this comparison is -3.56 points (95% *CI*: -6.62, -0.48). In contrast, though scores were also lower in Group 2 (>3-8mg/L), the association was not significant. Similarly, for the person and house drawing tasks, children in the two higher exposure groups (>3-8mg/L, and >8-15.5mg/L) had lower scores than those in the reference group, but this relationship was not statistically significant. Across ⁻⁻ exposure groups, males had a better drawing ability (particularly for the donkey and person (a_c/s_f) than females, though this association between scores and sex did not reach statistical significant significance (**Fig S3**).

3.3. Association of F^- in urine with drawing performance: We observed inverse associations between the F^- concentrations in urine and the drawing ability of children (a donkey, a house, and a person). Children's scores for the donkey traving again showed the strongest inverse correlation with F^- in urine (p=0.12) with a β of -0.15 and 95% *CI* (-0.43, 0.05) that explains 5% of the variance (**Fig 3**). Inverse relationships are also observed between urinary F^- and drawing scores for a person ($r^2=0.03$, p=0.24) and a house ($r^2=0.01$, p=0.49), similarly to the drinking water analysis; however, these relationships are not significant.

3.4. Association between exposure to F⁻ and children's CANTAB PAL tasks

Out of the six PAL tasks that children completed (**Table S1**), scores for four (PALTEA, PALNPR, PALMETS, and PALFAMS) are associated with F^- concentrations in drinking water (**Table 6**). Increased F^- in drinking water is significantly associated with the number of PAL total errors adjusted (PALTEA) made by the children (β =1.2, 95%CI: 0.32, 2.1; **Fig 4A**). Fluoride in drinking water is also inversely correlated with the PAL number of patterns reached (PALNPR) (β =-0.1, 95%CI: -0.19, -0.01); **Fig 4B**) and the number of times the correct box is chosen on first attempt (PALFAMS) when

recalling pattern location (β =-0.21 95%CI: -0.42, 0.01); *p*=0.06). Finally, the mean number of attempts needed to successfully complete a stage (PALMETS) is also negatively correlated with F⁻ concentration (β =-0.075, 95%CI: 0.16, 0.007); *p*=0.07). When controlling for covariates (children's grade level, BMI, As and Pb in drinking water, anemic appearance), however, only the PATEA association with F⁻ concentration remains significant. In a sensitivity analysis, with grade level replaced by age among the covariates, the association follows the same trend with a mildly lower effect size (**Table S4**). Similar trends were observed in unadjusted and adjusted relationships between F⁻ in urine and CANTAB PAL tasks for PALTEA, PALN' α , and PALFAMS. In unadjusted and adjusted categorical analysis (**Table S2**), significant differences were found between the lowest (<3mg/L) and highest (<8-15.5mg/L) drinking water concration groups, for the PALTEA, PALNPR and PALFAMS tasks.

3.5. Association between exposure to F⁻ and the difficulty of CANTAB PALTEA tasks

Fluoride concentration in drinking water mat, have larger effects as the level of task difficulty (i.e., the number of boxes) increases. The highest average number of errors by the children were measured in the most difficult task (8-box) among those children exposed to the highest F^- in water (i.e., >8-15.5mg/L) (**Fig 5**). The lowest number of errors were made for the lowest task difficulty (2-box) among those exposed to <3n.q/L of F^- . There was no statistically significant interaction between F^- in drinking water and the task difficulty (number of PALTEA boxes) on the total errors made by the children (**Fig 5**).

4. Discussion and conclusion

In this study, we assessed the association between chronic exposure to naturally-occurring F^- in drinking water and cognitive function in school-aged children, as measured using two distinct types of assessments: a simple drawing task with familiar objects, and the CANTAB PAL tests. The sample was recruited from 8 communities primary exposed to chronic F^- ranging from 0.41 to 15.5mg/L in the MER. These communities have relatively homogenous populations with similar lifestyles and stable residency, but the residents of different villages use community-based drinking

water sources that vary in their F⁻ levels. We hypothesized that measures of cognitive performance would decline with exposure to elevated F^- concentrations. Accordingly, we found adverse associations of F⁻ exposures in drinking water with children's drawing and CANTAB task performance. The strongest and most significant negative impacts were observed for the more challenging drawing task—a donkey (Fig 1A). It is observed that children struggled more when drawing a donkey than a house or a person, which may be indicative of a greater challenge accessing working memory for this task. In contrast, children appeared to have an easier time drawing a person or a house, and associations between drawing performance and F⁻ exposures were correspondingly weaker. Consistent with the negative esso, jations between drawing skill and F⁻ exposure, children drinking from wells in communities with higher F⁻ levels performed worse in CANTAB PAL tasks that are used to test new learning and memory, and especially the PAL total errors adjusted measure. It was also observed that 'nother F⁻ levels were related to higher deficits in the more difficult PALTEA tasks (i.e., increasing number of boxes from 2 to 8) (Fig 5). The PAL test targets hippocampal function by measuring visual memory and new learning (Barnett et al., 2016; de Rover et al., 2011). A study by Choi et al (2015) found that measured working memory using the Wechsler Intelligence Scale for Children-Revised (WISC-IV) in children was negatively associated with dental fluorosis (a marker of arly life F⁻ exposure during critical periods of tooth development, the first 8 years) and V(ec.)sle is total and backward digit span tests. Goodman et al., (2022) also reported that visual-spatic¹ and perceptual reasoning abilities may be more impacted by F⁻ exposure as compared to verbal abilities.

In previous related studies, drawing (e.g., a person) has often been used as a nonverbal screening measure of cognitive ability that may indicate visual sensory input and neuromuscular output (Kamphaus and Pleiss 1991; Kamphaus et al. 1993; Imuta et al. 2013; Reynolds and Hickman 2004; Abell et al., 1996; 2001). A study by Panesi and Morra (2016) assessed dog drawing in relation to executive function and working memory and found that these two parameters jointly accounted for 58.3% of the variance of dog drawing skill. Moreover, working memory individually accounted for the

largest variance (15.4%), whereas executive function accounted for 4.4%. The interaction of these two predictors was then responsible for the remaining 38.5% of this joint variance. Evidence of the role of working memory and executive function in drawing flexibility was also reported by Morra 2005 and Barlow et al. 2003. Moreover, environmental factors can impede these aspects of cognition and drawing. For example, a study by Guillette et al. (1998) observed impairments in memory, social interaction, creativity, drawing ability, and motor skills in a population of Mexican children exposed to pesticides relative to a comparable group living in an unexposed crea. Most dramatically, pesticide exposed children scored more poorly in a "draw a person" task, with h may indicate lower cognitive ability or poor visuomotor coordination.

In our study, grade level (or age) was positively associated with drawing ability, which is also consistent with prior literature (Cox 1993; Panesi and Mr. ra 2016). Owing to our sampling approach, however, which aimed to balance sex and age with a rod across communities, the children in each community are similar, such that sex- a d age-related effects cannot explain the variation in observed outcomes across communities. Other possible confounders include As and Pb, which are known to be neurotoxic contaminants, or these were found to be at low levels in drinking water from the sample communities and in particles of children's urine. The concentrations of As and Pb in drinking water ranged between 0.92 and 21.9µg/L (mean:7.3±6.83µg/L), and 0.001 to 0.73µg/L (mean: 0.23±0.27µg/L), respectively. Anemic appearance, as diagnosed from clinical signs of anemia (e.g., pallor on ccr, unctiva), was observed in 45.5% of the children, and is known to impair motor and mental development in infants, children, and adolescents (Burden et al., 2007; Lazoff 2007, Lam et al., 2017). In regression analysis the association of anemic appearance, and As and Pb in water and urine did not significantly correlate with performance measures, however, this exploratory study relied on a relatively small sample and used cross sectional data to proxy for longterm exposure. In addition, the purposeful recruitment of children to obtain representative age and sex distributions around specific community wells limits the representativeness of the sample. As a result, the study may not be viewed as providing a definitive analysis of F^{\neg} 's neurotoxicity in children.

Nonetheless, the similar sociodemographic and lifestyles in these communities minimizes the risk of confounding by variables that may be correlated with exposures and cognitive performance measures. An important additional limitation was the small number of sample communities and wells. In particular, when adjusting the standard errors for clustering within wells/communities, the statistical significance of the association between water F^- and the PALTEA task performance scores was reduced from *p*=0.034 to *p*=0.09, emphasizing the need to increase the number of wells and study participants to obtain greater statistical power.

Other limitations include a lack of control of parental variables such as maternal age, educational level of parent, socio-economic status, and assessment of chemical mixture models for better exposure and effect characterization other potential neucrotesticants (e.g., As, Pb), and elemental deficiencies such as iodine and iron that may modify cognition (Lam et al., 2017). For some urine biomarker measures that were collected as spot camples, we accounted for dilution using urine specific gravity, to reflect actual F⁻ exposure from drinking water.

5. Conclusion: Our findings suggest that here are cognitive impairments among children exposed to higher F⁻ concentrations, evaluated using figure drawing performance and validated CANTAB cognitive tools. This study also successfully demonstrated the use of language and culture neutral CANTAB testing in a rural thic pian sample of children for the first time. Thus, CANTAB can feasibly be administered in this and other similar rural African contexts (as also shown by Chetty-Mhlanga et al., 2022, 2018; Nkhoma et al., 2013). While this exploratory study adds evidence and concern about the potential neurotoxicity of elevated F⁻ exposure, more studies are critically needed to better establish neurodevelopmental impacts of a range of F⁻ exposures from gestation to adulthood, using rigorous study designs and advanced methodologies including mixture models for exposure and effect characterization. Such studies would help provide concrete evidence to inform leaders and policy makers on the need for effective approaches to mitigate environmental exposures to F⁻, including in F⁻ endemic geographic settings such as the study areas where alternative water sources

are limited, or to establish the threshold levels at which such exposures become toxic, and specifically, inform the growing controversy over the safety of water fluoridation.

Acknowledgements: We thank all the children and parents who participated in this study and for their valuable time, and the local water bureaus for their help in recruiting them as well as guiding us during the field work. The content expressed in this paper is the responsibility of the authors and does not necessarily reflect the official views of the funding agency. We also acknowledge the funding sources from the PI's Internal Institutional Start-up Fund, and National Institute of Environmental Health Sciences (NIEHS's) career development grant (K99/R00 ES023472) that made the research possible.

Data Availability: Data will be made available on reasonable request.

Conflict of interest: The authors declare they have the actual or potential competing interests.

Authors contributions: Tewodros Rango Jor ebc. Design of study concept, conducted the field work, and administered the figure drawing and CANTAB cognitive assessment, data analysis and interpretation, and wrote the manuscript

Marc Jeuland, Arti Shankar, and Amy *W*olfe: Statistical data analysis, and critical revision of the manuscript for important intellectual content.

Nati Phan: Figure drawing a segment, and critical revision of the manuscript for important intellectual content.

Tekle-Haimanot, and Einiyam Ayele: Field work assistance, and revision of the manuscript for important intellectual conter :.

References

- Ayoob, S., & Gupta, A.K. (2006). Fluoride in drinking water: A review on the status and stress effects. Critical Reviews in Environmental Science and Technolog, مج(6), 433-487.
- Aggeborn L, Oehman M. (2021). The effects of fluoride in drinking vater. J. Polit. Econ. 129, 465–491. https://doi.org/10.1086/711915.
- Abell SC, Von Briesen PD, Watz LS (1996). Intellectual evaluations of children using human figure drawings: An empirical investigation of two methods. *Journal of Clinical Psychology*, *52*(1): 67-74.
- Abell SC, Wood, W, Liebman, SJ (2001). Children's human figure drawings as measures of intelligence: The comparative validity of three coring systems. Journal of Psychoeducational Assessment, 19(3), 204-215.
- Agency for Toxic Substances and Disease Registry (ATSDR). (2003). Toxicological Profile for Fluorides, Hydrogen Fluoride, and Fluorine. Retrieved from https://www.atsdr.cdc.gov/ToxProfiles/ asp?id=212&tid=38
- Aikman KG, Belter RW, Finch Jr, AJ (195.). Human figure drawings: Validity in assessing intellectual level and academic achievement. Journal of Clinical Psychology, 48(1), 114-120.
- Ayoob S, Gupta AK (2006). Fluoride in Urinking Water: A Review on the Status and Stress Effects. Critical Reviews in Environmen I Science and Technology, 36(6), 433-487.
- Bittencourt LO, Dionizio A, Ferreira MKM, Aragão WAB, Cartágenes SC, Puty B, Maia C do SF, Zohoori FV, Buzalaf M.^Ak, Lima RR. (2023). Prolonged exposure to high fluoride levels during adolescence to adu:hood elicits molecular, morphological, and functional impairments in the hippocampus Crincep 13, 11083. https://doi.org/10.1038/s41598-023-38096-8
- Barnett JH, Blackwell AD, Sahakian BJ, & Robbins TW (2016). The Paired Associates Learning (PAL) Test: 30 Years of CANTAB Translational Neuroscience from Laboratory to Bedside in Dementia Research. Current Topics in Behavioral Neurosciences, 28, 449–474. 10.1007/7854_2015_5001.
- Burden MJ, Westerlund AJ, Armony-Sivan R, Nelson, CA, Jacobson SW, Lozoff B, et al. (2007). An event-related potential study of attention and recognition memory in infants with iron-deficiency anemia. Pediatrics 120, e336–e345.
- Barlow CM, Jolley RP, White DG, Galbraith D (2003). Rigidity in children's drawings and its relation with representational change. Journal of Experimental Child Psychology, 86(2), 124-152.
- Bhatnagar M, Rao P, Sushma J, Bhatnagar R. Neurotoxicity of fluoride: neurodegeneration in hippocampus of female mice. Indian J Exp Biol. 2002; 40(5):546–54.
- Bashash M, Marchand M, Hu H, Till C, Martinez-Mier EA, Sanchez BN, ... Téllez-Rojo MM (2018). Prenatal fluoride exposure and attention deficit hyperactivity disorder (ADHD) symptoms in children at 6–12 years of age in Mexico City. Environment International, 121, 658-666.
- Bashash M, Thomas D, Hu H, Martinez-Mier EA, Sanchez BN, Basu N, . . . Hernández-Avila M. (2017). Prenatal Fluoride Exposure and Cognitive Outcomes in Children at 4 and 6-12 Years of Age in Mexico. Environmental health perspectives, 125(9), 097017-097017.

- Barberio AM, Quinonez C, Hosein FS, and McLaren L. (2017). Fluoride exposure and reported learning disability diagnosis among Canadian children: Implications for community water fluoridation. Can J Public Health, 108(3), e229-e239.
- Broadbent JM, Thomson WM, Ramrakha S, Moffitt TE, Zeng J, et al., (2015). Community water fluoridation and intelligence: prospective study in New Zealand. Am J Public Health, 105(1), 72-76.
- Brooks M (2009). Drawing, Visualisation and Young Children's Exploration of "Big Ideas". International Journal of Science Education, 31(3), 319-341.
- Chetty-Mhlanga S, Viglietti P, Röösli M, Dalvie MA. (2022). Maternal drinking behavior and coexposure from smoking during and after pregnancy in relation to the neurocognitive function of school-children in the rural Western Cape. Neurotoxicology;88:36-43. doi: 10.1016/j.neuro.2021.10.010.
- Chetty-Mhlanga, S., Basera, W., Fuhrimann, S., Probst-Hensch, N., Delport, S., Mugari, M., ... & Dalvie, M. A. (2018). A prospective cohort study of sch. ol-going children investigating reproductive and neurobehavioral health effects due to environmental pesticide exposure in the Western Cape, South Africa: study protocol. BMC public health, 18(1), 1-13.
- Choi AL, Sun G, Zhang Y, Grandjean P (2012). Developmer tal ,'uoride neurotoxicity: a systematic review and meta-analysis. Environmental healt perspectives, 120(10), 1362-1368. doi:10.1289/ehp.1104912
- Choi AL, Zhang Y, Sun G, Bellinger DC, Wang K, Yai, X, Li JS, Zheng Q, Fu Y, Grandjean P. Association of lifetime exposure to fluoride and cognitive functions in Chinese children: a pilot study. Neurotoxicol Teratol. 2015 Jan-Feb;4⁻, 3-101. doi: 10.1016/j.ntt.2014.11.001.
- Das K, Monda NK (2016). Dental fluorosis and Cincary fluoride concentration as a reflection of fluoride exposure and its impact on 'Cincarel and BMI of children of Laxmisagar, Simlapal Block of Bankura District, W.B., In via. Environmental Monitoring and Assessment, 188(4), 218.
- de Rover M, Pironti VA, McCabe JA, Acusta-Cabronero J, Arana FS, Morein-Zamir S, Sahakian BJ. (2011). Hippocampal dysfunction, in patients with mild cognitive impairment: a functional neuroimaging study of *e* visuospatial paired associates learning task. Neuropsychologia.;49(7):20(0–20/0.
- Dong L, Yao P, Chen W, Li P, Shi X. (2018) Investigation of dental fluorosis and intelligence levels of children in drinking worter related endemic fluorosis area of Xi'an. Chinese Journal of Epidemiology, 37(1), 45–40.
- Ebersbach M, Hagedorn, F: (2011). The role of cognitive flexibility in the spatial representation of children's drawing's. Journal of Cognition and Development, 12(1), 32-55. doi:10.1080/15246272.2011.539526
- Ge, Y., Chen, L., Yin, Z., Song, X., Ruan, T., Hua, L., ... & Ning, H. (2018). Fluoride-induced alterations of synapse-related proteins in the cerebral cortex of ICR offspring mouse brain. Chemosphere, 201, 874-883.
- Goodman CV, Bashash M, Green R, Song P, Peterson KE, Schnaas L, Mercado-García A, Martínez-Medina S, Hernández-Avila M, Martinez-Mier A, Téllez-Rojo MM, Hu H, Till C. (2022). Domain-specific effects of prenatal fluoride exposure on child IQ at 4, 5, and 6-12 years in the ELEMENT cohort. Environ Res;211:112993. doi: 10.1016/j.envres.2022.112993.
- Grandjean P. (2019). Developmental fluoride neurotoxicity: An updated review. Environmental Health, 18(1), 110.
- Grandjean P, Landrigan, PJ (2014). Neurobehavioural effects of developmental toxicity. The Lancet Neurology, 13(3), 330-338.
- Guillette, EA, Meza MM, Aquilar MG, Soto AD, Garci IE. (1998). An anthropological approach to the evaluation of preschool children exposed to pesticides in Mexico. Environmental health perspectives, 106(6), 347-353.

- Hauser R, Meeker JD, Park S, Silva MJ, Calafat, AM. (2004). Temporal variability of urinary phthalate metabolite levels in men of reproductive age. Environmental health perspectives, 112(17), 1734-1740. doi:10.1289/ehp.7212
- Imuta K, Scarf D, Pharo H, Hayne H (2013). Drawing a close to the use of human figure drawings as a projective measure of intelligence. PloS one, 8(3), e58991-e58991.
- Jiang C, Zhang S, Liu H, Guan Z, Zeng Q, Zhang C, . . . Wang A (2014). Low glucose utilization and neurodegenerative changes caused by sodium fluoride exposure in rat's developmental brain. NeuroMolecular Medicine, 16(1), 94-105.
- Jiang, S., Su, J., Yao, S., Zhang, Y., Cao, F., Wang, F., ... & Xi, S. (2014). Fluoride and arsenic exposure impairs learning and memory and decreases mGluR5 expression in the hippocampus and cortex in rats. PLoS One, 9(4), e96041.
- Jolley R (2008). Children's understanding of the dual nature of pictures. In Drawing and the nonverbal mind: A life-span perspective. (pp. 86-103). New York, NY, US: Cambridge University Press.
- Kamphaus RW (1993). Clinical assessment of children's intelligence: A handbook for professional practice: Allyn & Bacon.
- Kamphaus, RW, Pleiss K. L. (1991). Draw-a-Person techni jue.: Tests in search of a construct. Journal of School Psychology, 29(4), 395-401.
- Karimzade S, Aghaei M, Mahvi A (2014). Investigation of intelligence quotient in 9–12-year-old children exposed to high-and low-drinking water fluc ide in West Azerbaijan Province, Iran. Fluoride, 47(1), 9-14.
- Lam LF, Lawlis TR. Feeding the brain The effacts of micronutrient interventions on cognitive performance among school-aged children: A systematic review of randomized controlled trials. Clin Nutr. 2017 Aug;36(4):1007-1015 doi: 10.1016/j.clnu.2016.06.013.
- Lazoff B, 2007. Iron deficiency and child de elogment. Food Nutr Bull. 28(4):S560-71.
- Lee, J., Han, Y. E., Favorov, O., Tommerdahi, M., Whitsel, B., & Lee, C. J. (2016). Fluoride induces a volume reduction in CA1 hippchampal slices via MAP kinase pathway through volume regulated anion channels. Experimental neurobiology, 25(2), 72.
- Liu F, Ma J, Zhang H, Liu P, Liu YP, Xing E, Dang YH (2014). Fluoride exposure during development affects both cognition and er notion in mice. Physiology & Behavior, 124, 1-7.
- Luciana M, Nelson, CA (2002). Seessment of Neuropsychological Function Through Use of the Cambridge Neuropsychological Testing Automated Battery: Performance in 4-to 12-Year-Old Children. Developmental Neuropsychology, 22(3), 595-624.
- MacPherson S, Arbuckle TE, Fisher M (2018). Adjusting urinary chemical biomarkers for hydration status during programmer. Journal of Exposure Science & Environmental Epidemiology, 28(5), 481-493.
- Malin AJ, Till C (2015). Sposure to fluoridated water and attention deficit hyperactivity disorder prevalence among children and adolescents in the United States: an ecological association. Environmental health: a global access science source, 14, 17-17.
- Morra S (1994). Issues in Working Memory Measurement: Testing for M Capacity. International Journal of Behavioral Development Int J Behav Dev, 17, 143-159.
- Mullenix PJ, Denbesten PK, Schunior A, Kernan WJ (1995). Neurotoxicity of sodium fluoride in rats. Neurotoxicology and Teratology, 17(2), 169-177.
- Fray PJ, Robbins, TW (1996). CANTAB battery: Proposed utility in neurotoxicology. Neurotoxicology and Teratology, 18(4), 499-504.
- Nkhoma, O. W., Duffy, M. E., Davidson, P. W., Cory-Slechta, D. A., McSorley, E. M., Strain, J. J., & O'Brien, G. M. (2013). Nutritional and cognitive status of entry-level primary school children in Zomba, rural Malawi. International journal of food sciences and nutrition, 64(3), 282-291.
- National Toxicology Program (2016). NTP Research Reports. In NTP Research Report on Systematic Literature Review on the Effects of Fluoride on Learning and Memory in Animal Studies: Research Report 1. Research Triangle Park (NC): National Toxicology Program.

- Niu, R., Chen, H., Manthari, R. K., Sun, Z., Wang, J., Zhang, J., & Wang, J. (2018). Effects of fluoride on synapse morphology and myelin damage in mouse hippocampus. Chemosphere, 194, 628-633.
- Pereira M, Dombrowski PA, Losso EM, Chioca LR, Da Cunha C, Andreatini R (2011). Memory Impairment Induced by Sodium Fluoride is Associated with Changes in Brain Monoamine Levels. Neurotoxicity Research, 19(1), 55-62.
- Panesi S, Morra S (2016). Drawing a dog: the role of working memory and executive function. J. Exp. Child Psychol. 152, 1–11.
- O'Mullane DM, Baez RJ, Jones S, Lennon MA, Petersen PE, Rugg-Gunn AJ, Whelton H, Whitford GM. (2016). Fluoride and Oral Health. Community Dent Health. 33(2):69-99.
- Medjedovic E, Medjedovic S, Deljo D, Sukalo A. (2015). Impact of Fluoride on Dental Health Quality." Materia Socio-Medica 27 (6): 395–98.
- Reynolds CR (1978). The McCarthy drawing tests as a group instrument. Contemporary Educational Psychology, 3(2), 169-174.
- Reynolds, C. R., & Hickman, J. A. (2004).Draw-a-person intercetual ability test for children, adolescents, and adults examiner's manual. Austin, TX: Pip-eta.
- Ron M, Singer L, Menczel J, Kidroni G (1986). Fluoride concentration in amniotic fluid and fetal cord and maternal plasma. European Journal of Obstatruce & Gynecology and Reproductive Biology, 21(4), 213-218.
- Rango TG, Petrini R, Stenni B, Bianchini G, Slejko F, Be cal ıva L, Ayenew T (2010). The dynamics of central Main Ethiopian Rift waters: Evidence rour δD, δ¹⁸O and ⁸⁷Sr/⁸⁶Sr ratios. *Applied Geochemistry*, *25*(12), 1860-1871.
- Rango TG, Kravchenko J, Atlaw B, McCornick F Jeuland M, Merola B, Vengosh A (2012). Groundwater quality and its health in point: An assessment of dental fluorosis in rural inhabitants of the Main Ethiopian Rif. *Evivironment International, 43*, 37-47.
- Rango TG, Vengosh A, Dwyer G, Bianchini C (2013). Mobilization of arsenic and other naturally occurring contaminants in ground vater of the Main Ethiopian Rift aquifers. *Water Research*, *47*(15), 5801-5818.
- Rango TG, Vengosh A, Jeuland M, iekle-Haimanot R, Weinthal E, Kravchenko J, Paul C, McCornick P (2014). Fluoride e posure from groundwater as reflected by urinary fluoride and children's dental fluorosis in the Main Ethiopian Rift Valley. *Science of The Total Environment*, 496, 188-197.
- Rango TG, Vengosh A, Jeuland N, Whitford GM, Tekle-Haimanot R. (2017). Biomarkers of chronic fluoride exposure in groundwater in a highly exposed population. Science of the Total Environment, 525. . .11
- Rango TG, Paul CJ, Jeuand M, Tekle-Haimanote R., 2019. Biomonitoring of metals and trace elements in urine of central Ethiopian populations. Int J Hyg Environ Health, 222 (3) 410-418.
- Rango TG, Jeuland M, Tekle-Haimanot R, Shankar A, Alemayehu B, Assefa G, Whitford G, Wolfe, A. (2020). Bone quality in fluoride-exposed populations: A novel application of the ultrasonic method. *Bone Reports*, *12*, 100235.
- Shivarajashankara, Y. M., Shivashankara, A. R., Bhat, P. G., Rao, S. M., & Rao, S. H. (2002). Histological changes in the brain of young fluoride-intoxicated rats. Fluoride, 35(1), 12-21.
- Sharma D, Singh A, Verma K, Paliwal S, Sharma S, Dwivedi J (2017). Fluoride: A review of preclinical and clinical studies. Environmental Toxicology and Pharmacology, 56, 297-313.
- US NRC (2006). Fluoride in Drinking Water: A Scientific Review of EPA's Standards. Washington, DC: The National Academies Press.
- US EPA 2001. The Technical Fact Sheet: Final Rule for Arsenic in Drinking Water contains requirements and compliance dates, the health effects associated with exposure to arsenic, the costs, benefits, and number of systems affected by the rule. Read the Arsenic Rule Technical Fact Sheet. EPA 815-F-00-016, January 2001. https://nepis.epa.gov/Exe/ZyPdf.cgi?Dockey=20001XXE.txt. Assessed July 19, 2023.

- US Centers for Disease Control and Prevention (US CDC). Ten Great Public Health Achievements-United States, 1900–1999. Morb Mortal Wkly Rep. 1999;48:241–3.
- US Environmental Protection Agency (US EPA). (2011). EPA and HHS Announce New Scientific Assessments and Actions on Fluoride: Agencies Working Together to Maintain Benefits of Preventing Tooth Decay while Preventing Excessive Exposure. Retrieved from https://archive.epa.gov/epapages/newsroom_archive/newsreleases/86964af577c37ab285257 811005a8417.html
- US Environmental Protection Agency (U.S. EPA). (2002). Integrated Risk Information System, Fluorine (soluble fluoride, CASRN 7782-41-4). Retrieved from https://cfpub.epa.gov/ncea/iris/iris documents/documents/subst/0053 summary.pdf
- U.S. DHSS-FP [U.S. Department of Health and Human Services Federal Panel on Community Water Fluoridation]. (2015). U.S. Public Health Service Recommendation for Fluoride Concentration in Drinking Water for the Prevention of Dental Caries. Public Health Reports, 130(4), 318-331.
- Valdez Jiménez L, López Guzmán OD, Cervantes Flores ivi, Costilla-Salazar R, Calderón Hernández, J, Alcaraz Contreras Y, Rocha-Amador, D. O (2017). In utero exposure to fluoride and cognitive development delay in infants. Neuro oxicology, 59, 65-70.
- Valdez-Jimenez L, Soria Fregozo C, Miranda Beltran ML, Cuticmez Coronado O, Perez Vega, M. I. (2011). Effects of fluoride on the central nervous system. Neurologia, 26(5), 297-300.
- Williams TO, Fall AM, Eaves, RC, and Woods-Groves, S. (2006). The Reliability of Scores for the Draw-A-Person Intellectual Ability Test for Chi'dren Adolescents, and Adults. Journal of Psychoeducational Assessment, 24(2), 137- 1-4.
- Whitford, G. M., Whitford, J. L., & Hobbs, S. H (2003). Appetitive-based learning in rats: lack of effect of chronic exposure to fluoride. Noticology and teratology, 31(4), 210-215.
- WHO (2000). Obesity: Preventing and Mar aging the Global Epidemic, Report of a WHO Consultation. WHO Technical Report Series No. 894. Geneva, Switzerland: World Health Organization. 252 p. Retrieved from: https://apps.who.int/iris/handle/10665/42330
- WHO (2019). Inadequate or excess flucrico: A major public health concern (Preventing disease through healthy environment; WHO/CED/PHE/EPE/19.4.5). Geneva, Switzerland: Department of Public Health Environmental and Social Determinants of Health, World Health Organization. 8 p. Retrie rec. from: https://www.who.int/publications/i/item/WHO-CED-PHE-EPE-19.4.5
- WHO, 2017. Guidelines for Drinking-Water Quality: Fourth Edition Incorporating the First Addendum. Geneva: World Health Organization; 2017.
- Yan N, Liu Y, Liu S, Cco C Wang F, Wang Z, Xi S (2016). Fluoride-induced neuron apoptosis and expressions of inflammatory factors by activating microglia in rat brain. Mol Neurobiol, 53(7), 4449-4460.
- Yong G (2015) Draw a person as an exploratory medium. Retrieved from http://georgeyonge.net/sites/georgeyonge.net/files/VanNiekerk-Eval ch9.pdf
- Zohouri FV, Swinbank CM, Maguire A, Moynihan PJ. 2006. Is the fluoride/creatinine ratio of a spot urine sample indicative of 24-h urinary fluoride? Community Dent Oral Epidemiol 34(2):130-

138, PMID: 16515677, 10.1111/j.1600-0528.2006.00269.x

 Table 1. Scoring criteria developed to assess the quality of each drawn object.

| Criteria score (parts scale | e + overall appearance) | |
|-----------------------------|-------------------------|----------------------|
| A person (11 scale) | A donkey (8 scale) | A house (6 scale) |
| Two hands (1) | Four legs (1) | Cross on the top (1) |
| Two legs (1) | Head (1) | Roof (1) |

| Head (1) | Ears (1) | Grass cover (1) |
|-----------------------------|-----------------------------|-----------------------------|
| Fingers (hand+foot) (1) | Eyes (1) | Door (1) |
| Ears (1) | Mouth (1) | Two windows (1) |
| Nose (1) | Hair (1) | Wall (1) |
| Eyes (1) | Tail (1) | |
| Mouth (1) | Neck (1) | |
| Hair (1) | | |
| Neck (1) | | |
| Shoulder (1) | | |
| Overall appearance (0 to 4) | Overall appearance (0 to 4) | Overall appearance (0 to 4) |

Table 2. Statistical descriptions of anthropometric values, measured concentrations of F^- , As, and Pb from 8 groundwater wells, demographic, and lifestyle factors of the children who performed the drawing tasks.

| | | | Pe | c, ntil | ,s | | |
|--|----|-------|------------------|---------|------------------|------|-----------|
| | N | Min | 25 th | 50" | 75 th | Max | Mean±SD |
| Anthropometric measures | | | | | | | |
| Age | 68 | 5 | 8 | 10 | 12 | 14 | 10.0±2.44 |
| Weight (kg) | 68 | 16.5 | 2°.9 | 25.3 | 33.1 | 48.2 | 28.2±7.45 |
| Height (m) | 68 | 1.04 | 1.24 | 1.31 | 1.47 | 1.59 | 1.34±0.14 |
| BMI (kg/m²) | 68 | 12 3 | 14.3 | 15.2 | 16.2 | 21.1 | 15.4±1.57 |
| Children's grade level | 68 | C | 1.0 | 1.0 | 3.0 | 6.0 | 1.97±1.44 |
| Biomarkers of F ⁻ Exposures | | | | | | | |
| Water F ⁻ , As, and Pb concentrations | | | | | | | |
| Individuals water intake (liter/day) | 68 | 0.3 | 0.9 | 0.9 | 1.13 | 1.5 | 0.91±0.23 |
| Individuals F ⁻ intake (mg/day) | 68 | 0.123 | 3.22 | 6.48 | 9.59 | 23.3 | 7.24±4.93 |
| Individuals Dose (mg/kg bw/day) | 68 | 0.005 | 0.124 | 0.23 | 0.35 | 1.15 | 0.27±0.21 |
| F ⁻ in community wells (mg/L) | 8 | 0.41 | 3.47 | 7.6 | 10.7 | 15.5 | 7.55±4.79 |
| As in community wells (µg/L) | | 0.92 | 2.57 | 5.41 | 10.8 | 21.9 | 7.3±6.83 |
| Pb in community wells (µg/L) | 8 | 0.001 | 0.01 | 0.1 | 0.44 | 0.73 | 0.23±0.27 |
| Urinary F ⁻ , As, and Pb concentrations | | | | | | | |
| F in urine (n, 1) | 48 | 0.54 | 3.54 | 6.34 | 9.1 | 15.7 | 6.44±4.0 |
| As in urine 'ug/L | 47 | 1.46 | 5.02 | 6.77 | 15.8 | 49.1 | 12±12 |
| Pb in urite (ug) | 47 | 0.2 | 0.61 | 0.95 | 1.23 | 5.76 | 1.05±0.84 |
| Age distribution by F exposul grc ips | | | | | | | |
| -omg/L | 17 | 7 | 8 | 10 | 11.5 | 14 | 10±2.12 |
| > 3-8mg/L | 25 | 5 | 7.5 | 9 | 12 | 13 | 9.9±2.6 |
| >o-15.5mg/L | 26 | 6 | 8 | 10 | 12.3 | 14 | 10.2±2.6 |
| Education levels by F ⁻ exposure groups | | | | | | | |
| <3mg/L | 17 | 0 | 1 | 1 | 3.5 | 6 | 1.88±1.76 |
| >3-8mg/L | 25 | 0 | 1 | 2 | 3 | 5 | 1.84±1.43 |
| >8-15.5mg/L | 26 | 0 | 0 | 3 | 3.25 | 5 | 2.15±1.78 |
| Weight by F ⁻ exposure groups | | | | | | | |
| <3mg/L | 17 | 20.6 | 23.0 | 24.7 | 32.3 | 48.2 | 28.3±7.82 |
| >3-8mg/L | 25 | 16.5 | 22.9 | 25 | 33 | 42.7 | 27.7±6.91 |
| >8-15.5mg/L | 26 | 18.2 | 20.8 | 27.1 | 34.4 | 44.4 | 28.6±7.85 |
| Height by F ⁻ exposure groups | | | | | | | |
| <3mg/L | 17 | 1.22 | 1.27 | 1.31 | 1.41 | 1.56 | 1.35±0.10 |
| >3-8mg/L | 25 | 1.16 | 1.23 | 1.3 | 1.43 | 1.56 | 1.33±0.12 |
| >8-15.5mg/L | 26 | 1.04 | 1.22 | 1.33 | 1.49 | 1.59 | 1.35±0.16 |
| BMI by F ⁻ exposure groups | | | | | | | |
| <3mg/L | 17 | 13 | 17 | 15.4 | 16.3 | 18.5 | 15.4±1.4 |
| >3-8mg/L | 25 | 12.3 | 14.5 | 15.5 | 15.9 | 19 | 15.3±1.4 |

| >8-15.5mg/L | 26 | 13 | 17 | 15.4 | 16.3 | 18.5 | 15.4±1.4 |
|--|-----------|----|----|------|------|------|----------|
| Sex distribution by F ⁻ exposure groups | | | | | | | |
| <3mg/L (n=17): Male/Female | 9/8 | | | | | | |
| >3-8mg/L (n=25): Male/Female | 12/13 | | | | | | |
| >8-15.5mg/L (n=26): Male/Female | 11/15 | | | | | | |
| Lifestyle Factors | n | | | | | | |
| Anemic appearance – present | 33(48.5%) | | | | | | |
| Anemic appearance – absent | 35(51.5%) | | | | | | |
| Current smoking habit- present | 0 (0%) | | | | | | |
| Current smoking habit – absent | 68 (100%) | | | | | | |
| Current toothpaste use – present | 12(17.6%) | | | | | | |
| Current toothpaste use – absent | 56(82.4%) | | | | | | |

Note: The urinary F⁻, As, and Pb concentrations of individuals from two of the eight community wells were excluded because the tested well water was only used intermittently in those settings and may not accurately represent current exposure.

| Table 3. | Statistical | descriptions | of concentratio | ns of | F^- | fron | 8 | groundwater | wells, | and | children's |
|-----------|-------------|--------------|-----------------|--------|-------|------|---|-------------|--------|-----|------------|
| drawing s | scores and | CANTAB PA | L task performa | nce so | core | es. | | | | | |

| | | |) P€ | ercentil | es | | 10.000 |
|---|------|-----|--------------|------------------|------------------|-------|------------|
| | N | Mn | <u>יי</u> 5" | 50 th | 75 th | Max | Mean±SD |
| BIOMARKER OF F ⁻ EFFECT (drawing performance) | | | | | | 1 | |
| Donkey drawing total scores at F exposure groups | | 1 | | | | | |
| 0.41-15.5mg/L | 8 | 0.5 | 1.5 | 4 | 7.75 | 11.5 | 4.76±3.39 |
| <3mg/L | 11 | 1 | 3.5 | 5.5 | 9.75 | 11 | 6.18±3.43 |
| >3-8', y, ". | 1 25 | 0.5 | 1.5 | 4 | 7.5 | 11.5 | 4.64±3.47 |
| >8-15 mg_ | 26 | 0.5 | 1 | 3.25 | 6 | 11 | 3.96±3.10 |
| Person drawing total scores at F ⁻ exposure groups | | | | | | | |
| 0.41-15.5mg/L | 68 | 0.5 | 4.63 | 8 | 11 | 13.5 | 7.88±3.63 |
| 3mg/L | 17 | 0.5 | 5 | 8.5 | 12 | 13.5 | 8.59±3.46 |
| こ。8rng/L | 25 | 0.5 | 5 | 9 | 11.3 | 12.3 | 7.98±3.46 |
| >8 15.5mg/L | 26 | 0.5 | 5 | 8 | 11 | 13.5 | 7.33±3.78 |
| House drawing total scores at F ⁻ exposule groups | | | 1.000 | | | 5.555 | |
| 0.41-15.5mg/L | 68 | 0.5 | 3.63 | 5.5 | 7.0 | 10.0 | 5.2±2.47 |
| <3mg/L | 17 | 1 | 5 | 6.0 | 7.25 | 10 | 6.12±2.28 |
| >3-8mg/L | 25 | 0.5 | 3.75 | 5.0 | 6 | 8.5 | 4.88±2.10 |
| >8-15.5mg/L | 26 | 0.5 | 2.63 | 5.25 | 7 | 9.0 | 4.92±2.83 |
| BIOMARKER OF F EFFECT (C AN AB PAL performance) | | | | | | | |
| a. PALTEA (PAL Total Erro. ~ (A viur ced)) | | | | | | | |
| 0.41-15.5mg/L | 74 | 5 | 18 | 33.5 | 47.5 | 69 | 34.1±18.8 |
| <3mg/L | 20 | 5 | 13 | 18.5 | 34 | 62 | 24.0±16.7 |
| >3-8mg/L | 26 | 9 | 20.8 | 34 | 48 | 68 | 35.7±18.1 |
| >8-15.5mg/L | 28 | 10 | 21.5 | 41 | 56.8 | 69 | 39.7±18.6 |
| PALNPR (PAL Number of Patterns Reached) | | 1.7 | | | | | |
| 0.41-15.5mg/L | 74 | 2 | 6 | 8 | 8 | 8 | 6.6±1.89 |
| <3mg/L | 20 | 4 | 8 | 8 | 8 | 8 | 7.4±1.31 |
| >3-8mg/L | 26 | 2 | 6 | 8 | 8 | 8 | 6.46±2.1 |
| >8-15.5mg/L | 28 | 2 | 4 | 6 | 8 | 8 | 6.14 ±1.96 |
| PALMETS (PAL Mean Errors to Success) | | | | | | | |
| 0.41-15.5mg/L | 69 | 0 | 0 | 2 | 3.5 | 6 | 2.04±1.69 |
| <3mg/L | 20 | 0 | 1.25 | 2 | 4 | 5 | 2.4±1.43 |
| >3-8mg/L | 23 | 0 | 0 | 2 | 4 | 6 | 2.1±1.95 |
| >8-15.5mg/L | 26 | 0 | 0 | 2 | 3 | 5 | 1.73±1.64 |
| PALFAMS (PAL First Attempt Memory Score) | | | | | | | |
| 0.41-15.5mg/L | 74 | 0 | 4.75 | 8 | 12 | 17 | 8.15±4.54 |
| <3mg/L | 20 | 3 | 8.25 | 11.5 | 14 | 15 | 10.5±3.68 |
| >3-8mg/L | 26 | 0 | 4.75 | 7.5 | 11.3 | 17 | 7.46±4.50 |

>8-15.5mg/L 28 0 2.25 7.5 10.8 15 7.11±4.67

Note that PALTEA= PAL Total Errors (Adjusted); PALNPR= PAL Number of Patterns Reached; PALFAMS= PAL First Attempt Memory Score; PALMETS= PAL Mean Errors to Success.

 Table 4. Multivariable linear regression between children's object drawing scores and F⁻ exposure in drinking water

drinking water including other covariates.

| | 5-14 years old | | | | | | |
|----------------------|----------------------|----------------|-----------------|----------------|----------------------|--|--|
| | Unadjus | Adju | usted | | | | |
| | | | | β ^a | p-value | | |
| a. Donkey drawing | β (95%CI) | R ² | <i>p</i> -value | | R ² =0.39 | | |
| Water F ⁻ | -0.2 (-0.37, -0.03) | 0.075 | 0.024 | -0.2 | 0.030 | | |
| b. House drawing | | | | | R ² =0.44 | | |
| Water F ⁻ | -0.082 (-0.21, 0.05) | 0.024 | 0.21 | -0.003 | 0.96 | | |
| c. Person drawing | | | | | R ² =0.34 | | |
| Water F ⁻ | -0.09 (-0.28, 0.09) | 0.015 | 0.32 | J.15 | 0.13 | | |

Abbreviation: β , regression coefficient; *CI*, confidence interval.

^aAdjusted for sex, grade level, BMI, As and Pb in drinking water and anemic appearance.

| Table 5. Associations between F | concentration grut | ups in drinking | water and c | hildren's ob | ject |
|---------------------------------|--------------------|-----------------|-------------|--------------|------|
| drawing scores. | | | | | |

| Fluoride concentration in | Rela | Relative object drawing scores | | | |
|-------------------------------------|----------------------|--------------------------------|-----------------------------------|---------|--|
| drinking water (mg/L) | Crude, β (95% μ." | P value | Adjusted, ^a β (95% Cl) | P value | |
| For donkey | | | | | |
| Water F ⁻ exposure group | R ² =0.07 | | R ² =0.40 | | |
| Group 1 (< 3) | Reference | | Reference | | |
| Group 2 (>3–8) | -1.54 (· 3.c, 0.55) | 0.07 | -2.88 (-5.7, -0.08) | 0.29 | |
| Group 3 (>8–15.5) | -2.2 (-40.17) | 0.03 | -3.56 (-6.62, -0.48) | 0.024 | |
| For house | | | | | |
| Water F ⁻ exposure group | P ² -0.05 | | R ² =0.46 | | |
| Group 1 (< 3) | Refe ence | | Reference | | |
| Group 2 (>3–8) | -1.24 (-2.77, 0.30) | 0.11 | -0.46 (-2.46, 1.54) | 0.64 | |
| Group 3 (>8–15.5) | - 1.21 (-2.74, 0.31) | 0.21 | -0.26 (-2.44, 1.91) | 0.81 | |
| For person | | | | | |
| Water F ⁻ exposure group | R ² =0.02 | | R ² =0.35 | | |
| Group 1 (< 3) | Reference | | Reference | | |
| Group 2 (>3–8) | -0.60 (-2.90, 1.68) | 0.59 | -1.53 (-4.7, 1.65) | 0.33 | |
| Group 3 (>8–15.5) | -1.28 (-3.55, 0.99) | 0.26 | -2.77 (-6.23, 0.69) | 0.11 | |

 β , regression coefficient; CI, confidence interval.

^aAdjusted for sex, children's grade level, BMI, As and Pb in drinking water, anemic appearance.

Table 6. Associations between F⁻ concentrations in drinking water and children's performance in CANTAB PAL tasks.

| 5-14 years old | |
|----------------|--|
| | |

| | Unadjusted | | | Adjusted | | | |
|---|--------------------------|----------------|-------------|----------------------|-----------------|--|--|
| | | | | β (95%Cl) | p- valu e | | |
| b. PALTEA (PAL Total Errors (Adjusted) | β (95%Cl) | R ² | p- value | R ² =0.12 | | | |
| Water F ⁻ | 1.2 (0.32, 2.1) | 0.09 3 | 0.008 | 1.32 (0.05, 2.6) | 0.05 | | |
| b. PALNPR (PAL Number of Patterns Reached) | | | | R ² =0.10 | | | |
| Water F | -0.1 (-0.19, -0.01) | 0.06 4 | 0.03 | -0.1 (-0.22, 0.014) | 0.11 | | |
| c. PALFAMS (PAL First Attempt Memory Score) | | | | R ² =0.08 | | | |
| Water F | -0.21 (-0.42, 0.01) | 0.04 8 | 0.06 | -(.19 (-0.51, 0.13) | 0.23 | | |
| d. PALMETS (PAL Mean Errors to Success) | | | 5 | R ² =0.10 | | | |
| Water F | -0.075 (-0.16, 0.007) | 0.04 | 10.97 | -0.016 (-0.13, 0.09) | 0.78 | | |

β, regression coefficient; CI, confidence interval. ^aAdjusted for children's grade level, BMI, sex, As and Pr in drinking water, anemic appearance. Note that PALTEA =PAL Total Errors (Adjusted); PALNPR= PAL : ur ber of Patterns Reached; PALFAMS= PAL First Attempt Memory Score; PALMETS= PAL Mean Errors in Success.



Fig 1. Linear regression plots showing the association between F^- in drinking water-and children's (5 to 14 years old; n=68) drawing ability scores for: **A**) a donkey (r²=0.075, *p*=0.024); **B**) a house (r²=0.023, *p*=0.21), and a person (not shown) (r²=0.015, *p*=0.32). **C**) integrating all objects (a donkey, a house and a person) drawing scores (r²=0.027, *p*=0.019).

Journal Prendrock



Fig 2. Representative children's drawings of a donkey for children in different F^- exposure groups. The figures were selected by averaging the scores at e_i ch community and picked a drawing close to the mean score. Note that the label on each figure represents water concentration of F^- (mg/L) / drawing score.



Fig 3. Linear regression plots showing the association between \overline{r} in urine and children's (5 to 14 years old; n=48) drawing scores in the donkey task (r²=0.051, μ =0.12).

Journal Prove



Fig. 4. Linear regression picts showing the association between F⁻ in drinking water and **A**) children's PALTEA (r=0.30, p=0.008), and **B**) PALNPR (r=0.25, *p*=0.03) performance



Fig 5. Linear regression plots showing the effect of increasing F^- in drinking water on mean CANTAB PAL total errors made by the children for tasks featuring different numbers of PAL boxes. Note: The interaction of F^- and PAL cifficulty levels was not significantly related to the mean PAL total errors made by the sample children (p=0.085).

Declaration of conflict of interest

The authors declare they have no actual or potential competing interests.

provent of the second s ind a low

Highlights

- Cognitive ability was assessed in children exposed to a range of F⁻levels in drinking water.
- The study successfully implemented the CANTAB test to children residing in rural Ethiopia.
- Water F⁻ levels were negatively associated with children's drawing, and CANTAB's memory and learning tests.
- Children exposed to high F⁻made more errors as CANTAB task difficulty increased.
- The findings add urgency to further study the potential neurotoxicity of low and high F⁻ in drinking water.

Journal Pression