



Title	Lead poisoning in children from townships in the vicinity of a lead-zinc mine in Kabwe, Zambia
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2 Zambia

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21 *Extensive childhood Pb poisoning in Zambia's Kabwe mining town may have serious*  
22 *health effects on the children ranging from neurological deficits to deaths.*

23

24 Abstract

25 Childhood lead poisoning is a serious public health concern worldwide. Blood lead levels  
26 exceeding 5 µg/dL are considered elevated. In Kabwe, the capital of Zambia's Central  
27 Province, extensive Pb contamination of township soils in the vicinity of a Pb-Zn mine and  
28 posing serious health risk to children has been reported. We investigated BLLs in children  
29 under the age of 7 years in townships around the mine; where blood samples were  
30 collected and analysed using an ICP-MS. Almost all of the sampled children had BLLs  
31 exceeding 10 µg/dL. Children in these areas could be at serious risk of Pb toxicity as 18%  
32 of the sampled children in Chowa, 57% (Kasanda) and 25% (Makululu) had BLLs  
33 exceeding 65 µg/dL. Eight children had BLLs exceeding 150 µg/dL with the maximum  
34 being 427.8 µg/dL. We recommend that medical intervention be commenced in the  
35 children with BLL exceeding 45 µg/dL.

36 **Keywords:** Children; lead poisoning; Pb-Zn mine; Kabwe, Zambia.

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38

## 39 **1. Introduction**

40 Childhood lead (Pb) poisoning is a serious public health concern worldwide (Tong et al.  
41 2000). Exposure to Pb affects multiple organ systems resulting in numerous morphological,  
42 biochemical and physiological changes that include hematological disorders, nervous  
43 system disturbances and impairment of liver and kidney functions (Lockitch 1993; Al-  
44 Saleh 1994; Canfield et al. 2003; Needleman 2004). Young children are particularly  
45 vulnerable to Pb exposure and poisoning. This is because young children frequently  
46 explore their environment via hand-to-mouth and object-to-mouth activities; behaviors that  
47 are likely to increase Pb intake in children from polluted environments such as house dust  
48 or yard soils (Calabrese et al. 1997; Manton et al. 2000). Biological factors also play a  
49 significant role in increased Pb uptake in children as the average fractional gastrointestinal  
50 absorption of Pb is much greater in infants and young children than in adults (Ziegler et al.  
51 1978). Moreover, Pb absorption is increased in the presence of nutritional deficiencies such  
52 as iron and calcium, which are more common in children than in adults (Bradman et al.  
53 2001). Children are also more vulnerable to Pb poisoning compared to adults as the central  
54 nervous system is most sensitive to Pb toxicity during developmental stages (Bellinger  
55 2004; Lidsky and Schneider 2003). Although the effects of Pb on the nervous system in  
56 adults tend to reverse after cessation of exposure (Baker et al. 1985), effects in children  
57 tend to persist (Needleman et al. 1990).

58 Lead concentration in whole blood (BLL) is the main biomarker used to monitor  
59 exposure and has been widely used in epidemiological studies (CDC 2009). The Centers  
60 for Disease Control and Prevention (CDC 2012) recently revised the blood lead “level of  
61 concern” from 10 to 5  $\mu\text{g}/\text{dL}$  in response to reports that BLLs  $< 10 \mu\text{g}/\text{dL}$  can cause  
62 neurological abnormalities such as decreased intelligence quotient (IQ) in children

63 (Canfield et al. 2003). Therefore, a threshold below which Pb does not result in  
64 neurological deficits has not been determined (Needleman 2004). However, individuals  
65 differ widely in the BLL at which signs of Pb toxicity appear, with some individuals  
66 seeming well at a BLL that in others results in encephalopathy or even death (Bellinger  
67 2004). The detrimental effects of elevated BLLs in the range of 10 to 45  $\mu\text{g}/\text{dL}$  are usually  
68 subclinical and may include neurodevelopmental impairment (CDC 2002). Generally,  
69 BLLs  $> 10 \mu\text{g}/\text{dL}$  in children are considered elevated and it has been recommended that  
70 chelation therapy be initiated at levels  $\geq 45 \mu\text{g}/\text{dL}$  (CDC 2002; Needleman 2004). At  
71 higher BLLs  $> 60 \mu\text{g}/\text{dL}$ , clinical symptoms such as abdominal pain and arthralgia become  
72 visible in children (Needleman 2004). Moreover, it has been reported that high BLLs  
73 exceeding  $100 \mu\text{g}/\text{dL}$  can cause encephalopathy, convulsions, coma and death, especially  
74 in children (CDC 2002; TNO 2001).

75 In the last decade, BLLs in children have reduced significantly in a number of developed  
76 countries following the phasing out of leaded gasoline (Wilhelm et al. 2006). However,  
77 childhood Pb toxicity continues to be a major public health problem in most developing  
78 countries. In Africa, major sources of childhood Pb poisoning include Pb mining and  
79 smelting, paint and battery recycling (Nriagu et al. 1996; Mathee et al. 2007). The recent  
80 Pb poisoning disaster in Nigeria, where more than 400 children died leaving numerous  
81 others with long-term neurological impairment including blindness and deafness, was  
82 attributed to gold ore-mining and processing, especially that metals were processed in their  
83 dwellings (Blacksmith Institute 2011, Dooyema et al. 2012; Lo et al. 2012). In Kabwe  
84 Town, the capital of Zambia's Central Province, extensive Pb contamination of township  
85 soils in the vicinity of a Pb-Zn mine has been reported and poses a serious health risk to  
86 children in these townships (Tembo et al. 2006; Nakayama et al. 2011). In an earlier study,

87 Pb poisoning and cases of encephalopathy were recorded in children from a township in  
88 the vicinity of the Pb-Zn mine in Kabwe (Clark APL, unpublished data).

89 Despite extensive Pb pollution in Kabwe, comprehensive studies of Pb exposure and  
90 poisoning in children in the vicinity of the mine are rare. In animal studies however, high  
91 concentrations of Pb were reported in wild rats (Nakayama et al. 2011; Nakayama et al.  
92 2013) as well as blood and edible organs of cattle (Yabe et al. 2011; Ikenaka et al. 2012)  
93 and chickens (Yabe et al. 2013) reared in the vicinity of the mine in Kabwe. Therefore, the  
94 objectives of the current study were to investigate BLLs in children under the age of 7  
95 years in townships around the Pb-Zn mine in Kabwe and to identify children with BLLs  
96 that require medical intervention.

97

## 98 **2. Materials and methods**

99

### 100 *2.1 Sampling sites*

101 Kabwe town, the provincial capital of Zambia's Central Province, is located at about  
102 28°26'E and 14°27'S. Kabwe has a long history of Pb-Zn mining. The mine operated  
103 almost continuously from 1902 to 1994 without addressing the potential risks of metal  
104 pollution. Dense fumes rich in Pb and other metals were emitted from smelters and they  
105 polluted the environment in the surrounding communities extensively (Tembo et al. 2006).  
106 Despite closure of the mine, scavenging of metal scraps from the abandoned tailings and  
107 wastes stored on the mine has continued to serve as a source of metal pollution, especially  
108 dusts emanating from the mine dumps.

109 In the current study, blood samples were collected from children at health centers  
110 located in Chowa, Kasanda and Makukulu townships, in May-June of 2012. Kasanda  
111 Township lies west to the mine and its center is about 2.2 km from the smelter (Fig. 1).

112 However, some households in Kasanda are within 1 kilometer of the mine. Makululu  
113 Township is a large squatter compound that lies adjacent and to the west of Kasanda  
114 Township. These two townships are affected by dust emanating from the mine as the  
115 prevailing winds most of the time blow from the east to the west. Most houses in Makululu  
116 are made of mud brick walls, mud floors and thatched roofs. Moreover, lots of dust is  
117 emitted by vehicles as roads in the township are not tarred. Many households in the  
118 township use well water in addition to communal water taps and there are high levels of  
119 poverty in the community. Chowa Township is equally close to the mine as Kasanda but is  
120 least affected by dust as it lies on the windward side of the mine. In contrast to Makululu,  
121 houses in Kasanda and Chowa are made of concrete bricks and use indoor tap water.  
122 Children from these townships were selected because soil samples in these townships are  
123 highly polluted with Pb (9-51188 mg/kg) and other metals (Nakayama et al. 2011).

## 124 *2.2 Blood collection*

125 The study was approved by the University of Zambia Research Ethics Committee  
126 (UNZAREC) and the Ministry of Health, Zambia. After informed and written consent was  
127 obtained from the children's parents or guardians, blood samples were collected by  
128 qualified laboratory technicians at Chowa, Kasanda and Makululu clinics. Before sampling  
129 commenced, an awareness campaign about the research activities was conducted by  
130 community health workers in each township to encourage parents/guardians to take their  
131 children under the age of 7 to the health centers for sample collection. To avoid sample  
132 contamination, all blood collection supplies were kept in plastic ziploc storage bags before  
133 sample collection. For each child, data on the age, sex and residential area were recorded.  
134 Blood up to 10 mL was collected from the cubital vein of each child, after careful cleaning  
135 and sanitization of the venipuncture site with an ethanol swab to avoid contamination, into

136 plain blood collection tubes for Pb analysis. The blood samples were immediately stored in  
137 freezers at -20 °C after sampling and then transported in cooler boxes on dry ice to the  
138 laboratories of the Kabwe District Health Offices and Kabwe Provincial Veterinary Offices  
139 where they were again stored at - 20 °C. After obtaining the material transfer agreement  
140 (MTA) clearance from the Zambia National Health Research Ethics Committee (NHREC),  
141 the blood samples were transported to Japan in cooler boxes on dry ice and analyzed for  
142 metal concentrations in Laboratory of Toxicology, Graduate School of Veterinary  
143 Medicine, Hokkaido University.

#### 144 *2.3 Sample preparation and metal extraction*

145 All laboratory materials and instruments used in metal extraction were washed in 2 %  
146 nitric acid (HNO<sub>3</sub>) and oven dried. The metal was extracted in blood samples using  
147 microwave digestion system (Speedwave MWS-2; Berghof) according to the  
148 manufacture's instruction. Metal extraction was done as recommended by Schweitzer and  
149 Cornett (2008). Briefly, 1 mL of each blood sample was placed in prewashed digestion  
150 flasks, and 5 mL of 60 % nitric acid (Kanto Chemical) and 1 mL of 30 % hydrogen  
151 peroxide (Kanto Chemical) were added. After digestion in the microwave for 52 minutes  
152 and temperatures of up to 190 °C, the digested samples were transferred into plastic tubes.  
153 The volume was then made up to 10 mL with bi-distilled and de-ionized water (Milli-Q).

#### 154 *2.4 Metal analysis*

155 Blood Pb concentrations were analyzed by Inductively Coupled Plasma-Mass  
156 Spectrometer (ICP-MS; 7700 series, Agilent technologies, Tokyo, Japan). The precision  
157 and accuracy of the applied analytical method was evaluated by analyzing the recovery  
158 rate using digested blood samples and spiking Pb standard solutions. Using this method, a  
159 good recovery of 97% was obtained. Certified Reference Materials, DORM-3 (Fish protein,

160 National Research Council of Canada, Ottawa, Canada) and DOLT-4 (Dogfish liver,  
161 National Research Council of Canada, Ottawa, Canada) were used to evaluate recoveries.  
162 Replicate analysis of these reference materials also showed good recoveries (95-105%).  
163 Instrument detection limit was 0.001 µg/L.

## 164 *2.5 Statistical analysis*

165 The data of BLLs were log transformed to stabilize variances. Statistical analysis was  
166 performed using JMP version 9 (SAS Institute, USA). The data are presented as mean,  
167 median and minimum-maximum values in µg/dL, wet weight. A stacked histogram was  
168 used to analyzed blood Pb accumulation trends in Kasanda and Makululu as well as in  
169 boys and girls. Stepwise multiple linear regression analyses on log-transformed data were  
170 used to estimate the influence of area, sex and age (0 – 3 years and 4 – 7 years old) on  
171 BLLs. Correlations between age and BLL were analyzed by both linear and quadratic  
172 regression analysis. Samples from Chowa were not included in the comparisons due to  
173 smaller sample size compared to Kasanda and Makululu. A *p*-value of less than 0.05 was  
174 considered to indicate statistical significance.

175

## 176 **3. Results**

177

### 178 *3.1 Blood lead levels (BLLs)*

179 A total of 246 blood samples were collected from children, up to 7 years old, at Chowa  
180 (*n* = 17 samples), Kasanda (*n* = 100) and Makululu (*n* = 129) health centres.  
181 Concentrations of Pb in blood samples are shown in Table 1.

182

183 As shown in Table 2, all of the sampled children had BLLs exceeding the guideline value  
184 that raise ‘health concerns’ (5 µg/dL). Numbers of children exceeding guideline values for

185 initiating chelation therapy (45  $\mu\text{g/dL}$ ), toxicity level (65 - 149  $\mu\text{g/dL}$ ) and levels  
186 associated with encephalopathy and death ( $> 150 \mu\text{g/dL}$ ) are also shown.

### 187 *3.2 Blood Pb accumulation patterns*

188 Using a stacked histogram, blood Pb accumulation patterns in children from Kasanda  
189 and Makululu as well as concentration differences between boys and girls in the two  
190 townships were analysed (Figure 2). Blood accumulation differences were highlighted as  
191 the highest BLLs were seen in younger children (0 – 3 years) than children aged 4 – 7  
192 years (Figure 3).

193

### 194 *3.3 Age and Sex differences*

195 Stepwise multiple linear regression analyses were performed on log-transformed data to  
196 estimate the influence of independent variables (age as continuous variable, sex  
197 represented as 0 for girls and 1 for boys, location (area) represented as 0 for Makululu and  
198 1 for Kasanda) on BLLs (Table 3). Concentrations in children from Kasanda were higher  
199 than levels in children from Makululu ( $p < 0.05$ ). There was no difference in the BLLs  
200 between boys and girls from Kasanda whereas in children from Makululu, BLLs were  
201 higher ( $p < 0.05$ ) in boys than girls. Younger children aged 0 - 3 years accumulated higher  
202 concentrations of Pb in blood than children aged 4 – 7 years in both Kasanda and Makululu  
203 ( $p < 0.05$ ).

204

205 Combining the data of Kasanda and Makululu, significant negative correlations between  
206 age and BLL were observed by both linear and quadratic regression analysis. Peak BLLs  
207 were observed around the age of 2 years (data not shown).

208

## 209 **5. Discussion**

210

211 The current study has demonstrated alarming childhood Pb poisoning in Zambia's Kabwe  
212 town, revealing serious Pb exposure in the children under the age of 7 years in townships  
213 surrounding the closed Pb-Zn mine. The study analysed BLLs in children because it is well  
214 established that children are more vulnerable to Pb poisoning and sensitive to its  
215 neurotoxic effects than adults (Lidsky and Schneider 2003). All of the sampled children in  
216 the current study had indications of Pb poisoning, with BLLs exceeding the 5 µg/dL "level  
217 of concern" set by CDC (2012). Moreover, the current study revealed that children in these  
218 townships could be at serious risk of Pb toxicity as 18% of the sampled children in Chowa,  
219 57% (Kasanda) and 25% (Makululu) had BLLs exceeding 65 µg/dL; the threshold widely  
220 considered to result in Pb toxicity (CDC 2002; Needleman 2004). Of the 246 children in  
221 the current study, 8 had BLLs exceeding 150 µg/dL, up to 427 µg/dL.

222 These findings agreed with reports in an earlier study before closure of the mine, where  
223 mean BLLs of 37 - 107 µg/dL were recorded in children from Kasanda Township (Clark  
224 APL, unpublished data). Of the 91 children between the ages of 1 - 2 years that were  
225 attended to at Kasanda clinic in the earlier study, 89% were reported to have accumulated  
226 BLLs > 60 µg/dL (Clark APL, unpublished data) compared to 61% of the sampled children  
227 from the same clinic in the current study. Therefore, there could be no difference between  
228 the severity of Pb poisoning during active mining period and almost 20 years after closure  
229 of the mine. Higher BLLs than the current study were recorded in children under the age of  
230 5 years in Zamfara State in Nigeria, where the affected families processed metals in their  
231 dwellings (Blacksmith Institute 2010; Dooyema et al. 2012; Lo et al. 2012). In the study by  
232 Dooyema et al. (2012), BLLs exceeding 10 µg/dL were reported in all the 204 sampled

233 children in Nigeria. In children from Nigeria, mean BLLs (107.5 – 153.3 µg/dL) were  
234 higher than mean BLLs in the current study (39 – 82.2 µg/dL). However, the maximum  
235 BLL of 445 µg/dL recorded in children from Nigeria was comparable to that of the current  
236 study (427.8 µg/dL). Although data on mortalities due to Pb poisoning in Kabwe are scarce,  
237 clinical signs consistent with Pb poisoning such as anemia, small stature and weakness  
238 were observed in children from the sampled areas during the current study. In Nigeria, over  
239 400 children were reported to have died of Pb poisoning (Blacksmith Institute 2011,  
240 Dooyema et al. 2012; Lo et al. 2012). Findings in the current study were higher than BLLs  
241 in children from an urban population in Kinshasa, Democratic Republic of Congo, where  
242 mean BLLs of 9.9 µg/dL and maximum concentrations of 49.3 µg/dL were recorded  
243 (Tuakuila et al. 2013). Moreover, BLLs in the current study were higher than mean BLLs  
244 (16.38 µg/dL) in children in the vicinity of Pb mines and sheltering plants in China (Lin et  
245 al. 2011). When compared to most European countries where the median BLL in the  
246 general population is below 5 µg/dL (Taylor et al. 2007), it is evident from the current  
247 study that levels of Pb poisoning in Kabwe, Zambia are alarming.

248 When the severity of Pb poisoning among the townships was compared in the current  
249 study, the mean BLL in children from Kasanda (82.2 µg/dL) was higher ( $p < 0.05$ ) than  
250 Makululu (57.1 µg/dL). Kasanda and Makululu were subjected to atmospheric Pb pollution  
251 emanating from the neighbouring mine as they are located on the western side of the mine,  
252 which is in the direction of the prevailing winds. However, the difference in BLLs in  
253 children from the two townships could be attributed to distance from the mines. Although  
254 all these townships were close to the mine, some households in Kasanda (even Chowa)  
255 were within 1 kilometre of the mine and the abandoned mine dumps hence most of the  
256 polluted dust settles in Kasanda Township. Despite being further away from the mine

257 compared to Kasanda, Makululu Township, the largest shanty compound in Zambia  
258 equally poses a serious threat as roads, dwellings and house floors are dusty. Therefore,  
259 more children in Makululu Township could be at risk of Pb poisoning due to poverty and  
260 poor living conditions.

261 There was no gender difference in BLLs between boys and girls in Kasanda Township.  
262 This finding was in agreement with observations in the Democratic Republic of Congo  
263 (Tuakuila et al. 2013). However, trends in blood Pb accumulations between boys and girls  
264 were observed in the current study as boys in Makululu Township accumulated higher  
265 BLLs ( $p < 0.05$ ) than girls in the same township. The same was observed when data of  
266 both Kasanda and Makululu were combined. Different behaviours between boys and girls  
267 could be one of the factors contributing to this difference as boys are likely to cover more  
268 distance away from home and play near the mine dumps than girls. When children in the  
269 current study were grouped according to age, it was observed that younger children  
270 between the ages of 0 – 3 years accumulated higher BLLs than their older counterparts (4 –  
271 7 years). Significant negative correlation between age and BLL supported this finding.  
272 Similarly, younger children (1 – 2 years) in the Democratic Republic of Congo  
273 accumulated higher BLLs than older children (Tuakuila et al. 2013). Therefore, findings in  
274 the current study emphasized the increased susceptibility of younger children to the health  
275 risks of Pb pollution.

276 Earlier studies also observed that BLLs tend to peak at around 2 years of age (Koller et  
277 al. 2004). This observation is not unexpected as this period encompasses both the onset of  
278 independent ambulation and the time when a child's oral exploration of the environment  
279 including hand-to-mouth or object-to-mouth behaviour (pica) is greatest. This exposure  
280 pathway of children has been well documented in other studies (Lanphear and Roghmann

281 1997; Lanphear et al. 2002). It has been established that children typically ingest an  
282 average of 50 mg/day of soil (Stanek and Calabrese 1995). However, this amount can  
283 exceed 5 g a day in the case of pica (Mielke and Reagan 1998), with some children having  
284 been reported to ingest 25-60 g during a single day (Calabrese et al. 1997). Given that  
285 maximum Pb concentration in soils in the vicinity of the mine in Kabwe is about 50,000  
286  $\mu\text{g/g}$  or 50 mg/g (Nakayama et al. 2011), it means that children who ingest about 5 - 60 g  
287 of soil/day in the vicinity of the mine in Kabwe would ingest 250 - 3000 mg of Pb/day.  
288 Since the permissible tolerable weekly intake (PTWI) of Pb is 25  $\mu\text{g/Kg}$  of body weight  
289 per week (WHO 1987), concentrations of Pb ingested by children through pica in Kabwe  
290 mining area could be high.

291 The current study has demonstrated that childhood Pb poisoning in Zambia's Kabwe  
292 mining town is among the highest in the world, especially in children under the age of 3  
293 years. Lead exposure among children is associated with developmental abnormalities  
294 including impaired cognitive function, reduced intelligence, impaired hearing and reduced  
295 stature (Canfield et al. 2003; Jusko et al. 2008). Although reports of clinical cases and  
296 deaths due to Pb poisoning among children in Kabwe are rare, the findings of the current  
297 study indicate that more studies need to be done in order to clearly establish the health  
298 effects of Pb poisoning in children exposed to Pb pollution in the townships around the  
299 mine in Kabwe. This is important because BLLs in all of the sampled children in the  
300 current study exceeded 5  $\mu\text{g/dL}$ . In children, it has been established that neurobehavioral  
301 effects such as decrease in IQ may occur at BLLs  $< 10 \mu\text{g/dL}$  (Canfield et al. 2003).  
302 Moreover, BLLs of 40 - 60  $\mu\text{g/dL}$  are considered to be markedly elevated, resulting in  
303 distinct neurobehavioral effects (TNO 2005). Since 18 % of the sampled children from  
304 Chowa, 57 % (Kasanda) and 25 % (Makululu) in the current study had markedly elevated

305 BLLs exceeding 65 µg/dL, it would not be surprising to observe neurological effects of Pb  
306 poisoning in the exposed children. Although this is the first published study evaluating Pb  
307 poisoning in Kabwe, it was earlier reported that during the mining period between 1971 to  
308 1973, cases of suspected Pb poisoning with encephalopathy occurred among children aged  
309 10 to 30 months living in the township of Kasanda (Clark APL, unpublished data).  
310 Therefore, the children in Chowa, Kasanda and Makululu townships should be closely  
311 monitored to enable early detection of clinical signs related to Pb toxicity and medical  
312 intervention.

313

## 314 **6. Conclusions**

315 Given that Pb poisoning among children in Kabwe was extensive, it is recommended that  
316 chelation therapy be commenced in the children with BLL exceeding 45 µg/dL prior to the  
317 onset of symptoms to reduce morbidity and prevent mortality in the affected children. This  
318 can be achieved for each child by devising and implementing an individualized plan of  
319 follow-up, especially for those children with extremely high BLLs. Interrupting the process  
320 of Pb poisoning through early detection and intervention can prevent children from dying  
321 or suffering severe permanent effects of Pb toxicity such as persistent seizures and mental  
322 retardation. Moreover, urgent interventions are required to reduce Pb exposure in the  
323 affected townships. This can be done through community-based programs to educate the  
324 affected communities about the health effects of Pb, sources of Pb and practical ways of  
325 reducing Pb exposure in their homes and communities.

326

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342 **Conflict of interest**

343 The authors declare no conflicts of interest.

344 **References**

- 345 Al-Saleh I. 1994. Biochemical and clinical consequences of lead poisoning. *Med Res Rev*  
346 14:415-486.
- 347 Baker EL, White RF, Pothier LJ, Berkey CS, Dinse GE, Travers PH, et al. 1985.  
348 Occupational lead neurotoxicity: improvement in behavioral effects after reduction of  
349 exposure. *Br J Ind Med* 42:507-516.
- 350 Bellinger DC. 2004. Lead. *Pediatrics* 113:1016-1022.
- 351 Blacksmith Institute. 2011. Project Completion Report: Nigeria Lead Poisoning Crisis.  
352 Available: [http://www.blacksmithinstitute.org/nigerian-lead-poisoning-crisisblacksmith-](http://www.blacksmithinstitute.org/nigerian-lead-poisoning-crisisblacksmith-report.html)  
353 [report.html](http://www.blacksmithinstitute.org/nigerian-lead-poisoning-crisisblacksmith-report.html).
- 354 Bradman A, Eskenazi B, Sutton P, Athanasoulis M, Goldman LR. 2001. Iron deficiency  
355 associated with higher blood lead in children living in contaminated environments.  
356 *Environ Health Perspect* 109 :1079-1084.
- 357 Calabrese EJ, Stanek EJ, James RC, Roberts SM. 1997. Soil ingestion: a concern for acute  
358 toxicity in children. *Environ Health Perspect* 105:1354-1358.
- 359 Canfield RL, Henderson Jr CR, Cory-Slechta DA, Cox C, Jusko TA, Lanphear BP. 2003.  
360 Intellectual impairment in children with blood lead concentrations below 10 µg per  
361 deciliter. *N Engl J Med* 348:1517-1526.
- 362 Centers for Disease Control and Prevention (CDC). 2012. Low Level Lead Exposure Harms  
363 Children: A Renewed Call for Primary Prevention. Report of the Advisory Committee  
364 on Childhood Lead Poisoning Prevention of the Centers for Disease Control and  
365 Prevention. Atlanta, Ga. [online]. Available at URL:  
366 [www.cdc.gov/nceh/lead/acclpp/final\\_document\\_030712.pdf](http://www.cdc.gov/nceh/lead/acclpp/final_document_030712.pdf).

367 Center for Disease Control and Prevention (CDC). 2009. Fourth National Report on Human  
368 Exposure to Environmental Chemicals. Available at URL:  
369 <http://www.cdc.gov/exposurereport/pdf/FourthReport.pdf>.

370 Dooyema CA, Neri A, Lo YC, Durant J, Dargan PI, Swarthout T, et al. 2012. Outbreak of  
371 fatal childhood lead poisoning related to artisanal gold mining in northwestern Nigeria,  
372 2010. *Environ Health Perspect* 120: 601-607.

373 Ikenaka Y, Nakayama SMM, Muroya T, Yabe J, Konnai S, Darwish WS, et al. 2012. Effects  
374 of environmental lead contamination on cattle in a lead/zinc mining area: changes in  
375 cattle immune systems on exposure to lead in vivo and in vitro. *Environ Toxicol Chem*  
376 31: 2300-2305.

377 Jusko TA, Henderson CR Jr, Lanphear BP, Cory-Slechta DA, Parsons PJ, Canfield RL. 2008.  
378 Blood lead concentrations < 10 µg/dL and child intelligence at 6 years of age. *Environ*  
379 *Health Perspect* 116:243-248.

380 Koller K, Brown T, Spurgeon A, Levy L. 2004. Recent developments in low-level lead  
381 exposure and intellectual impairment in children. *Environ Health Perspect* 112:987-994.

382 Lanphear BP, Roghmann KJ. 1997. Pathways of lead exposure in urban children. *Environ Res*  
383 74: 67-73.

384 Lanphear BP, Hornung R, Ho M, Howard CR, Eberly S, Knauf K. 2002. Environmental lead  
385 exposure during early childhood. *J Pediatr* 140: 40-47.

386 Lidsky TL, Schneider JS. 2003. Lead neurotoxicity in children: basic mechanisms and clinical  
387 correlates. *Brain* 126:5-19.

388 Lin S, Wang X, Tak Sun Yu I, Tang W, Miao J, Li J, et al. 2011. Environmental lead  
389 pollution and elevated blood lead levels among children in a rural area of China. *Am J*  
390 *Public Health* 101:834-841.

391 Lo YC, Dooyema CA, Neri A, Durant J, Jefferies T, Medina-Marino A, et al. 2012.  
392 Childhood lead poisoning associated with gold ore processing: a village-level  
393 investigation-Zamfara State, Nigeria, October-November 2010. *Environ Health*  
394 *Perspect* 120:1450-1455.

395 Lockitch G. 1993. Perspectives on lead toxicity. *Clin Biochem* 26:371-381.

396 Manton WI, Angle CR, Stanek KL, Reese YR, Kuehnemann TJ. 2000. Acquisition and  
397 retention of lead by young children. *Environ Res* 82:60- 80.

398 Mathee A, Rollin H, Levin J, Naik I. 2007. Lead in paint: three decades later and still a hazard  
399 for African children? *Environ Health Perspect* 115: 321-322.

400 Mielke HW, Reagan PL. 1998. Soil is an important pathway of human lead exposure. *Environ*  
401 *Health Perspect* 106 (suppl 1):217-229.

402 Nakayama SMM, Ikenaka Y, Hamada K, Muzandu K, Choongo K, Teraoka H, et al. Metal  
403 and metalloid contamination in roadside soil and wild rats around a Pb-Zn mine in  
404 Kabwe, Zambia. *Environ Pollut* 159: 175-181.

405 Nakayama SMM, Ikenaka Y, Hamada K, Muzandu K, Choongo K, Yabe J, et al. 2013.  
406 Accumulation and biological effects of metals in wild rats in mining areas of Zambia.  
407 *Environ Monit Assess* 185: 4907-4918.

408 Needleman HL, Schell A, Bellinger D, Leviton A, Allred E. 1990. The long-term effects of  
409 exposure to low doses of lead in childhood: an 11-year follow-up report. *N Engl J Med*  
410 322:83- 88.

411 Needleman H. 2004. Lead poisoning. *Annu Rev Med* 55:209-222.

412 Nriagu JO, Blankson ML, Ocran K. 1996. Childhood lead poisoning in Africa: a growing  
413 public health problem. *Sci Total Environ* 1 81:93-100.

414 Schweitzer L, Cornett C. 2008. Determination of heavy metals in whole blood using  
415 Inductively-Coupled Plasma Mass Spectrometry: a comparison of microwave and  
416 dilution techniques. *The Big M* 4: 75-83.

417 Stanek EJ, Calabrese EJ. 1995. Daily estimates of soil ingestion in children. *Environ Health*  
418 *Perspect* 103:276-285.

419 Taylor A, Angerer J, Arnaud J, Claeys F, Kristiansen J, Mazarrasa O, et al. 2007. Differences  
420 in national legislation for the implementation of lead regulations included in the  
421 European directive for the protection of the health and safety of workers with  
422 occupational exposure to chemical agents (98/24/EC). *Int Arch Occup Environ Health*  
423 80: 254-264.

424 Tembo BD, Sichilongo K, Cernak J. 2006. Distribution of copper, lead, cadmium and zinc  
425 concentrations in soils around Kabwe town in Zambia. *Chemosphere* 63:497-501.

426 The Netherlands Organization for Applied Scientific Research (TNO). 2001. Risks to health  
427 and the environment related to the use of lead in products. Report STB-01-39; TNO-  
428 MEP, Apeldoorn, the Netherlands.

429 Tong S, Von Schirnding YE, Prapamontol T. 2000. Environmental lead exposure: a public  
430 health problem of global dimensions. *Bull World Health Organ* 78:1068-1077.

431 Tuakuila J, Lison D, Mbuyi F, Haufroid V, Hoet P. 2013. Elevated blood lead levels and  
432 sources of exposure in the population of Kinshasa, the capital of the Democratic  
433 Republic of Congo. *J Expo Sci Environ Epid* 23:81-87.

434 Wilhelm M, Schulz D, Schwenk M. 2006. Revised and new reference values for arsenic,  
435 cadmium, lead, and mercury in blood or urine of children: basis for validation of human  
436 biomonitoring data in environmental medicine. *Int J Hyg Environ Health* 209:301-305.

437 World Health Organization (WHO) (1987): Toxicology evaluation of certain food additives  
438 and contaminants (WHO food additives Series, No.21), Cambridge University Press,  
439 223-255.

440 Yabe J, Nakayama SMM, Ikenaka Y, Muzandu K, Ishizuka M, Umemura T. 2011. Uptake of  
441 lead, cadmium, and other metals in the liver and kidneys of cattle near a lead-zinc mine  
442 in Kabwe, Zambia. *Environ Toxicol Chem* 30:1892-1897.

443 Yabe J, Nakayama SMM, Ikenaka Y, Muzandu K, Choongo K, Mainda G, et al. 2013. 2013.  
444 Metal distribution in tissues of free-range chickens near a lead-zinc mine in Kabwe,  
445 Zambia. *Environ Toxicol Chem* 32:189-92.

446 Ziegler EE, Edwards BB, Jensen RL, Mahaffey KR, Fomon SJ. 1978. Absorption and  
447 retention of lead by infants. *Pediatr Res* 12:29-34.

448

449 **Figures legends:**

450 Fig. 1. A map of Kabwe showing different geographic areas and sampling sites

451

452 Fig. 2. Stacked histogram showing blood lead accumulation trends in children from Kasanda  
453 (46 boys and 54 girls) and Makululu (59 boys and 70 girls) townships of Kabwe, Zambia.

454

455 Fig. 3. Histogram showing blood lead accumulation trends in younger (0 – 3 years) and older  
456 (4 – 7 years) children from Kasanda and Makululu townships of Kabwe (Zambia).

Table 1.

Mean age (year) and BLLs ( $\mu\text{g/dL}$ ) of children from Chowa, Kasanda and Makululu townships in vicinity of the Pb-Zn mine in Kabwe, Zambia

Township	Mean age	Sample size	Arithmetic mean BLL	Median	Minimum	Maximum
Chowa	5.76	$n = 17$	39.0	39.3	15.6	79.7
Kasanda	3.65	$n = 100$	82.2	74.9	5.40	427.8
Makululu	4.51	$n = 129$	57.1	51.1	9.40	388.7

$n$  = Number of samples

Table 2.

Numbers of children (under the age of 7 years) with elevated BLLs from the sampled townships in Kabwe

Reference limits	Chowa ( <i>n</i> = 17)	Kasanda ( <i>n</i> = 100)	Makululu ( <i>n</i> = 129)
< 5 µg/dL	0	0	0
5 - 44 µg/dL – elevated levels	8	27	50
45 - 64 µg/dL – initiate treatment	7	15	44
65 - 149 µg/dL – toxicity level	2	50	33
> 150 µg/dL – encephalopathy, death	0	8	2

*n* = Number of children sampled

Table 3.

Blood lead accumulation differences (age, sex and site) by stepwise multiple linear regression analyses in children from Kasanda and Makululu townships in Kabwe.

Kasanda and Makululu						
Parameter	Estimate	nDF	SS	F Ratio	p value (Prob>F)	
Intercept	1.898	1	0.00	0.00	1.0	
Age	-0.035	1	1.05	15.84	<b>9.35E-05</b>	
Sex{F-M}	-0.050	1	0.56	8.44	<b>0.004</b>	
Area{Makululu-Kasanda}	-0.048	1	0.48	7.25	<b>0.008</b>	
Kasanda						
Parameter	Estimate	nDF	SS	F Ratio	p value (Prob>F)	
Intercept	1.936	1	0.00	0.00	1.0	
Age	-0.033	1	0.41	4.36	<b>0.039</b>	
Sex{F-M}	-0.060	1	0.35	3.77	0.055	
Makululu						
Parameter	Estimate	nDF	SS	F Ratio	p value (Prob>F)	
Intercept	1.861	1	0.00	0.00	1.0	
Age	-0.038	1	0.66	14.3	<b>0.0002</b>	
Sex{F-M}	-0.043	1	0.22	4.79	<b>0.030</b>	

Bold indicate significant ( $p < 0.05$ ), nDF: number of degrees of freedom for a term, SS: Sequential Sum of Squares

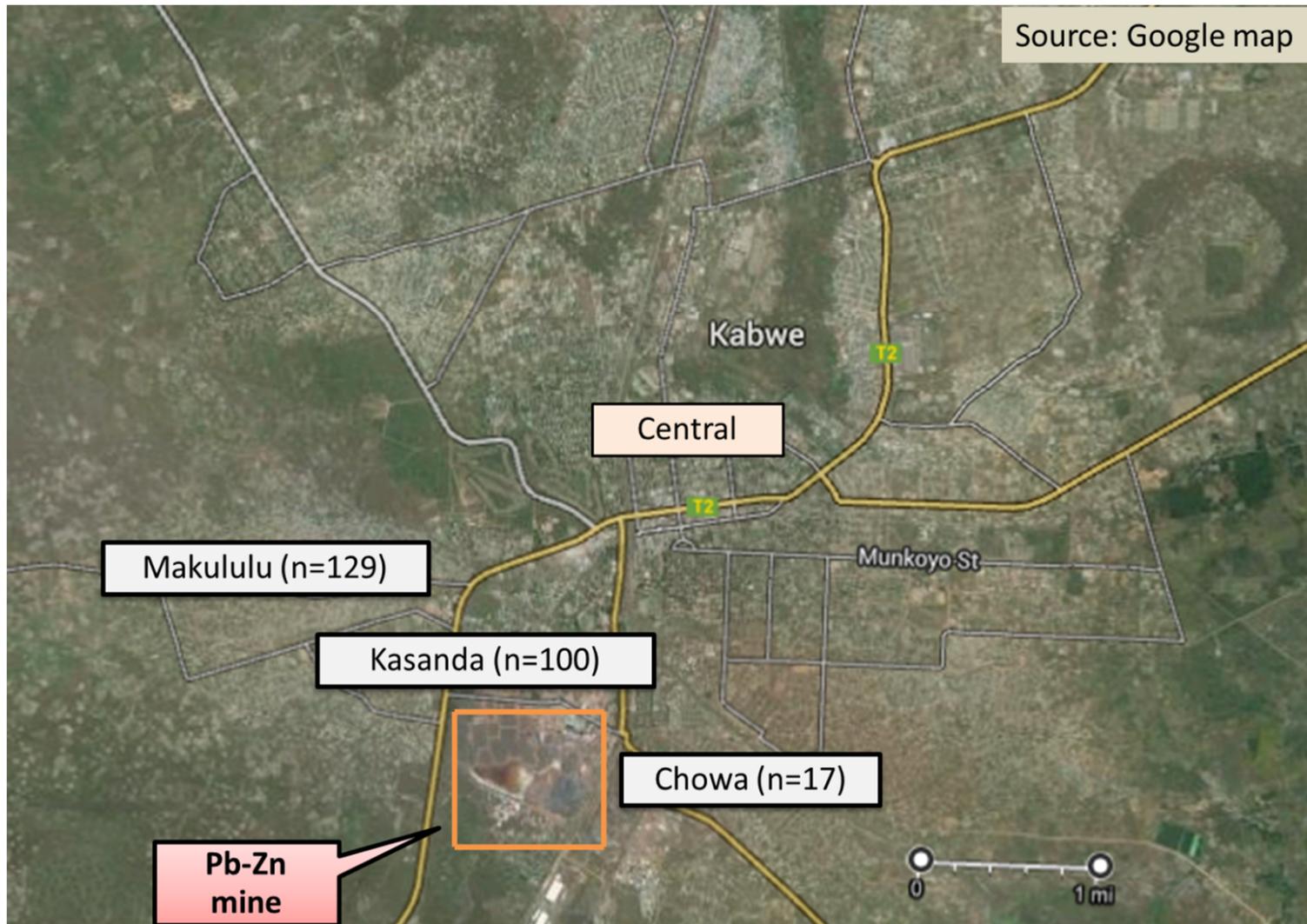


Fig. 1

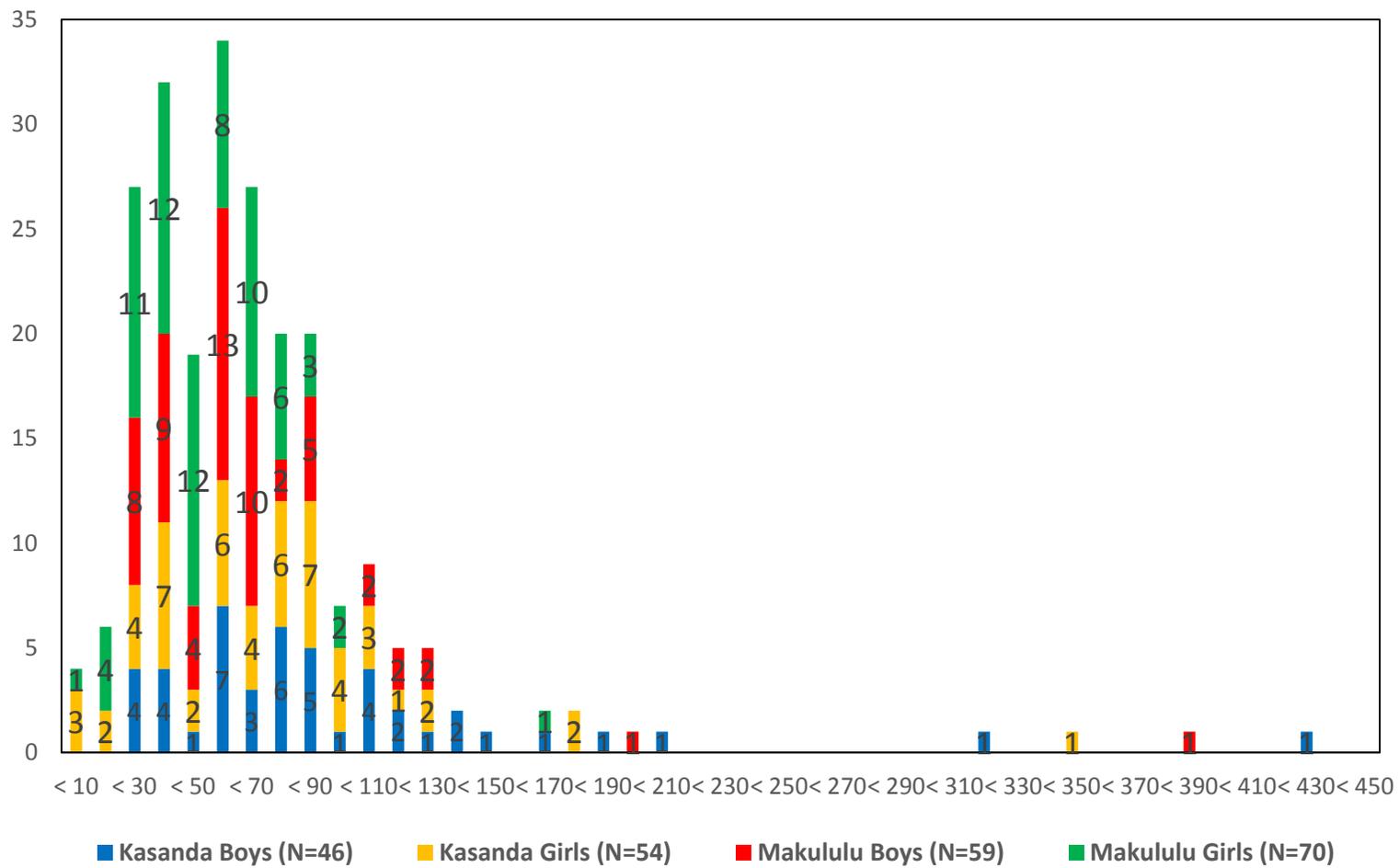


Fig. 2

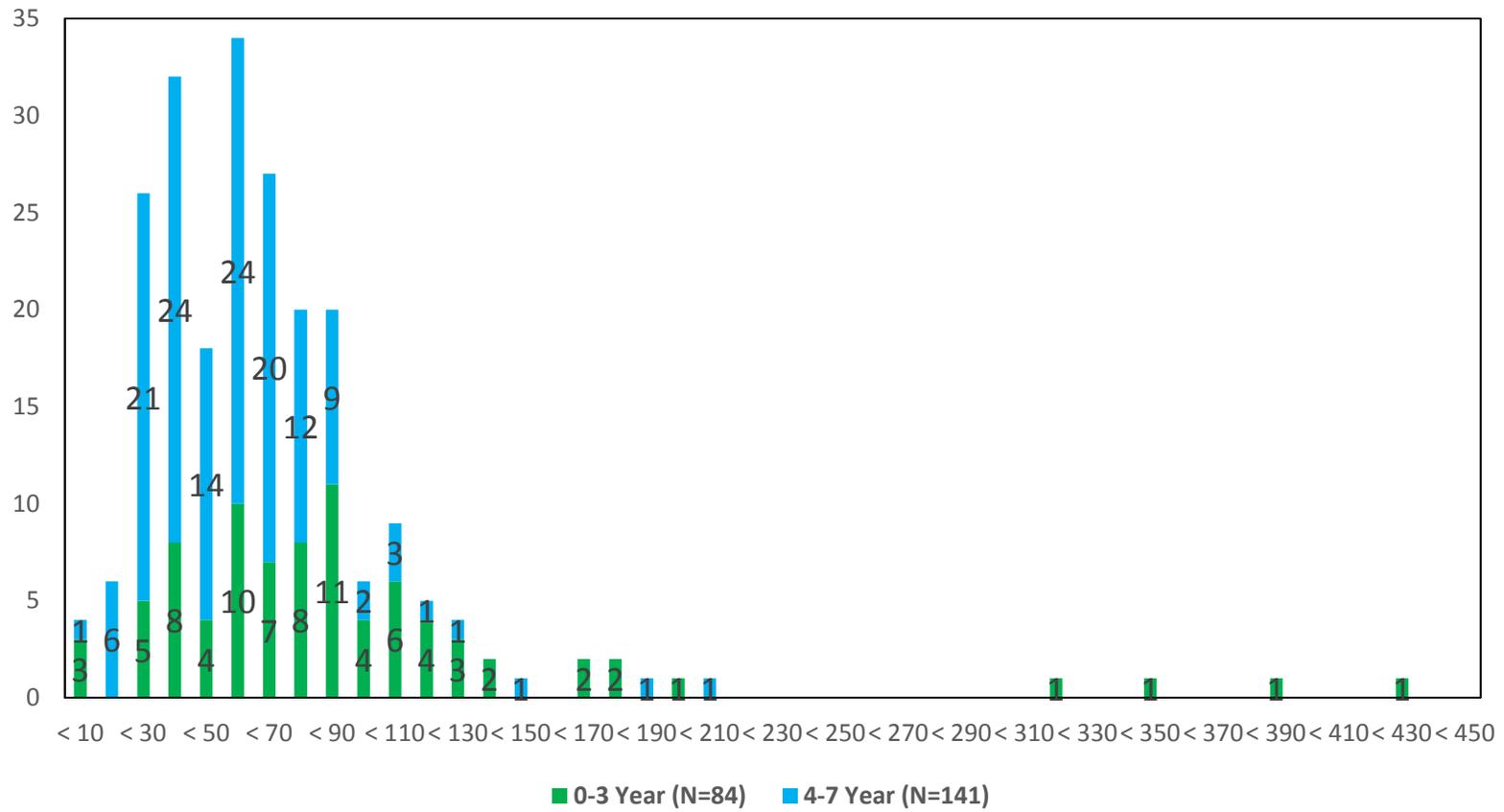


Fig. 3