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Effect of sewerage on the contamination of soil with pathogenic *Leptospira* in urban slums

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Manuscripts

1 **Effect of sewerage on the contamination of soil with pathogenic *Leptospira* in urban slums**

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20 **ABSTRACT**

21 Leptospirosis is an environmentally-transmitted zoonotic disease caused by pathogenic
22 *Leptospira spp.* that affects urban and rural poor communities world-wide. In urban slums,
23 leptospirosis is associated with precarious sanitary infrastructure. Yet, the role of sewerage in the
24 reduction of the environmental contamination with pathogenic *Leptospira* has not yet been
25 explored. Here, we conducted a survey of the pathogen in soils surrounding open and closed
26 sewer sections in six urban slums in Brazil. We found that the presence and concentration of
27 pathogenic *Leptospira* was lower in soils adjacent to conventionally closed sewers, when
28 compared to their open counterparts. However, no difference was observed in community closed
29 sewers. We also found that human fecal markers (BacHum) were positively associated with
30 pathogenic *Leptospira* even in closed sewers and that rat presence was not predictive of the
31 presence of the pathogen in soils suggesting that site-specific rodent control may not be
32 sufficient to reduce the environmental contamination with *Leptospira*. Overall, our results
33 indicate that sewerage expansion to urban slums may help reduce the environmental
34 contamination with the pathogen and therefore reduce the risk of human leptospirosis.

35

36 **Keywords**

37 Leptospirosis, sewer, public health, environment, fecal pollution

38

39 **Synopsis**

40 Sewerage construction in urban slums may reduce the presence and concentration of pathogenic
41 *Leptospira*, thus decreasing the risk of human exposures.

42

43 INTRODUCTION

44 Leptospirosis is a neglected zoonotic disease that affects urban and rural communities
45 worldwide¹ with an estimated annual burden of over a million cases and approximately 60,000
46 deaths.² Its clinical manifestations range from asymptomatic or a mild flu-like illness to severe
47 disease such as Weil's disease and pulmonary hemorrhagic syndrome for which fatality rates are
48 higher than 10% and 50%, respectively.^{3,4} Leptospirosis is caused by pathogenic spirochetes
49 from the genus *Leptospira*. Pathogenic *Leptospira* thrive in the kidneys of a wide variety of
50 animals, some of which are chronic carriers, and are released with the urine into the
51 environment at high concentrations^{5,6} where they can survive for extended time.^{7,8} Human
52 infection occurs through contact with previously contaminated water and soil or by exposure of
53 cuts and abraded skin with animal urine, making leptospirosis an environmentally-transmitted
54 disease.¹

55 Leptospirosis has historically been an occupational disease related to livestock raising,
56 mining, rice farming and other agricultural activities,⁹ but in the last 30 years it has emerged as
57 an epidemic in urban communities surrounding cities in developing countries.¹⁰⁻¹³ In these
58 neglected settings, poverty, precarious housing and trash accumulation create the ecological
59 conditions for the proliferation of rodents, particularly *Rattus norvegicus*, which are the primary
60 reservoirs of pathogenic *Leptospira* in urban environments.^{14,15} Extreme weather events and
61 seasonal periods of heavy rainfall increase the presence of the pathogen in the environment^{16,17}
62 and the likelihood of human exposure to contaminated water, soil and mud due to inadequate
63 sewer and storm drainage infrastructure.^{18,19} Indeed, cross-sectional and prospective
64 epidemiological studies have identified open sewers and drainage as risk factors for *Leptospira*
65 infection in urban slums.¹⁹⁻²² As the population living in urban slums is predicted to reach 2

66 billion by 2025²³, the burden of leptospirosis is only expected to increase.¹⁸ There is, therefore,
67 an urgent need to develop control measures for leptospirosis in resource-poor urban settings.

68 Sanitary interventions to close open sewers are an alternative to reduce exposures to
69 environmental sources of *Leptospira*^{16,24}, given the lack of efficacious vaccines for human use
70 ^{1,25} and the limited success of rodent control strategies due to regrowth after extermination ^{26,27}.
71 Sewerage construction is widely recognized to reduce the incidence of viral, bacterial and
72 parasitic diseases ^{28–30}. However, its effect on the reduction of pathogenic *Leptospira*
73 contamination has not been examined. Here, we aimed to determine the effect of conventional
74 and community-based sewer closings in the environmental contamination with pathogenic
75 *Leptospira*. To this end, we performed a cross-sectional study in soils surrounding open and
76 closed sewer sections in six Brazilian urban slums. The evaluation of the effect of sewerage in
77 preventing environmental contamination with the pathogen is critical to inform public health
78 interventions aimed to reduce the burden of leptospirosis in these neglected urban communities.

79

80 MATERIALS AND METHODS

81 Study sites

82 We conducted this study in six sites located in five urban slum settlements (*favelas*) in
83 the periphery of the city of Salvador (Brazil). The incidence of severe leptospirosis in urban
84 slums in Salvador is ca.19.8 cases per 100,000 inhabitants.²⁰ The communities studied were Pau
85 da Lima (sites 1 and 6), Sete de Abril (site 2), Campinas de Pirajá (site 3), Tancredo Neves (site
86 4) and Nova Constituinte (site 5) (Fig. 1A). All these slums have similar characteristics of
87 poverty, overcrowding, marginalization, poor quality housing and lack of reliable sanitation
88 infrastructure than other slum settlements in Brazil and other developing countries.^{23,31}

89 Specifically, the precarious sanitation system results in untreated sewage and storm water
90 drainage flowing through open sewers across these communities. In each community, we
91 selected one site containing a sewer with contiguous open and closed sections (Fig 1B and 1D).
92 Closed sections were classified as conventional or community-based depending on the type of
93 closing. Conventional closings (sites 1, 2, 3 and 4) were built by the local government sewage
94 company by digging trenches and placing sewer mains to which every house drain was
95 connected (Fig 1D). Conventional closings isolated the sewer and prevented sewage from
96 contaminating the surrounding environment. Community closings (sites 5 and 6) had been
97 performed informally by the local dwellers and consisted of wood planks or concrete boards
98 placed on top of the open sewer (Fig 1E). Community interventions prevented major spills from
99 the sewer but did not avoid leaking or major overflowing during rainfall events.

100

101 **Figure 1.** Distribution of sampling sites in the study area and typology of sewer closing. **A)** Map
102 of the city of Salvador (Brazil) with the locations of the six urban slum communities where soil
103 collections were performed: Pau da Lima (sites 1 and 6), Sete de Abril (site 2), Campinas de
104 Pirajá (site 3), Tancredo Neves (site 4) and Nova Constituinte (site 5). **B)** An open sewer section
105 in site 1. **C)** Soil sampling points (red dots) in soil at open sewer section. **D)** Conventionally
106 closed sewer section in site 2. **E)** Community closed sewer section in site 5. **F)** Soil sampling
107 points (red dots) in soil at closed sewer section

108



109

110 **Sampling design and sample collection**

111 At each site, open and closed section areas containing exposed soil within a 12 m
112 distance to the main sewer were demarcated, georeferenced and entered in a GIS database.
113 Polygons of 150 m² to 220 m² were drawn in each closed and open area and 24 collection points
114 were randomly selected using a packing density of 0.4 with corresponding minimum distances
115 between collection points for each area. (Fig 1C and 1F). Because of size constraints, only 16
116 collection points were selected in site 6. In total 272 collection points were selected, 136 in open
117 and 136 in closed sewer areas.

118 Samples were collected in the first week of December of 2018. At each collection point,
119 an area of ~400 cm² was cleared from surface rocks and vegetation debris, and ~25g of
120 subsurface soil were collected, stored in aseptic containers, transported to the laboratory and
121 processed within 4h of collection as described previously with minor modifications.³² Briefly, 40

122 mL of sterile double-distilled was added to each 5g sample and shaken with a horizontal vortex
123 adaptor at maximum speed for 2 min. Samples were centrifuged at 100 rcf for 5 min, the
124 supernatant recovered and centrifuged at 12,000 rcf for 20 min at room temperature. The
125 supernatants were discarded, and the pellets frozen at -20°C. In addition to the soil samples, two
126 paired 40-mL sewage samples were collected at the end of the closed and open sections in each
127 site. Sewage samples were processed as described previously.¹⁶

128

129 **Quantification of pathogenic *Leptospira* and human fecal markers.**

130 DNA was extracted from the frozen pellets within 3 days after processing using DNA
131 Easy PowerSoil kit (Qiagen) in batches of 20 samples and stored at -80 °C. An extraction blank
132 (sterile double-distilled water) was included to each batch to control for cross-contamination.

133 Pathogenic *Leptospira* was quantified using a TaqMan assay targeting the *lipL32* gene as
134 described previously.³² To determine the levels of human fecal contamination, we used the
135 BacHum TaqMan qPCR assay.³³ Calibration curves were included in each qPCR plate for with
136 concentrations of standard ranging from 2×10^2 to 2×10^9 GEq/mL. Samples were run in
137 duplicate and included non-template controls in each plate row to control for contamination.
138 qPCR inhibition was monitored using a an Internal Amplification Control (IAC) plasmid in
139 singleplex reactions as described previously¹⁶ for *lipL32* and testing at least two sample
140 dilutions for BacHum. For more details on cycling parameters, primer, probe and bovine serum
141 albumin (BSA) concentrations, calibration curves and tests for inhibition, see the Supporting
142 Information.

143

144 **Rat activity monitoring**

145 To evaluate the rat presence in the sampling sites during soil collections, we used a track
146 plate method that had previously showed high correlation with rat infestation measures and
147 trapping of rats to population exhaustion approaches.³⁴ Forty-eight track plates were placed in
148 each of the demarcated areas described above on the day of soil collection. Plates were randomly
149 distributed within each polygon with a packing density of 0.4 with corresponding minimum
150 distances between them (1.33 ± 0.73 m). Each site contained 96 track plates (48 in the
151 surroundings of the open section of the sewer and 48 in the closed section), for a total of 576
152 plates. Track plates were evaluated daily over the course of two days for evidence of rat activity
153 through the identification of footprints, scrapes, and tail slides and scored using a binary variable
154 (presence/absence of rat marks on a plate) and a continuous variable (the intensity of marks on
155 plates).³⁴ In addition, environmental rodent surveys were carried out at each sampling site by
156 looking for variables associated with rodent infestation and water or harborage sources for
157 rodents: pavement, soil, mud vegetation, trash, food, water, building material, rubble, others
158 animals and rat feces.³⁵

159

160 **Data treatment**

161 Samples were considered positive when both qPCR replicates showed amplification up to
162 a C_T of 40. Samples with a single positive reaction were submitted to an additional qPCR run in
163 duplicate. If in this second run the sample amplified in either of the replicates, it was considered
164 positive. The genomic equivalents (GEq) per reaction in all positive qPCR replicates were
165 averaged, normalized by the amount of soil or water processed, and \log_{10} -transformed to obtain
166 concentrations in \log_{10} GEq/g or mL. For the purpose of statistical analysis, soil samples with

167 concentrations below the limit of detection were considered to have a concentration equivalent to
168 the limit of detection of the lip132 qPCR assay in soil samples (2GC/g).³²

169

170 **Statistical analysis**

171 We built mixed generalized linear models (GLMMs) with binomial and gamma error
172 structure to investigate the probability of presence (binomial) and concentration of pathogenic
173 *Leptospira* in soil (continuous in log₁₀) and their association with sewer status (open /closed) and
174 type of closing (open, conventional or community-based).³⁶ We also included other covariates
175 such as distance to the sewer, soil moisture, presence of rats and human fecal markers (BacHum),
176 and a randomization factor for the sampling site. The modeling approach was carried out in two
177 stages. First, we built univariate models between all the variables and added an interaction
178 structure between them to understand how the presence and concentration of *Leptospira* in soil
179 varied. Variables with a p-value below 0.1 in univariate analyzes were included in the multivariate
180 analyzes, subsequently performed. Various multivariate statistical models were generated, and the
181 model with the lowest AIC (Akaike's Information Criterion) and $\Delta AIC < 2$ was selected as the best
182 model using the *dredge ()* function of the R MuMIn package.^{37,38} We estimated the odds ratios
183 (ORs) associated with the probability of *Leptospira* presence in soil and the rates (β coefficients)
184 for the *Leptospira* concentration model in soil. The analyzes were performed in R 3.3.1³⁹, and we
185 applied a significance level of $p < 0.05$.

186

187 **RESULTS**

188 **Presence of *Leptospira* DNA in soil samples**

189 We collected a total of 272 soil samples and 24 sewage samples in the six sites studied
 190 and tested them for the presence of pathogenic *Leptospira* DNA. Overall, 68 soil samples
 191 (25.0%) were positive for *Leptospira* DNA with more samples positive in soils surrounding the
 192 open sewer sections (31.6% [24.4–39.9%, 95% CI] than in their closed counterparts (18.4%
 193 [12.7%–26.8%, 95% CI] (Table 1). Among the 68 positive samples, the geometric mean
 194 concentrations and count range of *Leptospira* DNA was 3.3 [2.00–1.62×10³] GEq/g and 4.2 [2.0
 195 – 52.6] GEq/g in open and closed sections, respectively (Table 1, Fig. 1 and Suppl. Fig 1). The
 196 highest proportion of positive samples was detected in the open section of Site 1 (54.2%, 13 of
 197 24), whereas the lowest was found in the closed section of Site 2 (0%, 0 of 24) (Table 1).
 198 Interestingly, while we observed a relative reduction in the percentage of positive samples in
 199 most conventionally closed sites (sites 1, 2 and 3) compared to open sites, no reduction was
 200 observed in community closed sites (sites 5 and 6).

201

202 **Table 1.** Occurrence and concentration of pathogenic *Leptospira* in soils surrounding open and
 203 closed sewers in the six Brazilian urban slums. Closed sewers are classified based on the type of
 204 closing: conventional or community-based.

Site and type of closing	Pathogenic <i>Leptospira</i>		Pathogenic <i>Leptospira</i>	
	positivity rate (n and %) ^a		concentration (mean log ₁₀ and SD)	
	Open sewer	Closed sewer	Open sewer	Closed sewer
Conventional				
Site 1 – Pau da Lima 1	13 (54.2%)	2 (8.3%)	1.04 (0.70)	0.57 (1.19)
Site 2 – Sete de Abril	3 (12.5%)	0 (0.0%)	0.61 (0.82)	0.00 (0.00)
Site 3 – Campinas de Pirajá	8 (33.3%)	3 (12.5%)	-0.08 (0.74)	0.54 (0.04)

Site 4 – Tancredo Neves	4 (16.7%)	5 (21.7%)	1.04 (1.99)	0.48 (0.30)
Total conventional	28 (29.1%)	10 (10.5%)	0.67 (1.05)	0.51 (0.44)
Community				
Site 5 – Nova Constituinte	11 (45.8%)	11 (45.8%)	0.20 (0.62)	0.70 (0.56)
Site 6* – Pau da Lima 2	4 (26.7%)	4 (25.0%)	0.35 (0.45)	0.67 (0.72)
Total community	15 (38.4%)	15 (37.5%)	0.24 (0.57)	0.69 (0.57)
Overall	43 (31.9%)	25 (18.5%)	0.52 (0.93)	0.62 (0.52)

^a24 samples were collected in open and closed areas from sites 1-5.

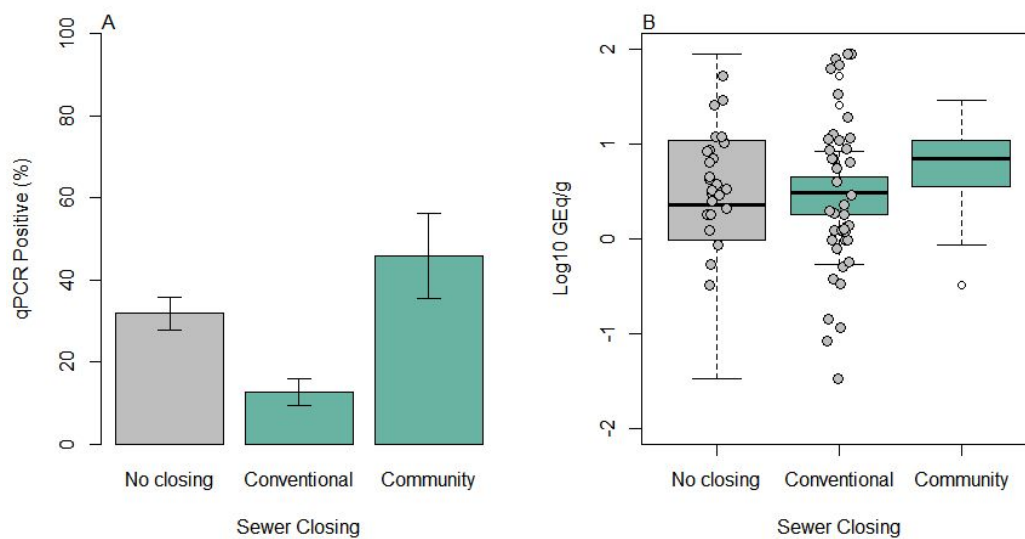
*16 samples were collected in open and closed areas from site 6.

205

206 **Fig 2.** Distribution of pathogenic *Leptospira* based on lipI32 qPCR in soils surrounding open and
 207 closed sewers. **A)** Occurrence by the type of sewer closing (mean percentage and standard
 208 error). **B)** Overall concentration of pathogenic *Leptospira* by the type of sewer closing
 209 (geometric mean and 95% confidence interval). Open sewers are denoted in gray and closed
 210 sewers in green.

211

212



213

214 Presence and concentration of human fecal pollution markers

215 We detected the human fecal pollution markers (BacHum) in 56.3% (153 of 272) of the
 216 soil samples collected (Suppl. Table 1). The presence of the marker was slightly higher in open
 217 than in closed areas of the sewers (58.8% [50.4%-66.8%] and 53.3% [45.0%-61.4%],
 218 respectively). The highest proportion of positive samples occurred in the open sections of sites 1
 219 and 6 (91.7% and 100%, respectively), whereas the lowest was found in the open and closed
 220 section of site 4 (8.3% and 16.7%, respectively). Among the 153 positive samples, the geometric
 221 mean concentrations and count range of BacHum was 3.04×10^3 [21.4 - 2.41×10^7] GEq/g and
 222 1.26×10^3 [66.4 – 8.06×10^3] GEq/g in open and closed sections, respectively (Supplemental Table
 223 2).

224 Presence of rats

225 We observed a higher presence and activity of rats as measured by tracking boards in the
 226 closed sections (12.2% and 29.2%; $p=0.0139$) when compared to the open sections of the sewers
 227 (9.3% and 18.1%; $p<0.001$). Tracking plates placed within the area of the closed section of the
 228 sewer had higher rat presence, 12.2 % (± 8.3) vs. 9.3% (± 9.8) $p=0.0139$, and higher percent rat

12

229 activity, 29.2% (± 6.9) vs. 18.1% (± 19.2) $p < 0.001$, than the tracking plates placed near the open
230 section of the sewer. We did not find significant differences between the open/closed status of
231 the sewer and the number of animals ($p = 0.4484$), number of rat holes ($p = 1.0000$), pavement
232 ($p = 0.4902$), soil ($p = 0.1138$), mud ($p = 0.5271$), vegetation ($p = 1.0000$), trash ($p = 1.0000$), food
233 ($p = 0.5271$), water ($p = 1.000$), building material ($p = 0.4902$), rubble ($p = 0.5271$), and rat feces
234 ($p = 1.000$).

235

236 **Sewage samples**

237 Pathogenic *Leptospira* was present in 17 of 24 the sewage samples collected with a
238 geometric mean and count range of 124 [20-1,545] GEq/mL. In all collection sites (before and
239 after sewage closing), at least one of the two paired samples collected was positive, indicating
240 that sewage was a frequent source of pathogenic *Leptospira* in all sites.

241

242 **Predictors of *Leptospira* DNA presence and concentration**

243 The univariate models found significant associations between the presence and
244 concentration of pathogenic *Leptospira* in soil with the status of the sewer (open/closed), the
245 type of closing, distance to other nearby open sewers, rat activity and the concentration fecal
246 human markers (BacHum) (Supp. Table 2 and 3). However, only two covariates remained
247 significant in the multivariate final models: type of sewer closing and presence of BacHum fecal
248 pollution markers (Table 2 and Supp. Table 4). First, soil samples collected in areas surrounding
249 sewers closed by the local government were more than 3 times less likely (inverse OR 3.44, 95%
250 CI: 1.66-8.33) to contain pathogenic *Leptospira* than soils collected in open areas overall. In
251 contrast, the presence of pathogenic *Leptospira* was not significantly different in soils

252 surrounding community closed sewers than in those adjacent to open sewers. Similarly, the
 253 logistic model using *Leptospira* concentration as outcome indicated that soils surrounding
 254 conventionally closed sewers contained a lower load of pathogenic *Leptospira* (0.82 log₁₀ units
 255 less, or approximately 6 times less). Furthermore, the logistic model showed that BacHum
 256 markers were significantly associated with the presence of pathogenic *Leptospira*. For every
 257 log₁₀ unit increase in BacHum concentration, the chances of finding a positive *Leptospira* sample
 258 increased by 15%. Likewise, the concentration of pathogenic *Leptospira* in positive samples was
 259 higher in those samples that also contained BacHum markers (Table 2). Notably, none of the
 260 other variables included in the model (rat presence and activity, soil moisture, distance to open or
 261 closed sewer and proximity to other open sewers) were found to be significantly associated with
 262 pathogenic *Leptospira* in the multivariate models. In summary, our model revealed that type of
 263 closing and BacHum markers were important predictors of presence and concentration of
 264 pathogenic *Leptospira* in soil.

266 **Table 2-** Final multivariate logistic and linear mixed models on the probability of finding a positive
 267 sample and log₁₀ concentration for *Leptospira* DNA. (**)*p* = 0.001 (***)*p* = 0.0001

Predictors	Logistic model for probability		Model for concentration	
	OR	CI	Estimates (β)	CI
(Intercept)	0.40***	0.21 – 0.70	-2.6***	-3.07 – -2.16
Type of closing				
No closing (Ref.)	–	–	–	–
Conventional	0.29***	0.12 – 0.60	-0.82**	-1.33 – -0.30
Community	1.09	0.46 – 2.55	0.19	-0.54 – 0.93

Fecal human markers				
Concentration fecal human markers (log ₁₀ GE/mL)	1.15**	1.04– 1.26	0.11**	0.04 – 0.18

268

269 **DISCUSSION**

270 In this study, we compared the presence and concentration of pathogenic *Leptospira* in
 271 soils surrounding open and closed sewer sections in six Brazil urban slums. We found that
 272 pathogenic *Leptospira* occurred in both areas but was more prevalent in soils adjacent to open
 273 sewer sections, although the concentration was generally low. More importantly, our results
 274 show that soils in conventionally closed sewers have a reduced presence of the pathogen, as
 275 opposed to community-closed sewers. These results have important implications for future
 276 public health and sewerage development in urban slums.

277 The soil contamination with pathogenic *Leptospira* was lower in soils adjacent to
 278 conventionally-closed sewers than open sections, but no reduction was observed in community-
 279 based closings (Table 1 and Fig. 2). Conventional sewer closings completely canalize sewage,
 280 isolating it from the surrounding environment and preventing spills and overflow during heavy
 281 rainfall events. Since sewage is a recognized source of *Leptospira* as evidenced by this and
 282 previous studies^{16,24,40}, its canalization may eliminate spillage contaminations in soil. However,
 283 the imperfect closure of community-based interventions could still allow sewage to contaminate
 284 adjacent soils. Moreover, despite the reduction observed in conventionally closed sections, the
 285 pathogen could still be detected in 3 of the 4 sites sampled. This suggests that the presence of
 286 pathogenic *Leptospira* contamination in these soils may not have its origin exclusively in the
 287 adjacent open sewers.

288 We identified human fecal markers (BacHum) as a predictor of presence and
289 concentration of pathogenic *Leptospira* in soils. A previous study in streams from Hawaii, also
290 found a positive correlation of pathogenic *Leptospira* concentrations and fecal pollution markers
291 (*Bacteroidales* and *Clostridium perfringens*)⁴¹. Considering that the major sources of human
292 fecal pollution in this environment are open sewers, human fecal markers are likely a surrogate
293 for the distance to the sewer in open sections. However, in closed sections, the correlation of
294 BacHum and pathogenic *Leptospira* indicates that there are other sources of fecal pollution and
295 of pathogenic *Leptospira*. Previous studies have hypothesized that intense rainfall events may
296 mobilize pathogenic *Leptospira* and human pollution markers occurring in soils in higher
297 elevated areas and transport them with the storm run-off to lower areas^{32,42,43}, where sewers are
298 located. Notably, the construction of conventional sewers does not canalize storm water, and
299 thus, run-off may still contribute to the contamination observed in conventionally closed sewer
300 sections. Therefore, the association of pathogenic *Leptospira* and human fecal markers is likely a
301 combination of the effect of sewer proximity and storm run-off. This may explain repeated
302 contamination events in the areas surrounding the sewer, favoring *Leptospira* survival and
303 increasing the risk of exposure of humans and animals to contaminated environments.^{43,44}

304 The concentrations of pathogenic *Leptospira* in the collected soils were generally low in
305 the six urban areas studied (mean $\log_{10} 0.56 \pm 0.8$ and $2.00 - 1.62 \times 10^3$ GEq/g) (Fig 1). This finding
306 is consistent with previous studies that have reported low concentrations of the pathogen in soils
307 and waters in high-risk environments.^{16,24,32,45} Besides the sewer and run-off contribution, the
308 presence and concentration of the pathogen in soil is related to its survival and long-term
309 persistence ability^{8,46} which is affected by the soil type, composition and physicochemical
310 characteristics. For instance, soils rich in nutrients such as iron, manganese, copper and nitrate

311 have been shown to be a positive risk factor for the presence of *Leptospira*, just as wetter soils
312 and basic pH can increase the survival of this pathogen⁴⁷⁻⁴⁹. Interestingly, a soil sample in site 4
313 contained a particularly high concentration of pathogenic *Leptospira* (1.62×10^3 GEq/g), which
314 indicates that hot-spots of the pathogen occur in the urban slum environment. Yet, the highly
315 heterogenic distribution of the pathogen in soil³² and the cross-sectional nature of our sampling
316 strategy may have prevented the identification of these high-concentration areas and determine
317 their origin and temporal dynamics. Since the human infectious dose is still unknown, more
318 studies are needed to determine the significance of these heterogenic distribution of the pathogen
319 in human infection dynamics.

320 Unexpectedly, rat presence and activity were not important factors to predict the presence
321 or concentration of the pathogen in soils. Rats are the main animal reservoir of pathogenic
322 *Leptospira* in urban slums^{5,50,51} and rat presence is commonly reported as a factor for
323 leptospirosis infection.^{35,52} Open sewers offer an ideal ecosystem for the proliferation of rodents
324 by providing burrowing areas, access to water and food sources. Counterintuitively, our results
325 suggest that the contamination of soils close to sewers is more related to the type of sewer
326 closing than to the presence and activity of rats. Therefore, rat control strategies alone such as
327 rodenticide campaigns, may not be effective in reducing the presence of the pathogen in the
328 sewer environment^{53,54} and should be combined with sewerage construction.

329 This study was limited by its cross-sectional design. Because *Leptospira* soil
330 contamination may be variable over time and, specifically, around rainfall events, future
331 prospective studies are needed to investigate the effect of sewer closing in the presence and
332 concentration of the pathogen. In addition, the high heterogeneity of urban slum environments
333 and diversity of community-based closings limit our ability to make wide generalizations of the

334 effects observed in this study. Furthermore, although a higher environmental presence of
335 pathogenic *Leptospira* is intuitively linked to a higher risk of infection, epidemiological studies
336 are needed to determine how sewerage interventions and the reduction of the environmental
337 burden of the pathogen affect the dynamics of leptospirosis infection and disease. These future
338 studies will also need to determine whether community interventions, despite not reducing the
339 environmental burden of *Leptospira*, may still decrease human infection. As community
340 interventions are cheaper and easier to implement in neglected communities, more research is
341 needed to understand their potential role in disease transmission.

342 Despite these limitations, taken together our results suggest that conventional sewer
343 systems may be an important, but not exclusive strategy to reduce the presence and concentration
344 of the pathogen in the environment. The closure of sewers could reduce the niches for the
345 environmental distribution and dissemination of pathogenic *Leptospira* subsequently decreasing
346 pathogenic *Leptospira* exposures in these neglected communities and eventually reducing human
347 leptospirosis. This adds to the body of evidence that sewerage reduces exposures to a wide
348 number of human pathogens and therefore, supports the expansion of sewer systems in urban
349 slums to help decrease the burden of leptospirosis and other environmentally-transmitted
350 infectious diseases.

351

352 **Conflict of interests**

353 The authors declare no competing financial interest.

354

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358

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