

Bound within Boundaries: How Well Do Protected Areas Match Movement Corridors of Their Most Mobile Protected Species?

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Abstract

1. Conserving and managing large portions of land to connect wildlife reserves is increasingly used to maintain and restore connectivity among wildlife populations. Boundaries of such conservation areas are often determined based on expert opinion and socio-political constraints, yet the extent to which they match species' movement corridors is rarely examined. This is mainly due to a lack of data, particularly on wide-ranging movement behavior such as dispersal. Nevertheless, empirically assessing the adequacy of protected areas is key for the implementation of targeted management actions and efficient use of limited conservation funds.
2. Between 2011 and 2019, we collected high-resolution GPS movement data on 16 dispersing African wild dog (*Lycaon pictus*) coalitions from a free-ranging population in the Kavango-Zambezi Transfrontier Conservation Area (KAZA-TFCA). Spanning five countries and 520'000 km² the KAZA-TFCA is the world's largest transboundary conservation area and a prime example for international conservation efforts. We used integrated step selection analysis to estimate habitat preferences of dispersers and to create a permeability surface for the entire KAZA-TFCA. We compared landscape permeability across different regions within the KAZA-TFCA as well as outside its boundaries. Lastly, we calculated least-cost paths and corridors to verify that major movement routes were adequately encompassed within the KAZA-TFCA.
3. Permeability within the boundaries of the KAZA-TFCA was more than double compared to areas outside it. Furthermore, we observed a five-fold permeability difference among the five KAZA-TFCA countries. We further showed that major movement corridors of wild dogs run within the KAZA-TFCA, although some minor routes remained outside formally protected areas.
4. Differences in permeability were mainly caused by different degrees of human activities across regions, which hampered dispersal. Rivers, swamps or open water also limited dispersal, while other landscape features had a limited effect.
5. *Synthesis and Applications:* In this study, we showed how pertinent dispersal data of a highly mobile species can be used to empirically evaluate the adequacy of already-existing or planned protected areas. Furthermore, observed regional differences in landscape permeability highlight the need for a coordinated effort towards maintaining or restoring connectivity, especially where international effort is required.

1 Introduction

2 Connectivity among subpopulations is a crucial pre-requisite for many species to thrive and
3 persist (Fahrig, 2003). Accordingly, preserving and protecting movement corridors between
4 wildlife reserves has become an utmost task for conservation management (Doerr et al., 2011;
5 Rudnick et al., 2012), resulting in an ever-growing number of large and often transboundary
6 protected areas. While boundaries of such areas are often drawn according to expert opin-
7 ion and socio-political needs, subjective assessments have revealed deficiencies in the past
8 (Clevenger et al., 2002; Pullinger and Johnson, 2010). Thus, an empirical assessment of the
9 adequacy of already-existing or planned protected areas using pertinent animal movement
10 data is paramount for targeted use of valuable and scarce conservation funds (Pullinger and
11 Johnson, 2010).

12 In recent years, a growing body of research has used animal relocation data to identify
13 movement corridors and assess connectivity at large scales (e.g. Chetkiewicz et al., 2006;
14 Squires et al., 2013; Elliot et al., 2014). Identification of potential corridors typically relies
15 on the estimation of permeability surfaces, which return the ease or willingness at which
16 the focal species traverses a specific landscape (Sawyer et al., 2011). Such surfaces are
17 created based on a species' habitat preferences, which can be quantified using a suite of
18 selection functions (Zeller et al., 2012). Specifically, habitat preferences are estimated by
19 comparing spatial covariates (e.g. environmental and anthropogenic) at locations visited
20 by the animal to the same spatial covariates at randomly selected locations (Zeller et al.,
21 2012). Importantly, selection functions rely on adequate landscape and relocation data that
22 are representative of the process being studied (Diniz et al., 2019). For instance, relocation
23 data collected on dispersing individuals has been shown to outperform data collected on
24 resident individuals in the detection of large-scale movement corridors (Elliot et al., 2014;
25 Diniz et al., 2019). Nevertheless, dispersal data is inherently difficult to collect and remains
26 scarce in the connectivity literature (Vasudev et al., 2015). As such, most permeability
27 surfaces upon which movement corridors are identified are created using relocation data
28 collected on resident individuals. This introduces severe biases and substantially reduces
29 the power to reveal meaningful movement corridors, for dispersing individuals have different
30 needs and drives compared to resident individuals (Elliot et al., 2014; Cozzi et al., 2020).
31 Such biases have limited our ability to meaningfully assess the effectiveness of protected
32 areas in securing connectivity for their protected species.

33 One initiative that aims at restoring and enhancing connectivity across large scales is the
34 Kavango-Zambezi Transfrontier Conservation Area (KAZA-TFCA), which constitutes the

35 world's largest transfrontier conservation area, spanning over 520'000 km² and five coun-
36 tries (www.kavangozambezi.org). While the KAZA-TFCA was originally set to facilitate
37 movements of African elephants (*Loxodonta africana*; Tshipa, 2017), it is also key to the con-
38 servation of other wide-ranging species such as African wild dogs (*Lycaon pictus*; Woodroffe
39 and Sillero-Zubiri, 2012; Cozzi et al., 2020), lions (*Panthera leo*; Elliot et al., 2014; Cushman
40 et al., 2018), and cheetahs (*Acinonyx jubatus*; Weise et al., 2017). To date, however, few
41 studies have attempted to assess the adequacy of the KAZA-TFCA using relevant global
42 positioning system (GPS) relocation data of its protected species at the appropriate spatial
43 scale (Elliot et al., 2014; Tshipa, 2017). Thus, how well the boundaries of the KAZA-TFCA
44 reflect natural movement patterns and dispersal corridors of its most mobile protected species
45 is virtually unknown.

46 Across the KAZA-TFCA, the African wild dog (*Lycaon pictus*) represents a highly mobile
47 and endangered flagship species for conservation efforts. Once widespread across the entire
48 Sub-Saharan continent, wild dogs have been widely extirpated through human persecution,
49 habitat destruction, and disease outbreaks (Woodroffe and Sillero-Zubiri, 2012). As a result,
50 the species has become one of Africa's most endangered large carnivores, and currently
51 only survives in small, spatially scattered subpopulations (Woodroffe and Sillero-Zubiri,
52 2012). Within these subpopulations, wild dogs form cooperative breeding packs of up to
53 thirty individuals (Creel and Creel, 2002), whose social structure is strongly governed by the
54 process of dispersal (McNutt, 1996; Behr et al., 2020). Both males and females disperse from
55 their natal pack, either alone or in same-sex dispersing coalitions, and search for unrelated
56 mates and a suitable territory to settle (McNutt, 1996; Cozzi et al., 2020; Behr et al., 2020).
57 During dispersal, wild dogs can cover several hundred kilometers (Masenga et al., 2016;
58 Woodroffe et al., 2019; Cozzi et al., 2020). Despite the importance of dispersal for the long-
59 term viability of this species, little empirical information is available on habitat selection and
60 potential movement barriers during dispersal. The few studies that have collected dispersal
61 data have shown that dispersers quickly move over large distances, avoid human-dominated
62 landscapes and areas densely covered by trees, but prefer proximity to water (Masenga et al.,
63 2016; Woodroffe et al., 2019; O'Neill et al., 2020; Cozzi et al., 2020).

64 Here, we collected GPS relocation data on 16 dispersing wild dogs in as many dispersing
65 coalitions from a free-ranging population in northern Botswana and analyzed it to assess the
66 adequacy of the KAZA-TFCA in securing connectivity. We estimated the degree of selection
67 or avoidance for environmental and anthropogenic landscape features and used the obtained
68 habitat preferences to predict a permeability surface spanning the entire KAZA-TFCA. We

69 then investigated how landscape permeability varies regionally and internationally and com-
70 pared permeability within and outside the KAZA-TFCA boundaries. Finally, we calculated
71 least-cost paths and corridors to identify major movement routes and to verify that these
72 are successfully covered by the KAZA-TFCA.

73 **2 Methods**

74 **2.1 Study Area**

75 The study area (centered at -17°13'9"S, 23°56'4"E; Figure 1a) was outlined by a rectangu-
76 lar bounding box stretching over 1.3 Mio km² and encompassing the entire KAZA-TFCA
77 (Figure 1b). The KAZA-TFCA lies in the basins of the Okavango and Zambezi rivers and
78 includes parts of Angola, Botswana, Namibia, Zimbabwe, and Zambia. With a total area of
79 over 520'000 km² it constitutes the earth's largest transboundary conservation area and is
80 characterized by diverse landscapes, including savanna, grassland, and dry or moist wood-
81 land habitats. Rainfall in the study area is seasonal and lasts from November to March.
82 The KAZA-TFCA also comprises the Okavango Delta, which represents a highly dynamic
83 hydrological flood-pulsing system (McNutt, 1996; Wolski et al., 2017). The extent of the
84 flood in the delta greatly changes within and between years depending on the amount of rain
85 that descends from the catchment areas in Angola and reaches the distal ends of the delta
86 between July and August (Figure S4). The flood drastically affects surrounding landscapes,
87 so that during maximum extent (ca. 12'000 km²) the delta becomes a patchy conglomerate
88 of swamps, open water, and islands, whereas these structures run dry when the flood re-
89 tracts to its minimum extent (ca. 5'000 km²; Wolski et al., 2017). Despite 36 national parks
90 (NPs) and other protected areas, there is considerable human influence in some regions of
91 the KAZA-TFCA, mainly originating from farms, high human density, and road traffic.

92 **2.2 GPS Relocation Data**

93 We used a population of free-ranging African wild dogs inhabiting the Okavango Delta in
94 northern Botswana as a source population for dispersing individuals. This population has
95 been extensively studied since 1989 (McNutt, 1996; Cozzi et al., 2013, 2020; Behr et al.,
96 2020). Between 2011 and 2019, we systematically collected GPS relocation data on 16 coali-
97 tions of dispersing African wild dogs (7 female and 9 male coalitions). Candidate dispersing
98 individuals were identified based on criteria reported in Behr et al. (2020), immobilized ac-
99 cording to protocols described in Osofsky et al. (1996), and fitted with GPS/Satellite radio

100 collars (*Vertex Lite*; *Vectornic Aerospace GmbH, Berlin*) while still with their natal pack.
101 All procedures were undertaken and supervised by a Botswana-registered wildlife veterinarian
102 during dispersal, GPS collars were programmed to record GPS relocation data every
103 4 hours and to regularly transmit them via Iridium satellite system to a base station.

104 Because we were interested in dispersal behavior only, we discarded any GPS data col-
105 lected while individuals were still with their natal packs and after settlement in a new
106 territory (Cozzi et al., 2020). We identified the exact time of emigration and settlement
107 based on direct field observations and through visual inspection of the net squared displace-
108 ment (NSD) metric. NSD quantifies the Euclidean distance of a relocation to a reference
109 point (Börger and Fryxell, 2012), which in our case was the center of the dispersing coali-
110 tion’s natal home range. Thus, dispersal was deemed to have started when a coalition had
111 left its natal home range and continued until the NSD metric remained stationary, implying
112 that the coalition had successfully settled (Figure S1). In total, we collected 4’169 GPS
113 relocations during dispersal (Figure S2 & Table S1), resulting in an average of 260 locations
114 per dispersing coalition (min = 37, max = 729). In our analysis, we did not differentiate
115 between male and female dispersing coalitions, for previous research found little differences
116 between sexes during dispersal (Woodroffe et al., 2019; Cozzi et al., 2020).

117 2.3 Spatial Covariates

118 To investigate habitat preferences of dispersing wild dogs, we used a set of geo-referenced
119 covariates (Figure 2) that we aggregated in the categories *land cover* (which included water
120 cover, distance to water, shrubs/grassland cover, and tree cover), *protection status* (pro-
121 tected vs. unprotected), and *human pressure* (which included human influence, presence of
122 roads, and distance to roads). For each of these covariates we prepared spatial raster layers
123 from freely available online services or from remotely sensed satellite imagery. To ensure a
124 consistent resolution (i.e. cell-size or grain) across covariates, we coarsened or interpolated
125 all layers to match a resolution of 250m x 250m. For further details on the preparation of
126 each covariate, see Appendix A.3. We performed processing and manipulation of data as
127 well as all spatial and statistical analyses using R, version 3.6.1 (R Core Team, 2019).

128 2.4 Habitat Selection Model

129 We used an integrated step selection function (iSSF; Avgar et al., 2016) to investigate
130 dispersers’ selection or avoidance for the above-mentioned spatial covariates. That is, we
131 paired each realized step (i.e. the connecting line between two consecutive GPS relocations;

132 Turchin, 1998) with 24 random steps. We generated random steps by sampling turning
 133 angles from a uniform distribution $U(-\pi, +\pi)$ and step lengths from a gamma distribution
 134 that was fitted using realized steps (Avgar et al., 2016). A realized step and its 24 associated
 135 random steps formed a stratum that received a unique identifier. Along each step, we
 136 extracted the above-mentioned covariates (Table S3), standardized extracted values using a
 137 z-score transformation, and checked for correlation using Pearson's Correlation Coefficient
 138 r . None of the covariates were overly correlated ($|r| > 0.6$; Latham et al., 2011) and we
 139 retained all of them for modeling. Our habitat selection model then assumed that dispersing
 140 wild dogs assigned a selection score $w(x)$ of the following exponential form to each realized
 141 and random step (Fortin et al., 2005):

$$w(x) = \exp(\beta_1 x_1 + \beta_2 x_2 + \dots + \beta_n x_n) \quad (\text{Equation 1})$$

142 The selection score $w(x)$ of a step depended on its associated covariates (x_1, x_2, \dots, x_n) , as
 143 well as on the animal's preferences for these covariates $(\beta_1, \beta_2, \dots, \beta_n)$. To estimate habitat
 144 preferences (i.e. the β 's), we used mixed effects conditional logistic regression analysis
 145 as suggested by Muff et al. (2020). We implemented their method using the R-package
 146 *glmmTMB* (Brooks et al., 2017) and used dispersing coalition ID to model random intercepts
 147 and slopes. We defined two movement metrics, namely the cosine of the turning angle
 148 ($\cos(ta)$) and the log of the step length ($\log(sl)$), as core covariates and ran stepwise forward
 149 model selection based on Akaike's Information Criterion (AIC; Burnham and Anderson,
 150 2002) for all other covariates. The inclusion of movement metrics served to reduce biases
 151 in estimated habitat preferences that may have arisen due to movement preferences (Avgar
 152 et al., 2016). To validate the predictive power of the most parsimonious habitat selection
 153 model, we ran k-fold cross-validation for case-control studies as described in Fortin et al.
 154 (2009) (details in Appendix A.5).

155 2.5 Permeability Surface

156 Using the most parsimonious habitat selection model, we predicted a permeability surface
 157 spanning the entire extent of the KAZA-TFCA. That is, we applied Equation 1 to our
 158 spatial covariates and calculated the selection score $w(x)$ for each raster cell. Because
 159 our representation of water was dynamic (to properly render the pulsing behavior of the
 160 Okavango Delta) we collapsed all dynamic water maps into a single static map using areas
 161 that were covered by water in at least 10% of the cases. Using the resulting static map we
 162 also calculated a layer returning the distance to water. To reduce the influence of outliers

163 in predicted permeability scores we followed Squires et al. (2013) and curtailed predicted
164 scores between the 1st and 99th percentile of their original values. To compare permeability
165 across different regions, we normalized the permeability surface to a range between 0 (most
166 impermeable) and 1 (most permeable). We then determined median permeability within
167 and outside the KAZA-TFCA, within and outside formally protected areas, and within each
168 of the five KAZA-TFCA countries.

169 **2.6 Least-Cost Paths and Corridors**

170 To identify movement corridors of dispersing wild dogs, we specified source points and
171 calculated factorial least-cost paths (LCPs) as well as factorial least-cost corridors (LCCs)
172 among them (Elliot et al., 2014). We generated source points by overlaying the study area
173 with a regular grid of points spaced at 100 km. We only considered those points that
174 fell within protected areas > 700 km², which conforms with home-range requirements of
175 African wild dogs reported in Pomilia et al. (2015). Finally, we defined centroids as source
176 points for those protected areas > 700 km² that were not assigned any source points from
177 the regular grid. In total, we generated 68 source points, which resulted in 2'278 unique
178 pairwise combinations and therefore 2'278 unique LCPs and LCCs. We computed factorial
179 LCPs and LCCs between source points using the R-package *gdistance* (for further details see
180 Appendix A.6). After computation, we tallied overlapping LCPs and LCCs, respectively,
181 into single connectivity maps.

182 **3 Results**

183 **3.1 Habitat Selection Model**

184 Our most parsimonious habitat selection model ($\Delta AIC > 2$ than any alternative model;
185 Table S4) retained the covariates *water*, *distance to water*, *trees*, *shrubs/grassland*, and
186 *human influence*, beside the fixed covariates *cos(ta)* and *log(sl)* (Figure 3a). Parameter
187 estimates showed that dispersing wild dogs moved in a directional and fast manner, as
188 indicated by a positive selection for small turning angles, i.e. high *cos(ta)* ($\beta = 0.14$; 95%
189 CI = 0.07 to 0.21) and longer steps, i.e. high *log(sl)* ($\beta = 0.06$, 95% CI = 0.02 to 0.09).
190 Dispersers avoided moving through water ($\beta = -0.52$, 95% CI -0.77 to -0.26) but selected
191 for locations in its vicinity, although the latter effect was not significant ($\beta = -0.32$, 95% CI
192 = -0.72 to 0.08). Dispersers avoided areas that were densely covered by trees ($\beta = -0.31$,
193 CI = -0.46 to -0.15) and preferred areas covered by shrubs/grassland ($\beta = 0.25$, 95% CI =

¹⁹⁴ 0.07 to 0.42). Finally, dispersers avoided areas that were influenced by humans ($\beta = -0.41$,
¹⁹⁵ 95% CI = -0.78 to -0.05).

¹⁹⁶ Results from the k-fold cross-validation suggested that our prediction was significant and
¹⁹⁷ robust, as highlighted by the fact that the 95%-CIs intervals of $\bar{r}_{s,realized}$ and $\bar{r}_{s,random}$ did
¹⁹⁸ not overlap (Figure 3b). Likewise, the significant correlation between ranks and correspond-
¹⁹⁹ ing frequencies for realized steps suggested a good fit between predictions and observations
²⁰⁰ (Figure 3b).

²⁰¹ **3.2 Permeability Surface**

²⁰² Our prediction of landscape permeability revealed substantial differences across regions in
²⁰³ the study area (Figure 4). Comparisons of median permeability values (Table 1) showed
²⁰⁴ that permeability inside the KAZA-TFCA is more than two times as high as permeability
²⁰⁵ outside it. Permeability varies by country, with a five-fold permeability difference among
²⁰⁶ them. Angola and Botswana are characterized by comparably highly permeable landscapes,
²⁰⁷ Zimbabwe and Zambia are relatively impermeable, and Namibia ranges in between the two
²⁰⁸ extremes (Table 1). Visual inspection of our covariate layers indicated that high permeability
²⁰⁹ in Angola and Botswana is mainly caused by a combination of low human influences, low
²¹⁰ tree cover, high shrubs/grassland cover, and a close distance to water. Although swamps,
²¹¹ wetlands, and permanent water themselves provide little permeability, their surroundings
²¹² act as strong attractants to dispersers. The low permeability that characterizes Zambia and
²¹³ Zimbabwe, on the other hand, is mainly caused by substantial human influences. Albeit the
²¹⁴ KAZA-TFCA covers most permeability hot-spots, several highly permeable regions remain
²¹⁵ uncovered by its borders. Across all countries, protected areas provide roughly double the
²¹⁶ permeability of unprotected landscapes (Table 1).

²¹⁷ **3.3 Least-Cost Paths & Least-Cost Corridors**

²¹⁸ Our least-cost analysis revealed three major movement corridors of which all were well-
²¹⁹ contained within the KAZA-TFCA boundaries (Figure 5). One major corridor runs SE-NW
²²⁰ and connects the Okavango-Linyanti ecosystem in Botswana with Luengue-Luiana NP in
²²¹ Angola. A second corridor runs W-E between Chobe NP in Botswana and Zimbabwe's
²²² Hwange NP. A third major corridor runs NE-SW, completely across unprotected areas, and
²²³ connects Kafue NP in Zambia with more central regions of the KAZA-TFCA. Several minor
²²⁴ corridors branch off from these three major corridors; these include a southward connection
²²⁵ between the Okavango-Linyanti and the Central Kalahari Game Reserve, a southwesterly

226 corridor connecting Luengue-Luiana NP with Namibia's Khaudum NP, and a northeasterly
227 extension of the Hwange corridor into Zimbabwe's Matusadona NP. According to our predic-
228 tions, the landscapes in the Okavango-Linyanti region are the highest frequented dispersal
229 routes within the KAZA-TFCA (Figure 5b). Our model did not detect any significant direct
230 corridors between Zimbabwe and Zambia or Zambia and Angola, and only a very limited
231 W-E direct connection between the Okavango region and Namibia's Khaudum NP. Except
232 for the corridor into the Central Kalahari National Park, our model did not detect any
233 significant connectivity outside the boundaries of the KAZA-TFCA. Furthermore, we found
234 little to no direct connectivity between peripheral points; that is, most paths and corridors
235 connecting two adjacent peripheral points run through more central regions before heading
236 towards their destination at the periphery (Figure 5).

237 4 Discussion

238 We used GPS relocation data collected on dispersing African wild dogs to investigate whether
239 their main movement corridors are contained within the boundaries of the world's largest
240 transboundary conservation area, namely the KAZA-TFCA. Our analysis suggests that the
241 KAZA-TFCA indeed encompasses all major corridors of African wild dogs, demonstrating
242 the potential value of such an initiative. We thus exemplified how pertinent dispersal data
243 of a highly mobile species can be used to assess the adequacy of already existing or planned
244 protected areas. Our approach is neither limited to the African wild dog, nor to our study
245 area and thus applicable to any study system. All covariates used throughout this study are
246 readily available on a global scale and many of them are likely to be important determinants
247 of movement behavior, landscape permeability, and connectivity for other species (Thurfjell
248 et al., 2014; Zeller et al., 2012). Interestingly, our predicted network of least cost-paths and
249 corridors for African wild dogs shows surprising similarities to corridors of dispersing lions
250 inhabiting the same ecosystem (Elliot et al., 2014; Cushman et al., 2018). This not only
251 reinforces confidence in our own predictions but also suggests potential synergies for the
252 conservation of these two, and possibly more, species. Expanding our analytical framework
253 to additional species will likely yield important insights on the consistency of inter-specific
254 movement corridors, thus highlighting areas that are exceptionally valuable for the conser-
255 vation of several species.

256 Our results emphasize that human influences constitute some of the main barriers to
257 connectivity among wild dog populations. This conforms to findings on dispersing wild
258 dogs from eastern Africa (Masenga et al., 2016; O'Neill et al., 2020) but conflicts with

259 findings from South Africa by Davies-Mostert et al. (2012), who reported a high willingness
260 of dispersers to cross human-dominated landscapes. We believe that such differences are due
261 to the unavailability of alternative routes through natural landscapes, which may have forced
262 dispersers in South Africa to cross human dominated landscapes despite a strong aversion to
263 do so. In this regard, our representation of dispersal corridors and the resulting connectivity
264 appear conservative, as dispersers may be able to make the best out of a bad situation and
265 cross landscapes characterized by considerably unfavorable conditions (Palomares et al.,
266 2000; Elliot et al., 2014). Nevertheless, successful conservation of this species relies on
267 policymakers' and local authorities' willingness and ability to provide and conserve natural
268 areas that remain free from anthropogenic pressures. This is not only paramount in light
269 of increasing connectivity and facilitating dispersal, but also in terms of reducing human-
270 caused mortality during dispersal. In fact, previous studies have shown that human-caused
271 mortality represents a major threat to wild dogs' ability to disperse (Woodroffe et al., 2019;
272 Cozzi et al., 2020).

273 Besides human influences, we identified water as additional obstacle to dispersal. This
274 corroborates earlier studies showing that water bodies are almost impenetrable to resident
275 packs (Abrahms et al., 2017) and only infrequently crossed by dispersing individuals (Cozzi
276 et al., 2020). An accurate and dynamic representation of water is thus imperative and
277 particularly relevant in seasonal or flood-pulsing ecosystems such as the Okavango Delta.

278 Although dispersers avoided moving through water, they selected locations in its vicinity.
279 This preference may be caused by the occurrence of prey close to water (Bonyongo, 2005).
280 For the same reason, however, competitors such as lions, spotted hyenas, and resident wild
281 dogs may also use areas close to water (Valeix et al., 2010), thereby occasionally forcing
282 dispersing wild dogs to move into prey-poorer areas away from water. Given the influence
283 that resident conspecifics, competitors, and prey can have on dispersers (Cozzi et al., 2018;
284 Armansin et al., 2019) future studies should strive to collect and incorporate intra- and
285 interspecific relationships into analyses of landscape connectivity.

286 Locally, we identified the Okavango-Linyanti region as a potential dispersal hub through
287 which dispersing wild dogs gain access to more peripheral regions of the KAZA-TFCA. It
288 appears that the absence of human activities, the central position within the KAZA-TFCA,
289 and the presence of relatively impermeable water bodies (e.g. Okavango Delta, Linyanti
290 Swamp) funnel dispersal movements, resulting in a highly frequented corridor. The key
291 role of the Okavango-Linyanti region for overall connectivity within the KAZA-TFCA thus
292 calls for actions to secure its protection status in the future. While the region is currently

293 a Wildlife Management Area, it has neither the status of a National Park nor that of a
294 Game Reserve. A similar case of non-formally protected but key dispersal landscape is
295 represented by the area south of Kafue NP in Zambia, for which a disruption of its main
296 and narrow dispersal corridor would result in considerable isolation of its subpopulations.
297 We also revealed a potential southwards corridor between the Okavango-Linyanti ecosystem
298 and the Central Kalahari National Park. Elliot et al. (2014) and Cushman et al. (2018)
299 identified a similar corridor for dispersing lions, suggesting that upholding and protecting a
300 link between those ecosystems is pivotal. Some areas through which the corridor runs are
301 neither part of the KAZA-TFCA nor profit from any form of protection status. In fact,
302 human presence and activities along the national road that longitudinally traverses this
303 corridor may limit realized connectivity (Cozzi et al., 2020).

304 Our approach of identifying movement corridors based on pre-defined start and end
305 points implicitly assumes that individuals know the end point of their dispersal journey and
306 that they have almost complete knowledge of associated movement costs (Panzacchi et al.,
307 2016). Since dispersers often move into unknown territory, this may not necessarily be the
308 case (Abrahms et al., 2017; Cozzi et al., 2020). However, specification of pre-defined end
309 points might not be necessary, as the parametrized iSSF model can be used as mechanistic
310 movement model to simulate dispersal events from known source points, yet without re-
311 stricting the domain of potential end points (Signer et al., 2017). Consequently, movement
312 corridors would emerge more naturally as the result of a myriad of simulated dispersal events.
313 While a simulation-based approach is conceptually straightforward, it requires a comprehen-
314 sive mechanistic understanding of dispersal movements, which is conditional on our ability
315 to collect additional dispersal data and adequately represent the landscape through which
316 individuals move.

317 Our work shows how dispersal data of a highly mobile species can be used to identify
318 movement corridors and to assess the adequacy of protected areas. In our case, the predicted
319 movement corridors of African wild dogs were well contained within the boundaries of the
320 world's largest transboundary conservation area, namely the KAZA-TFCA, suggesting that
321 it will significantly contribute to the long-term viability of this species. Moreover, our
322 connectivity network allowed revealing potential dispersal hubs through which dispersers
323 gain access to more remote regions of the study area. Finally, our investigations showed
324 that human influence constitutes one of the main barriers to dispersal and substantially
325 reduces landscape connectivity. Successful conservation of wide-ranging species, such as
326 exemplified by the African wild dog, will therefore be contingent on the willingness of local

327 authorities, policymakers, and land managers to preserve areas that remain free from human
328 strains. Ultimately, our work contributes to the growing field of connectivity studies and
329 provides and easily expandable framework for assessing the adequacy of already-existing or
330 planned protected areas.

331 **5 Authors' Contributions**

332 D.D.H., D.M.B., A.O. and G.C. conceived the study and designed methodology; D.M.B.,
333 G.C., and J.W.M. collected the data; D.D.H. and D.M.B. analysed the data; G.C. and A.O.
334 assisted with modelling; D.D.H., D.M.B., and G.C. wrote the first draft of the manuscript
335 and all authors contributed to the drafts at several stages and gave final approval for pub-
336 lication.

337 **6 Data Availability**

338 GPS movement data of dispersing coalitions will be made available on dryad at the time of
339 publication.

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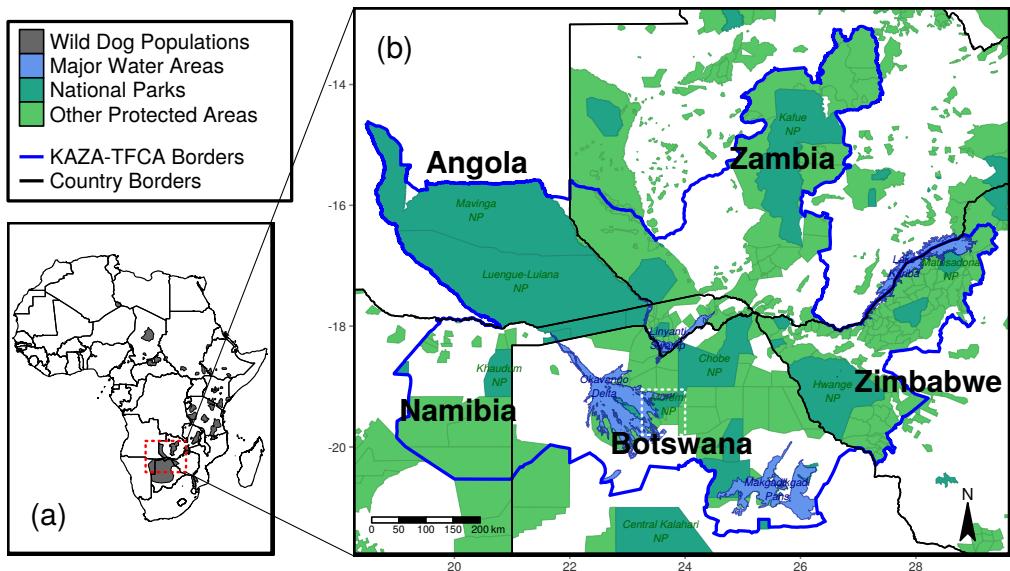


Figure 1: Overview of our study area. (a) The red dotted rectangle depicts the study area, which was confined by a bounding box encompassing the entire KAZA-TFCA. Gray areas indicate remaining wild dog populations according to the IUCN (Woodroffe and Sillero-Zubiri, 2012). (b) The white rectangle illustrates the area within which dispersing coalitions were collared. Since Game Reserves in Botswana virtually serve the same purpose as National Parks, we use the terms interchangeably for this region.

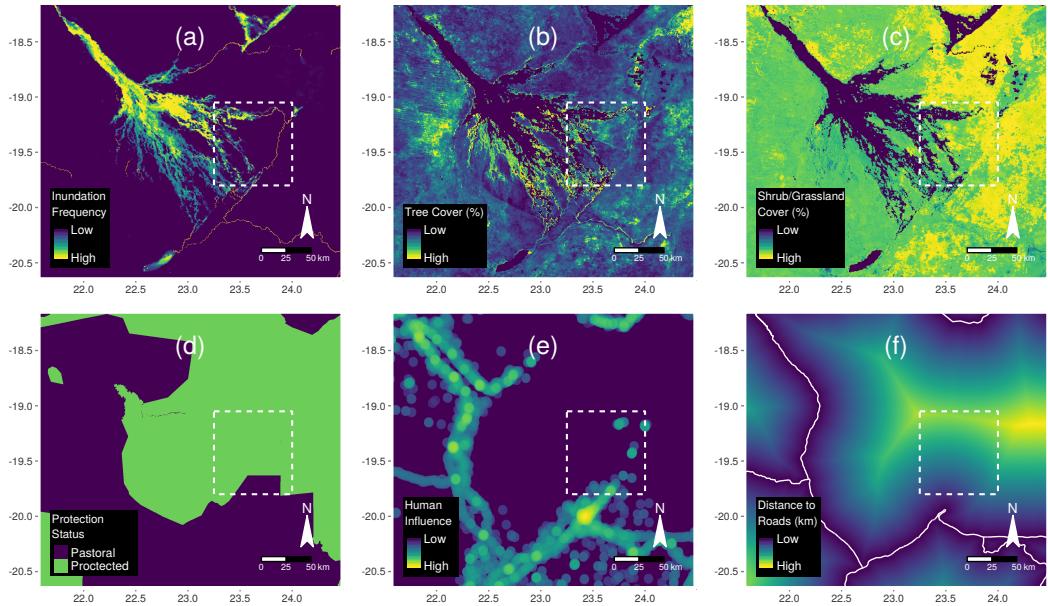


Figure 2: Overview of spatial covariates that we included in our models. We prepared all covariates for the entire study area but for better visibility we only plot them for the surroundings of the Okavango Delta. The white rectangle in each plot depicts the area within which dispersing coalitions were collared. (a) Averaged layer of all dynamic (binary) water maps. (b) Percentage cover of trees. (c) Percentage cover of shrubs/grassland. Anything that was not covered by trees or shrubs/grassland was deemed to be bare land. (d) Protection status of the area. (e) Human influence proxy composed of human density, farms, and roads. (f) Distance to nearest road (white lines depict actual roads).

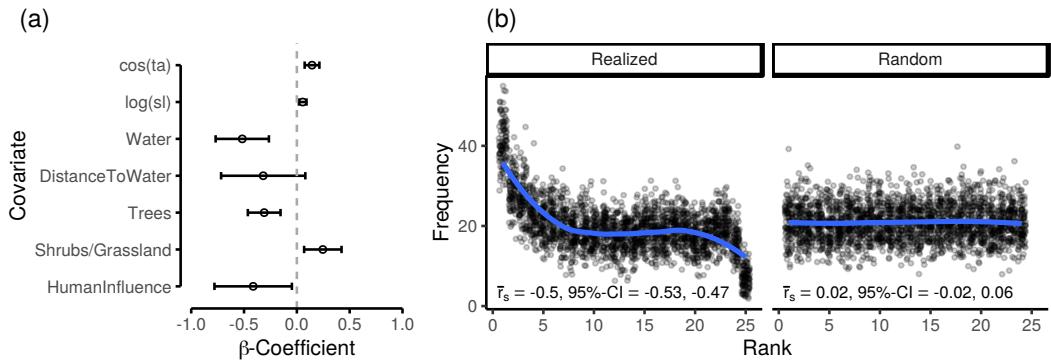


Figure 3: (a) Estimated selection coefficients from the most parsimonious habitat selection model. Negative coefficients indicate avoidance of a covariate, positive coefficients selection of a covariate. Whiskers delineate the 95%-CIs for estimated parameters. (b) Results from the k-fold cross validation for case-control studies. The left graph shows rank frequencies of *realized* steps according to predictions, whereas the right graph shows rank frequencies of *randomly selected* steps according to predictions. \bar{r}_s indicates the mean correlation coefficient resulting from 100 repetitions of the k-fold cross validation. The blue smoothing line was fitted using a locally weighted polynomial regression and serves to aid the eye in detecting the trends. Correlation coefficients suggest that our prediction was significant and robust, evidenced by the fact that the confidence intervals of $\bar{r}_{s,realized}$ and $\bar{r}_{s,random}$ did overlap and by the fact that there was strong and significant correlation between ranks and associated frequency for realized steps.

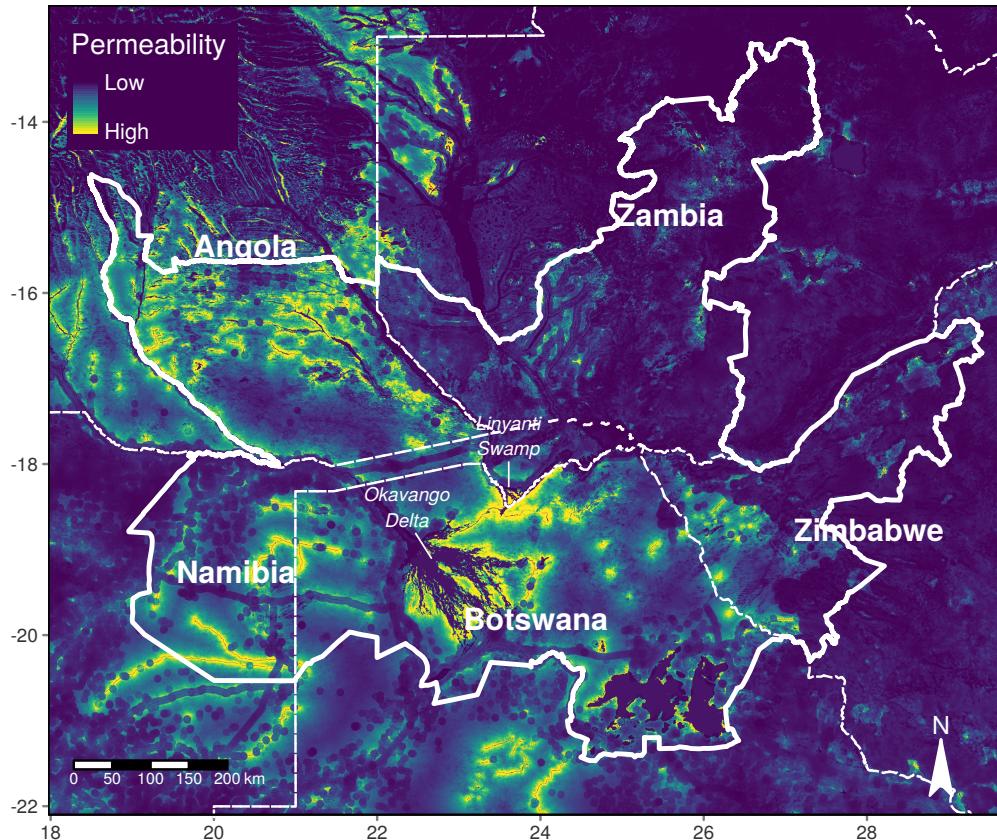


Figure 4: Predicted permeability surface for the extent of the KAZA-TFCA. Permeability was predicted by calculating selection scores $w(x) = \exp(\beta_1 x_1 + \beta_2 x_2 + \dots + \beta_n x_n)$ for each raster cell based on the raster cell's underlying covariates (x_i) and estimated habitat preferences (β_i). Areas that dispersers find easy to traverse are depicted in bright colors. Bold white lines delineate the borders of the KAZA-TFCA, whereas dashed white lines show country borders.

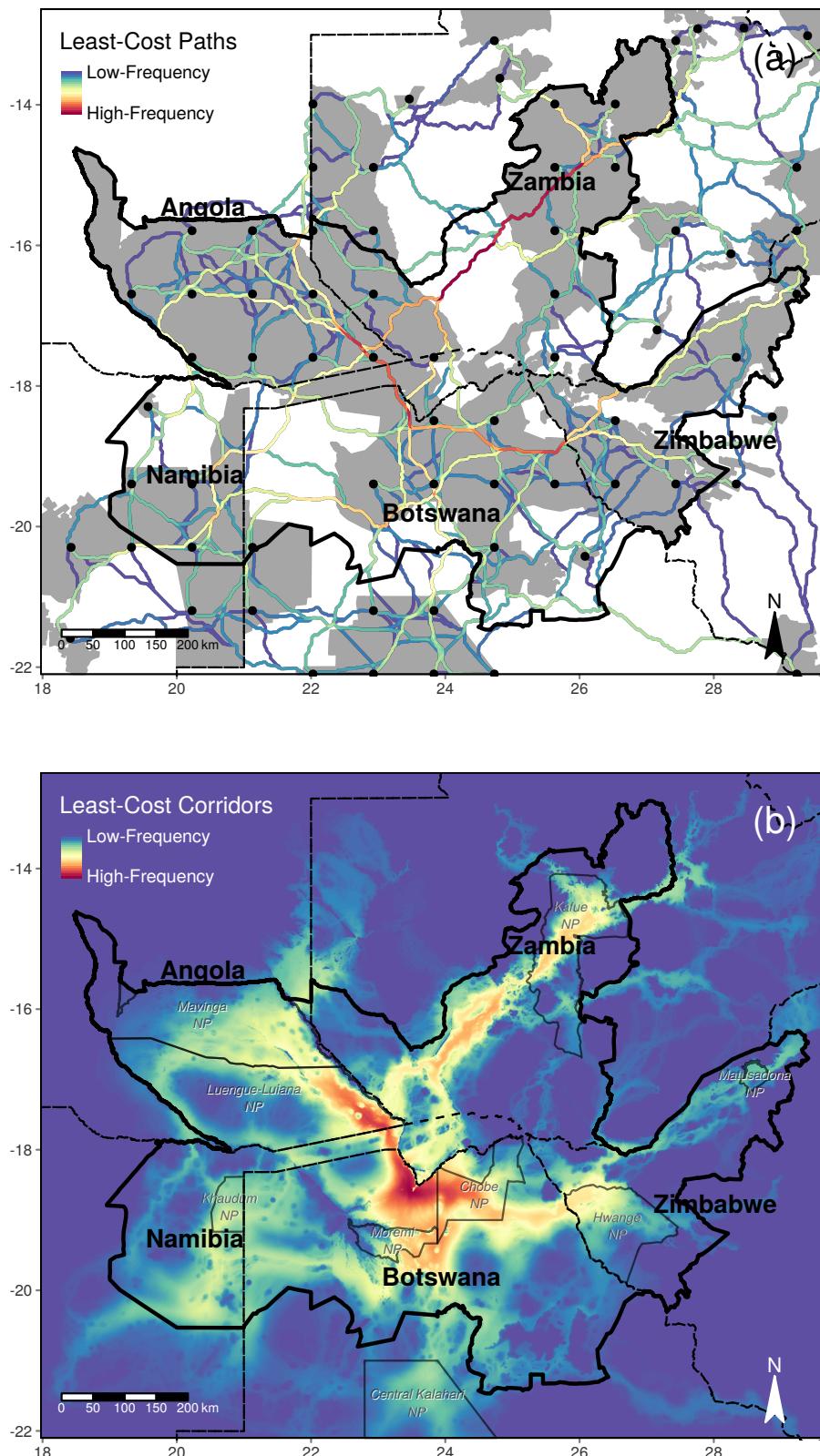


Figure 5: (a) Source points (black dots) and corresponding least-cost paths leaving from protected areas (light grey). Note that only contiguous protected areas covering more than 700 km² are depicted. Continuous thin black lines indicate the borders of the KAZA-TFCA, whereas dashed black lines delineate country-borders. (b) Least-cost corridors between the same source points as illustrated in subfigure (a). For ease of spatial reference, we also labeled some national parks (NPs, in dark-grey).

Table 1: Comparison of median permeability (interquantile range in brackets) across countries, separated into areas within and outside the KAZA-TFCA, as well as within and outside formally protected areas. High values indicate high permeability, whereas low values correspond to low permeability.

Country	KAZA-TFCA		Protection Status		
	Inside	Outside	Protected	Pastoral	Overall
Angola	0.36 (0.41)	0.12 (0.32)	0.36 (0.41)	0.12 (0.33)	0.20 (0.39)
Botswana	0.25 (0.30)	0.15 (0.16)	0.28 (0.35)	0.15 (0.18)	0.19 (0.25)
Namibia	0.22 (0.30)	0.13 (0.18)	0.24 (0.30)	0.11 (0.15)	0.16 (0.24)
Zambia	0.05 (0.09)	0.03 (0.06)	0.04 (0.10)	0.03 (0.05)	0.03 (0.07)
Zimbabwe	0.07 (0.16)	0.06 (0.05)	0.08 (0.17)	0.05 (0.05)	0.06 (0.07)
Overall	0.16 (0.30)	0.07 (0.15)	0.15 (0.30)	0.07 (0.15)	0.10 (0.22)