

# Rango de Hogar y uso de Hábitat de Carpinchos en Pastizales recién invadido en el Chaco Seco de Paraguay

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**Introduction:** Deforestation is one of the primary causes of species decline worldwide. Some species, however, can take advantage of change and expand their distribution, putting endemic species in danger of extinction, changing the composition of biotic communities, and altering ecosystems.

**Methods:** From June 2005 to July 2007, using radiotelemetry collars and implants we investigated home range, habitat use and body temperature dynamics of six capybaras (*Hydrochoerus hydrochaeris*). The study took place in a deforested area of the Central Dry Chaco region of Paraguay recently invaded by the species.

**Results:** The results show that the average home range of capybaras was 183 ha, similar to reported home ranges of capybaras in the Brazilian Pantanal. Within the study area, capybaras used and selected water significantly more than Chaco forest and pasture land, and had lower than expected use of shrub forest. Overall, capybaras were located 95% of the time < 500 m from permanent water, with the greatest distances from water occurring at 3:00 and 5:00 AM. Average year round body temperature was 36.15 °C. There was a significant positive correlation between body temperature and distance from water, and a significant negative correlation between distance from water and Chaco ambient temperature.

**Discussion:** These findings suggest that capybaras use water to thermoregulate. The results show how anthropogenic habitat modification has allowed capybaras to thrive in a harsh xeric environment, and assists us in understanding how capybaras can expand into deforested areas. This study provides valuable information for the future management of the species in the Dry Chaco region of Paraguay.

**Key words:** Capybara, Chaco region, habitat use, home range, invasive species, thermoregulation.

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## Resumen

La deforestación es una de las principales causas de la desaparición de especies en todo el mundo. Algunas especies, sin embargo, pueden aprovechar los cambios y ampliar su distribución, poniendo a las especies endémicas en peligro de extinción, cambiando la composición de las comunidades bióticas y alterando los ecosistemas. Entre junio de 2005 y julio de 2007 se determinó el rango de hogar, el uso del hábitat y la temperatura corporal de seis carpinchos (*Hydrochoerus hydrochaeris*), a través de collares e implantes de radiotelemedría en una área deforestada de la región del Chaco Central del Paraguay invadida recientemente por carpinchos. Nuestros resultados muestran que el rango de hogar promedio de los carpinchos fue de 183 ha, similares a los descritos en el Pantanal brasileño. Dentro del área de estudio los carpinchos utilizaban el agua significativamente más que su disponibilidad, seguido del bosque chaqueño y tierras de pastoreo, y utilizaban en un rango menor de lo esperado los malezales. En general, los carpinchos se encontraban el 95% del tiempo a < 500 m de fuentes permanentes de agua, con las mayores distancias del agua a las 3:00 y las 5:00 hs. El promedio anual de la temperatura corporal fue de 36.15 °C. Se encontró una correlación positiva significativa entre la temperatura corporal y la distancia de la fuente de agua, y una correlación negativa significativa entre la distancia del agua y la temperatura ambiente del Chaco. Estos hallazgos sugieren que el carpincho efectivamente utiliza el agua para regular su temperatura. Nuestros resultados muestran cómo la modificación del hábitat por el hombre ha permitido al carpincho prosperar, y nos ayudan a entender cómo pudieron invadir y utilizar las áreas deforestadas. Este estudio proporciona información valiosa para el manejo futuro de la especie en la región del Chaco Seco de Paraguay.

**Palabras clave:** Carpincho, Chaco, especies invasoras, hábitat, rango de hogar, termorregulación.

## Introduction

Anthropogenic habitat destruction and land fragmentation are the main causes of biodiversity loss worldwide (National Research Council 2001; Fahrig 2003), followed by the introduction and expansion of exotic species placing endemic species at risk of extinction (Aguirre and Tabor 2008), changes in the composition of biotic communities, and altered ecosystem services (Vitousek *et al.* 1997; Lockwood *et al.* 2007). However, not only can exotic species benefit from land use change, native species can take advantage and expand into ecosystems where they historically did not occur. The expansion of native species has been driven by land use and land management changes such as agriculture, game exploitation, predator and poaching controls (Acevedo *et al.* 2006). For example, coyotes (*Canis latrans*) were historically restricted to central North America, but have expanded and colonized most of the North American continent in less than two centuries (Fener *et al.* 2005). Many more native species are known to have taken advantage of anthropogenic land modification and expanded their range (coypu, Guichón *et al.* 2003; Steven *et al.* 2004).

Capybara (*Hydrochoerus hydrochaeris*) is the world's largest living rodent with an average weight of 50 kg (Mones and Ojasti 1986; Nowak 1991). They are semi-aquatic

mammals, extensively distributed throughout most of South America, from Colombia south into central Argentina, Paraguay and Uruguay (Redford *et al.* 1990; Ojasti and Robinson 1991). Capybaras inhabit open flooded grasslands as well as densely vegetated areas around permanent water sources such as ponds, rivers, marshes, and swamps.

Although closely associated with aquatic environments, capybaras use water primarily as a place of refuge, and most normal activity is on land (Schaller 1976). Nevertheless, year-round water is essential for capybaras to be present in an area. Capybaras live in herds ranging from a pair to complex groups of several adults of both sexes and their offspring (Azcarate 1980; Schaller and Crawshaw 1981; Mones and Ojasti 1986; Herrera 1992). These giant rodents are efficient grazing herbivores, feeding primarily on aquatic plants and grasses that grow in or near water (Gonzalez and Parra 1972; Macdonald 1981; Lord and Lord 1988; Quintana and Malvarez 1994, 1998; Barreto and Herrera 1998). Capybaras have been documented to share grazing areas with livestock and it is not uncommon for populations of capybaras to benefit from pasture management, predator control and provisioning of water by ranchers (Mones and Ojasti 1986; Ojasti and Robinson 1991; Quintana 2003). In certain regions of Brazil, capybaras have become an agricultural pest due to changes in habitat as a result of ranching and agriculture (Paschoaletto *et al.* 2003).

Capybara ecology, home range and habitat use have been previously studied throughout the core of their distribution, in habitats such as tropical forest and seasonally flooded plains where the species is locally abundant (Cordero and Ojasti 1977, 1981; Alho *et al.* 1987; Herrera and Macdonald 1989; Lord 1991; Herrera 1992; Quintana and Malvarez 1994; Barreto and Herrera 1998; Quintana 2002, 2003). By contrast, our study area is in the central Gran Chaco ecosystem of Paraguay at the periphery of its historical distribution. Recent deforestation in this region has facilitated the expansion of this species from more mesic habitat into the central Chaco (Campos-Krauer and Wisely 2011), where populations densities are low (Brooks 1998).

The central Chaco is dominated by thick tropical xeric thorn forest characterized by the trees Quebracho Blanco (*Aspidosperma quebracho-blanco*) and Guaimipire (*Ruprechtia triflora*), cacti and bromeliads (Holdridge 1969; Gorham 1973; Bucher 1982; Redford *et al.* 1990; Glatzle 1999; Mereles 2013). It experiences large variations in temperature between summer and winter with temperature during the summer reaching highs of 47 °C, and below freezing during winter; with an annual mean temperature of 24 °C to 25.5 °C (Savaria-Toledo 1993). Annual precipitation ranges between 600 and 1000 mm y (Gorham 1973, Adámoli *et al.* 1990). As much as 80% of the annual precipitation occurs from November to May (Eidt 1968). Precipitation and standing water diminish greatly or even disappear completely during the colder dry season that extends from June to October (Hueck 1966; Gorham 1973).

Due to these harsh characteristics of the Paraguayan Chaco ecosystem, it has remained mostly undeveloped and immune to modern anthropogenic land modification until recent years. The rate of deforestation has dramatically increased since the 1970s (Benirschke *et al.* 1989). Currently, approximately 1,000 ha of native forest are cleared daily and transformed into pastureland, with an estimated 3.5 M hectares cleared in the last decade (Huang *et al.* 2009; Caldas *et al.* 2013). To support a growing cattle population, approximately one pond is constructed for every 100 ha of pasture to

maintain year-round water. The conversion of dense xeric thorn forest to pasturelands and the availability of year-round water supply have greatly altered the resource potential of the ecosystem.

The Chaco region is an area in which capybaras have been present for thousands of years. However, much of the best habitat for capybaras is located in the southern Humid Chaco region along the shorelines of rivers and in marshes. It is clear that capybaras are taking advantage of land use change to expand their range to the central Chaco region (Campos-Krauer and Wisely 2011). Understanding the factors and mechanisms that have allowed this expansion requires a detailed understanding of how capybaras colonize areas on a fine scale. To this end, we examined movement patterns, habitat use, and thermal ecology in several groups of capybaras in the central Chaco region to better understand the ecological drivers of the species' range expansion. In this paper, we describe and compare home range and habitat use of capybaras in the Chaco and compare our results with observations in other studied regions. This information will be valuable to forecasting future distributions and vulnerabilities in light of human activities.

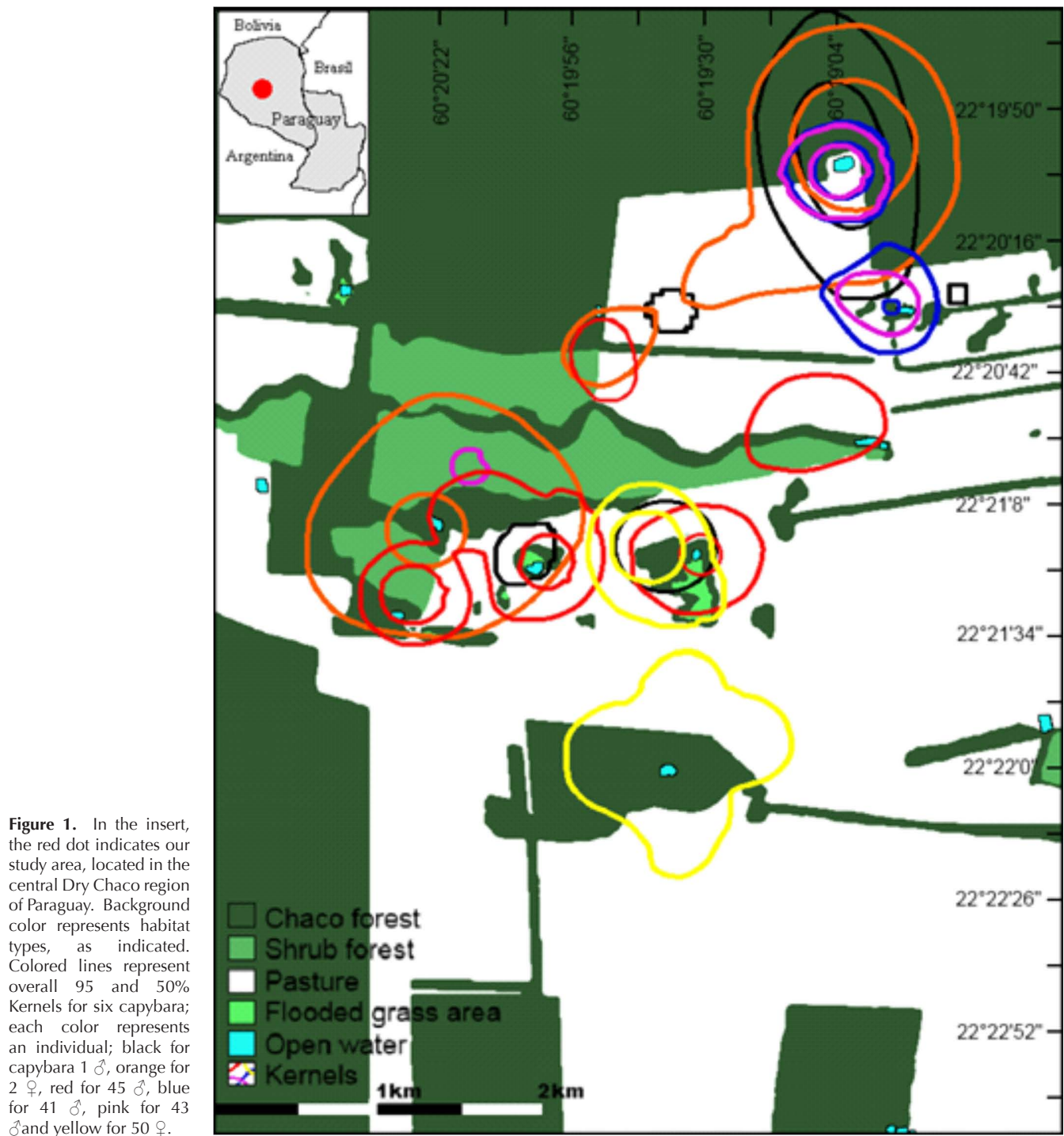
## Material and Methods

**Study area.** The research was carried out from June 2005 to October 2007, at Fortín Toledo, Boquerón Dept., Paraguay (-22°21'05" N, -60°19'35" W; Fig. 1). The study was carried out in and around a biological reserve of approximately 180 ha surrounded by private ranches which add up to a total area of 3,500 ha. Land cover in the study area consisted primarily of open water plus four vegetation types: 1) Chaco forest is virgin tropical xeric thorn forest characterized by thick understory vegetation with an abundance of bromeliads (*Bromelia* spp. and *Dyckia* spp.). Trees are less than 15 m tall with many spines, dominated by tree species such as quebracho (*Schinopsis quebracho-colorado*), bottle tree (*Chorisia insignis*), palo santo (*Bulnesia sarmientoi*), mistol (*Zizyphus mistol*) and verde olivo (*Cercidium praecox*). This habitat is distributed across the landscape as remnant forest islands surrounded by pasture or secondary growth. 2) Shrub Chaco forest consists of secondary forest regrown from 25 year-old, introduced pastures. This habitat is characterized by a wide variety of thorny shrubs, dominated by *Mimosa* spp., *Acacia* spp. *Prosopis alba*, and *P. nigra*. Cacti are also diverse and abundant, with taxa such as *Opuntia*, *Cleistocactus* and several tree cacti such as *Cereus* spp. Within this habitat there is little to no understory vegetation, due mostly to soil compaction. 3) Exotic pasture is cleared forest characterized by non-native grasses dominated by gatton panic (*Panicum maximum*) and to a lesser extent star grass (*Cynodon dactylon*). These areas have few trees remaining and are divided into large paddocks in which cattle are maintained year-round with a water source. 4) Seasonally flooded grassy areas, small wetlands which occur as natural depressions, which will hold water during the rainy season, but dry out during the dry season. These areas have a mix of introduced and native grass species (*Cynodon* spp.) together with palm trees (*Copernicia alba*). In addition to natural wetlands, areas surrounding man-made ponds had a high abundance of grass.

Areas of open water from man-made ponds were present in each pasture paddock in the study area. Each reservoir was at least 625 m<sup>2</sup> and 1.5 m in depth and many were covered by aquatic plants such as *Eichhornia crassipes* and *Pistia stratiotes*. Ponds were



generally built without modifying the surrounding forest, which was left to prevent eolic sand deposits and to provide shade for cattle. These forest patches and ponds remained as islands most of the time surrounded by pasture. During the dry season, standing water was found only in these man made ponds.



*Capture and radiotelemetry.* From June 2005 to August 2007, we located and monitored a total of 6 capybaras during different lengths of time (Table 1). Radio collars (Telonics, Inc., Mesa, Arizona 85202, USA) were placed on four capybaras (2 adult males, 2 adult

females) and intra-abdominal implants with temperature sensors (Telonics, Inc) were placed in two capybaras (1 adult male and 1 juvenile male). The small sample size reflects the low density and difficulty of capture of capybaras in this study area and is representative of the patchily distributed populations within this region.

Animals were trapped with box-style live traps and chemically immobilized using a combination of Ketamine HCl (Ketalar, 4.7 mg/kg), and Tiletamine HCl/Zolazepam HCl (Telazole, 1.17 mg/kg; Kreeger *et al.* 2002). Drugs were delivered intramuscularly by a blowgun (Telinject, U.S.A., Inc., Agua Dulce, CA) using 3 cc plastic darts (Telinject) or a standard 3 cc pole syringe. Animals were considered fully immobilized when they did not respond to external stimulus. Each animal was sexed, weighed and marked with ear-tags. All captured animals received a general examination to evaluate body condition, external parasites and possible wounds. Individuals were classified as juvenile (less than one year old) if < 25 kg, and animals > 25 kg were considered to be at least one year old, and classified as young adults or adults (Ojasti 1971). Each transmitter was equipped with a mortality sensor which activated when animals had not moved for > 6 hours. All procedures were in compliance with the Secretaría del Ambiente, Paraguay; Kansas State University Institutional Animal Care and Use Committee (Protocol No. 2362); and guidelines approved by the American Society of Mammalogists (Gannon *et al.* 2007).

Independent telemetry locations (White and Garrott 1990) were estimated using triangulation with a minimum of three vectors per location. We marked and georeferenced > 50 fixed reference locations every 500 m in the study area from which a vector bearing was collected. Vehicles were used to move between reference locations to reduce triangulation time. Three element folding yagi antenna (Advanced Telemetry Systems, Inc., Isanti, Minnesota), attached to a 3 m long aluminum pipe were used to maximize signal reception. Locations were estimated using a minimum of three vectors by using the maximum likelihood estimation option (Lenth 1981) in program LOAS (LOAS™ 2005, Ecological Software Solutions LLC). We calculated an average location error from reference collars placed in random locations approximately 1 m from the ground. Animals were located at six hour intervals with the location time moving ahead one hour daily to cover a 24 hour period every 12 days. Readings were taken from capture until the animal was no longer located or died (Table 1).

Observers were able to detect animal movement due to differences in signal strength during data collection at a single vector. We recorded movement patterns from two capybaras (capy 41 and 43) with internal implants using this approach. Direct observations ( $n = 193$ ) and patterns of signal reception other than a consistent pulse intensity were considered to be an moving animal. These data reflected the time of day during which animals were moving. Movement patterns were analyzed as the number of individuals active per hour of day over the total number of locations for each hour of the day.

*Home range analysis.* We determined individual home range size using 95%, 75% and 50% fixed Kernel home ranges (Worton 1989), using the Animal Movement Analysis extension to ArcView, version 3.2 (ESRI, Environmental Systems Research Institute, Table 2). All animals had > 70 locations and a minimum of two months of data (Table 1); however, of six animals, only three survived or retained their radio tag covering a period

of time that included parts or the total wet and dry season, allowing us to estimate wet and dry season home ranges. We compared average home range size during the wet and dry season and compared the average of our home range estimates to other published estimates of home range for capybaras using a Student's *t*-test (Zar 1996). We calculated the distance from the estimated location to the nearest permanent water source using ArcView.

**Table 1.** First column: name of capybara trapped, column 1 and 2 period of time during which it was monitored, 3 sex, 4 weight, 5 group to which it belonged, 6 transmitter type, 7 total location points, 8 final report of the animal.

Name	1	2	3	4	5	6	7	
capy 1	07/03/2005	14/10/2005	Male	82 kg	1	Collar	71	Collar destroyed
capy 2	21/09/2005	13/12/2005	Female	56 kg	1	Collar	185	Disappeared
capy 45	03/02/2006	09/08/2006	Male	52 kg	3	Collar	405	Disappeared
capy 43	06/02/2006	28/2/2007	Male	17 kg	2	Implant	435	Disappeared
capy 41	28/06/2006	10/10/2007	Male	64 kg	2	Implant	600	Killed by puma
capy 50	06/10/2007	28/8/2007	Female	56 kg	3	Collar	85	Disappeared

*Habitat use analysis.* We compared utilized to available habitat to determine which habitats were preferred or avoided. Habitat types were identified and delimited from high resolution satellite images (eMap International, Boulder Colorado, USA). We created polygon shape files by hand corresponding to the 5 land cover types by using Geographic Information System (GIS). We delimited the study area as a 5 x 7 km rectangle (3,500 ha) which incorporated the six minimum convex polygon (MCP) home range estimations (Mohr 1947) of our six study animals (Fig. 1). We carried out a compositional analysis to identify proportion of utilized and available habitat types, and evaluated whether they were used significantly more or less than expected by chance at two scales of use (Aebischer *et al.* 1993). First, we compared the proportion of habitat in the study area (the available habitat) with the proportion of habitat in each MCP home range (the used habitat). Second, we carried out a finer scale analysis by comparing the estimated proportion of habitat found in each MCP home range (the available habitat) to the proportions of observed animal locations within each habitat type (the used habitat) using ArcView, version 3.2 (Environmental Systems Research Institute). When no locations were recorded in a particular habitat type, a value of 0.01% replaced the zero as suggested by Aebischer *et al.* (1993).

To test the null hypothesis of random use, we calculated the difference in log-ratios between matching pairs of used and available habitats. This hypothesis was tested by a multivariate analysis of variance (MANOVA). If the hypothesis of random use was rejected, we then ranked the habitat types in order of relative use. For this analysis, we calculated the mean log-ratio difference for all possible pairs of habitat types, and compared them to zero using a *t*-test (Aebischer *et al.* 1993). Compositional analysis makes it possible to rank habitat types in order of relative use, although in our case, seasonal (dry and wet) sample sizes were too small to produce clear patterns of significant differences between ranks. Statistical tests were carried out with program SAS 9.1 (SAS Institute, Inc., Cary, North Carolina).

*Body temperature analysis.* Two capybaras received internal radio-implants with temperature sensitive pulse rates. The temperature sensor monitored body temperature in the proximity of the implant. Special circuitry provided approximately 0.1 °C resolution; a self calibration pulse interval eliminated calibration drifts due to aging and battery voltage changes over time. Transmitters were calibrated at the factory and retested at the field station to maximize accuracy. Temperature data were recorded as the number of pulses per minute. We also collected ambient temperature, humidity and wind direction for each triangulation animal for the two animals with implants. We assessed the relationship of body temperature with distance from water, ambient temperature, time of day and season using a multivariate analysis of variance (MANOVA) and conducting posterior univariate linear regressions implemented by program JMP-IN 4.0.4 (SAS 2001, SAS Institute Inc., Cary, North Carolina).

## Results

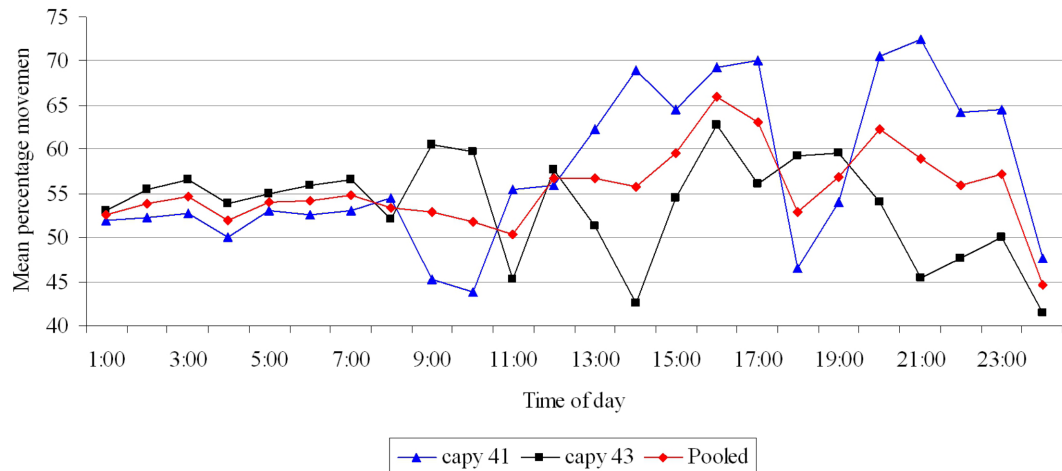
*Home range analysis.* Over the study period we collected total of 1,781 locations with a minimum of 71 and a maximum of 600 locations per animal; with an estimated mean location error rate of  $(41 \pm 12 \text{ m}, n = 9; \text{Table 2})$ . We found that there were patterns of spatial overlapping among individual capybara location points, between capybaras 95% kernel home range (Table 3). We were only able to estimate temporal overlap among four capybaras. We found 82% temporal and similar spatial overlap between capybaras 43 and 41 demonstrating that they were in the same group. We found zero temporal overlap between capybaras 43 and 41 with capybaras 50 and 45. Thus, we infer that capybaras 50 and 45 were from a separate group. Between capybaras 50 and 45, we were not able to estimate temporal overlap because they were radiomarked at different times during the study; however, we did find a spatial overlap of 19% and 80%, which suggests that these animals were likely from the same group. Similarly, we were not able to estimate temporal overlap between capybaras 1 and 2, because contact with them was lost before other animals were radiomarked.

Based on our spatial and temporal overlap data and confirmed by direct observation we were able to confirm that two capybaras (43 and 41) were in the same group. From observational data we found the group was formed by one adult male (41), one untagged adult female, and four juveniles including 43. We found that capybaras 1 (adult male) and 2 (adult female) had a high percent of spatial overlap between them, as did capybaras 43 and 41. These data suggest that capybaras 1 and 2 were possibly part of the same group, but that they either dispersed, or were depredated. We further infer that their home range was reoccupied by the group dominated by capybara 41. On the other hand, capybara 45 (adult male) had little overlap with any other capybara during the tracking period. This male was observed most of the time alone. However, on a few occasions it was observed together with a second capybara but with no juveniles. Home range analysis indicated that he was not part of the previously described group. This animal disappeared before capybara 50 (adult female) was trapped, which made it impossible to corroborate any temporal overlap between them. However, these animals had high spatial overlap, which suggests that this female may have been the one observed with capybara 45.



In general, the great majority of capybaras were observed ( $n = 304$  direct observations) near or on the shore of permanent ponds singly or in small groups, up to a maximum of six animals  $2.4 (\pm \text{SD } 1.4, n = 304)$ . Capybaras were often observed resting in the shade or walking near the ponds. Activity patterns of two capybaras revealed that daytime activity was higher than at night, with peaks of activity at 16:00 and 20:00 hrs (Fig. 2).

**Figure 2.** Long-term mean percentage activity by time of day for capybaras 41 and 43 individually and pooled in the Central Dry Chaco region (based on  $n = 1035$  locations).



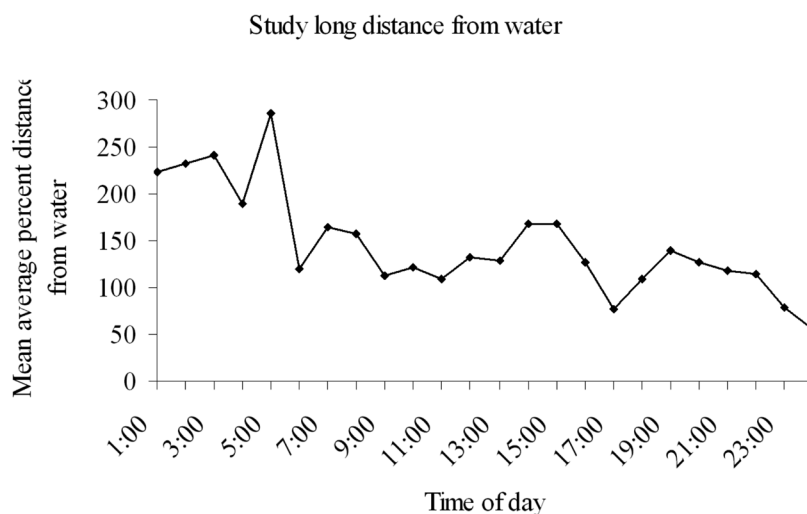
Using measures of adaptive kernels, the average 95% kernel home range size was 183 ( $\pm \text{SD } 54$  ha,  $n = 6$ ), the 75% kernel = 64 ha,  $\pm \text{SD } 22$  ha, and the 50% kernel =  $28 \pm \text{SD } 9$  ha (Table 2). We found no significant difference in home range size between seasons. Across seasons, the mean distance from water for all capybaras was 126 ( $\pm \text{SD } 4$  m,  $n = 1,781$ ), with the distance from water increasing from 1:00 to 6:00 AM ( $t = 2.01$ ,  $P < 0.01$ ; mean 1 = 214, Mean 2 = 132 (Fig 3). We found that 95 % of the locations were < 500 m from a permanent water source, 84 % were < 250 m, 64 % were < 100 m and 46 % were < 50 m from water.

**Table 2.** Overall capybara home range estimates.

Name	MCP	95% Kernel	75% Kernel	50% Kernel
capy 1	437	193	75	42
capy 2	492	421	168	67
capy 45	737	198	58	24
capy 43	484	52	17	10
capy 41	997	62	27	10
capy 50	352	176	40	16
Mean	583 ha	183 ha	64 ha	28 ha

*Habitat use analysis.* The average MCP home range size for all capybaras was 583 ( $\pm \text{SD } 97$  ha,  $n = 6$ ). Using our five land cover classifications (Chaco forest, shrub Chaco forest, introduced pasture, seasonally flood wetland and open water), we compared habitat use at two scales: the overall study area and within the MCP home range. Our results suggest that capybara did not establish home ranges at random ( $\Lambda = 0.023$ ,  $\chi^2 = 74.16$ ,  $P < 0.0001$ ). We assigned a ranking of a maximum of 4 for the habitat that was used most significantly and a minimum of 0 for the habitat that was significantly least used (Table 4). At the study

area scale habitat types were used in the following hierarchal sequence; Chaco forest > introduced pasture > seasonally flood wetland > open water > shrub Chaco forest (Table 4A). Shrub Chaco forest was used significantly less than available, followed by water and flooded grass area which were used significantly less in proportion to their availability. It is important to clarify that water surface and flooded grass area made up a comparatively small proportion of the overall study area. The two highest-ranked habitats, Chaco forest and introduced pasture, were used significantly more than their availability within the study area, and Chaco forest was selected over pasture. On the smaller scale of home range, overall use of the five habitat types based on proportion of radio locations in each habitat type differed significantly from the proportion of habitat available within the MCP home ranges ( $\Lambda = 0.037$ ,  $\chi^2 = 44.31$ ,  $P < 0.0001$ ). The ranking matrix indicated that the following habitat type followed sequence; open water > Chaco forest > pasture > flooded grass area > shrub forest (Table 4B). Within the home range, water was the most highly preferred habitat. Chaco forest and pasture had no detectable difference in use from availability. On the other hand, flooded grass area was significantly less used than available and finally; shrub forest was used less than its availability, suggesting that capybaras were avoiding this habitat type.



**Figure 3.** Long-term capybara average percent distance from water in the Central Dry Chaco region of Paraguay. Based on six capybaras and  $n = 1781$  locations.

*Body temperature analysis.* The mean ambient temperature during the study was  $28^{\circ}\text{C}$ , with an average day time temperature during dry season of  $27.26^{\circ}\text{C}$  ( $\pm$  SD  $7.04^{\circ}\text{C}$ ,  $n = 359$ ) and night  $24.64^{\circ}\text{C}$  ( $\pm$  SD  $5.66^{\circ}\text{C}$ ,  $n = 173$ ). The mean daytime wet season temperature was of  $31.88^{\circ}\text{C}$  ( $\pm$  SD  $6.10^{\circ}\text{C}$ ,  $n = 316$ ) and night  $29.52^{\circ}\text{C}$  ( $\pm$  SD  $4.50^{\circ}\text{C}$ ,  $n = 165$ ). We found no significant difference between wet and dry day time ambient temperatures ( $P = 1.96$ ; DF 673).

The mean body temperature for capybaras was  $36.15^{\circ}\text{C}$  ( $\pm$  SD  $0.27^{\circ}\text{C}$ ,  $n = 1021$ ), with no significant difference between dry and wet season. We found a significant positive correlation between body temperature and distance of the animal from water ( $r^2 = 0.004$ ,  $P = 0.02$ ;  $F = 4.51$ , DF = 1008). Likewise, we found significant negative correlation between ambient temperature and distance in which the animal was located from water ( $r^2 = 0.02$ ,  $P < 0.0001$ ;  $F = 29.01$ , DF = 1017; Fig. 4).

## Discussion

The study was conducted in a newly invaded area in which capybara population densities were low, thus our sample size was representative of the population at the leading edge of this newly occupied area (Brooks 1998; Campos-Krauer and Wisely 2009). Based on our direct observations and overlapping home range data, we confirmed that we were observing three groups of capybara. All groups were small, ranging from a pair to a family group of six. Each group had two to three ponds within their home ranges, near which they remained the majority of the time. Individuals within a group moved together to feed, rest or hide in surrounding habitats and always maintained the pond as the focal point. We found some spatial overlap but little temporal overlap between individuals from different groups. The exception was capybara two (female) who had high spatial overlap with two groups, and made occasional short solo trips to a neighboring pond.

**Table 3.** Asymmetrical matrix of percent data point spatial overlap between six capybaras monitored in the study area; columns correspond to percent overlap of corresponding capybara with different capybara.

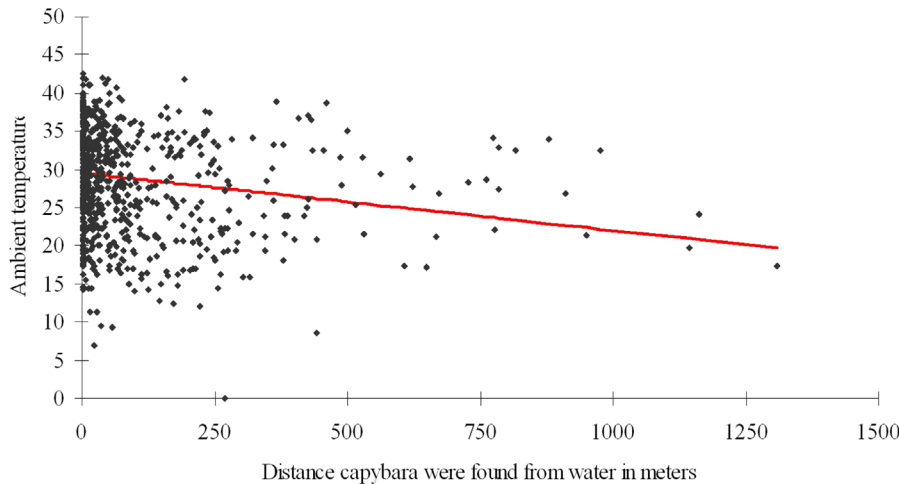
	capy 1	capy 2	capy 45	capy 43	capy 41	capy 50
capy 1		66	1	49	49	6
capy 2	56		20	29	29	0
capy 45	38	79		0	0	19
capy 43	83	90	12		84	1
capy 41	78	94	9	86		2
capy 50	0	1	80	0	0	

Activity patterns were similar to those of other populations in Venezuela (Herrera and Macdonald 1989) and Brazil (Alho *et al.* 1987). We found the highest activity of capybaras during the crepuscular hours with little movement during the middle of the day or at night. During the height of their movement, distance from water was less than 150 m indicating that movement was mostly in habitat directly surrounding or in ponds (Fig. 3). On the other hand, movement activity diminished during the night and distance from water increased. Groups moved during late afternoon and early night away from water ponds, occasionally moving to a different water pond or more commonly to a bedding area in the dense Chaco forest where they would spend the night. It appears that capybaras are using the cover of the darkness and the dense Chaco forest as protection during the night, moving closer to water the next morning (Fig. 3).

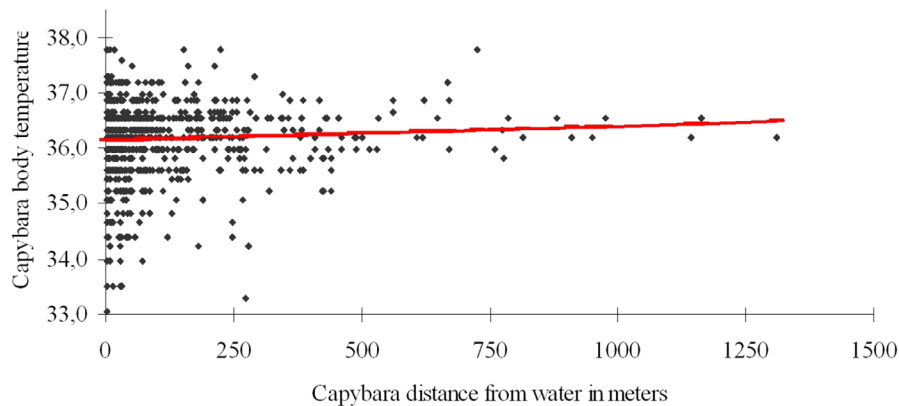
Capybara populations in the study area were not intensely persecuted by humans; occasional poaching was possible but rare. However, capybaras remained cautious of humans and as soon as human presence was detected, they would silently move toward the nearest ponds. Only when an animal was surprised by a human or a predator would they loudly jump in the water. Interestingly, capybaras remained in the water only for a few minutes where they would hide under aquatic vegetation. Shortly after entering the water, they would emerge and escape into the dense Chaco forest using it as the final hiding spot.

Predators such as jaguar (*Panthera onca*) and puma (*Puma concolor*) were common in the region and study area. One radiocollared capybara was confirmed to have been killed by a puma at the shore of a pond. Tracks and signs of predators were not uncommon surrounding the ponds. Similarly, the great majority of ponds in the study area had caiman (*Caiman yacare*) that are known to depredate juvenile and subadult capybara (Lord 2009). Their presence and the small size of some of the ponds in our study area could explain why capybaras tended to bed down in forested areas during the night, and use water as an escape mechanism for only a short while.

A. Ambient temperature vs capybara distance from water



B. Capybara body temperature vs distance from water



**Figure 4.** **A)** Scatterplot and linear regression of ambient temperature vs. capybara distance from water ( $y = 29.4909 - 0.0072 (x)$ ,  $R^2 = 0.02$ ,  $P < 0.001$ ). **B)** Scatterplot and linear regression of capybara body temperature vs. distance from water ( $y = 36.1138 + 0.003 (x)$ ,  $R^2 = 0.004$ ,  $P = 0.02$ ).

The overall average estimated 95% Kernels of 183 ha were similar to the estimates of home range for populations of capybara in the Brazilian Pantanal, with ranges of 200 ha (Schaller and Crawshaw 1981) and 196 ha (Alho *et al.* 1987, 1989). These home range sizes were considerably larger than those found in Venezuela by Herrera and Macdonald (1987) with home ranges of 10 to 16 ha with much higher population densities. Smaller home ranges and higher densities in the llanos of Venezuela could be due to the higher productivity of the grasslands than in the Pantanal or the Gran Chaco (Herrera and Macdonald 1989).

In general, capybaras were found close to man made ponds. In the Gran Chaco, capybara presence and survival directly depends on the availability of permanent water sources and introduced pastureland (Campos-Krauer and Wisely 2011). Natural permanent water sources in the region are scarce, and the great majority dry out during the dry season. Man made ponds are reliable year round sources of water that are regularly distributed throughout pasturelands. As a result, capybara home ranges in the central dry Chaco include two or three ponds separated by approximately 1 km, that are typically used by multiple groups, but only by one group at any given time. Despite the considerable distance covered by capybara, 95 % of all location points were < 500 m from a permanent water source demonstrating the importance of these ponds for capybara. A tight association with water is in agreement with previous reports (Barreto and Herrera 1998; Azcarate 1980; Cordero and Ojasti 1981; Murphey et al. 1985; Mones and Ojasti 1986; Herrera and McDonalds 1989; Lord 1991; Quintana and Malvarez 1994, 1998). Although we found no significant difference between wet and dry season home ranges when we considered all animals, two capybaras which were members of a group of six had larger home ranges during the dry season than during the wet season. Expansion during the dry season could be due to a need for additional food resources to support the group as productivity declined during the dry season. The dry season home ranges of solitary or paired capybaras were smaller than those of a larger group.

Considering that the capybara is a semi-aquatic rodent, it was not unexpected that open water surface was a preferred habitat type. Also preferred, however, was the Chaco forest. Chaco forest surrounded the great majority of ponds in the study area; we believe that the dense vegetation of the forest plays a significant role for capybara survival in the region. The forest likely serves as a cool shaded area during the hot summer and provides shelter from the cold south winds during winter, as well as suitable protection from predators. Introduced pastureland was also greatly preferentially used, providing extensive high quality forage year round for capybaras. Without pasturelands, capybara foraging habitat would be confined to small, periodically flooded natural pasture, or the vegetation immediately surrounding man made ponds. Interestingly, capybaras avoided shrub Chaco forest. The lack of understory vegetation provided few food resources and little protection against extreme climatic conditions and predators, making this habitat risky for capybaras.

Average body temperature of capybaras in our study (36.15 °C) was similar to body temperatures (36.0 to 36.6 °C) found by López-Barbella (1982). Body temperature was positively correlated with distance from water. These findings suggest that capybaras use water to thermoregulate as well as for predator avoidance. Our interpretation is further supported by the fact that capybaras ventured further away from water only when ambient temperatures were low (Fig. 4A, B). Although capybaras have been anecdotally reported to use water for thermoregulation, this is the first study providing evidence for this phenomenon through the use of internal temperature sensors. Other semi aquatic rodents such as the European beaver (*Castor fiber*), American beaver (*Castor canadensis*), and the Australian water rat (*Hydromys chrysogaster*) also use water as a thermoregulatory tool and have developed physiological adaptations to optimize the benefits of water high cooling capacity (Steen and Steen 1965; Hart 1971; Fanning and Dawson 1980).



Similarly, the capybara has evolved skin and hair that enhances the ability of water to regulate body temperature. Capybara epidermis is covered in folds of tissue which creates a larger surface area; additionally, the hair emerges at an acute angle and is clumped in groups of three or four follicles with the clumps spaced relatively sparsely which speeds the drying and body cooling process consequently reducing the body temperature (Pereira *et al.* 1980).

<b>A</b>						
	Open water	Flooded grass area	Pasture	Shrub forest	Chaco forest	Rank
Open water		-	---	+	---	1
Flooded grass area	+		-	+	-	2
Pasture	+++	+		+++	---	3
Shrub forest	-	-	---		---	0
Chaco forest	+++	+	+++	+++		4
ChF > P > FGA > W > ShF						
<b>B</b>						
	Open water	Flooded grass area	Pasture	Shrub forest	Chaco forest	Rank
Open water		+++	+++	+++	+++	4
Flooded grass area	---		-	+	-	1
Pasture	---	+		+++	-	2
Shrub forest	---	-	---		---	0
Chaco forest	---	+	+	+++		3
W > ChF > P > FGA > ShF						

**Table 4.** A. Ranking matrices for the overall comparison of habitat type use from minimum convex polygon home range versus habitat availability in the entire study area. B. Ranking matrices for capybara based on comparing proportion of radio locations for each animal in each habitat type within MCP home ranges per season. Each average element in the matrix was replaced by its sign. Sing represents level of significant deviation from random at  $P < 0.05$ . A value of 4 corresponds to the highest significantly used habitat, a value of 0 correspond to the lowest significant used habitat type.

Worldwide, anthropogenic land use change and fragmentation are implicated in the establishment of invasive species and expansion of native ones (Lockwood *et al.* 2007; Peterson and Vieglais 2001). Capybaras expanded their range to the Central Dry Chaco after large-scale deforestation occurred for ranching (Campos-Krauer and Wisely 2011).

It appears that ample forage and suitable cover were created when Chaco forest was fragmented. Perhaps more important to the expansion of capybaras into this region was the creation of regularly distributed, permanent water sources in this otherwise xeric habitat. As a taxon, rodents are highly successful invaders. They exhibit demographic traits necessary for expansion and invasion such as high fecundity, short generation time and opportunistic breeding which can provide advantages over local fauna. Rodents cause losses to harvest and can serve as vectors and host for diseases, and are the most significant crop pests (Singleton *et al.* 1999, Chu *et al.* 2003). Where they have invaded they often further alter habitat, examples being the American beaver (*Castor canadensis*; Fasanella *et al.* 2010) and the coypu (*Myocastor coypus*; Guichon *et al.* 2003). With

voracious appetites and high reproductive rates, these species are of great concern world wide.

The capybara also appears to be a species capable of taking advantage of anthropogenic land management and transformation. Large scale land cover change from Chaco forest to a patchwork of pasture and forest appears to be a critical factor allowing the capybara to expand its distribution to these regions. The central dry Chaco region has been intensively transformed during the last 70 years. This habitat transformation has drastically changed the structure and distribution of Chaco forest, creating conditions which have allowed the capybara to expand its range into an area previously inhospitable to it. Establishment in this region, however, is driven by the artificial provision of water.

The expansion of the capybara into the central dry Chaco is a clear consequence of the great land transformation that is affecting the area and its appearance should alert those responsible that the Gran Chaco ecosystem is being irreparably altered. Expanding species can outcompete native species and alter ecosystem services. Indeed, the presence of capybaras in the central Chaco may already be altering the epizootiological landscape of the region. Capybaras are reservoir hosts of *Trypanosoma evansi*, a protozoan parasitic infection that infects both domestic and wild mammals (Franke *et al.* 1994). At present, capybara population numbers are still small and have not been linked to any agricultural damage or disease reservoir in the central dry Chaco, but as livestock and capybara densities increase in this region (Campos-Krauer and Wisely 2011), the potential for disease outbreak will also augment. This study increases our understanding of the extent of the spatial distribution and ecological requirements of capybara which will be essential for effective management of this newly established species.

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