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DISTRIBUTION AND POSTBREEDING ENVIRONMENTAL RELATIONSHIPS OF NORTHERN LEOPARD FROGS (*RANA [LITHOBATES] PIFIENS*) IN WASHINGTON

Stephen S. Germaine^{1,3} and Dave W. Hays²

ABSTRACT.—Northern leopard frogs (*Rana [Lithobates] pipiens*) are considered sensitive, threatened, or endangered in all western states and western Canadian provinces. Historically present in eastern Washington in 6 major river drainages, leopard frogs are now only known to occur at 2 localized areas in the Crab Creek drainage in Grant County. During the summers of 2002–2005, we surveyed both areas to document extent of leopard frog distributions and to describe habitat and vertebrate community characteristics associated with leopard frog site occupancy. At Gloyd Seeps, 2 juvenile leopard frogs were observed in a total of 8.2 person-days of searching along a 5-km stream reach. At Potholes Reservoir, we surveyed 243 wetland sites in 7 management units known to have been occupied by leopard frogs during the 1980s. We confirmed leopard frog presence at only 87 sites (36%) in 4 management units. Site occupancy models for individual ponds indicated that, compared to unoccupied sites, occupied sites had slightly greater pond depths, less tall emergent vegetation, more herbaceous vegetative cover, and fewer neighboring ponds containing nonnative predatory fish. Models developed at the 1-km² scale indicated that occupied areas had greater average midsummer pond depths, fewer ponds occupied by bullfrogs (*Rana [Lithobates] catesbeiana*) and carp (*Cyprinus carpio*), and more herbaceous vegetation surrounding ponds. The Gloyd Seeps population now appears defunct, and the Potholes Reservoir population is in sharp decline. Unless management actions are taken to reduce nonnative fish and bullfrogs and to enhance wetland vegetation, leopard frogs may soon be extirpated from both sites and possibly, therefore, from Washington.

Key words: northern leopard frog, *Rana (Lithobates) pipiens*, nonnative fish, bullfrog, *Rana catesbeiana*, population declines, predation, site occupancy, wetland vegetation, Washington.

Despite widespread geographic distribution in North America, the northern leopard frog (*Rana [Lithobates] pipiens*; hereafter leopard frog) is now considered a sensitive, threatened, or endangered species in all western states and western Canadian provinces (see Alberta Wildlife Act 1996, Washington Fish and Wildlife Commission 2000, Conservation Data Centre of British Columbia 2001, COSEWIC 2002) and has been petitioned for listing as a federal candidate species (Anonymous 2006). In Washington, leopard frogs were elevated to endangered status after surveys of the 17 known historic locations confirmed occupancy at only 2 sites (Leonard et al. 1999). Historically present in the Columbia, Crab Creek, Pend Oreille, Snake, Spokane, and Walla Walla river drainages, leopard frogs are now only known in Washington at the Gloyd Seeps and Potholes Reservoir areas of the Crab Creek drainage in Grant County.

Leopard frogs disappeared from most sites in Washington by the mid-1980s (Leonard et

al. 1999), and timing of declines was similar in neighboring states and provinces. In Alberta, leopard frog populations declined during the late 1970s and have not recovered (Kendell 2003), and only one known population remains in British Columbia (Adama et al. 2004). In the northwestern United States, leopard frogs may have been extirpated from Oregon (St. John 1985, Stebbins 2003), and populations have declined in Idaho and Nevada (Groves and Peterson 1992, Panik and Barrett 1994, Koch et al. 1997).

Primary putative factors affecting leopard frogs include disease, habitat fragmentation and loss, artificial hydrologic manipulation, aquatic contaminants, bullfrogs (*Rana [Lithobates] catesbeiana*), and nonnative fish (Hayes and Jennings 1986, Hecnar and M'Closkey 1997a, Kendell 2003). In western North America, multiple factors appear responsible (Corn 1994), with habitat alteration and exotic species introductions considered the greatest threats (Richter et al. 1997). However, none of these

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studies were conducted in Washington. Our objectives were therefore to document the distribution of northern leopard frogs at the Gloyd Seeps and Potholes Reservoir areas and relate site occupancy to local vegetative, hydrologic, and nonnative-predator characteristics.

METHODS

Study Area

Gloyd Seeps (GS) and Potholes Reservoir (PR) lie 24 km apart along Crab Creek, Grant County, Washington, with Moses Lake Reservoir and its namesake town located between them. Both sites are on land administered by the Washington Department of Fish and Wildlife (WDFW), which administers Potholes Reservoir Unit jointly with the USDI Bureau of Reclamation (BOR). The BOR manages Potholes Reservoir as an agricultural water source under Columbia Basin Project (USDI-BOR 2002) directives. The mandate to provide irrigation constrains managers' ability to manipulate water levels to benefit native amphibians such as leopard frogs.

Wetlands at GS consisted of about 10 linear km along Crab Creek containing several small (<1-km-long) impoundments. Wetland vegetation was primarily low emergent and moisture-tolerant herbaceous species growing sporadically in narrow (2–10-m) linear bands along the creek. Lentic, shallow areas containing emergent vegetation that might serve as potential breeding sites were present but rare.

Wetlands at PR included an 8.8-km reach of Crab Creek and several hundred small ponds ranging in size from <0.1 to 30 ha surface area. Wetland vegetation at both areas included emergent (*Carex*, *Eleocharis*, *Juncus*, *Scirpus*, *Schoenoplectus*, and *Typha* spp.), grass-forb (*Polygonum*, *Bidens*, and *Xanthium* spp. and *Lactuca serriola*), and willow (*Salix* spp.) vegetation in both shrub and tree form. Reed canarygrass (*Phalaris arundinacea*), common reed (*Phragmites* sp.), and purple loosestrife (*Lythrum salicaria*) were rare but present.

At PR we surveyed ponds in 7 hydrologically distinct management units (USDI-BOR 2002), each of which had at least one record of a post-1980 leopard frog sighting (Leonard et al. 1999, McAllister et al. 1999, WDFW Heritage Database 2005). Units surveyed included Dunes, Lower Crab Creek, North Potholes Reserve, Peninsula North, Peninsula South,

Upper Crab Creek, and Upper West Arm. At PR in 2002, we focused surveys on areas most recently occupied by leopard frogs and on adjacent wetlands. During 2003–2004, we overlaid random sampling points on digital aerial photographs that were spatially rectified and high resolution, and we surveyed the wetland nearest each point. Because wetlands at GS were primarily linearly oriented along Crab Creek, we searched all of them, especially focusing on areas where leopard frogs had been observed since 1995 and on areas of suitable-looking habitat based on published descriptions (Dole 1965, Hine et al. 1981, Kendell 2002).

Leopard frog egg-laying occurs in late April to early May in our area (S. Germaine personal observation). This period coincided with annual rapid water impoundment at PR, and water levels rose ≥ 1 m during the period of egg-mass deposition and early larval development. During this time, most ponds surface-connected to large fish-bearing water bodies, and water levels rose into upland vegetation. During 1995–2004, annual high-water levels at PR occurred between 12 March and 15 June. Drawdown began immediately, and water levels had dropped 3–4 m by the time of larval metamorphosis, which occurred in mid-July in our area. Annual low-water levels occurred between 2 September and 23 October. During low water, mud and sand flats were exposed, and many shallow ponds dried completely.

Carp (*Cyprinus carpio*), rainbow trout (*Oncorhynchus mykiss*), brown trout (*Salmo trutta*), brown bullhead (*Ameiurus nebulosus*), walleye (*Stizostedion vitreum*), and bluegill (*Lepomis macrochirus*) were present in both areas. Large-mouth bass (*Micropterus salmoides*), smallmouth bass (*Micropterus dolomieu*), pumpkinseed (*Lepomis gibbosus*), black and white crappie (*Pomoxis nigromaculatus* and *Pomoxis annularis*), and yellow perch (*Perca flavescens*) were also present at PR (Washington Department of Fish and Wildlife unpublished data). In addition, mosquitofish (*Gambusia affinis*) were present in at least 2 isolated ponds in this area. None of these species are locally native.

Sampling

To determine leopard frog occupancy at each wetland, we conducted time-constrained visual-encounter surveys (Crump and Scott 1994, Olson et al. 1997, Kendell 2002) at both areas from July through early October of

2002–2005 (2002–2004 at PR, 2003–2005 at GS). During these months, juvenile leopard frogs were easy to observe (Kendell 2002, Simmons 2002), and bullfrogs were easily detectable via calling activity into early August, with bullfrog larvae and juveniles visibly detectable through September. We surveyed during daytime by slowly walking along waterlines and concentrating primarily on floating vegetative mats, shallow water with low emergent vegetation, and terrestrial areas of low emergent or herb/forb vegetation on moist to saturated soils. During summer, these areas were frequently used by leopard frogs (Dole 1965). Leopard frogs were seldom found in high (≥ 1 m) grassy vegetation (Merrell 1977), and Merriam (2002) observed leopard frogs 87% of the time in microsites having no overstorey vegetation (Hine et al. 1981). We therefore searched less-optimal areas (e.g., tall emergent vegetation) only after searching in focal habitat types and as time permitted. During searches, we methodically swept long-handled nets through vegetation to increase the likelihood of flushing frogs. At large ponds and sites permanently connected to the creek or reservoir, we surveyed 200–300 m of shoreline adjacent to the random-point origin. Time spent actively surveying at each site was generally 20–45 person-minutes, with variation among sites due to variation in wetland size; each wetland was searched at a uniform rate.

We also noted presence of bullfrogs and visually searched through the water column for fish from multiple vantage points around each pond. When we observed fish, we identified them to the most precise taxonomic level possible. In addition, we noted whether focal ponds connected at least seasonally to permanent water bodies (e.g., Crab Creek, Potholes Reservoir, or a large reservoir in the North Potholes Reserve), all of which contained non-native predatory fish and bullfrogs. We also noted whether focal ponds were ephemeral or permanent and whether they connected annually to other permanent ponds. We estimated maximum pond depth by noting high-water marks on shoreline vegetation. We measured actual pond depths by wading to pond center, and we indexed water clarity by inserting a white, 2.54-cm-diameter pole 0.5 m into each pond. Index values were clear (visibility > 50 cm), tannin colored (white pole became tea colored), slightly murky (visibility 20–50 cm

along pole), or murky (visibility < 20 cm). In cases where water depth was < 0.5 m, observers classified ponds as clear if objects on the bottom could be seen with fine detail, slightly murky if the bottom could be seen but with poor detail, and murky if the bottom could not be seen. Aided by high-resolution aerial photography, we walked pond perimeters and visually estimated percent vegetative cover to the nearest 5% in the following classes: tall emergent, low emergent, herbaceous, woody scrub-shrub, submerged or floating aquatic vegetation, and open water–exposed mudflats. All wetland-associated vegetation occurring below the ecotone with upland shrubsteppe vegetation was included in estimates.

Analyses

We related leopard frog site occupancy to environmental attributes by contrasting vegetative, vertebrate, and hydrologic characteristics of individual ponds using binary logistic regression and an information-theoretic multi-model comparison approach (Burnham and Anderson 2002). This calculation was done at 2 spatial scales: in individual ponds and in 1-km² areas within which individual pond data had been pooled. We chose a 1-km² spatial scale because juvenile leopard frogs may disperse up to 1 km soon after metamorphosis (Dole 1971, Seburn et al. 1997) and because radio-telemetered adult leopard frogs traveled this distance while migrating from breeding ponds to overwintering ponds in a related study (S. Germaine unpublished data). This scale also facilitated a logical organizational unit in GIS.

Because our survey season encompassed the period of rapid juvenile dispersal, we considered it likely that dispersing juvenile frogs would occasionally be encountered at ponds poorly suited to them. To reduce analytical noise associated with these observations, when developing models at the pond scale, we adjusted our logistic cutpoint to less than or ≥ 1 frog observed per 15 minutes of survey effort, following pseudospecies methods described by Hill et al. (1975) and Jongman et al. (1995). When developing models at the 1-km² scale, we used true presence (observed)–absence (not observed) to model wetland characteristics of grid cells.

Candidate predictor variables in the logistic regression model included the vegetative and hydrologic variables described above, plus

an enumeration of the neighboring ponds containing bullfrogs, carp, and predatory fish. We generated this number by counting how many of the 6 nearest neighboring ponds (inclusive of focal ponds) contained each type of nonnative vertebrate. We required ponds to be ≤ 1 km from focal ponds but made no further effort to account for variability in distance between ponds. Within 1-km² cells, we generated average and maximum values for each candidate predictor variable.

At each spatial scale, we evaluated the following competing models: a full global model, 1–2 reduced global models, a hydrologic-vegetative model, a hydrologic–nonnative vertebrate model, a vegetative–nonnative vertebrate model, a hydrologic model, a vegetative model, and a nonnative vertebrate model. The pond-scale full global model contained the variables pond depth, tall emergent vegetative cover, open water–exposed mud, herbaceous vegetative cover, bullfrog abundance, and fish abundance; the 1-km²-scale global model contained the variables pond depth, water clarity, maximum observed tall emergent vegetative cover, maximum observed herbaceous vegetative cover, bullfrog abundance, and carp abundance. While each independent variable in our study was chosen based on *a priori* information about its potential to influence leopard frog distributions, we screened each variable prior to model development by requiring *t* scores ≥ 0.7 when grouped on the dependent variable. We assessed correlations between candidates; and in all pairwise instances of $r \geq 0.60$, we retained only the variable thought to be of greater importance, based on other studies. For each model other than the full global model, we assessed within-model combinations of variables for confounding and interaction and kept only subsets of variables that fit the data as well as each initial model did based on log-likelihood G scores at $\alpha = 0.05$ (Hosmer and Lemeshow 1989).

We used the second-order Akaike's information criterion (AIC_c) values to compare models within each spatial scale because AIC_c is adjusted for small sample sizes (Burnham and Anderson 2002). Low AIC_c scores indicated models that used relatively few variables to achieve high fit to the data relative to other models being assessed. We ranked models by calculating Akaike weighted probabilities (w_i) for each score. Akaike weights rank models from best (high scores) to poorest using an

objective proportional ranking factor that sums to 1 over all models (Burnham and Anderson 2002). Once optimal models were identified at each spatial scale, we reran our logistic regression analyses and generated prediction-success tables to determine the classificatory success of each model using SYSTAT software (SYSTAT 2004).

RESULTS

At GS in 2003, we observed 2 juvenile leopard frogs in 0.5 person-days while searching a 0.5-km length of Crab Creek. In 2004 and 2005, we surveyed approximately 9.1 km along the creek in 7.7 person-days and confirmed no leopard frogs. In 2004, one suspect leopard frog exhibiting long zigzag jumps (Stebbins 2003) was observed but not captured for verification.

At PR we surveyed 243 individual wetland ponds within a 41-km² area in the 7 management units during 2002–2004. Fifty-five ponds (23%) were surveyed in multiple years, resulting in a total of 302 surveys. For ponds surveyed multiple times, we kept only the information from the survey with the highest abundance of leopard frogs. Leopard frogs were present at 87 PR ponds.

Because of the paucity ($n = 2$) of leopard frogs at GS, we collected detailed habitat data and modeled environmental relationships only at PR. At PR, distribution of leopard frogs and nonnative vertebrates varied among management units, as did the number of seasonally surface-connected ponds. Percent pond occupancy by leopard frogs was highest in the Upper Crab, Lower Crab, and Peninsula North management units, and leopard frogs were rare or absent elsewhere. Bullfrogs were largely absent from Upper Crab, Lower Crab, and Peninsula North but were present in $>25\%$ of all ponds surveyed in each management unit in the west half of the study area and in $>60\%$ of ponds in the Dunes and Upper West Arm units. Carp and nonnative sportfish were abundant in all units except Peninsula North, where they were entirely absent. More than 75% of ponds in all management units except for the Upper Crab and Peninsula North units contained multiple nonnative vertebrate types.

We collected environmental data at 222 surveyed sites throughout PR, and 8 variables met criteria for inclusion in pond-specific model building: surface connectivity, pond depth (at

TABLE 1. Akaike's information criteria parameters and log-likelihood scores for 8 logistic regression models characterizing environmental factors associated with leopard frog pond occupancy at Potholes Reservoir Wildlife Area, WA, 2002–2004. Best models, based on differences in weighted AIC values, shown in boldface. Variable definitions: PD = pond depth (at time of survey); CONNECT = degree of pond isolation; TE = tall emergent vegetative cover; OWEX = open water or exposed mudflat containing $\leq 10\%$ vegetative cover at water or soil surface; HE = live herbaceous vegetation; BFab = bullfrog abundance; Fab = nonnative predatory fish abundance.

Model: \pm coefficients(variables)	K	$\log_e(L)$	AIC	ΔAIC_c	w_i
Global reduced: 0.68(PD) – 0.04(TE) + 0.02(HE) – 0.27(Fab)	6	-55.29	123.13	0.00	0.68
Global: 0.69(PD) – 0.04(TE) + 0.00(OWEX) + 0.01(HE) – 0.27(BFab) – 0.26(Fab)	7	-54.49	125.94	2.81	0.17
Hydro + Veg reduced: 0.41(PD) – 0.03(TE) + 0.02(HE)	4	-58.13	126.64	3.51	0.12
Hydro + Exotics reduced: 0.62(PD) – 0.26(Fab)	3	-60.44	129.13	6.00	0.03
Hydrology: 0.43(CONNECT) + 0.58(PD)	3	-62.97	134.19	11.06	0.00
Veg + Exotics reduced: – 0.03(TE) – 0.29(Fab)	3	-67.04	142.32	19.18	0.00
Veg reduced: 0.02(HE)	2	-69.31	144.76	21.62	0.00
Exotic Vertebrates reduced: – 0.23(Fab)	2	-71.10	148.33	25.20	0.00

TABLE 2. Akaike's information criteria parameters and log-likelihood scores for 5 logistic regression models characterizing environmental factors associated with leopard frog occupancy of 1-km² wetland areas at Potholes Reservoir Wildlife Area, WA, 2002–2004. Best model, based on both differences in AIC scores and weighted AIC, shown in boldface. Variable definitions: pond depth = average pond depth; clarity = average water clarity; maxTE = maximum value of tall emergent vegetative cover; maxHE = maximum value of live herbaceous vegetation; BFab = average bullfrog abundance; CAab = average carp abundance.

Model: \pm coefficients(variables)	K	$\log_e(L)$	AIC	ΔAIC_c	w_i
Global reduced: 3.47(pond depth) + 0.02(maxHE) – 0.84(BFab) – 0.35(CAab)	6	-18.49	52.19	0.00	0.76
Global: 2.35(pond depth) + 0.45(clarity) – 0.01(maxTE) + 0.02(maxHE) – 0.71(BFab) – 0.44(CAab)	8	-17.36	55.36	3.17	0.16
Nonnative Vertebrates: – 0.68(BFab) – 0.17(CAab)	4	-23.95	56.96	4.77	0.07
Hydrology: 0.54(pond depth) + 0.29(clarity)	4	-26.95	63.04	10.85	0.00
Vegetation: 0.02(maxTE) + 0.02(maxHE)	4	-26.04	61.13	8.94	0.01

time of survey), percent area covered by tall emergent, herbaceous, or woody-stemmed vegetative types, percent area of open water-exposed mudflat, abundance of bullfrogs, and abundance of predatory fish. Variables for woody-stemmed and herbaceous vegetation were highly correlated (Pearson $r = -0.97$). We considered herbaceous vegetation more important to leopard frogs (Dole 1965, Merrell 1977, Hine et al. 1981) and therefore removed woody-stemmed vegetation from further analysis. Surface connectivity and predatory fish abundance were also correlated (Pearson $r = 0.64$), so we excluded connectivity from all models containing both hydrologic and nonnative vertebrate variables. We didn't find any interaction terms that improved model fit to the data.

Based on the w_i values, greater empirical support existed for the reduced global model than for any other model evaluated (Table 1; Burnham and Anderson 2002). The reduced global model was significant ($\chi^2 = 17.25$, df =

4, $P = 0.002$) and correctly classified 87.7% of occupied sites but had poor success (23.4%) in classifying sites where few or no frogs occurred.

Based on the variables in the reduced global model, sites containing leopard frogs during the postbreeding summer season had the following characteristics (Fig. 1): greater pond depths (median 0.5 m; interquartile range 0.1–1.0 m), less tall emergent vegetative cover (median 10%; 10%–20%), more herbaceous vegetative cover (median 80%; 30%–90%), and few fish (median index value 1; 0–4 neighboring pond[s] containing nonnative fish).

Six variables met criteria for inclusion in model-building at the 1-km² scale: average pond depth, water clarity, maximum percent cover of tall emergent and herbaceous vegetation, and abundances of bullfrogs and carp. Again, no interaction terms improved the model's fit to the data. Weighted AIC (w_i) values indicated overwhelming support for the reduced global model (Table 2).

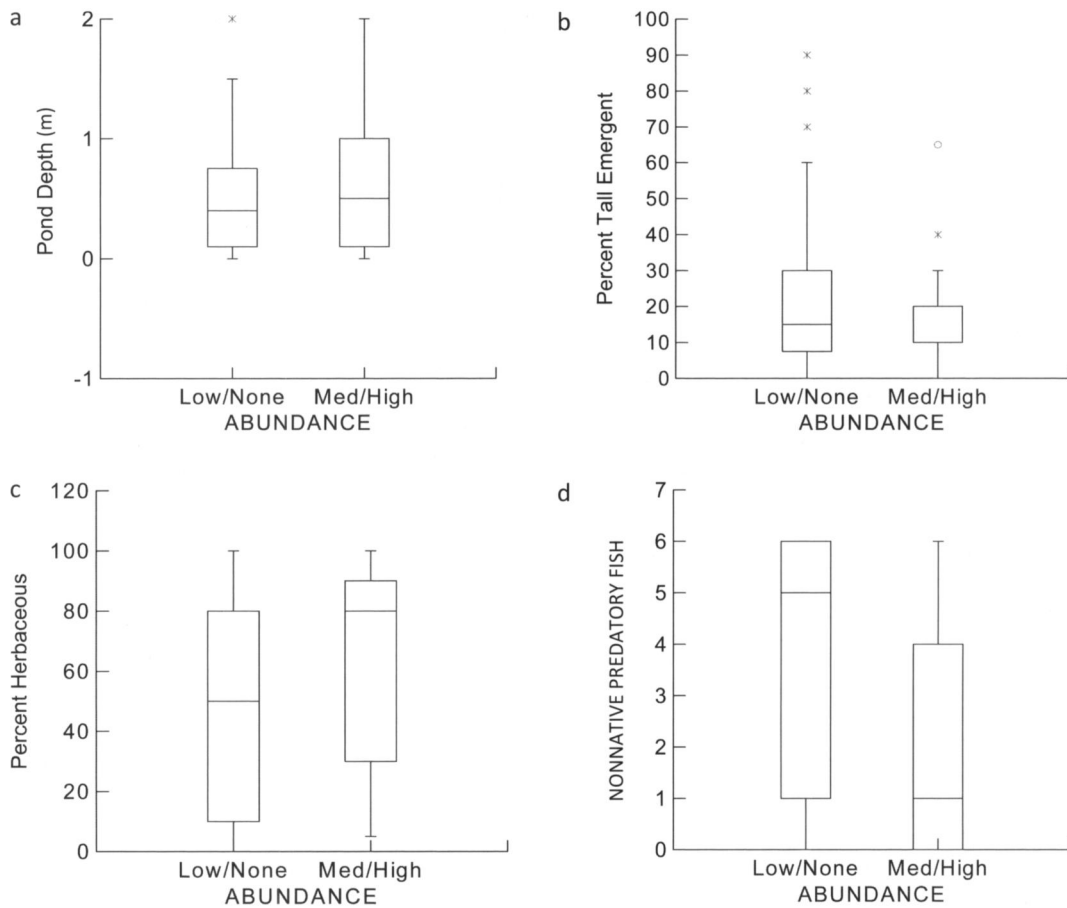


Fig. 1. Median values and 25th and 75th percentile values (box ends) for variables included in the most parsimonious of 5 models evaluated for ability to classify ponds by leopard frog abundance in Potholes Reservoir Wildlife Area, Washington, 2002–2004. Panels depict (a) pond depth (m), (b) percent area composed of tall emergent vegetation, (c) percent area covered in herbaceous vegetation, and (d) number of neighboring ponds (range 0–6) containing nonnative predatory fish. Asterisks (*) represent outlier values lying beyond $1.5 \times$ hinge-spread range.

The reduced global model for the 1-km² scale was significant ($\chi^2 = 21.14$, $df = 4$, $P < 0.001$), and correctly classified 72.5% of occupied areas and 69.8% of unoccupied areas. Areas occupied by leopard frogs during the postbreeding summer season had deeper average water depths, greater maximum amounts of herbaceous vegetative cover, and fewer ponds occupied by bullfrogs and carp than did areas where we found no leopard frogs (Fig. 2).

DISCUSSION

Leopard frog populations have clear trends and associations with nonnative vertebrates in our study area. We found leopard frog distributions at both GS and PR severely reduced rela-

tive to 10–20 years ago. At GS we documented no reproduction and observed only 2 leopard frogs during 3 years of surveying. The PR population now appears to be present in only 4 (57%) of the 7 management units in which it occurred during the 1980s. In 1 of the 4 PR units still occupied (the North Potholes Reserve Unit), leopard frogs were present at only 7% of the sites surveyed and co-occurred in all instances with both bullfrogs and nonnative fish.

At both spatial scales, strong empirical support existed for models containing vegetative, nonnative vertebrate, and hydrologic variables, suggesting that a suite of environmental factors influenced leopard frog distributions during summer. Also, our logistic regression models demonstrated that leopard frogs were

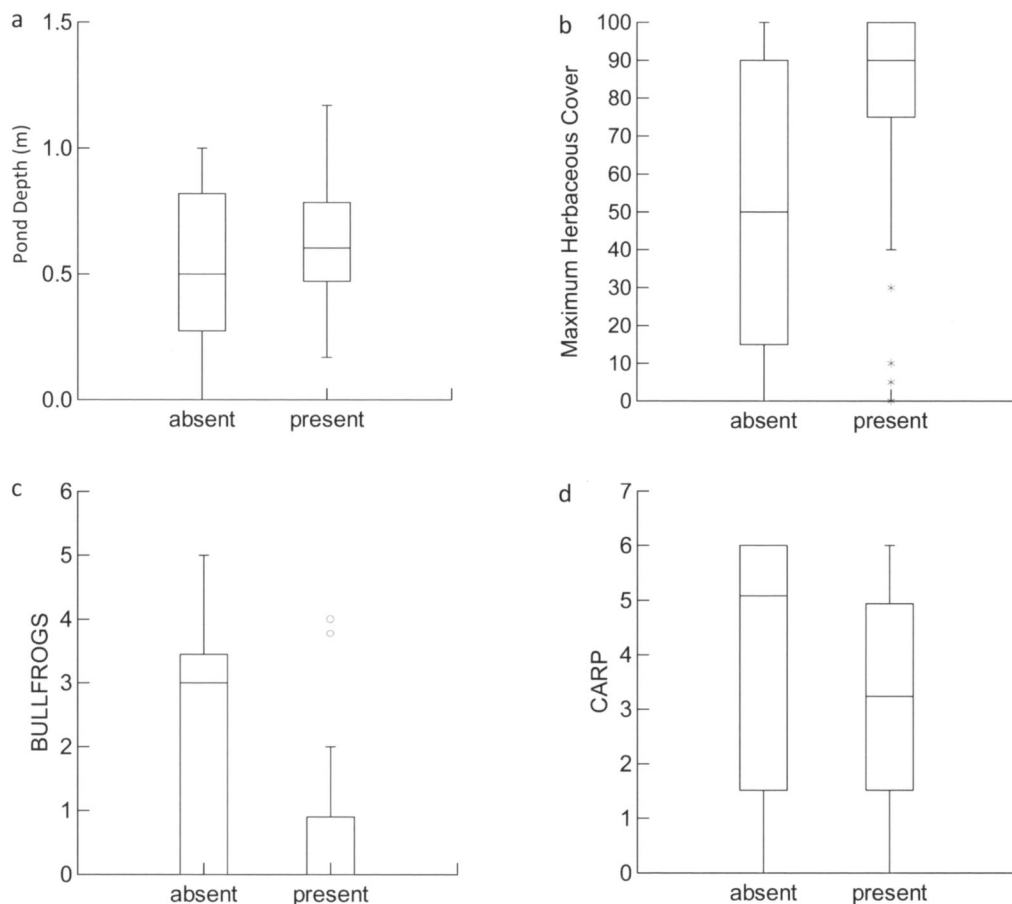


Fig. 2. Median values and 25th and 75th percentile values (box ends) for variables included in the most parsimonious of 5 models evaluated for ability to classify 1-km² wetland areas by leopard frog presence/absence in Potholes Reservoir Wildlife Area, Washington, 2002–2004. Panels depict (a) average pond depth (m), (b) maximum value of herbaceous vegetative cover, (c) average number of ponds occupied by bullfrogs, and (d) average number of ponds containing carp. Asterisks (*) represent outlier values lying beyond $1.5 \times$ hinge-spread range. Circles (o) represent outlier values lying beyond $3 \times$ hinge-spread range.

negatively associated with both bullfrogs and nonnative predatory fish.

While correlative studies do not prove cause and effect, a large body of published literature exists associating nonnative sportfish and bullfrogs with declines in native amphibian populations. Fish prey upon all life stages of ranid frogs, and heterogeneous fish assemblages are capable of causing local extinctions and altering distribution patterns (Petranka 1983, Hayes and Jennings 1986, Hecnar and M'Closkey 1997a, Bradford et al. 1993). Hine et al. (1981) concluded that fish predation could substantially reduce a leopard frog population, and multiple studies have described optimal leopard frog breeding ponds as free of nonnative fish

(Hine et al. 1981, Knutson et al. 2004). Most introduced fish species in the Pacific Northwest are aggressive and efficient generalist predators with whom native amphibians are not well adapted to coexist (Kats et al. 1988, Werner and McPeck 1994, Skelly 1996). Monello and Wright (1999) found a variety of amphibians absent from ponds containing introduced fish, with only the bullfrog able to reproduce successfully in fish-bearing ponds.

Bullfrogs also have a strong influence on amphibian community structure (Hecnar and M'Closkey 1997b, Laufer et al. 2007). Bullfrog colonization of the Upper West Arm and North Potholes Reserve units at PR occurred during the 1980s (R. Friesz and J. Tabor, Washington

Department of Fish and Wildlife, personal communication), which closely preceded leopard frog declines. We found leopard frogs almost absent from these units. Bullfrogs prey readily on other ranids, including leopard frogs, and consume both larval and newly metamorphosed stages (McAlpine and Dilworth 1989, authors' personal observation). Smith (1977) found that up to 80% of bullfrog diets consisted of frogs. Bullfrogs also may displace ranid frogs competitively (Kiesecker and Blaustein 1997, Kupferberg 1997) and may be more tolerant of disturbed environments (Kruse and Francis 1977, Hayes and Jennings 1986). Kupferberg (1997) found that yellow-legged frogs (*Rana boylei*) were less abundant in areas where bullfrogs became well established and documented that presence of bullfrog tadpoles resulted in a 48% reduction in survivorship of *R. boylei* tadpoles.

Herbaceous vegetation loaded in models at both spatial scales and was more abundant at occupied sites, suggesting high relative importance of herbaceous vegetation to leopard frogs during summer. We commonly found adult leopard frogs in moist terrestrial areas covered by low emergent and herbaceous vegetation. Adult leopard frogs become highly terrestrial in summer and may travel long distances (>3 km) from water under appropriate conditions (Breckenridge 1944, Merrell 1977, Knutson et al. 2004). Factors important to adults during summer include prey abundance, herbaceous cover, and moisture (Dole 1965, Merrell 1977, Seburn et al. 1997, Smith 2003). However, we observed that most herbaceous habitat occurred in small patches (≤ 0.04 ha) and was restricted to within 10–20 m of pond margins, suggesting that this important seasonal habitat component may be a limiting factor for leopard frogs in our system.

During summer, juvenile leopard frogs frequent the margins of more-permanent bodies of water (Merrell 1970), and this behavior may have contributed to our observation that occupied ponds tended to be slightly deeper than unoccupied ones. Also, pond depths were negatively correlated with the likelihood of a pond drying prior to completion of metamorphosis, and this may have contributed to our observation of occupancy at deeper ponds. We observed juveniles most frequently on floating vegetative mats and in terrestrial and aquatic areas containing low emergent vegetation (e.g.,

Eleocharis, *Carex*, and *Scirpus* spp.). In a telemetry study of leopard frogs at a natural site in Minnesota, Pember et al. (2002) observed leopard frogs in natural grass, aquatic emergent, and wet meadow vegetation 83% of the time during summer.

Finally, we advise caution when interpreting the coefficients of our tall emergent vegetation model. We focused surveys primarily on areas of herbaceous and short emergent vegetation because leopard frogs frequented these areas during summer (Dole 1965) and because leopard frogs were seldom found in grassy vegetation a meter or more high (Merrell 1977). Also, in a radiotelemetry study in nearby Idaho, Merriam (2002) observed leopard frogs 87% of the time in microsites having no overstory vegetation. We therefore spent relatively little time searching in tall emergent vegetation and recognize the possibility that we occasionally may have failed to detect leopard frogs in these areas. For this reason, we advise that any efforts to reduce tall emergent vegetation to favor leopard frogs be closely combined with intensive effects monitoring.

Management Implications

Our results suggest that removal of nonnative fish and bullfrogs will benefit northern leopard frogs. In California, *Rana muscosa* population densities increased 40-fold following removal of nonnative fish (Knapp et al. 2007), and the tadpole stage appeared to be the most vulnerable life stage (Vredenburg 2004). Petranks (1983) found densities of *Ambystoma texanum* larvae 16-fold greater in years when predatory fish were absent from breeding reaches of streams. Adams (1999) found fish to have a greater negative impact on *Rana aurora* than did bullfrogs. In systems where water levels are artificially managed (such as ours), some leeway exists for water-level manipulation even within irrigation constraints. Fish removal could be accomplished via drawdowns, by pumping isolated ponds dry, or with applications of Rotenone® during late summer or fall, after leopard frog larvae have metamorphosed and when water levels are already at annual lows.

Reducing bullfrog larvae may facilitate leopard frog coexistence (Doubledee et al. 2003), but such reduction should be carefully monitored because survival and growth of bullfrog larvae can be density dependent (Govindarajulu

2004). Stage-specific population modeling by Govindarajulu et al. (2005) suggests that if efforts are undertaken to remove bullfrog larvae, methods with potential to achieve near-complete removal, such as drawdowns and rotenone application, should be used.

At the pond scale, we recommend reducing tall emergent vegetative cover to $\leq 20\%$ (based on our interquartile-range estimates of occupied sites) of the total cover along the perimeters of ponds. This can be accomplished most efficiently by burning or mechanical removal and, to avoid mortalities, should be done during winter hibernation or early spring when frogs are at aquatic-breeding positions. In place of tall emergent vegetative cover, herbaceous vegetative cover should be encouraged to compose 30%–90% of the vegetation along the perimeters of ponds used as summer habitat (see Fig. 1).

At the 1-km scale, depths of breeding ponds (which may not be the same ponds as oversummering ponds) should be maintained at ≥ 0.5 m until metamorphosis is completed in mid-July, after which we encourage drawdowns at ponds containing nonnative fish or bullfrog larvae. Patches of herbaceous vegetation exceeding 0.4 ha should be encouraged and could be brought about simultaneously with burning and other actions designed to reduce tall emergent vegetation. Herbaceous patches should be implemented first at ponds known to be used by leopard frogs during summer. Finally, reducing the overall number of ponds containing bullfrogs, carp, and nonnative sport fish will benefit leopard frogs. These management actions should be coupled with continued monitoring at a level sufficient to evaluate their effectiveness for conserving northern leopard frog populations in Washington and elsewhere.

Optimally, the enactment of any of these recommendations should be coupled with a monitoring effort designed to assess whether such a management action achieves predefined response levels for both the direct management targets (e.g. tall emergent vegetation) and leopard frogs (Morrison et al. 2001). Also, the management plan should be cast in an adaptive framework, where monitoring information directs decisions about whether a particular action should be continued, altered, or replaced by another management action (Block et al. 2001).

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