Gharial nesting in a reservoir is limited by reduced river flow and by increased bank vegetation

Gaurav Vashistha1, Ninad Avinash Mungi2, Jeffrey W. Lang3, Vivek Ranjan2, Parag Madhukar Dhakate4, Faiyaz Ahmad Khudsar5 & David Kothamasi1*

The gharial (Gavialis gangeticus Gmelin) is a fish-eating specialist crocodylian, endemic to south Asia, and critically endangered in its few remaining wild localities. A secondary gharial population resides in riverine-reservoir habitat adjacent to the Nepal border, within the Katarniaghat Wildlife Sanctuary (KWS), and nests along a 10 km riverbank of the Girwa River. A natural channel shift in the mainstream Karnali River (upstream in Nepal) has reduced seasonal flow in the Girwa stretch where gharials nest, coincident with a gradual loss of nest sites, which in turn was related to an overall shift to woody vegetation at these sites. To understand how these changes in riparian vegetation on riverbanks were related to gharial nesting, we sampled vegetation at these sites from 2017 to 2019, and derived an Enhanced Vegetation Index (EVI) from LANDSAT 8 satellite data to quantify riverside vegetation from 1988 through 2019. We found that sampled sites transitioned to woody cover, the number of nesting sites declined, and the number of nests were reduced by > 40%. At these sites, after the channel shift, woody vegetation replaced open sites that predominated prior to the channel shift. Our findings indicate that the lack of open riverbanks and the increase in woody vegetation at potential nesting sites threatens the reproductive success of the KWS gharial population. This population persists today in a regulated river ecosystem, and nests in an altered riparian habitat which appears to be increasingly unsuitable for the continued successful recruitment of breeding adults. This second-ranking, critically endangered remnant population may have incurred an "extinction debt" by living in a reservoir that will lead to its eventual extirpation.

Freshwater habitats occupy less than one percent of the Earth’s surface, yet a tenth of all known species inhabit these areas, including a third of all vertebrate species1. Intense human pressures threaten the rich biodiversity in freshwater environments worldwide2,3, particularly the megafauna species which are at greater risk of declines and extinctions, than their smaller taxonomic counterparts4. The main threats include overexploitation, dam construction, habitat degradation, pollution and species invasion5. For example, dams not only interrupt river channel connectivity, but also have profound effects on riverine landscapes6,7. The major driver or grand structuring factor of river ecosystems is the natural flow regime, pulsating seasonally with floods and droughts8–10. River studies, firmly grounded in landscape-level perspectives11,12, are increasingly focused on dynamic models and management strategies that help predict restoration outcomes13–16.

For river-adapted habitat specialists, threats associated with loss of channel connectivity, altered flow regimes, and water extraction schemes are often direct and immediate and include increased harvest, restricted foraging opportunities, or loss of aquatic habitats. If the resultant impacts of these freshwater infrastructures, such as dams and irrigation canals, remain unaddressed, they can lead to species’ reductions, fragmentation, local extirpations (e.g., Indus River dolphins17), and regional extinctions (e.g., gharial in the Indus18). Spatially, freshwater megafauna show the largest range contractions, approaching 99%, in the Indomalaya realm, higher than other regions19.

Natural flood events can result in channel shifts altering river discharge dynamics, and ultimately affect species composition, distribution and abundance. A recent example occurred within the Karnali River basin, the third largest riverine system originating in Nepal and flowing into India. In 2010, after monsoon floods, the active

1Laboratory of Soil Biology and Microbial Ecology, Department of Environmental Studies, University of Delhi, Delhi 110007, India. 2Wildlife Institute of India, Dehradun, Uttarakhand 248003, India. 3Gharial Ecology Project, Madras Crocodile Bank Trust, Mamallapuram, Tamil Nadu 603104, India. 4Conservator of Forests, Western Circle, Haldwani, Uttarakhand 263139, India. 5Biodiversity Parks Program, Centre for Environmental Management of Degraded Ecosystems, University of Delhi, Delhi 110007, India. 6email: dkothamasi@es.du.ac.in
mainstream of the Karnali River shifted at a natural bifurcation from the east Geruwa (Girwa in India) channel to the west Karnali (Kaudiyala in India) channel (Fig. 1), resulting in attendant changes in river depth and flow from the east to the west channel. As a result of this natural channel shift, the relict Ganges river dolphin (*Platanista gangetica*) population shifted from east to west channel, but in doing so, moved from a protected stretch to one in which fishing and irrigation activities predominated20,21.

Just downstream, across the Indo-Nepal border, gharial (*Gavialis gangeticus*) inhabits a protected stretch of Girwa River (20 km in length; = east branch of the Karnali River), within the Katerniaghat Wildlife Sanctuary (KWS). This small breeding population in the KWS is highly ranked globally, second only to the much larger Chambal population18. The species is a habitat specialist, formerly abundant in large rivers across south Asia. In particular, gharials nest on sand substrates such as mid-river sandbars and high sandbanks, adjacent to deep water pools22. The natural channel shift in the Karnali Basin in 2010, described above, has resulted in reduced river flow in the Girwa stretch where the gharials nest. As a consequence, there has been a marked increase in woody vegetation at previously utilized nesting sites, and a concomitant decrease in gharial nesting (Fig. 2C).

Decades of river studies have established that 1) the riparian environment is disturbance-driven, and 2) the main disturbing element is river flow fluctuation23. Floods continuously shape and rework river landscapes, and erosion and deposition create open banks and bars24,25. But, in regulated rivers, these forces are greatly reduced or absent26,27. To date, the most relevant studies dealing with processes and mechanisms have focused on downstream vegetation below dams23,28–30, but recent research aimed at understanding riparian vegetation dynamics in upstream deltas and backwaters of reservoirs is particularly informative31,32.

In this study, our goal is to investigate how the recent changes in riverine habitats are related to gharial nesting patterns in the Katerniaghat Wildlife Sanctuary (KWS), and how these features ultimately relate to natural and regulated flow regimes, within the Karnali Basin from Nepal into India. Here, we document (1) the long term trends and recent patterns of gharial nesting in the KWS, (2) the long term trends and recent changes in nesting habitats, and investigate, (3) how gharial nesting in the KWS may be limited proximally by an increase in woody vegetation, related to the recent channel shift, and ultimately by reservoir setting in a regulated river. Finally, we briefly discuss, (4) conservation strategies to maintain and enhance the KWS gharial population.
Results

Gharial population size, composition, and nesting (1975–2020). Population counts of gharial and the number of nests located in the Katerniaqhat Wildlife Sanctuary (KWS) on the Girwa River are summarized in Table 1, based on the available primary sources referenced. The number of reproductive adults increased from less than 10 in the late 1970s to close to 50 in recent years (2016–2020; Table 1). Notably, there was a concomitant increase in nesting, from less than 5 nests in the late 1970s to more than 30 in recent years (Table 1). Importantly, for most of the past four decades, although nesting increased, recruitment as indicated by the numbers of juvenile and/or yearlings in the smaller size classes, has shown little evidence of anticipated increases, based on either the observed increase in natural nesting and/or the numbers of captive-reared gharial added to the resident population via periodic releases of >1800 gharials (Table 1).

Gharial nesting site distribution (2015–2019). Eight nest sites were confined to a 9 km stretch of the Girwa River within the KWS. Four sites were located on river banks (N1–3; N8) and the other four on mid-river sandbars (N4–7; Table 2, and Supplementary Data, S1 map and S2 natural history notes). In 2015, 5 of the 8 total sites had nests, increasing to 6 in 2016, and then to 7 in 2017. Then, the trend toward increasing nest sites reversed. In 2018, 3 sites were no longer used (N5, 7, 8), but a new site (N6) on an exposed mid-river sandbar was used. In 2019, only two sites (N1, N4) had nests.

Figure 2. Nesting site of gharial in different habitats. (A) A gharial nesting site on Chambal river in National Chambal sanctuary, Madhya Pradesh. This site represents a typical sandy river bank which gharials naturally prefer for nesting. (B) A gharial nesting site in Girwa river in Katerniaqhat before 2010 flood-channel shift. (C) Gharial nesting site N4 in Girwa river in 2018, post channel shift. (Image credits: (A) Suyash Katdare; (B) R. Whitaker).
From 2015 through 2019, the majority of nesting occurred at these two key sites, N1 and N4. Considering all 8 sites used during the five years, only these two sites had nests consistently year after year. Overall, N1 showed a tendency to flood, with reduced sand cover from 2017 through 2019 (N1, Fig. 5). N1 had the most nests in 2019, when nesting was restricted to only these two sites (N1 and N4), and was absent at the other sites (N2–3, N5–7; Table 2). A marked increase in woody cover (shrubs and trees) was evident at N4, coincident with a noted decrease in the number of nests, from 21 to 15 to 6, in 2017, 2018, and 2019, respectively (Table 2; Supplementary data, Figure S3).

### Habitat types based on vegetation cover

Vegetation at gharial nesting sites consisted of three grasses (*Phragmites karka*, *Saccharum spontaneum*, and *Typha* sp.), two herbs (*Euphorbia hirta* and *Agaratum conyzoides*), four shrubs (*Tamarix* sp., *Lantana camara*, *Ricinus communis* and *Calotropis gigantea*) and three tree species (*Adina cordifolia*, *Wrightia tinctoria* and *Bombax ceiba*). Open areas were mixtures of sand, water, and silt. There was an increase of 6% from non-woody to woody cover ($R^2 = 0.06 \pm 0.04$), but not in the sand cover ($R^2 = -0.45 \pm 1$). But otherwise, there were no substantial changes detected in the proportions of these categories at each site (Supplementary data, Figure S3).

### Habitat types demarcated using enhanced vegetation index

An Enhanced Vegetation Index (EVI) was derived from high resolution Landsat 8 satellite images of vegetation patterns, as these related to the five habitat types associated with the observed gharial nest sites sampled in 2015 to 2019. The corresponding EVI values for the habitat types were: $-1 < \text{water} < 0$; $0 < \text{sand} < 0.08$; $0.05 < \text{grass} < 0.13$; $0.13 < \text{shrub} < 0.215$ and $0.215 < \text{trees} < 0.325$ (Fig. 3A). When the above-derived EVI threshold were used to categorise the EVI from 1988 to 2019 into water, sand, grass, shrubs and trees, the average classifying accuracy across these classes was 66%. When the three combined types (water, sandy grasslands, and woody vegetation) were used to derive EVIs from 1988 to 2019, the overall accuracy of classifying increased to 94%, 91% and 88% for water, woody vegetation and sandy grasslands respectively (Fig. 3B).
Habitat–nesting relationship (2017–2019). The high classification accuracy indicated above validated the long-term habitat classification of gharial nests. The relation between EVI and the number of gharial nests indicated a strong preference for sandy and grass habitat types, and a sharp decline in numbers of nests with increasing EVI, representing woody vegetation ($R^2 = 0.52$, $p = 0.0004$) (Fig. 4).

Long term habitat dynamics (1988–2019). Time-series analyses of the eight nesting habitats revealed long-term consistent seasonal fluctuations between water (EVI < 0) to sandy grasslands (0 < EVI < 0.13). These nesting habitats changed gradually or abruptly to woody cover (EVI > 0.13). The dynamics and shift point to the woody condition differed among sites. However, across all sites, a consistent change occurred with the flood-related channel shift in 2010. Specifically, sites were either converted to water followed by woody cover, or converted directly to woody cover (Fig. 5). Sites N2 and N3 had the highest shifts with a single prominent shift in 2010, after which a gradual shift from sandy grasslands to woody cover was observed. Site N4 changed from sandy grasslands to woody cover in 1998 and from woody cover to water in 2010, after which it gradually changed into sandy grasslands and in 2013 to woody cover until 2019. A similar pattern was observed at site N5, where it shifted from sandy grasslands to water in 2010 and gradually into woody cover from 2013 until 2019. In 2019, at both sites, there was a sudden shift back to sandy grasslands. This sudden shift from woody cover to sandy grasslands at N4 and N5 was due to an active intervention by the Forest Department to remove vegetation on these sites.

Site N1 had the least overall shift, with two prominent change points during 2010 (sandy grasslands to water) and in 2015 (water to sandy grasslands). Site N6 and N7 were mostly submerged in water until 2013, after which the variance expanded due to oscillations among water, sandy grasslands, and woody cover. At site N8, mostly sandy grasslands until 2008, there was a gradual shift to woody cover in subsequent years (Fig. 5).

Discussion

Our study illustrates how reduced river flow, precipitated by a flood-related, natural channel shift adversely affected gharial nesting by promoting vegetation that impeded nesting at riverbank sites where nesting occurred in previous years. Specifically, from 2015 through 2019, the number of nesting sites declined by 70% and the number of nests were reduced by 46%. These decreases in nesting were coincident with a marked increase in woody vegetation at the established nesting sites used previously, related to the overall flow reduction in the east channel (Girwa = Geruwa), as the mainstream of the Karnali River upstream in Nepal shifted to the west channel (Kaudiyala). The observed effects were not immediate, but were observed subsequent to a major flood event in 2010. As a direct result of the flood, the Karnali River below the Chisapani Gorge shifted from the east channel (Geruwa), flowing into India as the Girwa, to the west channel of the Karnali, flowing into India as the...
Kaudiyala (Fig. 1). Prior to the flood event, the Geruwa had higher discharge and depth than the Karnali, and after the flood, the opposite was observed. In addition, subsequent to 2012, large scale water diversions from the Karnali west channel have occurred. In early 2011, within the Katerniaghat Wildlife Sanctuary (KWS), a 10 km stretch of the mainstream Girwa River, immediately below the Indo-Nepal border at Kothighat, still contained locally high densities of gharial, estimated at about 45 + adults. Nesting at that time was estimated at about 25 + nests (Table 1). In KWS, the number of active nest sites reduced from 7 in 2017 to 2 in 2019. During the study period, 6 out of 8 nesting sites were dominated by native and invasive woody species (e.g., Lantana camara), making them inaccessible for nesting. As a consequence, the number of gharial nests that were detected also declined in 2016 from 35 to 19 in 2019 (Table 1).

Our analysis of the vegetation changes at known nesting sites, by monitoring the dynamics between water, sandy grasslands and dense woody vegetation on the ground as well as via remote sensing, revealed that these habitats had fluctuated along the river banks in response to high water pulses from periodic floods. The riverine landscape experienced major flood events around 1995, 2000, 2008 and 2010. After the 1995 and 2000 floods, most of these sites gradually changed into sandy grasslands and dense vegetation, until they were submerged by another extreme flood in 2008 that removed the dense vegetation. However, these vegetation changes ceased after the 2010 flood. In subsequent years, the sampled areas were eventually converted into densely packed woody vegetation, without the periodic reversals to water, sand or grass stage. With exception of one site predominantly covered by grass (N7), all of the other gharial nesting sites were covered by woody vegetation (Figs. 2C, 5).

However, discussing habitat dynamics using moderate resolution data is challenging, so additional documentation of the shift to woody vegetation would be instructive. For instance, a single pixel of satellite imagery may constitute more than 75% water and 25% sandy grassland, but because of the classification method it is classified as water. In another scenario, a pixel might constitute > 90% vegetation and is classified as woody. In both of these settings, classified as either “water” or “woody,” if small patches of sand were present, these may have been suitable sites for nesting gharial, but such patches would have not been detected at the resolution level used in this study.

Therefore, our estimate of nesting habitat and its dynamics require refinement with high resolution satellite data. Moreover, for a large-ranging aquatic species known to utilize long stretches of free flowing rivers, our study area was relatively small (< 20 km). A small sampling area with just eight nest sites, precludes general comments about floodplain vegetation dynamics in an unregulated river system. Lastly, due to the absence of adequate hydrological data for the east (Girwa) vs. west (Kaudiyala) channels, a direct assessment of the relative channel discharges following the 2010 flood could not be done. However, upstream just across the border in Nepal, a small relictual population of Ganges river dolphins shifted from the east channel (Geruwa) to the west channel after the 2010 flood, coincident with documented depth changes in the channels, from deep to shallow in the east channel, and vice versa in the west channel. There were reductions in river depths in both channels after 2012 as a result of an increased irrigation demand upstream near the branch point at Chisapani, as well as increased water extraction from east channel to west channel within the braided inter-channel region of the Karnali Basin.

In other localities, gharial prefer to nest on high sandy banks, devoid of vegetation and adjacent to deep water, e.g. Chambal River nesting sites (Fig. 2A). In comparison with Chambal River sites, the nesting sites described...
here for KWS and characterized by fluctuating and higher EVI values (EVI > 0.13) present a completely different scenario, related to the altered riverine ecosystem associated with overall reduced river flow in the reservoir landscape. Habitats such as sand and grass (EVI < 0.13) provide easy access to such localities, compared to sites with woody vegetation (EVI > 0.13) (Fig. 2B). In this setting, the high water table related to the presence of the Girijapuri barrage and resultant reservoir generally promotes the growth and proliferation of riverside vegetation, particularly in the broad deltaic areas where inlet channels with reduced flow join the reservoir31,32. In the KWS, the recent reduction in river flow in the Girwa River related to the channel shift away from the mainstream Karnali has likely contributed to the loss of potentially suitable nesting sites in several additional ways. Included here is the loss of scouring action by floodwaters that would periodically clear riverbanks of vegetation, as well as the lack of "new" sediment deposition sites created by floodwaters, such as sand and/or gravel banks or bars that would create nesting sites de novo33,29,30.

Figure 5. Habitat dynamics of gharial nesting sites from 1988 to 2019 in KWS. The EVI (y-axis) below the blue line represents water submerged habitat, between blue and green line represents mosaic of sand and grass ("sandy grasslands"), and above the green line represents mixture of shrubs and trees ("woody vegetation"). A habitat shift was observed across most of the sites around the year 2010, from which point the habitat gradually converted into woody cover.
Nest site selection is an important factor that contributes to successful breeding in reptiles. Vegetation has several direct and indirect effects on the nesting of crocodylians. It can reduce accessibility to potential nesting area, influence hatching sex ratio, increase embryo mortality and change physical parameters of the substrate. Our observations indicate that the invasion of woody vegetation limits a gharial's access to potential sites in the KWS, making these overgrown sites unfit for nesting (Fig. 2C). It can further affect maternal care by restricting the female gharial's access to nests, resulting in death of fully grown hatchlings. During our sampling period, we found evidence of physical damage to shells of incubating eggs by vegetation roots that ultimately resulted in egg mortality. In addition, gharial, incubation temperature determines hatching survival as well as many important attributes, so suitable nesting substrate and other physical features of the nesting site play important roles in nest success.

In the KWS, the small remnant gharial population is vulnerable to limited habitat size, lack of suitable nesting sites, hatching predation (including egg poaching by local people) and poor hatching recruitment. Status reports, primarily winter basking counts of adults and nest numbers, provide snapshots of population numbers and composition. The total population size and nest numbers have gradually increased since 1975, but the increase is almost negligible compared to numbers of captive-reared, released gharial added periodically to the KWS population. Restocking with captive reared 'head-started' gharial, primarily juveniles, was initiated in 1979, and has continued through 2020, with the cumulative releases totaling 1852 to date (Table 1). But in reality, very few of the head-started animals released in KWS actually became residents there, as is clearly evident from low numbers of juvenile gharial in the population counts (Table 1).

The plausible reason for a chronic failure to recruit young animals is that most, if not all, are simply moving downstream below the reservoir dam, and are not able to move back upstream, once they cross the Girijapuri barrage gates and enter the Gaghara River. This situation applies to young individuals, both wild hatched residents as well as captive-reared releases. The loss of nesting habitat in the Gaghara river, that we have documented in this study, could result in local extinction of this species in the KWS. If this gharial population were not able to recruit successfully, as it appears may now be the case, its loss may well qualify as the unfortunate payment of an "extinction debt," e.g. a delayed species extinction as a consequence of an ecosystem perturbation, directly related to the dam and reservoir established in 1976. Poor recruitment despite the addition of hundreds of supplemented individuals is a serious problem confronting the persistence of this long-standing reservoir population.

Survival of the KWS gharial population may well depend on active and immediate intervention to reverse multiple factors contributing to this obvious lack of recruitment. Two management interventions were piloted at the KWS nesting sites recently: (1) vegetation removal by clearing vegetation on nesting banks, and (2) sand addition by shifting sand to extend river banks. One effective immediate solution could be providing artificial sandbanks by simply piling sand at the river's edge to simulate naturally occurring sandbanks that formed at KWS prior to the channel shift post-flood. This has proved remarkably successful in 2020 at KWS. Newly constructed sites where sand was added were readily utilized for nesting, resulting in nearly double the number of nests (Vashistha et al. 2021, Unpublished data; also see Table 1).

The long term solution will necessitate retention of juveniles within the KWS system, and their eventual recruitment into the adult breeding population. Unfortunately, there is little advantage for the resident gharials in the KWS to shift westward into the unprotected mainstream Kaudiyala channel where the river flow conditions might periodically fluctuate sufficiently to create suitable nesting habitat, because the west channel is intensively cultivated and fished. Upstream in the Karnali Basin, below Chisapani, gharial have largely been eliminated in recent years, with little to no evidence of breeding and only the occasional animal sighted between the Indo-Nepal border and Chisapani, in either channel within the past several decades. Movement upstream in the east channel across the Indo-Nepal border from KWS into the Geruwa is less likely due to the reduced flow since the 2010 flood, and the 7 km stretch of the Geruwa upstream from Kothiaghat is unprotected below the Bardiya National Park boundary. Our results suggest that one possible solution may involve a cooperative initiative by the two countries in the Karnali basin—India and Nepal—to develop a collaborative conservation agenda for this important riverine landscape while it is still largely intact and free-flowing.

**Methods**

**Study site.** Our study focused on 20 km of the Girwa river from Girijapuri barrage (28°16'21" N 81°05'13" E) to Indo-Nepal Border (28°22'02" N 81°12'05" E) in Katerniaghat Wildlife Sanctuary (KWS). This sanctuary has an actively breeding gharial population with 47 adults. This is the only breeding area remaining for gharial in the Karnali basin. The study site consists of extensive alluvial plains, hygrophilous grasslands and tropical moist deciduous forests. Girwa channel width was 403.18 ± 150.25 m during 2017–2019 and the river is interspersed with sand bars. Owing to freshwater demands for agriculture and human consumption along the river banks, infrastructures such as a barrage (Girijapuri barrage) and irrigation canals (Sharda Sahayak link canal and Saryu Nahar Irrigation) were built at the confluence of Girwa and Kaudiyala inside the KWS. The barrage marks the downstream boundary of KWS gharial population. It is opened at least twice each year, once for gate maintenance in April and again during the summer monsoon. The Girijapuri barrage presents a one way travel route for any gharials moving downstream, especially hatchlings and juveniles. Upstream movement across the barrage is not possible when the gates are closed. Hence the barrage likely plays a key role in the population dynamics of the gharials upstream in the KWS, but movement data to support this supposition is lacking.

**Gharial population data.** We assembled available data on gharial population counts and on nest counts from primary sources, published as well as unpublished (Table 1). Population counts for 2016–20 were conducted by boat surveys in January–February annually. The surveys started in Girwa river from Pathara, went...
which we sampled the habitat and compared the imagery with sampled habitat plots and with Google Earth imagery. First we trained our visual interpretation of the Google Earth imagery by using the visually validated a subset of EVI classified scenes for every year by visually comparing the classified habitat map the habitat types in this landscape. However, across the considered 31 years, Landsat satellite sensors and resolutions were used for each site. Using the aforementioned thresholds of classifying the EVI into five habitat types, the all available scenarios across different months over a period of 31 years. A total 357 scenes of different time periods 1988-2019 was used. For each sampled location, EVI values were compiled for series 4, 5, 6, 7 and 8 for the years 1988-2019. For analyses using vegetation cover, three habitat types were considered, consisting of "herbs", "woody" (shrubs and trees), and "sand" (sand and silt).

For EVI based classification, five habitat types (water, sand, grass, shrub, tree) were identified for this study. Water habitat consisted of plot submerged under water for at least 5 cm depth. Sand habitats were barren areas covered with sand and silt and lacking vegetation. Grass, shrub and tree habitats were areas that were covered by monocotyledon plants (excluding bamboo) or woody dicotyledon plants taller than 30 cm or woody dicotyledon plants taller than 200 cm respectively. Habitat plots were classified into different habitat types based on criteria mentioned in Table 3. The differential threshold used to determine the plot dominance was based on the relative contribution of different habitat types to the plot cover. For instance, even if a single tree was present in a plot (i.e. cover > 10%), it hinders the growth of grasses under it; while a relative dominance by grasses was possible only when more than a third of the plot (i.e. cover > 30%) was covered with grasses.

For understanding the habitat changes and its relation with the study species, it was important to correlate the habitat with satellite remote sensing data so as to monitor long-term changes. In order to establish a relationship between the sampled habitat plots and satellite data, we used high resolution Landsat 8 satellite images for the sampled locations to derive an Enhanced Vegetation Index (EVI) following Jensen60. The EVI for all images captured in March and April of every year were averaged to obtain an average EVI for every habitat plot for every year for the period 2015–2019. A random subset of 80% habitat plots was used to assign an average EVI value. This assignment provided the data variability in EVI values for the five habitat types. This data was used to classify the averaged EVI into these five habitat types using supervised classification in ERDAS Imagine Software (Version 2015). The remaining 20% habitat plots were used to validate the parity between habitat types recorded on ground to the habitat types mapped derived from EVI, using Kappa statistic61. Due to overlap in signature values of sand and grass, as well as shrubs and trees, we combined these five habitat types resulting in only three habitat types, i.e. water, sandy grasslands and woody vegetation.

Long-term habitat dynamics. Landsat EVI effectively captures long-term changes in vegetation cover at an optimal resolution. To estimate vegetation cover trends for the last three decades, EVI acquired from Landsat series 4, 5, 6, 7 and 8 for the years 1988–2019 was used. For each sampled location, EVI values were compiled for all available scenarios across different months over a period of 31 years. A total 357 scenes of different time periods were used for each site. Using the aforementioned thresholds of classifying the EVI into five habitat types, the EVI values were classified into the five types for each nesting site. This provided the first time series product of the habitat types in this landscape. However, across the considered 31 years, Landsat satellite sensors and resolution have changed substantially. This may affect EVI accuracy and band reflectances. To address this issue, we visually validated a subset of EVI classified scenes for every year by visually comparing the classified habitat map with Google Earth imagery. First we trained our visual interpretation of the Google Earth imagery by using the images of 2018, during which we sampled the habitat and compared the imagery with sampled habitat plots and

| SL. No | Habitat type | Criteria used for assigning an individual plot to a habitat type |
|--------|-------------|---------------------------------------------------------------|
| 1      | Water       | >75% of the plot’s surface was covered by water and the rest of the plot was either sand or grass |
| 2      | Sand        | >75% of the plot’s surface was barren sand and the rest of the plot was water or grass |
| 3      | Grass       | >30% of the plot’s surface was covered by grass and the rest of the plot was water or sand |
| 4      | Shrub       | >10% of the plot’s surface was covered by shrubs (woody dicotyledon plants taller than 30 cm) and the rest of the plot was water, grass or sand |
| 5      | Tree        | >10% of the plot’s surface was covered by trees (woody dicotyledon plants taller than 200 cm) and the rest of the plot was water, grass, shrub or sand |

Table 3. Habitat groupings, based on surface features measured in a habitat plot.
associated EVI values. Subsequently, we assessed at least one Google earth scene per sampled location per year to record the type of habitat. In total, there were 88, 67, 35, 33 and 37 observations for water, sand, grasses, shrubs and trees respectively. We matched this visually interpreted habitat type of each nesting site with the categories derived from the closest classified EVI. Parity was assessed using Kappa statistic.

Post validation, the habitat categories were compared across the temporal scale from 1988 to 2019 to visualize the seasonality and shifts in the habitat dynamics of each nesting site. We used this time series model of the habitats to identify timeline points after which a shift was observed in mean and/or variance of the habitat dynamics, using the R package CHANGEPOINT62. The point, following which there was no return to prior dynamics of the seasonality and shifts in the habitat dynamics of each nesting site. We used this time series model of the habitat dynamics and habitat nesting based on the visual observations of the habitat dynamics and potential regime shift. EVI, a derivative of habitat type and satellite imagery assessment, was compared with the nesting effort. The average EVI of the nesting season was compared with the number of actual nests. Trial nests were not included in the assessment. For comparison, we assessed the fit of linear, exponential and logarithmic lines. The line with highest R-squared value was considered as the best representation of the relationship between gharial nest number and the EVI. Precision of the fit was assessed by 1000 bootstrap replications in R version 3.3.166, wherein we selected 80% of observation points each time, assessed the p-value and reported the average p-value.

**Ethics statement.** This study was based on non-invasive sampling such as boat surveys, vegetation sampling and satellite imaging. No animal handling was involved and therefore the study was not assessed by an animal ethics committee. All necessary permissions for entry into protected areas were obtained from the Environment, Forest and Climate Change Department, Uttar Pradesh (EFCCD), vide letter no. 2522/23-2-12(G) dated 31st March 2016.

Received: 14 September 2020; Accepted: 12 February 2021
Published online: 26 February 2021

**References**

1. Strayer, D. L. & Dudgeon, D. Freshwater biodiversity conservation: Recent progress and future challenges. *Freshw. Sci.* **29**, 344–358 (2010).

2. Dudgeon, D. *et al.* Freshwater biodiversity: Importance, threats, status and conservation challenges. *Biol. Rev.* **81**, 163–182 (2006).

3. Reid, A. J. *et al.* Emerging threats and persistent conservation challenges for freshwater biodiversity. *Biol. Rev.* **94**, 849–873 (2019).

4. He, F. *et al.* Freshwater megafauna diversity: Patterns, status and threats. *Divers. Distrib.* **24**, 1395–1404 (2018).

5. He, F. *et al.* Disappearing giants: A review of threats to freshwater megafauna. *WIREs Water* **4**, e1208 (2017).

6. Nilsson, C. & Berggren, K. Alterations of riparian ecosystems caused by river regulation: Dam operations have caused global-scale ecological changes in riparian ecosystems. How to protect river environments and human needs of rivers remains one of the most important questions of our time. *BioScience* **50**, 783–792 (2000).

7. Nilsson, C., Reidy, C. A., Dynesius, M. & Revenga, C. Fragmentation and flow regulation of the world’s large river systems. *Science* **308**, 405–408 (2005).

8. Nilsson, C. & Svedmark, M. Basic principles and ecological consequences of changing water regimes: Riparian plant communities. *Environ. Manag.* **30**, 468–480 (2002).

9. Lyle, D. A. & Poff, N. L. Adaptation to natural flow regimes. *Trends Ecol. Evol.* **19**, 94–100 (2004).

10. Junk, W. J. & Wantzen, K. M. The flood pulse concept: New aspects, approaches and applications—an update. in Proceedings of the Second International Symposium on the Management of Large Rivers for Fisheries (eds. Welcomme, R. L. & Petz, T.) 117–149 (Bangkok: Food and Agriculture Organization and Mekong River Commission, FAO Regional Office for Asia and the Pacific, 2004).

11. Wiens, J. A. Riverine landscapes: Taking landscape ecology into the water. *Freshw. Biol.* **47**, 501–515 (2002).

12. Benda, L. *et al.* The network dynamics hypothesis: How channel networks structure riverine habitats. *Bioscience* **54**, 413–427 (2004).

13. Poff, N. L. Beyond the natural flow regime? Broadening the hydro-ecological foundation to meet environmental flows challenges in a non-stationary world. *Freshw. Biol.* **63**, 1011–1021 (2018).

14. Castro, J. M. & Thorne, C. R. The stream evolution triangle: Integrating geology, hydrology, and biology. *River Res. Appl.* **35**, 315–326 (2019).

15. Palmer, M. & Ruhi, A. Linkages between flow regime, biota, and ecosystem processes: Implications for river restoration. *Science* **365**, eaau2087 (2019).

16. Van Looy, K. *et al.* The three Rs of river ecosystem resilience: Resources, recruitment, and refugia. *River Res. Appl.* **35**, 107–120 (2019).

17. Braulik, G. T., Arshad, M., Noureen, U. & Northridge, S. P. Habitat fragmentation and species extirpation in freshwater ecosystems; causes of range decline of the Indus River Dolphin (Platanista gangetica minor). *PLoS ONE* **9**, e101657 (2014).

18. Lang, J., Chowfin, S. & Ross, J. *P. Gavialis gangeticus.* in The IUCN Red List of Threatened Species 2019: e.T8966A3148543. [https://doi.org/10.2305/IUCN.UK.2019-1.RLTS.T8966A3148543.en](https://doi.org/10.2305/IUCN.UK.2019-1.RLTS.T8966A3148543.en).

19. He, F. *et al.* The global decline of freshwater megafauna. *Glob. Change Biol.* **25**, 3883–3892 (2019).

20. Khanal, G. *et al.* Irrigation demands aggravate fishing threats to river dolphins in Nepal. *Biol. Conserv.* **204**, 386–393 (2016).

21. Paudel, S., Timilsina, Y. P., Lewis, J., Ingerson, T. & Inaswali, S. R. Population status and habitat occupancy of endangered river dolphins in the Karnali River system of Nepal during low water season. *Mar. Mammal Sci.* **31**, 707–719 (2015).

22. Whithaker, R. & Basu, D. The Gharial (*Gavialis gangeticus*): A review. *J. Bombay Nat. Hist. Soc.* **79**, 531–548 (1983).

23. Vesipa, R., Camporeale, C. & Ridolfi, L. Effect of river flow fluctuations on riparian vegetation dynamics: Processes and models. *Adv. Water Resour.* **110**, 29–50 (2017).
24. Merritt, D. M. & Cooper, D. J. Riparian vegetation and channel change in response to river regulation: a comparative study of regulated and unregulated streams in the Green River Basin, USA. *Regul. Rivers: Res. Mgmt.* 16, 543–564 (2000).
25. Littrell, J. J., Bechtold, J. S., O’Keefe, T. C., Pelt, R. V. & Naiman, R. J. Dynamic patch mosaics and channel movement in an unconfined river valley of the Olympic Mountains. *Freshw. Biol.* 51, 523–544 (2006).
26. Braatne, J. H., Rood, S. B., Goater, L. A. & Blair, C. L. Analyzing the impacts of dams on riparian ecosystems: A review of research strategies and their relevance to the Snake River through Hells Canyon. *Environ. Manag.* 41, 267–281 (2008).
27. Merritt, D. M., Scott, M. A., LeRoy, P. N., Auble, G. T. & Lytle, D. A. A. Theory, methods and tools for determining environmental flows for riparian vegetation: Riparian vegetation-flow response guilds. *Freshw. Biol.* 55, 206–225 (2010).
28. Poff, N. L. & Zimmerman, J. K. Ecological responses to altered flow regimes: A literature review to inform the science and management of environmental flows. *Freshw. Biol.* 55, 194–205 (2010).
29. Miller, K. A., Webb, J. A., de Little, S. C. & Stewardson, M. J. Environmental flows can reduce the encroachment of terrestrial vegetation into river channels: A systematic literature review. *Environ. Manag.* 52, 1202–1212 (2013).
30. Tonkin, J. D., Merritt, D. M., Olden, J. D., Reynolds, L. V. & Lytle, D. A. Flow regime alteration degrades ecological networks in riparian ecosystems. *Nat. Ecol. Evol.* 2, 86–93 (2018).
31. Liro, M. Dam reservoir backwater as a field-scale laboratory of human-induced changes in river biogeomorphology: A review focused on gravel-bed rivers. *Sci. Total Environ.* 651, 2899–2912 (2019).
32. Volke, M. A., Johnson, W. C., Dixon, M. D. & Scott, M. L. Emerging reservoir delta-backwaters: Biophysical dynamics and riparian biodiversity. *Ecol. Monogr.* 89, e01363 (2019).
33. Choudhury, S. Seasonal habitat use and resource partitioning between two sympatric crocoddilian populations (Gavialis gangeticus & Crocodylus palustris) in Katerniaghat Wildlife Sanctuary, India. *Master's thesis submitted to Saurashtra University, Rajkot, Gujarat, India* (2011).
34. MacClune, K. et al. Urgent case for recovery: What we can learn from the August 2014 Karnali River floods in Nepal. in *Technical Report*. Zurich Insurance Group Ltd, Zurich, ISET-International, Boulder 1–44 (2015).
35. Lang, J. W. & Kumar, P. Behavioral ecology of gharial on the chambal river, India. in *Crocodiles*. Proceedings of the 22nd Working Meeting of the IUCN-SSC Specialist Group. 42–52 (IUCN, Gland, 2013).
36. Lang, J. W. & Kumar, P. Chambal gharial population project-2010. in *Crocodiles*. Proceedings of the 24th Working Meeting of the IUCN-SSC Specialist Group. 136–148 (IUCN, Gland, 2016).
37. Gladfelter, S. R. Training rivers, Training people: Interrogating the making of disasters and the politics of response in Nepal’s lower Karnali River basin. Master’s thesis, University of Colorado (2017). https://broadresilience.net/resources/item/training-rivers-train-ing-people-interrogating-the-making-of-disasters-and-the-politics-of-response-in-nepals-lower-karnali-river-basin.
38. Kolbe, J. J. & Janzen, F. J. Impact of nest-site selection on nest success and nest temperature in natural and disturbed habitats. *Ecology* 83, 269–281 (2002).
39. Brown, G. P. & Shine, R. Maternal nest-site choice and offspring fitness in a tropical snake (Tropidonophis mairii, Colubridae). *Ecology* 85, 1627–1634 (2004).
40. López-Luna, M. A., Hidalgo-Mihart, M. G., Aguirre-León, G., González-Ramón, M. D. C. & Rangel-Mendoza, I. A. Effect of nesting environment on incubation temperature and hatching success of Morelet's crocodile (Crocodylus moreletii) in an urban lake of southeastern Mexico. *J. Therm. Biol.* 49, 66–73 (2015).
41. Calverley, P. M. & Downs, C. T. The past and present nesting ecology of Nile crocodiles in Nidumo Game Reserve, South Africa: Reason for concern?. *J. Herpetol.* 51, 19–26 (2017).
42. Somaweera, R., Brien, M. L., Platt, S. G., Manolis, C. & Webber, B. L. Direct and indirect interactions with vegetation shape crocodile ecology at multiple scales. *Freshw. Biol.* 64, 257–268 (2019).
43. Lang, J. W. & Andrews, H. V. Temperature-dependent sex determination in crocoddilians. in *Ess. Zool.* 270, 28–44 (1994).
44. Andrews, H. V. & Whitaker, R. Captive breeding and reproductive biology of the Indian Gharial Gavialis gangeticus (Gmelin), in *Crocodiles. Proceedings of the 17th Working Meeting of the IUCN-SSC Crocodile Specialist Group*. 401–411 (IUCN, Gland, 2004).
45. Rhen, T. & Lang, J. W. Phenotypic effects of incubation temperature in reptiles. in *Temperature-dependent sex determination in vertebrates* (eds. Valenzuela, N. & Lance, V. A.) 90–98 (Smithsonian Books, Washington, 2004).
46. Singh, V. P. Status of the gharial in Uttar Pradesh and its rehabilitation. *J. Bombay Nat. Hist. Soc.* 75(3), 668–683 (1979).
47. Basu, D. The gharial of Katerniaghat. *Sanctuary* 11, 36–43 (1991).
48. Srivastava, A. K. The biology of Indian gharial, Gavialis gangeticus, with special reference to its behaviour. PhD thesis submitted at University of Lucknow, Uttar Pradesh, India (1981).
49. Singh, V. P. Evaluation of gharial rehabilitation U.P. forestry project. Report prepared for biodiversity research, aided by World Bank, 1–49 (2003).
50. Andrews, H. V. Status of the Indian gharial, conservation action and assessment of key locations in North India. Unpublished report to Cleveland Metro Park. 1–8 (2006).
51. Whitaker, R. The gharial: Going extinct again. *Iguana* 14, 23–33 (2007).
52. Chaudhari, S. Gharial reproduction and mortality. in *Iguana* 15, 130–153 (2008).
53. Converse L. *Katerniaghat Gharial Project 2008–2009. Report of Preliminary Findings*. A Report to GCA and James Cook University, Australia. 1–8 (2009).
54. Das, A., Basu, D., Converse, L. & Choudhury, S. C. Herpetofauna of Katerniaghat Wildlife Sanctuary, Uttar Pradesh, India. *J. T. T.* 4, 2553–2568 (2012).
55. Choudhury, S., Choudhury, B. C. & Gopi, G. V. Differential response to disturbance factors for the population of sympatric crocoddilians (Gavialis gangeticus and Crocodylus palustris) in Katerniaghat Wildlife Sanctuary, India. *Aquat. Conserv.* 27, 946–952 (2017).
56. Kussussari, M. et al. Extinction debt: A challenge for biodiversity conservation. *Trends Ecol. Evol.* 24, 564–571 (2009).
57. Figueredo, L., Krauss, J., Steffan-Dewenter, I. & Sarmento Cabral, J. Understanding extinction debts: Spatio-temporal scales, mechanisms and a roadmap for future research. *Ecography* 42, 1973–1990 (2019).
58. Bashyal, A. et al. Gharials (Gavialis gangeticus) in Bardia National Park of Nepal: Population, habitat, and threats. *Aquat. Conserv. (in press).*
59. Grill, G. et al. Mapping the world’s free-flowing rivers. *Nature* 569, 215–221 (2019).
60. Jensen, J. R. Remote sensing of the Environment: An Earth Resource Perspective (Pearson Prentice Hall, Upper Saddle River, 2007).
61. Cohen, J. A. Coefficient of agreement for nominal scales. *Educ. Psychol. Meas.* 20, 37–46 (1960).
62. Killick, R., Haynes, K. & Eckley, I. A. Changepoint: An R package for changepoint analysis. R package version 2.2.2 (2016). https://CRAN.R-project.org/package=changepoint
63. Carpenter, S. R. & Kinne, O. Regime Shifts in Lake Ecosystems: Patterns and Variation, Vol. 15 (International Ecology Institute, Oldendorf/Luhe, 2003).
64. Whited, D. C. et al. Climate, hydrologic disturbance, and succession: drivers of floodplain pattern. *Ecology* 88, 940–953 (2007).
65. Heffernan, J. B. Wetlands as an alternative stable state in desert streams. *Ecology* 89, 1261–1271 (2008).
66. R Core Team. *R: A Language and Environment for Statistical Computing*. (R Foundation for Statistical Computing, Vienna, 2013). http://www.R-project.org/.
Acknowledgements
R. Whitaker (Madras Crocodile Bank Trust) and the late D. Basu studied and conserved the Katerniaghat gharial population for decades prior to our study, and shared their valuable observations and insights with us. We thank the Environment, Forest and Climate Change Department, Govt. of Uttar Pradesh, India for providing the necessary permissions and logistic support during the field work. GV was supported by Rufford Foundation small grants program (Project: 19031-1 and 24452-2), IDEAWILD and Gharial Ecology Project. DK received the DU-DST PURSE Grant Phase II. JWL received major funding for the Gharial Ecology Project from the City of Prague/Prague Zoo, Winter CROCfest 2018, Wildlife Conservation Society, Los Angeles County Zoo, Montgomery Zoo (Alabama), and the Berlin Zoo.

Author contributions
G.V., N.A.M., J.W.L. and D.K. conceptualized the study. G.V. and V.R. conducted the field work. N.A.M. collected and analysed the satellite imagery data. J.W.L. compiled the population count data. G.V. and N.A.M. wrote the original manuscript draft. D.K. and J.W.L. reviewed and revised the original draft. D.K., J.W.L., F.A.K. and P.M.D. arranged all the necessary permissions and funding for the field work. All authors have read and approved the manuscript for publication.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-021-84143-7.
Correspondence and requests for materials should be addressed to D.K.
Reprints and permissions information is available at www.nature.com/reprints.
Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021