# The taxonomy of Catocala nupta (Linnaeus, 1767) and its allies, with description of a new species (Lepidoptera: Noctuidae) 

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#### Abstract

The taxonomy of the Catocala nupta (L.), 1757 complex is analyzed using corresponding characters of the male genitalia, wing pattern, and COI 5 ' mitochondrial DNA, and is presently found to contain only two additional species: Catocala concubia Walker, [1858], and a new species Catocala benedeki described herein. Three new lectotypes are designated, and 12 status changes for existing names are introduced ( 10 new synonyms, 2 revised statuses).


Key words: Lepidoptera, Noctuidae, Catocala, new species, synonymy, Pakistan, China, India

## Introduction

Catocala nupta is the type species of the diverse Holarctic moth genus Catocala Schrank, and is one of approximately 40 species with red/pink-banded hindwings whose larvae feed on Salicaceae (willows and poplars). The taxonomy and nomenclatural histories of the Salicaceae-feeding Catocala are problematic (see Gall \& Hawks, 2002, 2010, 2015 for overview of the Nearctic taxa; comparable recent assessments of the Palearctic fauna are lacking). However, C. nupta is part of a morphologically well-defined species group, herein named the C. nupta species group. This group includes 7 currently recognized Palearctic species: C. nupta, C. concubia, Catocala szechuena Hampson, 1913, Cactoala amnonfreidbergi Kravchenko, Speidel, Witt, Mooser, Seplyarsky, Saldaitis, Junnila \& Muller, 2008, Catocala adultera Menetries, 1856, Catocala neglecta Staudinger, 1888, and Catocala laura Saldaitis, Ivinskis \& Speidel, 2008, as well as most Nearctic Salicaceae feeders (the four exceptions being Catocala amatrix (Hubner, [1813]), Catocala cara Guenée, 1852, Catocala carissima Hulst, 1880, and Catocala concumbens Walker, [1858]). These species are highly homogeneous in their genitalic structure. Exemplary synapomorphies include: left valve saccular extension along ventral margin of cucullus $2-3$ times as long as wide (Figs. 41, 55-61); right valve saccular extension (same); dorsal side of right valve costa with a distinct notch just anterior of midpoint (Fig. 54); large patch of sclerotization surrounded by membrane in the anterior half of the right cucullus (Figs. 41, 55-61); ventral side of left clasper convex proximally and concave distally, dorsal side of left clasper convex proximally and concave distally (Figs. 41, 55-61) (except in C. neglecta and C. laura); and vesica with numerous small sclerotized plates adjacent to ventral aedeagus hood (sometimes fused together) (Fig. 62).

Among the seven Palearctic species noted above, C. nupta, C. concubia, C. szechuena and C. amnonfreidbergi are the most closely similar in wing pattern, and are referred to herein as the C. nupta complex. These four are all readily distinguishable from the differently patterned C. neglecta, C. laura, C. adultera, and all of the related Nearctic species. Below we summarize the results of our studies on the Catocala nupta complex, using a combination of genitalic, wing pattern, and COI 5' mitochondrial DNA character systems (see Table 1, Appendix $1)$.

TABLE 1. Variable COI 5' characters among C. benedeki, C. concubia, \& C. nupta. Bold font character states in larger font size are unique to one species among these three; underlined character states are infraspecific polymorphisms.

|  | 49 | 85 | 88 | 274 | 325 | 335 | 343 | 433 | 460 | 520 | 547 | 548 | 574 | 607 | GenBank |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Catacala benedeki 6015 India | A | T | I | C | T | G | C | A | C | A | T | C | T | C | GU678880 |
| Catacala benedeki 9004 Pakistan | A | T | I | C | T | G | C | A | C | A | T | C | T | C | HQ970472 |
| Catacala benedeki 10249 Pakistan | A | T | C | C | T | G | C | A | C | A | T | C | T | C | KT960834 |
| Catacala benedeki 20430 Pakistan | A | T | C | C | T | G | C | A | C | A | T | C | T | C | KT960832 |
| Catacala benedeki 20431 Pakistan | A | T | C | C | T | G | C | A | C | A | T | C | T | C | KT960833 |
| Catacala benedeki 20436 Pakistan | A | T | C | C | T | G | C | A | C | A | T | C | T | C | KTPL0835 |
| Catocala concubia 20432 China | A | T | T | T | T | G | T | A | T | A | T | T | T | T | KT690828 |
| Catocala concubia 2121 China | A | T | T | T | T | G | T | A | T | A | T | T | T | T |  |
| Catocala concubia 2122 China | A | T | T | T | T | G | T | A | T | A | T | T | T | T |  |
| Catocala concubia 8000 China | A | T | T | T | T | G | T | A | T | A | T | T | T | T | HM426442 |
| Catocala nupta 9564 China | A | I | T | T | I | G | C | A | C | A | C | C | I | C |  |
| Catocala nupta 10250 China | A | I | T | T | I | G | C | G | C | A | C | C | I | C | KT960831 |
| Catocala nupta 20439 China | A | I | T | T | I | G | C | G | C | A | C | C | I | C | KT960829 |
| Catocala nupta 8002 China | A | I | T | T | I | G | C | $\underline{\mathrm{G}}$ | C | A | C | C | I | C | HM426444 |
| Catocala nupta 8004 E Russia | A | I | T | T | I | G | $\underline{C}$ | G | C | A | C | C | I | C | HN426446 |
| Catocala nupta 8071 China | A | I | T | T | I | G | C | G | C | A | C | C | I | C | HM426488 |
| Catocala nupta 8003 China | A | I | T | T | I | A | C | G | C | A | C | C | I | C | HM426445 |
| Catocala nupta 9218 Japan | $\underline{C}$ | I | T | T | I | G | C | G | C | A | C | C | C | C | JF860521 |
| Catocala nupta 9219 E Japan | C | I | T | T | I | G | C | G | C | A | C | C | C | C | JF860522 |
| Catocala nupta 5625 E Russia | C | $\underline{\text { C }}$ | T | T | I | G | $\underline{\text { C }}$ | G | C | A | C |  | $\underline{C}$ | C |  |
| Catocala nupta 8005 E Russia | C | $\underline{C}$ | T | T | I | G | $\underline{C}$ | $\underline{\mathrm{G}}$ | C | A | C | C | C | C | HN426447 |
| Catocala nupta MM15851 Finland | A | I | T | T | C | G | I | A | C | G | C | C | I | C | HM876626 |
| Catocala nupta 20019 Kyrgyzstan | A | I | T | T | C | G | I | A | C | A | C | C | I | C | KT960830 |
| Catocala nupta 20394 Kyrgyzstan | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta 2332 Slovakia | A | I | T | T | $\underline{C}$ | G | T | A | C | A | C | C | I | C |  |
| Catocala nupta MM04216 Finland | A | I | T | T | $\underline{C}$ | G | I | A | C | A | C | C | I | C | HM872333 |
| Catocala nupta MM04750 Finland | A | I | T | T | C | G | T | A | C | A | C | C | I | C | HM872531 |
| Catocala nupta 8001 W Russia | A | I | T | T | C | G | I | A | C | A | C | C | I | C | HM426443 |
| Catocala nupta BC ZSM Lep 21863 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C | JF415760 |
| Catocala nupta BC ZSM Lep 29037 Germany | A | I | T | T | $\underline{C}$ | G | I | A | C | A | C | C | I | C | JF415759 |
| Catocala nupta BC ZSM Lep 64831 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta BC ZSM Lep 64832 Germany | A | I | T | T | C | G | I | A | C | A | C | c | $I$ | C |  |
| Catocala nupta BC ZSM Lep 70164 Germany | A | I | T | T | C | $\underline{\text { G }}$ | I | A | C | A | C | C | I | C |  |
| Catocala nupta BC ZSM Lep 72632 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta BC ZSM Lep 75217 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta BC ZSM Lep 78585 Germany | A | I | T | T | $\underline{C}$ | G | I | A | C | A | C | C | $\mathrm{I}$ | C |  |
| Catocala nupta BC ZSM Lep 80284 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta BC ZSM Lep R 21673 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta P21863 Germany | A | I | T | T | C | G | I | A | C | A | C | C | I | C |  |
| Catocala nupta TLMF Lep 09855 Austria | A | I | T | T | C | G | I | A | C | A | C | C | I | C | KM572102 |

## Materials and methods

Methods for data analysis, genitalic dissection, and terminology follow Kons and Borth (2015) and as described at http://www.lepidopterabiodiversity.com. Paul Hebert's lab at the University of Guelph sequenced the 5' region of COI as described in Hebert et al. (2003). Statistical analyses were performed in v9.4 of SAS for Windows (SAS Institute, Cary, NC). Acronyms for institutional and private collections are as follows: AFM, Alessandro Floriani (Milan, Italy); ASV, Aidas Saldaitis (Vilnius, Lithuania); NHM, Natural History Museum (London, England); KIS, Katsumi Ishizuka (Saitama, Japan); MNHU, Museum fur Naturkunde (Berlin, Germany); RJB, Robert J. Borth (Milwaukee, USA); SEHU, Hokkaido University Museum (Hokkaido, Japan); ZSM, Zoologische

Staatssammlung, München (Munich, Germany); ZFMK, Zoologisches Forschungsmuseum, A. Konig (Bonn, Germany); WIGJ, World Insect Gallery (Joniškis, Lithuania).

## Systematic part

We find that in the Catocala nupta complex, only C. nupta and C. concubia are diagnosable by any discrete morphological characters. Exemplary differences in upperside wing pattern that separate C. nupta from $C$. concubia include: the medial red band on the hindwing is nearly concolorous throughout its length in C. nupta, but is noticeably paler white with only sparse red scaling along the leading edge of the hindwing in C. concubia; the inner hindwing margin has red/dark hairs in C. nupta, but red orange hairs in C. concubia; and the forewing of $C$. concubia has scattered pale violet tinted scales, and C. nupta lacks these where a bluish tint is sometimes evident. Specimens we have measured of C. concubia average 38.0 mm ( $\mathrm{sd}=0.7 \mathrm{~mm}, \mathrm{n}=5$ ) in forewing length (base to apex) compared to 33.2 mm in C. nupta ( $\mathrm{sd}=1.9 \mathrm{~mm}, \mathrm{n}=43$ ), and this size difference is significant ( $\mathrm{F}=33.91, \mathrm{df}=1 / 44, \mathrm{p}<$ 0.001 , by ANOVA controlling for sex). There are also several character state differences in COI 5' mtDNA sequences between $C$. concubia and $C$. nupta (Table 1). Since C. concubia has been treated inconsistently in the Palearctic literature as a synonym of C. nupta (e.g., Hampson 1913, Poole 1989, Kononenko 2010), a subspecies (Goater et al. 2003: 84), or a full species (e.g., Warren 1913, Goater et al. 2003: 83) we here reaffirm specific rank for C. concubia, stat. rev. This species occurs from Sichuan and Yunnan Provinces in China west to Punjab in northern India.

We also find no justification for subdividing C. nupta into further infraspecific taxa. Catocala nupta is the most geographically widespread of the Palearctic Catocala species, ranging from Portugal and England in the west to Russia and Japan in the east. The wing pattern of C. nupta exhibits considerable inter- and infrapopulational variation, but our research shows this variation to be continuous, and the differences in COI $5^{\text {‘ }}$ haplotypes seen within C. nupta do not correspond to known morphological differences (Table 1; see Diagnosis section below).

In addition, we have identified an additional diagnosable species from the northern region of the Indian subcontinent close to $C$. concubia that lacks an available name, which we describe as new.

## Catocala benedeki Borth, Kons, Saldaitis \& Gall sp. nov.

(Figs 1-5, 38-55, 62-69, 77, 78, 85-96)

Type material. Holotype: Male (Fig. 1), Pakistan, Karakoram Mts., Deitar Valley, 3000 m, 27.VII.2011, leg. Balázs Benedek (Dissection No. 2015HLK:2120; DNA No. 10249-270711-PA) preserved in ASV collection, later to be deposited in the WIGJ.

Paratypes: (Figs 3-4), PAKISTAN: 1 male, Karakoram Mts., Deitar Valley, 3000 m, 27.VII.2011, leg. Balázs Benedek (Dissection No. 2014HLK:2046; DNA No. 20430-COI-14) (collection of (coll.) RJB); 1 male Karakoram Mts., Deitar Valley, 3000 m, 27.VII.2011, leg. Balázs Benedek (coll. AFM); 1 male Kashmir, Karakoram Mts., Deitar Valley, 3000 m, 29.VI.2014, leg. Balázs Benedek (coll. RJB); 1 female, Himalaya Mts., Bubin Valley, 3000 m, 29-31.VII.2011, leg. Balázs Benedek (Dissection No. 2014HLK:2081; DNA No. 20431-COI-14) (coll. RJB); INDIA: 1 male, Prov. Uttar Pradesh, Garwahl Himal, Gangotri, 3000 m, 4.1X.1997, L. Nadai (Dissection No. JB1071) (coll. AFM); 1 female, Kashmir, PirPanJar, Tatakutti, Astar, 4000m, 11.VIII. 2012, leg. T. Surumaki (coll. KIS); 4 males, S. Kashmir, Tatakutti Area, PirPanJal Mts., Mt. Ashitar, 3800m, 2-3.VIII.2013, leg. T. Surumaki (coll. KIS).


FIGURES 1-12. Imagos of Palearctic nupta group: C. benedeki, C. concubia, C. nupta, C. adultera, C. neglecta \& C. laura.


FIGURES 13-22. Imagos of high altitude Catocala from India \& China: C. concubia, C. prolifica \& C. fuscinupta.


FIGURES 23-37. Imagos of C. nupta types compared to similar sequence vouchers.

Diagnosis. The wing pattern of C. benedeki (Figs 1-4) resembles the closely related C. concubia (Figs 6a-7, 13-20) and C. nupta (Figs 8, 9, 23-37). Both Catocala fuscinupta Hampson, 1913 (Fig. 22) and Catocala prolifica Walker, [1858] (Fig. 21) are superficially similar in wing pattern (but are dissimilar in genitalia), and can be readily separated by the ventral hindwing median band, which is distinctly curved basally between veins Rs and M2 in $C$. benedeki, and not so in either C. fuscinupta or C. prolifica. The following characters enable the separation of $C$. benedeki, C. concubia, and C. nupta. Forewing upperside: C. benedeki has mottled grey and white forewings with strong contrasts in maculation, whereas C. concubia has a more diffused, less distinct pattern (especially in males) with an olive and brownish cast. The smaller C. nupta is highly variable: some specimens are grey with white, but they have lower densities of white mottling giving the wings a smoother, less powdery appearance. Catocala benedeki has pale violet tinted scales, which are present to a lesser extent in C. concubia, but are lacking in C. nupta where a bluish tint is sometimes evident. Hindwing upperside: In $C$. benedeki the inner hindwing margin has dense pale tan hairs contrasting against the red background color, C. nupta has red or red and dark hairs, and $C$. concubia has red orange hairs. The area posterior to vein Rs between the medial and marginal black bands is pale white with sparse red scaling in $C$. benedeki; this area is also pale in C. concubia with slightly denser red orange scaling, but in C. nupta there is dense red scaling with much less pale contrast. Wing size: The average forewing length (base to apex) in measured specimens of $C$. benedeki is $42.0 \mathrm{~mm}(\mathrm{sd}=1.0 \mathrm{~mm}, \mathrm{n}=3)$, versus 38.0 mm ( $\mathrm{sd}=0.7$ $\mathrm{mm}, \mathrm{n}=5$ ) in C. concubia, and 33.2 mm ( $\mathrm{sd}=1.9 \mathrm{~mm}, \mathrm{n}=43$ ) in $C$. nupta. The differences among species are significant ( $\mathrm{F}=46.36, \mathrm{df}=2 / 45, \mathrm{p}<0.001$, by ANOVA controlling for sex, pairwise species comparisons $\mathrm{p}<0.05$ using Tukey least squares means). Male genitalia: While we compared all of the male genitalic characters covered in the description (below) among C. benedeki ( $\mathrm{n}=2$ ) (Figs $38-55,62-69,77,78$ ), C. concubia ( $\mathrm{n}=2$ ) (Figs 56, 57, $71,72,79,80$ ), and C. nupta ( $\mathrm{n}=4$ ) (Figs 58-61, 73-76, 81-84), the only genitalic characters found to vary between these three species were minor but consistent differences in vesica diverticula 6 and 7 . While purported differences in the length, shape, or sclerotization of valve costa have been suggested as a basis for subdividing C. nupta into additional species or subspecies (e.g., Kravchenko et al. 2008) we found no evidence these characters have diagnostic value at the species level within the Catocala nupta complex (Figs 41, 55-61). The following vesica characters are best viewed when the vesica is orientated in lateral aspect with the ventral aedeagus hood on top. In C. benedeki and C. concubia diverticulum 6 has two shallow but distinct subdiverticula (Figs 76-79), whereas $C$. nupta has three (Figs 80-83); the equivalent of subdiverticulum 6 b in C. benedeki and C. concubia is subdivided in C. nupta (Figs 80-83). Catocala benedeki and C. nupta have two distinct subdiverticula on diverticulum 7 (Figs 67-69, 72-75) whereas in C. concubia diverticulum 7 is expanded on the anterior side in the equivalent position but there is no clear bifurcation (Figs 70-71). In C. benedeki the ventral base of subdiverticulum 7a is closer to the anterior base of diverticulum 7 than in C. nupta (Figs 68-69 versus Figs 72-75). In C. benedeki and C. nupta diverticula 6 and 7 diverge from near the posterior base of diverticulum 6 and the anterior base of diverticulum 7, creating a distinct separation near the base; the diverticula touch or nearly touch farther distally where diverticula 7 expands laterally on the anterior side (Figs 67-69, 72-75). However, in C. concubia diverticula 6 and 7 touch throughout the basal half of their length or more (Figs 70-71). When viewed laterally with a slight lateral tilt such that diverticulum 7 appears as wide as possible, subdiverticulum 7 b is distinctly narrower in $C$. benedeki compared to C. nupta (Figs 68-69 versus 74-75; in Figs 72-73 diverticulum 7 is slightly tilted from where it appears widest, causing diverticulum 7 b to appear narrower). Female Genitalia: No diagnostic characters are known between $C$. benedeki, C. concubia, and C. nupta. COI 5' Mitochondrial COI 5' DNA: Catocala benedeki can be diagnosed from all sequenced Catocala species by the following unique combination of six character states: 274(C), 343(C), 460(C), 547(T), 548(T) and 607(C). There are two consistent character state differences between C. benedeki and C. nupta, and five between C. benedeki and C. concubia (see Table 1).

Description. Head. Vertex with predominantly grey scales, peppered with darker grey, white, and tan scales. Frons predominately grey but peppered with white and tan scales. Labial palp basal segment almost exclusively white, with sparse grey scales on the lateral side; middle segment predominately grey with a peppering of white scales increasing in density distal to proximally; terminal segment predominately grey with a peppering of white scales. Antennae dorsally and laterally predominantly covered by grey scales with scattered white scales, except for scape and pedicel which are predominately white with sparse grey scales.

Thorax. Patagia predominantly grey with a dense scattering of white and tan scales. Tegulae with heavily mottled with grey and white scales, narrow irregular band along inner margin mottled with tan and darker grey scales. Elsewhere a mix of grey, tan, and white scales, sometimes with a diffuse grey and white inverted "U"
pattern. Paired tufts of hair on posterior mesothorax predominately tan with some grey. Ventrally with dense pale tannish-white hairs.

Wings (Figs 1-4). Wingspan: Forewing length (base to apex) of holotype male 41 mm ; forewing length of paratypes: male $(\mathrm{n}=1) 42 \mathrm{~mm}$, female $(\mathrm{n}=1) 43 \mathrm{~mm}$. Forewing Shape: length of FW base to apex divided by length of FW apex to anal angle: holotype male 1.77 , paratype male 1.68 , paratype female 1.75 . Forewing upperside: Background color predominantly grey, mottled with white scales creating a powdery appearance, many white scales with a distinct violet tint under natural light. Two small areas where white scales dominate: on the basal side of the reniform and orbicular spots, and between the postmedial and subterminal lines between veins M2 and M3. Reniform spot oblong, basal side smoothly curved, distal side irregular to slightly dentate, primarily with dark greyish-black scales, diffuse lighter grey scales in center. Reniform border with a mix of pale brown and light grey scales, rimmed with a smooth basally curved greyish black basal border, fused with the medial band on anterior and posterior sides of reniform. Basal dash absent. Basal line sharp greyish-black posterior of discal cell, comprised of an anterior distally curved loop and a posterior fairly straight line. Antemedial line distinctly double, dark grey to greyish black borders with pale whitish-grey between, borders variably sharp or diffuse. Antemedial line comprised of three distinct distally curved loops: posterior loop (below vein 2 A ) protruding basally on anterior side; second (medial) loop spanning between veins 2 A and lower margin of discal cell; third loop irregular, from lower discal cell margin to costa, distal border doubly looped, basal border a single jagged loop. Median line distinct the full length of the wing except where broken by the open base of the subreniform spot, black, sharp anterior to reniform and posterior to vein Cu 2 , generally diffuse between. Median line irregularly undulated to dentate, double between costa and anterior base of subreniform, the two sections widely separated anterior of vein R4, fused together at distal anterior corner of reniform and intersection of veins R4 and R5, then narrowly separated and fusing back together at anterior base of subreniform. Postmedial line black, sharp, distinctly contrasting, bordered distally by a sharp to diffuse narrow band of pale whitish grey. Postmedial line undulations: below vein 2 A convex loop, extended as far basal as edge of reniform at vein 2 A ; between Cu 2 and 2 A singly or doubly dentate; subreniform open, but pale band on distal side of postmedial line fused; tooth between veins Cu 1 and Cu 2 with apex close to vein Cu 2 ; shallow tooth between veins M 3 and Cu ; two dentate distally protruding teeth between veins M1 and M3 with a concave division between them across vein M2; concave and angling basally between veins R5 and M1, then sharply turned basally along vein R4 roughly perpendicular to costa and thickened as small black patch slightly distal to the outer border of the reniform. Subterminal line broad and diffuse pale whitish grey, a series of undulated to dentate, distally protruding chevrons with diffuse to indistinct black distal border. Wing margin with series of diffuse to sharp black, straight to slightly concave bars or patches between each pair of veins from R4 to CuA2. Farther distally at the extreme margin, a sinusoidal, broken, thin dark grey line between veins R4 and 2A, with the distal protruding convex loops transecting the veins and basally protruding concave loops between the veins. Fringe peppered pale whitish grey and darker grey, the darker grey progressively increasing in density posterior to anterior, without patches along veins. Hindwing upperside: Background color reddish (with a slight pink tinge) posterior to vein Rs, predominately pale white anterior to vein Rs, peppered with a variable amount of the reddish background scales. Black median band prominent, distal side with three distal undulations and smoothly angled along vein M2, basal side a fairly smooth simple distal curve; bulged basally between costa and vein M2, and bulged distally between veins M2 and Cu1, oblong patch with no distinct curvature in cell CuA 2 ; thin, diffuse, and discontinuous between vein 2A and inner margin. Marginal black band thick and fairly smoothly curved anterior to vein Cu 2 , deep concave gouge in interspace between veins Cu 2 and 2 A , each side of gouge with triangular basally pointing projection with apices on veins 2 A and CuA 2 , thin and narrow at anal angle. Band of dense pale tan hairs along inner margin, contrasting against background colour, some darker grey hairs mixed in, tan and grey hairs along the veins basal to the median band. Fringe thick and clean, bright white, some small patches of pale white intruding into the marginal black band between the veins such that the distal edge of the marginal band is undulated between M1 and CuA2. Apical patch prominent, a paler white than the fringe, basal edge with two basally protruding undulations divided by vein Rs. Anterior margin with a thin band of white. Forewing underside: Background color bright, clean white. Marginal band thick and black with sharp margins, except on distal side at apex, where it blends into a greyish white area at apex; outer side with two broad undulations divided by vein M2, a distally protruding triangular indentation at the veinlet posterior of vein CuA 2 , a smaller triangular indentation of white along vein 2 A , small slivers of white intruding along some of the other veins. Termen with the thin undulating line present dorsally, but more diffuse and becoming progressively
less distinct posteriorly. Median band wide and black; sharply contrasting but with some whitish diffusion along the edges; both sides angle distally along vein M2, progressively narrowing between vein M2 and inner margin, tapering to a blunt point at inner margin; a distally protruding triangular tooth at veinlet posterior of vein CuA2, with the tooth apex on the veinlet. Basal band diffuse and lighter black; both margins irregular; extends between anterior margin of discal cell and vein 2 A , but costa anterior to discal cell and inner margin posterior to vein 2 A white with sparse black diffusion. Incomplete diffuse marginal black band between basal and median bands between the veinlet posterior of vein CuA 2 and 2 A , becomes more diffuse distally and partially fuses with median band. White background color basal to basal band with a pale tan tint. Hindwing underside: Marginal black band similar to upperside but slightly narrower and with more diffused margins, and not extending anterior of vein $\mathrm{Sc}+\mathrm{R} 1$ where this area is white with a peppering of black scales. Medial black band of dissimilar shape to upperside; relatively thin anterior to vein Rs, slightly narrowing anteriorly, fairly straight but with diffused and slightly irregular margins; distinctly curved basally between veins Rs and M2; widest at intersection with vein M2, strongly tapering between veins M2 and CuA2 with an undulated distal edge; similar to upperside in cell CuA2, but with a faint but distinct distal curve; absent between vein 2A and inner margin. Small, diffuse, black discal spot on the veinlet. Background color: basal to medial band: white anterior to posterior margin of discal cell, pale red with some white diffusion posteriorly; distal to medial band: white anterior to vein CuA 1 , pale red with some white diffusion posteriorly. Fringe, apical patch, and extreme outer margin similar to upperside.

Legs. Similar in male and female except male mesotibia has a hair pencil groove on the inner side and is slightly wider. Foreleg: Profemur without laterally flattened apical spine. Protibia unspined, but with small convex sulcus with radiating spines near basal extremity on the inner side. Protibial flange in shallow ovuloid pit, ventral margin of flange with dense row of short setae. Protarsomeres $1-4$ with three ventral rows of large triangular spines, and two rows of minute hair-like curved spines between them; protarsomere 5 with four rows of large triangular spines, with two rows of minute hair-like spines in-between. Irregularly spaced hair-like spines present on lateral sides of tarsomeres. Pretarsus simple. Arolium convex on basal side, widening distally with lateral margins concave and then convex, distal margin slightly concave, sclerotization predominately dark brown to blackish, most intense along basal and basal-lateral sides, translucent greyish on distal and distal-lateral margins. Midleg: Mesofemur unspined, mesotibia with a single row of fourteen to fifteen large spines on outer side, hairpencil groove on inner side. Tarsal spination/pretarsus/arolium as in proleg. Hindleg: sclerotization pattern typical for Catocala, with femur sclerotized throughout, mesotibia translucent white except at base, metatarsomere 1 translucent white except at apex, remaining tarsomeres sclerotized throughout. Metafemur unspined, metatibia with three subapical spines on outer side near ventral margin, two thicker spines ventral to one much thinner spine. Tarsal spination/pretarsus/arolium like proleg except metatarsomere 1 with a few extra smaller spines scattered between the three rows of ventral spines.

Abdomen. Scale pattern grey dorsally, white and pale tan ventrally. Cuticle (Figs 51-53, 92-94): male tergites $1-6$ as shown in Fig. 51, sternites $2-7$ as shown in Fig. 52, terminal tergite (left) and sternite (right) as shown in Fig. 53. Female tergites $1-6$ as shown in Fig. 92, sternites $2-6$ as shown in Fig. 93, tergite 7 (Fig. 94) with strongly convex sides, convex posterior side, anterior side with shallow concave depressions on each side and fairly straight in the middle.

Male genitalia (Figs 38-55, 62-69, 77, 78). Capsule (Figs 38-41, 48): Juxta and vinculum strongly fused with valvae, vinculum weakly fused with tegumen, vinculum arms laterally expanded and weakly fused midventrally, diaphragma membranous except for juxta/anellus. Valvae (Figs $41 \& 55$ ): asymmetrical. Outer surfaces densely covered with elongate tan colored hairs and scales except for anterior portion of sacculus (Fig. 39); inner surface of "cucullus" (or the membranous valvae structure in the equivalent position) with shorter scales and hairs along ventral margin. Sacculus with triangular posterior extension at fusion with cucullus, approximately twice as long as wide; inner side densely covered with elongate setae (Figs $41 \& 55$ ). Additional elongate setae scattered along posterior margin of sacculus on inner side. Ventral inner sides of sacculus with concave indentation along margin of clasper base (Fig. 38). Left cucullus clear and membranous. Right cucullus with an ovuloid patch of sclerotization between clasper and ventral margin of cucullus, additional sclerotization bordering the cucullus medially, otherwise clear and membranous (Figs $41 \& 55$ ). Cucullus with scattered elongate setae on inner surface along ventral margin, densest anteriorly. Left costa (Figs $41 \& 55$ ) heavily sclerotized, widest medially, narrowest posteriorly; ventral margin concave anteriorly, convex medially, and concave posteriorly; dorsal margin strongly convex anteriorly, more weakly concave medially, then weakly convex, concave, and convex posteriorly (anterior


FIGURES 38-54. Male genitalia and cuticle of $C$. benedeki Holotype; 55-61: valvae (inner) of C. benedeki, C. concubia and C. nupta.


FIGURES 62-67. Everted vesica at different angles for C. benedeki.


FIGURES 68. Everted vesica of $C$. benedeki (lateral aspect with ventral aedeagus hood orientated up); 69: Diverticulum 7 of C. benedeki (lateral aspect with ventral aedeagus hood orientated up); 70: Diverticulum 7 of C. benedeki (lateral aspect with ventral aedeagus hood orientated down); 71-72: Diverticulum 7 of C. concubia (lateral aspect with ventral aedeagus hood orientated up); 73-76: Diverticulum 7 of C. nupta (lateral aspect with ventral aedeagus hood orientated up) [Note: In Fig. 73 the apex of 7 a is partially inverted inward making it appear shorter]. Solid arrows point to the boundaries of subdiverticulum 7a; dashed arrows point to the anterior base of diverticulum 7. Note: It is impossible to photograph diverticulum 7 in this aspect at exactly the same angle between specimens, and some apparent minor differences in shape within species are incidental to minor differences in the angle between the photographs.
to posterior); costa extends distinctly beyond cucullus where it projects ventro-posteriorly. Right costa (Figs 41 \& 55) dissimilar to left; distinct thickened anterior section terminates anterior of clasper apex, posterior to this the "costa" may be modified sclerotized cucullus, sclerotization less intense than on left costa in this area; ventral margin concave anteriorly, weakly convex medially, weakly concave posteriorly; dorsal margin strongly convex anteriorly, beyond thickened area convex overall but exact shape somewhat variable; extends slightly posteriorly beyond membranous cucullus, but shorter than left costal extension. Dorsally left costa smooth and narrow, right costa with a distinct notch slightly anterior to the clasper apex (Fig. 54, arrow). Claspers asymmetrical (Figs 41 \& 55). Left clasper ventral side convex basally, concave medially-distally, convex at apex; dorsal side convex basally, concave distally. Right clasper ventral margin concave, dorsal margin also concave but relatively less so. Left clasper base much wider than left (Figs $41 \& 55$ ), left clasper but not right curved ventrally at apex (Figs 41 \& 55), both clasper apices curved inward but right apex more strongly so (Fig. 38). Clasper apices covered with scattered minute short setae, apices with knob-like expansions (Fig. 20). Ventral and dorsal margins of each clasper base with patches of elongate setae, with scattered shorter setae extending distally along ventral margin. Juxta (Fig. 49): Two elongate slightly asymmetrical lobes, right lobe slightly shorter, both narrowest posteriorly, progressively widening anteriorly, with a narrow band of darker sclerotization along inner margin. Lobes narrowly fused to anellus at posterior apex, barely touching each other near posterior end but not fused together. Pitted pattern of anellus not extending to juxta lobes. Anellus (Fig. 49): Lobes fused together throughout and appearing as a single sclerotized plate, strongly asymmetrical with right lobe larger and wider than left. Outer margin of left lobe strongly concave but with small convex protrusion medially, outer margin of right lobe concave, posterior apex narrowly rounded, anterior edges of both lobes convex. Band of sclerotized dense shallow depressions (pits) of variable lateral width throughout longitudinal midline. Uncus (Figs 45-46): Tubular, progressively gradually narrowing distally, posterior margin strongly convex, anterior margin strongly concave; terminating in heavily sclerotized curved spine, laterally appearing pointed apically but narrowly rounded in dorsoventral view. Extensive lateral setae throughout length, most not longer than width of uncus but longest setae 2-3 X width of uncus and located in medial and subapical areas. Tuba analis (Fig. 44-45): Membranous except for scaphium, scaphium an elongate rectangular plate terminating slightly dorsally to the uncus apex. Aedeagus (Figs 42-43, 47): Translucent throughout. Coecum of similar width to adjacent aedeagus, weakly bent. Aedeagus bent at posterior margin of coecum, and strongly bent before ventral extension over vesica, fairly straight in-between. Left flank of posterior ventral extension ("hood" over everted vesica) with a concave gouge and a convex posterior-lateral expansion (Fig. 67), right flank weakly convex (Figs $62 \& 67$ ), apex concave medially and convex on edges (Fig. 62). All of the four sclerotized chords sometimes present on the ventral hood in Catocala are clearly present (Fig. 67): left outer chord (LOC) extends almost to the apex of the hood (Figs $62 \& 67$ ); left inner chord (LIC) terminates well anterior of apex, roughly parallel with LOC at terminus (Figs 62 \& 67); right inner chord (RIC) terminates about even or just anterior to anterior side of concave depression on left flank of hood, merges with right outer chord (ROC) on both ends (Fig. 67); ROC gradually weakens posteriorly along outer edge of hood with no clear terminus (Fig. 62). Ductus ejaculatorius (Fig. 50): Slender region with distinct bend just before scoop-shaped region, bent over at about 180 degrees. Scoop-shaped region strongly convex on outer side with prominent concave gouge basally, inner side convex at base, then strongly concave. Vesica (Figs 62-66, 68, in part Figs 69-84): Vesica diverticulum 1 trilobed, concealed underneath diverticula 2 at most angles but best seen in ventral view with the hood slightly tilted to the left (Fig. 62), or in anterior aspect with the focal plane underneath diverticulum 2; 1a the most elongate of the three lobes, fang-shaped with outer side convex, inner side concave, and apex narrowly rounded; 1 b and lc finger-like with convex apex, 1b about half as wide as 1c (Fig. 62). Diverticulum 2 (Figs 63-64 \& 66) lacking subdiverticula, elongate and gradually tapering to a narrowly rounded apex, strongly bent in two places. Diverticulum 3 not clearly bifurcate but wide with convex bulges on each side, fairly straight in-between (Figs 6364). Diverticulum 4 clearly bifurcate with two distinct convex bulges on each side and concave in-between (Figs 63-64). Diverticulum 5 with three lobes, the central ( 5 a ) a broadly rounded convex bulge; the left ( 5 b ) somewhat fang-shaped, narrowly rounded (almost pointed) apically, with the left side convex and the right side concave; the right ( 5 c ) a broadly rounded shallow convex bulge barely distinct from 5 a and separated by a shallow convex to nearly straight separation (Figs 63-64). Diverticulum 6 large and broad, with two shallow but discernible convex lobes at apex (Figs 68, 77-78). Diverticulum 7 distinctly bilobed with posterior/distal lobe (7b) longer and narrower than anterior lobe (7a), 7b protruding from posterior corner, 7a protruding from anterior side, posterior side of 7 roughly parallel to anterior side of diverticulum 6 , anterior base of 7 diverging from base of 6 with a distinct separation, but touching or nearly so farther distally where 7a bulges out (Figs 68-70). Diverticulum 8


FIGURES 77-84. Diverticulum 6 of C. benedeki (Figs 77-78), C. concubia (Figs 79-80), and C. nupta (Figs 81-84) in lateral aspect with the ventral aedeagus hood orientated up. In C. nupta diverticulum 6 has three shallow but distinct subdiverticula; in C. benedeki and C. concubia there are two. Solid arrows point to the dorsal margin of subdiverticulum 6 a ; dashed arrows point to the anterior edge of subdiverticulum 6c. Note: It is impossible to photograph diverticulum 6 at exactly the same angle between specimens, and some apparent differences in shape within species are due to minor differences in the angle between the photographs, especially between Figs 79 and 80. FIGURE 85. Habitat of C. benedeki: Kashmir, Lower Bubin Valley, 3000 m.


FIGURES 86-96: Female genitalic structures and abdominal cuticle of $C$. benedeki (HLK:2081).
broad but shallow convex bulge curving around right and apical sides of ventral aedeagus hood, several times as wide as high (Figs $62 \& 68$ ). Diverticulum 9 a simple convex bulge (Figs $63 \& 65$ ). Diverticulum 10 a broad simple convex bulge wider than high (Figs $62 \& 65$ ). Diverticulum 11 not discernible. Diverticulum 12 a prominent convex bulge (Figs 62-63, 66) overlapping most of the ventral aedeagus hood when viewed in the flattest ventral aspect (Fig. 67). Diverticulum 13 a prominent, fairly narrow convex bulge, longer higher than wide, paralleling the distal margin of diverticulum 3 (Fig. 64). Much of vesica covered with minute inward projecting triangular teeth, proximal teeth on diverticulum 8 bordering the right flank of the aedeagus hood larger, sclerotized, some with the sclerotization fused together into a narrow triangular plate (Fig. 62).

Female genitalia ( $\mathrm{n}=1$ ) (Figs 86-96): Papillae analis (Figs 87, 89, \& 96): Transparent except for a broad band of light dorsal sclerotization on each papillus. Longest setae at base, projecting posterior/outward. Shorter setae throughout papillae project posterior/outward or perpendicular. Apices and medial area densely covered with short setae (shorter than width of papillus) in addition to less dense longer setae of variable lengths. Papillae curved such that dorsal side strongly convex, ventral side convex proximally and distally, strongly concave medially, apex broadly rounded. Papillae of similar width throughout most of length, widest proximally and subapically and weakly constricted medially, progressively tapering apically. Intersegmental membrane between papillae and segment 8 (Fig. 87): Narrowing anterior to posterior, anterior end approximately 1.4 times width of anterior end.

Length/width at anterior end $=1.5$. Segment A8 (Fig. 87): Ventrally extending well posterior of lamella antevaginalis (only slightly overlapping on anterior side), shape as shown in Fig. 87; dense, elongate, posteriorly projecting setae encircling A8 posterior margin and much of dorsal and lateral sides, absent in area of ventral midline; thin heavily sclerotized U-shaped band connecting posterior margins of plates on ventral side. Intersegmental membrane between lamella and segment 8 on ventral side (Fig. 87): Slightly curved (concave), moderately sclerotized and with minute spiculations (Fig. 91), posterior end of sclerotization broader with convex sides, anterior end tubular and narrower with straighter sides. Lamella antevaginalis (LAV) (Fig. 87): Posterior margin broadly convex; anterior margin convex on inner side, slightly concave on outer side, roughly perpendicular to the midline. Slit in LAV along ventral midline with an ovuloid expansion throughout its length; sides sclerotized and thickened anteriorly. Antrum (Fig. 87): Sclerotized throughout except a small area around juncture with corpus bursae. Sides fairly straight to weakly convex, widest and most convex at anterior end of slit in LAV. Ductus bursae (Fig. 87): Rectangular and strongly dorso-ventrally flattened with a sclerotized plate on each side, posterior approximately $2 / 3$ strongly curved ventrally, anterior approximately $1 / 3$ twisted about 90 degrees to left of ventral. Corpus bursae (Fig. 87): Posterior section with longitudinal wrinkles, elongate, about 4 times as long as wide at anterior base (ventral aspect), sides convex, widening anterior to posterior. Anterior section densely covered with minute inverted teeth, globular and nearly spherical. Ductus seminalis (Fig. 92): Total length approximately 12 mm if straightened out. Coiled basal section about 3.5 mm straightened out, with about five coils. Distal section about 8.5 mm , oblong bulla seminalis about 1.7 mm . Collaterial gland complex ( $($ Figs 87,89 \& 96) 86, 88, \& 90 ): Terminology follows Mitter (1987). Adjoining differentiated canals of receptacle duct with one coil basal to the vesicle; abrupt transition to undifferentiated section in basal third of vesicle; vesicle unsclerotized, expanded relative to preceding coil, singly curved (Fig. 90). Utriculus and lagena elongate (Fig. 86), utriculus with many narrow longitudinal grooves throughout. Collaterial gland elongate and tubular, gradually widening basal to distal and then slightly widening into a two pronged sack from which the paired glands arise (Fig. 88). Oviductus communalis as shown in Fig. 86, typical of Catocala nupta complex. Vagina globular, almost spherical (Fig. 86).

COI 5' Mitochondrial DNA: the C. benedeki holotype has the following sequence for COI 5' positions 1 658; a single position was polymorphic among six sequenced specimens: 88 (C\&T).

AACTTTATATTTTATTTTTGGAATTTGAGCAGGAATAGTAGGAACTTCATTAAGATTATTAATTCGAGCT GAATTAGGTAATCCTGGCTCTTTAATTGGAGATGATCAAATTTATAATACTATTGTTACAGCTCATGCTTT TATTATAATTTTTTTTATAGTTATACCAATTATAATCGGAGGATTTGGTAATTGATTAGTACCTTTAATATT AGGAGCTCCTGATATAGCTTTTCCTCGTATAAATAATATAAGTTTTTGACTTCTACCCCCCTCATTAACTT TATTAATTTCAAGAAGAATTGTAGAAAATGGAGCAGGAACTGGATGAACAGTTTATCCCCCTCTTTCTT CTAACATTGCTCATAGAGGTAGTTCAGTAGATTTAGCTATTTTTTCCCTACATTTAGCTGGAATTTCTTC AATTTTAGGAGCTATTAATTTTATTACTACAATTATCAATATACGATTAAATAATTTAATATTTGATCAAAT ACCTTTATTTATTTGAGCTGTTGGAATTACTGCATTTCTTCTTCTTCTTTCTCTACCAGTATTAGCTGGAG CTATTACTATACTTCTAACTGATCGAAATTTAAATACTTCCTTTTTTGATCCTGCTGGAGGAGGAGATCC TATTTTATATCAACATTTATTT.

Biology and distribution. The larval host of C. benedeki is unknown, but related species whose host plant usages are documented all feed on various Salicaceae. The new species is found in the Karakoram Mountains, where willow and poplar thickets occur along watercourses up to 3000 m (Bessarabov \& Allan 2017). Abbas et al. (2014) reported Salix alba L. and Salix denticulata Bayoow from the Haramosh Valley in Central Karakoram National Park, which is located approximately 25 km from where the holotype of C. benedeki was collected. We note that Hampson (1913:87) tabulates C. concubia specimens at the NHM from "Murree (Harford), 2才, 2 中" (now in Kashmir). We show one of these specimens (Fig. 5, collected 1887, Murree Hills, ex. Harford collection) which matches $C$. benedeki in wing pattern.

Etymology. Catocala benedeki is named for Balázs Benedek who collected most of the type series during four trips to Kashmir.

## Nomenclature of the Catocala nupta complex

The identity of $C$. nupta, pivotal to the genus, had been considered settled since the early 1800 s despite the fact that
typification occurred only recently. Mikkola and Honey (1993: 141) located two suitable syntypes in the Linnaean collection, and selected a female from Germany collected by Schreber as lectotype (Fig. 33; see http://linneanonline.org/insects.html). They also suggested the remaining unlabeled male paralectotype (collected by Brander) could be from North Africa i.e., "Barbaria" as stated in the original description. "Barbaria" is an unusual locality for C. nupta given that among red-hindwing Salicaceae feeders only Catocala elocata Esper, 1787 and Catocala oberthuri Austat, 1879 have otherwise been recorded from what is present day Morocco, Algeria, and Tunisia (see reviews by Oberthur 1918, Rothschild 1920, Rungs 1981; Arahou 2008). The number of original C. nupta types was not indicated, and Linnaeus' "Barbaria" might suggest a specimen or label attribution error, or even a mixedtaxon type series. Fortunately, Mikkola and Honey anchored C. nupta to prevailing usage and limited the type locality.

We are aware of 34 names that have been tabulated in the synonymy of C. nupta, only four of which were described originally as full species: Catocala concubina Borkhausen, 1792, Catocala unicuba Walker, [1858], C. concubia, and Catocala nozawae Matsumura, 1911. The location of Borkhausen's types and collection remain unknown (Stafleu \& Cowan 1976), but we consider that C. concubina has been appropriately treated as a synonym of C. nupta (both were illustrated by Hubner [1803]: pl. 69, figs 329, 330; see also Godart 1837, Herrich-Schaffer 1845, Guenée 1852, Staudinger \& Wocke 1861). Walker's holotype of C. unicuba (Fig. 14) falls within the normal range of variation of C. nupta, and C. unicuba has been appropriately treated as a synonym of C. nupta (see Hampson 1894, Poole 1989, Kononenko 2010). Walker's holotype of C. concubia (Fig. 13) represents a species separable from C. nupta, as discussed above. Matsumura's (1911: 89, Pl. XXXVII, Fig. 1) illustration and English description of C. nozawae are both consistent with C. nupta, although Matsumura noted the hindwing was not red but "yellowish white" and that "it resembles somewhat to Catocala lara Butl. [sic] but much smaller and the markings quite others." At the SEHU is a single male labeled by Matsumura as type of C. nozawae (Fig. 37). This specimen has whitish hindwings that appear to be albinic and/or faded from exposure to light, consistent with comparison having been made to the larger C. lara Bremer, 1861 (which does have a yellowish-white hindwing band). It also bears a lectotype label dated 1977 by Sugi. It is known that Sugi affixed lectotype labels to Matsumura lepidopteran types but did not publish designations (M. Owada, in litt. 2017), and we have been unable to locate a publication in which Sugi designated a lectotype for C. nozawae. The Japanese segment of the original description states "body length 1 sun [=30 mm], expanse 1 sun $4 \mathrm{bu}[=42 \mathrm{~mm}]$. This was collected by Mr. Shunjiro Nozawa in the Sapporo area in Meiji-23-year [=1890], seemed to be very rare." In addition, the entry for C. nupta nozawae in Sugi (1959: 142) states "the subspecific name nozawae, which was given to an abnormal individual with whitish underwings instead of red, is nomenclaturally available, and japonica Mell, named to a normal form, is a junior synonym of it." All the above evidence suggests that C. nozawae was described from only one specimen, and we therefore consider the SEHU male to be the holotype by monotypy. Catocala nozawae has recently been treated as a synonym (e.g., Poole 1989, Kononenko 2010) or subspecies (e.g., Goater et al. 2003) of C. nupta, and since the C. nozawae holotype is an otherwise typical specimen of C. nupta, we hereby reaffirm Catocala nozawae syn. rev. as a synonym of $C$. nupta.

The other 30 names in the synonymy of C. nupta were described in infraspecific contexts. Thirteen are explicitly infrasubspecific aberrations: "flava" Schultz, 1906, "mutilata" Schultz, 1906, "dilutior" Schultz, 1909, "fida" Schultz, 1909, "alterata" Warren, 1913, "brunnescens" Warren ,1913, "languescens" Warren, 1913, "rubridens" Warren, 1913, "xanthophaea" Schawerda, 1925, "salmonea" Cokayne, 1946, "quasiinterrupta" Schnaider, 1949, and "nigra" Cockayne, 1951 (the infrasubspecific "nigrata" Lempke, 1966 is an unnecessary replacement for the infrasubspecific "nigra" Cokayne, 1951, nec "nigra" Lempke, 1949). Another 8 form names (e.g., "f. n.") are clearly infrasubspecific based on analysis of the original descriptions: "guiartii" Lambillion ,1905, "confusa" Oberthur, 1912, "grisescens" Hanneman, 1917, "nigrescens" Hanneman, 1917, "victoria" Woskressensky, 1927, "nigra" Lempke, 1949, "sanguinea" Lempke, 1949, and "variegata" Lempke, 1949.

The two varietal names Catocala nupta var. obscurata Oberthur, 1880 and Catocala nupta v. nuptialis Staudinger, 1901 have been treated as available since geographical context appears in their original descriptions. The holotype of C. nupta var. obscurata, syn. nov. (Fig. 32) at the NHM is a melanized but otherwise typical specimen of C. nupta (melanics occasionally occur throughout the species' geographic range), and the lectotype of C. nupta var. nuptialis, syn. nov. (Fig. 23) designated by Ishizuka (2015: 100) at the MNHU is a typical specimen of C. nupta. The name nuptialis Staudinger 1901 is preoccupied by the Nearctic species Catocala nuptialis Walker, [1858], with the replacement name for nuptialis Staudinger, 1901 being centrasiatica Kusnezov, 1903 (the name
ottostaudingeri Ishizuka, 2015 is a subsequent and hence unnecessary replacement name for nuptialis Staudinger, 1901). The name centrasiatica Kusnezov syn. nov. is thus a synonym of C. nupta.

The remaining five names in the Catocala nupta synonymy were described as subspecies: Catocala nupta ssp. kansuensis Bang-Haas, 1927, Catocala nupta ssp. clara Osthelder, 1933, Catocala nupta ssp. japonica Mell, 1936, Catocala nupta ssp. likiangensis Mell, 1936, and Catocala nupta ssp. alticola Mell, 1942. The lectotype of C. nupta ssp. kansuensis syn. nov. (Fig. 26) designated by Ishizuka (2010: 105) at the MNHU is a typical specimen of C. nupta. The holotype of C. nupta ssp. japonica syn. nov. (Fig. 35) is also at the MNHU, and is a small but otherwise typical specimen of C. nupta. The original description for C. nupta ssp. clara states "M[arasch und Umgebung] VIII. [19]30 einige leidlich frische Stücke" [="Marasch and environment. August 1930 Some tolerably fresh specimens"], and to clarify application of the name, we hereby designate a male at the ZSM as LECTOTYPE for C. nupta ssp. clara (Fig. 29). The type locality is hereby restricted to the Taurus Mountains near Kahramanmaras (formerly Marash) in south central Turkey on the basis of the lectotype labels (="Syr. sept./Taurus/ Marasch"). The lectotype of C. nupta ssp. clara syn. nov. is a typical specimen of C. nupta. Examination of the types for C. nupta ssp. likiangensis and C. nupta ssp. alticola demonstrates these to be specimens of C. concubia, not C. nupta (Mell stated that his likiangensis and alticola were larger than C. nupta, which is consistent with the size difference between C. nupta and C. concubia). The original description for C. nupta ssp. likiangensis lists 27 specimens, and to clarify application of the name, we hereby designate a male at the ZFMK as LECTOTYPE for $C$. nupta ssp. likiangensis (Fig. 17). The type locality is hereby restricted to Lijiang (=Likiang), Yunnan Province, China, on the basis of the lectotype labels ("Li-kiang ca. $2000 \mathrm{~m} /$ Prov. Nord-Yuennan"). The name C. nupta ssp. likiangensis syn. nov. is a synonym of C. concubia. The original description for C. nupta ssp. alticola lists several localities but does not enumerate specimens. Mell stated that $C$. nupta ssp. alticola had slightly more white on the hindwing than C. nupta ssp. likangensis, but this falls within the normal range of variation in C. concubia. To clarify application of the name, we hereby designate a male at the ZFMK as LECOTYPE for C. nupta ssp. alticola (Fig. 15). The type locality is hereby restricted to the Jinsha River valley, Batang, Sichuan Province, China on the basis of the lectotype labels ("Batang. (Tibet). Im Tal/des Yangtze (ca. 2800 m )" [=in valley of Yangtse]; Jinsha River is the name of the upper stretches of the Yangtze River). The name C. nupta ssp. alticola syn. nov. is a synonym of C. concubia.

Both C. szechuena and C. amnonfreidbergi were described as species, and have not previously been placed in synonymy. The name C. szecheuna appears not to have been analyzed: Mell (1936: 71) devotes one terse sentence to it in his otherwise detailed study of the Asian Catocala fauna, Goater et al. (2003: 83) simply include it in their checklist of Palearctic species, and the name is absent in Kononenko (2010) and Ishizuka (2011). Hampson's description of C. szechuena states it has "a narrow black medial band," in contrast to C. concubia which has "an obliquely curved medial black band from costa to vein 1, expanding towards costa and its outer edge excurved at vein 5." We have found the width of the black medial band to be quite variable within C. nupta and related taxa. The holotype of C. szechuena (Fig. 19) is at the NHM. The medial band of the C. szechuena holotype represents the narrowest end of continuous variation exhibited by $C$. concubia, and we place the name $C$. szechuena syn. nov. as a synonym of $C$. concubia.

Catocala amnonfreidbergi was described as a distinct species on the basis of a single male specimen from Israel, and was diagnosed as separable by having the "finger of left valva much longer than in C. nupta; black band of hindwing less angled, more curved." The genitalia of the holotype (illustrated in Kravchenko et al. 2008) are within the normal range of genitalic variation that we have found in our dissections of C. nupta, and the hindwing band is similarly typical of C. nupta. Hence we place the name C. amnonfreidbergi syn. nov. as a synonym of $C$. nupta. Kravchenko et al. (2008) also observed that the "Japanese taxon (C. nozawae =C. japonica) is very unusual as the left valva is without a sclerotised costa and finger, whereas the right valva has a sclerotised costa with a short finger," and suggested this might represent a character of species-level distinction. Our male C. nupta dissections from Japan have a sclerotized left costal extension that is longer than the right, and this holds for our C. nupta dissections from other geographic locations. In this regard, we note that sometimes the left costal extension can be broken off (for example, if specimens were placed in envelopes when the apex of the male genitalic capsule was protruding) and such specimens can misleadingly appear to lack an extension of the left costa.

## Checklist of the Catocala nupta complex:

Catocala benedeki Borth, Kons, Saldaitis \& Gall, 2017
Catocala concubia Walker, [1858] stat. rev.
=szechuena Hampson, 1913 syn. n.
= nupta likiangensis Mell, 1936 syn. n.
=nupta alticola Mell, 1942 syn. n.
Catocala nupta (Linnaeus, 1767)
= concubina Borkhausen, 1792
= unicuba Walker, [1858]
=nupta obscurata Oberthür, 1880 syn. n.
=nupta nuptialis Staudinger, 1901 syn. n. (preocc. by nuptialis Walker, [1858])
=nupta centrasiatica Kusnezov, 1903 syn. n. (nom. nov. for nuptialis Staudinger, 1901)
=nozawae Matsumura, 1911 syn rev.
=nupta kansuensis Bang-Haas, 1927 syn. n.
=nupta clara Osthelder, 1933 syn. n.
=japonica Mell, 1936 syn. n.
= amnonfreidbergi Kramvchenko et al., 2008 syn. n.
=ottostaudingeri Ishizuka, 2015 (unnecessary nom. nov. for nuptialis Staudinger, 1901)

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| APPENDIX 1: Data for Catocala concubia and C. nupta sequence and dissection vouchers |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | Dissection |  | Statel | Collection |  |  | Elev |  |  | COI-5P |
| Species name | Sample ID | No. HLK: | Country | Province | Date | Lat | Lon | (m) | Collector | Collection | base pr. |
| C. concubia | 2121-Yale | 2121 (male) | China | Sichuan | 10-Jul-2013 | 31 | 98 | 3600 |  | NRCV | $658[0 \mathrm{n}]$ |
| C. concubia | 2122-Yale | 2122 (female | China | Sichuan | 10-Jul-2013 | 31 | 98 | 3600 |  | NRCV | $658[0 \mathrm{n}]$ |
| C. concubia | 8000-050802-CH |  | China | Yunnan | 5-Aug-2002 | 25.7 | 101.6 | 2000 | A. Floriani | RJB | $658[0 \mathrm{n}]$ |
| C. concubia | 20432-CO1-14 | 2047 (male) | China | Yunnan | 19-Jun-2009 | 28 | 99.7 | 3356 | B. Benedek | RJB | $658[0 \mathrm{n}]$ |
| C. concubia | 8067-280700-TI |  | China | Tibet | 28-Jul-2000 | 29.02 | 89.258 | 3800 | D. Bruna | NRCV | $307[0 \mathrm{n}]$ |
| C. nupta | Non DNA Voucher | 051 (female) | Russia |  | 14-Aug-2003 | 55.1 | 61.4 | 256 |  | RJB | N/A |
| C. nupta | 2332-120706-SL |  | Slovakia |  | 12-Jul-2006 | 47.97 | 18.106 | 109 | L. Misko | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | Non DNA Voucher | 1555 (female | Lithuania |  | 2003 |  |  |  | V. Pacevicius | RJB | N/A |
| C. nupta | 20394-150913-KY |  | Kyrgyzstan |  | 15-Sep-2013 | 42.1 | 77.6 | 1700 | S. Toropov | NRCV | $658[0 \mathrm{n}]$ |
| C. nupta | 20019-050812-KY |  | Kyrgyzstan |  | 05-Aug-2012 | 42.74 | 77.67 | 1632 | S. Toropov | NRCV | $658[0 \mathrm{n}]$ |
| C. nupta | 8001-062408-RU | 2097 (male) | Russia | Chelyabinsk | 24-Jun-2008 | 55.2 | 61.4 | 2100 |  | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 20439-280814-CH |  | China |  | 28-Aug-2014 | 31.8 | 103.73 | 1600 | Floriani \& Saldaitit | AFM | $658[0 \mathrm{n}]$ |
| C. nupta | 9564-150807-CH | 2062 (male) | China | Shaanxi | 15-Aug-2007 | 33.9 | 109.1 | 1600 | D. Nilsson | RJB | $8[0 \mathrm{n}]$ |
| C. nupta | 8003-150807-CH |  | China | Shaanxi | 15-Aug-2007 | 33.9 | 109.1 | 1600 | D. Nilsson | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 8002-150807-CH | 714 (male) | China | Shaanxi | 15-Aug-2007 | 33.9 | 109.1 | 1600 | D. Nilsson | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 10250-240911-CH |  | China | Sichuan | 24-Sep-2011 | 33.32 | 103.93 | 2100 | A. Floriani | AFM | $658[0 \mathrm{n}]$ |
| C. nupta | $8071-210807-\mathrm{CH}$ |  | China |  | 21-Aug-2007 | 25 | 102 | 3400 | S. Murzin | RJJ | $658[0 \mathrm{n}]$ |
| C. nupta | $8005-150704-\mathrm{RU}$ |  | Russia | Primorsky Krai | 15-Jul-2004 | 43.2 | 131.5 | 170 | Aniskovich | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 5625-250702-RU |  | Russia | Primorsky Krai | 25-Jul-2002 | 45 | 134 | 170 | Aniskovich | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 8004-160502-RU |  | Russia | Primorsky Krai | 16-May-2002 | 44.82 | 131.9 | 180 | K. Kolesnichenko | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 9218-190798-JA | 2096 (male) | Japan |  | 19-Jul-1998 | 36.2 | 137.97 | 594 | Takamura | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | 9219-190798-JA |  | Japan |  | 19-Jul-1998 | 36.2 | 137.97 | 594 | Takamura | RJB | $658[0 \mathrm{n}]$ |
| C. nupta | TLMF Lep 09855 |  | Austria | Vorarlberg | 26-Sep-10 | 47.27 | 9.64 | 468 |  |  | $658[0 \mathrm{n}]$ |
| C. nupta | Mm04216 |  | Finland | Aland Islands | 07-Oct-2006 | 59.96 | 20.012 | 0 | M. Mutanen |  | $658[0 \mathrm{n}]$ |
| C. nupta | MM04750 |  | Finland | Finland Proper |  | 60.44 | 22.202 | 15 | M. Mutanen |  | $658[0 \mathrm{n}]$ |
| C. nupta | MM15851 |  | Finland |  |  | 60.41 | 26.819 | 0 | M. Mutanen et. al. |  | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 29037 |  | Germany | Bavaria | 14-Aug-2001 | 47.9 | 12.851 | 440 | A. Haslberger | ZSM | $613[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 78585 |  | Germany | Bavaria | 15-Sep-2013 | 48.26 | 11.545 | 481 | A. Hausmann | ZSIM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep R 21673 |  | Germany | Bavaria | 13-Aug-1983 |  |  | 450 | F. Stamer | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 80284 |  | Germany | Schleswig-Holste | 05-Sep-1995 | 54.3 | 10.433 | 34 | Dr. D. Kolligs | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 64832 |  | Germany | Sachsen |  | 51.16 | 13.2 | 250 | M. Franzen | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 72632 |  | Germany | Saarland | 07-Sep-2012 | 49.19 | 7.0581 | 321 | H. Martin | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 64831 |  | Germany | Sachsen |  | 51.16 | 13.2 | 250 | M. Franzen | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 75217 |  | Germany | Sachsen | 18-Sep-2012 | 50.83 | 12.983 | 371 | S. Erlacher et al. | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 70164 |  | Germany | Bavaria | 17-Jul-2012 | 48.15 | 11.334 | 480 | A. Hausmann | ZSM | $658[0 \mathrm{n}]$ |
| C. nupta | BC ZSM Lep 21863 |  | Germany | Bavaria | 05-Sep-1996 | 48.26 | 11.545 | 480 | A. Hausmann | ZSM | $658[0 \mathrm{n}]$ |

