Structural genomic changes underlie alternative reproductive strategies in the ruff (*Philomachus pugnax*)

Sangeet Lamichhaney^{1,11}, Guangyi Fan^{2,3,11}, Fredrik Widemo^{4,11}, Ulrika Gunnarsson¹, Doreen Schwochow Thalmann^{5,6}, Marc P Hoeppner^{1,7,10}, Susanne Kerje¹, Ulla Gustafson⁵, Chengcheng Shi², He Zhang², Wenbin Chen², Xinming Liang², Leihuan Huang², Jiahao Wang², Enjing Liang², Qiong Wu², Simon Ming-Yuen Lee³, Xun Xu², Jacob Höglund⁸, Xin Liu² & Leif Andersson^{1,5,9}

The ruff is a Palearctic wader with a spectacular lekking behavior where highly ornamented males compete for females¹⁻⁴. This bird has one of the most remarkable mating systems in the animal kingdom, comprising three different male morphs (independents, satellites and faeders) that differ in behavior, plumage color and body size. Remarkably, the satellite and faeder morphs are controlled by dominant alleles^{5,6}. Here we have used whole-genome sequencing and resolved the enigma of how such complex phenotypic differences can have a simple genetic basis. The Satellite and Faeder alleles are both associated with a 4.5-Mb inversion that occurred about 3.8 million years ago. We propose an evolutionary scenario where the Satellite chromosome arose by a rare recombination event about 500,000 years ago. The ruff mating system is the result of an evolutionary process in which multiple genetic changes contributing to phenotypic differences between morphs have accumulated within the inverted region.

Independents constitute 80–95% of male ruffs and strive to defend territories on leks^{1–3,7}. Independent males show a spectacular diversity in the color of their ruff and head tufts (**Fig. 1a**). Satellites are slightly smaller than independents, usually show white ruff and white tufts (**Fig. 1a**) and constitute 5–20% of males^{7,8}. Satellites are non-territorial and display submissive behavior, allowing independent males to dominate them at leks (**Fig. 1a**, middle). Independents clearly recognize satellites as a different kind of male and behave differently with satellites than they do with other independents (see URLs for a link to a video showing the reproductive strategies of the three male morphs). Both independents and satellites may benefit from their interaction by attracting females⁷. The faeder is a rare third morph (<1% of male ruffs) mimicking females by its

smaller size and female-like plumage^{1,9,10} (**Fig. 1a**). These disguised males appear on the leks where they attempt to gain access to females that are ready to mate.

A high-quality genome assembly was established using genomic DNA from an independent male kept at the Helsinki zoo in Finland. We estimated the genome size to be 1.23 Gb and generated 139.3 Gb of Illumina HiSeq 2000 sequencing data using fragment libraries with insert sizes ranging from 250 bp to 20 kb (**Supplementary Figs. 1–3** and **Supplementary Table 1**). The N50 scaffold size was as high as 10.0 Mb (**Supplementary Table 2**).

We generated $\sim 8 \times$ genome coverage on the basis of 2×125 -bp paired-end reads from 15 independent and nine satellite males, all from a single location. A screen based on the fixation index (F_{ST}) comparing independents and satellites identified a single highly differentiated 4.5-Mb region on scaffold 28 (Fig. 1b). Independents and satellites clustered as two genetically distinct groups in a phylogenetic tree based on this region (Fig. 1c, left). In contrast, there was no significant differentiation between these groups when the tree was constructed on the basis of the rest of the genome (Fig. 1c, right). We hypothesized that the large region of strong differentiation might reflect the presence of an inversion. We used BreakDancer¹¹ to screen for structural changes and identified a 4.5-Mb inversion present in satellites and overlapping perfectly the differentiated region (Fig. 1d). PCR-based sequencing confirmed a proximal breakpoint at 5.8 Mb and a distal breakpoint at 10.3 Mb and identified a 2,108-bp insertion of a repetitive sequence at the distal breakpoint. A diagnostic test (Fig. 1e) showed that all satellites were heterozygous for the inversion and all 112 independents except five were homozygous for the wild-type sequence (Table 1); the latter five were heterozygous for the inversion and most likely reflect phenotype misclassifications in the field. The inversion was also found among adult females and young birds (Table 1).

¹Science for Life Laboratory, Department of Medical Biochemistry and Microbiology, Uppsala University, Uppsala, Sweden. ²BGI-Shenzhen, Shenzhen, China.
³State Key Laboratory of Quality Research in Chinese Medicine, Institute of Chinese Medical Sciences, University of Macau, Macao, China. ⁴Department of Wildlife, Fish and Environmental Studies, Swedish University of Agricultural Sciences, Umeå, Sweden. ⁵Department of Animal Breeding and Genetics, Swedish University of Agricultural Sciences, Umeå, Sweden. ⁵Department of Animal Breeding and Genetics, Swedish University of Agricultural Sciences, Umeå, Sweden. ⁵Department of Animal Breeding and Genetics, Swedish University of Agricultural Sciences, Institute National de la Recherche Agronomique, AgroParisTech, Génetique Animale et Biologie Intégrative, Jouy-en-Josas, France. ⁷Bioinformatics Infrastructure for Life Sciences, Uppsala University, Uppsala, Sweden. ⁸Department of Ecology and Genetics, Uppsala University, Uppsala, Sweden. ⁹Department of Veterinary Integrative Biosciences, Texas A&M University, College Station, Texas, USA. ¹⁰Present address: Institute of Clinical Molecular Biology, Christian Albrechts University of Kiel, Kiel, Germany. ¹¹These authors contributed equally to this work. Correspondence should be addressed to L.A. (leif.andersson@imbim.uu.se) or X. Liu (liuxin@genomics.cn).

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The *Independent* allele clearly represents the ancestral state, as the inversion disrupts conserved synteny among birds (**Fig. 1f**).

The single faeder in our material was also heterozygous for the 4.5-Mb inversion (Table 1). We sequenced this individual to 30× coverage, and F_{ST} analysis indicated striking genetic differentiation between the faeder and both independents and satellites within the inverted region (Fig. 2a). The differentiation between the faeder and independents was equally strong across the 4.5-Mb region, whereas the pattern of differentiation between the faeder and satellites was a mirror image of the pattern between satellites and independents (Fig. 2a). We phased haplotypes using Beagle¹² and constructed haplotype trees separately for region A showing high F_{ST} between satellites and independents and for region B showing low FST between satellites and independents (Fig. 2a,c). In region A, the Satellite and Faeder chromosomes were closely related and divergent from Independent chromosomes. In contrast, in region B, the Satellite and Independent chromosomes were more closely related, whereas the Faeder chromosome was divergent (Fig. 2c, bottom). Because we only had access to a single faeder, we genotyped the entire material (>200 birds) using two SNPs diagnostic for the Faeder chromosome. The Faeder chromosome should not be present in independent or satellite males but should occur at a low frequency in adult females and young birds. The results confirmed this prediction, as we identified the Faeder chromosome in only a single adult female (Supplementary Table 3). This female was clearly an outlier with regard to body size (Supplementary Fig. 4), consistent with the observation that females heterozygous for the Faeder allele are smaller than other females⁶.

We examined the genetic consequences of the inversion and searched for candidate mutations that might contribute to phenotypic differences. This is challenging because the inverted region contains about 90 genes (**Supplementary Fig. 5** and **Supplementary Table 4**). First, we note that the inversion disrupts the *CENPN* gene (encoding centromere protein N) (**Fig. 1d**). The inversion may be recessive

lethal, as data from human cells¹³ and zebrafish¹⁴ with mutations in the orthologous gene show that *CENPN* inactivation has severe deleterious effects. In fact, ruff pedigree data have confirmed that the inversion is recessive lethal¹⁵. Birds heterozygous for the *Satellite* allele must have about 5% higher fitness to maintain an allele frequency of about 5% in compensating for the lethality of the homozygote. Second, we identified a large number of missense mutations present on *Satellite* and/or *Faeder* chromosomes (**Supplementary Table 5**). Third, BreakDancer¹¹ and depth-of-coverage analysis identified three deletions ranging in size from 3.3 to 17.6 kb (**Fig. 2d**) present in the heterozygous condition in all satellites and in the faeder but not in independent males (**Supplementary Table 3**). Two of these (5.2 kb and 17.6 kb in length) delete evolutionarily conserved sequences (**Fig. 2e** and **Supplementary Fig. 6**), and all three cluster in the vicinity of *HSD17B2* (hydroxysteroid (17-β) dehydrogenase 2) and

 Table 1 Results of a diagnostic test for the presence of a 4.5-Mb inversion associated with alternative mating strategies in ruff

	Inversion ^a		
Phenotype class	_/_	+/-	Total
Males			
Independent	107	5	112
Satellite	0	24	24
Faeder	0	1	1
Young	18	4	22
Total for males	125	34	159
Females			
Adults	52	5	57
Young	28	3	31
Total for females	80	8	88
Total across sexes	205	42	247

^aTest based on the inversion breakpoint at 5.8 Mb in scaffold 28: -, no inversion; +, inversion.

LETTERS



present in satellites and faeders within the 4.5-Mb inversion are all located in the vicinity of *HSD17B2* and *SDR42E1*. Two duplications (magenta) are only present in faeder. (e) The 5.2-kb deletion shows 50% reduced sequence coverage in satellites (top); the deletion encompasses evolutionarily conserved sequences among birds¹⁸ (bottom). (f) Proposed evolutionary history of the three alleles underlying the mating system in ruff.

SDR42E1 (short-chain dehydrogenase/reductase family 42E, member 1) (Fig. 2d). HSD17B2 and SDR42E1 both have important roles in the metabolism of sex hormones: HSD17B2 catalyzes conversion of the 17 β -hydroxy forms of estrogen and androgens (including testosterone and dihydrotestosterone) into their less active 17-keto forms¹⁶. We postulate that one or more of these deletions constitute *cis*-acting regulatory mutations that alter the expression pattern of HSD17B2 and/or SDR42E1 and contribute to phenotypic differences among male morphs. We identified two deletions and two duplications that were unique to the faeder and may thus contribute to the faeder phenotype (**Supplementary Fig. 7**).

There is a striking diversity in plumage color among male ruffs (**Fig. 1a**). One of the most obvious candidate genes for variation in pigmentation, *MC1R* (encoding melanocortin 1 receptor), is located within the inverted region at position 10.2 Mb (**Supplementary Fig. 5**). Whole-genome sequencing showed that satellites are heterozygous for four derived *MC1R* missense mutations (encoding p.Val105Leu, p.Arg149His, p.His207Arg and p.Arg303Trp) at residues that are conserved among birds and mammals (**Fig. 3**). We performed Sanger sequencing of the single *MC1R* exon in all satellites, the faeder

and a subset of the independents. This analysis confirmed complete association between the Satellite allele and these four missense mutations, whereas the faeder was heterozygous for the variant encoding p.His207Arg and three other missense mutations (Supplementary Table 6). The p.His207Arg substitution most likely has functional consequences because the same variant is associated with light color in the red-footed booby¹⁷. We propose that the MC1R allele on Satellite chromosomes, possibly together with altered metabolism of sex hormones, underlies the white color of ornamental feathers in satellites. To be causal, this allele must have a dominant effect, as satellites are always heterozygous. This implies a dominant-negative effect or, more likely, a combination of regulatory and coding changes leading to overexpression of a variant form of MC1R specifically in ornamental feathers. The latter mechanism would explain why satellite males, despite their spectacular light color during the breeding season (Fig. 1a), are almost indistinguishable from independents outside the breeding season and why females carrying the Satellite allele are not markedly lighter in color than other females.

We estimated the time since divergence of the *Satellite* and *Independent* alleles on the basis of region A, showing high F_{ST} between

Figure 3 Four amino acid substitutions encoded by *MC1R* missense mutations associated with the *Satellite* allele. Missense variants detected in satellites are represented by red circles, and a selected set of variants associated with plumage variation in other birds²⁹ is represented by yellow circles. Lower left, the amino acids associated with the *Independent* allele are conserved among birds and mammals.

the two types of males, to 3.87 ± 0.15 million years ago (Fig. 2a,b), using the nucleotide divergence (1.4%) and estimated mutation rates for birds¹⁸. A similar estimate of 4.09 \pm 0.16 million years ago was obtained for divergence of the Faeder and Independent alleles on the basis of sequence divergence for the entire 4.5-Mb region. There can be no recombination between the Satellite and Faeder chromosomes, as the Faeder/Satellite genotype is not viable. Furthermore, an inversion is expected to cause suppression of recombination within the inverted region between the wild-type and mutant alleles. Comparison of the faeder and independents across the inversion is consistent with this lack of recombination, as F_{ST} values are

equally strong across the 4.5-Mb region (Fig. 2a). Remarkably, the strong differentiation between satellites and independents is disrupted in two regions (1.5 and 0.7 Mb in size) that show $F_{\rm ST}$ values that are much lower than those for other segments of the inversion but still markedly higher than the background level (Fig. 2a). We postulate that the Satellite chromosome arose by one or two rare recombination events between an Independent and a Faeder-like chromosome (Fig. 2f). This would explain why the pattern of genetic differentiation between the Faeder and Satellite chromosomes is a mirror image of the differences between the Independent and Satellite chromosomes (Fig. 2a). A similar rare recombination event between an inversion and a wild-type chromosome created a third allele at the Rose-comb locus in chicken¹⁹. Our model predicts that the pairwise genetic distances between the Satellite and Independent chromosomes for region B should all be equal, as they reflect divergence since the recombination event happened. Our data are consistent with this hypothesis (Supplementary Table 7). Similarly, the pairwise genetic distances between the Satellite and Faeder chromosomes in region A are also equal (Supplementary Table 7). We estimated that this recombination event occurred 520,000 \pm 20,000 years ago, on the basis of sequence divergence (0.2%) between the Satellite and Independent chromosomes in region B (Fig. 2b). We constructed 5-kb and 10-kb mate-pair libraries from one satellite male, but analysis of these data did not identify any additional inversions in the vicinity of the 4.5-Mb inversion.

The genetic basis for the satellite and faeder morphs constitutes a combination of genetic changes that have accumulated within the inverted region over a period of about 3.8 million years. This resembles the situation in white-throated sparrows in which at least two pericentric inversions involving about ~100 Mb are associated with altered plumage (brown-and-tan stripes versus white-and-black stripes on the crown) and altered territorial and parental behavior^{20,21}. A recent study suggested that sequence differences in the promoter of *ESR1* (encoding estrogen receptor α) are causally related



to behavioral differences between morphs²². When an inversion is associated with a complex phenotype, it is challenging to pinpoint causal mutations because many sequence polymorphisms within the inverted region show an equally strong association to the phenotype. An additional complication in ruff is that recessive variants located within the inverted region will never be exposed to purifying selection because homozygosity for the inversion is lethal. Only sequence variants on the Satellite and Faeder chromosomes that show some degree of dominance can contribute to phenotypic differences among the morphs. We propose that the inversion itself caused the first phenotypic effects that constituted the starting point for an evolutionary process eventually resulting in the current mating system in ruff. An inversion may cause phenotypic effects as a result of changes affecting the coding sequence or the regulation of genes, primarily in the vicinity of the breakpoints, as illustrated by the Rosecomb inversion in chicken where translocation of MNR2 leads to ectopic expression of the MNR2 transcription factor and altered comb development¹⁹. Similarly, translocation of CYB5B, located at the breakpoint at 10.3 Mb in ruff (Fig. 1d), may cause altered expression of this gene that has a role in the biosynthesis of glucocorticoids and sex steroids²³. Furthermore, it appears highly plausible that one or more of the three deletions in the near vicinity of HSD17B2 and SDR42E1 lead to altered metabolism of testosterone and other steroids, which may affect both behavior and plumage. In fact, independents have higher circulating levels of testosterone, whereas satellites and faeders have higher concentrations of androstenedione¹⁵. A possible explanation for this difference is that upregulation and/or ectopic expression of HSD17B2 lead to conversion of testosterone into androstenedione in individuals heterozygous for the Satellite or Faeder allele, a model consistent with the dominant inheritance of these alleles. This is a testable hypothesis because it predicts allelic imbalance in HSD17B2 expression in satellites and faeders.

Our study has demonstrated how an inversion followed by subsequent accumulation of several adaptive changes within the inverted region led to the evolution of a spectacular mating system comprising three alleles at a single locus maintained by balancing selection. The presence of an inversion that allowed the evolution of a non-recombining 'supergene' (ref. 24) was critical for this process. Other examples where inversions are associated with complex phenotypes include mimicry in butterflies^{25–27} and colony organization in fire ants²⁸.

URLs. NCBI Sequence Read Archive (SRA), http://www.ncbi.nlm. nih.gov/sra/; RepeatMasker, http://repeatmasker.org/; Picard, http:// broadinstitute.github.io/picard/; video of ruff reproductive strategies, http://www.scilifelab.se/research/scientific-highlights/ruff.

METHODS

Methods and any associated references are available in the online version of the paper.

Accession codes. The Illumina reads have been submitted to the Sequence Read Archive (SRA) under accession SRA266458. The assembly and annotation are available under accession PRJNA281024. DNA sequences reported in this manuscript have been submitted to GenBank under accessions KT202232–KT202235 and KT428875.

Note: Any Supplementary Information and Source Data files are available in the online version of the paper.

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AUTHOR CONTRIBUTIONS

L.A., J.H. and F.W. conceived the study. F.W. and J.H. collected all material and phenotype data. G.F., M.P.H., C.S., H.Z., W.C., X. Lia, L.H., J.W., E.L., Q.W., S.M.-Y.L., X.X. and X. Liu contributed to genome assembly and annotation. S.L., G.F. and L.A. were responsible for bioinformatics analysis of genome resequencing data. U. Gunnarsson, D.S.T., S.K. and U. Gustafson performed experimental work. L.A. and S.L. wrote the manuscript with input from other authors. All authors approved the manuscript before submission.

COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

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ONLINE METHODS

Sample collection and DNA extraction. Blood was collected form the wing vein of a captive male ruff (further referred to as the reference individual) kept at the Helsinki zoo, Finland, and mixed with EDTA as anticoagulant. DNA was immediately isolated using a standard salt precipitation method. The quantity and quality of the sample were evaluated using the Qubit dsDNA BR assay (Life Technologies) as well as by pulsed-field gel electrophoresis (CHEF Mapper XA, Bio-Rad).

The samples for whole-genome resequencing and further genetic analysis were collected from a ruff population that was studied during the breeding seasons of 1990–2002 on the island of Gotland in the Baltic Sea (57° 10' N, 18° 20' E)^{30–32}. In each year, males and females were caught on leks using cannon nets. In addition, females and newly hatched young were caught on nests. Individuals were ringed, and morphological measurements including tarsus length (in mm) and wing length (in mm; maximum chord) were collected (all by F.W.). A blood sample was drawn from the wing or brachial vein of each bird and later used for DNA extraction. The birds were released after handling. Birds were caught and handled according to ethical permits and permissions guiding Swedish research and animal welfare (Stockholms Södra Djurförsöksetiska Nämnd S54-99).

Male strategies were determined from plumage pattern and from behavioral observations at leks for ringed independents and satellites⁷. Most males (>90%) in nuptial plumage were scored as independents or satellites from their plumage alone. Whether female-colored birds were faeder males could only be preliminarily determined from morphological measurements and later verified through sexing using a DNA test³³.

Genome assembly. We generated 139.32 Gb of high-quality next-generation sequencing data with fragment lengths ranging from 250 bp to 20 kb for the reference individual using Illumina HiSeq 2000 sequencing (**Supplementary Table 1**). We estimated the ruff genome size using *k*-mer analysis to be about 1.23 Gb (**Supplementary Fig. 1**), suggesting genome sequence coverage of 113.7-fold. Using SOAPdenovo (version v2.04)³⁴, we obtained an assembly spanning 1.25 Gb, with contig N50 and scaffold N50 sizes of 106.46 kb and 10.00 Mb, respectively (**Supplementary Table 2**); 96.3% of the assembly was non-gap sequences. The distribution of sequencing depth, calculated on the basis of reads from all sequencing libraries, and the distribution of GC content are presented in **Supplementary Figures 2** and **3**, respectively.

Annotation of the region associated with an inversion. A preliminary annotation for the inverted region was generated with the Maker package (version 2.31-8)³⁵. We first composed a set of high-confidence reference sequences from UniProt by selecting bird proteins that were classified as full length and supported by either proteomics or transcriptomics (74,138 sequences). As additional input, we collected available 454 sequencing data from the SRA available under accession SRA049313. Reads were assembled with the Trinity package (release 2014-07-07)³⁶ into 16,746 transcripts. Finally, to improve the accuracy of the annotation process, we modeled new repeat sequences from a preliminary genome assembly and used this library in combination with a curated repeat library for vertebrates included in the RepeatMasker package. From these data, we generated two complementary gene builds: one based directly on the aligned sequences to most accurately reflect the evidence data ('evidence build') and a second set of gene models seeded from ab initio (de novo) gene predictions generated by the chicken reference profile model, included with the Augustus package (version 2.7)³⁷. Both builds were compared and reconciled for the target region using the WebApollo curation platform³⁸. Functional annotation of candidate transcript models was performed through similarity searches³⁹ against the UniProt/ SWISS-PROT reference protein set (downloaded May 2014) in combination with the prediction of functional motifs and domains via the InterProScan package (release 5.7-48.0)⁴⁰.

Whole-genome resequencing and SNP calling. Sequencing libraries (average fragment size of about 500 bp) were constructed for 15 independents, nine satellites and one faeder, and 2×125 -bp paired-end reads were generated using Illumina HiSeq 2000 sequencers. For the independents and satellites, we generated ~8-fold coverage based on high-quality reads, after strict filtering

of low-quality and adaptor-contaminated reads, and ~30-fold coverage was generated for the single faeder male. The genomic reads were mapped against the ruff genome assembly using Burrows-Wheeler Aligner (BWA)⁴¹ (version 0.6.2) with default parameters. PCR duplicates were filtered from the alignments using Picard. Further, we performed base quality recalibration and indel realignment using the Genome Analysis Toolkit (GATK)⁴² and performed SNP discovery and genotyping across the 25 samples according to GATK bestpractices recommendations^{43,44}. Low-quality SNP calls were filtered out by an in-house filtering pipeline that excluded a SNP if it did not satisfy the threshold of a combination of various quality parameters (for example, SNP quality, base quality, mapping quality, haplotype score, Fisher strand bias, minimum read depth and maximum read depth). Thresholds were chosen on the basis of the distribution of each of these parameters from the raw variant calls.

Genome-wide screen for genetic differentiation among male morphs. We divided the genome into non-overlapping 15-kb windows and estimated the genetic divergence (F_{ST}) between independents and satellites using VCFtools (v.0.1.11)⁴⁵. The 4.5-Mb region in scaffold 28 that showed strong genetic differentiation between independents and satellites was further subdivided into smaller windows (5 kb in length) to refine the pattern of differentiation. A similar analysis was carried out when comparing data from the single faeder male and the other morphs.

Phylogenetic analysis. We used PLINK⁴⁶ to calculate pairwise genetic distances between individuals. These distances were used to generate neighbor-joining trees with PHYLIP⁴⁷. Phased haplotypes for the 4.5-Mb inversion region were generated using Beagle¹², and these haplotypes were used to estimate Jukes-Cantor corrected nucleotide distances among the *Independent*, *Satellite* and *Faeder* alleles with PHYLIP⁴⁷. The net frequency of nucleotide substitutions (d_A) was calculated according to the method of Nei⁴⁸, and the time since divergence (t) of alleles was calculated as $t = d_A/2\lambda$, where λ is the genomic substitution rate. As a substitution rate in ruff is not yet available, we used the substitution rates estimated for each of 48 other bird genomes¹⁸ to calculate a confidence interval for t, and the data are presented as a box plot.

Structural variants. We used the paired-end sequence data for detection of structural variants with BreakDancer¹¹. Information on read pairs that mapped with unexpected separation distances or orientation was used to predict inversions, insertions, duplications and deletions. Two-sided Fisher's exact test was used to identify structural variants showing significant frequency differences between morphs. Sequence alignments around the detected structural variants were also manually inspected using the Integrative Genomics Viewer (IGV)⁴⁹, to exclude false positives. Normalized read coverage was compared between morphs to check the consistency of the deletions predicted by BreakDancer¹¹. We also generated 5-kb and 10-kb mate-pair data from a single satellite individual to confirm the structural variants detected using paired-end data.

Functional annotation of genetic variants. We used SNPeff $(v.3.4)^{50}$ to annotate the genetic variants and categorized the variants into coding (synonymous and nonsynonymous), upstream/downstream and intronic/ intergenic classes.

Diagnostic PCR tests, Sanger sequencing and SNP genotyping. PCR was used to amplify the regions around the identified inversion and deletion breakpoints and the *MC1R* coding sequence; all primer sequences are given in **Supplementary Table 8**. Amplified fragments were either analyzed by agarose gel electrophoresis or subjected to Sanger sequencing using standard methods. DNA sequences were analyzed using CodonCode Aligner 5.1.4 software. Four TaqMan SNP genotyping assays (Life Technologies) diagnostic for the *Satellite* or *Faeder* haplotype were designed (**Supplementary Table 8**). Standard TaqMan allele discrimination assays were performed using an Applied Biosystems 7900 HT real-time PCR instrument.

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