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Two Lineages of KLRA with Contrasting Transcription Patterns Have Been Conserved at a Single Locus during Ruminant Speciation

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Cattle possess the most diverse repertoire of NK cell receptor genes among all mammals studied to date. Killer cell receptor genes encoded within the NK complex and killer cell Ig-like receptor genes encoded within the leukocyte receptor complex have both been expanded and diversified. Our previous studies identified two divergent and polymorphic KLRA alleles within the NK complex in the Holstein–Friesian breed of dairy cattle. By examining a much larger cohort and other ruminant species, we demonstrate the emergence and fixation of two KLRA allele lineages (KLRA*01 and -*02) at a single locus during ruminant speciation. Subsequent recombination events between these allele lineages have increased the frequency of KLRA*02 extracellular domains. KLRA*01 and KLRA*02 transcription levels contrasted in response to cytokine stimulation, whereas homozygous animals consistently transcribed higher levels of KLRA, regardless of the allele lineage. KLRA*02 mRNA levels were also generally higher than KLRA*01. Collectively, these data point toward alternative functional roles governed by KLRA genotype and allele lineage. On a background of high genetic diversity of NK cell receptor genes, this KLRA allele fixation points to fundamental and potentially differential function roles. *The Journal of Immunology, 2020, 204: 000–000.

Natural killer cells are a population of large granular lymphocytes with essential functions in immunity, cancer, and reproduction (1). NK cells are early responders to infection, particularly against viruses, and are capable of recognizing and killing infected and transformed host cells as well as initiating subsequent immune responses through the secretion of cytokines. This cytokine release helps to drive adaptive immune responses led by B and T cells, which can be augmented by direct interactions between NK cells and other APCs or restricted by direct interactions with T cells (2–4). NK cells also play a fundamental role in human reproduction during the formation of the placenta through cooperation with the extravillous trophoblast (1).

In humans and rodents, NK cells are a highly heterogeneous population with different functions, specificities, and activation thresholds (5, 6). This functional diversity is largely driven by the variegated expression of polygenic and polymorphic germline encoded receptors, a majority of which recognize polymorphic determinants of MHC class I. These receptors can be either activating or inhibitory, and the balance of signals received when these receptors are cross-linked with their ligand determines the functional potential and activation status of an individual NK cell. This NK cell receptor and MHC class I diversity is now known to influence differential outcomes to several viral infections and is associated with a number of autoimmune diseases (7–11).

The diversification of NK cell receptor gene families has occurred independently several times during mammalian evolution. There are two classes of NK cell receptors in mammals that are known to have diversified: the killer cell Ig-like receptor (KIR) family located within the leukocyte receptor complex (LRC), and the structurally unrelated killer cell lectin-like receptors (KLR) located within the NK complex (NKC). A diverse KIR gene repertoire is a feature of prosimian primates, including humans, apes, and Old and New World monkeys. In contrast, lemurus have expanded the KLR/C/D gene families (12), whereas rodents and horses have expanded the KLR family (13, 14). Cattle were thought to be unique in having expanded LRC and NKC gene families, the KIR (15), and the KLR/C/D (16), respectively. However, recent studies have confirmed other ruminant species have also expanded both NK cell receptor gene complexes (17, 18).

The ligands for a majority of the primate KIR and rodent KLRA are the highly diverse classical MHC class I (19). These receptor–ligand systems are considered equivalent to each other in that they generate NK cell functional diversity in a rare example of convergent evolution. Although the ligands for these receptors in...
other species have yet to be confirmed; considering the genomic synteny, homologous gene expansion mechanisms, and gene sequence conservation, it seems likely that these other diversification events are also examples of convergent evolution being driven by selection to produce NK cell functional heterogeneity. We have previously shown that the KIR and KLRC/D are differentially transcribed between cattle NK cells and that this mRNA expression is influenced by the MHC genotype (20). This offers strong evidence that this highly diverse receptor gene repertoire in cattle also creates functional diversity.

Against this background of NKC and LRC gene expansion, cattle KLRA appears to have remained functional and mono-
genomic, but there are at least two distinct gene lineages (21, 22). Two novel KLRA transcripts were identified that significantly differed between the extracellular lectin domains from the genome reference sequence, creating the potential for differential ligand binding and functional diversity (22). The KLRA receptors are homodimeric C-type lectin type II transmembrane glycoproteins. In mice, this includes at least 23 members (Ly-49A-W), with different mouse strains having a specific complement of genes (23). Not all these genes are functional; of those that are, most are inhibitory, several are polymorphic, and they generally demonstrate broad recognition of MHC class I or recognize defined class I subsets, as is the case with KIR in primates (24–28).

To further understand if cattle KLRA diversity was indicative of limited gene expansion or a highly polymorphic gene, we confirmed the structure and polymorphism of the KLRA locus in cattle and closely related species and examined KLRA lineage frequency within two cohorts of Holstein–Friesian bulls used as pedigree sires for dairy farmers. To assess the potential functional impact of KLRA diversity, we analyzed the relative NK cell mRNA expression of both allele lineages within and between individuals with different KLRA genotypes under cytokine stimuli. Cattle are an essential food-producing species subject to intense selective breeding in a global effort to increase food production while protecting the climate. Understanding the immunogenetic diversity of livestock species is a fundamental part of this effort by underpinning breeding strategies that reduce disease burden and increasing vaccine efficacy.

Materials and Methods

Ethics statement

Peripheral blood samples from Holstein–Friesian cattle (Bos taurus) were collected in accordance with the U.K. Animal (Scientific Procedures) Act, 1986, and approved by either The Pirbright Institute’s Animal Welfare and Ethics Committee, or The Roslin Institute’s Animal Welfare and Ethics Committee. The Chillingham cattle samples were taken from animals culled for welfare reasons. Blood sampling of Kuchinoshima-Ushi cattle was carried out in accordance with the Regulations for Animal Experiments in Nagoya University and the Guidelines for the Care and Use of Laboratory Animals by the Tokyo University of Agriculture. For all the other species, blood samples taken during necessary veterinary interventions were kindly supplied by The Zoological Society of London.

Transfection, flow cytometry, and quantification of KLRA surface expression

The full coding sequence of KLRA*01 and KLRA*02 were amplified using cDNA from animals previously identified as being homozygous using primers sense Ly-49HindIII (5′-CTGCTGAATTACAGTGTGTAATCAAGAAGTG-3′) and antisense XhoI (5′-GGATATCAGGAGTTTTATTGAAGCAATC-3′). PCR products were cloned into TOP10 Escherichia coli using the TOPO cloning system (Thermo Fisher Scientific, Renfrew, U.K.) and Sanger sequenced to identify clones with no errors. Inserts were digested from identified clones using HindIII and XhoI and ligated into the pcDNA6/V5-His vector (Thermo Fisher Scientific) and electroporated into P815 cells using standard protocols. Expanding clones were grown under blastocidin selection and picked for further expansion after 3–5 d. Cultures derived from the clones were screened by flow cytometry using an Ab against the V5 tag (V8012; Merck, Dorset, U.K.). Expression was confirmed in positive clones by Western blot using the V5 Ab to detect a protein of the correct size.

KLRA PCR amplification from genomic DNA

The genomic sequences of KLRA from a range of ruminant species (cattle, African buffalo, Bactrian camel, European bison, gembok, giraffe, greater kudu, lowland anoa, moose, red deer, water buffalo, and yak) were aligned. Primers (Sigma-Aldrich, Gillingham, U.K.) were designed against unique, allele-specific regions of KLRA*01 and KLRA*02, and for KLRA*01/02, a region shared with KLRA*02. Primers were selected in regions conserved between all relevant species: sense BotaKLRA1_SSP1 5′-GGCAAGTCCAATTCGTG-3′, sense BotaKLRA2_SSP2 5′-GCCAAAGTCACAATTCGTG-3′, and antisense KLRA 5′-CCCAAGAAGAACCTGACACTC-3′. These primers were used, where appropriate, to genotype all animals in this study for the presence/absence of KLRA*01, KLRA*02, and KLRA*01/02. PCR amplification of genomic DNA was performed with GoTaq (Promega) for 40 cycles using a minimum of 20 ng DNA. The complete amplified KLRA*02 genomic DNA sequence was submitted to GenBank (accession number MH991705; https://www.ncbi.nlm.nih.gov/nuccore/MH991705).

Genomic enrichment of the cattle KLRA locus, mapping, and variant calling

The variant data used to analyze single nucleotide polymorphisms (SNPs) in this manuscript were generated from enriched genomic DNA samples produced in (29). Briefly, fragmented genomic DNA was used to generate indexed Illumina TruSeq sample libraries, which were then enriched using a custom set of SeqCap EZ oligonucleotide probes (Roche Sequencing). Four libraries were pooled and sequenced on an Illumina MiSeq (2 × 250-bp paired-end reads). In addition to variant calling using SAMtools, the SAMtools phase function was used on the BAM file for each mapping (30), which is a relatively conservative estimate of different haplotypes, as it requires evidence for phasing within individual reads.

Isolation of cattle NK cells from peripheral blood

NK cells were positively selected from Holstein–Friesian PBMCs by labeling with anti-CD335 mAb recognizing bovine NCR1 (AbD Serotec, Kidlington, U.K.) and rat anti-mouse IgG1 MicroBeads (Miltenyi-Biotec, Bergisch Gladbach, Germany) to purities >85%. After washing in RPMI 1640, cells were either cultured with cytokines or frozen for RNA extraction.

Cytokine stimulation assays

NK cells were isolated as described above and cultured in RPMI 1640 supplemented with 10% heat-inactivated FCS, 10 mM HEPES, and 50 μM 2-ME (Sigma-Aldrich with 10 U/ml penicillin/streptomycin) (Microbion, Kidlington, U.K.) and analyzed using FCS Express v3 (De Novo Software, Los Angeles, CA). A minimum of 25,000 events were collected, and variant calling was performed by matching isotype control samples.

RNA isolation, cDNA synthesis, and quantitative PCR analysis of cattle KLRA transcription

Total RNA was isolated from 3 × 10⁶ cytokine-stimulated NK cells or NK cell clones using TRIzol reagent (Life Technologies) according to
manufacturers’ instructions. cDNA was synthesized using SuperScript II with oligo (dT)18 primers (both Life Technologies). Quantitative PCR was used to assess the mRNA levels of cattle KLRA. Assays were performed using Luminaris HiGreen low ROX qPCR Master Mix (Thermo Fisher Scientific) with 250 ng cDNA per reaction. Thermocycling conditions were 2 min at 50°C, 10 min at 95°C before 40 cycles of 15 s at 95°C, and 1 min at 60°C. An additional heating step was run from 60 to 95°C to obtain melting curves. The primers for ATP5B (sense: 5′-CTCTGCTGTGTGCTTATCA-3′; antisense: 5′-CAGGACATGCTTCAAGTCAATC-3′), EIF2B2 (sense: 5′-GAGGATCATTTCACTCAGGA-3′; antisense: 5′-CTGCTGCACAAATGACATG-3′), and SDHA (sense: 5′-GCTTTCCACGTCCATCA-3′; antisense: 5′-AGGCTTCTAGTCTCTCATAGTA-3′) were designed and validated in-house (obtained from Sigma-Aldrich) using the genorm software version 2.5.1 (2, 3). These primers were applied at a concentration of 300 nM. Primers for KLRA*01 and KLRA*02 were custom designed using Locked Nucleic Acid bases (Exigon A/S, Vedbaek, Denmark) and used at a concentration of 900 nM. All primer efficiencies lie within the range of 90–110%.

Statistical analysis

Statistical analyses were performed in R v. 2.15.2 (http://www.r-project.org). Expression levels for each animal were first normalized using three endogenous reference genes (ATP5B, EIF2B2, and SDHA) (20) and then against the mean normalized expression level across either all NCR1-positive samples (for the cytokine stimulation data) or all samples (for the ex vivo cultures). Differences in KLRA expression levels among ex vivo cell populations were analyzed using a linear mixed model for each gene of interest, with log2 normalized expression levels as the response variable, cell population (PBMC, NCR1+, or NCR1−), and comparison (ex vivo or in vivo cell) as fixed effects and animal as a random effect. Differences in KLRA expression levels between cell population (PBMC, NCR1+, or NCR1−) and comparison (ex vivo or in vivo cell) were analyzed using a linear mixed model for each gene of interest, with log2 normalized expression levels as the response variable, cell population (PBMC, NCR1+, or NCR1−), and cell population (ex vivo or in vivo cell) as fixed effects and animal as a random effect. For both analyses, significant (i.e., p < 0.05) factor levels were compared using Tukey honest significant differences.

Results

The two divergent allele lineages of cattle KLRA are the products of a single locus

Cattle contain a unique and expanded NK cell receptor gene repertoire that creates diversity in the NK cell population. We previously reported that although there is only one KLRA gene in the cattle genome assembly (derived from a Hereford), Holstein–Friesian cattle possess two very distinct KLRA allele lineages: Ly-49*01 and Ly-49*02/3 (22). To examine the relationship between these divergent alleles, we used an overlapping exon to exon genomic DNA PCR strategy to amplify the entire KLRA gene—encoding Ly-49*02 and compared this to the known KLRA (Ly-49*01) within the genome assembly. Both genes are almost identical in length at ~15 kb, but 235 SNPs and several small indels create a pairwise sequence identity of 97%. PCR amplification of the full coding cDNA from 15 unrelated Holstein–Friesians never identified more than two alleles from one individual. This strongly implies that the sequences previously described as Ly-49*01 and Ly-49*02/3 are the divergent products of a single KLRA locus, subsequently referred to as the KLRA*01 and KLRA*02 allele lineages.

The coding sequence of both KLRA lineages is conserved despite genomic recombination

Full-length cDNA analysis from the 15 animals above only identified two new KLRA*01 alleles that both differed from KLRA*01 by a single synonymous substitution. This lack of minor variants was surprising considering the extent of polymorphism between KLRA*01 and -02. To study the entire locus in more detail, we used probe-based genome enrichment to pull down the complete KLRA gene and surrounding region of the cattle genome from 17 Holstein–Friesian cattle, three feral B. taurus (two Chillingham and one Kuchinoshima-Ushi), and three animals from the closely related B. indicus subspecies (two Nelore and one Sahiwal) that diverged from B. taurus 1.85 ± 0.15 million y ago (mya) (31). Included in the Holstein–Friesian cohort were 11 animals that had already been analyzed by PCR and were known to be KLRA*01 or KLRA*02 homozygous or heterozygous. The enriched genomic fragments were sequenced and mapped to the extended KLRA locus in the cattle genome. The overall high identity between KLRA*01 and KLRA*02 allowed the reads corresponding to either allele group to map with high confidence.

Enriched genomic sequence data from all the animals, including the feral B. taurus and B. indicus animals, mapped relatively uniformly over the entire KLRA locus and ~100 kb either side. Within the 14 kb KLRA gene, there were four intronic regions not represented by any sequence data, which in total accounted for <1.1 kb of the gene. These regions were all of low complexity or very repetitive and were not conducive to initial probe design. Of the 100 kb either side of the KLRA locus analyzed, a 64 kb region encompassing the KLRA locus contained a highly variable SNP pattern between animals (Fig. 1).

The allele content of each animal could clearly be determined based on the known exonic SNPs that distinguish both KLRA lineages. Based on mapped exon sequences alone, seven animals were homozygous for KLRA*01, thirteen were homozygous for KLRA*02, and five appeared to be heterozygous (Fig. 1). Data from Holstein–Friesian animals with a known KLRA allele content were entirely consistent with the predicted genotype, based on our previous cDNA analysis. However, the SNP patterns at the 5′ end of the gene indicated recombination had likely occurred. In an attempt to more accurately decipher the allele content from heterozygous animals and locate potential regions of recombination, the sequence data from each animal were phased to distinguish both alleles. This method uses SNPs between two genome copies to allocate an individual read to one allele or another. The number of SNPs over this region is relatively high in a whole-genome context. Accordingly phasing successfully separated KLRA*01 and KLRA*02 in animals that were previously known to be heterozygous except for animal 766. This animal was a twin and appeared to contain more than two KLRA alleles as well as other regions of the NKC. A vascular connection between the placentas of cattle twins and the sharing of genetic material is a well-known phenomenon, and therefore, this animal was excluded from further analysis. As the sequences were not entirely contiguous because of the gaps in the pulldown probe coverage, assembled regions were manually labeled as KLRA*01 or KLRA*02 based on identity to the known genomic sequences. The sequences of these contigs were then compared with the same assembled reads for the homozygous animals to provide confidence that they were from the correct KLRA allele lineage. It is, however, possible that a recombination point within one of these low complexity regions has been overlooked.

Domain-by-domain phylogenetic analysis with the phased allele sequences clearly highlighted two regions where recombination between KLRA*01 and KLRA*02 had occurred (Fig. 2). Analysis of SNP patterns revealed that, in the B. taurus animals, there was a break point within a 198-bp region of intron 1–100 bp upstream of exon 2 (positions 4518–4690 in accession number MH991705; https://www.ncbi.nlm.nih.gov/nuccore/MH991705) characterized by adenine cytosine repeats, and in B. indicus, the break point was within a 174-bp region in the middle of intron 2 (positions 1571315–1571118 in accession number KX592814; https://www.ncbi.nlm.nih.gov/nuccore/KX592814).
characterized by 66% adenine thymidine content. The same recombinant alleles were seen six times in the taurine and three times in the indicine cohort, indicating that these alleles are common in both groups. However, as the break point is in a different place between these species, these were likely independent gene conversion events. As some intronic sequence was missing, the phasing was not contiguous across the whole allele, but phylogenetic analysis of the data were entirely consistent with the alleles present at the 5′ end of the gene. It is, therefore, interesting that the consequence of this recombination in both species has been to increase the frequency of the KLRA*02 extracellular domains, as all these recombinant alleles essentially encode an allele of KLRA*02.

To examine the genomic context of this KLRA allele conservation on a background of recombination, we analyzed the phased sequences associated with each allele lineage up to 100 kb either side of the KLRA locus (Supplemental Fig. 1). Our analysis and the current genome builds do not locate any other genes within this region. Phylogenetic analysis of the 100-kb region centromeric of the KLRA gene divides the KLRA*01 and -*02 alleles into two distinct clades, with the recombinant alleles falling within KLRA*01, as expected from the domain-by-domain analysis. Within the KLRA*01 clade, there are three distinct branches, one containing all the B. indicus animals and another more divergent, but distinct, clade containing six Holstein–Friesian and one Kuchinoshima allele (Supplemental Fig. 1). The clear division between the KLRA*01 and -*02 alleles breaks down after the coding region, with less than half the allele sequences from each lineage cladding together. It is clear that the recombination in this region of the genome has been more frequent and complex.

The high resolution of the enrichment data made it possible to characterize allelic variation. In the B. taurus animals, at least seven KLRA*01 and four KLRA*02 minor alleles were present, with an average of 199 SNPs between each lineage, which is entirely consistent with our previous whole-gene amplification. All the KLRA*01 and KLRA*02 allele lineages sequenced in this study can be distinguished from each other between 213 and 223 SNPs over the entire 14 kb of the locus, 18 of which are in the exons, as previously reported (22), and remain completely invariable in this study. Within the Holstein–Friesian dataset, each allele lineage is remarkably conserved, with 16 and 20 variable nucleotide positions between KLRA*01 and KLRA*02 alleles respectively, none of which are located in the exons. Although there has clearly been diversification of at least two KLRA allele lineages, because they shared a common ancestor, they appear to have become fixed in the Holstein–Friesian population.

KLRA allele lineages predate cattle speciation and have been conserved for millions of years

Modern Holstein–Friesian dairy cattle genetics are dominated by relatively few elite bulls, and their offspring that have been selected largely for production traits. To establish the frequency of the KLRA lineages in representative bulls, we PCR typed DNA from a group of elite British and Canadian bulls. Genotyping by PCR using KLRA*01 and -*02 lineage–specific primers revealed a higher frequency of homozygous genotypes than expected from a random distribution (Fig. 3A). This was particularly striking in the Canadian bull cohort, in which 50% of all animals were homozygous for KLRA*01, with only 22% heterozygous. This suggests that there has been positive selection for KLRA homozygous animals, although it is not yet clear if this has been recent artificial selection of breeding bulls. This supports our previous data, as there does appear to be an advantage to maintaining both allele lineages in the population.

To better resolve the evolutionary history of KLRA allele diversification, we amplified intron 4 of the KLRA gene from representative ruminant species and took advantage of sequencing data from ruminant genome projects. Intron 4 is 2.2 kb and the
most variable between lineages. Phylogenetic analysis shows that well before the speciation of cattle, these lineages were clearly distinguishable in species that shared a common ancestor ∼17 mya and suggests that the KLRA*02 lineage may be more like the ancestral sequence (Fig. 3B). Although cattle are the only single species in which we have found evidence of both lineages, we cannot be certain this is unique because of the limited numbers of other species used in this study. However, it is clear that these lineages have persisted for at least 17 mya, and we found no evidence of gene duplication.
KLRA is predominantly transcribed by NCR1⁺ lymphocytes and responds to cytokine stimulation

Lineage conservation suggests an important functional role for KLRA on bovine lymphocytes. To confirm that both alleles have the potential to be expressed on the cell surface, P815 cells were transiently transfected with an expression construct containing KLRA*01 or KLRA*02 that incorporated the V5 epitope on the extracellular C-terminal domain. Flow cytometry confirmed that both allele lineages are trafficked to the cell surface by the intra-cellular machinery and are likely functional (Supplemental Fig. 2).

To explore expression further, mRNA from ex vivo PBMCs isolated from 14 Holstein–Friesian cattle of known KLRA genotype (12 were included in the genome enrichment experiment) was assessed for KLRA allele transcription using quantitative PCR. As KLRA receptors are typically found on NK cells in mice (32), ex vivo NCR1⁺ cells [the key differentiator of NK cells and a subset of NKT-like cells in cattle (33)] and NCR1⁺-depleted lymphocytes were also assessed. These analyses demonstrated that both KLRA lineages were transcribed, with mRNA significantly (p < 0.001) more likely to be transcribed by NK cells (Fig. 4A, 4B, Supplemental Table I). To interrogate whether KLRA lineage expression was likely to vary during an immune response, NCR1⁺ cells were stimulated using three different cytokine regimes. IL-2 and IL-15 primarily promote cell division and homeostasis, whereas IL-12 and IL-18 in synergy induce NCR1⁺ cells to secrete IFN-γ and produce cytotoxic granules. Surprisingly, KLRA*01 and KLRA*02 transcription levels contrasted (Fig. 4C, 4D, Supplemental Table II). Between stimulated samples, KLRA*01 transcription increased after incubation with IL-2 or IL-15 in comparison with IL-12/18. In contrast, KLRA*02 transcription was reduced by the same cytokines, with IL-12/18 having no significant effect. The two KLRA lineages show the opposite transcriptional response to IL-2 and IL-15 stimulation on NCR1⁺ cells.

KLRA lineage copy number correlates with transcription

Both KLRA*01 and KLRA*02 homozygous animals showed significantly higher transcription than heterozygous animals, immediately ex vivo and after cytokine stimulation, with homozygous animals transcribing 3.5- to 8-fold more than homozygous animals (Fig. 4A, 4B). To examine this and the variable expression between each KLRA lineage in more detail, mRNA from a previously generated set of 67 single-cell NCR1⁺ dilution cultures (20) was examined using quantitative PCR. These cultures were derived from six animals that were also included in the genome enrichment experiment. Both KLRA alleles were differentially expressed by individual dilution cultures within and between individuals with 7-fold and 16-fold differences in expression levels for KLRA*01 and KLRA*02, respectively (Fig. 5). This limited sample set also indicated a gene dosage effect may be apparent.
with dilution cultures from homozygous animals transcribing more KLRA than those from heterozygous individuals and higher transcription of KLRA*02. This all points to a functional difference between cells from homozygous and heterozygous individuals that may be linked to the overrepresentation of homozygous individuals in the Holstein–Friesian breeding bulls.

**Discussion**

This study examines KLRA complexity in a large and diverse cohort of cattle to develop a greater understanding of its potential importance in cattle immune responses. We previously described two distinct KLRA allele lineages at the cDNA level (22) and, in this study, found lineage divergence extends across the entire length of both nucleotide sequences at the gene level. Consistent with previous studies (21), we found no evidence of gene duplication in cattle, confirming these allele lineages are transcribed from a single KLRA locus. Lineage-segregating patterns of polymorphism were identified, including 18 stable coding region SNPs identical to those found previously (22). This provided strong evidence these lineages have become fixed during cattle evolution, suggesting they have emerged as beneficial and retain important functional roles. The yak and water buffalo genome sequences contain 17 and 9 of these 18 coding region SNPs, respectively, confirming that this allele diversification predates cattle speciation and has become fixed.

Phasing the enrichment data identified two recombination hot spots toward the 5′ end of the gene. It is interesting to observe that although the exact locations of these two breakpoints differ, they remain relatively close to one another within the gene. The net outcome of these independent conversion events is an enrichment of KLRA*02 extracellular domains, presumably ligand binding, in both recombinant allele groups. Fifteen out of the twenty-two
animals used in this study have at least one KLRA*02 lineage allele despite random sampling. Although the KLRA*02 lineage allele was also the most common in our previous study (22), analysis was restricted to Holstein–Friesian cattle. From selecting a more genetically diverse group of cattle in this study, we found KLRA*02 lineage alleles in every feral B. taurus and two of the three B. indicus animals examined, several of which were recombinants.

In the two groups of breeding bulls studied, we found strong evidence of allelic bias with a greater prevalence of homozygous animals of both KLRA allele lineages. A similar trend was apparent in the animals we randomly selected for genome enrichment. There is presumably a fitness benefit to both allele lineages persisting in the population, which may explain why the frequency of both homozygous genotypes remains consistently higher than expected. This frequency of homozygosity without any obvious loss of fitness may indicate both lineages encode receptors with equally important functions. Our data further demonstrate a higher amount of KLRA transcription in homozygotes, which is unrelated to the allele lineage. This trend is apparent in NCR and NCR1 (with or without cytokine stimulation) and dilution culture datasets, suggesting a potential advantage of a homozygous genotype.

The lineages differ in their ligand binding domains, suggesting they may recognize different ligands, and the discrete patterns of transcription we observed would support this. Relatively higher levels of KLRA*02 are found under steady-state conditions (ex-vivo), with this trend reversed following cytokine stimulation. It is tempting to speculate the two cattle lineages contribute to opposing immune functions, which exist to balance one another. Rodent KLRA genes recognize MHC class I molecules, and although both cattle KLRA lineages are structurally related to their murine equivalents, sequence similarity is low. It has yet to be shown in cattle whether KLRA bind MHC class I or have evolved different ligand specificities.

The transcriptional regulation of Ly-49/KLRA genes in mice has been well characterized (14). Inhibitory KLRA are expressed in a monoeallelic fashion (34), whereas biallelic expression of activating KLRA is apparent (35). Histone acetylation and DNA methylation of a downstream Ly-49 promoter region Pro-2 underlies the regulation of Ly-49a monoallelic gene expression in mice (36). Our data show that cattle have biallelic expression at the lineage level, as heterozygotes always express both inhibitory KLRA*01 and KLRA*02 alleles that invariably contain intact ITIM sequence motifs. Although our quantitative PCR data are unable to determine whether both minor alleles in a homozygote are expressed, biallelic expression in heterozygotes and the higher level of transcription observed in homozygotes suggest monoeallelic expression is unlikely in cattle.

Our wider analysis of KLRA in several divergent ruminant species revealed these lineages were present long before the speciation and subsequent domestication of cattle. This may be a rare example of trans-species polymorphism (37, 38). The apparent presence of only one KLRA lineage in the rest of the ruminant species we examined could be an artifact of low numbers of individuals or nonrandom deletion of a lineage. Regardless, these alleles have become fixed, and any intermediate alleles have either been lost through gradual selection or a recombination event. Although none of the other ruminant genomes we analyzed show more than one gene, there are species (human, horse, and pig) that only have KLRA*01-like genes, which may have emerged in a similar way. Further analysis of larger cohorts of the other ruminant species is needed to determine a more complete evolutionary history.

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Disclosures
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