

RESEARCH ARTICLE

# The 3'-Ja Region of the TCR $\alpha$ Locus Bears Gene Regulatory Activity in Thymic and Peripheral T Cells

Martina Kučerová-Levisohn<sup>1,2</sup>, Stefan Knirr<sup>1</sup>, Rosa I. Mejia<sup>1</sup>, Benjamin D. Ortiz<sup>1,2\*</sup>

**1** Department of Biological Sciences, City University of New York, Hunter College, New York, New York, United States of America, **2** Doctoral Program in Biology, City University of New York, Graduate Center, New York, New York, United States of America

\* [Ortiz@genectr.hunter.cuny.edu](mailto:Ortiz@genectr.hunter.cuny.edu)



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## Abstract

Much progress has been made in understanding the important *cis*-mediated controls on mouse TCR $\alpha$  gene function, including identification of the E $\alpha$  enhancer and TCR $\alpha$  locus control region (LCR). Nevertheless, previous data have suggested that other *cis*-regulatory elements may reside in the locus outside of the E $\alpha$ /LCR. Based on prior findings, we hypothesized the existence of gene regulatory elements in a 3.9-kb region 5' of the C $\alpha$  exons. Using DNase hypersensitivity assays and TCR $\alpha$  BAC reporter transgenes in mice, we detected gene regulatory activity within this 3.9-kb region. This region is active in both thymic and peripheral T cells, and selectively affects upstream, but not downstream, gene expression. Together, these data indicate the existence of a novel *cis*-acting regulatory complex that contributes to TCR $\alpha$  transgene expression *in vivo*. The active chromatin sites we discovered within this region would remain in the locus after TCR $\alpha$  gene rearrangement, and thus may contribute to endogenous TCR $\alpha$  gene activity, particularly in peripheral T cells, where the E $\alpha$  element has been found to be inactive.

## Introduction

The functional rearrangement and expression of the TCR $\alpha$  gene during T cell development results in cell surface  $\alpha\beta$  TCR complex emergence. These processes are tightly regulated in *cis* at the TCR $\alpha$  gene locus [1, 2]. DNase I hypersensitive sites (HS) located 3' of the C $\alpha$  constant region exons comprise a locus control region (LCR) that supports a great deal of TCR $\alpha$  transcriptional regulatory characteristics [3, 4]. In particular, the region of HS1 (which contains the E $\alpha$  transcriptional enhancer [5]) and the HS1 prime (HS1') element [6] is critical for normal TCR $\alpha$  gene rearrangement [7]. This region also very strongly increases transcription levels in T cells *in vivo* [6, 7], likely via epigenetic regulation of chromatin states [8, 9].

While the importance of the HS1/HS1' DNA region to the function of the TCR $\alpha$  locus is indisputable, limited TCR $\alpha$  gene recombination is observed in its absence. This is accompanied by TCR $\alpha$  transcription levels adequate to support maintenance of a peripheral T cell population

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with normal  $\alpha\beta$  TCR levels [7]. It has been speculated that the remaining elements of the TCR $\alpha$  LCR might explain this HS1/HS1' region-independent TCR $\alpha$  gene activity [7]. Nevertheless, heterologous reporter transgenes linked to the TCR $\alpha$  LCR, while mimicking the kinetics/levels of TCR $\alpha$  gene expression in thymocytes [10], are transcribed at lower than expected levels in peripheral T cells [4]. This phenomenon is congruent with prior data indicating that while the E $\alpha$ /HS1 element is active in thymocytes, its deletion from reporter transgenes had no impact on transgene mRNA levels in peripheral lymphoid organs [6]. Using completely different experimental models, a very recent report similarly concluded that, by multiple criteria, the E $\alpha$  element is inactive in peripheral T cells [11]. Together these reports strongly suggest the presence of additional TCR $\alpha$  gene regulatory elements in the locus outside of the LCR capable of maintaining transcription of the TCR $\alpha$  gene in peripheral T cells.

In contrast to heterologous, TCR $\alpha$  LCR-driven reporter genes, transgenic mice bearing cognate TCR $\alpha$  transgenes linked to the full TCR $\alpha$  LCR display normal levels of transgenic  $\alpha\beta$  TCR in peripheral T cells [12]. A major difference between the TCR $\alpha$  transgenes and the heterologous TCR $\alpha$  LCR reporter constructs previously analyzed is that the former include TCR $\alpha$  locus DNA sequences upstream of the LCR up to a SacI restriction site located near J $\alpha$ 3 [13]. The vast majority of this DNA region would remain present in the endogenous locus following functional TCR $\alpha$  gene rearrangement. We hypothesized that transcriptional control elements might be present in this DNA region between the J $\alpha$ 3-proximal SacI site and the LCR. In the present study, we examined this region for indications of gene regulatory activity.

We report the presence of an array of DNase I hypersensitive sites (HS) in a region of the mouse TCR $\alpha$  locus that ranges from the J $\alpha$ 2 segment to the C $\alpha$ 1 exon. We previously described a TCR $\alpha$  locus derived bacterial artificial chromosome (BAC) construct containing two reporter genes [14]. One, V $\alpha$  promoter-driven reporter lies upstream of the HS cluster in the orientation and position of the TCR $\alpha$  gene. The second gene reports the activity of the Dad 1 promoter that lies downstream of both the HS cluster and TCR $\alpha$  LCR. Deletion from this construct of a 3.9-kb region of TCR $\alpha$  locus DNA, that includes the J $\alpha$ 3-proximal SacI site and the identified HS clusters, impairs upstream, but not downstream reporter gene activity in transgenic mice. The deleted region is active in both thymocytes and peripheral T cells. The HS cluster discovered here lies in a region of the locus that would remain in all functionally rearranged TCR $\alpha$  alleles. Therefore, this novel regulatory region may play a role in endogenous TCR $\alpha$  gene activity. It may be especially important to maintaining TCR $\alpha$  mRNA levels in peripheral T cells.

## Materials and Methods

### Ethics Statement

Transgenic animal studies presented in this work have been reviewed and approved by the Hunter College Institutional Animal Care and Use Committee (protocol # BO 10/17-01). Animals are euthanized by carbon dioxide inhalation in conformance with American Veterinary Medical Association recommendations.

### TCR $\alpha$ /Dad1 bacterial artificial chromosome (BAC) dual-reporter constructs

The wild type dual-reporter BAC construct used in this study has been previously described [14]. The mutant BAC was engineered to delete a 3.9-kb region spanning from 38-bp 5' of a SacI site (located between J $\alpha$ 4 and J $\alpha$ 3) to 9-bp 3' of an EcoRV site within the C $\alpha$  constant region exon 1. BAC modifications utilized Red/ET recombination technology (Gene Bridges) [15].

## Transgenic mice

Wild type (76.2-kb) and mutant (72.3-kb) dual-reporter BAC fragments were released from the parent BAC pBACe3.6 vector backbone using NotI and FseI. Transgenic founders bearing intact integrants were identified by Southern blot and PCR screening. Founders were then outcrossed to C57BL/6 mice (Taconic) to establish transgenic mouse lines. The relative transgene copy numbers among the individual mouse lines were determined by multiple Southern blot experiments as described previously [16]. Four independent wild type (line 36, 42, 62 and 71) and four independent mutant reporter BAC transgenic mouse lines (line 4, 18, 25 and 30) are analyzed in this study.

## DNase I Hypersensitivity Assay

These experiments were carried out as previously described [17] using nuclei of MACS (Miltenyi) purified spleen T cells from C57BL/6 mice that were subjected to titrating amounts of DNase I (Worthington). NdeI restriction enzyme digestion was used to generate the 6.7-kb parent fragment of the TCR $\alpha$  locus examined. A 669-bp probe was generated by PCR using the parent BAC clone as a template and the following primers: forward: 5'-atggctgagggaaaggctctacg-3' and reverse: 5'-agaaaagtctctgggaactggtgc-3'. The probe was labeled with [ $\alpha$ -<sup>32</sup>P] dCTP and/or [ $\alpha$ -<sup>32</sup>P] dATP using the RadPrime DNA Labeling System (Life Technologies).

## Flow cytometric analyses

1x10<sup>6</sup> splenocytes were pretreated in 100  $\mu$ L of FACS staining medium (RPMI 1640 supplemented with 3% FBS and 10mM HEPES buffer) for 10 min at 4°C with 1 $\mu$ g of rat anti-CD16/32 (Clone 2.4G2, Life Technologies) to block Fc receptors. Afterwards, 0.2-.5  $\mu$ g of the Abs (from BD Biosciences) were added and incubated for 20 min at 4°C, followed by three washes with FACS staining medium. The Ab clones used were mouse anti-human CD2 (clone S5.2), mouse anti-rat CD2 (clone OX-34), and hamster anti-mouse TCR $\beta$  chain (clone H57-597). Samples were acquired using a FACSCalibur device (BD Biosciences) and the collected data was analyzed with FlowJo software (Tree Star).

## RNA Analysis

5–10  $\mu$ g of total thymic RNA was separated by agarose gel electrophoresis and transferred to neutral nylon membrane (Amersham) for Northern blot analyses. Probes for detection of reporter hCD2 and rCD2 transgene RNA were previously described [15]. A highly specific 18S rRNA 20-mer oligonucleotide [18] was used to control for loading and efficiency of transfer. PhosphorImager analyses were used to obtain normalized transgene expression levels (per copy). For quantitative (q)RT-PCR, RNA samples from MACS purified spleen T cells (>91% purity) were isolated using the RNeasy Mini Kit (Qiagen). The QuantiTect Reverse Transcription Kit (Qiagen) was utilized for cDNAs syntheses. Signals from hCD2-specific primers were normalized to endogenous TCR $\alpha$  as previously described [10]. Signals from rCD2-specific primers (fw: 5'-ccagtgcctgttcaggatacg-3', rev: 5'-ggagttcttctgctcttcagcc-3') were normalized to endogenous Dad1 expression levels. For this, a forward primer specific for exon 1 (that is not present in the BAC reporter): (5'-tgcagttcggtactgtctcc-3') was used with a reverse primer complementary to Dad1 exon 3 (5'-ggaaagtaaggctacagtggagg-3'). qRT-PCR experiments were carried out using DyNAmo HS SYBR Green qPCR Kit (New England BioLabs) and the ViiA7 system (Applied Biosystems).

## Results

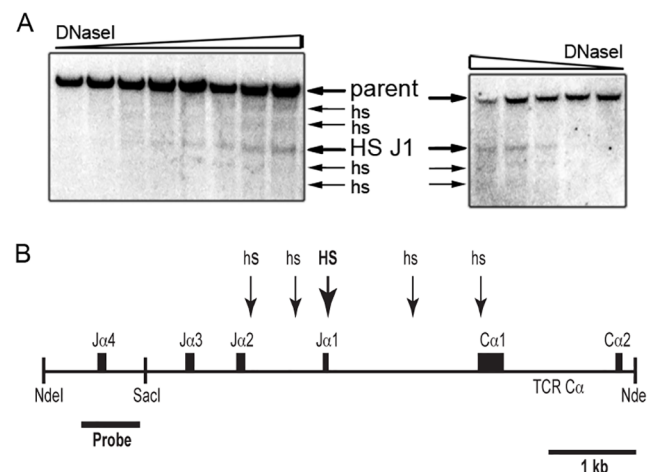
### A region of active chromatin in the 3'-J $\alpha$ region

We used DNase I hypersensitivity assays to examine the chromatin state of a 6.7-kb NdeI fragment of the TCR $\alpha$  locus containing the J $\alpha$ 1 to J $\alpha$ 4 segments and C $\alpha$  exons 1 and 2 (Fig 1). Of the four J $\alpha$  segments present in this genomic fragment, only J $\alpha$ 2 supports functional TCR $\alpha$  protein production [19]. To examine this region for HS in a way that minimizes the potential complication of V $\alpha$ -J $\alpha$  recombination-mediated production of multiple parent restriction fragments, we selected a probe that would only detect alleles at which V $\alpha$ -J $\alpha$  recombination occurred to a J $\alpha$  segment upstream of J $\alpha$ 3. To minimize the presence of non-functional J $\alpha$ 4 rearranged TCR $\alpha$  alleles in the DNase-treated genomic DNA samples, these assays were carried out using nuclei from isolated peripheral T cells.

A cluster of five HS was detected (Fig 1). The most prominent of these HS is proximal to the J $\alpha$ 1 segment (HS-J1), which is located near the center of the HS array. The other four HS are less prominent. In multiple, independent experiments, these four HS display variable intensity in comparison to each other. Two of these HS approximately localize to DNA in between the J $\alpha$ 2 and J $\alpha$ 1 segments. A third HS is found in the J $\alpha$ 1-C $\alpha$ 1 intron, while the fourth localizes to the C $\alpha$ 1 exon. Since J $\alpha$ 1 is a non-functional pseudo-J $\alpha$  segment [19], the region containing the HS array detected would remain present in all functionally rearranged TCR $\alpha$  alleles.

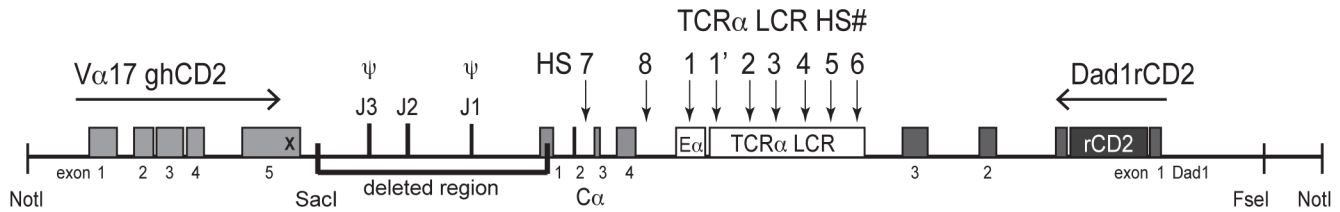
### Testing *cis*-acting TCR $\alpha$ gene regulatory activity in transgenic mice

We previously described a dual-reporter transgene based on a mouse TCR $\alpha$ /Dad1 locus BAC construct (Fig 2) [14]. At the 5' end of the BAC, a human CD2 (hCD2) reporter in genomic configuration is integrated in a position upstream of the C $\alpha$  exons. The hCD2 reporter gene in this construct is driven by a TCR $\alpha$  V-region promoter that was fused in frame with the hCD2 ATG start codon. In addition, a rat CD2 (rCD2) cDNA was fused in frame with the ATG start codon located in Dad1 exon 1. Thus, the rCD2 reporter gene is driven by the Dad1 promoter. To test the function of the above-described target region of the locus, we created a deletion mutant in the context of this dual-reporter BAC that removed a 3.9-kb SacI to EcoRV fragment



**Fig 1. DNase I hypersensitivity sites (HS) within the J $\alpha$ 2 to C $\alpha$ 1 region. (A)** Spleen T cell nuclei from C57BL/6N mouse were subjected to DNase I titration. Arrows indicate the positions of the 6.7-kb NdeI parent fragment and HS. Results from two independent, representative experiments are shown. **(B)** Scaled diagram of the approximate locations of detected HS (arrows). The thick arrow near J $\alpha$ 1 indicates the most prominent HS.

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**Fig 2. Diagram (not to scale) of the TCR $\alpha$ /Dad1 dual-reporter BAC construct [14].** Horizontal arrows indicate the orientation of the two reporter genes. A V $\alpha$ 17 promoter drives expression of a genomic human CD2 reporter gene (ghCD2). The rat CD2 reporter is driven by the Dad 1 promoter. Vertical arrows indicate the location of the HS of the TCR $\alpha$  LCR (including the TCR $\alpha$  enhancer, E $\alpha$ ). The 3.9-kb region deletion in the mutant BAC is marked and runs from 38-bp 5' of the Sacl site through 30-bp 5' of the end of C $\alpha$  exon 1.

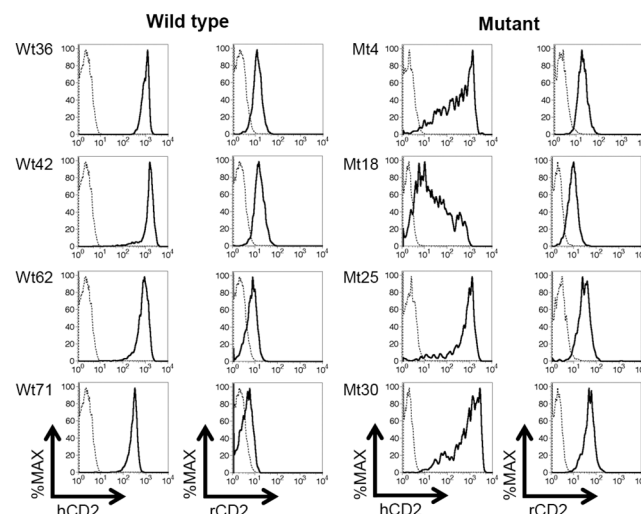
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stretching from upstream of J $\alpha$ 3 to C $\alpha$ 1 (target region). Both wild type and target region mutant BAC constructs were liberated from the vector backbone and used to create transgenic mice.

### Target region deletion impairs upstream, but not downstream reporter gene expression

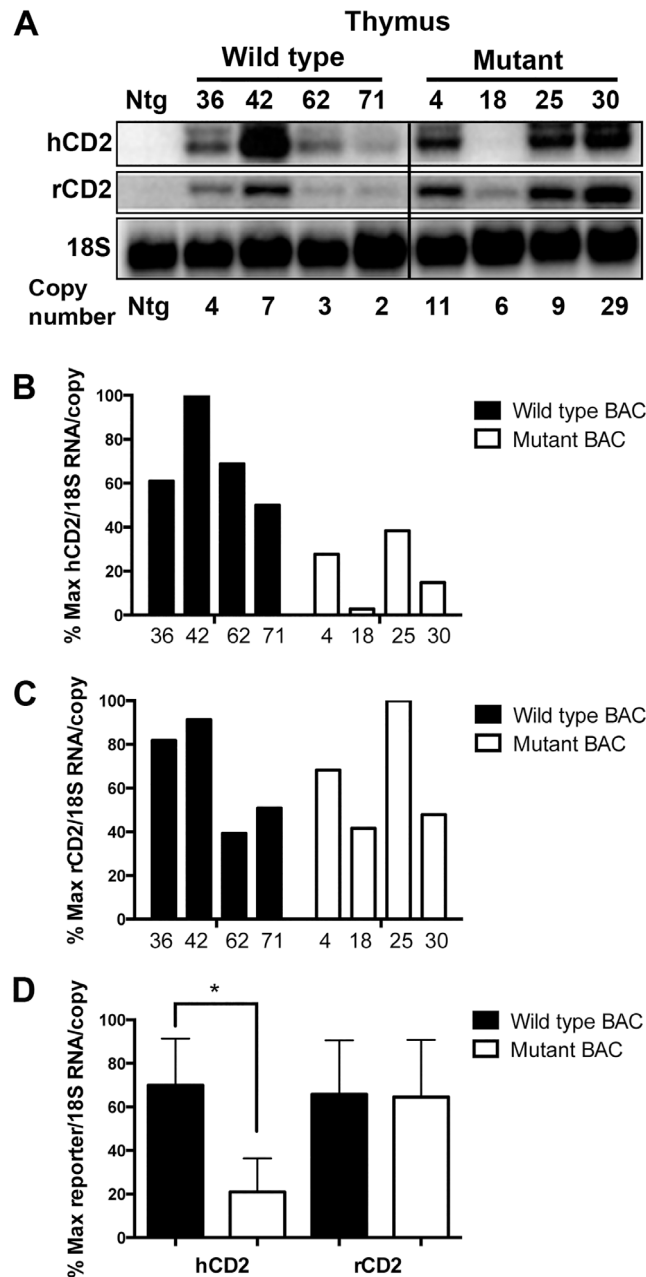
Flow cytometry analyses were carried out to initially assess the impact of target region-deletion on BAC reporter gene expression (Fig 3). Four wild type and four mutant reporter BAC transgenic mouse lines were analyzed. T cells from wild type reporter BAC transgenic mice expressed the upstream hCD2 reporter gene at high and relatively uniform levels on a per cell basis. In contrast, per cell hCD2 expression levels in the target region-deletion mutant BAC were more variegated in transgenic T cells. Expression of the downstream, rCD2 reporter gene was detected in all lines at low, but uniform levels in T cells. The deletion mutation did not appear to affect the uniformity of rCD2 expression levels.

We next analyzed reporter gene mRNA levels in T cells isolated from wild type and mutant BAC transgenic mice. Northern blot analyses of transgenic thymus RNA samples (Fig 4)



**Fig 3. Impaired hCD2 reporter expression in the absence of the deleted region.** Flow cytometry analyses of human CD2 (hCD2) and rat CD2 (rCD2) reporter gene expression in spleen T cells (TCR $\beta^+$ ) from the indicated, independent wild type (Wt) and mutant (Mt) dual-reporter BAC transgenic mouse lines. Reporter gene expression in transgenic (solid line) and non-transgenic control (dashed line) cells is shown.

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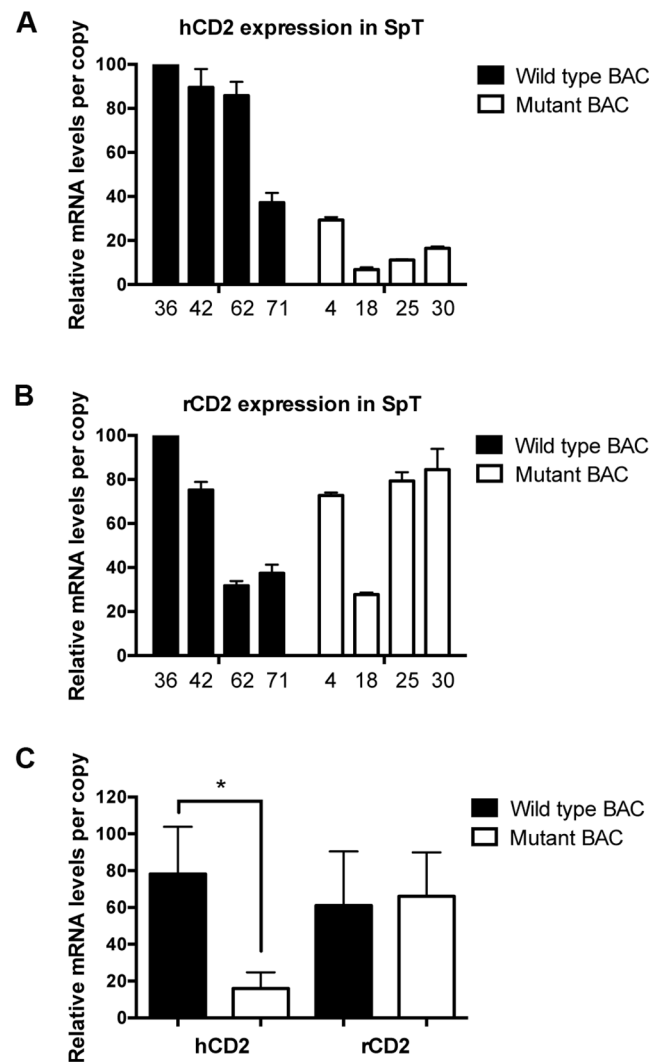


**Fig 4. Absence of the deleted region impairs upstream, but not downstream reporter gene expression in thymocytes.** (A) Northern blot analyses of human and rat CD2 reporter gene mRNA levels in thymocytes from the indicated lines of wild type and mutant reporter BAC transgenic mice. 18S rRNA signals are used as a loading control. Relative transgene copy number for each mouse line is indicated. The black line indicates excision of samples from the blot that are irrelevant to the present study. Panels B and C depict PhosphorImager analyses of the human CD2 (B) and rat CD2 (C) reporter mRNA signals detected by northern blots. The normalized mRNA levels (per transgene copy) from each wild type (black bars) and mutant (white bars) transgenic mouse line are graphed relative to each other (as % maximum). (D) Statistical analyses of the above data using the two-tailed students t test. Graph bars indicate the average ( $\pm$  S.E.) normalized mRNA levels among the lines. The asterisk indicates the statistical significance of the difference in hCD2 mRNA levels between wild type and mutant BAC ( $p = 0.012$ ). In contrast, no significant difference in rCD2 mRNA levels was detected ( $p = 0.942$ ).

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indicated that upstream hCD2 reporter mRNA levels (per transgene copy) driven by the mutant reporter BAC were, on average, about one-third of those observed from wild type reporter BAC. In contrast, downstream rCD2 reporter RNA levels were not affected by the mutation. Similar results were obtained from mRNA-level analyses of transgenic peripheral T cells (Fig 5). Per copy levels of hCD2 mRNA driven by the mutant BAC reporter were, on average, nearly five-fold lower than those observed in wild type BAC transgenic T cells. Once again, the levels of rCD2 reporter mRNA per transgene copy were not altered by the deletion



**Fig 5. The deleted region is functional in peripheral T cells.** qRT-PCR analyses of human (A) and rat (B) CD2 reporter gene mRNA levels (+/- S.E.) in isolated spleen T cells (SpT) from the indicated lines of wild type (black bars) and mutant (white bars) transgenic mice. Observed reporter mRNA levels per copy from each transgenic line are graphed relative to each other (as % maximum). hCD2 reporter mRNA levels were normalized to endogenous TCR $\alpha$  mRNA levels, and rCD2 expression were normalized to endogenous Dad1 mRNA levels using primers that detect sequences not present in the reporter BAC. Three experiments (S1 Table) were performed in duplicates. (C) Statistical analyses of the above data using the two-tailed students t test. The asterisk indicates the statistical significance of the difference in hCD2 mRNA levels between wild type and mutant BAC ( $p = 0.016$ ). In contrast, no significant difference in rCD2 mRNA levels was detected ( $p = 0.819$ ). Graph bars indicate the average (+/- S.E.) normalized mRNA levels among the lines.

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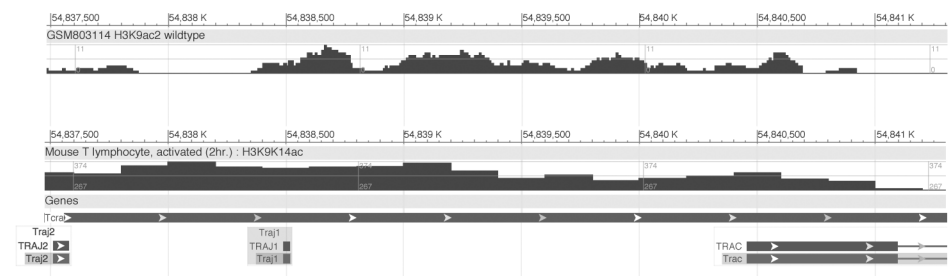
mutation. Overall the data demonstrate the function of the deleted region in regulating TCR $\alpha$ -based reporter transgene expression in both thymic and peripheral T cells.

## Discussion

In this report we describe evidence for the presence of a novel *cis*-acting regulatory DNA region in the mouse TCR $\alpha$  gene locus. It is probable that this activity resides within the region of active chromatin we identified between the J $\alpha$ 2 and C $\alpha$ 1 exons. This DNase hypersensitive region of DNA would remain present in the locus after virtually any functional V $\alpha$ -J $\alpha$  gene rearrangement event. Thus, this regulatory complex could contribute to TCR $\alpha$  gene expression in the thymus and/or periphery. The residual TCR $\alpha$  gene activity observed in E $\alpha$ /HS1/HS1' region knockout mice [7] adds to the rational basis for this notion.

In addition, an early report presented evidence of transcriptional enhancer activity in the J $\alpha$ -C $\alpha$  intronic region of the human TCR $\alpha$  gene locus [20]. The enhancer activity described in that report was weaker (~14-fold) than that reported for the 3' E $\alpha$  element discovered subsequently in the mouse TCR $\alpha$  locus (>100-fold) [5]. This later report presented evidence suggesting that the corresponding J $\alpha$ -C $\alpha$  intronic region of the mouse TCR $\alpha$  locus does not bear significant classical enhancer activity [5]. A contemporaneous study identifying the human counterpart to the 3'-E $\alpha$  enhancer did not include examination of the human J $\alpha$ -C $\alpha$  intron for enhancer activity [21]. In any case, all these prior studies utilized transiently transfected reporter gene bearing plasmids in T cell tumor lines. Here we identified *cis*-acting regulatory DNA 5' of the C $\alpha$  exons that is functional in the context of native T cell chromatin in whole mice. Publicly available data from chromatin immunoprecipitation-next generation sequencing (ChIP-Seq) experiments corroborate these findings. The J $\alpha$  region displays epigenetic signatures of active chromatin, including acetylated histone H3, in T cells. In particular, thymocyte chromatin displays discrete peaks of histone H3 acetylation in the DNA stretching from 5' of J $\alpha$ 1 into roughly the first third of the C $\alpha$ 1 exon. High histone acetylation levels across this DNA region are also observed in peripheral T cells (Fig 6).

A very recent report provided multiple lines of evidence indicating the inactivity of the E $\alpha$  element in peripheral T cells [11]. The conclusions of this report are consistent with a prior report indicating that the removal of E $\alpha$  from a TCR $\alpha$  LCR-driven reporter transgene has virtually no impact on transgene expression in the spleen [6]. Taken together, these findings may help explain why TCR $\alpha$  LCR driven transgenes are expressed at lower levels in peripheral T



**Fig 6. Signatures of active chromatin in the 3' J $\alpha$  region.** Visualization of histone H3 acetylation marks in CD4<sup>+</sup>, CD8<sup>+</sup> thymocytes (top track) and activated peripheral CD4<sup>+</sup> T lymphocytes (bottom track) assayed by ChIP-seq. Shown are screenshots of tracks obtained from publicly available data via the NCBI Epigenomics Browser. The region depicted spans the mouse TCR $\alpha$  gene locus DNA (chromosome 14) containing the J $\alpha$ 2 and J $\alpha$ 1 segments, and the C $\alpha$ 1 constant region exon. Top row: (Unpublished data). ChIP-seq in *Mus musculus*, strain 129SvJae x C57BL/6 (H3K9ac2). Accession number: ESX000004775. Bottom row: [22]. ChIP-seq in *Mus musculus*, strain C57BL/6 (H3K9K14ac). Accession number: ESX000001399.

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cells than those observed in thymocytes [4], because the TCR $\alpha$  LCR includes E $\alpha$  among its functional components [6]. Despite apparent E $\alpha$  inactivity, it is important to point out that the TCR $\alpha$  LCR, as a whole, remains active in the peripheral T cells, and this finding has been confirmed in combination with four different reporter genes, each bearing its own distinct promoter [4, 12, 15, 23]. Previous reports point to at least two distinct TCR $\alpha$  LCR sub-elements outside of E $\alpha$ , named HS1' [6] and HS6 [17, 24], that manifest activity in peripheral lymphoid organs. These elements may also contribute to TCR $\alpha$  mRNA levels in T cell subsets in which the E $\alpha$  element becomes inactive.

There is strong precedent for antigen receptor gene loci bearing multiple, important *cis*-acting enhancer-like elements located both 5' and 3' of their constant region exons. The IgH [25], Igk [26], TCR $\gamma$  [27], TCR $\delta$  [1] loci all display a version of this arrangement. The findings presented here would add the TCR $\alpha$  gene to this category. The literature has produced a consensus that the multiple *cis*-elements functioning within a particular gene locus can support both redundant and non-overlapping functions (e.g. [27–29]). A subset of these functions can have significant impact on immunity. A recent example of this comes from the IgH gene locus. The intronic E $\mu$  enhancer has long been known to play a key role in V-D-J recombination at the IgH locus [25]. But it was also found to have overlapping function with a 3' regulatory region complex [30]. Creation of a functional V-D-J knock-in/E $\mu$  knockout IgH allele bypassed E $\mu$ 's role in recombination to enable investigation of its subsequent functions in mice. These studies revealed a surprising impairment of allelic exclusion at the targeted locus in the absence of E $\mu$  [31]. The downstream consequences of this included faulty clonal selection and generation of autoimmune B cell clones [32]. These findings highlight the continuing importance of identifying and studying *cis*-acting elements that might, on the surface, appear either redundant or dispensable for antigen receptor gene function. The multiple genetic and epigenetic processes that occur at these complex gene loci regulate the assembly, timing, level and distribution of the proteins that form the basis of adaptive immunity. The present study adds a new region of the TCR $\alpha$  locus to the collection of potential sources of regulation of these important processes.

## Supporting Information

**S1 Table. Data from triplicate experiments (Fig 5).**  
(PDF)

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## Author Contributions

Conceived and designed the experiments: MKL SK BDO. Performed the experiments: MKL SK RIM. Analyzed the data: MKL RIM BDO. Wrote the paper: MKL BDO.

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