Mitochondrial metabolism is essential for invariant natural killer T cell development and function

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Edited by Douglas R. Green, St. Jude Children’s Research Hospital, Memphis, TN, and approved February 26, 2021 (received for review October 12, 2020)

Conventional T cell fate and function are determined by coordination between cellular signaling and mitochondrial metabolism. Invariant natural killer T (iNKT) cells are an important subset of “innate-like” T cells that exist in a preactivated effector state, and their dependence on mitochondrial metabolism has not been previously defined genetically or in vivo. Here, we show that mature iNKT cells have reduced mitochondrial respiratory reserve and iNKT cell development was highly sensitive to perturbation of mitochondrial function. Mice with T cell-specific ablation of Rieske iron-sulfur protein (RISP; T−Uqcrfs1−/−), an essential subunit of mitochondrial complex III, had a dramatic reduction of iNKT cells in the thymus and periphery, but no significant perturbation on the development of conventional T cells. The impaired development observed in T−Uqcrfs1−/− mice stems from a cell-autonomous defect in iNKT cells, resulting in a differentiation block at the early stages of iNKT cell development. Residual iNKT cells in T−Uqcrfs1−/− mice displayed increased apoptosis but retained the ability to proliferate in vivo, suggesting that their bioenergetic and biosynthetic demands were not compromised. However, they exhibited reduced expression of activation markers, decreased T cell receptor (TCR) signaling and impaired responses to TCR and interleukin-15 stimulation. Furthermore, knocking down RISP in mature iNKT cells diminished their cytokine production, correlating with reduced NFATc2 activity. Collectively, our data provide evidence for a critical role of mitochondrial metabolism in iNKT cell development and activation outside of its traditional role in supporting cellular bioenergetic demands.

Mitochondrial metabolism | NK cells | T cell development | CD1 | knockout mice

Cellular metabolic pathways are interwoven with traditional signaling pathways to regulate the function and differentiation of T cells (1–3). Upon activation, effector T cells display a marked increase in glycolytic metabolism even in the presence of ample oxygen, termed aerobic glycolysis (4). We have previously shown that despite increased aerobic glycolysis, T cell activation depends on mitochondrial metabolism for generation of reactive oxygen species (ROS) for signaling (5). As activated T cells progress to a memory or regulatory phenotype, they preferentially oxidize fatty acids to support mitochondrial metabolism, and enhanced fatty acid oxidation (FAO) and spare respiratory capacity (SRC) are essential to maintenance of their phenotype (6, 7).

CD1d-restricted invariant natural killer T (iNKT) cells are a unique subset of lymphocytes that exhibit a preactivated phenotype with rapid effector responses (8, 9). iNKT cells are capable of producing large amount of proinflammatory and antiinflammatory cytokines thus have broad immunomodulatory roles (8–10). Given that these cells are poised for rapid proliferation and cytokine production, we hypothesized that coordination of cellular signaling with cellular metabolism will be especially critical for optimal iNKT function. In support of this hypothesis, several studies suggest that modulation of cellular metabolism affects iNKT cell development and function. iNKT cell development is diminished upon deletion of the miR-181 a1b1 cluster, which regulates phosphoinositide 3-kinase signaling and decreases aerobic glycolysis (11, 12). In addition, T cell-specific deletion of Raptor (a component of mTORC1), a metabolic regulator, leads to defects in iNKT cell development and function (13, 14). Loss of folliculin-interacting protein 1 (Fnip1), an adaptor protein that physically interacts with AMP-activated protein kinase, also results in defective NKT cell development, and interestingly conventional T cells develop normally (15). Furthermore, a number of studies targeting bioenergetics processes or related molecules, like alteration of glucose metabolism, mitochondrial-targeted antioxidant treatment, and receptor-interacting protein kinase 3-dependent activation of mitochondrial phosphatase, showed significant effects on iNKT cell ratio and function (16–19). A recent study showed that iNKT cells are less efficient in glucose uptake than CD4+ T cells. Furthermore, activated iNKT cells preferentially metabolize glucose by the pentose phosphate pathway and mitochondria, instead of converting into lactate, since high lactate environment is detrimental to their homeostasis and effector function (20).

In conventional lymphocytes, mitochondria clearly play a role in coordination of cell signaling and cell fate decisions outside of production of energy (5, 21–23). During T cell activation

Significance

We show CD1d-restricted natural killer (NKT) cells have distinct metabolic profiles compared with CD4+ conventional T cells. Mature NKT cells have poor fatty acid oxidation and exhibit reduced mitochondrial respiratory reserve in the steady state. In addition, NKT cell development is more sensitive to alterations in mitochondrial electron transport chain function than conventional T cells. Using T cell-specific mitochondrial complex III ablation in mice, we further demonstrate that mitochondrial metabolism plays a crucial role in NKT cell development and function by modulating T cell receptor/interleukin-15 signaling and NFAT activity. Collectively, our data provide evidence for a critical role of mitochondrial metabolism in NKT cell development and activation, opening a new avenue for NKT cell-based immunotherapy by manipulating NKT cell metabolism.

Author contributions: X.W., A.K., L.C., N.S.C., and C.-R.W. designed research; X.W., A.K., L.C., Y.H., E.M., J.Z., L.A.S., and S.E.W. performed research; X.W., A.K., L.C., E.M., L.V., and C.-R.W. analyzed data; and X.W., A.K., L.C., E.M., L.V., and C.-R.W. wrote the paper.

The authors declare no competing interest.

This article is a PNAS Direct Submission.

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Published March 22, 2021.
mitochondria localize at immune synapses that T cells form with antigen-presenting cells (22). T cell receptor (TCR) stimulation triggers mitochondrial ROS (mROS) production as well as mitochondrial ATP production that are released at the immune synapses and are critical for Ca2+ homeostasis and modulation of TCR-induced downstream signaling pathways (22). We previously showed that mice with T-cell-specific deletion of Rieske iron sulfur protein (RISP), a component of mitochondrial complex III of the mitochondrial electron transport chain (ETC), are defective in antigen-specific T cell activation due to deficiency of mROS required for cellular signaling (5). Several recent studies showed that ROS or factors that affect ROS production are also important in iNKT cell development and effector functions (24–27). In addition, inhibition of mitochondrial oxidative phosphorylation (OXPHOS) by oligomycin has been shown to decrease survival and cytokine production by splenic iNKT cells (20). However, the requirement of mitochondrial metabolism for iNKT cell development and function has not been previously defined genetically or in vivo.

Here we showed that iNKT cells have comparable basal mitochondrial oxygen consumption to conventional T cells but displayed lower SRC and FAO, which are thought to impart cells with mitochondrial reserve under stress. Using Uqcrfs1−/−,CD4+CD8+Cre+ (hereafter referred as T-Uqcrfs1−/−) mice, we showed that abrogation of mitochondrial metabolism resulted in a cell-autonomous defect in iNKT cell development in thymus and periphery. The iNKT cells were able to proliferate but exhibited impaired activation, suggesting that they were not lacking bioenergetically but rather had aberrant TCR signaling in vivo, leading to altered expression of downstream factors required for their terminal maturation. Accordingly, T-Uqcrfs1−/− iNKT cells displayed lower T-bet and CD122 levels and did not respond to interleukin (IL)-15 stimulation. Knockdown of RISP in mature iNKT cells also limited NFATc2 translocation to the nucleus. Collectively, our data highlighted an important role of mitochondrial metabolism in modulating TCR signaling in vivo and regulating iNKT cell development and function.

Results

iNKT Cells Are Metabolically Distinct from CD4+ Conventional T Cells. While most conventional T cells circulate in the body in a naive resting state, NKT cells exhibit a preactivated effector phenotype similar to NKT cells (31, 32). We found that the frequencies of T effector memory (TEM) were comparable between T-Uqcrfs1−/− mice and control mice (8). To determine whether this translates to distinct metabolic programs, we first compared mitochondrial oxygen consumption rate (OCR) between splenic iNKT cells and CD4+ conventional T cells. We found that splenic iNKT cells and CD4+ T cells had comparable rates of basal OCR; however, consistent with a previous study (28), splenic iNKT cells had markedly reduced mitochondrial SRC, as indicated by the lower increase of OCR following uncoupling with fluorocarbonyl cyanide phenylhydrazone (FCCP) (Fig. 1 A and C). Given that iNKT cells are predominant in the liver, we further evaluated SRC in hepatic iNKT cells in comparison with hepatic CD4+ conventional T cells. Similar to splenic iNKT cells, hepatic iNKT cells had barely detectable SRC (Fig. 1 B and C). It is worth noting that hepatic CD4+ T cells exhibited lower SRC than splenic CD4+ T cells (Fig. 1C), indicating the unique tissue environment of the liver influenced metabolic profiles of resident T cells. However, the consistent patterns observed between splenic and hepatic iNKT cells suggest iNKT cells intrinsically possessed a lower SRC independent of organ distribution. SRC has previously been shown to be driven by FAO (4, 7). Therefore, we assessed ability to oxidize fatty acids by measuring OCR following treatment with the fatty acid palmitate-BSA. Both splenic and hepatic iNKT cells had significantly lower increase in OCR after treatment with palmitate-BSA compared to organ-matched CD4+ T cells and reduced sensitivity to etomoxir, an inhibitor of the rate-limiting step in FAO, suggesting reduced rates of FAO (Fig. 1 D and E). A reduction in SRC might be explained by a decrease in total mitochondrial mass. Consistent with previous reports (20, 28), we found iNKT cells had significantly lower mitochondrial mass compared to CD4+ T cells, as indicated by decreased MitoTracker Green staining and decreased relative mitochondrial DNA levels (Fig. 1 F, Upper, Fig. 1G, and SI Appendix, Fig. S1). iNKT cells also had lower mitochondrial membrane potential as shown by reduced tetramethylrhodamine, ethyl ester (TMRE) staining compared with CD4+ T cells (Fig. 1 F, Lower). Collectively, our data reveal that lower mitochondrial mass in iNKT cells is associated with reduced SRC and FAO, which might render them more susceptible or sensitive to mitochondrial injury or stress.

iNKT Cell Development Is More Sensitive to Mitochondrial Perturbation than Conventional T Cell Development. It has been previously reported that mitochondrial content and metabolic activity are dynamically regulated in iNKT cells (27, 29). Inhibition of mitochondrial OXPHOS blocks splenic iNKT cell proliferation and induces their apoptosis (20). To investigate the potential role of mitochondria in the development of NKT cells, we first set up thymic organ culture by culturing thymic lobes from 1-d-old B6 neonates in the presence of FCCP, a potent inhibitor of OXPHOS. In addition, we examined mitochondrial ROS diffusion into the cytosol by reducing mitochondrial membrane potential (5). In newborn thymic organ cultures, iNKT cells started appearing as early as day 2, reaching maximum frequency around day 8 (SI Appendix, Fig. S2A). Interestingly, FCCP treatment led to a significant reduction in iNKT cells in a dose-dependent manner but showed little effect on the development of CD4+ or CD8+ cells (Fig. 2 A–C). The impairment of iNKT cell development was not due to defects in survival or proliferation because low doses of FCCP (i.e., 2.5 μM and 5 μM) did not increase cell apoptosis or reduce cell proliferation (SI Appendix, Fig. S2 B and C). These results indicate iNKT cell development is more sensitive to alterations in mitochondrial ETC function than conventional T cells.

To further investigate the role of mitochondrial metabolism in iNKT cell development and function, we characterized iNKT cells in mice with a T cell-specific deletion of RISP (T-Uqcrfs1−/−) (5). RISP, a nuclear-encoded protein subunit of mitochondrial complex III, is required for transfer of electrons downstream of complex III as well as ROS production. Therefore, RISP-deficiency specifically affects mitochondrial production of ATP and complex III ROS (5). We backcrossed T-Uqcrfs1−/− mice into the B6 background and confirmed efficient ablation of RISP in T cells by immunoblotting (Fig. 3A). As reported previously, the total thymiccellularity and proportion of CD4+ and CD8+ DN (CD4+CD8−), DP (CD4−CD8+), and CD8+ T cells, in the spleen but not liver, was significantly reduced (SI Appendix, Fig. S3A, Fig. S3B, and Fig. S3C), which is consistent with our previous finding that mitochondria are essential for iNKT cell development. We also showed that the impairment of iNKT cells in T-Uqcrfs1−/− mice was due to defects in other T cell subsets selected by DP thymocytes. Indeed, the percentage of CD1d-restricted type II NKT cells in the liver (defined as CD4+CD8−NK1.1+CD1d/PBSS75 tetramer− T cells) (30), was significantly reduced in T-Uqcrfs1−/− mice (Fig. 3E).

We also assessed the effect of RISP-deletion on the development of mucosal associated invariant T (MAIT) cells, another innate-like T cell population that exhibits an effector-memory phenotype similar to NKT cells (31, 32). We found that the frequency and total number of MAIT cells decreased significantly in the thymus and peripheral tissues of T-Uqcrfs1−/− mice (Fig. 3 F and G). In contrast, T-Uqcrfs1−/− mice have normal numbers of conventional CD4+ T cells in the spleen and liver. The total numbers of CD8+ T cells in the spleen but not liver, was significantly reduced (SI Appendix, Fig. S3A, and B). In addition, T-Uqcrfs1−/− mice had normal number of Treg cells in the thymus while their numbers were decreased in the spleen.
were modestly reduced in the spleen and liver (SI Appendix, Fig. S3 C–E). This, RISP ablation (and hence complex III dysfunction) has profound impacts on innate-like T cell development, but only minimally affects conventional T cell development.

**Impaired iNKT Cell Development in T-Uqcrfs1−/− Mice Is Cell Intrinsic.** Having shown that iNKT cell numbers are decreased in T-Uqcrfs1−/− mice, we next questioned whether this defect was cell intrinsic or cell extrinsic. We found that the CD1d expression level on DP thymocytes, the cell type that mediates the positive selection of iNKT cells (9), is comparable between T-Uqcrfs1−/− and WT mice. Similar levels of CD1d expression were also observed in splenic B cells, T cells, and dendritic cells between these two groups (Fig. 4A). Furthermore, we found no difference in IL-2 production by iNKT cell hybridomas in responses to stimulation with either WT or T-Uqcrfs1−/− mice. Although ultimately T-Uqcrfs1−/− mice had lower Vα14 transcripts in total thymocytes and sorted DP thymocytes from WT and T-Uqcrfs1−/− mice, using qRT-PCR. The rearrangement of Vα4 to the distal Jα18 locus is known to require prolonged survival of DP thymocytes. Consistent with reduced iNKT cells in thymus, total thymocytes from T-Uqcrfs1−/− mice stems from a cell-autonomous defect in iNKT cells.

**RISP-Deficiency Affects iNKT Cell Maturation.** iNKT cells branch from the developmental pathway of conventional T cells at the DP thymocyte stage upon a productive Vα14Jα18 TCR-α gene rearrangement (9, 33). Transcriptome data revealed that T-Uqcrfs1−/− DP thymocytes had decreased expression of not only Uqcrfs1, but also several components of ETC, pointing to an overall decrease in the ETC functionality (SI Appendix, Fig. S4). To investigate how this defect affects the iNKT cell development, we first compared the Vα14Jα18 transcripts in total thymocytes and sorted DP thymocytes from WT and T-Uqcrfs1−/− mice, using qRT-PCR. The rearrangement of Vα4 to the distal Jα18 locus is known to require prolonged survival of DP thymocytes. Consistent with reduced iNKT cells in thymus, total thymocytes from T-Uqcrfs1−/− mice had lower Vα14Jα18 transcripts relative to WT counterparts. However, the levels of Vα14Jα18 transcripts in DP thymocytes were comparable between WT and T-Uqcrfs1−/− mice (Fig. S4). Hence, RISP deficiency does not alter the number of thymic iNKT cell progenitors by reducing the lifespan of DP thymocytes.

Following positive selection, iNKT cells progress through conventionally characterized stages: stage 0 (CD24hiCD44loNK1.1+), stage 1 (CD24hiCD44hiNK1.1+), and stage 2 (CD24loCD44hiNK1.1+) occur in the thymus, and terminal maturation to stage 3 (CD24hiCD44hiNK1.1+) occurs in the thymus or periphery (9, 10). In the thymus, we observed higher percentages of stages 0 to 2 iNKT cells in T-Uqcrfs1−/− mice compared to WT mice, while percentage of stage 3 iNKT cells were about fivefold lower (Fig. 5B and C). Although ultimately T-Uqcrfs1−/− mice had significantly

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Based on transcription factor expression profiles and cytokine production, three different sublineages of iNKT cells, namely T-bet+ NKT1, GATA3+ NKT2, and RORγt+ NKT17, have been described (10, 38). Due to significantly lower iNKT cell numbers in T-Uqcrfs1−/− mice, we found lower absolute numbers of all subsets detected by PLZF, RORγt, and T-bet expression (Fig. 5D and E). A higher proportion of PLZF+ cells, enriched for NKT2 and NKT cell precursors (39), was found in T-Uqcrfs1−/− mice compared to WT littermates. Based on CCR7 expression, the NKT cell precursors (CCR7+) appear to be the major population of PLZF− cells in T-Uqcrfs1−/− mice, while NKT2 cells (CCR7−) are the dominant PLZF+ cells in WT mice (Fig. 5F). Furthermore, the proportion of NKT1 (PLZF−T-bet+) was much lower in T-Uqcrfs1−/− mice compared to WT mice, suggesting complex III dysfunction led to a severe defect in interferon-γ (IFN-γ)-producing NKT1 subsets (Fig. 5D and E).

Residual iNKT Cells in T-Uqcrfs1−/− Mice Are Hyperproliferative but Undergo Enhanced Apoptosis. Since iNKT cells have a intrathymic proliferation wave before acquiring maturity, we evaluated if defective maturation of T-Uqcrfs1−/− iNKT cells was due to impaired cell proliferation. Contrary to our expectation, we found higher bromodeoxyuridine (BrdU) incorporation in residual iNKT cells in the thymus, spleen, and liver of T-Uqcrfs1−/− mice (Fig. 5G), indicating T-Uqcrfs1−/− iNKT cells underwent hyperproliferation. In the thymus, T-Uqcrfs1−/− iNKT cells proliferated normally during stages 0 to 2 but exhibited increased proliferation during stage 3 compared with WT iNKT cells (Fig. 5H). We did not observe any difference in BrdU incorporation in conventional T cells (SI Appendix, Fig. S7A). The fact that RISP deficiency does not reduce iNKT cell proliferation led us to examine whether it was affecting Uqcrfs1−/− iNKT cell survival. Indeed, we found iNKT cells in the thymus, spleen, and liver of T-Uqcrfs1−/− mice have higher percentage of Annexin V+ cells as compared with WT mice (Fig. 5I), indicating T-Uqcrfs1−/− iNKT cells are more prone to apoptosis. In thymic iNKT cells, the enhanced apoptosis in T-Uqcrfs1−/− iNKT cells was most profound at stage 2 (Fig. 5J). Consistent with previous reports, we did not detect significant differences in the survival of conventional T cells in T-Uqcrfs1−/− mice compared to WT counterparts (SI Appendix, Fig. S7B). Taken together, these results demonstrate that RISP deletion does not affect iNKT cell proliferation but leads to increased apoptosis of iNKT cells during their development and differentiation. However, impaired iNKT cell development in T-Uqcrfs1−/− mice might not be solely due to an intrinsic survival defect because transgenic overexpression of Bcl-xL in T-Uqcrfs1−/− mice does not restore iNKT cell population in either the thymus or the periphery (SI Appendix, Fig. S8).

RISP-Mediated Mitochondrial Metabolism Is Required for Optimal TCR Signaling. Strong TCR signals are suggested to be crucial for iNKT cell positive selection and the early stages of iNKT cell development (40–42). In T-Uqcrfs1−/− mice, we observed lower expression of Nur77, an orphan nuclear receptor used as a marker for TCR signaling strength (42), on CD40+CD80+ and DP thymocytes as compared to WT counterparts (Fig. 6A and B). This suggested that RISP-deficiency reduces the TCR signaling strength of developing thymocytes. T cell activation is known to elevate ROS and is associated with efflux of calcium ions from the endoplasmic reticulum (ER) into the cytosol (43). To further understand the effect of RISP-deficiency on TCR signaling strength, we examined the TCR-triggered calcium flux in DP thymocytes (containing iNKT cell precursors) from T-Uqcrfs1−/− and WT mice. DP thymocytes from T-Uqcrfs1−/− mice exhibited impaired TCR-driven ER calcium flux upon cross-linking of CD3. Moreover, after addition of Ca2+, store-operated Ca2+ influx in the T-Uqcrfs1−/− mice was also reduced (Fig. 6C). The magnitude of calcium flux

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**Fig. 2.** Disruption of mitochondrial metabolism affects iNKT cells development. Thymic lobes from 1-d-old B6 neonates were cultured in the presence of indicated doses of FCCP for 8 d. CD1d-PBS57 tetramer stained iNKT cells, CD4− and CD8− cells were detected by flow cytometry. (A) Representative dot plots from three independent experiments. (B) Bar graphs depict mean ± SEM of absolute numbers of CD4+ (Left) and CD8+ (Right) cells. *P < 0.05, **P < 0.01, and ***P < 0.001. Lower numbers of total thymic iNKT cells (Fig. 3D), they had comparable numbers of stage 0 iNKT cells. Differences were observed later, where T-Uqcrfs1−/− mice had significantly lower numbers of stages 1 to 3 iNKT cells compared to WT counterparts (Fig. 5C). These results indicate that RISP deletion-related mitochondrial dysfunction does not influence iNKT precursor selection (i.e., stage 0) but exerts its influence as early as the stage 0 to 1 transition, leading to significantly fewer mature iNKT cells.

When examining transcription factors, the expression of retinoic acid receptor-related orphan receptor γ (RORγt) and c-Myc, factors known to control iNKT cells development at a very early stage (34), was not significantly altered in T-Uqcrfs1−/− thymocytes (SI Appendix, Fig. S5 A and B). In addition, T-Uqcrfs1−/− thymic iNKT cells show no defect in the expression of promyelocytic leukemia zinc finger protein (PLZF) or early growth progression 2 protein (Egr2) (SI Appendix, Fig. S5C), two transcription factors essential to induce a multipotent, unbiased effector program that precedes terminal differentiation of iNKT cells (35–37). Thus, the defect in iNKT cell maturation is PLZF and Egr2 independent. In the spleen and liver, total numbers of iNKT cells from all three stages also decreased dramatically in T-Uqcrfs1−/− mice (SI Appendix, Fig. S6 A and B). The increased ratio of stage 1 and stage 2 iNKT cells in T-Uqcrfs1−/− thymus and periphery (Fig. 5C and SI Appendix, Fig. S6 A and B) suggested that maturation of residual T-Uqcrfs1−/− iNKT cells was also impaired.
in T-Uqcrfs1−/− mice was decreased by ~30% both at TCR cross-linking and at Ca2+ addition stages (Fig. 6D and E).

It has been shown that TCR signaling strength distinctly regulates T-bet expression (44). T-bet up-regulates IL-15Rβ (CD122) and increases iNKT cell responsiveness to IL-15 signaling (45, 46), which are essential for iNKT cell terminal maturation and homeostasis (47, 48). Notably, expression of T-bet (Fig. 6F and G) and CD122 (Fig. 6H and I) were significantly reduced in T-Uqcrfs1−/− thymic iNKT cells. In addition, IL-15–induced phosphorylation of STAT5 was impaired in T-Uqcrfs1−/− iNKT cells (Fig. 6J and K).

Fig. 3. RISP-deficiency affects the development of iNKT cells, type II NKT cells, and MAIT cells. (A) Immunoblot of RISP protein in purified TCR-β+CD4+ and TCR-β− splenocytes from T-Uqcrfs1−/− and WT mice (n = 2). (B–D) Lymphocytes were stained with CD1d/PBS57 tetramer or unloaded CD1d tetramer and mAb against TCR-β, CD4, and CD8. CD8− cells were gated for flow cytometry analysis (B). Bar graphs depict mean + SEM of percentage (C) and number (D) of iNKT cells in indicated organs (n = 10). (E) Bar graph depicts mean + SEM of percentage of type II NKT cells in liver lymphocytes (n = 5). (F and G) Lymphocytes from indicated organs and mice were stained with anti–TCR-β and MR1/5OP-RU tetramer. Bar graph depicts mean + SEM of percentage (F) and number (G) of MAIT cells in indicated organs (n = 5). ***P < 0.001; **P < 0.01; *P < 0.05.

Residual iNKT Cells in T-Uqcrfs1−/− Mice Have Defective Responsiveness to TCR Stimulation. To assess whether RISP deletion affects iNKT cell activation and function, we examined CD69 expression and cytokine production in T-Uqcrfs1−/− iNKT cells. We found that residual T-Uqcrfs1−/− iNKT cells in the liver had lower constitutive and TCR-inducible CD69 expression (Fig. 7A and B). Upon in vivo α-GalCer stimulation, T-Uqcrfs1−/− iNKT cells had a profound reduction in IFN-γ and IL-4 cytokine production compared with WT iNKT cells (Fig. 7C and D). Similar to
conventional T cells, iNKT cells down-regulate their TCR upon stimulation (49). We noticed that T-Uqcrfs1"−" iNKT cells had impaired TCR down-regulation while WT iNKT cells down-modulated their TCR normally upon α-GalCer stimulation (Fig. 7 E and F). This provided further evidence of impaired TCR signaling in T-Uqcrfs1"−" iNKT cells. Consistent with our in vivo observations, in vitro stimulation with anti-CD3/anti-CD28 resulted in significantly lower IFN-γ and IL-4 production by T-Uqcrfs1"−" iNKT cells compared to WT iNKT cells from the thymus and the spleen (Fig. 7G). As expected, T-Uqcrfs1"−" iNKT cells produced significantly reduced amounts of TCR-induced mROS as measured by MitoSox red (SI Appendix, Fig. S10 A and B). Consistent with our previous observation (5), splenic CD4+ T cells in T-Uqcrfs1"−" mice had lower TCR-induced mROS production (SI Appendix, Fig. S10 C and D) compared with WT controls. Thus, similar to conventional T cells, T-Uqcrfs1"−" iNKT cells were incapable of increasing mitochondrial complex III ROS upon activation, leading to impaired iNKT cell activation and function.

Knockdown of Uqcrfs1 in Mature iNKT Cells Results in Impaired TCR-Induced Cytokine Production. Since the periphery maturation of residual iNKT cells in T-Uqcrfs1"−" mice is impaired (SI Appendix, Fig. S6), we further investigated the role of RISP-dependent mitochondrial function in mature iNKT cells and CD4+ T cells that developed normally in a WT background. We performed nucleofection of splenocytes from Vu14Tg mice with small-interfering RNA (siRNA) against Uqcrfs1 or scrambled control (NC). We detected ~50% reduction in Uqcrfs1 transcripts upon transfection with Uqcrfs1 siRNA compared to NC siRNA (Fig. S4). The percentages and numbers of iNKT cells and CD4+ T cells (CD1d/PBS57 tetramer+) (Fig. 8 B and C) did not change significantly with ~50% knockdown of Uqcrfs1 transcripts. However, anti-CD3 stimulated IFN-γ production was reduced by ~40 to 50% in both iNKT cells and CD4+ T cells transfected with Uqcrfs1 siRNA compared with siNC-transfected counterparts (Fig. 8 D and E). Thus, RISP-mediated mitochondrial metabolism is essential for optimal TCR-induced cytokine production in mature iNKT and conventional CD4+ T cells.

To further determine the mechanism by which Uqcrfs1 affects cytokine production by iNKT cells, we infected an iNKT cell hybridoma (DN32.D3) with lentiviruses encoding control or Uqcrfs1-targeting short-hairpin RNAs (shRNAs), and specific gene knockdown was verified by immunoblotting. Uqcrfs1 was successfully knockdown with shUqcrfs1-1 and shUqcrfs1-2 compared to shNC (Fig. 8F); further experiments were carried out with shUqcrfs1-2 (greater than 90% reduction of RISP expression). Consistent with Uqcrfs1 knockdown results in primary NKT cells, Uqcrfs1 knockdown in DN32.D3 cells significantly reduced anti-CD3 or α-GalCer-stimulated production of cytokines (IL-2) compared with control shNC-expressing DN32.D3 cells (Fig. 8G). In addition, calcium flux in shUqcrfs1-expressing DN32.D3 cells was significantly reduced compared to that of shNC-expressing cells upon α-GalCer stimulation (Fig. 8 H and I). NFATC2 (or NFAT1) is a key regulator for T cell function and development (50). Hypoxia, which paradoxically increases mROS production from complex III, induces NFATC2 nuclear translocation (51). Therefore, we examined nuclear translocation of NFATC2 to determine whether abrogating Uqcrfs1 activity and thus decreasing mROS production affected iNKT cells similarly to what is known in conventional CD4+ T cells (5). Nuclear translocation of NFATC2 was significantly increased in α-GalCer-treated control cells, but not detected in Uqcrfs1 knockdown cells. Interestingly, the abundance of NFATC2 in the cytoplasmic compartment was also decreased in Uqcrfs1 knockdown cells compared to control NKT cells (Fig. 8 J and K). These data suggest that mROS production from complex III may contribute to cytokine production in mature iNKT cells by regulating NFAT activity.

Discussions

T cells utilize glycolysis and OXPHOS in different ratios during activation, anergy, and development (52). The fine-tuning of metabolic programming throughout a T cell’s lifetime is crucial for proper effector function. iNKT cells are a unique subset of “innate-like” T cells, which constitutively express T cell activation markers and rapidly produce both Th1 and Th2 cytokines upon stimulation. Our study and others (20, 27, 28) suggest that irrespective of organs examined, iNKT cells have lower mitochondrial content and SRC that may be developmentally regulated during their thymic ontogeny (27). We provide further link between poor FAO and lower basal SRC in iNKT cells (Fig. 1). Importantly, through direct pharmacological (Fig. 2) and genetic (Fig. 3) ablation of mitochondrial function, we provide direct evidence that intact mitochondrial respiration is essential for iNKT cell development. Our data revealed the critical role of mitochondrial complex III-related mitochondrial metabolism in regulating iNKT cell development and function through modulation of TCR signaling strength, NFATc2 translocation and T-bet/IL-15 signaling (Figs. 6–8).
The reduced FAO in iNKT cells suggests that they lack the ability to utilize fat as energy source and rely on other carbon sources for their basal OXPHOS (Fig. 1 D and E). In CD8\(^+\) memory T cells, FAO that occurs in mitochondria has been proposed to play a critical role in supporting increased SRC, which is necessary for providing extra energy required for their long-term survival and rapid recall responses (7). Though iNKT cells have a “memory-like” phenotype, a single dose of α-GalCer induces a long-term anergic state of iNKT cells (53). In addition, iNKT cells have been shown to be anergic after certain microbial infections and antigen exposure (54, 55). It has been proposed that T cell anergy is partially due to an inability of the cell to broadly coordinate metabolic pathways for growth and function (52). Thus, we speculate that acquisition of an anergic phenotype in iNKT cells might be correlated with lack of SRC and utilization of FAO for energy. Recent studies showed that high concentration of etomoxir could impact mitochondrial respiration (56, 57) besides inhibiting CPT1, the rate-limiting enzyme FAO. Thus, further investigation of the effect of FAO on NKT cell survival, responsiveness or anergy would require the use of genetic tools.

Metabolic regulators have been shown to affect iNKT cell development by indirectly altering mitochondrial homeostasis (15, 26, 27, 29). For example, T cell-specific loss of Hippo/Mst signaling resulted in altered mitochondrial homeostasis, metabolism, and impaired iNKT cell development (27). Deletion of Fnip1 resulted in lower mitochondrial mass in iNKT cell precursors, but increased mROS production that was associated with their enhanced apoptosis (15). Far from simply being an agent of DNA damage, mROS generation is essential for cellular metabolism and survival, as well as intracellular signaling in T cells (58). Mitochondrial complex III is a crucial component for mROS generation (59). Studies in T-\(\text{Uqcrfs}^{{1/2-}}\) mice lacking mROS production from complex III in T cells demonstrated that conventional CD4\(^+\) T cells required mROS for NFAT nuclear translocation and subsequent production of IL-2 (5). Recent studies have also demonstrated that mice lacking RISP specifically in CD4\(^+\) Treg cells displayed fatal hyperinflammatory disease early in life due to their functional inability to limit inflammation (60). It is important to note that while RISP-deficiency impacts the activation and function of conventional CD4\(^+\), CD8\(^+\), and Treg cells, it does not affect the development of these T cell subsets. In contrast, our studies show a marked dependence of iNKT cells on intact mitochondrial function and mROS for proper thymic development and function. As such, iNKT cells from T-\(\text{Uqcrfs}^{{1/2-}}\) mice are decreased in number, developmentally halted at the stage 0 to 1 transition, and functionally impaired. It is worth noting that the development of other innate-like T cells, such as type II NKT cells and MAIT cells, was also severely impeded in T-\(\text{Uqcrfs}^{{1/2-}}\) mice. In T-\(\text{Uqcrfs}^{{1/2-}}\) mice not only is mROS production reduced upon activation, but the activity of downstream transcription factors, such as Ets-1 (61), PLZF (36, 48), and T-bet (47, 50), is affected. Studies in T-\(\text{Uqcrfs}^{{1/2-}}\) mice have also demonstrated that mice lacking RISP specifically in CD4\(^+\) Treg cells are severely impaired in their ability to limit inflammation (60). It is important to note that while RISP-deficiency impacts the activation and function of conventional CD4\(^+\), CD8\(^+\), and Treg cells, it does not affect the development of these T cell subsets. In contrast, our studies show a marked dependence of iNKT cells on intact mitochondrial function and mROS for proper thymic development and function. As such, iNKT cells from T-\(\text{Uqcrfs}^{{1/2-}}\) mice are decreased in number, developmentally halted at the stage 0 to 1 transition, and functionally impaired. It is worth noting that the development of other innate-like T cells, such as type II NKT cells and MAIT cells, was also severely impeded in T-\(\text{Uqcrfs}^{{1/2-}}\) mice. In T-\(\text{Uqcrfs}^{{1/2-}}\) mice not only is mROS production reduced upon activation, but the activity of downstream transcription factors, such as Ets-1 (61), PLZF (36, 62), Egr2 (37, 63), c-Myc (64, 65), RORyt (66), and T-bet (47, 67). This suggests that there are multiple checkpoints unique to iNKT cell differentiation. WT and RISP-deficient iNKT cells
have shown that RISP-related mitochondrial metabolism may regulate multiple checkpoints in iNKT cell development by modulating TCR signaling strength and T-bet/IL-15 signaling.

First, defects in early iNKT cell development were correlated with decreased TCR signaling strength, as reflected by lower Nur77 expression and decreased calcium flux in DP thymocytes in T-Uqcrfs1$^{−/−}$ mice (Fig. 6). Second, Uqcrfs1$^{−/−}$ iNKT cells suffered high apoptosis during development and showed decreased IL-15 responsiveness (Figs. 5 f and 6 J and K). Third, functional iNKT cell development was disturbed in the T-Uqcrfs1$^{−/−}$ thymus, with T-bet$^{+}$ NKT1 subset particularly affected (Fig. 5 D and E). Thus, our data suggest that RISP-dependent mitochondrial metabolism is essential for optimal TCR signaling strength and subsequent T-bet/IL-15 signaling to support iNKT cell survival, maturation, and function. Interestingly, a recent study showed that altered mitochondrial metabolism due to Mst1 deficiency could affect IL-15 signaling leading to enhanced apoptosis of stage 3 (T-bet-expressing) iNKT cells. However, this effect appears to be independent of aberration in TCR signaling strength (27).

A recent study found that iNKT cells have lower glucose uptake compared to CD4$^{+}$ T cells, suggesting a quiescent phenotype in resting iNKT cells (20). However, activation and effector functions of iNKT cells were accompanied with enhanced OXPHOS (20) and iNKT cell proliferation, survival, and cytokine production were impaired upon OXPHOS inhibition. We had earlier reported that RISP-deficiency did not alter homeostatic proliferation of conventional T cell, but their antigen-induced expansion was affected (5). Our data and the report from Kumar et al. (20) imply that like conventional T cell OXPHOS is not required for homeostatic proliferation of iNKT cells but essential during TCR-induced proliferation and effector functions. However, intact mitochondrial metabolism is required for optimal TCR and IL-15 signaling, thus critical for integrating survival signals in iNKT cells during their development or effector functions. Furthermore, consistent with previous reports, we found that similar to conventional CD4$^{+}$ T cells, mature iNKT cells require OXPHOS and complex III ROS for optimal TCR-induced cytokine production, supporting a signaling role of mitochondria in both T cell types. A previous report showed that ROS regulates the inflammatory function of iNKT cells through PLZF and high levels of ROS in the peripheral NKT cells was primarily produced from NADPH oxidase but not mitochondria (24). However, we found the residual iNKT cells in the spleen and liver of T-Uqcrfs1$^{−/−}$ mice produced reduced amounts of activation-induced mROS and had a profound reduction in cytokine production upon TCR stimulation (Fig. S10 and Fig. 7), supporting a role of mROS in the activation and function of iNKT cells. The decrease in cytokine production was likely due to the impairment of both basal expression and nuclear translocation of NFAT, as shown in the RISP-knockdown NKT cell hybridoma DN32.D3 cells (Fig. 8 J and K). More work is necessary to better understand the mechanism by which RISP regulates the expression of NFAT in NKT cells.

Immune dysfunctions and increased susceptibility to infections have been reported in patients with primary mitochondrial disorders due to ETC dysfunction (23). It will be interesting to study if mitochondrial diseases and associated immune deficiencies due to impaired ETC functions are related to defects in iNKT cell development and function. Furthermore, as discussed above, lower SRC in iNKT cells might be correlated with their enhanced propensity toward an anergic phenotype. It would be valuable to investigate if drugs can be developed to induce mitochondrial biogenesis and enhance SRC in iNKT cells and circumvent anergy, as this might have important implications for iNKT cell-based therapeutic research and development.

**Materials and Methods**

**Mice.** B6, CD45.1 congenic B6 mice, Vv14-Jα18 transgenic (Vv14Tg) mice, TCR-α$^{−/−}$ mice, and Lck$^{−/−}$-Bcl-xL Tg mice were purchased from The Jackson Laboratory. Vv14Tg mice were crossed with TCR-α$^{−/−}$ to generate Vv14TgTCR-α$^{−/−}$.
mice. Uqcrfs1<sup>fl/fl</sup>;CD4-Cre<sup>+</sup> and Uqcrfs1<sup>fl/fl</sup> on B6 background were generated as described previously (5). Jx18<sup>–/–</sup> mice on B6 background were provided by Luc Van Kaer, Vanderbilt University, Nashville, TN. All animal work was approved by the Northwestern University Institutional Animal Care and Use Committee.

Antibodies and Mitochondrial Probes. The generation of CD1d-specific mAb 5C6 has been described previously (68). Allophycocyanin (APC) and Brilliant violet 421 (BV421) conjugated CD1d/PBS57-tetramers and APC conjugated MR1-5-OP-RU tetramers were obtained from the NIH tetramer core facility. The generation of CD1d-specific mAb 5C6 has been described previously (68). Allophycocyanin (APC) and Brilliant violet 421 (BV421) conjugated CD1d/PBS57-tetramers and APC conjugated MR1-5-OP-RU tetramers were obtained from the NIH tetramer core facility. Annexin V apoptosis detection kit and BrdU flow kit were purchased from BD Pharmingen. Fluorescently labeled antibodies against TCR-β, CD4, CD8<sup>a</sup>, CD3, CD44, NK1.1, CD69, CD24, CD244, NK1.1, CD69, CCR7, CD45.2, CD45.1, CD122, IL-4, -α, β, γ<sup>+</sup>/β<sup>−</sup>, -IFN-γ, Nur77, phosphor-STAT5, and biotinylated anti-PARP antibody were purchased from Biolegend, BD, eBioSciences, or Cell Signaling technology. MitoTracker, TMRE for and MitoSox for and MitoSox were purchased from ThermoFisher.

Surface Staining, Intracellular Staining, and Flow Cytometry Analysis. Single-cell suspensions from various organs were prepared and surface stained as described previously (69, 70). For IL-15-induced STAT5 phosphorylation, thymocytes, were stained with CD1d/PBS57 tetramer and incubated with or without IL-15 (100 ng/mL) for 20 min. Cells were fixed with 4% paraformaldehyde (PFA) for 10 min at room temperature followed by methanol permeabilization on ice for 30 min. For staining of transcription factors, cells were surface stained first, fixed, and permeabilized with the Foxp3 staining buffer set (eBiosciences) and stained with relevant antibodies. For in vivo stimulation assay, mice were injected intravenously with 5 μg of α-GalCer. Liver leukocytes were isolated after 1 h and surface stained with CD1d/PBS57 tetramer and anti-TCR-β followed by fixation with 4% PFA. Cells were washed and permeabilized with 0.15% saponin, and then stained with anti-anticytokine mAbs. For in vitro stimulation assays, lymphocytes were stimulated with plated bound anti-CD3 (5 μg/mL) and soluble anti-CD28 (5 μg/mL). Three hours later, monensin (0.01 mM) was added to each well and cells were incubated for another 3 h. Surface staining and intracellular cytokine staining were performed as above. For mitochondrial profiles detection, cells were incubated with 100 nM MitoTracker, 50 nM TMRE, or 5 μM MitoSox for 15 min in 37 °C incubator. Flow cytometry was performed with a FACS Canto II and analyzed using FlowJo software.

Metabolism Assays. Splenic CD4<sup>+</sup> T cells from B6 mice and iNKT cells from Vx14tgTCR-α<sup>−/−</sup> mice were stained with anti-CD4 and CD1d/PBS57-tetramer, respectively, and sorted using BD FACS Aria. Hepatic lymphocytes from WT and T<sup>−/−</sup> mice were positively selected for CD4<sup>+</sup> cells, respectively. Enriched cells were suspended in XF media (nonbuffered RPMI 1640 containing 20 mM glucose, 2 mM l-glutamine, and 1 mM sodium pyruvate) followed by measurement of OCR under basal conditions and in response to 200 μM etomoxir, 1 mM oligomycin, 1.5 μM FCCP, and 100 nM rotenone + 1 μM antimycin A (Sigma) using XF-96 Extracellular Flux Analyzer (Seahorse Biosciences).

Thymic Organ Cultures. Thymic lobes were harvested from 1-d-old B6 neonates, placed on nitrocellulose filters (Millipore), and cultured in DMEM supplemented with 20% FCS, glutamine, sodium pyruvate, nonessential amino acids, 2-mercaptoethanol, and antibiotics, in the presence or absence of indicated concentrations of FCCP at 37 °C for 8 d.

Immunoblotting. Splenocytes from WT and T<sup>Uqcrfs1−/−</sup> mice were stained with APC-anti mouse TCR-β, followed by anti-APC beads (Miltenyi). TCR-β<sup>+</sup> and TCR-β<sup>−</sup> fractions were purified with a magnetic column. Immunoblotting was performed using antibody against RISP (MitoScience, Abcam) or α-tubulin (Santa Cruz) or β-tubulin (Calbiochem), as described previously (5).
Antigen Presentation Assay. Thymocytes (5 × 10^5) isolated from WT and T-Uqcrfs1−/− mice were incubated with 5 × 10^4 iNKT cell hybridomas in the absence or presence of α-GalCer (100 ng/mL) for 24 h. Supernatants were collected and IL-2 production was quantified by ELISA.

Generation of BM Chimeras. BM cells from WT (CD45.1) and T-Uqcrfs1−/− (CD45.2) mice were depleted of T cells with anti–Thy-1 and rabbit complement. Mixed BM cells (1 × 10^7 cells, 1:1 ratio) were injected intravenously into Jα18−/− recipient mice that were irradiated with 900 rad. Lymphocytes
isolated from recipient mice were analyzed by flow cytometry 6 wk after transfer.

Real-Time PCR. Splenic CD4+ T cells from B6 mice and iNKT cells from Vsv14Tg mice were positively selected using MACS (Miltenyi). Genomic DNA and mitochondrial DNA were extracted with QIAamp DNA kit (Qiagen) for determination of mtDNA/nuclear DNA ratio. Total RNA from thymocytes, sorted DP thymocytes or splenocytes transfected with siRNAs were extracted using RNAeasy kit (Qiagen). Following reverse transcription, PCR was performed using SYBR Green PCR master mix with respective primers (SI Appendix, Table S1) and MyiQ real-time detection system (Bio-Rad). Each PCR was run in duplicate or triplicate and normalized to β-actin.

RNA-Sequing Library Generation and Data Analysis. RNA from FACS-sorted DP thymocytes was extracted with RNAeasy Mini Kit (Qiagen). Libraries were generated using the Illumina TruSeq preparation kit and sequenced on HiSeq 4000. Reads were analyzed through the Ceto pipeline (https://github.com/elbarton/NGSbarton) using STAR (71) for alignment on mm10 mouse genome, HTSeq (72) for read counting, and edgeR (73) for differential expression analysis, and gProfiler for gene enrichment analysis (74).

Proliferation and Apoptosis Assays. Mice were injected intraperitoneally with 100 μL of BrdU (10 mg/mL) and supplied with 0.8 mg/mL of BrdU in drinking water for 5 d. Thymocytes were enriched for iNKT cells by depleting CD8+ cells using anti-CD8α antibodies and rabbit complement, while splenocytes and liver leukocytes were enriched for iNKT cells using MACS negative selection (61). Enriched iNKT cells were surface stained with appropriate antibodies followed by staining with BrdU flow kit. To detect apoptotic cells, an annexin V-FITC labeling kit was used according to the manufacturer’s protocol. 

Calcium Signaling. Cells were incubated with Fluoro-4 (4 μM/mL) and Fura Red (9 μM/mL; Molecular Probes) for 45 min in the dark. After washing with Hank’s buffer, cells were blocked with 2.4g2 and stained with anti-CD3–biotin, anti-CD4–biotin (RM4-4), anti-CD4–PE (GK1.5), and anti-CD8–BV510 (53-6.7) for 15 min. Cells were then washed and resuspended in Ca2+-free DPBS and placed in 37 °C incubator. Cross-linking with streptavidin and 2 mM Ca2+ solution was added at the indicated time points.

Transient Knockdown of Uqcrfs1 in iNKT Cells. Splenocytes isolated from Vsv14Tg mice were stimulated with plate bound anti-CD3 (5 μg/mL) and anti-CD28 (2 μg/mL) for 16 h. Nucleofection of stimulated splenocytes with siRNA targeting Uqcrfs1 or scrambled control (Integrated Technologies) was performed following the manufacturer’s instructions (Nucleofection Kit; Lonza). Transfected cells were cultured in complete RPMI supplemented with 200 U/mL IL-2 for additional 3 d followed by assessment of intracellular IFN-γ production.

Stable Knockdown of Uqcrfs1 in DN32.D3 NKT Cell Line. DN32.D3 cells were transduced with lentivirus encoding Uqcrfs1-targeting or NC shRNA (Sigma). Stable knockdown cell line was generated through selection with 5 μg/mL puromycin-containing RPMI 1640 complete medium.

Nfatc2 Knockdown Translocation. DN32.D3 cell lines with stable knockdown of Uqcrfs1 were stimulated with α-GalCer (100 ng/mL) at 0, 1, 2, and 4 h. Isolation of nuclear and cytoplasmic fractions was performed according to the protocol of the Nuclear Extract Kit (Active Motif) followed by Western blotting. NFATC2 was detected by mAB 4G6-GS (Santa Cruz). HDAC1 (Santa Cruz), and α-tubulin (Calbiochem) were used as a loading control for nuclear and cytoplasmic fractions, respectively.

Statistical Analysis. All statistical analyses were performed using Prism software. Comparisons for two groups were performed by Student’s t test. Comparisons for more than two groups were calculated by one-way or two-way ANOVA, followed by Fisher’s least significant difference posttest. Results with a value of P < 0.05 were considered significant.

Data Availability. The data reported in this paper have been deposited in the Gene Expression Omnibus (GEO) database, https://www.ncbi.nlm.nih.gov/geo (accession no. GSE157283). All other study data are included in the article and SI Appendix.

Acknowledgments. We thank Dr. Paul Schumacker for the Uqcrfs1^1/1 mice; Dr. Albert Bendelac for DN32.D3 NKT cell hybridoma; Dr. Sebastian Joyce for N38.2C12 natural killer T cell hybridoma; the NIH tetramer core facility for CD1d and MR1 tetramers; NUSEq Core for library construction and sequencing, and Northwestern University Flow cytometry core facility for cell sorting services. This work was supported by NIH Grants R01 AI43407 and R01 A057460 to (C.-R.W.).

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