CLCNt2 Mediates Nitrate Content in Tobacco Leaf, Impacting the Production of Tobacco-Specific Nitrosamines in Cured Leaves

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Nitrate accumulation in tobacco (Nicotiana tabacum L.) leaf, particularly in the burley (BU) type, is a reservoir for the generation of nitrosating agents responsible for the formation of tobacco-specific nitrosamines (TSNAs). TSNAs are mainly produced via the nitrosation of alkaloids occurring during the curing of tobacco leaves. Additional formation of TSNAs may also occur during tobacco storage, leaf processing and in some circumstances via pyrosynthesis during combustion. Two TSNAs species, 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) and N-nitrosonornicotine (NNN) are found in the tobacco products and have been documented to be animal carcinogens. A previous study showed that decreasing the accumulation of nitrate in tobacco leaf via the overexpression of a deregulated form of nitrate reductase is efficient to reduce the production of TSNAs. We pursue in finding another molecular genetic target to lower nitrate in BU tobacco. Suppressing expression or knocking-out CLCNt2 has a direct impact on leaf nitrate and TSNA reduction in cured leaves without altering biomass. This study provides now a straight path toward the development of new commercial tobacco varieties with reduced TSNA levels by breeding of variants deficient in active CLCNt2 copies.

Keywords: CLCNt2, chloride channel, tobacco, NNN, NNK, nitrate transport, chloride/proton antiporters, tobacco-specific nitrosamines

INTRODUCTION

The International Agency for Research on Cancer has identified eight tobacco-specific nitrosamines (TSNAs) in tobacco and tobacco smoke (IARC, 2004 and the references therein). They are unique to tobacco and are present in smokeless tobacco, snuff, cigarettes, and electronic cigarette liquids. Two TSNAs have been classified as group 1 carcinogens: N-nitrosonornicotine (NNN) and 4-methyl-N-nitrosamino-1-(3-pyridyl)-1-butanone (NNK). For this reason, strategies for reducing TSNA levels in tobacco are being developed. To date, no genetic solution has been found to significantly reduce the levels of NNN. In contrast, a technique known as the Zyvert™ technology (Lusso et al., 2017) can reduce the levels of NNN and, specifically, its precursor nornicotine by knocking out three
Tobacco-specific nitrosamines are generated in tobacco through the nitrosation of alkaloids. The principal TSNA molecules found in tobacco products—NNK, NNN, N-nitrosoanatabine (NAT), and N-nitrosoanabasine (NAB)—are directly produced during the curing process through a reaction of nitrite or any other nitrogen oxide (NOx) species with nicotine (or its oxidized derivative pseudooxynicotine; Hu et al., 2015), nornicotine, anatabine, and anabasine, respectively (Burton et al., 1989; Staaf et al., 2005; Bovet et al., 2012). Bacterial populations present on the tobacco leaf surface are suspected to propagate during curing, thereby facilitating the reduction of nitrate to nitrite or other NOx species to catalyze the nitrosation of the secondary amines in alkaloids (Wienrik et al., 1995; Rundlöf et al., 2000). Post-curing storage of tobacco under improper conditions may also contribute to increased TSNA levels as the result of active nitrosation (Shi et al., 2013).

Plant genetics, agronomic practices, and curing conditions all contribute to the accumulation of TSNA levels from the two tobacco ancestors Nicotiana sylvestris and Nicotiana tomentosiformis (Sierró et al., 2013), were named CLCNt2-S and CLCNt2-T, respectively, according to a previous work published by Barbier-Brygoo et al. (2000). The purpose of this study was: (i) to determine the degree of nitrate reduction in BU plants subjected to suppressed expression using a RNAi approach and to consolidate the results via the knockout of CLCNt2 using CRISPR–Cas9 technology and (ii) to assess its impact on TSNA reduction in cured leaves and cigarette smoke.

**MATERIALS AND METHODS**

### Plasmid Constructs and Plant Materials

The specific DNA fragment selected for suppressing the expression of both copies of CLCNt2 (CLCNt2-S and CLCNt2-T) spans the exon–intron borders within coding exons 4 and 5: 5′-gtcatacatcaggtgtgtgtgtgtggttctcctctctcctcaggttcttcttctctcactttgcttcaggttctttcacctccatttt. This insert was cloned in both sense and antisense orientations using standard Agrobacterium tumefaciens (MMV) promoter and the 3′ nos terminator sequence of the nopaline synthase gene of Agrobacterium tumefaciens (Cheng et al., 1997) within a PMI-proprietary binary vector (WO2012098111).

For the suppressed expression, the BU tobacco variety TN90e4e5 was transformed using standard Agrobacterium-mediated transformation.
transformation protocols (Horsch et al., 1985). TN90e4e5 represents a selection from an ethylmethane sulfonate-mutagenized BU population that contains KO mutations in CYP82E4 and CYP82E5v2 (Lewis et al., 2010) to prevent nornicotine production and, therefore, NNN accumulation. Minimizing the nornicotine content in tobacco is highly desirable, because this compound serves as the direct precursor in the synthesis of NNN. Nornicotine is likely produced almost entirely via the N-demethylation of nicotine, in a process called nicotine conversion that is catalyzed by the enzyme nicotine N-demethylase (NND). Previous studies have identified CYP82E4 as the specific NND gene responsible for the unstable conversion phenomenon, and CYP82E5v2 as a putative minor NND gene (Lewis et al., 2010). Therefore, we used TN90e4e5 as background to silence the expression of CLCNt2 to confirm specifically the impact of potential reduced nitrate on the accumulation of all TSNA, particularly NNN which can also be generated in part by the accumulation of nornicotine. Ten independent T0 plants from each construct were assayed by quantitative reverse transcription PCR (qRT-PCR) to select transgenic lines. T1 seeds were harvested from three independent T0 plants that exhibited the strongest CLCNt2 silencing. T1 plants were grown in the greenhouse and selected for the presence of construct insertion in the genomic DNA by PCR with the following primers (5′–3′): MMV-F (gacgtctaatcccacttcgtc) and IPMS2-R (gacgtctaatcccacttcgtc).

**Phylogenetic Tree**

A multiple sequence alignment was generated using Clustal Omega (version 1.2.4, with the options --full--full--iter parameters), and a phylogenetic tree was constructed from these multiple sequence alignments by using FastTree (version 2.1.10, with the options-gamma-pseudo-spr4-mlacc2-slownni). The resulting phylogenetic tree was re-rooted on the longest branch and ordered on the leaf labels, while preserving the tree topology by using the nw_re-root and nw_order programs from newick_utils (version 1.6). The phylogenetic tree was visualized by using ete3 (version 3.1.1). Labels on the nodes are local support values calculated with the Shimodaira–Hasegawa test. The sequences mentioned are available at https://www.solgenomics.net/organism/Nicotiana_tabacum/genome.

**CLCNt2 Genomic PCR**

Specific primers were designed to amplify DNA fragments by using genomic DNA: CLCNt2-S (forward primer: aacatggttctacaccc; reverse primer: tcaagagatagagacaccacta) and CLCNt2-T (forward primer: aagcgtctaggcctactaatc; reverse primer: cacctccttaagaaagacccagc). As internal control (housekeeping gene), tubulin (TUB) transcripts were amplified using forward primer atttggtacttggtcccaac and reverse primer tctcatgctcaacttcagca. PCR amplification of genomic TUB, CLCNt2-S, and CLCNt2-T fragments was performed by using leaf DNA as matrix template isolated from leaves of the three main N. tabacum types BU (TN90), FC (K326), and OR (Basma Xanthi) and from the tobacco ancestors N. sylvestris and N. tomentosiformis cultivated in the greenhouse using DNeasy Plant Mini Kit (Qiagen, Hilden, Germany).

**RNA-Seq Analyses With Field Grown BU TN90**

Burley variety TN90 plants were grown in a Swiss field. After flowering, the petal, sepal, immature flower, stem, root, and upper, middle, and lower leaves were collected in triplicate and immediately frozen in liquid nitrogen. Total RNA was isolated from each tissue sample (n = 3) by using the RNeasy mini kit (Qiagen, Hilden, Germany). Before cDNA library construction, RNA purity and integrity were checked using the Nanodrop device (Qiagen, Hilden, Germany). mRNA was enriched from total RNA by using a polyA column to generate cDNA libraries for sequencing. All libraries were sequenced on an Illumina HiSeq-2500 sequencer (Illumina, San Diego, CA, United States) by using version 3 chemistry and flow cells with runs of 2×100 bases. The reads were cleaned using Trimmomatic (version .32; Bolger et al., 2014) and aligned to the genome using Hisat2 (version 2.0.1 beta; Kim et al., 2015). Cuffdiff2 (version 2.2.1; Trapnell et al., 2013) was used for calculating the FPKM values based on the aligned reads. The transcriptomes were derived by mapping RNA-Seq reads from each tissue to the following tobacco reference genome. For each plant, three biological replicates were obtained for each tissue. Previously published gene models were used as the basis for differential gene expression analysis (Sierro et al., 2014).

**qRT-PCR Analysis**

The effectiveness of the RNAi constructs in silencing CLCNt2 expression was assessed by qRT-PCR in a greenhouse environment. RNA was isolated from transgenic plants from each independent transformation event and their corresponding control plants, and qRT-PCR was performed using primers (5′ to 3′) F1 (tcccaacatgagccgcaaa) and R1 (aactgctgctagctctccaa) and an internal probe (acactttttctgactac). To measure isoform-specific expression of the two CLCNt2 isoforms in different tissues of the greenhouse-grown TN90 plants, the following primer pairs were used: Nt2S-F (cctttgccaggtccttacaca) and Nt2S-R (agttcagagactagcggc) for CLCNt2-S; and Nt2T-F (cttttgagctttcctatgtg) and Nt2T-R (agttcagagactagcct) for CLCNt2-T. The qRT-PCR reactions which were normalized against a tobacco actin gene by using the following primers: NtACT-F (ccttcagacgctcttata) and NtACT-R (aaactactcactgctgggattc). Each PCR well contained 0.1 μM of forward and reverse primers, 2 μl of cDNA, and 5 μl of iTaq Universal SYBR Green Supermix (Bio-rad) per a 10-μl reaction mix. The reaction mix was heated to 95°C for 3 min and then amplified over 40 cycles of 95°C for 5 s and 62°C for 15 s.

**Field Design**

All field evaluations were conducted at Clayton (NC, United States) using a randomized complete block design with individual plants as the experimental units (39–50 plants per genotype). Agronomic practices typical for BU tobacco production, including topping, were employed. The plants grew for 95 ± 9 days from transplant to harvest time, after which

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1https://www.solgenomics.net/organism/Nicotiana_tabacum/genome
they were cured for 56 ± 10 days. For field-grown samples, leaf tissues were analyzed for nitrate content at the North Carolina State University Tobacco Analytical Service Laboratory as previously described (Lu et al., 2021).

**Alkaloid and TSNA Analysis of Air-Cured Leaves**

Nicotine, nornicotine, anatabine, and anabasine levels in cured upper-stalk leaf samples were quantitated using a Perkin–Elmer AutosystemXL Gas Chromatograph in accordance with previously established protocols (Jack et al., 2007). Total alkaloid content was calculated as the sum of nicotine, nornicotine, anatabine, and anabasine levels. NNN, NNK, NAT, and NAB were quantified in accordance with “Method 1” of Morgan et al. (2004). Total TSNA levels represent the sum of NNN, NNK, NAT, and NAB levels.

**Cigarette Smoke TSNA Analysis**

Test cigarettes were handmade and filled with the electric cigarette injector “Powersmatic 2” by using standard cigarette tubes. The weight of the cut filler was varied (average, 800 mg) to obtain a resistance-to-draw corresponding to 120–140 mm water gauge. The cigarettes were subjected to machine smoking as outlined by Health Canada Method T-115 (1999). Mainstream smoke was collected on Cambridge filter pads and extracted with 100 mM ammonium acetate containing D4-NNN and D4-NNK standards. The extracts were analyzed by LC–MS/MS, liquid chromatography (LC) tandem mass spectrometry (MS), coupled with ESI and quantitated using D4-NNN and D4-NNK internal standards.

**KO Mutations in CLCNt2-S (nt2s) and CLCNt2-T (nt2t) by Using CRISPR–Cas9 Technology**

The CRISPR–Cas9 vector pGREB31 (Addgene) was used for targeted mutagenesis of the CLCNt2 genes. The backbone of pGREB31 is the CAMBIA vector pCAMBIA1300, which has the hptII gene as a selectable marker (hygromycin resistance). After cloning the sequences specifying the desired target site (5’-AGCTGGGTGAACTATATCG-3’) into the sgRNA cassette of pGREB31, the vector was introduced into Agrobacterium strain LBA4404, and T0 plants (TN90) were generated using standard transformation protocols (Horsch et al., 1985). For such KO mutations, we used a TN90 as background, instead of TN90e4e5, to measure the impact of the lack CLCNt2 copies in a tobacco type used in agronomy, the TN90e4e5 plants being not implemented in the field to date. Nuclease-induced mutations were screened by PCR amplification by using primers flanking the target site, followed by DNA sequence analysis of the amplification products. The primers used for amplifying the CLCNt2-S isoform in order to identify CRISPR-induced mutations were Fw 5’-ttggacagaagctctccatat-3’ and Rev 5’-ccctgcagcagtaggtgcaaa-3’.

**Statistical Analysis**

Descriptive statistics, such as mean, minimum, maximum, and variance values, were computed per location and per genotype for all compounds investigated in this study. Potential outliers were identified and removed upstream the statistical modeling. One-way fixed effects ANOVA was performed to assess the variation in the different compounds across the different genotypes. The restricted maximum likelihood (REML) objective function was fit with the Newton–Raphson optimizer to obtain the statistical results. For each genotype tested, the genotypic means and the differences between genotypic means (along with their respective CIs) were calculated for each compound. Then, the relative variations of individual compounds were obtained for all comparisons of interest. Relative variation was defined as the effect of a compound concentration in a plant of a given genotype compared with that in a wildtype (WT) plant. Homogeneity of variances across the different genotypes was checked using the Akaike Information Criterion (AIC), and, when necessary, an additional residual variance was added into the linear model with Satterthwaite approximation for degrees of freedom. SAS software 9.4 was used for the statistical treatments, including Procedure MIXED for the computation aspects. R software 4.0.2 and ggplot2 library were used together for producing the charts.

**RESULTS**

**CLCNt2-S and CLCNt2-T Are the Orthologs of AtCLCa and AtCLCb in Nicotiana tabacum**

Considering the vacuole as the main storage compartment for nitrate (Martinoia et al., 1981; Granstedt and Huffaker, 1982; van der Leij et al., 1998), and the AtCLCa (De Angeli et al., 2006) and AtCLCb (von der Fecht-Bartenbach et al., 2010) proteins as the tonoplast nitrate/proton antiporters involved in nitrate accumulation, our goal was to measure the impact of silencing or knocking out the tobacco orthologs of AtCLCa-b genes on nitrate storage in tobacco leaves. Using tobacco genome resources (Sierro et al., 2014), we were able to confirm that 17 genes encoding CLC proteins are located on the genome, as recently reported by Zhang et al. (2018). Ten NtCLC-like gene products cluster within the subfamily of Cl−/H+ antiporters, and the remaining seven proteins belong to the subfamily of Cl− channels.

A phylogenetic tree additionally populated with the seven and nine CLC proteins found in Arabidopsis thaliana (Marmagne et al., 2007) and Solanum lycopersicum, respectively, is shown in Figure 2A. This allowed us to assign a putative function to the tobacco CLC gene products, principally by sequence analogy subclustering with A. thaliana proteins. Eight of the tomato CLCs have duplicated copies in tobacco, one originating from each of the respective
ancestors of \textit{N. sylvestris} and \textit{N. tomentosiformis}. The exceptions are Solyc02g094060, which has no homolog in tobacco, and Solyc01g112300, which clusters with the tobacco proteins NtCLCeA-S, NtCLCeB-S, and NtCLCe-T (Figure 2A). Interestingly, the \textit{Arabidopsis} nitrate/proton vacuolar antiporters AtCLCa and AtCLCb (De Angeli et al., 2006; von der Fecht-Bartenbach et al., 2010) also have two orthologous proteins in tomato, Solyc02g0206880 and Solyc02g094060. Only the former isoform, however, appears to have been duplicated in tobacco. CLCNt2-S (\textit{N. sylvestris}-related copy) and CLCNt2-T (\textit{N. tomentosiformis}-related copy) correspond to NtCLC6 and NtCLC1, respectively, in the report by Zhang et al. (2018).

To confirm that the \textit{CLCNt2-S} and \textit{CLCNt2-T} genes are present in all three major commercial tobacco types—BU, FC, and OR—and in the respective \textit{N. sylvestris} and \textit{N. tomentosiformis} tobacco species, we performed PCR amplification of genomic DNA by using specific primers. The data confirm the presence of the \textit{N. sylvestris} and \textit{N. tomentosiformis} copies in the three major commercial types of tobacco (Figure 2B). RNA-seq data of the field-grown BU variety TN90 suggests that the \textit{CLCNt2-S} gene is more highly expressed than \textit{CLCNt2-T} in the main organs of the plant, with the highest expression in the petals and stem (Figure 2C). The results of qRT-PCR assays of the greenhouse-grown TN90 plants also support the conclusion that \textit{CLCNt2-S} is the more highly expressed of the two genes.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure2.png}
\caption{(A) Phylogenetic tree of NtCLCs (17 gene products) including the corresponding chloride channel (CLC) orthologs of \textit{Solanum lycopersicum} (nine gene products) and \textit{Arabidopsis thaliana} (seven gene products). The family of NtCLC proteins comprises both chloride/proton antiporters (Cl$^-$/H$^+$; 10 gene products) and CLCs (Cl$^-$; seven gene products). Gene product annotation: AtCLCa (AT5G40890); AtCLCb (AT3G27170); AtCLCc (AT5G49890); AtCLCd (AT5G26240); AtCLCe (AT4G04640); AtCLCh (AT1G56670); AtCLChg (AT5G23280); NtCLCeA-S (Ntab-TN90_AYMY-SS16622:116826-118446); NtCLCeB-S (Ntab-TN90_AYMY-SS16602:130486-145725); NtCLCe-T (Ntab-TN90_AYMY-SS14895:89895-95381); CLCNt2-S (Ntab-TN90_AYMY-SS8473:105857-110645); CLCNt2-T (Ntab-TN90_AYMY-SS15578:118926-119506); NtCLCeB (Ntab-TN90_AYMY-SS8106:77848-84317). (B) PCR amplification of CLCNt2-S and CLCNt2-T fragments by using leaf DNA as the matrix from the three main tobacco types—burley (BU), flue-cured (FC), and oriental (OR)—and from the tobacco ancestors \textit{Nicotiana sylvestris} (Ns) and \textit{Nicotiana tomentosiformis} (Nt). (C) RNA-seq data showing the expression of CLCNt2-S and CLCNt2-T copies in tissues of the main organs of mature field-grown BU plants (TN90 background), mean ($n=3$) calculated with a CI by Cuffdiff2 (version 2.2.1). Relative transcript accumulation of CLCNt2-S vs. CLCNt2-T in various tissues of greenhouse-grown TN90, according to qRT-PCR analysis ($n=3$, average and SD).}
\end{figure}
(Figure 2C), despite the differential not appearing as clear as observed with RNA-seq data.

**Suppressed Expression of CLCNt2-S and CLCNt2-T Affects Nitrate Content and TSNA Production**

We used a genetic approach to determine the impact of CLCNt2-S and CLCNt2-T on leaf nitrate content. For this purpose, we generated an RNAi construct to suppress the expression of CLCNt2-S and CLCNt2-T by incorporating a 407-bp DNA fragment exhibiting 98% identity to both CLCNt2 sequences. The resulting RNAi constructs were expressed under the control of the constitutive MMV promoter in BU tobacco line TN90e4e5, which harbors KO mutations in CYP82E4 and CYP82E5v2 (Lewis and Dewey, 2015) to prevent excess nonnicotine production and, therefore, variable NNN accumulation. To assess the effectiveness of the RNAi suppression, we performed qRT-PCR assays to measure the relative CLCNt2 transcript accumulation in several T0 individuals and selected the three plants that exhibited the strongest inhibition for further investigation. The cumulative levels of CLCNt2-S and CLCNt2-T transcripts were decreased by as much as 80% in the CLCNt2-RNAi lines (designated CLCNt2-1, CLCNt2-2, and CLCNt2-3) relative to WT controls (Table 1).

The CLCNt2-RNAi lines and WT control were grown in a field in the United States by using a randomized complete block design. No obvious phenotypic differences were observed between the WT and CLCNt2-RNAi lines grown in the field (Figure 3A). Plant biomass—determined by directly weighing plants freshly harvested by cutting at the base of the stalk—was not impacted by the suppressed expression of the CLCNt2 genes (Table 1). After each plant on the stalk was air-cured for 10 weeks in a dedicated barn in accordance with the standard practice for BU tobaccos,² the lamina of the fourth leaf from the top of each plant was analyzed for nitrate content. In this experiment, the nitrate content in the three independent CLCNt2-RNAi lines was reduced significantly (up to 68% lower than the nitrate content in WT plants; Figure 3B).

In a randomized subset of the same tobacco samples, we analyzed the impact of decreasing CLCNt2 expression on alkaloid and TSNA content. Interestingly, the alkaloid content was minimally impacted in the CLCNt2-RNAi lines (Figure 4A). In contrast, the TSNA levels in the CLCNt2-RNAi lines, individually and cumulatively, exhibited reductions ranging from 33 to 45% relative to the WT. The impact of CLCNt2 silencing was significant for each TSNA species (NNN, NNK, NAT, and NAB) as well as across the three RNAi lines tested (Figure 4B), which suggested that the nitrosation process is similarly impacted for each of the different alkaloid substrates, nicotine, nornicotine, anatabine, and anabasine.

We determined the TSNA levels in the mainstream smoke of cigarettes made from leaf material of the CLCNt2-RNAi and WT plants. The presence of individual TSNA in mainstream cigarette smoke is primarily due to the thermal release of preexisting TSNA in the cut filler; although, in some circumstances, pyrosynthesis (which involves TSNA formation in the aerosol or in the hot zone of the cigarette by nitrosation of alkaloids precursors) can additionally contribute to the TSNA yield in cigarette smoke (Jaccard et al., 2018). To investigate the effect of CLCNt2 suppression on TSNA transfer and/or synthesis in smoke, the cigarettes were made from pure lamina of CLCNt2-1 plants and WT controls collected from the same location over 2 successive years. Cut filler for cigarette production was generated by finely shredding the cured leaf lamina. Differences in environmental growth and curing conditions can lead to substantial differences in TSNA formation, even in tobaccos that share the same genetic background. This phenomenon was evident when the TSNA levels in cigarette smoke were compared between the 2 years (Table 2). For example, twice as much NNN was observed in the smoke from cigarettes produced from materials grown in year 1 than those grown in year 2, even though the smoke nicotine levels were quite similar between the 2 years. The relative differences between the smoke TSNA levels in the CLCNt2-1-RNAi and WT control; however, were similar across both trial years, with the exception of NAB levels, where the differential was greater in year 2 than in year 1 (Table 2). These data are consistent with the observed reduction in leaf nitrate levels in the transgenic lines, which was also relatively comparable between the two growing seasons: 72% reduction in year 1 and 68% in year 2 for line CLCNt2-1 (data not shown).

**CLCNt2-S and CLCNt2-T KO Impairs Nitrate Storage and TSNA Formation**

Although RNAi suppression is a powerful tool for assessing the consequences of knocking down transcript accumulation of a targeted gene(s), this approach cannot help to establish the effect of complete gene inactivation or assess the relative contribution of multiple genes sharing high sequence identity. To test the effects of complete disruption of CLCNt2-S and CLCNt2-T gene function, both individually and collectively, we created KO mutations in CLCNt2 gene copies in the BU background TN90 by using the CRISPR–Cas9 system. In the T0 generation, an individual was identified (designated CLCNt2#8) that was monoallelic for the same 2-bp deletion both at the CLCNt2-S and CLCNt2-T loci (Figure 5). Given

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²https://www.yumpu.com/en/document/view/11305787/burley-tobacco-production-guide-virginia-tech

**Table 1** qRT-PCR analyses of the CLC-RNAi lines (CLC2-1, CLC2-2, and CLC2-3) compared with the WT plant.

| Genotype | Gene expression (relative level) | Biomass (Kg FW/Plant) |
|----------|----------------------------------|-----------------------|
| WT       | 0.99 ± 0.01                      | 1.62 ± 0.34           |
| CLCNt2-1 | 0.21 ± 0.13                      | 1.49 ± 0.41           |
| CLCNt2-2 | 0.26 ± 0.07                      | 1.59 ± 0.39           |
| CLCNt2-3 | 0.25 ± 0.12                      | 1.59 ± 0.24           |

Average and SD from three analyses (p < 0.001); fold changes are expressed as delta–delta ct values (ΔΔCT). Biomass of fresh cut plants at harvest time from field experiment (see Figure 3); average and SD are shown (n = 39–50); no statistical differences found.
**FIGURE 3 |** (A) Field view of the CLCNt2-RNAi (CLC2-1, CLC2-2, and CLC2-3) and wildtype (WT) plant lines grown in a random design. All plant materials of each genotype were pooled from the same field location (between 39 and 47 viable plants per genotype). (B) Average nitrate content (%DW) in cured leaf lamina of the WT and CLC-RNAi lines (CLC2-1, CLC2-2, and CLC2-3); boxplots of genotype groups are shown ($n = 39–50$), with the difference of nitrate content observed between the WT and CLCNt2 lines being $-67.8\% (-51, -84.7\%); p < 0.0001^{***}$, as per the results of ANOVA with a CI of 90%.

**FIGURE 4 |** Alkaloid (nicotine, nornicotine, anatabine, and anabasine) and TSNA (NNK, NNN, NAT, and NAB) content in the cured leaf lamina of CLC-RNAi (CLC2-1, CLC2-2, and CLC2-3) and WT plants grown in the field. Data were collected from 10 plants randomly selected after curing. The charts represent the box plots of genotypic groups, with no significant differences in alkaloid content between the WT and CLCNt2 lines (A). For TSNA content (B), the difference observed in NNK levels between the CLCNt2-1 lines and WT was $-33.2\% (-10.6, -55.8\%); p = 0.018^{**}$; for NNN levels, the difference observed between the WT and CLCNt2 lines was $-44.5\% (-19.8, -69.1\%); p = 0.004^{***}$; for NAT levels, the difference observed between the WT and CLCNt2 lines was $-40.3\% (-14.4, -66.2\%); p = 0.013^{**}$; for NAB levels, the difference observed between the WT and CLCNt2 lines resulted in $-34.7\% (-4.7, -64.6\%); p = 0.056^{*}$. An ANOVA test with a CI of 90% was used.
TABLE 2 | Nicotine, NNK, NAT, and NAB content in the mainstream smoke of cigarettes made from the lamina of WT and CLCNt2-1 leaves.

|                | Year 1                        | Year 2                        |
|----------------|------------------------------|------------------------------|
|                | WT (mg/cig)  | CLCNt2-1 (mg/cig) | Percentage | p-value | WT (mg/cig)  | CLCNt2-1 (mg/cig) | Percentage | p-value |
| Nicotine       | 3.17±0.01   | 3.79±0.15  | 119%       | -        | 3.15±0.14   | 2.81±0.07  | -89%        | -        |
| NNN            | 564±12      | 274±10     | −51%       | <0.001   | 269±15      | 131±4       | −51%        | <0.001   |
| NNK            | 178±13      | 116±6      | −34%       | <0.001   | 144±7       | 86±1        | −40%        | <0.001   |
| NAT            | 520±32      | 219±6      | −58%       | <0.001   | 349±16      | 150±6       | −57%        | <0.001   |
| NAB            | 129±12      | 59±10      | −54%       | <0.001   | 65±2        | 19±1        | −71%        | <0.001   |

For 2 successive years, leaf materials from each genotype were pooled. The data shown represent the combined means and SD from four smoke analyses (n=4). All data are statistically relevant as per pairwise t-tests (p<0.001).

FIGURE 5 | CRISPR-Cas9-induced knockout (KO) mutations in CLCNt2-S and CLCNt2-T. The region of the WT CLCNt2 sequence that was targeted by the CRISPR-Cas9 construct is shown on top, with the mutant variants found in line CLCNt2#8 underneath. The 20-bp CRISPR-Cas9 target sequence is underlined, and the PAM site is in green type. Dashes represent bases that have been deleted. WT amino acids are shown in black type; unique amino acids predicted to be present on the C-terminus of the mutant truncated proteins because of frame shifts caused by the two deleted base pairs are shown in red type. Asterisks represent termination codons. Numbers in black correspond to the cDNA sequence beginning at the ATG start codon, and orange numbers indicate the amino acid position. Nucleotides and amino acids that are polymorphic between the CLCNt2-S and CLCNt2-T isoforms are shaded gray.

that these small deletions cause a frame shift that would result in the production of highly truncated CLCNt2 proteins that lack over 80% of the normally present amino acids, it can reasonably be assumed that the mutant gene products are incapable of functioning as nitrate transporters.

Plant CLCNt2#8 was self-pollinated, and over 200 T1 plants were screened by PCR using primers specific for the nptII selectable marker gene. Those plants that had lost the CRISPR transgene through segregation were genotyped by DNA sequence analysis in order to select plants homozygous for mutations in the individual CLCNt2-S (nt2s) and CLCNt2-T (nt2t) genes (nt2s/nt2T and Nt2T/nt2t) as well as the double-homozygous mutant genotype (nt2s/nt2t). A single line of plants carrying each individual mutation and two double-homozygous mutant lines were grown and evaluated in the field together with the WT as control. After harvest, plant biomass was determined by weighing the plants directly in the field before air-curing, in the same manner as that described above for the CLCNt2-RNAi plants. Plant biomass was not affected by the presence of the clcnt2-KO mutations in the Nt2S/nt2T or double-homozygous mutant lines. The homozygous mutant clcnt2-s line (nt2s/nt2T), however, unexpectedly displayed a statistically significant 25.9% reduction in biomass relative to the rest of the lines (Figure 6A). After curing, the leaf lamina from the same upper stalk position of each plant was analyzed for nitrate content. Despite the observation that CLCNt2-S transcripts were more prevalent than those corresponding to CLCNt2-T across every plant tissue examined (Figure 2C), neither single-mutant line displayed a nitrate phenotype that differed significantly from that of the WT control or each other (Figure 6B). In contrast, a significant 61.8% reduction in nitrate was observed in the two double-mutant lines (nt2s/nt2t), consistent with the data obtained with the CLCN2-RNAi lines (Figure 3B).

Alkaloids and TSNA were measured in the two independent double-KO lines (nt2s/nt2t). Surprisingly, the inactivation of both CLCN2 copies in tobacco also significantly impacted the alkaloid content in leaves (Figure 7A), which was not observed when CLCN2 transcript accumulation was knocked down in the CLCN2-RNAi lines (see Figure 4A). The decrease was particularly marked for anabasine, which showed a decrease of 45.2% relative to the WT levels. Lower nicotine, nornicotine, and anabasine levels were also found to be statistically relevant in the double-homozygous clcnt2-KO plants, with reductions ranging from 23.9 to 34.9% relative to the WT (Figure 7A). As expected, because of the low-nitrate phenotype, the TSNA content in leaves was also substantially affected in the clcnt2-KO plants. The levels of all four TSNA (NNN, NNK, NAT, and NAB) were clearly decreased in the two clcnt2 double-mutant lines (nt2s/nt2t), with a reduction of 44% in NNN, 35.7% in NNK, 62.8% in NAT, and 68.6% in NAB levels relative to the WT (Figure 7B). Despite the mean NNN values being lower, this measurement was not deemed significant owing to the exceptionally large variation measured among the plants, as attested by the minimum and maximum percentage values.
Such variations were likely due to the use of TN90 lines as background for developing the clcnt2-KO plants, instead of TN90e4e5 in case of the CLCNt2-RNAi plant materials. Indeed, the TN90e4e5 cultivar (Lewis and Dewey, 2015) is similar to the Zyvert™ varieties, having lower and more stable nornicotine levels than TN90. Cumulatively, the data show that suppressed expression or inactivation of CLCNt2 genes resulted in a nitrate-underaccumulation leaf phenotype that impacted the TSNA formation.

**DISCUSSION**

We went through the hypothesis that tobacco homologs of AtCLC2a and b may represent good targets for reducing nitrate, and thus TSNAs. Therefore, to test this hypothesis, we generated RNAi lines to target the CLCNt2 downregulation. Because the field results of the RNAi lines were promising, this justified going through the effort of knocking-out the two CLCNt2 gene copies using genome editing to precisely investigate the respective role of S and T homeologous copies of CLCNt2.

We have effectively demonstrated that decreasing the expression of CLCNt2-S and CLCNt2-T in tobacco limits the storage of nitrate in the leaf, thus reducing the propensity for TSNA formation in the cured lamina and in the smoke of cigarettes made from these tobaccos. Mechanistically, this is best explained by a model, whereby less nitrate is available for reduction by nitrate-reducing leaf surface microbes during the curing period, which limits the formation of nitrite and/or other NOx species that can catalyze alkaloid nitrosation.

In a previous study, we demonstrated that constitutive expression of a deregulated NR enzyme dramatically decreases the free nitrate levels in leaves to approximately 5.9% of that observed in the WT material (Lu et al., 2016). The strategy in this case was based on actively increasing the reduction, and thus assimilation, of free nitrate entering the cytosol of the cell. In the present study, inactivation of the two CLCNt2 copies, either by using a classical RNAi construct or through CRISPR–Cas9-mediated mutagenesis, led to a reduction in free nitrate levels to approximately 30–40% of those observed in WT leaves. On the basis of gene homology with *A. thaliana*, we can reasonably speculate that the CLCNt2 proteins are localized in tonoplast membranes and are responsible for storing nitrate reserves in tobacco leaf vacuoles. The absence of a tonoplast NO$_3^-$/H$^+$ exchanger likely prevents vacuolar storage in CLCNt2-RNAi or clcnt2-KO plants and, therefore, favors the reduction of free nitrate by the endogenous NR toward assimilation. It has been demonstrated that vacuoles from plants grown in the presence of nitrate likely do represent the major nitrate storage organelle in the plant, but not the only one, accounting for about 58% of the total cellular free nitrate content, according to one study (Granstedt and Huffaker, 1982). Stems and, in particular, xylem vessels may also serve as nitrate repositories (Dechorgnat et al., 2011). Our data indirectly suggest that a nitrate storage compartment(s) other than vacuoles could be present in the laminar tissue of tobacco leaves, possibly explaining the lower magnitude of the nitrate reduction in
tobacco plants deficient in CLCNt2 function. These plants have normal NR cycles of activation/inactivation compared with tobacco plants that overexpress a permanently activated NR (Lu et al., 2016). Alternatively, a portion of the nitrate remaining in CLCNt2-compromised plants may be due to the presence of additional transporters or NtCLCs whose function has yet to be elucidated. In Arabidopsis, two members of the broad-spectrum peptide transporter/nitrate transporter 1 (PTR/NRT1) family have been shown to be localized to the tonoplast, though their substrates could not be defined when expressed in heterologous systems (Weichert, et al., 2012).

Our data confirmed that decreasing tobacco plant free nitrate content has a direct impact on TSNA levels both in cured material and cigarette smoke. In our prior study, field-grown tobacco plants expressing the 35S:S523D-NR transgene exhibited greatly reduced levels of TSNs in cured material—corresponding to 44, 64, 65, and 32% lower levels of NNN, NNK, NAT, and NAB, respectively, compared with the levels in cured leaf from WT control plants—and thus an observed total TSNA reduction corresponding to 52% (Lu et al., 2016). In the CLCNt2-RNAi lines of the present study, we observed similar reductions in NNN and NAB levels in the cured leaf, while the reductions in NNK and NAT levels were less dramatic (Figure 4). Except in the case of NNN, the reductions in TSNA levels in the clcnt2-KO lines tended to be greater than those in the CLCNt2-RNAi materials. Given the similar nitrate-lowering efficiencies of the two CLCNt2 inhibition strategies, the enhanced TSNA reduction in the clcnt2-KO lines could not be attributed to these plants possessing lower levels of nitrate per se. Instead, it is likely that the reduced alkaloid phenotype observed in the double-homozygous clcnt2-KO lines accounted for the greater TSNA reduction, as both nitrate and alkaloid contents are highly correlated with TSNA levels (Lu et al., 2021). As nitrate is the fuel for such pathways, secondary metabolite pathways possibly rely more on stored resources than on directly assimilated ones. The absence of a similar trend with NNN is likely because we used the TN90e4e5 genotype that possesses nornicotine-stabilizing mutations in the RNAi-based study, as opposed to the TN90 background we chose for the KO experiments that lack these mutations.

The mainstream smoke from cigarettes made with tobaccos possessing the 35S:S523D-NR construct showed 76% lower total TSNA content than the mainstream smoke from WT
cigarettes (Lu et al., 2016); in contrast, the mainstream smoke from CLCNt2-RNAi cigarettes showed 52% lower total TSNA levels than WT cigarettes, a value close to that observed in the tobacco filler (48%). This suggests that, in comparison to 35S:S523D-NR cigarettes, cigarettes made of CLCNt2-RNAi tobacco generate a more substantial amount of TSNA by pyrosynthesis during the smoking process (Jaccard et al., 2018), which is consistent with the levels of nitrate measured in the respective tobacco fillers. Indeed, the extremely low nitrate content measured in the 35S:S523D-NR cut filler certainly more actively circumscribes the nitrosation process occurring during pyrosynthesis compared with what one would predict could occur in CLCNt2-RNAi cured materials upon combustion, given the greater levels of free nitrate in the latter.

It is necessary to inhibit the function of both of the tobacco CLCNt2 genes, CLCNt2-S and CLCNt2-T, to decrease the nitrate levels in WT plant leaf tissues, although CLCNt2-S is expressed at substantially higher levels than CLCNt2-T in the leaves of TN90 plants (Figure 2C). Consequently, either the level of CLCNt2 protein produced in the cell is greater than that required for storing excess nitrate in the vacuole, or a crosstalk exists between the two genes, where a deficiency in one may lead to upregulation of the other. The biomass of the controls of the CLCNt2-RNAi and clcnt2-KO trials is supposed to be constant. The observed differences capture the effects of different crop seasons and limit the direct comparison of CLCNt2-RNAi and clcnt2-KO plant biomass. Some level of redundancy exists between the two CLCNt2 genes sharing 98% protein sequence identity, nevertheless, the lower plant biomass of the single clcnt2-s compared to control or clcnt2-t suggests that the function of the two CLCNt2 genes may not be totally overlapping (Figure 6A). We have preliminary data suggesting that the flowering time is altered in mt2s/mt2T, but not in Nt2S/mt2T plants, in comparison to WT TN90—a phenomenon that could account for differences in plant biomass, as recently documented in our follow-up study on low-nitrate-containing tobacco plants possessing the 35S:S523D-NR gene (Lu et al., 2021). Elucidation of the molecular origin of this phenotype is currently an area of active investigation.

Surprisingly, unlike the results observed when the CLCNt2-RNAi knockdown lines were grown in the field, KO of both CLCNt2 genes resulted in a reduction in the alkaloid content, with anatabine being impacted to the greatest extent. Given that alkaloid production occurs in the root, these results suggest that CLCNt2 activity within this tissue plays a role, whether direct or indirect, in the production of tobacco alkaloids. In most crop fertilization regimens, nitrate is the primary source of nitrogen for biosynthesis of Asp, Orn, and Arg amino acids, from which pyridine amino acids are derived (Dewey and Xie, 2013). One model that could explain these results is that the absence of CLCNt2 function in the root hinders nitrate storage within root vacuoles, leading to a greater proportion of the nitrate being directly mobilized to the aerial tissues of the plant, where it cannot be routed toward alkaloid production. That we did not observe this phenomenon in CLCNt2-RNAi plants might be attributable to: (1) the RNAi-mediated suppression of CLCNt2 being less effective in the root than in the leaf and/or (2) the low levels of CLCNt2 protein still produced in knockdown plants, as opposed to that in KO plants, is sufficient to meet the needs for vacuolar sequestration of nitrate within root tissue. Alternative models could be envisioned whereby the disruption of nitrate storage pools in clcnt2-KO plants impacts the production of alkaloids, particularly anatabine, via alteration of signaling pathways that regulate alkaloid biosynthesis.

**CONCLUSION**

The plants produced in this study offer new materials for studying the origin of and relationships between nitrogen stocks and the nitrogen pathways leading to alkaloid synthesis, an area of investigation which has yet to be adequately addressed (Xi et al., 2008; Zenkner et al., 2019). From the perspective of practical application, this study provides a promising solution to the problem of TSNA accumulation in tobaccos. Although overexpression of a deregulated NR has also proven effective in decreasing TSNA levels via the reduction of cellular nitrate pools (Lu et al., 2016, 2021), that approach is mediated by a transgene and is thus subject to limitations that hinder commercial deployment. Breeding of variants deficient in active CLCNt2 copies, however, offers a straight path toward the development of new commercial tobacco varieties with reduced TSNA levels. Concerning NNN, the other main carcinogenic TSNA along with NNK (IARC, 2004), a suitable solution for reducing its levels is already available through the Zyvert™ technology, which aims to reduce the levels of its precursor alkaloid nornicotine (Lusso et al., 2017). One could foresee TSNA reduction being maximized in both the tobacco matrix and tobacco smoke through the development of commercial tobacco varieties that combine the CLCNt2-deficiency approach with the Zyvert™ technology.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are publicly available. RNA-Seq data can be found here: National Center for Biotechnology Information (NCBI) BioProject database under accession number PRJEB46762.

**AUTHOR CONTRIBUTIONS**

PC, JL, AH, HL, JS, YM, HM, SG, RL, LB, and RD contributed to conception and design of the study. LB and RD organized the database and wrote the first draft of the manuscript. SK performed the statistical analysis. LB, RD, HL, and SK wrote sections of the manuscript. All authors contributed to the article and approved the submitted version.

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The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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