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Mutating alfalfa *COUMARATE 3-HYDROXYLASE* using multiplex CRISPR/Cas9 leads to reduced lignin deposition and improved forage quality

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Alfalfa (*Medicago sativa* L.) forage quality is adversely affected by lignin deposition in cell walls at advanced maturity stages. Reducing lignin content through RNA interference or antisense approaches has been shown to improve alfalfa forage quality and digestibility. We employed a multiplex CRISPR/Cas9-mediated gene-editing system to reduce lignin content and alter lignin composition in alfalfa by targeting the *COUMARATE 3-HYDROXYLASE* (*MsC3H*) gene, which encodes a key enzyme in lignin biosynthesis. Four guide RNAs (gRNAs) targeting the first exon of *MsC3H* were designed and clustered into a tRNA-gRNA polycistronic system and introduced into tetraploid alfalfa via *Agrobacterium*-mediated transformation. Out of 130 transgenic lines, at least 73 lines were confirmed to contain gene-editing events in one or more alleles of *MsC3H*. Fifty-five lines were selected for lignin content/composition analysis. Amongst these lines, three independent tetra-allelic homozygous lines (*Msc3h-013*, *Msc3h-121*, and *Msc3h-158*) with different mutation events in *MsC3H* were characterized in detail. Homozygous mutation of *MsC3H* in these three lines significantly reduced the lignin content and altered lignin composition in stems. Moreover, these lines had significantly lower levels of acid detergent fiber and neutral detergent fiber as well as higher levels of total digestible nutrients, relative feed values, and *in vitro* true dry matter digestibility. Taken together, these results showed that CRISPR/Cas9-mediated editing of *MsC3H* successfully reduced shoot lignin content, improved digestibility, and nutritional values without sacrificing plant growth and biomass yield. These lines could be used in alfalfa breeding programs to generate elite transgene-free alfalfa cultivars with reduced lignin and improved forage quality.

KEYWORDS

CRISPR/Cas9, gene editing, lignin composition, alfalfa, forage quality, *MsC3H*, nutritional value, digestibility

Introduction

Alfalfa (*Medicago sativa*, L) is a perennial legume forage crop widely cultivated in the world for livestock feed as it provides highly nutritious forage (Singer et al., 2018; Chen et al., 2020). Alfalfa is grown for hay, silage, dry pellets, or used directly for grazing as a quality source of proteins for livestock (Zhou et al., 2011; Barros et al., 2019b). The fast-growing perennial growth habit of alfalfa makes it ideal for multiple harvests within one growing season (Singer et al., 2018; Wolabu et al., 2023). The herbaceous nature of alfalfa plants with abundant lignocellulosic biomass yields and the ability to thrive in a variety of soil conditions make it a promising next-generation source of feedstock for biofuels (Hisano et al., 2009; Tong et al., 2015). Alfalfa leaves are more nutritious than stems, since high lignification in stems is a limiting factor to cell wall digestion, and any improvement that helps to liberate nutrients embedded in stems during digestion results in overall benefits for forage animals (Lacefield, 2004; Shadle et al., 2007; Getachew et al., 2018; Wolabu et al., 2023). Therefore, lignin is one of the most important limiting factors for forage nutritional quality (Getachew et al., 2018; Singer et al., 2018; Wolabu et al., 2023). Forage digestibility is negatively correlated with lignin content, which limits the nutritional value of forage crops (Li et al., 2008). Normally, lignin binds with structural carbohydrates (cellulose and hemicellulose) and cell wall proteins, thus reducing nutrient availability (Capstaff and Miller, 2018). Hence, lignin severely affects the digestibility of proteins and other nutritive components (Capstaff and Miller, 2018). Previous studies showed a strong negative correlation between lignin content and sugars released by enzymatic hydrolysis, implying that low lignin content correlates with high released carbohydrate levels (Chen and Dixon, 2007; Ziebell et al., 2010; Tong et al., 2015; Park et al., 2017).

Lignin is one of the most abundant biopolymers and a key structural component crucial for integrity of cell wall and stem strength in plant growth and development (Hisano et al., 2009; Liu et al., 2018; Dixon and Barros, 2019; Vanholme et al., 2019). In plants, lignin plays an important role in protecting plants from adverse environmental stresses, including mechanical damage, drought, low temperature, and resistance to various pests, serving as a mechanical barrier (Li et al., 2008; Bhuiyan et al., 2009; Gallego-Giraldo et al., 2014; Liu et al., 2018; Dixon and Barros, 2019; Vanholme et al., 2019). In addition, lignin functions as a conduit for the transport of water and nutrients through the plant vascular system (Liu et al., 2018; Dixon and Barros, 2019; Vanholme et al., 2019).

Lignin is a phenolic polymer synthesized through the phenylpropanoid pathway, which involves multiple enzymes and intermediates (Dixon et al., 2001; Vanholme et al., 2010; Barros et al., 2019a). The pathway begins with the conversion of the

aromatic amino acid phenylalanine to cinnamic acid through the action of the enzyme phenylalanine ammonia-lyase (PAL) (Dixon and Paiva, 1995). Cinnamic acid is then converted into three major monomers, also known as monolignols, namely *p*-coumaryl alcohol, coniferyl alcohol, and sinapyl alcohol, through the concerted action of cinnamate 4-hydroxylase (C4H), 4-coumarate CoA ligase (4CL), hydroxycinnamoyl-CoA shikimate/quinic acid hydroxycinnamoyltransferase (HCT), *p*-coumaroyl shikimate/quinic acid 3-hydroxylase (C3H), caffeoyl-CoA 3-O-methyltransferase (CCoAOMT), ferulate 5-hydroxylase (F5H), caffeic acid/5-hydroxyferulic acid O-methyltransferase (COMT), cinnamoyl-CoA reductase (CCR), and cinnamyl alcohol dehydrogenase (CAD) (Boerjan et al., 2003; Hisano et al., 2009; Vanholme et al., 2010). These monolignols are transported to plant cell walls where they are oxidized by apoplastic peroxidases and laccases to form lignin polymers in different patterns (Vanholme et al., 2010; Barros et al., 2019a).

Coumarate 3-hydroxylase (C3H) belongs to a large family of cytochromes P450 and is one of the key enzymes in lignin biosynthesis (Reddy et al., 2005; Ralph et al., 2006; Ziebell et al., 2010; Tong et al., 2015). C3H uses *p*-coumaroyl shikimate as a substrate to direct the synthesis of guaiacyl and syringyl monolignols (Hisano et al., 2009; Ziebell et al., 2010; Ralph et al., 2012; Tong et al., 2015). Given its central role in the synthesis of S and G lignin, C3H has been targeted to reduce lignin content and alter composition in multiple plant species. Downregulation of *C3H* at the RNA level in alfalfa leads to reduced lignin content and altered composition with improved digestibility (Reddy et al., 2005; Ralph et al., 2006; Ziebell et al., 2010; Tong et al., 2015; Getachew et al., 2018). Downregulation of *C3H* in poplar and maize also leads to reduced lignin content and altered composition with improved saccharification efficiency for biofuel and pulping industries (Ralph et al., 2012; Fornalé et al., 2015).

Genome editing approaches, such as CRISPR/Cas9, have revolutionized the field of plant biotechnology through genome engineering, facilitating important trait-driven improvement in major crop species (Chen et al., 2020; Wolabu et al., 2020a, 2020b; Bottero et al., 2021; Zheng et al., 2021; Bao et al., 2022; Kumar et al., 2022; Nerkar et al., 2022; Singer et al., 2022; Wolabu et al., 2023). CRISPR/Cas9 has been employed to manipulate lignin content in various plant species to improve the efficiency of lignocellulose processing and enhance forage quality, digestibility, and saccharification efficiency (Park et al., 2017; Jang et al., 2021; Afifi et al., 2022; Roy et al., 2023; Wolabu et al., 2023; Laksana et al., 2024). A recent study that altered flowering time via multiplex-CRISPR/Cas9 edition of a florigen in alfalfa generated promising lines with improved biomass yield and quality (Wolabu et al., 2023). Editing the switchgrass *4CL* gene has also been shown to generate mutant lines with reduced lignin content and improved sugar release (Park et al., 2017). Genome-editing has also been applied in wood species for lignin manipulation. By targeting *CAFFEYOYL SHIKIMATE ESTERASE* (*CSE*) and *PHBMT1* in poplar, Zhao et al. (2021) and Jang et al. (2021) demonstrated that precise editing of targeted genes via CRISPR/Cas9 can improve plant lignocellulosic biomass production with less lignin. Recently, Sulis et al. (2023) reported the impact of an assembled multiplex CRISPR/Cas9

Abbreviations: tRNA, transfer RNA; TDN Total Digestible Nutrients; gRNA, guide RNA; RFV, Relative Feed Value; PAM, Protospacer Adjacent Motif; SNP, Single Nucleotide Polymorphism; CP, Crude Protein; C3H, Coumarate 3-hydroxylase; G, Guaiacyl; S, Syringyl; H, *p*-hydroxyphenyl; ADF, Acid Detergent Fiber; NDF, Neutral Detergent Fiber; IVTDM In Vitro True Dry Matter Digestibility.

genome editing strategy by targeting six genes, namely *C3H*, *CCoAOMT*, *AldOMT*, *PAL*, *C4H*, and *CAD*, in poplar. They used different multigenic approaches to edit these genes to improve the chemical and physical properties of the woody plant and achieve optimized fiber production for the pulping industry. Edited lines exhibited varying degrees of loss-of-function mutations in the target genes with biallelic loss-of-function editing of all target genes (Sulis et al., 2023). Sugarcane lignin content reduction (up to 51% with high S/G ratio) was achieved using the CRISPR/Cas9 genome editing system by targeting SoLIM transcription factors that are involved in the lignin biosynthesis pathway (Laksana et al., 2024). In this regard, modern genome editing approaches offer great opportunities to reduce lignin content and/or to alter composition to improve product quality without compromising plant growth and resilience. This is one of the breeding goals to improve forage nutritional quality and fiber production (Park et al., 2017; Wolabu et al., 2020a; Nerkar et al., 2022; Roy et al., 2023; Sulis et al., 2023; Wolabu et al., 2023).

In this study, four specific target gRNAs were designed to knock out the alfalfa *C3H* (*MsC3H*) gene using a previously optimized multiplex CRISPR/Cas9 system to manipulate lignin content and composition for forage quality improvement. The generated tetra-allelic homozygous *Msc3h* mutant lines showed significantly reduced lignin content and a low level of ADF and NDF with no significant difference in biomass yield but with improved digestibility and nutritional values. These lines could be integrated into alfalfa breeding programs to generate elite transgene-free alfalfa cultivars with reduced lignin and improved forage quality.

Materials and methods

Plant materials and growth conditions

Alfalfa (*Medicago sativa* L.) cultivar, Regen-SY4D, was used for genome editing and genetic transformation. CRISPR/Cas9-edited lines and the non-edited control (empty vector-EV) were grown in greenhouse under conditions of 22°/19°C day/night temperature, 16/8 h day/night photoperiod, 150 $\mu\text{mol.m}^{-2}\text{s}^{-1}$ light intensity, and 70%-80% relative humidity. All alfalfa lines analyzed in this study were maintained and vegetatively propagated via stem cuttings.

Multiplex gRNAs-CRISPR/Cas9 vector construction and plant transformation

To permanently mutate the alfalfa *C3H* (*MsC3H*) gene, we used a previously optimized efficient multiplex gRNAs-CRISPR/Cas9 system (Wolabu et al., 2020a) to generate tetra-allelic mutants in the complex alfalfa genome. To examine any potential genotypic SNPs that might influence the genome-editing efficiency due to mismatch among four alleles of *MsC3H*, the genomic DNA fragment of *MsC3H* spanning the target region was amplified from the Regen-SY4D genomic DNA by PCR and cloned into the pGEM-Teasy vector (Promega, Madison, WI, USA). At least twenty clones were randomly picked for Sanger sequencing to determine the sequence identity and presence of any

SNPs in the conserved coding regions of four alfalfa alleles. Four *MsC3H* gRNAs, gRNA1-CCATACCCACTTCCCATCAT, gRNA2-ACAAAACCTCTCAAACATCTA, gRNA3-TGAGGTTT TTGATATTGGTG, and gRNA4-TGGGGTATCATGGAAGAAGC upstream of respective PAM, AGG, TGG, AGG and TGG sites, were designed to target the first exon of the *MsC3H* gene using the web-based tool CRISPR-P (<http://cbi.hzau.edu.cn/cgi-bin/CRISPR>) (Lei et al., 2014). All designed gRNAs (spacers) were inserted between tRNA and gRNA scaffolds to be clustered in a tandem manner using the Golden Gate assembly method (Engler et al., 2008). The pGTR plasmid, which contains the tRNA-gRNA fragment, was used as a template to synthesize the polycistronic tRNA-gRNA (PTG) for construction of the multiplex tRNA-*MsC3H* spacer-gRNA (Xie et al., 2015; Wolabu et al., 2020a). First, the overlapping PCR products were purified using the Spin Column PCR Product Purification Kit (Promega, Madison, WI, USA) following the manufacturer's instruction (www.promega.com). The PCR products containing four *MsC3H*-spacers were then inserted into the optimized vector with the AtU6-tRNA-*MsC3H*-gRNAs-AtUBQ10-Cas9-pRGE31-bar backbone by digestion and ligation using Fok I (NEB) and BsaI (Wolabu et al., 2020a). Insertion of all multiplexed gRNAs into the CRISPR/Cas9 module was verified by DNA sequencing. The resulting binary vector, *MsC3H*-gRNA-CRISPR/Cas9, was transformed into the *Agrobacterium tumefaciens* strain, EHA105. *Agrobacterium*-mediated transformation of alfalfa was carried out following the protocol described earlier by Wolabu et al. (2020a) to generate CRISPR/Cas9-edited alfalfa *Msc3h* mutants.

Genotyping of CRISPR/Cas9-edited *Msc3h* lines

To identify and select CRISPR/Cas9-edited *Msc3h* lines, multiple rigorous genotyping steps were applied. First, the transgenic lines were screened by PCR for presence or absence of the *BAR* gene using *BAR* gene specific primers (PPT-F + PPT-R). The *BAR* gene-positive transgenic lines were subjected to a second PCR amplification of the target region spanning the designed gRNAs with *MsC3H*-specific primers, followed by Sanger sequencing. The amplified PCR products were purified after treating with enzymes Antarctic phosphatase and exonuclease I (Wolabu et al., 2020a) and sequenced using the Sanger sequencing method to determine the mutation signature (double peak in the electropherogram upstream of PAM) in at least one of the gRNAs (gRNA1, gRNA2, gRNA3, and gRNA4). Then, transgenic lines considered as putative heterozygous and/or chimeric were selected and subjected to phenotypic analysis for desirable traits. Furthermore, promising lines with desirable phenotypes were characterized for homozygosity by cloning the target regions spanning the gRNAs (gRNA1, gRNA2, gRNA3, and gRNA4) into the pGEM-T Easy vector. Twenty colonies were randomly picked for each mutant line, and each plasmid DNA was sequenced using the Sanger sequencing method to determine the nature of mutation events in each *Msc3h* line (homozygous, heterozygous, or chimeric mutation) by aligning the reads with the reference sequence using the SeqMan Pro 15.0.1 (DNASTAR software for life scientists)

(<https://www.dnastar.com/quote-request/>). All primer pairs used to genotype the plasmids and the transgenic lines in this study are listed in [Supplementary Table S2](#).

Tissue sampling and lignin composition analysis

For lignin analysis, stem samples were harvested at the early flower bud growth stage (the stage when 10% of the plants showed first flower buds). Pooled stem sections between node 4 and 7 from the bottom were collected and dried at a fixed temperature of $55^{\circ} \pm 5^{\circ}\text{C}$ for 10 days followed by grinding into powder using a Wiley mill. The ground samples were extracted with chloroform/methanol (2:1, v/v), methanol, and water. Lignin composition was determined using a modified thioacidolysis method as described previously ([Chen et al., 2021](#)), with 4,4'-ethylidene bisphenols as the internal standard. The total lignin contents are represented by the sum of total thioacidolysis monomer yields. Three technical replicates were used in the first lignin analysis of independent transgenic lines on a single plant basis (a total of 55 *Msc3h* lines). Based on the first record of lignin analysis and phenotyping of desirable agronomic traits, three promising candidate lines, *Msc3h-013*, *Msc3h-121*, and *Msc3h-158*, were selected for further analysis of lignin composition using six biological replicates.

Determination of forage nutritional quality

To determine the forage biomass yield and nutritional quality (digestibility and nutritional values) of three selected *Msc3h* lines, above-ground plant tissues (stems and leaves) were harvested when 10% of plants showed their first flower buds. Forage nutritional quality assessment was carried out at the Ag Services and Resources Core Facility (ASRC), Noble Research Institute, LLC. Sample grinding and processing were performed following procedures described by [Gou et al. \(2018\)](#). Forage nutritional quality was determined with the near infrared reflectance spectroscopy (NIRS) system using a Foss NIRS 6500 monochromator with a scanning range of 1100–2500 nm (Foss NIR System). The following forage nutritional quality parameters were measured: acid detergent fiber (ADF), neutral detergent fiber (NDF) in percentage of dry matter basis, total digestible nutrients (TDN), relative feed value (RFV), *in vitro* dry matter digestibility (IVTDMD), crude protein (CP) content, minerals including magnesium (Mg), potassium (K), calcium (Ca), and phosphorus (P) content.

Statistical analysis

At least six plants for each independent line were used as biological replicates for statistical analyses. Data shown in graph bars represent the mean $\pm$ SD. The analyses were performed using the Microsoft Excel software and the asterisks indicate significant differences based on Student's *t*-test (** $p < 0.001$, * $p < 0.01$, * $p < 0.05$). The relative advantage (%) of the tested lines over the

control (EV) was calculated as the performance of the tested line minus the control divided by the control and multiplied by 100 [(Line tested – EV)/EV $\times$ 100 = %].

Results

Generation, screening, and analysis of CRISPR/Cas9-mediated *Msc3h* mutant lines

In this study, we mutated the *Msc3H* gene in alfalfa by employing a multiplex CRISPR/Cas9 system ([Wolabu et al., 2020a](#)). Four gRNAs were designed in the first exon of *Msc3H* at 111 bp (gRNA1), 156 bp (gRNA2), 465 bp (gRNA3), and 572 bp (gRNA4) downstream of the translation start codon, segments containing no SNPs across all four alleles of *Msc3H*, using CRISPR-P 2.0 (<http://cbi.hzau.edu.cn/cgi-bin/CRISPR>) ([Lei et al., 2014](#)) ([Figure 1A, C](#)). One hundred and thirty *BAR* (selection marker gene) positive lines were generated via *Agrobacterium*-mediated transformation. To identify CRISPR/Cas9-edited mutant lines, the target region spanning all four gRNAs were amplified from all 130 lines using a pair of primers flanking the target region ([Supplementary Table S2](#)). Of the 130 transgenic lines, 73 lines were identified as putative *Msc3h* mutant lines with at least one mutation signature (double peaks in sequencing chromatograph) at one of the target gRNA sites by Sanger sequencing of the PCR amplicons ([Figure 1B](#)). Of the 73 putative candidates, 55 lines were selected for lignin analysis based on phenotypic evaluation of desirable agronomic traits. Based on the results of lignin analysis, 12 candidate lines with reduced lignin content and altered composition were propagated for further characterization and statistical validation. Finally, three *Msc3h* mutant lines (*Msc3h-013*, *Msc3h-121*, and *Msc3h-158*) that exhibited similar growth patterns as control, empty vector (EV) plants ([Figure 2F](#)) were selected for analysis of the nature of mutation events at the target sites prior to a deeper analysis of lignin content and composition ([Figure 3](#)). The results showed that mutant line *Msc3h-013* had an 8-bp deletion and a 1-bp “T” insertion in Allele 2 (A2) and Allele 3 (A3), respectively, at gRNA1, and 21-bp deletions in A1 and A3 upstream of the PAM of gRNA2 ([Figure 3](#)). Moreover, a deletion of 63 bp between gRNA1 and gRNA2 in A4 had occurred, removing the entire sequence between gRNA1 and gRNA2 ([Figure 3](#)). Similarly, 1–4 bp deletions and 1-bp insertion upstream of PAM at gRNA3 were detected in four different alleles of *Msc3h-013* ([Figure 3](#)). Consequently, the mutation events at gRNA3 of *Msc3h-013* were sufficient to cause 100% tetra-allelic homozygous mutation and knock out the gene at a single site. Overall, the combined multiple mutation events occurred at gRNA1, gRNA2, and gRNA3 made line *Msc3h-013* a 100% tetra-allelic homozygous mutant ([Figure 3, Supplementary Figure S1A](#)). Similarly, analysis of mutant line *Msc3h-121* showed 8- and 9-bp deletions in A2 and A4, respectively, along with a 1-bp “C to T” substitution in A1 at gRNA1 ([Figure 3](#)). *Msc3h-121* also showed 21-bp deletions (in A2 and A4) at gRNA2, 4-bp deletions (in A1 and A2) and a 1-bp “T” insertion (A3 and A4) at gRNA3

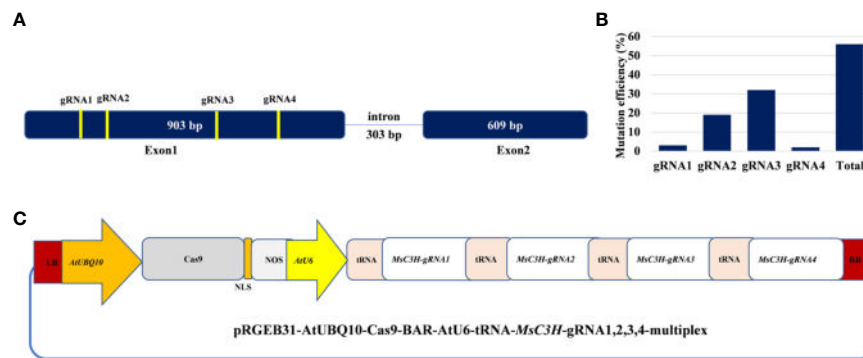


FIGURE 1

Schematic illustration of alfalfa *COUMARATE 3-HYDROXYLASE* (*Msc3H*) gene structure with designed multiplex gRNAs-CRISPR/Cas9 vector and genome editing efficiency in alfalfa. (A) *Msc3H* gene structure and four guide RNA sites with specific sequences in the coding region. (B) Mutation efficiency (%) of four guide RNAs at different target sites of *Msc3H*. (C) Illustration of the multiplex construct of *Msc3H*-gRNA1,2,3,4-CRISPR/Cas9 vector.

(Figure 3), making it a 100% tetra-allelic homozygous mutant. Therefore, the combined overall mutation events detected in *Msc3h-121* at three specific targets (gRNA1-3) led to a 100% tetra-allelic homozygous mutant (Figure 3, Supplementary Figure S1B). Diverse mutation events were also detected at gRNA1, gRNA2 and gRNA3 of the mutant line *Msc3h-158* (Figure 3, Supplementary Figure S1C). Mutation analysis indicated that 5-bp and 7-bp deletions (in A3 and A4) occurred at gRNA1 followed by 15-bp deletions (A1 and A2) and a 5-bp deletion (A3) at gRNA2 (Figure 3). At gRNA3, five different types of mutation events occurred across all four alleles of the *Msc3H* gene, including 1-bp deletion in A1, 2-bp deletion in A2, 3-bp deletion in A3, and 4-bp deletion in A4 (Figure 3, Supplementary Figure S1C), along with a 4-bp deletion and a 1-bp insertion in one of the alleles, indicating the chimeric nature of mutation events in *Msc3h-158*. Interestingly, the chimeric mutation also showed 100% homozygosity and the combined mutation events detected at gRNA1, gRNA2, and gRNA3

similarly produced 100% tetra-allelic homozygosity in the mutant line *Msc3h-158* (Figure 3, Supplementary Figure S1C). In summary, mutation event analysis indicated diverse mutation efficiencies and mutation events in gRNA1, gRNA2, and gRNA3 with complete mutations in all four alleles (copies) in the three selected mutant lines, *Msc3h-013*, *Msc3h-121*, and *Msc3h-158* (Figure 3, Supplementary Figure S1A-C). The overall mutation efficiency at gRNA3 was higher than that of any other gRNAs. The mutagenesis efficiency at gRNA3 was 32%, followed by 19% at gRNA2 and 4% at gRNA1, while gRNA4 had the lowest mutagenesis efficiency with only 2% (Figure 1B), which might be the reason why we did not detect mutations at gRNA4 in the selected three lines *Msc3h-013*, *Msc3h-121*, and *Msc3h-158* (Figure 3). All mutation events in these three lines occurred in the framework of designed gRNA1, 2, and 3 at 1-3 bp upstream of the PAM regions with a dominance of small deletions (1-9 bp). These results further highlight the usefulness of multiplex CRISPR/Cas9 system in maximizing the chances of

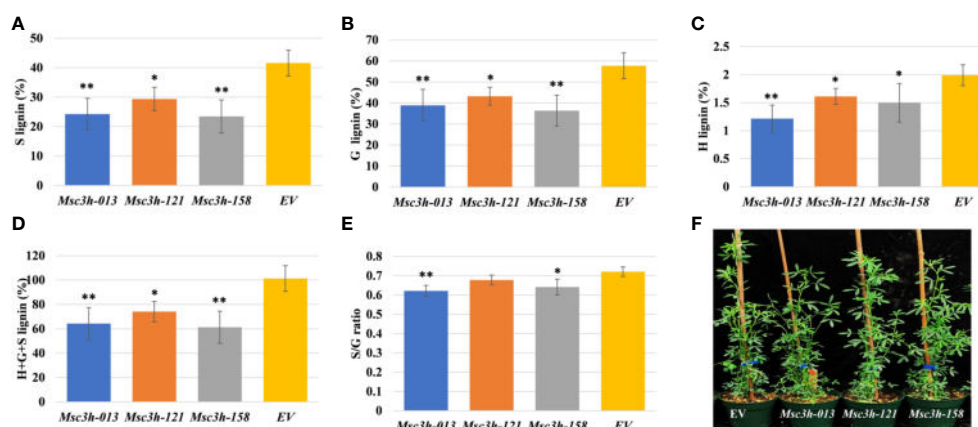


FIGURE 2

Analysis of lignin content and composition in stem tissues of three selected *Msc3h* mutant lines *Msc3h-013*, *Msc3h-121* and *Msc3h-158*. (A) S lignin content (%) in stem tissues of *Msc3h* lines and EV. (B) G lignin content (%) in stem tissues of *Msc3h* lines and EV. (C) H lignin content (%) in stem tissues of *Msc3h* lines and EV. (D) Total (H+G+S) lignin content (%) in stem tissues of *Msc3h* lines and EV. (E) S/G ratio in stem tissues of *Msc3h* lines and EV. (F) Phenotype of mutant lines *Msc3h-013*, *Msc3h-121*, and *Msc3h-158* with EV at early flower bud vegetative growth stage. Data represent mean values ($\pm$ SD; n = 6). Statistics was analyzed using Student's t-test (* p < 0.05; ** p < 0.01).

Genotype	Type of mutation events detected at target sites of <i>Msc3h</i> mutant lines				Overall mutation status
	gRNA1	gRNA2	gRNA3	gRNA4	
WT	CCATACCACTTCCCATCAT ⁴⁵⁶	ACAAACTCTCAACATCTA ⁷⁵⁶	TGAGGTTTTGATATGTCG ⁶⁵⁶	TGGGGTATCATGGGAAGC ⁷⁵⁶	Non-edited wildtype sequence
<i>Msc3h-013</i>	A1-CCATACCACTTCCCATCAT ⁴⁵⁶ A2-CCATACCCAC-----A3 ⁴⁵⁶ A3-CCATACCACTTCCCATCAT ⁴⁵⁶ A4-CCATACCACTTCCCATC-----TA ⁴⁵⁶	G-----21bp-----CTA ⁷⁵⁶ ACAAACTCTCAACATCTA ⁷⁵⁶ G-----21bp-----CTA ⁷⁵⁶	TGAGGTTTTGAT-----GTG ⁶⁵⁶ TGAGGTTTTGATATT-----GTG ⁶⁵⁶ TGAGGTTTTGATATTGTG ⁶⁵⁶ TGAGGTTTTGATATTGTG ⁶⁵⁶	TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶	100% tetra-allelic homozygous due to the combination of 1-45 bp deletions and 1 bp insertion at gRNA1, gRNA2 and gRNA3.
<i>Msc3h-121</i>	A1-CCATACCACTTCCCATCAT ⁴⁵⁶ A2-CCATACCA-----CAT ⁴⁵⁶ A3-CCATACCACTTCCCATCAT ⁴⁵⁶ A4-CCATACCA-----T ⁴⁵⁶	ACAAACTCTCAACATCTA ⁷⁵⁶ G-----21bp-----CTA ⁷⁵⁶ ACAAACTCTCAACATCTA ⁷⁵⁶ G-----21bp-----CTA ⁷⁵⁶	TGAGGTTTTGAT-----GTG ⁶⁵⁶ TGAGGTTTTGAT-----GTG ⁶⁵⁶ TGAGGTTTTGATATTGTG ⁶⁵⁶ TGAGGTTTTGATATTGTG ⁶⁵⁶	TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶	100% tetra-allelic homozygous due to the combination of 4-21 bp deletions, 1 bp insertion, 1 bp substitution at gRNA1, gRNA2 and gRNA3.
<i>Msc3h-158</i>	A1-CCATACCACTTCCCATCAT ⁴⁵⁶ A2-CCATACCA-----CAT ⁴⁵⁶ A3-CCATACCA-----CAT ⁴⁵⁶ A4-CCATACCACT-----TCAT ⁴⁵⁶	AC-----21bp-----CTA ⁷⁵⁶ AC-----21bp-----CTA ⁷⁵⁶ ACAAACTCTCAACATCTA ⁷⁵⁶ ACAAACTCTCAACATCTA ⁷⁵⁶	TGAGGTTTTGATATT-----GTG ⁶⁵⁶ TGAGGTTTTGATATT-----GTG ⁶⁵⁶ TGAGGTTTTGATATT-----GTG ⁶⁵⁶ TGAGGTTTTGATATT-----GTG ⁶⁵⁶	TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶ TGGGGTATCATGGGAAGC ⁷⁵⁶	100% tetra-allelic homozygous due to the combination of 1-9 bp deletions and 4 bp substitutions at gRNA1, gRNA2 and gRNA3. Chimeric mutation detected at gRNA3.

FIGURE 3
Molecular analysis of *Msc3H* mutation events (deletions/insertion/substitution) generated by multiplex gRNAs-CRISPR/Cas9 gene editing at four target sites of *Msc3H*. Mutation events (deletions, insertions, substitutions) of mutant lines *Msc3h-013*, *Msc3h-121*, and *Msc3h-158* occurred at each specific target site (gRNA1, gRNA2, gRNA3 and gRNA4) compared to wild type (WT). Deletions are indicated by red dashed lines; insertions or substitutions are indicated by red letters. PAMs are indicated by red underlined italicized letters. The four allelic copies are designated as red allele-1 (A1), allele-2 (A2), allele-3 (A3), and allele-4 (A4).

obtaining knockout or edited mutants in the target genes through a single transformation event.

Analysis of lignin content and composition in *Msc3h* mutant lines

To determine lignin content and composition (syringyl (S), guaiacyl (G), *p*-hydroxyl (H) units and total lignin) in all *Msc3h* mutants, we used stem samples consisting of internodes number 4, 5 and 6 counting from the bottom. Lignin deposition was determined in two phases by using the thioacidolysis method (Chen et al., 2021) for independent *Msc3h* mutant lines. In the first phase, 55 putative *Msc3h* mutant lines with various mutation events were analyzed to determine lignin content and composition compared to the non-edited EV control. The results showed a wide range of lignin composition, including the individual content of S, G, and H lignin units and the total lignin content. After performing lignin analysis on the 55 *Msc3h* lines, we selected 12 lines for vegetative propagation and further assessment of lignin content and composition and nutritional values (Supplementary Figure S2A). These twelve lines showed different agronomic traits with two of the lines having reduced growth while the others showed comparable plant height to the control (Supplementary Figure S2A). Finally, three of the 12 propagated lines (*Msc3h-013*, *Msc3h-121* and *Msc3h-158*), which were phenotypically comparable to EV and were tetra-allelic homozygous mutants, were selected and subjected to a second round of lignin content and composition analysis. Results showed that all three *Msc3h* lines had significantly reduced levels of S, G, H monomer yields, and total lignin compared to the EV control (Figure 2A-D). Remarkably, in the line *Msc3h-013*, the S, G, H monomer yields, and the total lignin content were reduced by 42%, 33%, 39%, and 36%, respectively, compared to EV (Figure 2A-D, Supplementary Table S1). Similarly, line *Msc3h-158* showed a reduction of 44%, 37%, 25% and 39% in S, G, H, and total lignin content, respectively, compared to the control. Line *Msc3h-121* showed a reduction of 29%, 25%, 20% and 27% in S, G, H, and total lignin content, respectively, compared to EV (Figure 2A-D, Supplementary Table S1). All three *Msc3h* mutant lines also

showed 14%, 7% and 11% relative advantage of S/G ratio for *Msc3h-013*, *Msc3h-121* and *Msc3h-158*, respectively, over EV (Figure 2E, Supplementary Table S1). In conclusion, disruption of *Msc3H* substantially reduced key lignin components in selected lines compared to the control (Figure 2A-E, Supplementary Table S1).

Analysis of forage digestibility and nutritional quality in *Msc3h* mutant lines

Given the significant reduction in lignin content in the selected *Msc3h* lines (*Msc3h-013*, *Msc3h-121*, and *Msc3h-158*), we evaluated the nutritional parameters of forage quality (ADF, NDF, TDN, RVE, IVTDMD, CP and macro-minerals) in whole shoot tissues of these lines at flower bud growth stage (Figure 4A-F, 5A-D). The results showed that all three mutant lines had significantly lower levels of ADF with a relative advantage of 7-11% over EV (Figure 4A, Supplementary Table S1). Likewise, *Msc3h-013* and *Msc3h-158* also showed significantly reduced level of NDF with 9-13% relative advantage over EV (Figure 4B, Supplementary Table S1). Although not statistically significant, the NDF content in *Msc3h-121* was lower than that in EV with a relative advantage of 8%. In a similar trend, the nutritional values (specifically, TDN and RFV) in the three lines were significantly higher than that in EV (with 9-13% and 14-23% relative advantage over the EV for TDN and RFV, respectively) (Figure 4C, D, Supplementary Table S1). Highly significant (*p* < 0.001) elevation of IVTDMD was also recorded in mutant lines *Msc3h-013* and *Msc3h-158* with a relative advantage of 8% and 10% over EV, respectively (Figure 4E, Supplementary Table S1), followed by *Msc3h-121* with a relative advantage of 5% over EV, which was not statistically significant (Figure 4E, Supplementary Table S1). We also analyzed nutritional parameters such as crude protein (CP) and mineral contents in the three *Msc3h* mutant lines. Interestingly, *Msc3h-013* showed a significantly higher CP content with a relative advantage of 30% over EV (Figure 4F). Similarly, *Msc3h-013* showed a significantly high level of P, Ca and Mg content with a relative advantage of 15%, 11% and 27%, respectively, over EV (Figure 5A, B, D). Although not statistically

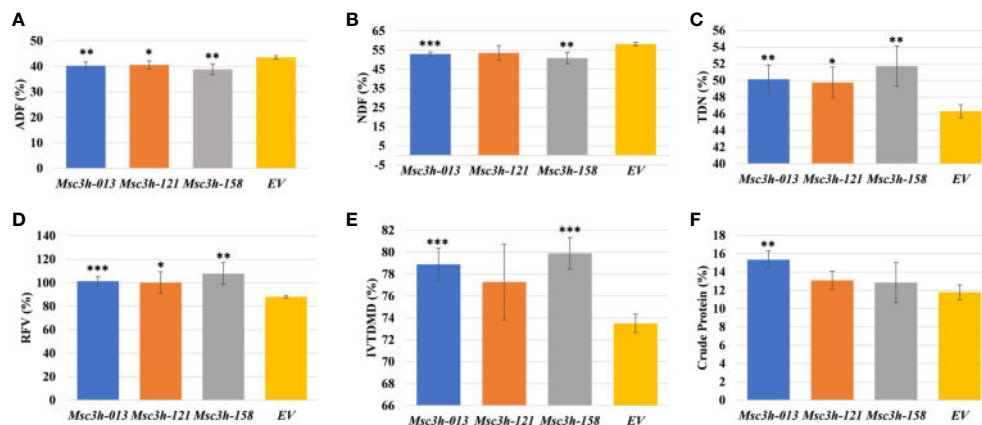


FIGURE 4

Analysis of forage quality in aboveground (leaves and stem) tissues of *Msc3h* mutant lines and EV. (A) Acid Detergent Fiber (ADF) content (%) in *Msc3h* lines and EV. (B) Neutral Detergent Fiber (NDF) content (%) in *Msc3h* lines and EV. (C) Total Digestible Nutrients (TDN) content (%) in *Msc3h* lines and EV. (D) Relative Feed Value (RFV) (%) in *Msc3h* lines and EV. (E) *In Vitro* True Dry Matter Digestibility (IVTDM) (%) in *Msc3h* lines and EV. (F) Crude protein content (%) in *Msc3h* lines and EV. Data represent mean values ($\pm$ SD; $n = 6$) and were analyzed statistically using Student's *t*-test (* $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$).

significant, *Msc3h-121* and *Msc3h-158* also showed high levels of CP and four microminerals (P, Ca, K, and Mg) compared to EV (Figure 4F, 5A–D). Overall, the mutant line *Msc3h-013* had the lowest lignin content, which was reflected by higher digestibility and nutritional values, followed by *Msc3h-158*, which ranked second based on lignin analysis (Figure 2A–E, 4A–E). In conclusion, these results indicate that the multiplex CRISPR/Cas9-mediated gene editing system generated *Msc3h* mutant lines with significantly reduced lignin content and improved digestibility and nutritional quality.

Furthermore, we analyzed the correlation between lignin content/composition (S, G, H, and total lignin) and forage quality parameters, including digestibility (ADF and NDF) and nutritional

values (TDN, RFV, IVTDM and CP) to evaluate the importance of each variable (Table 1 and Supplementary Figure S3A–H, Figure S4A–H, Figure S5A–H). Specifically, S lignin and total lignin showed a positive correlation with ADF with $R^2 = 0.52$ and 0.50 , respectively (Supplementary Figure S3A, D, Table 1), while S, G, and total lignin were positively correlated with NDF with $R^2 = 0.65$, 0.62 , 0.64 , respectively (Supplementary Figure S3E, F, H, Table 1). However, the correlation between H lignin and ADF and NDF was consistently weaker than other lignin parameters (Supplementary Figure S3C, G, Table 1). On the other hand, S lignin was negatively correlated with TDN ($R^2 = 0.52$), RFV ($R^2 = 0.62$), IVTDM ($R^2 = 0.75$) and CP ($R^2 = 0.62$) (Supplementary Figure S4A, E, S5A, E, Table 1). Similarly, G lignin and total lignin showed a moderate

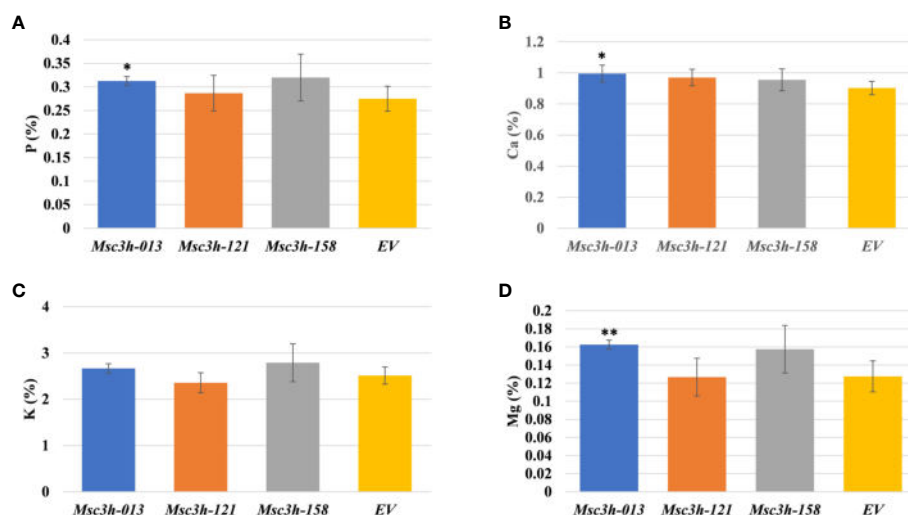


FIGURE 5

Analysis of forage macro-minerals contents in aboveground (leaves and stem) tissues of mutant lines *Msc3h-013*, *Msc3h-121* and *Msc3h-158* and EV. (A) Phosphorus (P) content (%). (B) Calcium (Ca) content (%). (C) Potassium (K) content (%). (D) Magnesium (Mg) content (%). Data represent mean values ($\pm$ SD; $n = 6$) and were analyzed statistically using Student's *t*-test (* $p < 0.05$; ** $p < 0.01$).

TABLE 1 Correlation analyses of lignin composition vs forage quality traits in alfalfa *Msc3h* lines.

Correlation	(R ²)	p-value	Status	Correlation (%)	(R ²)	p-value	Status
ADF x S-lignin	0.52	0.01	Positive	TDN x S-lignin	0.52	0.01	Negative
ADF x G-lignin	0.47	0.01	Positive	TDN x G-lignin	0.47	0.01	Negative
ADF x H-lignin	0.28	0.07	Positive	TDN x H-lignin	0.28	0.07	Negative
ADF x Total-lignin	0.50	0.01	Positive	TDN x Total-lignin	0.50	0.001	Negative
NDF x S-lignin	0.65	0.001	Positive	RFV x S-lignin	0.62	0.02	Negative
NDF x G-lignin	0.62	0.001	Positive	RFV x G-lignin	0.58	0.04	Negative
NDF x H-lignin	0.40	0.05	Positive	RFV x H-lignin	0.36	0.05	Negative
NDF x Total-lignin	0.64	0.001	Positive	RFV x Total-lignin	0.61	0.002	Negative
IVTDMD x S-lignin	0.75	0.001	Negative	CP x S-lignin	0.65	0.001	Negative
IVTDMD x G-lignin	0.76	0.001	Negative	CP x G-lignin	0.60	0.002	Negative
IVTDMD x H-lignin	0.56	0.004	Negative	CP x H-lignin	0.64	0.001	Negative
IVTDMD x Total-lignin	0.76	0.001	Negative	CP x Total-lignin	0.59	0.003	Negative

negative correlation with TDN, RFV, IVTDMD and CP (Supplementary Figure S4B, D, F, H, S5B, D, F, H, Table 1). Results of correlation analysis suggest that lignin content and composition directly alter forage nutrition quality. In summary, we generated *Msc3h* mutant lines with significantly reduced lignin content and improved digestibility and nutritional quality using the multiplex CRISPR/Cas9-mediated gene editing system.

Discussion

Forage digestibility is negatively affected by lignin deposition in the feedstock (Baucher et al., 1999; Reddy et al., 2005). A number of studies targeting genes involved in lignin biosynthesis, such as *COMT*, *CAD*, *C4H*, *C3H*, *HCT*, *F5H*, and *CCoAMT*, via RNA interference or antisense approaches have shown the value of reducing lignin content in increasing forage digestibility of alfalfa plants (Marita et al., 2003; Reddy et al., 2005; Ralph et al., 2006; Shadle et al., 2007; Ziebell et al., 2010; Tong et al., 2015; Getachew et al., 2018; Singer et al., 2018; Barros et al., 2019b). However, transgenic approaches such as RNA interference, antisense, or artificial microRNA, which all manipulate gene functions at the RNA level and depend on sustained expression of transgenes in each generation, may suffer from transgene instability or silencing under field conditions, making these approaches less predictable (Jung and Altpeter, 2016). In this study, with the objective of overcoming these issues, we employed a multiplex CRISPR/Cas9 gene-editing approach to mutate the alfalfa *C3H* gene (*Msc3H*) at the DNA level and analyze the impact of altering lignin content and composition on plant growth and forage digestibility. In three *Msc3h* mutant lines, namely *Msc3h-013*, *Msc3h-121* and *Msc3h-158*, small Indels just upstream of the PAM regions of respective gRNAs were the most frequent mutations (Figure 3, Supplementary Figure S1A-C). Furthermore, large fragment deletions of 21-63 bp between gRNA1 and gRNA2 also occurred in the mutant line *Msc3h-158*,

with 100% tetra-allelic homozygosity (Figure 3, Supplementary Figure S1C). Overall, analysis of editing events showed diverse mutation types in each specific target region with the highest mutagenesis efficiency in gRNA3 (Figure 1B). Consistent with these findings, similar mutation events with small Indels and some large, truncated fragments were observed in previous studies of alfalfa genome editing system targeting *MsPDS*, *MsPALM1*, *MsSGR*, *MsSPL8*, and *MsFTa1* genes (Chen et al., 2020; Wolabu et al., 2020a; Singer et al., 2022; Wolabu et al., 2023).

Remarkably, *Msc3h* mutant lines identified as tetra-allelic homozygous showed expected reduced lignin content and altered lignin composition in stem samples (Figure 2A-D). Interestingly, the results of thioacidolysis analysis showed wide ranges of lignin reduction in *Msc3h-013* and *Msc3h-158* followed by *Msc3h-121* with S lignin content lower than G lignin content (Figure 2A, B). Our results also showed a greatly reduced H lignin content, unlike previous reports on downregulation of the *C3H* gene using RNAi/antisense approaches (Chen and Dixon, 2007). The discrepancy of the H lignin contents between our study and the previous study could be attributed to the partial loss of function in the antisense/RNAi system and the complete loss of function in the gene-editing system. Overall, *Msc3h* mutant lines had significantly reduced lignin content compared to the control (Figure 2A-D, Supplementary Table S1). Furthermore, the three *Msc3h* mutant lines also exhibited reduced S/G ratio compared to EV plants. Previously, Reddy et al. (2005) reported the possibility of producing phenotypically normal alfalfa plants with strong downregulation in C3H activity and reduced lignification, which is in line with our *Msc3h* knockout results. The authors also suggested that there is no relationship between S/G ratios and digestibility, whereas the total lignin content is highly correlated with digestibility (Reddy et al., 2005). In summary, reduced components of lignin composition (monomeric units S, G, and H) were achieved in all three selected *Msc3h* mutant lines when compared to the EV control (Figure 2A-E, Supplementary Table S1).

The aim of this study was to reduce lignin content in alfalfa to improve forage digestibility and nutritional values by mutating the *MsC3H* gene. Small change in forage digestibility significantly influences animal performance efficiency (Shadle et al., 2007). The assessment of forage nutritional quality in the three *Msc3h* mutant lines (*Msc3h-013*, *Msc3h-121* and *Msc3h-158*) showed a significant impact of reducing lignin content in improving standard forage quality parameters (ADF, NDF, TDN, RVE, IVTDM, CP and macro-minerals) (Figure 4A–F, 5A–D). The three promising mutant lines consistently had lower levels of ADF and NDF and higher levels of TDN, RVE and IVTDM compared to EV (Figure 4A–E). This is consistent with previous reports that decrease in ADF and NDF is associated with decrease in lignin content (Shadle et al., 2007; Getachew et al., 2018; Wolabu et al., 2023). Crude protein and mineral content were also significantly increased, thus enhancing forage quality. Among the three lines, *Msc3h-013* showed significantly higher crude protein (CP) content with a relative advantage of 30% over EV (Figure 4F) and significantly higher levels of Mg (27%), P (15%) and Ca (11%) relative to EV (Figure 5A, B, D). Overall, the mutant line *Msc3h-013* exhibited the lowest lignin content with superior improvement in digestibility and nutritional values, but slightly lower dry matter yield (Supplementary Figure S2B). *Msc3h-121* and *Msc3h-158* showed desirable agronomic traits comparable to the control (EV) without a significant effect on dry matter (Supplementary Figure S2B). Reduced biomass yield has been reported when *C3H* expression is strongly downregulated in transgenic alfalfa lines (Chen and Dixon, 2007). Furthermore, Reddy et al. (2005) targeted three specific cytochrome P450 enzymes involved in lignin pathway and generated transgenic lines with different lignin content and composition in alfalfa, suggesting the importance of fine-tuning the level of *C3H* or *C4H* expression to eliminate the possibility of reduced growth. In a similar manner, Chen and Dixon (2007) used alfalfa transgenic lines expressing antisense constructs for downregulation of six genes (including *MsC3H*) to determine the relationship between lignin content/composition and chemical/enzymatic saccharification. However, our study targeted a single gene (*MsC3H*) for forage quality improvement. The difference in the results obtained in this study and those previously reported in relation to the content and composition of lignin and agronomic parameters of nutritional quality could be attributed to the technology used, the germplasm, environmental conditions, and/or developmental factors that influence the complex biosynthesis of monolignols. However, as a trade-off for the reduction in dry matter, *Msc3h-013* showed superior performance in digestibility that substantially benefits nutritional values, CP, and mineral contents. This line could be used as a valuable genetic resource in alfalfa breeding programs. Decreases in ADF and NDF are associated with decreases in lignin components (Shadle et al., 2007; Getachew et al., 2018; Wolabu et al., 2023). Recently, we employed the CRISPR/Cas9 system to target the *MsFTa1* gene in alfalfa to delay flowering and found that the resulting late flowering *Msfta1* mutant lines produced not only greater biomass, as expected, but also had reduced lignin and fiber content (Wolabu et al., 2023). Such findings suggest that it is

possible to design efficient gRNAs to knock out multiple genes with combined developmental traits, such as delayed flowering, delayed senescence, and reduced lignin traits to improve biomass yield and nutritional quality in alfalfa (Wolabu et al., 2020a, 2023).

In conclusion, we generated *Msc3h* mutant lines with significantly reduced lignin and enhanced forage digestibility and nutritional quality using the multiplex CRISPR/Cas9-mediated gene editing system. These promising *Msc3h* mutant lines could be useful genetic resources in alfalfa breeding programs for the development of transgene-free mutant lines and germplasm enrichment.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

Author contributions

TW: Formal Analysis, Investigation, Writing – original draft. KM: Formal Analysis, Writing – original draft. FC: Investigation, Writing – review & editing. IT: Writing – review & editing, Data curation. MU: Writing – review & editing, Conceptualization. MT: Writing – review & editing. LC: Writing – review & editing, Investigation. ZW: Conceptualization, Writing – review & editing. JW: Data curation, Project administration, Supervision, Writing – review & editing.

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Conflict of interest

The 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.

The author(s) declared that they were an editorial board member of Frontiers, at the time of submission. This had no impact on the peer review process and the final decision.

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Supplementary material

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fpls.2024.1363182/full#supplementary-material>

SUPPLEMENTARY FIGURE 1

Molecular analysis of mutation events in lines *Msc3h-013*, *Msc3h-121*, and *Msc3h-158*. (A) Mutation events occurred in *Msc3h-013* at gRNA3 with one to four nucleotide deletions and one nucleotide insertion. (B) Mutation events occurred in *Msc3h-121* at gRNA3 with one to four nucleotide deletions and one nucleotide insertion. (C) Mutation events occurred in *Msc3h-158* at gRNA3 with one to four nucleotide deletions and one nucleotide substitution. All three promising *Msc3h* lines showed tetra-allelic homozygosity at gRNA3, generating 100% tetra-allelic homozygous mutations in *Msc3H*. Red arrows show PAM (protospacer adjacent motif) sequences.

SUPPLEMENTARY FIGURE 2

(A) Phenotype of twelve selected *Msc3h* mutant lines at flower bud vegetative stages showing different agronomic traits as compared to EV. Tetra-allelic homozygous lines with reduced lignin content and composition were propagated for replicated statistical analysis. (B) Analysis of dry weight (g) of three selected promising candidate *Msc3h* mutant lines vs EV.

SUPPLEMENTARY FIGURE 3

Correlation analysis of lignin composition of stems with forage quality traits in *Msc3h* mutant lines. (A) S lignin (%) x ADF (%). (B) G lignin (%) x ADF (%). (C) H lignin (%) x ADF (%). (D) H+G+S lignin (%) x ADF (%). (E) S lignin (%) x NDF (%). (F) G lignin (%) x NDF (%). (G) H lignin (%) x NDF (%). (H) H+G+S lignin (%) x NDF (%).

SUPPLEMENTARY FIGURE 4

Correlation analysis of lignin composition of stems with forage quality traits in *Msc3h* mutant lines. (A) S lignin (%) x TDN (%). (B) G lignin (%) x TDN (%). (C) H lignin (%) x TDN (%). (D) H+G+S lignin (%) x TDN (%). (E) S lignin (%) x RFV (%). (F) G lignin (%) x RFV (%). (G) H lignin (%) x RFV (%). (H) H+G+S lignin (%) x RFV (%).

SUPPLEMENTARY FIGURE 5

Correlation analysis of lignin composition of stems with forage quality traits in *Msc3h* mutant lines. (A) S lignin (%) x IVTDMD (%). (B) G lignin (%) x IVTDMD (%). (C) H lignin (%) x IVTDMD (%). (D) H+G+S lignin (%) x IVTDMD (%). (E) S lignin (%) x CP (%). (F) G lignin (%) x CP (%). (G) H lignin (%) x CP (%). (H) H+G+S lignin (%) x CP (%).

SUPPLEMENTARY TABLE 1

Relative advantage (%) of lignin and nutrition quality in alfalfa *Msc3h* mutant lines over EV (control).

SUPPLEMENTARY TABLE 2

Sequences of primers used in this study.

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