APPLICATION OF SHORT TANDEM TARGET MIMIC (STTM) TECHNIQUE FOR FUNCTIONAL ANALYSIS OF MICRO-RNA396 IN TRANSGENIC POPLAR TREES

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APPLICATION OF SHORT TANDEM TARGET MIMIC (STTM) TECHNIQUE 
FOR FUNCTIONAL ANALYSIS OF MICRO-RNA396 
IN TRANSGENIC POPLAR TREES

By
Surattana Boonsai

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This thesis has been approved in partial fulfillment of the requirements for the Degree of MASTER OF SCIENCE in Biological Sciences.

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Preface

In this study, the experiments were designed together by Dr. Chandrashekhar Joshi, Dr. Guiliang Tang and myself. The PCAMBIA1300STTM396a construct was made by Dr. Haiping Liu. All transgenic plants were generated and analyzed by the author of this thesis.
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First, I would like to thank my advisor, Dr. Chandrasekhar Joshi for all the advice and support for almost three years during my MS studies at Michigan Tech. I also would like to thank my thesis co-advisor, Dr. Guiliang Tang for his advice during experimental design and my thesis committee member Dr. Rupali Datta for her valuable help during my research work and thesis writing. I also would like to thank Biological Sciences staff and faculty for their help during my research. I wish to thank the Royal Thai Government and the Biological Sciences department for the financial support.

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Last but not least, I would like to thank my family for always supporting me all the way. Also, my Thai friends at Michigan Tech are also my family for supporting me and helping me during my Michigan Tech life.
Abbreviations

**Ath-GRF** Arabidopsis thaliana-Growth regulating factor

**Bra-GRF** Brassica rapa subsp. Pekinensis- growth regulating factor

**GRF** growth regulating factor

**MiR** microRNA

**Osa-GRF** Oryza sativa-Growth regulating factor

**Ptr-GRF** Populus trichocarpa-Growth regulating factor

**STTM** short target tandem mimic

**Sly-GRF** Solanum lycopersicum-Growth regulating factor

**WT** wild type

**Zma-GRF** Zea mays- Growth regulating factor
Abstract

Short Tandem Target Mimic (STTM) is a commonly used technique for functional studies of a number of genes in several plant model systems. However, very little is known about application of STTM technique in tree species. In this study, STTM was applied to knock down microRNA396 (miR396) in transgenic poplar trees for the first time. STTM396 expression resulted in dramatic decrease in miR396 expression levels leading to taller plants with larger leaves and larger leaf cell size. Additionally, an expression analysis of growth regulating factor genes (GRFs) that are members of miR396 target gene family showed up-regulation of GRF07 gene suggesting that miR396 might be negatively regulating growth and cell expansion in leaves via GRF07 suppression. On the contrary, down-regulation of GRF06 and GRF18 was observed in the same transgenic trees suggesting the complexity of gene regulatory relationship between expression of miR396 and GRFs in poplar. Remarkably, GRF10 and GRF17 were barely expressed in leaves tissues suggesting they might be expressed in other tissues in a tissue-specific manner. Overall, these results suggest that the STTM is an effective approach for functional analyses of miRNAs in poplars, a model tree species. In addition, miR396 may be playing some important roles in growth and cell expansion of leaves via regulating expression of GRF genes in poplars.
Chapter 1 Introduction

1.1 Overview and Significance

MicroRNAs (miRNAs) are short, non-coding RNAs consisting of 20-24 nucleotides (nt) that can silence gene expression of target genes via messenger RNA (mRNA) degradation and/or mRNA translational repression (Pareek et al., 2015). In plants, miRNAs have been reported to play important roles in various biological processes including in growth and development (Li and Zhang, 2016).

For example, miR396 is one of the most conserved miRNAs among plant species (Sun, 2012). Computational prediction suggests that miR396 targets expression of growth regulating factors (GRFs), which are plant-specific transcriptional factors (https://plantgrn.noble.org/psRNATarget/analysis). Because miR396 as well as GRFs are highly conserved among plants, miR396 are known to regulate cell growth and proliferation via regulating expression of GRF genes in various plants such as Arabidopsis, rice, tomato, maize, and Chinese cabbage but little is known about miR396 and GRFs in tree species (Liang et al., 2014; Gao et al., 2015; Cao et al., 2016; Zhang et al., 2018; Wang et al., 2014).

Poplar is an excellent tree model system for forest genetics and woody plant research due to its rapid growth rate, well-studied small genome and capability to be transformed using Agrobacterium-mediated gene expression vectors (Tuskan et al., 2006). In order to study miRNAs functions, Short Tandem Target Mimic (STTM) is a relatively new technique to block functions of miRNAs (Tang et al., 2012). STTM has two mimic miRNA binding sites, which partially complement with target miRNA. STTM has been used to block miRNAs effectively in animals and several plant species (Tang et al., 2012) but has been applied to study miRNAs in trees only a few times (Su et al., 2018; Kumar, 2019). Therefore, applying STTM to study miR396 and GRFs in poplar will elucidate functions of miR396 and GRF gene family and examine the efficiency of STTM in poplar trees.
1.2 MicroRNAs

MicroRNAs (miRNAs) are 20-24 nt short endogenous non-coding small RNAs that can inactivate target mRNA and silence gene expression (Reinhart et al., 2002). Briefly, primary miRNA (Pri-miRNA) is encoded from a miRNA gene that forms a hairpin loop structure in the nucleus. Then, Pri-miRNA is modified through several steps by Drosha-DGCR8 and exported to cytoplasm by HASTY, a homolog of an Exportin 5 (Exp5) in animals. Afterward, it is incorporated into Dicer and Argonaute proteins to form RNA-induced silencing complex (RISC) and bind to target mRNA. In plants, the RISC can either degrade target mRNA directly (mRNA degradation) if it can match with the target mRNA perfectly or it can suppress translation of the target mRNA (inhibition of translation) in case of a partial complement (Bartel, 2004; Axtell et al, 2011) as shown in Figure 1.1.

![Diagram of plant miRNAs biogenesis and function](image)

**Figure 1.1**: Diagram of plant miRNAs biogenesis and function (Adapted from Ma et al., 2009; Axtell et al., 2011).

Plant miRNAs play important roles in growth and development (Li and Zhang, 2016). For example, in *Arabidopsis*, miR160 controls root growth and gravitropism by negatively regulating *auxin response factor (ARF)* 10, *ARF16* and *ARF17* (Khan et al., 2011). The miR390 controls growth and senescence through *ARF2* and *ARF3* expression regulation (Mecchia et al., 2013). In addition, miRNAs
are involved with hormone signaling pathways. For example, miR165/166 regulate auxin (IAA) content and sensitivity by enhanced expression of auxin biosynthesis genes in *Arabidopsis* (Jia et al., 2015). Several miRNAs are important in biotic and abiotic stress responses (Khraiwesh et al., 2012). MiR397 positively regulates cold tolerance via C-repeat binding factor (CBF) dependent pathway (Dong and Pei, 2014). Also, miR402 regulates seed germination timing and cold tolerance (Kim et al., 2010).

MicroRNA396 (miR396) is an evolutionarily conserved miRNA among monocot and dicot plants. So far miR396 has 60 members of miR396 from 21 plant species are known (Sun, 2012). According to microRNA database (http://www.mirbase.org), miR396 has two mature types in *Arabidopsis* and tomato, while seven members in poplar, nine members in rice, and eight members in maize (Griffiths-Jones et al., 2006). Alignment of miR396s in *Arabidopsis*, maize, rice, tomato, and poplar show two miR396 groups that have identical mature sequence (Figure 1.2). Remarkably, both mature types of miR396 have only one nucleotide difference at their 3’ end (G vs U) (Figure 1.3).
** Figure 1.2: ** Alignment of miR396s in Arabidopsis, poplar, rice, maize, and tomato.

<table>
<thead>
<tr>
<th>miR396 Allele</th>
<th>Sequence</th>
<th>Length</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Group B</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ath-miR396b-5p</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>ptc-miR396c</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>ptc-miR396e-5p</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>ptc-miR396d</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>osa-miR396c-5p</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>zma-miR396f-5p</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>zma-miR396e-5p</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
<tr>
<td>sly-miR396b</td>
<td>UUCCACAGCUUUUCUUUGAACUU-</td>
<td>21</td>
</tr>
</tbody>
</table>

| **Group A**  |          |        |
| sly-miR396a-5p | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| zma-miR396c | UUCCACAGCUUUUCUUUGAACUG- | 22 |
| zma-miR396d | UUCCACAGCUUUUCUUUGAACUG- | 22 |
| osa-miR396g | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| osa-miR396h | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| osa-miR396d | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| osa-miR396e-5p | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| osa-miR396f-5p | UUCCACAGCUUUUCUUUGAACUG- | 22 |
| ptc-miR396g-5p | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| zma-miR396h | UUCCACAGCUUUUCUUUGAACUG- | 21 |
| zma-miR396g-5p | UUCCACAGCUUUUCUUUGAACUG- | 21 |

** Figure 1.3: ** MiR396 group A and B show one nucleotide difference at the end of 3’ (G vs U).
Tomato (*Solanum lycopersicum*) has two mature miR396 types including miR396a and miR396b, which have only one nucleotide difference at the end of 3’ (G vs U) but miR396a show dramatically higher expression than miR396b in cotyledon, flower, sepal, and fruit suggesting that miR396a might play more important roles in tomato growth and development than miR396b. Furthermore, STTM396a/396a-88 construct could knock down both miR396a and miR396b in tomato (Cao et al., 2016). In *Arabidopsis*, there are two miR396 genes in miR396 family including miR396a and miR396b encoding two mature miRNAs, which are different in the last nucleotide (G in miR396a and U in miR396b). In *Populus trichocarpa*, miR396 family has seven genes (miR396a-g). Mature sequence of ptc-mir396a and ptc-miR396b are identical to ath-miR396a whereas mature sequence of ptc-miR396c is identical to ath-miR396b. In addition, mature sequence of ptc-miR396f and ptc-miR396g have one and two nucleotides variation compared to ath-miR396a respectively (Figure 1.2).

MiR396 has been studied extensively in *Arabidopsis*, rice and some other crop plants such as tomato and maize. For example, miR396 targets *Arabidopsis thaliana-growth regulating factors* (ath-GRFs) gene family, which are transcription factors for cell proliferation in leaves (Rodriguez et al., 2010). Also, miR396 negatively regulates size of flower, sepal, fruit and cotyledon through suppressed GRFs expression in tomato (Cao et al., 2016). In addition, Peng et al. observed larger tomato with bigger flowers, leaves, and fruit when miR396 was knocked down (Peng et al., 2018). Besides, over-expression of ptc-miR396c in tobacco resulted in cotyledon fusion, lack of shoot apical meristem, delayed and altered floral organ specification. The results from this heterologous study suggested a role of miR396 in floral organs specification (Baucher et al., 2013). Nevertheless, functions of poplar-miR396a-b remain unexplored.
1.3 Growth Regulating Factors

Growth regulating factors (GRFs) are a small plant-specific transcription factor family. The main common character of GRFs is containing two conserved regions including QLQ and WRC domain, which are in N-terminal part of GRFs (Omidbakhshfard et al., 2015). The QLQ domain can be found in all eukaryotes, while WRC is a plant-specific domain (Van der Knaap et al., 2000).

According to PlantTFDB database (http://planttfdb.cbi.pku.edu.cn), GRFs family includes 9 proteins in Arabidopsis, 12 proteins in rice and tomato, 14 proteins in maize, and 19 proteins in poplar (Table 1.1). Computational prediction, based on complementarity between miRNA and sequence of the target genes, suggested that 7 out of 9 ath-GRFs, 8 out of 12 osa-GRFs, 8 out of 12 sly-GRFs, 12 out of 14 zma-GRFs, and 18 out of 19 ptc-GRFs are targeted by miR396 (psRNATarget) (https://plantgrn.noble.org/psRNATarget/analysis).

GRFs have been studied experimentally in several plant species. For example, knocking down miR396 resulted in up-regulated osa-GRF6 and increased yield in rice through inflorescence development, altered auxin biosynthesis and auxin-response factors, and branch and spikelet development (Gao et al., 2015). In addition, over-expression of osa-GRF1 in Arabidopsis resulted in impaired stem growth suggesting that it plays regulatory roles in GA-induce stem elongation (Van der Knaap et al., 2000). In Arabidopsis, over-expression of miR396 caused abnormal flowers and down-regulation of all ath-GRFs (ath-GRF1-9) (Liang et al., 2014). In Chinese cabbage (B. rapa L ssp. pekinensis), 15 bra-GRFs out of 17 were expressed mostly in young leaves but fewer bra-GRF genes were expressed in old leaves. Bra-GRF8 showed the largest expression difference between young and old leaves while bra-GRF16 was strongly expressed in roots (Wang et al., 2014). However, the functions of GRFs in poplar remain unclear.
Table 1.1: Number of *Ptc-GRFs* genes and transcription factors in poplar.

<table>
<thead>
<tr>
<th>Gene name</th>
<th>Transcription factor ID</th>
<th>Description(^a)</th>
<th>Chromosome</th>
<th>Length(^b)(aa)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Ptc-GRF01</em></td>
<td>Potri.001G082700.1</td>
<td>GRF 8</td>
<td>Chr01</td>
<td>514</td>
</tr>
<tr>
<td></td>
<td>Potri.001G114000.1</td>
<td>GRF 10</td>
<td>Chr01</td>
<td>203</td>
</tr>
<tr>
<td><em>Ptc-GRF02</em></td>
<td>Potri.001G114000.2</td>
<td>GRF 10</td>
<td>Chr01</td>
<td>178</td>
</tr>
<tr>
<td><em>Ptc-GRF03</em></td>
<td>Potri.001G132600.1</td>
<td>GRF 4</td>
<td>Chr01</td>
<td>343</td>
</tr>
<tr>
<td><em>Ptc-GRF04</em></td>
<td>Potri.001G169100.1</td>
<td>GRF 5</td>
<td>Chr01</td>
<td>347</td>
</tr>
<tr>
<td><em>Ptc-GRF05</em></td>
<td>Potri.002G115100.1</td>
<td>-</td>
<td>Chr02</td>
<td>468</td>
</tr>
<tr>
<td><em>Ptc-GRF06</em></td>
<td>Potri.003G065000.1</td>
<td>GRF 5</td>
<td>Chr03</td>
<td>343</td>
</tr>
<tr>
<td></td>
<td>Potri.003G100800.1</td>
<td>GRF 3</td>
<td>Chr03</td>
<td>348</td>
</tr>
<tr>
<td></td>
<td>Potri.003G100800.2</td>
<td>GRF 3</td>
<td>Chr03</td>
<td>337</td>
</tr>
<tr>
<td></td>
<td>Potri.003G100800.3</td>
<td>GRF 3</td>
<td>Chr03</td>
<td>340</td>
</tr>
<tr>
<td></td>
<td>Potri.003G100800.4</td>
<td>GRF 3</td>
<td>Chr03</td>
<td>343</td>
</tr>
<tr>
<td><em>Ptc-GRF08</em></td>
<td>Potri.003G118100.1</td>
<td>GRF 10</td>
<td>Chr03</td>
<td>202</td>
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<tr>
<td><em>Ptc-GRF09</em></td>
<td>Potri.006G115200.1</td>
<td>GRF 4</td>
<td>Chr06</td>
<td>378</td>
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<tr>
<td><em>Ptc-GRF10</em></td>
<td>Potri.006G143200.1</td>
<td>GRF 1</td>
<td>Chr06</td>
<td>386</td>
</tr>
<tr>
<td><em>Ptc-GRF11</em></td>
<td>Potri.007G007100.1</td>
<td>GRF 1</td>
<td>Chr07</td>
<td>615</td>
</tr>
<tr>
<td><em>Ptc-GRF12</em></td>
<td>Potri.012G022600.1</td>
<td>GRF 4</td>
<td>Chr12</td>
<td>581</td>
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<td>Potri.012G022600.2</td>
<td>GRF 4</td>
<td>Chr12</td>
<td>536</td>
</tr>
<tr>
<td><em>Ptc-GRF13</em></td>
<td>Potri.013G077500.1</td>
<td>GRF 4</td>
<td>Chr13</td>
<td>341</td>
</tr>
<tr>
<td><em>Ptc-GRF14</em></td>
<td>Potri.014G007200.1</td>
<td>GRF 1</td>
<td>Chr14</td>
<td>608</td>
</tr>
<tr>
<td><em>Ptc-GRF15</em></td>
<td>Potri.014G012800.1</td>
<td>-</td>
<td>Chr14</td>
<td>610</td>
</tr>
<tr>
<td><em>Ptc-GRF16</em></td>
<td>Potri.014G071800.1</td>
<td>GRF 9</td>
<td>Chr14</td>
<td>459</td>
</tr>
<tr>
<td></td>
<td>Potri.015G006200.1</td>
<td>GRF 4</td>
<td>Chr15</td>
<td>448</td>
</tr>
<tr>
<td></td>
<td>Potri.015G006200.3</td>
<td>GRF 4</td>
<td>Chr15</td>
<td>412</td>
</tr>
<tr>
<td><em>Ptc-GRF17</em></td>
<td>Potri.015G006200.4</td>
<td>GRF 4</td>
<td>Chr15</td>
<td>436</td>
</tr>
<tr>
<td><em>Ptc-GRF18</em></td>
<td>Potri.018G065400.1</td>
<td>GRF 1</td>
<td>Chr18</td>
<td>383</td>
</tr>
<tr>
<td><em>Ptc-GRF19</em></td>
<td>Potri.019G042300.1</td>
<td>-</td>
<td>Chr19</td>
<td>329</td>
</tr>
</tbody>
</table>

\(^*\) Potri.003G118100.1 is non-miR396 target.

\(^a\) Description was obtained from NCBI.

\(^b\) Length was referred to the translated Pt-GRF proteins.

Data were obtained from http://planttfdb.cbi.pku.edu.cn
Figure 1.4: Ptc-GRFs genes distribution in poplar genome. The chromosome number is indicated at the bottom of each chromosome representation.

*Ptc-GRFs* genes are distributed unevenly in poplar genome as shown in Figure 1.4, which is similar to the previous results in *Arabidopsis* (Kim et al., 2003), tomato (Choi et al., 2004), and rice (Zhang et al., 2008). In poplar, the chromosome 01 has four *ptc-GRFs* (01-04) genes, whereas chromosome 14 has three genes (*GRF14-16*). In addition, two genes each are located on chromosome 03 and 06. Furthermore, one gene each is positioned on chromosome 02, 07, 12, 13, 17, 18 and 19.

1.4 Short tandem target mimic (STTM)

Since plant miRNAs were found in 2002 by several research groups (Llave et al., 2002; Mette et al., 2002; Park et al., 2002; Reinhart et al. 2002;), some approaches have been used to identify and characterize miRNAs. Functional study of genes traditionally relied on reverse genetics to knock down genes of interest and observe loss of functions or altered phenotypes. However, it was difficult to
knockdown miRNAs due to their small size and each miRNA has many target genes. After the discovery of target mimic (TM), short tandem target mimic (STTM) was developed to target small RNAs (Tang et al., 2012).

STTM is a synthetic, short, non-coding RNA (~100 nt). The STTM construct is made as described earlier (Tang et al., 2012) and is expressed by CaMV 35S promoter. The STTM transcript consists of a stem-loop linker at the middle and two small RNA binding sites on both sides. The stem-linker may play a role in STTM stability meanwhile the binding sites partially complement with small RNAs, except CUA region, which enhance STTM to trap small RNAs without being cleaved by them (Figure 1.5). Additionally, the 48-88 nt spacer with a weak hair-pin loop structure is required for STTM. STTM can target miRNA functions by degrading miRNAs directly and suppressing mRNA translation (Tang et al., 2012).

STTM has been used to block miRNAs successfully in fungi, animals and several plants. STTM can knockdown several fungal miRNAs in *Metarhizium robertsii* and *Aspergillus flavus* and leads to up-regulation of target genes involving development and conidial production (Wang et al., 2019). In addition, miR30-d expression was drastically reduced in STTM30-d mouse cells, demonstrating that STTM also functions efficiently in animal cells (Tang et al., 2012). In *Arabidopsis*, STTM165/166 reduced miR165/166 expression level and increased HD-ZIP III transcription factors genes expression, their target genes, leading to a series of developmental changes (Yan et al., 2012). In addition, knocking down miR160 in tomato using STTM160 showed decrease in miR160 expression levels and up-regulation of *SIARF10A*, *SIARF10B* and *SIARF17*, its target genes. Also, STTM160 tomato plants displayed changes in ovary patterning, floral organ abscission and lateral organ outgrowth (Damodharan et al., 2016). Nonetheless, STTM approach has recently been applied to study miRNAs in trees only a few times before (Su et al., 2018; Kumar, 2019).
1.5 Poplars (*Populus* sp.)

Several characteristics of trees, especially having a large size and long life span, differentiate them from other model plants and make molecular and cellular levels study challenging. *Populus* is a good model tree genus suitable for forest genetics due to its modest genome, rapid growth, and compatibility with DNA transformation by *Agrobacterium tumefaciens*. *Populus* includes ~30 species including aspens, cottonwoods, and poplars. *Populus trichocarpa* or black cotton wood was selected as the tree model species for genome sequencing in 2006. Genome sequencing uncovered that it has 19 chromosomes with compact 500 Mbp-big genomes. There is a total of 169 identified miRNA genes representing 21 families (Tuskan et al., 2006).
1.6 Aims of the study and hypothesis

Overall, miR396 has been reported to play roles in growth and development of plant species via regulation of GRFs target gene expression. Since ptc-miR396a has never been studied in any tree species and STTM has been applied to study miRNAs in tree species only a few times, in this study, we applied STTM to study functions of miR396a in poplar. Aims of this study include to investigate efficiency of STTM in poplars and to study functions of ptc-miR396a especially in growth and development through expression of GRF genes. We hypothesized that STTM396a will knock down miR396a expression in transgenic poplar trees and will show decreased miR396a expression level. Also, expression level changes of GRFs, miR396a target genes, are expected. In addition, we expected to see phenotypes changes of poplar in growth and development.
Chapter 2 Materials and Methods

2.1 Plasmid construction, plant transformation and growth condition

The PCAMBIA1300STTM396a construct was made and introduced into *Agrobacterium tumefaciens* EHA105 by Dr. Liu (Liu, 2017). The STTM396a was made based on mature sequence of miR396a as shown in Figure 1.3, group A. and can be expressed as the transcript shown in Figure 2.1. The *Agrobacterium* strain containing STTM396a was cultured with Rifampicin and Kanamycin as selection markers. Positive colonies were verified for an inserted DNA by PCR using B4 and B5 primers before using *Agrobacterium* for plant transformation. In this study, *Populus tremula x Populus alba* clone 717-1B4 was used as wild type (WT) control and to generate transgenic plants. The *Agrobacterium* carrying STTM396a construct was used to transform wild type poplar plants by leaf disk transformation method previously described by (Tsai et al. 1994; Liu et al., 2012).

Briefly, wild type leaves were cut to allow *Agrobacterium* infection through several steps. Then, callus formation was induced after growing the infected leaves on callus induction media (CIM) for three to four weeks under light and dark cycle for 16 and 8 hours at 25 °C in a growth chamber. Next, calluses were moved to shoot induction medium (SIM) and shoot elongation medium (SEM) respectively for three to four weeks each to allow shoots to grow and elongate. Each shoot that grown out from separate callus was considered as an independent line and was moved to root induction medium (RIM) for another three to four weeks. Each line that started rooting was propagated on Murashige and Skoog (MS) media for another four weeks. Hygromycin (10 µg/ml) was used as a selection marker for STTM plants.

Then, DNA of both wild type and transgenic plants were isolated and transgene insertion was verified by PCR using the B4 and B5 primers before moving plants from artificial media to soil. Plants were acclimatized in soil and
watered as needed for two to three weeks before moving to greenhouse for further experiments. In greenhouse, plants were grown under light and dark cycle for 16 and 8 hours, respectively at 23-28 °C, watered every day and applied regular fertilizer supply every two weeks.

![Diagram of STTM396 transcript binding with miR396 through STTM396a binding sites]

**Figure 2.1:** Transcript of STTM396a.
The STTM396 transcript binds with miR396a through STTM396a binding sites.

### 2.2 Phylogenetic analysis

The protein sequence of 9 ath-GRFs and 19 ptc-GRFs were obtained from plant transcription factor database or PlantTFDB (http://planttfdb.cbi.pku.edu.cn). Then all 28 protein sequences were aligned using ClustalX 2.1 (Larkin et al., 2007) before performing phylogenetic tree analyses. A phylogenetic tree was generated with OMEGA 7.0 (Hall, 2013) program using neighbor-joining (NJ) method and 1,000 replicates of bootstrapping test.

### 2.3 Total RNA isolation and stem-loop RT-qPCR

PsRNATarget (https://plantgrn.noble.org/psRNATarget/analysis) was used to select predicted miR396 target genes for RT-qPCR. All 19 ptc-GRFs were
ranked by setting at 25 for maximum energy to unpair the target site (UPE) and at 3 for maximum expectation for complementarity. The first five GRFs including GRF07 (Potri.003G100800.1), GRF10 (Potri.006G143200.1), GRF17 (Potri.015G006200.1), GRF18 (Potri.018G065400.1) and GRF06 (Potri.003G065000.1) were chosen to represent predicted ptc-miR396 target genes.

Leaves were collected and stored in liquid nitrogen. Total RNA was isolated using TRIzol (Invitrogen, USA) method. The stem-loop RT primers and qPCR primers for miR396 were designed according to the method developed by Kramer (Kramer, 2011). The 1 µg total RNA was reverse transcribed using a high-capacity cDNA Archive kit (Applied Biosystems, USA). Applied Biosystems step one machine was applied to perform stem-loop qPCR for miRNA (Varkonyi-Gasic et al., 2007) and qPCR for mRNA using SYBR green PCR master mix (Applied Biosystems, USA). Actin was used as an internal control for qPCR. All primers are listed in Table 2.1. All gene expression analysis was performed using three biological replicates or more.

Table 2.1: Primers for real-time PCR and stem-loop real-time PCR.

<table>
<thead>
<tr>
<th>Primer name</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>miR396 Forward</td>
<td>ACAAATGCAGCTGGGTTCCACAGCTTTCTTT</td>
</tr>
<tr>
<td>miR396 Reverse</td>
<td>ATGGAGAGACCTGTAGTCTTTA</td>
</tr>
<tr>
<td>Potri.003G100800.1 GRF07 Forward</td>
<td>GCAGAGGGTATCCAAGCTAATC</td>
</tr>
<tr>
<td>Potri.003G100800.1 GRF07 Reverse</td>
<td>ATGGAGAGACCTGTAGTCTTTA</td>
</tr>
<tr>
<td>Potri.006G143200.1 GRF10 Forward</td>
<td>ATCCCTCCGAATCTTCTTT</td>
</tr>
<tr>
<td>Potri.006G143200.1 GRF10 Reverse</td>
<td>ATCCCTCCGAATCTTCTTT</td>
</tr>
<tr>
<td>Potri.015G006200.1 GRF17 Forward</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Potri.015G006200.1 GRF17 Reverse</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Potri.018G065400.1 GRF18 Forward</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Potri.018G065400.1 GRF18 Reverse</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Potri.003G065000.1 GRF06 Forward</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Potri.003G065000.1 GRF06 Reverse</td>
<td>GCTTCCCAAACCCTATCT</td>
</tr>
<tr>
<td>Stem-loop universal Reverse</td>
<td>CTGGTGATTTCCACGCGTCGAGA</td>
</tr>
<tr>
<td>B4</td>
<td>AACCTCCCGGATTTCCATGCGAGA</td>
</tr>
<tr>
<td>B5</td>
<td>AACTCCCGGATTTCCATGCGAGA</td>
</tr>
<tr>
<td>Actin Forward</td>
<td>ATGGCCGATTTCCAGATTTCCATGAG</td>
</tr>
<tr>
<td>Actin Reverse</td>
<td>TCTGCCCAATTCCACCATGACA</td>
</tr>
</tbody>
</table>
2.4 Microscopy

Wild type and transgenic leaves were cross-cut at the middle into 0.4 cm pieces and immersed in cold fixative solution containing formaldehyde, acetic acid, and ethanol overnight at 4 °C. Then, fixed samples were dehydrated and embedded in wax. Next, five micrometers thick section were crosscut consecutively with 100 micrometers between level by microtome and mounted on slides (performed by Sharon Juntilla at Histotech lab, Portage Hospital, Hancock). Slides were dewaxed with xylene and ethanol series before stained with safranin and observed under light microscope (Adapted from Johansen, 1940). To investigate cell proliferation and cell expansion change in cellular level, cell size and cell number were measured within the same four areas of leaves cross section for three pictures each using ImageJ program (https://imagej.nih.gov/ij/).

2.5 Growth measurements

After plants were moved to the greenhouse for one week, growth data including estimated height (cm), estimated stem girth (mm), leaf count (n), and leaf length (cm) were measured every week.

2.6 Statistical analysis

In this study, SPSS 25.0 software (https://www.ibm.com/analytics/spss-statistics-software) was used to conduct statistical significance analysis of the data. Student T-test was used to analyze two data sets. One asterisk (p≤0.05), two asterisks (p≤0.01), and three asterisks (p≤0.001) were displayed.
Chapter 3 Results

3.1 Phylogenetic tree of GRFs

Figure 3.1: Phylogenetic tree of AthGRFs and PtcGRFs.

Predicted GRF protein sequences from Arabidopsis (Ath-GRF) and poplar (Ptc-GRF) indicated PtcGRF19 and AthGRF9 are outgroup. GRFs can be divided into 6 groups. The phylogenetic tree also suggested that some of GRFs are identical including PtcGRF10 and PtcGRF18, PtcGRF04 and PtcGRF06, PtcGRF03 and PtcGRF07, PtcGRF13 and PtcGRF19, PtcGRF11 and PtcGRF14, PtcGRF05 and PtcGRF15, PtcGRF12 and PtcGRF17, PtcGRF02 and PtcGRF08 in poplar. In Arabidopsis, there are a few GRFs that are identical including
AthGRF1 and AthGRF2 and AthGRf3 and AthGRF4. These could be a result from genome duplication in poplar. Also, those identical GRFs might play redundancy roles in plants. According to the phylogenetic tree, AthGRFs and PtcGRFs can be classified into six group including group I (AthGRF6, AthGRF5, PtcGRF10, PtcGRF18, PtcGRF04, PtcGRF06, PtcGRF03, PtcGRF07, PtcGRF13, PtcGRF19), group II (AthGRF1, AthGRF2, PtcGRF11, PtcGRF14, PtcGRF05, PtcGRF15), group III (AthGRF3, AthGRF4, PtcGRF09), group IV (PtcGRF01, AthGRF8, AthGRF7, PtcGRF12, PtcGRF17), group V (PtcGRF02, PtcGRF08), group VI (PtcGRF16, AthGRF9). According to those six group, group VI including PtcGRF16 and AthGRF9 is the out group suggesting they might play similar role in poplar and Arabidopsis. Group V includes only PtcGRF02 and PtcGRF08 implying that they might play some specific role only in poplar but not in Arabidopsis.

3.2 MiR396 was effectively knocked down in poplar by STTM

MiR396 was knocked down effectively in poplar by STTM. STTM396 resulted in dramatically decreased relative expression level of miR396 in four lines of STTM396 plants, #3 #5 #6 #9, compared to one line in wild type (Figure 3.2 A). The predicted miR396 target genes showed GRF07 up-regulation (Figure 3.2 C), down-regulation of GRF06 and GRF18 (Figure 3.2 B and 3.2 F). Besides, another two targets genes, GRF10 and GRF17, were barely expressed in both wild type and STTM plants (Figure 3.2 D and 3.2 E).
Figure 3.2: MiR396 was effectively knocked down in poplar by STTM. (A-F) RT-qPCR analysis of miR396 and five predicted target genes in WT and four independent lines of STTM plants. Binding sites between miR396 and its target genes, which are in coding region of QLQ domain, are shown. Asterisk displayed statistically significant difference applying student paired t-test (*p≤0.05, **P≤0.01, ***p≤0.001). The error bars indicated mean ± SD.

3.3 Blockage of miR396 induced taller plants with larger leaves

After plants were moved to the greenhouse, growth measurement was recorded every week. Estimated leaf length from base to tip of the first two leaves from the bottom of wild type and five lines of transgenic plants were tracked.
STTM396 plants showed significantly larger leaf length in both positions compared to one in wild type (Figure 3.3-3.4 and 3.6 B).

**Figure 3.3:** Length of the first leaf from the bottom in centimeters (cm). Comparison between wild type and STTM396 plant leaves over seven weeks.

**Figure 3.4:** Length of the second leaf from the bottom in centimeter (cm). Comparison between wild type and STTM396 plants over seven weeks shown.
Figure 3.5: Estimated height in centimeter (cm) of wild type and STTM396 plants over seven weeks.

Remarkably, transgenic plants were notably taller but had smaller stem girth than in wild type plants (Figure 3.5, 3.6 A and 3.7). To explain this phenomenon, more lines of transgenic plants should be examined. Besides, leaf count data...
showed inconsistent trends in STTM plants (Figure 3.8). More lines of transgenic plants are also required to conclude this result.

**Figure 3.7:** Estimated stem girth in millimeter (mm) of wild type and STTM396 plants over seven weeks.

**Figure 3.8:** Leaf count (n) of wild type and STTM396 plants over seven weeks.
3.4 STTM396 plants have altered leaf cell size and cell number

To investigate cell proliferation and cell expansion change in cellular level, cross-section of wild type and STTM396 leaves were observed under bright field microscope. STTM396 resulted in larger cell size and smaller cell number in all four areas (Figure 3.9, 3.10 and 3.11) suggesting that altered leaves size was a result from cell expansion rather than cell proliferation.

![Figure 3.9: Transverse section of wild type and STTM396 leaves, particularly at mid vein region.](image)

![Figure 3.10: STTM396 resulted in smaller cell number in STTM leaves. Cell number measurement in WT and STTM leaves. Asterisk displayed statistically significant difference applying student paired t-test (*p≤0.05, **P≤0.01, ***p≤0.001). The error bars indicated mean ± SD.](image)
Figure 3.11: STTM396 resulted in larger cell size in STTM leaves. Cell size measurement in WT and STTM leaves. Asterisk displayed statistically significant difference applying student paired t-test (*p≤0.05, **P≤0.01, ***p≤0.001). The error bars indicated mean ± SD.
Chapter 4 Discussion

4.1 MiR396 was effectively knocked down in poplar by STTM

MiR396 is one of the highly conserved miRNAs among plant species. Mature sequence of miR396a in Arabidopsis, rice, and maize and miR396a and miR396b in poplar are identical (http://www.mirbase.org). Their targets are members of GRFs gene family, which are plant-specific transcription factors. Poplar has 19 GRFs, whereas Arabidopsis, rice, and maize has 9, 12, and 14 GRFs respectively. In addition, GRFs genes are distributed unevenly in poplar genome, mostly on chromosome 01 and 14 (Kim et al., 2003; Choi et al., 2004; Zhang et al., 2008).

STTM has been used to knockdown small RNAs successfully in fungi, animals, and several plants species (Wang et al., 2019; Tang et al., 2012; Yan et al., 2012). Here, we show that STTM can also disrupt miRNA functions effectively in poplar, which represents a tree species. Remarkably, STTM396 resulted in 78-98% suppression of miR396 expression levels (Figure 3.2 A).

4.2 Blockage of miR396 induced taller plants with larger leaves

STTM396 produced taller transgenic plants (Figure 3.6A and 3.7) with larger leaves (Figure 3.6 B, 3.7 and 3.8). These results suggest that ptc-miR396 might be negatively playing a role in leaf growth and development of poplar. This conclusion is consistent with other studies in Arabidopsis and tomato. (LiuKanpp et al., 2009; Cao et al., 2016) suggesting that miR396 as a negative regulator of plant growth.

4.3 STTM396 plants have altered leaf cell size and cell number

Furthermore, organ size is basically determined by cell expansion and/or cell proliferation (Rodriguez et al., 2016). To understand causes of organs enlargement, leaf histology was examined. Leaf transverse sections at the midrib
region showed larger cell sizes but smaller cell number in STTM396 leaves as compared to wild type control (Figure 3.9-3.11). These results indicate that the leaf enlargement may be a result of cell expansion rather than cell proliferation. In stem, STTM396 plants showed increased in height but reduced girth than control plants. These could be resulted from cell expansion or cell elongation as in rice and Arabidopsis. Previous studies provided some clues about miR396 regulatory functions through its target genes, GRFs. For example, Van der Knaap et al. (2000) suggested that osa-GRF1 may be involved with stem elongation in rice. Also, suppression of osa-GRFs resulted in dwarf plants. (Kuijt et al., 2014).

However, processes of stem growth and development in tree species are likely to be more complicated than monocot plants and other herbaceous dicot plants. Furthermore, a careful histological study of transgenic poplar stems expressing STTM396 is required to explain these phenotypes. Therefore, the regulation of cell proliferation and expansion by miR396 is complicated and might be different among plant species and even in different tissues.

4.4 Expression level of GRFs changed in STTM396 plants

Mir396 target genes expression analysis in transgenic poplars expressing STTM396 showed that GRF07 was up-regulated but GRF06 and GRF18 were down-regulated in STTM plants. These results suggested that miR396 normally plays a role as negative regulator of growth and cell expansion through GRF07 suppression. Additionally, GRF06 and GRF18 may be not only regulated by miR396 but also controlled by another gene or another miRNA as shown in Figure 4.1 in a much stronger manner. Also, qRT-PCR analysis suggests that GRF10 and GRF17 are barely expressed in leaves tissues suggesting that these two genes may be expressed in other tissues for example stem instead of leaves. Similarly, bra-GRFs were expressed in a tissue-specific manner in Chinese cabbage. Bra-GRF1, 4, 6, 8, 10, 15, and 17 were expressed in young leaf higher than in other tissues (Wang, 2014). However, another 14 ptc-GRFs are required to be studied
for a better understanding of role of various GRFs in poplar growth. Also, over-expression of GRF07 might help to confirm the roles of GRF07 in poplars.

![Proposed schematic relations between miR396 and GRFs affecting growth and development in poplar.](image)

**Figure 4.1**: Proposed schematic relations between miR396 and GRFs affecting growth and development in poplar.

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**4.5 Conclusion**

In this study, STTM was applied to knock down miR396 in poplar. STTM396 resulted in dramatic suppression of miR396 expression levels and taller plants with larger leaves with larger cell size. Up-regulation of GRF07 and down-regulation of GRF06 and GRF18 were observed in STTM396 poplars suggesting that ptc-miR396 might be negatively regulating growth and development through GRF07 gene suppression. Besides, GRF06 and GRF18 may not be only controlled by miR396 but probably be regulated by another miRNA or gene in a much stronger manner. Remarkably, GRF10 and 17 were barely expressed in leaves tissue suggesting that they might be expressed in other specific tissues.

In conclusion, these results indicate that STTM is an effective approach to knock down miRNAs in poplars, a tree species. Additionally, ptc-miR396a may play role in growth and development of poplar especially organ size regulation supporting the observation that miR396 acts as a negative regulator in plant growth
and development. Furthermore, ptc-miR396 mediate ptc-GRFs expression is a complex network. They might be expressed and function in tissue-specific manner.

### 4.6 Future studies

In the future, stem GRFs should be studied for a better understanding of interrelationship between miR396 and GRFs in poplars. To complete the whole picture of miR396 and ptc-GRFs functions, the rest of the 14 GRFs should be also studied in both leaves and stems of transgenic STTM396 poplar plants. Furthermore, importance of miR396 and ptc-GRFs in reproductive organs should also be explored in the future. This could be done via their ectopic expression in other plant model species such as Arabidopsis or tobacco because flowering in poplars takes many years.
References


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