An Overview on Advances in Cotton Genome and Regulation of Fiber Development

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Abstract: In the evolution of flowering plants major driving forces for the evolution is polyploidy and whole-genome duplication. The genus Gossypium contains 45 diploid (2n = 26) and six tetraploid (2n = 52) species. During evolution, polyploidy (allo-tetraploidy) enhanced the fiber traits in allo-tetraploids as compared to diploids, resulting in higher fiber yield and quality. The advancement of G. hirsutum genome into AD-genome (allo-tetraploid) brings by the polyploidization event between diploid progenitors of A- and D-genome. The genome size of G. raimondii and G. arboreum is 880 Mb and 1,700 Mb respectively reveals that the size of D-genome is lesser than the genome size of A-genome. Initiation and elongation are the best understood developmental stages at the transcriptional level among different stages of fiber development. However, scanty knowledge is available about the SCW stage-specific transcriptional regulation and its detailed molecular mechanism. Since specific promoters regulates transcription in fiber cells hence the knowledge of novel fiber specific regulatory elements along promoters and their interacting transcription factors might be a better tool to be used in the genetic manipulation of fiber quality parameters.

Keywords: Gossypium, transcription regulation, promoter, transcription factor, cotton genome, regulatory elements

1. Introduction

The cotton fiber is a single cell structure initiated on the epidermal surface of an ovule and eventually undergoes into four developmental stages namely: initiation, elongation, secondary cell wall (SCW) synthesis and maturation (Basra and Malik 1984). A single mature cotton fiber consists of the thin outer primary cell wall (PCW), an inner thick secondary cell wall (SCW), and a central lumen. In spite of being a valuable resource for the textile industries cotton fiber is an excellent model to study the cell differentiation and development. Cotton fiber is an important raw material for the textile industry. Economy of the world and daily human life is affected by the world wide cotton production. The genus Gossypium contains 45 diploid (2n=26) and six tetraploid (2n = 52) species (Hawkins et al., 2006 Grover et al., 2015). The genus, Gossypium is a perennial shrub belongs to malvaceae family, grown mainly for the seed hair (textile fiber). The spinnable fiber is produced by two tetraploids (G. hirsutum and G. barbadense) and two diploids Gossypium species (G. herbaceum and G. arboreum).

In the evolution of eukaryotic organism's specially in flowering plants the main driving force for the evolution is polyploidy and whole-genome duplication (Soltis et al., 2014; Hegarty et al., 2008; Otto et al., 2007; Jiao et al., 2011). During evolution, polyploidy (allo-tetraploidy) enhanced the fiber traits as compared to diploids, resulting in higher fiber yield and quality (Chen et al., 2007; Paterson et al., 2012; Li et al., 2014; Guan et al., 2014). The *G. hirsutum*

is an allo-tetraploid (AD-genome) species evolved through the polyploidization event between diploid progenitors of Aand D-genome during the course of evolution (Page et al., 2013). D- genome is dominating than A- genome inferred from the origin of AD genome, transposable elements (TEs) and reduction in the genome size (AD) resulted from allopolyploidization (Li et al., 2015) during course of evolution of *Gossypium* spp. The *G. hirsutum*, in which fiber length is approximately 25 to 34 mm, is most widely cultivated cotton overall and more than 90% of lint fiber of world's total cotton production is obtained from *G. hirsutum* (Lin et al., 2009). Thus, the *G. hirsutum* tends to become the intensive target of basic and applied research.

2. Evolutionary divergence of Gossypium

The phylogenetic analysis revealed *Gossypium arboreum* and *Gossypium raimondii* evolved from T. cacao (Li et al., 2014; **Figure 1**). The differences in the size of *Gossypium arboreum* (AA; 2n = 26) and *Gossypium raimondii* (DD; 2n = 26) genome are due to the insertions of long terminal repeats (5 million years back).

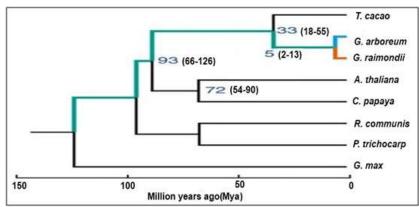


Figure 1: The schematic diagram of the *G. arborium*, *G. raimondii* and six other genomes descended from common eudicot genome ancestors. Number represent the evolutionary time in million years (Mya) (Source: Li et al., 2014).

Allo-tetraploid cotton (AADD) evolved from the natural hybridization between A and D genome species and has split into six species, including the widely cultivated *G. barbadense* (AD2) and *G. hirsutum* (AD1). The *Gossypium spp.* comprises of nine different genome groups (A, B, C, D, E, F, G and K). The genomes sizes of different genome groups (A, B, C, D, E, F, G and K) vary due to the lineage-specific proliferation of retro-transposons (Hawkins et al., 2006). The genome size of *G. raimondii* and *G. arboreum* is 880 Mb (Hendrix et al., 2005; Paterson et al., 2012; Wang et al., 2012) and approximately 1,700 Mb (Li et al., 2014)

respectively implies that the genome size of D genome is less than the genome size of A genome. Genome duplication, retro-transposons and polyploidization events during course of evolutionary process in *Gossypium* spp. including flowering plants and many crop plants results in divergence of genome into different sub-genomes (Renny-Byfield et al., 2014; Gong et al., 2013). However, the molecular mechanism of occurrence of allopolyploids and the fiber traits of cotton remains limited.

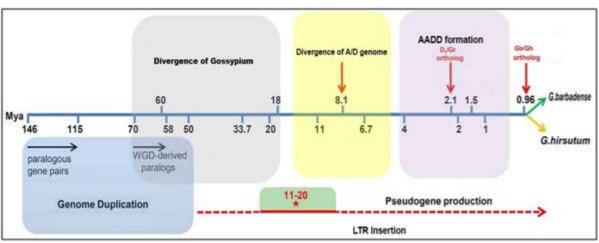


Figure 2: The schematic presentation of the evolution of allotetraploid (AADD) genome of cotton

G. hirsutum (AD1) and *G. barbadense* (AD2) splits from allotetraploid cotton (AADD) which evolved from A- and D- genome. Numbered axis represent time in million years (Mya) and evolutionary events are represented by arrows and encircled in boxes. Red star indicates pseudogene production. *Gr: G. raimondii; Gb: G. barbadense; Gh: G. hirsutum,* LTR: Long terminal repeats (Source: Liu et al., 2015).

3. Expression Pattern of Genes in Cotton Fiber Development

A number of reports have been generated to identify the genes related to fibre development. People have used microarray to identify the genes involved in fibre development of domesticated, mutant and poor fibre quality cotton cultivars (Li et al., 2002, Wu et al., 2007; Yang et al.,

2006, Nigam et al., 2014). The differentially expressed genes (*DEGs*) during different stages of fiber development are required for fiber differentiation and development. Several genes from developing cotton fiber have been cloned and characterized (John & Crow, 1992; John & Keller, 1995; John & Keller 1996; Reinhart et al., 1996; Ma et al., 1995, 1997; Song & Allen, 1997; Smart et al., 1998; Whitaker & Triplett, 1999; Orford & Timmis, 2000; Liu et al., 2000). So far, only few of these genes are known to involved in the biosynthesis of cell wall components have been identified.

Initiation and elongation stage of fiber development is better understood. Some of the characterized initiation and elongation-specific regulator and genes include; *MYB25* (Machado et al., 2009; Wu et al., 2006), *MYB25-like* (Walford et al., 2011), *Sucrose synthase* (*Sus*) (Ruan et al.,

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2003), GbPDF1 (Deng et al., 2012), GhH2A12 (Hao et al., 2014), IAA (Zhang et al., 2011), GhMYB2 (Wang et al., 2004), GhHD1 (Walford et al., 2012) and GhJAZ2 (Hu et al., 2016), GhFAnnxA (Zhang et al., 2016), GhCaM7-like (Cheng et al., 2016), GhMYB109 (Pu et al., 2008) and GhPK6 (Zhang and Liu, 2016) etc. During fiber development fiber cells undergo dramatic changes in gene expression that lead to massive changes in the level of proteins and metabolites important for fiber development (Mansoor & Paterson, 2012). A detailed microarray analysis of fiber development in contrasting genotypes of G. hirsutum suggested about the transcriptional reprogramming during fiber development (Nigam et al., 2014). Some fiber genes are reported to be constitutively expressing whereas others are reported to express during the specific stages (John & Crow, 1992; Delmer et al., 1995; John & Keller, 1995; Ma et al., 1997; Song & Allen, 1997; Smart et al., 1998; Loguerico et al., 1999; Whittaker & Triplett, 1999; Orford & Timmis, 2000; Xu et al., 2008; Nigam et al., 2014).

With the recent availability of progenitors genome sequences (Paterson et al., 2012; Wang et al., 2012; Li et al., 2014), more focus on cotton genomic studies is now shifted towards gaining insight into the mechanistic details. The study of mechanisms involved in regulation of critically important genes and metabolic networks at various levels is of immediate need to utilize the rapidly increasing cotton genomic resources for the betterment of cotton fiber.

4. Involvement of **Promoter** Fiber in Development

Several promoters have been reported that regulate gene expression in cotton fibers during the initiation and elongation stages (Delaney et al., 2007; Hussey et al., 2011; Larkin et al., 1996; Li et al., 2002; Li et al., 2005; Ma et al., 1995; Ni et al., 2008; Song and Allen, 1997; Wang et al., 2004; Wu et al., 2009). The promoter of GaRDL1, GhTUB1 and GhMYB25 are active early during the initiation stage (Li et al., 2002; Machado et al., 2009; Wang et al., 2004), while the glucuronosyl transferase promoter (PGhGlcAT1), has been shown to be highly active during fiber elongation (Wu et al., 2007). The fiber-specific lipid transfer protein (FSltp4) promoter has also been reported to be active during fiber elongation and has been shown to be functional during the synthesis of fiber cutin (Delaney et al., 2007). GhACTIN1 is another gene expressed in fiber and its promoter has been shown to be elongation-specific (Li et al., 2002). Promoters for the genes, GhLTP3, GhDET2, GaMYB2, and GhMYB109, have also been reported to be active during initiation and some of them continue to be active until the elongation stage (Liu et al., 2000; Luo et al., 2007; Pu et al., 2008; Wang et al., 2004). There are other promoters, such as P_{GhRING1}, that are active in all the stages of cotton fiber development, starting from initiation to the SCW stage (Ho et al., 2010). Among different developmental stages of the fiber, initiation is the best understood at the transcriptional level. However, there are scanty knowledge about the SCW stage-specific transcriptional regulation and its detailed molecular mechanism. Recently SCW stage specific GhGDSL promoter have been cloned and characterized in cotton transgenic lines and showed enhanced GUS and GFP expression at SCW biosynthesis stage (19-25DPA) of fiber development. In this study through yeast one hybrid interaction (Y1H) it has been shown that GhGDSL promoter interact with GhMYB1 which is the regulator of SCW stage specific expression in Gossypium hirsutum (Yadav et al., 2017).

Efforts to understand and manipulate fiber development would be greater by an improved knowledge of the promoter elements and transcription factors that regulate fiber-specific gene expression. Promoters are DNA sequences that determine the expression level, time and tissue specificity of gene. Genetic engineering for improved agronomic traits requires specific promoters for the modification of fiber properties (John, 1998). The promoters of fiber-specific genes may regulate gene function and fiber development by altering transcription of fiber cells.

| | Table 1: Some conserve motif sequences of promoter | | | | | |
|-------|--|----------------|--|-------------------------------|--|--|
| S.No. | Conserve | Motifs ID | Putative Functions | References | | |
| | Sequences | | | | | |
| 1 | CW8G | CARGCW8GAT | A variant of CArG, binding site for AGL15 | Plant Physiol (2006) | | |
| 2 | ACACNNG | DPBFCORE | Age & developmental regulation | Mol Cell Biol (2012) | | |
| 3 | GATA | GATABOX | Tissue-specific promoter activity of legumin gene | Mol Gen Genet (1989) | | |
| 4 | GCCGCC | GCCCORE | Light & SA-inducible expression | Plant Mol Biol (1995) | | |
| 5 | GRWAAW | GT1CONS | Transcriptional activation of the rbcS 3A gene | EMBO (1988) | | |
| 6 | GGTTAA | GT1CORE | Pathogen & salt-induced SCaM4 gene expression | Plant Physioly (2004) | | |
| 7 | GAAAAA | GT1GMS | Regulation of expression of histone H3 gene | Plant Cell Physioly (1989) | | |
| 8 | ACGTCA | HEXMOTIF | Light-regulated transcription | Plant Mol Biol (1995) | | |
| 9 | GATAA | IBOXCORE | Light responsive transcription of psadb gene | Plant Journal (2002) | | |
| 10 | TAAATGYA | L1BOXAT | Consensus sequence for plant introns | Nucleic Acids Research (1986) | | |
| 11 | WAACCA | MYB1AT | Trichome regulation, phospholipid signaling (GLABRA2) & epidermal differentiation Plant Ce | | | |
| 12 | YAACKG | MYB2CONSAT | Transcriptional activators in abscisic acid signaling | Plant Cell (2003) | | |
| 13 | CNGTTR | MYBCORE | Transcriptional activators in abscisic acid signaling | Plant Cell (2003) | | |
| 14 | AACGG | MYBCORE | Regulate flavonoid biosynthesis in stress | EMBO (1995) | | |
| 15 | GGATA | MYBST1 | Cell cycle regulation | Plant Mol Biol (2002) | | |
| 16 | GGATA | MYBST1 | Function as a transcriptional activator | EMBO (1994) | | |
| 17 | CATGTG | MYCATERD1 | Function in dehydration stress and dark-induced senescence | Plant Cell (2004) | | |

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| 18 | GANTTNC | EECCRCAH1 | Regulate light & tissue specific expression | Plant Cell (1990) |
|----|----------|-------------|--|-----------------------------|
| 19 | CACATG | MYCATRD22 | Regulation of ABA-induced transcription | Plant Cell (1997) |
| 20 | CANNTG | MYCCONSAT | Phenylpropanoid biosynthesis genes | Plant Cell (2005 |
| 21 | ACTTTA | BBF1ARR | Tissue-specific expression & auxin induction | Plant Cell (1999) |
| 22 | AGAAA | POLLEN1AT52 | Play role as polyadenylation signal for plant genes | Nucleic Acids Res (1987) |
| 23 | ACTCAT | PREATPRODH | Pollen-specific transcription | Plant Mol Biol (1998) |
| 24 | CATGCA | RYREPEAT | Required for phytochrome regulation | Plant Cell (1996) |
| 25 | ATGGTA | S1FBOX | ABA-responsive complex for seed-specific expression | Plant Mol Biol (1999) |
| 26 | RTTTTTR | SEF4MOTIF | Developmental regulation of a house-keeping gene | Mol Cell Biol (1993) |
| 27 | GGGCC | SORLIP2AT | Nuclear factors interact with soybean embryo factor | Plant Cell (1989) |
| 28 | TTATCC | SREATMSD | Phytochrome A regulated expression, circadian regulation | Plant Physiol (2003) |
| 29 | GAGAC | SURECOREAT | Circadian regulation | Plant Physiol (2003) |
| 30 | TAAAG | TAAAGSTKST1 | Regulate gene expression in axillary bud outgrowth | Plant Physiol (2005) |
| 31 | TATTTAA | TATABOX | Regulation of a light-regulated gene | Plant Physiol (1995) |
| 32 | T-3ATATA | TATABOX | Binding site for OsTBP2 in pal promoter | Plant Cell (2002) |
| 33 | TATCCA | TATCCAOS | TATA and MYB binding site in GapC4 promoter | Biochem Biophys Acta (2001) |
| 34 | ATTC-3GC | TE2F2NTPCNA | Required for high-level expression of alpha-Amylase | Plant Cell Physiol (2001) |
| 35 | TGACGT | TGACGTV | Involved in transcriptional activation | Plant Journal (2002) |
| 36 | TGAC | WRKY | Phyto hormonal signalling GA & ABA | EMBO (1999) |
| 37 | TGACY | WBox | Transcription activation | Biochim Biophys Acta (2004) |
| 38 | CAAT | CAATBox | Act cooperatively with HSEs to increase promoter activity | Plant Physiol (2002) |
| 39 | GATA | GATABox | Required for light regulation and tissue specific expression | Plant cell (1990) |

Several promoters have been shown to drive fiber-specific gene expression in transgenic plant (Rinehart et al., 1996, Wu et al., 2009, Wu et al., 2007, Delaney et al., 2007, Li et al., 2002, Harmer et al., 2002, Botha et al 2011, Ni et al., 2008, Pyee and Kolattukudy 1995, Wang et al., 2004). Fiber characteristics have also been manipulated by coupling fiber-specific promoters to specific transgenes that may improve fiber quality (John and Keller 1996, Rinehart et al., 1996).

5. Transcription regulation

Promoters from some of the fiber specific genes have been isolated and their activities have been evaluated in transgenic plants. The conserve *cis*-regulatory elements help in transcription of genes in fiber cells (**Table 1**). Several of these promoters are proven to be applicable in genetic engineering of fiber. Trichomes in *Arabidopsis* and tobacco show a number of structural and genetic similarities to cotton fibers (Wang et al., 2004, Humphries et al., 2005).

Because promoter analysis in cotton is limited by the resources and extended timeline (minimum 12 months), the activity of cotton fiber-specific promoters has been also monitored in the heterologous model systems (Hsu et al., 1999, Liu et al., 2000, Wang et al., 2004, Wu et al., 2006).

The developing cotton fiber is an excellent model for unraveling the fundamental process of plant cell growth, differentiation, cell wall biogenesis and number of fiber mutants is available for exploring the regulatory mechanism of fiber growth and developmental processes. Recently much of work has been done for promoter characterization specific to fiber tissue in transgenic plants (**Table 2**).The promoter of *LTP6* gene was shown to drive expression of *gus* gene specifically in tobacco leaf trichome cells and may contain similar regulatory factors for tissue specific expression of gene in fiber cells (Hsu et al., 1999). Recently, the promoter of *GhTUB1* gene was shown to direct *gus* expression in transgenic cotton fiber cells (Li et al., 2002).

| Table 2: Some Trichome/Fiber s | necific promoter | expression stages and | I nutative function | in transgenic plants |
|--------------------------------|------------------|-------------------------|---------------------|----------------------|
| Table 2. Some Thenome/Ther s | pecific promoter | , expression stages and | i putative function | in transgeme plants |

| S.No. | Promoter | Expression stage | Putative function | References |
|-------|-----------|-------------------------------|----------------------------|-----------------------|
| 1 | GhLTP6 | Initiation & Elongation | Fiber morphogenesis | Hsu et al., 1999 |
| 2 | GhGlcAT1 | Fiber elongation | Cell wall synthesis | Wu et al., 2007 |
| 3 | GhACTIN1 | Elongation | Fiber elongation | Li et al., 2002 |
| 4 | GhE6 | Elongation | Fiber growth | John & Crow 1992 |
| 5 | FSltp4 | Fiber elongation | Synthesis of fiber cutin | Delaney et al., 2007 |
| 6 | GhLTPxGH3 | Initiation & Elongation | Fiber morphogenesis | Ma et al., 1995 |
| 7 | AtTSG | Initiation | Fiber morphogenesis | Ni et al., 2008 |
| 8 | GhMYB109 | Initiation- Elongation | Fiber formation | Wang et al, 2008 |
| 9 | GaRDL1 | Initiation | Fiber formation | Wang et al., 2004 |
| 10 | AtGL1 | Initiation | Fiber morphogenesis | Larkin et al., 1993 |
| 11 | FbL2A | Elongation and wall synthesis | Fiber growth | Rinehart et al., 1996 |
| 12 | GhLTP3 | Initiation & Elongation | Fiber morphogenesis | Liu et al., 2000 |
| 13 | GhTUB1 | Early stage | Fiber initiation | Li et al., 2002 |
| 14 | GhGlcAT1 | Elongation | Fiber development | Wu et al., 2007 |
| 15 | GhTUA9 | Elongation | Fiber growth & development | Li et al., 2007 |
| 16 | GaMYB2 | Initiation & elongation | Fiber development | Wang et al., 2004 |
| 17 | GhMYB25 | Initiation | Fiber formation and growth | Machado et al., 2009 |
| 18 | GaHOX | Initiation & Elongation | Fiber formation | Guan et al., 2008 |

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| 19 | GhCesA4 | Elongation & SCW | Cell wall morphogenesis | Wu et al., 2009 |
|----|---------|---------------------------|-------------------------|-------------------|
| 20 | GhSUS3 | Initiation | Fiber morphogenesis | Ruan et al., 2009 |
| 21 | GhRING1 | Initiation to SCW | Fiber formation | Ho et al., 2010 |
| 22 | GhXTH1 | Elongation | Fiber growth | Lee et al., 2010 |
| 23 | GhSCFP | Initiation and Elongation | Fiber development | Hou et al., 2008 |
| 24 | GhDET2 | Initiation & Elongation | Fiber formation | Luo et al., 2007 |
| 25 | GhGDSL | SCW | Cell wall biosynthesis | Yadav et al.,2017 |

The promoter of a cotton lipid transfer protein gene, FSltp4, was isolated and shown to direct fiber-specific transcription in cotton. In transgenic tobacco, this promoter was strongly active in foliar trichomes. Deletion analysis of the promoter identified an AT-rich 84 bp fiber specificregion (FSR) necessary for activity solely in the fiber cells. Cotton fiber proteins that bind the FSR were isolated using a yeast onehybrid assay (Delaney et al., 2007). There are several CesA genes which belongs to multi-gene family have been reported for primary and secondary cell wall cellulose synthesis in Arabidopsis (Fagard et al., 2000). In Comparison to these cotton homologs such as GhCesA1, GhCesA2, and GhCesA4, were expressed at high levels in cotton fibers during secondary cell wall synthesis while GhCesA3 gene expression was observed during both primary and secondary cell wall biogenesis stages (Kim and Triplett, 2001). GhCesA4 promoter activity was conferred by GUS activity in transgenic tobacco plant (Liu et al., 2000; Wu et al., 2007). In the study of GhGlcAT1 promoter (Wu et al., 2007) it was observed that it direct temporal, spatial and inducible expression of a reporter gene in transgenic tobacco and shows transcriptional regulation in transgenic plants. *Ltp6* and *Ltp3* promoter activity were conferred in transgenic tobacco (Ma et al., 1997). The transgenic plants demonstrated highly specific gus expression in the leaf and stem trichomes, with some expression in the vascular tissue and leaf margin. Deletion assay were performed to find out the promoter regions necessary for trichome specific gene expression (Hsu et al., 1999, Liu et al., 2000) and with removal of the basal promoter elements CAAT and TATA boxes, Ltp6 promoter resulted in a progressive reduction in GUS activity therefore basal promoter activity ceased in transgenic plant (Hsu et al., 1999). Recent experiment on RDL1 promoter provide envisage that cotton and Arabidopsis use similar transcription factors for regulating trichomes and GaMYB2 may be a key regulator of cotton fiber development (Wang et al 2004). Thus, promoter regulatory regions and interacting factors may be employed for the improvement of cotton fiber trait.

6. Conclusion

Cotton fiber is one of the important commercial crops worldwide and its development takes place in the stage specific manner. The regulation of cotton fiber developmental stages is very important for fiber quality parameters. Promoter is crucial region of gene which plays an important role in the fiber-specificity and regulation of timing of expression of genes. The promoters of these genes are of prime interest because they control transcription in the fiber cell and may be used in the genetic manipulation of fiber quality. Promoter allows the expression of transgenes targeted to the fibers and avoid detrimental effects on growth and phenotype elsewhere within the plant. This will help in developing new strategies to better utilize the promoter for the regulation of gene expression during fiber developmental processes and ultimately in improving the cotton fiber quality and yield.

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