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# Organization of cellulose synthase complexes involved in primary cell wall synthesis in *Arabidopsis thaliana*

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In all land plants, cellulose is synthesized from hexameric plasma membrane complexes. Indirect evidence suggests that in vascular plants the complexes involved in primary wall synthesis contain three distinct cellulose synthase catalytic subunits (CESAs). In this study, we show that CESA3 and CESA6 fused to GFP are expressed in the same cells and at the same time in the hypocotyl of etiolated seedlings and migrate with comparable velocities along linear trajectories at the cell surface. We also show that CESA3 and CESA6 can be coimmunoprecipitated from detergent-solubilized extracts, their protein levels decrease in mutants for either CESA3, CESA6, or CESA1 and CESA3, CESA6 and also CESA1 can physically interact *in vivo* as shown by bimolecular fluorescence complementation. We also demonstrate that CESA6-related CESA5 and CESA2 are partially, but not completely, redundant with CESA6 and most likely compete with CESA6 for the same position in the cellulose synthesis complex. Using promoter- $\beta$ -glucuronidase fusions we show that CESA5, CESA6, and CESA2 have distinct overlapping expression patterns in hypocotyl and root corresponding to different stages of cellular development. Together, these data provide evidence for the existence of binding sites for three distinct CESA subunits in primary wall cellulose synthase complexes, with two positions being invariably occupied by CESA1 and CESA3, whereas at least three isoforms compete for the third position. Participation of the latter three isoforms might fine-tune the CESA complexes for the deposition of microfibrils at distinct cellular growth stages.

Cellulose microfibrils are synthesized from a multiprotein complex inserted into the plasma membrane. These “rosette” complexes consist of six globules, each of which contains multiple cellulose synthase catalytic subunits (CESAs). These complexes migrate in the plasma membrane along microtubules, propelled by the polymerization of the  $\beta$ -1,4-glucan chains (1).

Plant CESA genes are members of multigene families. *Arabidopsis* has 10 CESA isoforms that, based on sequence comparison with other plant species, can be classified into six orthologous groups (2). Mutational analysis shows that these six groups of isoforms have nonredundant functions in cellulose synthesis. Mutants for three isoforms (CESA4, CESA7, and CESA8) show defects in cellulose synthesis specifically in secondary walls (3–5). Microarray data show that the mRNAs for the three genes are coregulated (6, 7). The three proteins are expressed in the same cell types during secondary cell wall deposition, and co-immunoprecipitation (IP) experiments show that all three proteins interact (3). Although the interactions remain to be validated *in vivo*, these data strongly suggest that at least in these cells the complexes contain three isoforms. Mutants for isoforms CESA1, CESA3, and CESA6 have cellulose defects in primary cell walls (8–11). The three genes are also coregulated at the mRNA level (12). It is not known, however, whether the corresponding proteins are expressed in the same cells at the same time. It is also not known whether the three CESAs are

present in different populations of complexes or are part of the same complex. Interestingly, a class of mutations in the C terminus of either CESA3 (*cesa3<sup>ixr1</sup>*) or CESA6 (*cesa6<sup>ixr2</sup>*) (isoxaben-resistant) confers increased resistance to the cellulose inhibitor isoxaben (13, 14). The simplest explanation for the existence of two nonredundant resistance loci is that isoxaben recognizes an epitope associated with the CESA3- and CESA6-containing complex. Null mutants for CESA1 and CESA3 are gametophytic lethal (39), indicating the essential nature of the genes. In contrast, CESA6 null mutants show a relatively mild phenotype, which might be explained by the existence of CESA2, CESA5, and CESA9, which are closely related to CESA6 (12). Partial redundancy between CESA6-related isoforms also might explain the lower isoxaben resistance conferred by *cesa6<sup>ixr2</sup>* compared with *cesa3<sup>ixr1</sup>*. In this study, we initially focused on the isoxaben targets CESA3 and CESA6. We raised specific antibodies against both proteins and constructed functional GFP fusions. We show that CESA3 and CESA6 are expressed in the same cells at the same time and migrate with comparable velocities along linear trajectories at the cell surface during early dark-grown seedling development. The levels of both proteins decrease in mutants for either CESA1, CESA3, or CESA6. Using co-IP experiments, we further show that at least CESA3 and CESA6 interact with each other, and using bimolecular fluorescence complementation (BiFC) experiments, we show that CESA1, CESA3, and CESA6 can interact *in vivo*. Through the analysis of single, double, and triple mutants, we also show that CESA5 and CESA2 are partially redundant with CESA6 and most likely compete with CESA6 for the same binding site in the complex. Finally, we show that CESA6-related isoforms display distinct expression patterns during cellular differentiation.

## Results

**Simultaneous Accumulation and Similar Subcellular Localization of CESA3 and CESA6 in Developing Dark-Grown Seedlings.** The existence of two nonredundant isoxaben-resistant loci suggested that CESA3 and CESA6 are part of the same complex. To investigate this idea,

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The authors declare no conflict of interest.

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Abbreviations: CESA, cellulose synthase catalytic subunit; BiFC, bimolecular fluorescence complementation; IP, immunoprecipitation; YFP, yellow fluorescent protein; GUS,  $\beta$ -glucuronidase.

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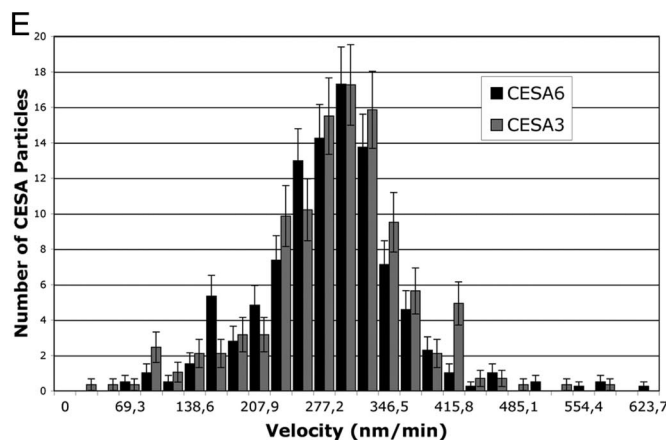
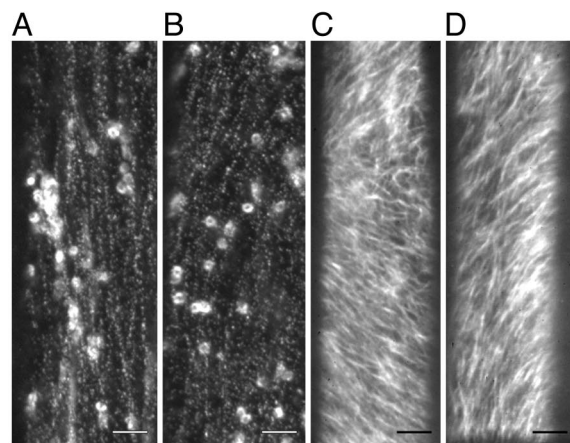
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we raised isoform-specific antibodies against the N-terminal predicted cytoplasmic domain of each protein. The antibodies were enriched by using a two-step immunopurification procedure, and their specificity was shown on immunoblots with the corresponding N-terminal fragments of CESA3, CESA6, and CESA1 produced in *Escherichia coli* (see *Methods*). The absence of cross-reactivity was also shown in the co-IP experiments described below. It is not excluded, however, that anti-CESA6 antibodies cross-reacted with more closely related isoforms CESA2, CESA5, or CESA9.

CESA3 and CESA6 were undetectable in imbibed seeds and appeared after germination [supporting information (SI) Fig. 5]. The immunostained bands showed a size of  $\approx 120$  kDa, which is the predicted molecular mass for the CESA proteins. The amount of both proteins increased simultaneously in growing seedlings to reach a maximum at 96 h postimbibition, suggesting a coregulation at the protein level in developing seedlings.

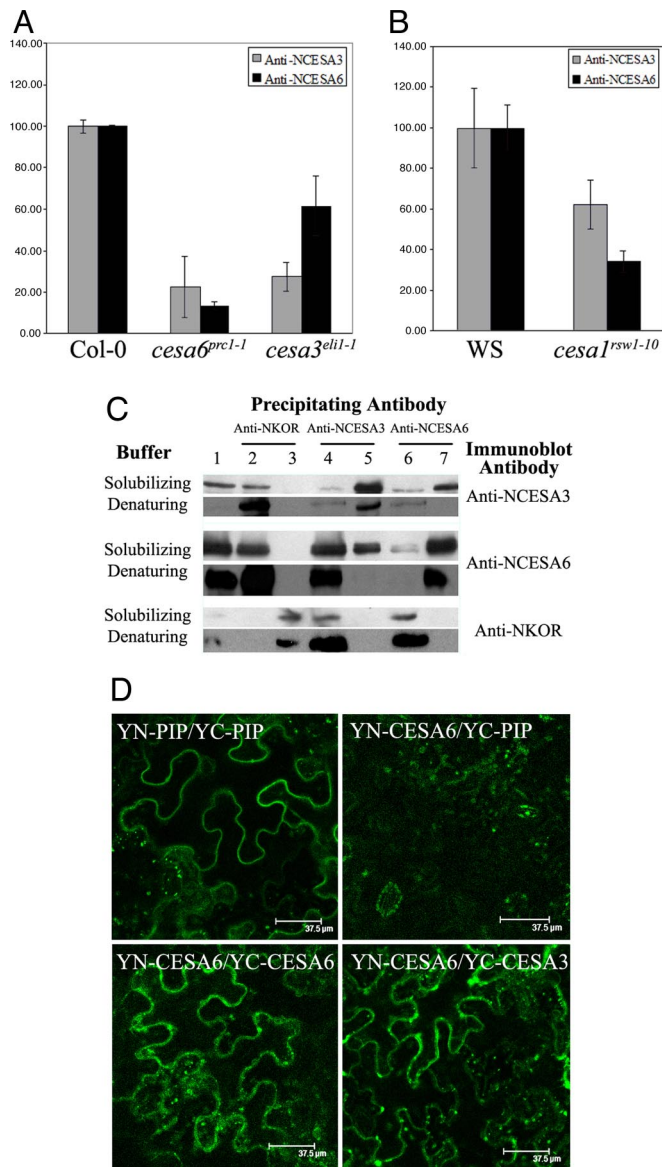
We next investigated the cellular and subcellular localization of CESA3 and CESA6. GFP-CESA3 and GFP-CESA6 were expressed from their own promoters in the respective *cesa3<sup>3le5</sup>* and *cesa6<sup>prc1-1</sup>* mutant backgrounds. These constructs were biologically functional as shown by the complementation of the short hypocotyl phenotype of the mutants (SI Fig. 6). Both GFP-CESA3 and GFP-CESA6 could be observed in epidermal and cortical cells of 3-day-old dark-grown hypocotyls (Fig. 1 *A–D*). In addition, GFP-CESA3 and GFP-CESA6 fluorescence showed a similar subcellular distribution. Both proteins accumulated in similar intracellular compartments and formed rows of fluorescent particles at the cell surface as described for yellow fluorescent protein (YFP)-CESA6 (1). Among these compartments one could recognize 1.3- $\mu$ m doughnut-shaped compartments as described for YFP-CESA6 (1) and smaller highly motile compartments. Time-lapse spinning disk microscopy showed bidirectional migration of surface particles for both fusion proteins. The average velocity of the particles in elongating epidermal cells was 272 nm/min (range: 53 to 622 nm/min) and 277 nm/min (range: 40 to 566 nm/min) for GFP-CESA6 and GFP-CESA3, respectively (Fig. 1*E*). These velocities are similar to those reported for YFP-CESA6 (1, 15).

**CESA3 and CESA6 Protein Levels Are Reduced in *cesa1*, *cesa3*, and *cesa6* Mutants.** If distinct CESA isoforms are present in the same complex, one can expect that reducing the protein levels of one isoform will also affect the levels of the other isoforms. To investigate this notion, we quantified the relative abundance of CESA3 and CESA6 in 4-day-old dark-grown seedlings in *cesa1*, *cesa3*, and *cesa6* mutants. *cesa6<sup>prc1-1</sup>* is a null allele (10), whereas *cesa3<sup>eli1-1</sup>* (11) and *cesa1<sup>rsw1-10</sup>* (16) are both leaky alleles. All three mutants show a short hypocotyl and are deficient for cellulose in their primary cell walls (SI Fig. 7) (17). CESA3 and CESA6 proteins accumulated to similar levels either in Col-0 or WS backgrounds (Fig. 2 *A* and *B*). As expected, in *cesa6<sup>prc1-1</sup>* the anti-CESA6 antibody detected only a few percent of the wild-type levels (Fig. 2*A*). The residual signal may be related to cross-reaction with related CESA subunits, such as CESA2 or CESA5. Interestingly, CESA3 protein levels were also significantly reduced in this mutant. In a similar way, in *cesa3<sup>eli1-1</sup>* not only the level of CESA3 was reduced but also that of CESA6 (Fig. 2*A*). The less severe reduction of CESA6 compared with CESA3 in this mutant may be related to the altered properties of the remaining mutant CESA3 protein. Because we did not dispose of specific antibodies for CESA1, we could not analyze directly the CESA1 protein levels in the three mutants. However, in *cesa1<sup>rsw1-10</sup>* we observed that levels of CESA3 and CESA6 were also diminished (Fig. 2*B*). Together, these data show that the normal accumulation of each of the CESA3 and CESA6 proteins requires the presence of normal levels of both CESA1, CESA3, and CESA6, which is consistent with the presence of these three proteins in the same complex.



**Fig. 1.** Visualization of similar distribution and dynamics of GFP-CESA3 and GFP-CESA6. (*A–D*) Functional GFP-CESA3 and GFP-CESA6 fusion proteins expressed, respectively, in mutant backgrounds *cesa3<sup>3le5</sup>* and *cesa6<sup>prc1-1</sup>* label similar intracellular compartments and complexes that migrate with comparable velocities at the cell surface. Spinning disk microscope images (*A* and *B*) and projection of a time series (10 min, 30 s, *C*; 10 min, *D*) of epidermal cells at the top of a dark-grown hypocotyl expressing GFP-CESA3 (*A* and *C*) or GFP-CESA6 (*B* and *D*). (Scale bars: 4  $\mu$ m.) (*E*) Distribution of the velocities of the surface particles.

**CESA3 Coimmunoprecipitates with CESA6 but Not with KOR1.** To investigate whether CESA3 and CESA6 physically interact, we carried out co-IP experiments on detergent-solubilized protein extracts from 4-day-old dark-grown seedlings (Fig. 2*C*). As expected, anti-CESA3 antibodies precipitated CESA3 both in non-denaturing and denaturing conditions (Fig. 2*C*, lane 5). Interestingly, CESA6 also coimmunoprecipitated with CESA3 in non-denaturing conditions, whereas no CESA6 signal was observed in denaturing conditions. The latter observation also confirms the absence of cross-reactivity of anti-CESA6 with CESA3. Similar results were obtained with the reciprocal experiment in which anti-CESA6 antibodies were used for the co-IP. Again, CESA6 precipitated in both non-denaturing and denaturing conditions, whereas CESA3 only coprecipitated in non-denaturing conditions. These results show that a stable interaction existed between CESA3 and CESA6 in Triton X-100-solubilized extracts and that this interaction was disrupted in denaturing buffer. None of the CESAs were immunodetected in the absence of the primary antibody (CESA3 or CESA6) (data not shown). Using the anti-CESA6 antibody, we observed a second band on the immunoblot. This band may correspond to a truncated CESA6 isoform or a CESA6-related isoform. We also investigated the interaction of CESA3 and CESA6



**Fig. 2.** CESA3 and CESA6 are in the same cellulose synthase complex. (A and B) CESA3 and CESA6 protein levels, measured in dark-grown seedlings by quantitative immunoblotting, are reduced in mutants *cesa6<sup>prc1-1</sup>*, *cesa3<sup>eli1-1</sup>* (A) and *cesa1<sup>rsw1-10</sup>* (B). Protein levels are expressed as percentage of wild-type levels, and error bars are SDs of three biological repeats. (C) CESA3 coimmunoprecipitates with CESA6 and vice versa, but not with membrane-bound cellulase KOR1. IP was carried out in nondenaturing or denaturing conditions. Shown are total Triton X-100-solubilized extracts (lane 1), IP with anti-NKOR, unbound (lane 2) and bound (lane 3) fractions; with anti-NCESA3, unbound (lane 4) and bound (lane 5) fractions or with anti-NCESA6, unbound (lane 6) and bound (lane 7) fractions. Immunoblotting was carried out with anti-NCESA3 anti-NCESA6 or anti-NKOR antibodies. (D) BiFC in *N. benthamiana* leaf epidermal cells shows *in vivo* CESA6 homodimer and CESA3/CESA6 heterodimer formation. (Upper Left) YN-PIP/YC-PIP-positive control. (Upper Right) YN-CESA6/YC-PIP. (Lower Left) YN-CESA6/YC-CESA6. (Lower Right) YN-CESA6/YC-CESA3. Other CESA combinations are shown in SI Fig. 8.

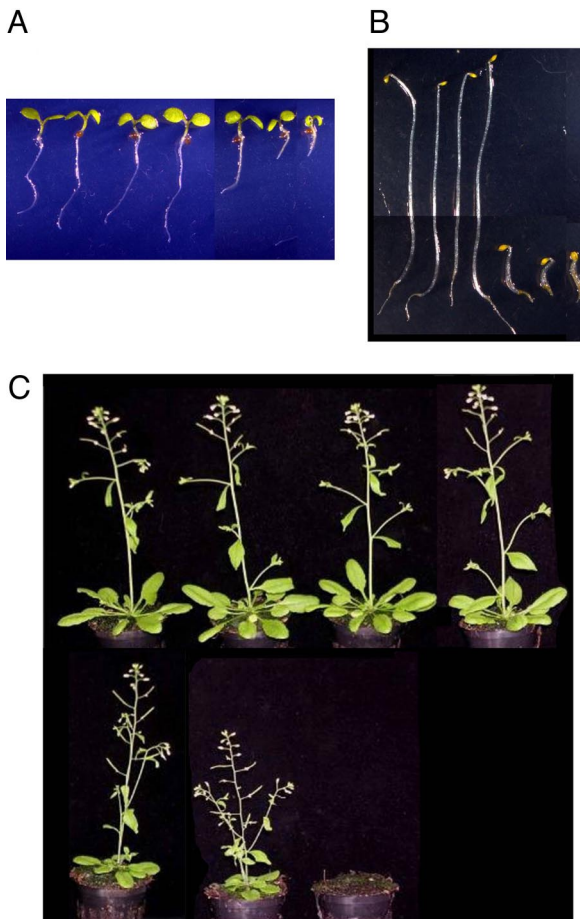
with the membrane-bound cellulase KOR1, which is also required for normal cellulose synthesis (18). KOR1 protein was detected only in the unbound fraction and not in the immunoprecipitates with anti-CESA3 or CESA6 antibodies. These results indicate that KOR1 did not stably interact with CESA3 nor with CESA6 in detergent-solubilized extracts and underscore the specificity of the observed CESA3–CESA6 interaction.

**Visualization of CESA Homodimers and Heterodimers in *Planta* Using BiFC.** To confirm the co-IP results, we studied the capacity of CESA isoforms to interact *in vivo* by using the BiFC technique (19). The N-terminal and the C-terminal fragments of YFP were both fused to the N terminus of the coding sequences of CESA1, CESA3, and CESA6. As shown in SI Fig. 6, GFP fusions at this position do not interfere with the functionality of at least CESA3 and CESA6. Combinations of chimeric genes expressed from the cauliflower mosaic virus 35S promoter were transiently expressed in *Nicotiana benthamiana* leaves using *Agrobacterium* infiltration. As a positive control, we used the aquaporin PIP2–1 (20). Aquaporins are known to form homotetramers in the plasma membrane (21). As expected, coexpression of YN-PIP2–1 and YC-PIP2–1 yielded a YFP signal (Fig. 2D), whereas no signal was detected by expression of either YN-PIP2–1 or YC-PIP2–1 alone (data not shown). The fluorescence emission spectrum of the signal indeed confirmed that the signal was derived from a functional YFP (SI Fig. 8B). We next investigated whether the fusion proteins derived from the three CESA proteins could homodimerize. As shown in Fig. 2D and SI Fig. 8, a fluorescent signal was observed for YN-CESA3/YC-CESA3 and YN-CESA6/YC-CESA6 combinations. Fluorescent signals were observed both at the cell surface and in intracellular compartments, similar to that of CESA3 or CESA6 fused to intact YFP (SI Fig. 8C). For YN-CESA1/YC-CESA3 and YN-CESA3/YC-CESA3, the signal was always weaker, suggesting that this protein homodimerizes less efficiently (SI Fig. 8E). We also analyzed the YN-CESA1/YC-CESA1 combination, which also reconstituted the YFP fluorescence (SI Fig. 8D). For all three CESA fusions, no fluorescent signal was detected by expression of either YN-CESA or YC-CESA alone (data not shown).

To investigate whether heterodimers could also be formed, we coexpressed YN-CESA3/YC-CESA1, YN-CESA1/YC-CESA3, YN-CESA6/YC-CESA1, YN-CESA1/YC-CESA6, YN-CESA3/YC-CESA6, or YN-CESA6/YC-CESA3 (SI Fig. 8 and Fig. 2D). We observed that YFP fluorescence was reconstituted for all of the combinations, suggesting that all three isoforms can form a complex. As a negative control, we coexpressed PIP2–1 chimerae with corresponding CESA constructs (Fig. 2D). No signal was detected with combinations of either YN-PIP2–1 or YC-PIP2–1 for all of the CESA fusions tested, suggesting that PIP2–1 and CESA proteins were unable to interact despite the presence of both proteins in the plasma membrane. In conclusion, all three isoforms showed interactions *in vivo*, confirming the co-IP results for CESA3 and CESA6.

**Partial Redundancy Among CESA6-Related CESA Isoforms.** The mild phenotype of *CESA6* null alleles as compared with strong mutant alleles of *CESA1* or *CESA3* and the lower resistance to isoxaben of *cesa6<sup>ivr2-1</sup>* mutant than *cesa3<sup>ivr1-1</sup>* or *cesa3<sup>ivr1-2</sup>* (13, 14) might be caused by partial redundancy with other CESA isoforms.

CESA2, CESA5, and CESA9 are closely related to CESA6. *CESA9* is only expressed during embryogenesis, as shown by public microarray data and promoter- $\beta$ -glucuronidase (GUS) fusions (data not shown), and was not considered in this study. We first studied single T-DNA insertion mutants for *CESA2* or *CESA5*. Mutant light-grown seedlings and adult plants were indistinguishable from the wild type (Fig. 3A and C). However, the dark-grown hypocotyl of *cesa2* was slightly shorter than that of the wild type (Fig. 3B and Table 1). The analysis of the cell wall of dark-grown mutant hypocotyls with FTIR microspectroscopy failed to detect differences with the wild type as shown by the tight clustering of the mutants with wild-type controls (SI Fig. 7). The FTIR spectra of double *cesa2/cesa5* mutants also clustered with those of the wild type. Interestingly, compared with the single *cesa2* mutant, the dark-grown hypocotyl of the double mutant was slightly longer. In addition, roots of light-grown *cesa2/cesa5* seedlings were slightly longer, and adult plants were slightly larger than the wild type (Fig. 3A and C). In contrast, mutant combinations with *cesa6* showed a very different picture. *cesa2/cesa6* double mutants showed an



**Fig. 3.** Partial redundancy between CESA6-related CESAs. Five-day-old light-grown (A), dark-grown seedlings (B), and 32-day-old greenhouse-grown plants (C) of (from left to right) Col-0, *cesa2*, *cesa5*, *cesa2/cesa5*, *cesa2/cesa6<sup>prc1-1</sup>*, and *cesa5/cesa6<sup>prc1-1</sup>* are shown.

enhanced phenotype compared with *cesa6* as shown by the reduced hypocotyl length in the dark, the reduced root length in the light, and the strongly dwarfed and bushy adult *cesa2/cesa6* plants (Fig. 3). *cesa5/cesa6* double homozygotes were seedling lethal. Interestingly, the *cesa5* allele showed a strong dosage effect in a homozygous *cesa6* background as shown by the 1:2:1 segregation of the dark-grown hypocotyl into three size classes within the F<sub>2</sub> population of a *cesa6<sup>-/-</sup>;cesa5<sup>+/-</sup>* heterozygote (SI Fig. 9). To construct the triple mutant, we crossed *cesa2<sup>-/-</sup>cesa5<sup>-/-</sup>* with *cesa2<sup>-/-</sup>cesa6<sup>-/-</sup>*. Among the *cesa2<sup>-/-</sup>cesa6<sup>-/-</sup>* F<sub>2</sub> seedlings only the *cesa5<sup>+/-</sup>* seedlings were viable. The *cesa5<sup>+/-</sup>* or *cesa5<sup>-/-</sup>* seedlings either bleached and died or survived as very small sterile plants (data not shown).

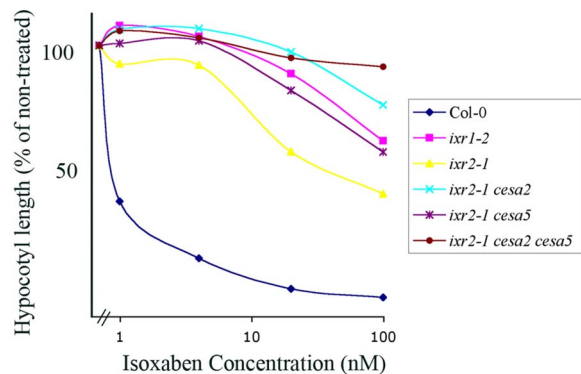
**CESA6-Related CESAs Are Targets for Isoxaben.** We next investigated whether the lower isoxaben resistance of *cesa6<sup>ixr2-1</sup>* compared with

**Table 1. Hypocotyl lengths of 5-day-old dark-grown seedlings**

Type	Mean	SD	n	Student's t test*
Col-0	16.26	1.19	67	
<i>cesa2</i>	14.24	1.21	57	9.33 <sup>†</sup>
<i>cesa5</i>	15.39	1.06	70	1.67
<i>cesa6</i>	3.41	0.34	69	85.6 <sup>†</sup>
<i>cesa2/cesa5</i>	16.95	1.34	69	3.18 <sup>†</sup>
<i>cesa2/cesa6</i>	2.12	0.18	71	96 <sup>†</sup>

\*All comparisons were made to Col-0.

<sup>†</sup>Significantly different;  $P = 0.01$ .



Isoxaben concentration	Col-0		<i>ixr1-2</i>		<i>ixr2-1</i>	
	mean	SD	mean	SD	mean	SD
0 nM	100	8,99	100	22,29	100	23,06
100 nM	6,87	1,07	64,84	16,54	45,17	12,97

Isoxaben concentration	<i>ixr2-1 cesa2</i>		<i>ixr2-1 cesa5</i>		<i>ixr2-1 cesa2 cesa5</i>	
	mean	SD	mean	SD	mean	SD
0 nM	100	17,89	100	12,47	100	15,74
100 nM	78,02	14,54	60,63	11,23	92,2	13,13

**Fig. 4.** CESA6, CESA2, and CESA5 are targets for isoxaben. (Upper) The effect of isoxaben on hypocotyl length: dose-response curve. Hypocotyl length is expressed in percentage of the untreated control. Average values of  $\approx 35$  individuals are shown. (Lower) For clarity, SDs of untreated and 100 nM isoxaben-treated seedlings are shown.

*cesa3<sup>ixr1-2</sup>* was caused by the presence of CESA6-related proteins in the cellulose synthase complexes of *cesa6<sup>ixr2-1</sup>* plants. To this end, we quantified the resistance to isoxaben of all double and triple mutant combinations of *cesa2* and *cesa5* with *cesa6<sup>ixr2-1</sup>* (Fig. 4) by measuring the hypocotyl length of 5-day-old dark-grown seedlings on increasing concentrations of isoxaben. As observed previously, *cesa6<sup>ixr2-1</sup>* was  $\approx 50$ -fold more resistant to isoxaben than the wild type, and *cesa3<sup>ixr1-2</sup>* was 5-fold more resistant than *cesa6<sup>ixr2-1</sup>*. Interestingly, adding *cesa5* increased the resistance of *cesa6<sup>ixr2-1</sup>* to that of *cesa3<sup>ixr1-2</sup>*, adding *cesa2* caused a  $>10$ -fold increase in resistance, and adding both *cesa2* and *cesa5* rendered the seedling insensitive to 100 nM isoxaben. The increased resistance was not caused by increased CESA3 or CESA6 protein levels in the mutants (data not shown). Together, these data show that all three CESA6-related isoforms are targets for isoxaben, suggesting that they compete for the same binding site within the cellulose synthase complex.

**Distinct Expression Patterns of CESA6-Related Isoforms During Cellular Differentiation.** Using promoter-GUS fusions we studied the expression patterns of CESA6, CESA2, and CESA5 genes in dark-grown hypocotyls and roots of 72-h-old seedlings (SI Fig. 10A). At this stage, different phases of cell expansion are represented from the top to the base of the hypocotyl or from the tip to the base of the root. Promoter CESA5-GUS was weakly expressed only in the unexpanded cells in the apical hook of the hypocotyl. Promoter CESA6-GUS expression occurred throughout the hypocotyl and root, peaking in the growth acceleration zone just below the apical hook and in the cell elongation zone in the root. Promoter CESA2-GUS expression largely overlapped with that of promoter CESA6-GUS, but in the root, peaked in more mature cells. The GUS expression patterns in the root were consistent with the microarray data for CESA2, CESA5, and CESA6 on flow-sorted root protoplasts (22). In conclusion, the three partially redundant CESA genes mark distinct cellular growth stages in hypocotyl and root.

**CESA5 Is Only Partially Redundant with CESA6.** We showed that CESA5 is partially redundant with CESA6 and most likely competes for the same site in the complex. The incomplete redundancy

can be caused by the partially overlapping expression patterns or intrinsic differences in the protein structure and function. To investigate this, we expressed *CESA5* from the *CESA6* promoter in a *cesa6<sup>prc1-1</sup>* null background. Three independent transformants showed incomplete restoration of the hypocotyl growth defect (SI Fig. 10B). On the contrary, expressing *CESA6* from the same promoter sequence completely restored the wild-type phenotype to *cesa6<sup>prc1-1</sup>* (data not shown). The results show that *CESA5* cannot entirely substitute for *CESA6*, suggesting that both isoforms have specialized roles in the deposition of cellulose in expanding cells.

## Discussion

Several lines of evidence presented in this article show that *CESA3*, *CESA6*, and most likely *CESA1*, are in the same protein complex in dark-grown seedlings. This idea was already suggested by the coexpression of transcripts for *CESA1*, *CESA3*, and *CESA6* (12, 23), the existence of isoxaben-resistant alleles for both *CESA3* and *CESA6* (13, 14), and the observation that the complexes involved in secondary wall synthesis also contain three nonredundant *CESA* isoforms (3). Here, we showed that *CESA3* and *CESA6* proteins follow the same accumulation kinetics in developing seedlings, functional GFP-fusion proteins are expressed in the same cells and occupy similar subcellular compartments, and the GFP-fusion proteins migrate at the cell surface with comparable velocities. Although we did not observe *CESA1* directly, the coexpression at the mRNA level strongly suggests that *CESA1* is also expressed in the same cells. We further showed that mutations in either *CESA1*, *CESA3*, or *CESA6* cause a decrease in protein levels of *CESA3* and *CESA6*, which is compatible with the idea that the assembly of the complex is prevented in the absence of one component, and that *CESA3* and *CESA6* coimmunoprecipitate with each other but not with *KOR1*. It remains possible however, that the antibodies coimmunoprecipitated not only the *CESA* complex but also detergent-resistant membrane (DRM) fractions that contain other membrane proteins. This is unlikely the case because *KOR1*, which also has been detected in DRMs (24), did not coimmunoprecipitate with *CESA3* or *CESA6*. Similar observations have been reported for the secondary cell wall *CESAs* (3, 25). Finally, using BiFC we show that *CESA3*, *CESA6*, and *CESA1* can interact *in vivo*.

One might question the validity of the BiFC technique for the study of the interaction of membrane proteins because false positives might arise as a result of proximity of noninteracting proteins confined within membranes. However, highly specific interactions between membrane proteins have been demonstrated using this technique in other systems (26–28). Also, the absence of an interaction with the plasma membrane protein PIP2-1 makes this possibility unlikely. Although we tested only pairwise combinations of *CESA* isoforms, a positive signal does not exclude indirect interactions, perhaps through the formation of complexes with endogenous tobacco proteins. Collectively, these results provide strong evidence for the presence of three *CESA* proteins in the same complex.

The strong phenotypes of *cesa1* and *cesa3* mutants demonstrate the absence of redundancy with other *CESAs*. We show in this article that the relatively mild phenotype of *cesa6* instead reflects partial redundancy with two related isoforms, *CESA2* and *CESA5*, which presumably can replace *CESA6* in the complex. We also show that all three *CESA6*-related isoforms are isoxaben targets, which explains the lower resistance of *cesa6<sup>ivr2-1</sup>* to isoxaben compared with *cesa3<sup>ivr1</sup>*, because *CESA3* does not compete with other isoforms for its position in the complex.

The different expression patterns of *CESA5*, *CESA6*, and *CESA2* during hypocotyl and root growth suggests that the composition of the cellulose synthase complex changes at different stages of cellular development. The partial complementation of the *cesa6<sup>prc1-1</sup>* phenotype, through the expression of *CESA5* under control of the *CESA6* promoter, also shows that the isoforms are functionally specialized. Similar observations have been made for

*CESA2* and *CESA6* (39). We previously showed that distinct stages of cellular development can be distinguished from the top to the base of the dark-grown hypocotyl (29). It is conceivable that properties of the cellulose synthase complexes change to cope with varying constraints associated with cellulose deposition at successive growth stages. It will be interesting to see to what extent the three *CESA6*-related isoforms confer distinct properties to the terminal complexes, such as the velocity and the density of the complexes, the orientation of the trajectories, interaction with the cytoskeleton, and coupling to cell elongation.

## Methods

**Plant Material, *In Vitro* Growth Conditions, and Genetic Analysis.** *Arabidopsis* wild-type Col-0 and mutant lines were grown as described (30) at 20°C without sucrose. See SI Table 2 for mutants and T-DNA insertion lines used in this study. Plants were grown in the dark as described (29) or in a 16-h light and 8-h dark cycle. Homozygous lines for *cesa2* and *cesa5* were screened for the simultaneous presence of a PCR product amplified from the left T-DNA border and absence of a PCR product from the genomic sequence using primers flanking the T-DNA insertion (SI Table 3). Double and triple *cesa* mutants were genotyped by PCR. Promoter-GUS lines for *CESA2*, *CESA5*, and *CESA6* (described in ref. 31) were kindly provided by M. Doblin (University of Melbourne, Melbourne, Australia). The length of hypocotyls fixed with 0.2% formaldehyde were measured by using Optimas 4.1 software as described (29).

**Production of Polyclonal Antibodies Against *CESA3* and *CESA6*.** The N-terminal regions upstream of the first transmembrane domains of *CESA3* and *CESA6* were amplified (primers in SI Table 3) and cloned into the EcoRI site of pGEX3X and pGEX2T (Amersham Biosciences, Piscataway, NJ), respectively. GST-fusion proteins were purified on GHS-Sepharose 4B resin according to the manufacturer's recommendations (Amersham Biosciences). Rabbit antibodies against the GST-fusion polypeptides were produced by BioGenex (Berlin, Germany). GST-fusion proteins were coupled to CNBr-activated Sepharose 4F according to the supplier's protocol (Amersham Biosciences). The serum was incubated with the antigen-Sepharose matrix, and the unbound fraction was extensively washed from the resin according to the manufacturer's recommendations. Antibodies were eluted with 0.2 M glycine (pH 2), 0.15 M NaCl and neutralized with 1.5 M Tris-HCl, pH 8.8. To further improve the specificity of the sera, the pooled fractions eluted from the *CESA3*-Sepharose resin were loaded on the *CESA6*-Sepharose resin, and the unbound fraction on this matrix was used as purified *CESA3* antibody. The antibodies raised against *CESA6* were prepared in the same manner. The specificity of the *CESA3* and *CESA6* antibodies was verified by immunoblotting against the GST-fusion proteins (SI Fig. 11A).

**Quantification of Protein Levels.** Samples corresponding to 50 dark-grown seedlings were collected at different times and ground in 50  $\mu$ l of extraction buffer (4 M urea and 100 mM DTT). Twenty-five microliters of modified Laemmli buffer (without bromophenol blue) was added, and the samples were boiled for 5 min and centrifuged at 10,000  $\times$  g for 10 min at room temperature. Protein content of the supernatant was quantified by using the RC DC Protein Assay method (Bio-Rad, Hercules, CA). Aliquots of 20  $\mu$ g protein were then analyzed by 8% SDS/PAGE (32) and immunoblotted according to standard protocols. The purified anti-*CESA* sera were used in 1:1,000 dilution, and the signals were measured by fluorescent detection of AlexaFluor 488 goat anti-rabbit (used at 1/3,000; Molecular Probes–Invitrogen, Carlsbad, CA) and quantified with ImageGauge software (Fuji, Tokyo, Japan). Fifty *Arabidopsis* seedlings grown in liquid medium (30) for 48 h in the dark were labeled with a mixture of <sup>35</sup>S-labeled methionine and [<sup>35</sup>S]cysteine (0.2 MBq; PerkinElmer Life Science, Albany, NY) for 4 h.

Seedlings were then extensively washed and used for IP. Radiolabeled proteins were separated by 8% SDS/PAGE (32) and visualized by phosphorimaging (Fuji FLA 5000).

**Co-IP Experiments.** Two grams of 3-day-old dark-grown seedlings was ground in liquid nitrogen, and the powder was transferred to either 6 ml of solubilization buffer [50 mM Tris·HCl, pH 8/150 mM NaCl/2% Triton X-100, containing 1% (wt/vol) polyvinylpyrrolidone (Polyclar AT), plant protease inhibitor mixture and phosphatase inhibitor mixture 1 and 2 (Sigma, St. Louis, MO) used, respectively, at a 100- and 200-fold dilution] or denaturing buffer [50 mM Tris·HCl, pH 8/150 mM NaCl/1% Nonidet P-40/0.5% deoxycholate/0.1% SDS, containing 1% (wt/vol) polyvinylpyrrolidone (Polyclar AT) and the mixture of inhibitors]. A cleared cell lysate was obtained by centrifugation at  $8,000 \times g$  for 10 min and filtration of the supernatant through Miracloth. An aliquot of 2 mg of protein was supplemented with purified polyclonal antisera at 1:50 dilutions, and IPs were carried out essentially according to established protocols (33). Immunoprecipitated proteins bound on Protein A Sepharose CL-4B beads (Amersham Biosciences) were boiled in  $2\times$  Laemmli buffer, and proteins were analyzed by 8% SDS/PAGE (32) and immunoblotted using the ECL Luminescent Detection System (Amersham Biosciences). Autoradiography of the proteins precipitated in denaturing conditions with either anti-CESA6 or anti-KOR1 out of extracts from  $^{35}\text{S}$ -labeled seedlings confirmed the specificity of the respective antibodies for proteins of the expected molecular masses (respectively, 120 and 72 kDa; SI Fig. 11B) also when used in IP experiments.

**Plant Expression Vectors.** For plant expression vectors see SI Text.

**Transient Expression in *N. benthamiana*.** Leaves of 3-week-old plants were transformed by infiltration as described (34). YFP fluorescence was detected 3 days after infiltration by using the 514-nm laser line of a SP2 AOBS confocal laser scanning microscope (Leica, Solms, Germany,) equipped with an argon laser. To check the YFP reconstitution, spectral analysis was performed with the 496-nm laser line. For BiFC experiments, all constructs fluorescence was detected at the same photo-multiplier tube (PMT) settings (760),

except for the negative interactions for which the PMT was increased up to 880.

**Imaging of GFP-CESA and Quantification of Velocity.** Seedlings expressing GFP-CESA3 and GFP-CESA6 were cultured as described (35). Three-day-old etiolated seedlings were analyzed on an Axiovert 200M confocal microscope (Zeiss, Thornwood, NY) equipped with a Yokogawa CSU22 spinning disk, Zeiss  $\times 100/1.4$  N.A. oil objective, and Andor EMCCD DV885 camera (Plateforme d'Imagerie Dynamique, Institut Pasteur, Paris). GFP was excited at 488 nm by a diode pumped solid-state laser, and fluorescence emission was collected through a 505/555-nm band-pass filter (Semrock, Rochester, NY). GFP-CESA velocities were quantified by using the manual tracking plugin (F. Cordelières) in ImageJ (W. Rasband, National Institutes of Health, Bethesda, MD) ( $n = 307$  for CESA3;  $n = 393$  for CESA6).

**GUS Staining.** Seventy-two-hour-old dark-grown seedlings expressing promoter CESA-GUS fusion were submerged in GUS buffer as described (36), infiltrated three times (3 min) under vacuum, and incubated at  $37^\circ\text{C}$  for 1 or 2 h. Seedlings were washed three times with 70% ethanol.

**FTIR Microspectroscopy.** Four biological replicates of wild-type or mutant dark-grown seedlings were analyzed by FTIR microscopy as described (37). The collected spectra were baselined and normalized as described, and statistical analysis was performed by using Student's *t* test (38).

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