Discolored1 (DSC1) is an ADP-ribosylation factor-GTPase activating protein required to maintain differentiation of maize kernel structures

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INTRODUCTION

The maize kernel is a single-seeded fruit composed of the triploid endosperm, the diploid embryo, and the maternally derived pericarp, pedicel, and placenta. At maturity the maize embryo comprises a primary root axis and a relatively precocious shoot axis that may develop up to six leaf primordia. During germination the endosperm nurtures the embryo as it matures into a seedling. Development of the clonally distinct endosperm and embryo ensues following double fertilization of the female gametophyte. The fertilized central cell gives rise to the triploid endosperm, which becomes cellularized 4 days after pollination (DAP; Olsen, 2001). Four distinct structures differentiate to form the maize endosperm, and include: the basal endosperm transfer layer (BETL); the embryo surrounding region (ESR); the starchy endosperm; and the aleurone. Developing at the base of the kernel, the BETL facilitates the transfer of maternally derived nutrients and photosynthates from the placenta into the developing endosperm (Brink and Cooper, 1947; Kiesselbach and Walker, 1952). A distinguishing feature of the BETL is cell wall projections that increase the surface area of the plasma membrane, and thus facilitate the transport function of these cells (Thompson et al., 2001). The ESR forms around the earliest-staged embryo (the proembryo) and has a predicted signaling function between the embryo and the early endosperm compartments but is not present in the mature endosperm (Schel et al., 1984; Opsahl-Ferstad et al., 1997). Functioning as energy reserves, starchy endosperm cells accumulate starch and proteins that will nurture development of the germinating seedling (Duvik, 1961). Cuboidal-shaped aleurone cells form a single-cell layer that encompasses the starchy endosperm; during germination the aleurone functions to digest the stored energy reserves in the kernel (Becraft and Yi, 2011). Aleurone cells are clonally related to the starchy endosperm and can re-differentiate into starchy endosperm in the absence of an as yet unidentified signal that is required to maintain aleurone-specific cell fate (Becraft and Asuncion-Crabb, 2000).

The diploid embryo is derived from the fertilized egg cell, and at maturity is composed of the shoot and root apical meristems, the scutellum, five to six foliar leaves protected by the sheathing coleoptile, and a primary root protected by the sheathing coleorhiza (Abbe and Stein, 1954). Landmark events during embryo development (Kaplan and Cooke, 1997) include formation of the pre-meristematic embryo proper, the establishment of the shoot apical meristem (SAM), and the elaboration of three distinct varieties of shoot lateral organs (the scutellum, the coleoptile, and foliar leaves). Historically, these key events in maize embryo development have been described in nine discrete stages (Abbe and Stein, 1954) and include: (1) the proembryo, before the meristem is formed; (2) the transition stage, when the SAM is established; (3) the coleoptile stage, after the SAM has formed and initiates the coleoptile; and (4) stages L1-L6, wherein the SAM initiates up to six foliar leaves (Abbe and Stein, 1954; Poethig et al., 1986).

Defective kernel (dek) mutations condition defects in both embryo and endosperm development, and thus are useful genetic tools to study kernel development (Neuffer and Sheridan, 1980; Scanlon et al., 1994). Several dek mutants have shared defects in

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embryonic epidermal patterning and in the maintenance of the endosperm aleurone layer (Becraft et al., 1996, 2002; Becraft and Asuncion-Crabb, 2000; Kessler et al., 2002; Lid et al., 2002; Shen et al., 2003). Several genes underlying these *dek* mutations are implicated to function in cell-to-cell signaling during endosperm differentiation. *Dek1* encodes a trans-membrane domain protein with a cytoplasmic calpain-like protease (Lid et al., 2002), and *cr4* encodes a tumor necrosis factor-like receptor kinase (Becraft et al., 1996); both DEK1 and CR4 are required to maintain aleurone cell fate. In contrast, *sal1* functions to inhibit aleurone cell fate during kernel development and encodes an E class vacuolar sorting protein (Shen et al., 2003), which implicates an essential role for endomembrane trafficking in endosperm differentiation.

Endomembrane vesicle trafficking involves the intracellular and intercellular transport of cellular cargo, including proteins, cell wall pectins, structural sterols, receptors, lipids, and signaling molecules, from one membrane-bound compartment to another (Cosgrove, 1997; Takai et al., 2001; Samaj et al., 2005). Vesicle trafficking is regulated in part by the activity of ARF-GTPases, which cycle between active and inactive forms that correlate with vesicle formation and dissociation, respectively (Nie and Randazzo, 2006). Active ARF-GTPases associate with GTP and are membrane-bound during vesicle formation; inactive ARF-GTPases associate with GDP in the cytosol and function during vesicle dissociation. These cyclic activities of ARF-GTPases are regulated by ARF-GEFs that catalyze the exchange of GDP for GTP, and by ADP-ribosylation factor-GTPase activating proteins (ARF-GAPs) that catalyze the subsequent hydrolysis of GTP-bound ARFs (Chardin et al., 1996; Scheffzek et al., 1998; Goldberg, 1999). In *Arabidopsis*, endomembrane cycling of the PINFORMED (PIN) family of auxin efflux proteins requires the activities of both the ARF-GEF Gnom/Emb30 (GN) and the ARF-GAP vascular network defective3 (VAN3)/Scarface (SFC; Geldner et al., 2003; Koizumi et al., 2005; Sieburth et al., 2006; Naramoto et al., 2010). Mutations in *GN* and *VAN3/SFC* give rise to mutants that have defects in embryo development or vascular differentiation, respectively (Geldner et al., 2003; Koizumi et al., 2005; Sieburth et al., 2006). ARF-GAP domain1 (AGD1), a second ACAP-type ARF-GAP characterized in *Arabidopsis*, functions in signaling pathways that remodel the actin cytoskeleton and direct membrane trafficking to maintain polarized root hair growth (Yoo et al., 2008, 2012). The characterization of these ARF-GAPs in *Arabidopsis* indicates that there is a wide diversity of ACAP-type ARF-GAP function and cargo specificity.

Previously, the maize *dek* mutation *discolored1* (*dsc1*) was identified in a *Mutator* (*Mu*) transposon-mutagenized population and named for the shrunken, brown phenotype of homozygous mutant kernels (Scanlon et al., 1994). Recessive mutations in *DSC1* condition inviable kernel phenotypes, and transposon-tagging identified a *Mu*-inserted genomic DNA fragment from the 5′ UTR region of the *dsc1*-Reference (*dsc1*-R) mutation (AF006498; Scanlon and Myers, 1998). *DSC1* encodes an ARF-GAP protein and accumulates in kernels harvested after 6 DAP and in seedling roots and shoots. Transient expression assays in *N. benthamiana* leaf tissue show that YFP-tagged DSC1 proteins co-localize with the trans-Golgi network/early endosomes and with the plasma membrane. Taken together, these data reveal that DSC1 functions in endomembrane trafficking between the trans-Golgi and the plasma membrane, and is required for the maintenance of differentiated cell types in the maize kernel.

**MATERIALS AND METHODS**

**PLANT MATERIALS**

The *dsc1*-R mutation arose from a *Mu* transposon line (Scanlon et al., 1994) and was introgressed into a B73 background for at least six generations before harvesting kernels used in phenotypic and gene expression analyses. The *dsc1*-C06, *dsc1*-H02, and *dsc1*-B09 alleles were identified after screening the trait utility system for corn (TUSC), a *Mu* transposon-mutagenized population (Meeley and Briggs, 1995). Primers for this screen can be found in Table A1 in Appendix.

**HISTOLOGICAL ANALYSES AND IN SITU HYBRIDIZATIONS**

For histological analyses, wild type and discolored mutant kernels were harvested 6 DAP to 20 DAP and fixed overnight in FAA (37% formaldehyde: ethanol: glacial acetic acid: water at 10:50:5:35). The kernels were dehydrated in an ethanol/tert-butyl alcohol series, embedded in paraplast, and 10 μm thick sections were stained with either safranin O-fast green or safranin O-orange G as previously described (Ruzin, 1999). For *in situ* hybridizations, wild type and discolored mutant kernels were harvested 6 DAP to 20 DAP, fixed in FAA, dehydrated, embedded in paraplast, sectioned, and hybridized with gene specific probes as previously described (Jackson, 1991). Primers used to make probes can be found in Table A1 in Appendix. All samples were imaged using the Zeiss Axio Imager Z1-Apomte microscope (Thornwood, New York) and Zeiss Axiovision release 4.6 software.

**IDENTIFICATION OF THE FULL LENGTH *dsc1* TRANSCRIPT AND GENE EXPRESSION ANALYSIS**

The Invitrogen Superscript III One Step RT-PCR Platinum Taq HiFi kit was used to clone *dsc1*. Briefly, tissue segments including the vegetative SAM were cut from 10 day old B73 seedlings. Total RNA was isolated using the RNeasy Plant Mini Kit (Qiagen) and poly(A) RNA was isolated by the Oligotex mRNA mini kit (Qiagen). Prior to RT-PCR, total RNA was isolated from harvested wild type and discolored mutant kernels (6 DAP, 8 DAP, 12 DAP, 14 DAP, 16 DAP, 18 DAP) and hybridized with gene specific probes as previously described (Prescott and Martin, 1986) with some modifications. Following the five minute incubation on ice with chloroform/isoamyl alcohol (24:1), samples were centrifuged for 10 min at 4°C. After transferring the aqueous phase to a new tube, 1 mL of TRIzol (Invitrogen) was used to extract RNA following the manufacturer’s protocol. Total
RNA was extracted from whole 14 day old seedlings and from the upper third of the emerging leaf blade from 14 day old seedlings grown on soil, and roots from 14 day old seedlings grown on 0.02% agar using TRIzol (Invitrogen) according to the manufacturer’s protocol. Superscript III (-Invitrogen) was used to synthesize cDNA from 1 μg of RNA treated with DNasel (Invitrogen). SYBR-green (Quanta) methodology combined with gene specific primers (Table A1 in Appendix) as described in (Zhang et al., 2007) was used in the quantitative RT-PCR analysis. Relative gene expression normalized to 18 s rRNA was determined using the 2−ΔΔCT method as described in Livak and Schmittgen (2001).

TRANSIENT EXPRESSION ASSAYS
The Gateway Recombination Cloning System (Invitrogen) was used to clone the DSC1 ORF into the pEarleyGate104 destination vector as described (Earley et al., 2006). Details about the organelle marker constructs are previously described (Kohler et al., 1997; Boevink et al., 1998; Reisen and Hanson, 2007; Geldner et al., 2009). Electroporation was used to transform agrobacterium strain C58C1 with the N-terminal 35S-YFP<DSC1 fusion construct, the control construct (empty pEarleyGate104 vector), and the organelle marker constructs (35S-ERD2<GFP, 3553535AMV-COXIV<GFP, 35S-DsRed<CAT, UBPQ10-mCherry<VTI12, UBPQ10-mCherry<Got1p; Kohler et al., 1997; Boevink et al., 1998; Reisen and Hanson, 2007; Geldner et al., 2009). Transmants from individual construct lines were grown overnight at 28°C in 2 mL of LB medium containing 50 μg/mL kanamycin and 5 μg/mL tetracycline. After centrifugation, the cells were resuspended in 10 mM MgCl2 to an OD of 0.5 and incubated at room temperature for 2–4 h. For localization of YFP-tagged DSC1 and the control construct 35S-YFP, N. benthamiana leaves were infiltrated as described in Goodin et al. (2002) and observed using epifluorescence microscopy between 48 and 72 h later. Images were obtained on the Zeiss Axio Imager Z1-Apottome microscope (Thornwood, New York) and Zeiss Axiovision release 4.6 software. For co-localization assays, equal parts of the suspensions transformed with each one of the five organelle markers were individually mixed with the suspension transformed with YFP-tagged DSC1 prior to incubation at room temperature. N. benthamiana leaves were infiltrated as described in Goodin et al. (2002) and observed using confocal microscopy between 48 and 72 h later. Imaging of fluorescent proteins was performed using a Leica TCS-SP5 confocal microscope (Leica Microsystems, Exton, PA, USA) using either 10× or 40× objectives (NA 0.4 or 0.85, respectively). Images were obtained sequentially to separate signal from the two channels and were later superimposed. Time lapse series were collected non-sequentially. All images were taken using either a blue argon ion laser (Ar) or a diode pumped solid state laser (DPSS). Excitation and emission parameters are presented in Table A2 in Appendix. Leica LAS-AF software (version 1.8.2) was used to process all images.

PIN1A TRANSPORT ASSAYS
Plants heterozygous for dsc1-R were crossed with ZmPIN1a-YFP transgenic individuals (Gallavotti et al., 2008). The resulting progeny were planted, screened for the dsc1-R allele and YFP, grown to maturity, and self-pollinated. Ears were harvested the same day kernels were removed for live imaging. Embryos were dissected and put on culture media as described in Scanlon et al. (1997). A BFA (Sigma-Aldrich) stock solution was diluted in DMSO and added to liquid culture media to make a final concentration of 100 μM BFA. Mock treatments were made by adding the same amount of DMSO (minus BFA) to liquid culture media. Harvested embryos were incubated in culture media containing BFA or the mock treatment for at least 4 h before confocal microscopy image analysis. Images were collected non-sequentially as described above.

RESULTS
DSC1 IS REQUIRED TO MAINTAIN DIFFERENTIATION OF EMBRYO AND ENDOSPERM STRUCTURES
Self-pollinated plants heterozygous for the dsc1-R mutation segregate mutant kernels with aberrations in both embryo and endosperm development (Scanlon et al., 1994; Scanlon and Myers, 1998). Wild type kernels are yellow at 12 DAP, where upon discolored mutant kernels are white and smaller than wild type siblings (Figure 1A). Embryo structures are not discernible in mutant kernels dissected after 18 DAP, and the reduced endosperm development fails to fill the kernel space (Figures 1B,C). At maturity, all discolored mutant kernels are brown, misshapen, and embryo lethal (Figure 1D).
Detailed phenotypic analyses of discolored mutant kernels harvested at different time-points following pollination show that development of both the embryo and endosperm is normal, albeit delayed, before eventual kernel abortion and tissue disintegration. For example, whereas late proembryo-staged embryos are harvested from wild type embryos at 8 DAP (Figures 2A,B), mutant embryos harvested from the same ear comprise far fewer cells than wild type siblings and are still encased within the ESR (Figures 2G,H). Wild type embryos harvested at 12 DAP are in stage L2, having already elaborated the scutellum, the coleoptile, and the first foliar leaf (Figures 2C,D). In contrast, 12 DAP discolored mutant sibling embryos are stalled at the early transition stage (Figures 2I,J). Three foliar leaf primordia (i.e., stage L3) are initiated in wild type embryos harvested at 16 DAP (Figures 2E,F), whereupon discolored mutant embryos exhibit a variable range of phenotypes, including embryos retarded at stage L1 (Figures 2K,L) and others in which identification of embryo developmental stage is impossible owing to tissue degeneration (Figures 2M–P). In agreement with the mutant embryo phenotypes described above, molecular markers for shoot meristem maintenance [knotted1 (kn1)], vasculature development [ran binding protein2 (ranbp2)], and scutellum initiation [Zea mays yabby1 (zyb1)] showed normal transcript accumulation in transition-staged discolored mutant embryos (Figures 3A–F; Smith et al., 1995; Juarez et al., 2004). Likewise, degenerated later-staged discolored mutant embryos do not accumulate kn1 or zyb1 transcripts after 16 DAP (Figures 3G,H).

Similar to the development discolored mutant embryos, the differentiation of aleurone, BETL, and starchy endosperm cell types is also delayed and eventually aborted in discolored mutant kernels. For example, all these endosperm-specific structures are fully differentiated in 8 DAP wild type kernels. Aleurone cells assume their distinctive cuboidal shape within a single layer surrounding the perimeter of the starchy endosperm (Figure 4A), and three layers of highly extended BETL cells develop in the base of the endosperm, immediately juxtaposed to the maternally
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derived placenta (Figure 4B). In contrast, differentiated aleurone cells are not identified in the discolored endosperm at 8 DAP (Figure 4E), and just a single layer of BETL cells form at the base of the mutant endosperm (Figure 4F). By 16 DAP however, both discolored mutant and wild type sibling kernels have fully differentiated endosperm structures, including an anatomically distinct aleurone layer surrounding the enlarged, vacuolated cells of the starchy endosperm, and a BETL comprising three cell layers (Figures 4C,D,G,H). Finally, discolored mutant endosperm development is again aberrant by 20 DAP, wherein undifferentiated cells are observed within the mutant aleurone and BETL layers and other cells are degenerated and of non-descript identity (Figures 4I–L).

**dsc1 ENCODES AN ADP-RIOSYLATION FACTOR-GTPase ACTIVATING PROTEIN**

Previously, a 3,808 bp *Mu1*-inserted 5′ genomic fragment of the *dsc1* locus on chromosome 4S was identified by transposon-tagging analysis of the *dsc1-R* mutation (Scanlon and Myers, 1998). Three additional *Mu* insertion alleles of *dsc1* (*dsc1-H02, dsc1-C06, and dsc1-B09*) obtained from the TUSC (Meeley and Briggs, 1995) failed to complement the *dsc1-R* mutation (Figure 5A), and thus provided further confirmation that the *dsc1* locus has been cloned. Alignment of the ~2.4 KB genomic DNA fragment of the *dsc1* clone to the sequenced maize genome (MaizeSequence.org Release 5b.60; Schnable et al., 2009) identified the predicted full length gene, the 2.472 KB full length transcript, the boundaries of the 17 predicted introns, and the 823 amino acid sequence of the predicted *dsc1* gene product. The sequence of the *dsc1* cDNA was confirmed by RT-PCR using primers anchored within the *dsc1-R* transposon-inserted genomic DNA fragment described above (Figures 5A,B). PCR analyses utilizing *dsc1* gene-specific primers (Table A1 in Appendix) and a primer targeted to the *Mu* transposon termini (*MuTIR; Table A1 in Appendix*) identified the *Mu*-insertion sites of the four *dsc1* mutant alleles. As indicated in Figure 5A, the *dsc1-R* and *dsc1-H02* alleles harbor 5′ *Mu* transposon insertions located 233 bp and 283 bp upstream of the *dsc1* start codon, respectively. In contrast, the *dsc1-C06* and *dsc1-B09* alleles contain *Mu*-insertions within the first intron, 284 bp and 250 bp downstream from the *DSC1* start codon. Notably, no exon insertions were identified among our *dsc1* mutant alleles. The predicted 823 amino acids of the DSC1 protein encode a putative ARF-GAP protein comprising a BIN-ampiphysin-RVS (BAR) domain, a pleckstrin homology motif, and two ankyrin repeats, in addition to an ARF-GAP domain (Figure 5B). Both the BAR and pleckstrin homology domains are implicated in membrane interactions (Hurley, 2006), whereas ankyrin repeats are known to function in protein-protein interactions (Inoue and Randazzo, 2007). ARF-GAPs comprise a highly conserved group of proteins within the eukaryotes, and function during the regulation of vesicle trafficking (Vernoud et al., 2003; Jiang and Ramachandran, 2006; Inoue and Randazzo, 2007). Taken together, the DSC1 protein is predicted to function during endomembrane trafficking in maize.

A total of 43 predicted maize genes encode an ARF-GAP domain, including *dsc1* and a close paralog (95% nucleotide identity/98% amino acid identity) located on chromosome 1 (GRMZM5G872204, designated here as *dsc2*). In adherence to the mammalian classification system, nine predicted maize ARF-GAP proteins (including DSC1 and DSC2) belong to the ACAP subgroup, comprising ARF-GAPs that function during post-Golgi transport (Jackson et al., 2000; Miura et al., 2002; Nie et al., 2003; Randazzo and Hirsch, 2004). The ARF-GAP domain of DSC1 is homologous (66% identity) to that of the *Arabidopsis* protein VAN3 (Koizumi et al., 2005; Sieburth et al., 2006), which regulates endomembrane trafficking of the auxin efflux protein PIN1, although yet another maize ARF-GAP (GRMZM2G059225) is more similar to VAN3 (69% identity). ClustalW alignments of the ARF-GAP domains of DSC1, DSC2, VAN3, and a human ACAP-type ARF-GAP (ACAP2) are shown in Figure 5C.

RT-PCR identified *dsc1* transcript accumulation during multiple stages of kernel development (6DAP, 8 DAP, 12 DAP, 14 DAP, 16 DAP, and 18 DAP) and in 14-day-old seedling shoots and roots, but not in the fully differentiated distal tips of emerged seedling leaves (Figure 6A). Notably, *dsc1* transcript accumulation in 12 DAP, early transition-staged discolored mutant kernels is reduced to less than 0.3 times the level found in wild-type transition-staged seed harvested at 8 DAP (Figure 6B). Furthermore, transcripts of the kernel patterning genes *dek1* and *cr4* both accumulate to less
Development of endosperm tissues is delayed in dscl-R mutant kernels, prior to eventual tissue degeneration. Sagittal sections of (A–D) non-mutant and (E–L) dscl-R mutant kernels harvested at 8 DAP (A,B,E,F), 16 DAP (C,D,G,H), and 20 DAP (I,L). (I) The lack of staining due to the presence of dense cytoplasm and the round cell morphology indicate that the aleurone layer has not fully differentiated in this 20 DAP kernel. (J) The aleurone and the starchy endosperm cells in this 20 DAP kernel have already degenerated. (K) The arrow points to undifferentiated cells in the aleurone cell layer in this kernel harvested 20 DAP. (L) The arrows point to misshapen basal endosperm transfer layer cells that lack cell wall ingrowths at the base of this 20 DAP kernel. All sections were stained with Safranin-O and Fast Green for the exception of (K), which is stained with Safranin-O and Orange G. AL, aleurone; BETL, basal endosperm transfer layer; DAP, days after pollination; PE, pericarp; PL, placenta; SE, starchy endosperm. Scale bars represent 100 μm.

DSC1 co-localizes with the trans-Golgi network/early endosomes and the plasma membrane

Yellow fluorescent protein (YFP)-tagged DSC1 constructs were generated to determine the subcellular localization of DSC1 in the N. benthamiana leaf (Earley et al., 2006). Epifluorescence and confocal microscopic imaging of infiltrated transgenic leaf sectors reveals 35S-YFP<DSC1 accumulation within distinct intracellular compartments, and as foci at the plasma membrane (Figures 7A,D,G,J,M; Figures A1A,C and A2A,D,G in Appendix). 35S-YFP<DSC1 labeled bodies also actively move around the cell, as evidenced by time series taken of cells to illustrate the intracellular movement of 35S-YFP-tagged DSC1 (Movie S1 in Supplementary Material). In contrast, N. benthamiana leaves expressing control 35S-YFP construct that lacked the DSC1 open reading frame show fluorescent signal within the nucleus, or dispersed within the cytoplasm (Figures A1E,G in Appendix).

The lipophilic, styryl dye FM4-64 (Invitrogen) fluoresces in hydrophobic environments, and is commonly utilized to trace endocytotic activities in cells (Bolte et al., 2004). Once internalized FM4-64 labeled membranes successively co-localize with the trans-Golgi network/early endosomes, prevacuolar compartments, and ultimately with the tonoplast (vacuolar membrane; Bolte et al., 2004; Geldner et al., 2009). Co-localization assays with several organelle-specific markers and the endocytic tracer FM4-64 were performed to identify more precisely the intracellular localization of YFP-tagged DSC1 protein. These studies confirmed that 35S-YFP<DSC1 does not co-localize with the...
FIGURE 5 | dscl encodes an ARF-GAP. (A) The full length dscl transcript is 3.194 kb. Positions of Mu transposon insertions in individual dscl mutant alleles with images of their kernel phenotypes from failed complementation crosses are indicated. Hash marks denote unknown lengths of genomic fragments. (B) The DSC1 protein comprises 823 amino acids and is composed of a BAR, a PH, an ARF-GAP, and two ANKYRIN domains. (C) ClustalW mediated amino acid alignment of the ARF-GAP of DSC1 with that of the two maize paralogs on chromosomes 1 (DSC2, GRMZM5G872204) and 7 (GRMZM2G059225), the Arabidopsis protein VAN3/SFC/AGD3 (NP196834), and ACAP2 (AAH60767) from Homo sapiens. Amino acid percent identity to DSC1 ARF-GAP domain is in parentheses.
FIGURE 6 | Transcript accumulation of *dscl*. (A) RT-PCR of *dscl* transcript accumulation in developing maize kernels (6–18 DAP), 14-day-old seedlings, 14-day-old roots, and leaf blade tips. (B) qRT-PCR analysis of *dscl*, *dekl*, *cr4*, and *sall* in 8 DAP non-mutant and *dscl-R* mutant transition-staged kernels. Accumulation was normalized to 18S rRNA. Each experiment utilized three biological replicates. Error bars denote SE. Primers were designed to the 3′ end of the *dscl* transcript and can be found in Table A1 in Appendix.

mitochondrial marker 35S35SAMV-COXIV<GFP, the peroxisome marker 35S-DsRed<catalase, auto-fluorescent chloroplasts, the cis-Golgi marker 35S-ERD2<GFP; or with the Golgi stacks marker UBQ10-mCherry<Got1p (Figures A2A–I in Appendix; Figures 7A–F; Kohler et al., 1997; Boevink et al., 1998; Reisen and Hanson, 2007; Geldner et al., 2009). However, in some instances 35S-YFP<DSC1 labeled bodies localize next to the cis-Golgi and Golgi stacks markers (Figures 7A–F). Recurrently, 35S-YFP<DSC1 labeled bodies co-localize with the trans-Golgi network/early endosome marker UBQ10-mCherry<VTI12 (Figures 7G–I), in addition to FM4-64 labeled intracellular compartments and the plasma membrane (Figures 7J–O; Geldner et al., 2009). Taken together, the cellular mobility of YFP-tagged DSC1 labeled compartments and their co-localization with both FM4-64 and a marker for the trans-Golgi network/early endosome illustrate that DSC1 functions in endomembrane trafficking.

ZmPIN1A ENDOMEMBRANE TRANSPORT IS NOT DISRUPTED IN DISCOLORED MUTANT EMBRYOS

In *Arabidopsis*, the ARF-GAP VAN3 functions during the transport of PIN1 from the plasma membrane to the recycling endosome (Sieburth et al., 2006). YFP-tagged ZmPIN1a localization was observed in wild type and discolored mutant embryos to determine if the endomembrane transport of ZmPIN1a is disrupted by the *dscl-R* mutation. ZmPIN1a<YFP localizes to the plasma membrane in both mutant and non-mutant 14 DAP embryos (Figures 8A–D). After treatment with Brefeldin A, which blocks vesicle cycling from the endosome to the plasma membrane (Steinmann et al., 1999; Geldner et al., 2001), ZmPIN1a<YFP accumulates in endosomal compartments in both wild type and discolored mutant embryos and no differences in size or number of ZmPIN1a<YFP-tagged endosomal compartments are observed (Figures 8E,F).
FIGURE 7 | YFP-tagged DSC1 in transient expression assays in N. benthamiana leaves. (A,D) 35S-YFP<DSC1 bodies do not co-localize with (B) cis-Golgi marker 35S-ERD2<GFP ([C], merged), or (E) the Golgi marker UBO10-mCherry<Gotlp ([F], merged). (G,J,M) 35S-YFP<DSC1 bodies do co-localize with some of the intracellular compartments labeled with (H) the trans-Golgi/early endosomal marker UBO10-mCherry<VTI12 ([I], merged) and (K,N) the endocytic tracer FM4-64 in mesophyll and epidermal cells ([L,O], merged). Arrows point to compartments that co-localize with YFP-tagged DSC1. Arrowheads point to YFP-tagged DSC1 compartments at the plasma membrane. Scale bars represent 50 μm.

DISCUSSION
Defective kernel mutants are phenotypically variable (Sheridan and Neuffer, 1980; Scanlon et al., 1994), such that the wide range in developmental progression observed in the dsc1-R homozygous mutant endosperm and embryo is not unusual, even after more than six introgressions into the B73 inbred background. Analyses of dsc1 transcript accumulation in equivalently staged discolored mutant and wild type kernels reveal that dsc1 transcripts are reduced in dsc1-Rmutants. Another previously reported phenotype observed in maize dek mutants is the relatively normal, albeit retarded, embryo, and endosperm development of the early-staged kernel, followed by kernel abortion and reabsorption.
of most or all kernel tissues (Clark and Sheridan, 1988; Scanlon et al., 1997; Fu et al., 2002). In this way, discolored mutant embryos initiate a SAM, vascular tissue, and a scutellum, whereas some mutant embryos progress to the coleoptilar stage or stage L1 before embryo structures completely degenerate (Figures 2G–P). Likewise, the discolored mutant endosperm forms at least some fully differentiated BETL and aleurone cell types prior to kernel abortion. The delay in discolored mutant endosperm development is evidenced by the delay in forming the aleurone cell layer and the presence of only one file of BETL cells (Figures 4E,F). dsc1 and dsc2 display similar transcript accumulation patterns in previous transcriptomic analyses performed on mixed maize tissues (Sen et al., 2009; Sekhon et al., 2011); partial genetic redundancy of these paralogous ACAP/ARF-GAP proteins early in kernel development may explain why discolored mutant kernels can progress, albeit at a retarded rate, through at least the first stages of kernel development. Perhaps the most interesting and informative aspect of the discolored mutant phenotype is the appearance of undifferentiated cells in the aleurone cell layer, prior to kernel abortion, and tissue degeneration (Figure 4K). This represents a mosaic phenotype and a failure to complete differentiation of the aleurone layer. This striking phenotype reaffirms the conclusions drawn from previous mosaic analyses of maize kernel development (Becraft and Asuncion-Crabb, 2000), in that determination of endosperm cell fate requires a differentiation signal that must be maintained until late stages in kernel development. Our data also implicate a role for DSC1 in the maintenance of endosperm cell fate.

In a heterologous transient expression system, the fluorescently tagged DSC1 protein exhibited intracellular motility and localized to the plasma membrane as well as the trans-Golgi network/early endosomes of N. benthamiana leaves (Figures 7G–O; Movie 1). These data suggest that the DSC1 ARF-GAP protein regulates endomembrane transport of an unknown cargo between the plasma membrane and the trans-Golgi network/early endosomes. Additional co-localization assays could be performed to determine if DSC1 co-localizes with late endosomes or prevacuolar compartments. In addition, localization of ZmPINa-YFP in maize embryos before and after Brefeldin A treatment reveals that DSC1 is not required for trafficking of this auxin efflux transporter from the plasma membrane to endosomal compartments. Taken together, these data suggest that the DSC1 ARF-GAP regulates transport of an unidentified cargo that is required to maintain differentiation of maize kernel tissues. Interestingly, genetic data suggest that SAL1, a class E vacuolar sorting protein, is antagonistic to DEK and CR4, two maize proteins that were initially identified in an intercellular signaling pathway required for the maintenance of endosperm cell fate (Becraft and Asuncion-Crabb, 2000; Becraft et al., 2002; Lid et al., 2002; Shen et al., 2003). However, no direct interaction has yet been demonstrated. We show that DSC1 function is required for normal accumulation of dekland cr4 mRNA, although transcript accumulation of sal1 is not significantly disrupted in discolored kernels (Figure 6B). These data suggest that DSC1 either functions downstream of SAL1, or in an unrelated signaling pathway. Likewise, these data are consistent with DSC1 function upstream of the CR4/DEK1 signal transduction pathway, however no genetic or biochemical evidence of this interaction is currently available. Our analyses of DSC1 further implicate a role for endomembrane trafficking during maize kernel development, and future studies will investigate the identity of the specific cargo implicated during DSC1 function and its specific role in maize kernel development.

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SUPPLEMENTARY MATERIAL

The Movie S1 for this article can be found online at http://www.frontiersin.org/Plant_Evolution_and_Development/10.3389/fpls.2012.00115/abstract

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### Table A1 | Primers utilized in this study.

| Gene name          | Accession      | Forward primer             | Reverse primer             |
|--------------------|----------------|----------------------------|----------------------------|
| **IN SITU HYBRIDIZATION PROBES** |                |                            |                            |
| knotted1           | GRMZM2G017087  | ACAAGGTGGGGGCACCA          | TCGGTCTCTCCTCCGCTA          |
| Zea mays yabby14   | GRMZM2G005353  | CGACCTCAACCAGCGTCT         | GAGCTCCTCCTGAGTTTGC         |
| ran binding protein2| GRMZM2G094353  | GAACAGGAAGCCAGGAGACT       | CAGTGCAAGTAGTTTCTGAGGT      |
| **FULL LENGTH TRANSCRIPT** |                |                            |                            |
| dsc1               | CATGCATTTGCGCAAGCTGTACTCG  |                           |  |
| actin              | TGTCAGGGACATCAAGAGAA        |                           |  |
| dsc1-qRT-PCR       | CTTCACCATGTTGAAAGTCTCTAGA  |                           |  |
| 18s rRNA           | CTGTCGCAAGCCTTACTGACTT     |                           |  |
| defective kernel1  | TGTTTCGGAACCTACACTAAT      |                           |  |
| crinkly4           | TCTCTAACCACACTCTCGCT       |                           |  |
| supernumery aleurone1 | ACCGAGCAGCTACTACACT       |                           |  |
| **RT-PCR/qRT-PCR** |                |                            |                            |
| dsc1-RT-PCR        | CTTACCACTCTGTGAAAGTCTCTAGA|                           |  |
| actin              | TGTCAGGGACATCAAGAGAA        |                           |  |
| dsc1-qRT-PCR       | CTTCACCATGTTGAAAGTCTCTAGA  |                           |  |
| 18s rRNA           | CTGTCGCAAGCCTTACTGACTT     |                           |  |
| defective kernel1  | TGTTTCGGAACCTACACTAAT      |                           |  |
| crinkly4           | TCTCTAACCACACTCTCGCT       |                           |  |
| supernumery aleurone1 | ACCGAGCAGCTACTACACT       |                           |  |
| **DSC1 SUBCELLULAR LOCALIZATION** |                |                            |                            |
| dsc1               | CACCATGCATTTGCGCAAGATGAT   |                           |                            |
| **TUSC SCREEN**    |                |                            |                            |
| MuTIR              | AGAGAAGCACAAGCCACCAWCAGCTCY|                           |                            |
| DO146621           | TCAACGCTCTAACCACACTCCAGTTAC|                           |                            |
| DO146618           | CTCTCCTTCCCTCCCCAAGGAAG   |                           |                            |

### Table A2 | Confocal microscopy parameters.

| Fluorescent protein | Laser | Excitation (nm) | Emission range (nm) | Additional fluorescent proteins used in assay | Assay          |
|---------------------|-------|-----------------|---------------------|-----------------------------------------------|----------------|
| GFP                 | Ar    | 488             | 496 to 513          | YFP/chlorophyll                               | Transient expression |
| Chlorophyll         | Ar    | 488             | 672 to 690          | GFP/YFP/Red                                   | Transient expression |
| Chlorophyll         | Ar    | 488             | 664 to 718          | mCherry/FM4-64/YFP                             | Transient expression |
| YFP                 | Ar    | 488             | 526 to 609          | GFP/chlorophyll                               | Transient expression |
| YFP                 | Ar    | 488             | 525 to 536          | DsRed/chlorophyll                             | Transient expression |
| YFP                 | Ar    | 488             | 522 to 555          | mCherry/FM4-64/chlorophyll                    | Transient expression |
| YFP                 | Ar    | 488             | 524 to 583          | N/A                                           | PINI a transport   |
| DsRed               | Ar    | 514             | 589 to 620          | YFP/chlorophyll                               | Transient expression |
| mCherry             | DPSS  | 561             | 582 to 632          | YFP/chlorophyll                               | Transient expression |
| FM4-64              | DPSS  | 561             | 582 to 632          | YFP/chlorophyll                               | Transient expression |
FIGURE A1 | Subcellular localization of YFP-tagged DSC1 and 35S-YFP control construct. DSC1 localizes to punctate structures in (A) mesophyll and (C) epidermal cells. The control construct has nucleo-cytoplasmic localization in (E) mesophyll and (G) epidermal cells. (B,D,F,H) Bright field images were taken of the cells located directly to the left. Images were obtained using epifluorescence microscopy.
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FIGURE A2 | YFP-tagged DSC1 transient expression assays in N. benthamiana. (A,D,G) 35S-YFP<DSC1 bodies do not co-localize with (B) chlorophyll auto-fluorescence [(C), merged], (E) the mitochondria marker 35S35SAMV-COXIV>GFP [(F), merged], or (H) the peroxisome marker 35S-DsRed<catalase [(I), merged]. Chlorophyll auto-fluorescence is shown in each of the merged images (C,F,I). Scale bars represent 50 μm. CHL, chlorophyll.