Novel Regulation of Skp1 by the Dictyostelium AgtA α-Galactosyltransferase Involves the Skp1-binding Activity of Its WD40 Repeat Domain*

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Background: AgtA mediates terminal α-galactosylation of Skp1, but genetic studies show an unexpectedly profound role in oxygen sensing.

Results: AgtA exhibits salt-sensitive Skp1-binding activity that competes with other Skp1 interactions but subsequently activates α-galactosylation that alters Skp1 binding. AgtA regulates development through non-enzymatic mechanisms.

Conclusion: AgtA modulates Skp1 interactions by a self-limiting glycosylation mechanism.

Significance: Polypeptide recognition by a glycosyltransferase mediates non-enzymatic functions.

The role of Skp1 as an adaptor protein that links Cullin-1 to F-box proteins in E3 Skp1/Cullin-1/F-box protein (SCF) ubiquitin ligases is well characterized. In the social amoeba Dictyostelium and probably many other unicellular eukaryotes, Skp1 is modified by a pentasaccharide attached to a hydroxyproline residue near its C terminus. This modification is important for oxygen-sensing during Dictyostelium development and is mediated by a HIF-α type prolyl 4-hydroxylase and five sequentially acting cytoplasmic glycosyltransferase activities. Gene disruption studies show that AgtA, the enzyme responsible for addition of the final two galactose residues, in α-linkages to the Skp1 core trisaccharide, is unexpectedly critical for oxygen-dependent terminal development. AgtA possesses a WD40 repeat domain C-terminal to its single catalytic domain and, by use of domain deletions, binding studies, and enzyme assays, we find that the WD40 repeats confer a salt-sensitive second-site binding interaction with Skp1 that mediates novel catalytic activation in addition to simple substrate recognition. In addition, AgtA binds similarly well to precursor isoforms of Skp1 by a salt-sensitive mechanism that competes with binding to an F-box protein and recognition by early modification enzymes, and the effect of binding is diminished when AgtA modifies Skp1. Genetic studies show that loss of AgtA is more severe when an earlier glycosylation step is blocked, and overexpressed AgtA is deleterious if catalytically inactivated. Together, the findings suggest that AgtA mediates non-enzymatic control of unmodified and substrate precursor forms of Skp1 by a binding mechanism that is normally relieved by switch-like activation of its glycosylation function.

Skp1, an adaptor protein for the Skp1/Cullin-1/F-box protein (SCF) 3 class of E3 ubiquitin ligases (1), is modified by a novel pentasaccharide attached to a hydroxyproline residue near its C terminus in the social amoebe Dictyostelium discoideum and likely other protists (2–4). AgtA was previously characterized as the α-galactosyltransferase catalyzing addition of the 4th sugar, resulting in formation of Galα1,3Fucα1,2Galβ1,3GlcNAcα1–4O-Pro-Skp1 (5). Each of the enzymes required to build the trisaccharide form of Skp1 that serves as the AgtA substrate are known (Fig. 1A) (2). Recent studies have indicated that AgtA also mediates the addition of the final sugar, another α-linked Gal residue, to complete the glycan. 4

AgtA regulates the execution of a developmental process in Dictyostelium known as culmination. Culmination refers to the transition in the starvation-induced developmental program in which the migrating slug reorganizes into a fruiting body consisting of dormant spores perched atop a cellular stalk. Environmental O2 is one of several regulators of culmination, and substantial evidence indicates that the Skp1 prolyl 4-hydroxylase, an O2-dependent enzyme required for formation of the 4-hydroxyproline to which the glycan is attached, is an O2-sensor for culmination. Further genetic analysis has revealed that PgtA, a bifunctional diglycosyltransferase that catalyzes addition of the second and third sugars, and AgtA, modulate the response to Skp1 prolyl hydroxylation (see pathway scheme in Fig. 1A). Remarkably, the effect of interrupting the AgtA locus is nearly as severe as blockade of hydroxylation itself as opposed to disruption of PgtA, which shows only a modest effect (6). Because biochemical and genetic evidence indicates that Skp1 is the only protein substrate of the hydroxylation/glycosylation

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3 The abbreviations used are: SCF, Skp1-Cullin-1-F-box protein; FGGn-pNP, 1-octyl 4-nitrophenyl; FGGn-pNP, 1,2Galβ1,3GlcNAcα1–1-pNP; GT, glycosyltransferase; TCA, trichloroacetic acid; PhyA, prolyl 4-hydroxylase; TEV, tobacco etch virus.
4 Z. Chino, C. M. West, and G. J. Boons, unpublished studies.
pathway (2, 4, 7, 8), modification of Skp1 by PgtA to convert the monosaccharide to the trisaccharide was proposed to reverse the effect of hydroxylation, and terminal α-galactosylation by AgtA was proposed to liberate Skp1 from this inhibition (2). However, the mechanism by which AgtA regulates Skp1 function remains unclear because the protein contains, in addition to its N-terminal catalytic domain, a C-terminal sequence of 7 predicted WD40 repeats, expected to fold into a β-propeller (5, 9, 10). Furthermore, the initial purification of AgtA from Dictyostelium suggested partial co-purification with Skp1 and similar levels of expression, which is unusual for an enzyme/substrate pair (11). Although the overexpressed catalytic domain rescued Skp1 α-galactosylation and O2-sensing in the agtA deletion strain (6), the high level of overexpressed protein might mask a critical function of the WD40 repeats, which represent one of the most common protein domains in nature and often mediate protein-protein interactions (9, 12).

Most eukaryotic GTs are type 2 membrane proteins consisting of a single catalytic domain connected by a linker region to an N-terminal transmembrane domain and a short cytoplasmic extension (13), but AgtA and the other Skp1 GTs lack these domains and exist as soluble cytoplasmic proteins. Although the WD40 repeats of AgtA are unprecedented for known GTs, other GTs possess SH3 (14), TPR (15), and lectin (16) domains. In some instances, these so-called “add-on” domains (17) have been implicated in acceptor substrate recognition, as has the linker region of a type 2 membrane GT (18). Thus the AgtA C-terminal domain may contribute to Skp1 recognition, although it is noted that the earlier GTs in the pathway, the αGlcNAcT (Gnt1) and PgtA (see Fig. 1A), do not rely on add-on domains to achieve exquisite selectivity for Skp1 in vitro (7) and in the cell (4, 6, 8). Here we investigate the functionality of the AgtA WD40 repeat domain in the recognition of Skp1 by AgtA. Our investigations show that the modification by AgtA of Skp1 depends on the WD40 repeats by a mechanism that not only involves second site recognition but novel catalytic activation. Furthermore, Skp1 recognition by AgtA is competitive in vitro with F-box protein (FBP) binding required for SCF complex assembly, and interferes with early Skp1 processing steps in cells. The importance of a non-enzymatic function of AgtA is supported by the blockade of culmination which occurs in a double knock-out with the first Skp1 GT Gnt1 in which the substrate form of Skp1 is not assembled. The findings are interpreted in terms of a binding activity that titrates the interaction of Skp1 with FBPs and other binding partners and is relieved by execution of its glycosylation function.

**EXPERIMENTAL PROCEDURES**

*Dictyostelium Growth and Development*—Cells were grown axenically in HL-5 medium, and assayed for O2-dependent development as described previously (19). Briefly, cells were deposited on filters in PDF buffer and developed in the dark for 42 h under the indicated O2 concentrations with the balance made up with N2. Development was assessed by fruiting body formation by daily counting in a hemacytometer. Stalk morphology was imaged in the DAPI channel of an epifluorescence microscope in the presence of Calcofluor White ST to reveal cellulosic cell walls (19). Cell proliferation was assayed by counting cells during logarithmic growth in HL-5 medium by daily counting in a hemacytometer.

*Dictyostelium Strains*—Normal (Ax3), agtA− (HW420), and gnt1− (gnt1.3/HW503) strains were described previously (4). The gnt1−/agtA− double knock-out with the first Skp1 GT Gnt1 in which the substrate form of Skp1 is not assembled. The findings are interpreted in terms of a binding activity that titrates the interaction of Skp1 with FBPs and other binding partners and is relieved by execution of its glycosylation function.

*Preparation and Purification of Recombinant AgtA and AgtA Domains*—The protein coding sequences for AgtA-FL (amino acids 2–649) was excercped from pCR4TOPOcagta (5) and selection of clones in the presence of G418, yielding strain HW530. The corresponding plasmid encoding catalytically inactive AgtA(D132A) was derived in the same manner (4) and introduced into HW532. These pVSC-derived plasmids are designed to stably overexpress DpGnt1 (the ortholog from *D. purpureum*) under control of the prosopore cell-specific cotB promoter.

Overexpression of His6-AgtA-FL in agtA− strain HW420 was achieved by electroporation of pVS(His)agta (5) and selection of clones in the presence of G418, yielding strain HW530. The corresponding plasmid encoding catalytically inactive AgtA(D132A) was constructed as described below, and used to generate HW532. These pVSC-derived plasmids are designed to stably overexpress His6-AgtA under control of the semi-constitutive discoidin γ promoter.

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Non-enzymatic Binding Function of a Skp1 Glycosyltransferase

WD40 was coexpressed with the protein chaperones GroEL and GroES (pGroELS was the gift of Xiaqiang Wang, Noble Foundation, OK). 1-Liter cultures were induced with 0.3 mM isopropyl 1-thio-β-D-galactopyranoside overnight at 22 °C, pelleted, and resuspended in 40 ml of 0.1 M Tris-HCl (pH 8.2), 5 mM benzamidine, 0.5 μg/ml of peptatin A, 5 μg/ml of apro- 
tinin, 5 μg/ml of leupeptin, 0.5 mM PMSF, and 1 ml of mg/ml of 
lysozyme, lysed using a French Press at 18,000 p.s.i., and 
centrifuged at 100,000 × g for 1 h. The supernatant (S100) was 
applied to a 1-ml GE HiTrap Ni2+ column equilibrated in 20 
M Tris (pH 7.8), 0.5 M NaCl, and 5 mM imidazole. Protein was 
eluted using a 0.005–1.0 M gradient of imidazole in the same 
buffer. The His6-tagged proteins eluted at single peaks that 
were further purified on a HiLoad Superdex 200 size exclusion 
column in 50 mM Tris-HCl (pH 8.0), 0.2 M EDTA, 5 mM 
β-mercaptoethanol.

Substrates—Fuc1 para-nitrophenol (F-pNP) was from Sigma, Fuc1,2Galβ1 octyl (FG) was the gift of Monica Palcic, 
Fuc1,2Galβ1,3GlcNAc1 para-pNP (FGpNP-pNP) was 
prepared as before (11), and FGGn-4(N-Ac)Hyp(DR25)-NH2 
(FGGn-Hyp) and Ac-CIKNDFT(FGGn-)HypEEEEQIRK-NH2 
(FGGn-peptide) were synthesized by Zoiasha Chinyo and Geert- 
Jan Boons (CCRC, University of Georgia).4 Skp1 isoforms were 
prepared recombinantly without epitope tags in 
E. coli (25), and 
purified essentially to homogeneity as described elsewhere (5).

Fbs1 — A plasmid encoding guinea pig His6Fbs1 (also known 
as Ocp1) was provided by M. Henzl (University of Missouri). 
His6Fbs1 was purified essentially to homogeneity from E. coli 
as previously described (20).

AgtA Enzyme Assays—α-GalT activity was assayed as de- 
scribed previously (11). Reaction progress was monitored by the 
transfer of [3H]Gal from UDP-[3H]Gal to synthetic gly- 
cosides or Skp1. The standard reaction was prepared in 50 mM 
HEPES-NaOH (pH 7.2), 10 μM UDP-[3H]Gal, 50 mM NaCl, 2 
mM MnCl2, and 5 mM DTT, in 20 
μl for protein or peptide 
substrates and 50 μl for other substrates. Reactions were initi- 
atated by the addition of enzyme and incubated for 30 min at 
22 °C. Conditions were varied as noted, and incubation times 
and enzyme levels were adjusted to ensure that measurements 
were carried out under initial velocity conditions.

UDP-[3H]Gal was diluted with non-radioactive UDP-αGal to 
achieve final concentrations. UDP-[6-3H]Gal was obtained from 
American Radiochemical Corporation (20 mCi/μmol) or 
Amersham Biosciences. The latter, a generous gift from Joel 
Shaper (Johns Hopkins School of Public Health), was originally 
synthesized in 1984 and stored according to the manufacturer’s 
recommendation. UDP-[6-3H]Gal was repurified by chromatography over a Q-anion exchange column as described (21), and 
further purified by adsorption to a 150-mg Carbograph 
cartridge (Grace-Alltech) pre-equilibrated with 0.1% TFA (v/v). 
UDP-[6-3H]Gal was eluted with 30% acetonitrile in 0.1% TFA. 
Its properties were indistinguishable from that of recently pre- 
pared material in pilot enzyme assays.

Reactions using substrates F-pNP, FG-octyl, and FGGn-pNP were stopped by addition of 1 ml of ice-cold 1 mM sodium 
EDTA (pH 8.0). Reaction mixtures were applied to a 0.5-ml 
high-cap SepPak C18 cartridge (Grace-Alltech), washed six 
times with 5 ml of H2O, and eluted with 5 ml of MeOH. Samples 
were added to 8 ml of Biosafe II scintillation counting fluid 
(RPI) and radioactivity was quantitated using a Beckman L6500 
scintillation counter. Blank activities determined from re- 
actions lacking substrates were subtracted.

Reactions using Skp1 as a substrate were stopped by addition of 80 μl of ice-cold 10 mM sodium EDTA (pH 8.0), 20 μl of 100 
mg/ml of BSA, and 1 ml of 10% (v/v) TCA to precipitate pro- 
teins. The precipitate was collected on a glass fiber filter under 
vacuum, and washed 4 times with 1.5 ml of ice-cold 10% TCA 
and 4 times with 1.5 ml of acetone. The filter was added to 15 ml of 
Biosafe NA scintillation fluid (RPI) and counted as above.

For activation or inhibition studies, purified Skp1 isoforms 
were added in advance. All data were analyzed by non-linear 
regression using GraphPad Prism.

PhyA and Gnt1 Enzyme Assays—Activities of the purified 
enzyme proteins with Skp1 acceptor protein preparations 
were assayed as described (7) in the presence of Fbs1. Brief- 
lly, Gnt1 reactions were monitored by TCA precipitation of 
Skp1 to determine incorporation of [3H]GlcNAc from UDP- 
[3H]GlcNAc. PhyA reactions were monitored using the same 
Gnt1 reaction based on its specificity for the HO-Skp1 product 
of the PhyA reaction. Gnt1 activity was assayed in cell extracts 
as described (4).

Mass Spectrometry of AgtA Reaction Products—Reactions 
containing F-pNP, FG-octyl, and FGGn-pNP were carried to 
near completion, adsorbed to a SepPak C18 eluted in MeOH, 
dried, and analyzed by MALDI-TOF-MS on a Bruker Ultraflex 
II, in positive ion mode using dihydroxybenzoic acid as the 
matrix (19).

Size Exclusion Chromatography—Protein-protein interac-
tions were monitored chromatographically on a Pharmacia 
3.2/10 Superdex 200 column equilibrated in 50 mM HEPES- 
NaOH (pH 7.2), 50 mM NaCl (or 150 mM if indicated), 2 mM 
MnCl2, 5 mM DTT on a Pharmacia SMART System. Protein 
samples were pre-mixed in this same buffer for 30–60 min at 
22 °C before analysis at a flow rate of 100 μl/min. Protein elu- 
tion was monitored by A280 and Western blot analysis of 
column fractions.

Disuccinimidy Suberate (DSS) Cross-linking—His6-AgtA FL 
and Skp1 were dialyzed exhaustively at 4 °C against 50 mM 
HEPES (pH 7.2), 2 mM DTT and co-incubated, each at 600 μM, 
in 50 mM HEPES (pH 7.2), 2 mM MnCl2, 2 mM DTT, 10 μM 
UDP-Gal, 0–250 mM NaCl for 1.5 h at 22 °C. DSS (Thermo) 
was dissolved in dry DMSO at 12.5 mM, diluted H2O to 1 mM, 
and introduced to the preincubated AgtA/Skp1 mixture at a 
final concentration of 250 μM. Cross-linking reactions were 
incubated for 10 min in the dark at 22 °C before quenching by 1 M 
Tris-HCl (pH 7.4) to a final concentration of 75 mM.

SDS-PAGE and Western Blotting—Samples were separated 
by SDS-PAGE and Western blotted using mAb 4E1 for Skp1, 
UOK87 for unmodified Skp1, UOK85 for HO-Skp1, and 
UOK72 or UOK101 for AgtA, as described (4). Alexa 680 fluo-
rescence was imaged in an Odyssey Li-Cor scanner, and images 
were adjusted using the Levels control in Adobe Photoshop so 
that it, or a subregion as indicated, utilized the full brightness 
range available without saturation.

AgtA Antiserum—His6-AgtA-FL and His6-AgtA-CAT were 
used to immunize separate female New Zealand White rabbits
as described previously for other rabbit Abs (4). The exsanguination of UOK72 (His_{6}AgtA-FL) and UOK101 (His_{6}AgtA-CAT) were used at a 1:1000 dilution in Western blotting.

RESULTS

Importance of the WD40 Repeat Domain for AgtA Enzyme Activity—His_{6}-tagged full-length AgtA (AgtA-FL) was purified essentially to homogeneity from E. coli (Fig. 1C) and kinetically analyzed using a highly recombinant purification preparation of the natural substrate, Fucα1,2Galβ1,3GlcNAcα1-Skp1 (FGGn-Skp1). The reaction exhibited Michaelis-Menten characteristics with an apparent $K_m$ of 0.76 μM (Fig. 2A) and an apparent $k_{cat}$ of 0.11 s⁻¹. The apparent $K_m$ compares with a value of 6 μM previously exhibited by a partially purified native preparation from Dictyostelium (11). Further characterization revealed two component reactions (Fig. 1A): addition of αGal-1 to the 3-position of Fuc (4), and addition of αGal-2 to αGal-1 at an undertermined position. It is not known which step is rate-limiting for the overall reaction, and the impact of Skp1 dimerization (20) is also unknown.

To assess the importance of the C-terminal WD40 repeat domain for catalytic activity, the N-terminal catalytic domain (AgtA-CAT) and the His_{6}-tagged C-terminal domain (AgtA-WD40) were separately expressed in E. coli (Fig. 1B) and purified (Fig. 1C). In initial assays conducted with small sugar acceptor substrates, AgtA-CAT exhibited substantial activity comparable with that of full-length AgtA-FL, and AgtA-WD40 had no activity. To compare initial reaction velocities, acceptor substrates were tested near their $K_m$ values as previously determined by studies of partially purified AgtA from Dictyostelium (11), and enzyme protein levels were adjusted to yield similar incorporation for the simplest substrate, Fuc-pNP (F-pNP). Both AgtA-FL and AgtA-CAT exhibited substantially greater activity toward Fucα1,2Galβ1-ocetyl (FG-ocetyl), and substantially lower activity toward Fucα1,2Galβ1,3GlcNAcα1-pNP (FGGn-pNP) (Fig. 2B). This trend was similar to that of partially purified native AgtA (11), indicating that this unusual pattern of relative activity, in which the longer and more native glycan is less active, was a property of the interaction between the substrates and the catalytic domain itself. MALDI-TOF-MS analysis of the products from reactions carried to near completion revealed that only a single Gal residue was added by any of the enzyme preparations to the small glycan substrates (Fig. 2C), confirming previous observations (11).

AgtA-FL was almost 10-fold more active toward a 400-fold lower concentration of FGGn-Skp1 compared with FG-octyl (Fig. 2B). In contrast, AgtA-CAT was severely deficient in modification of the natural FGGn-Skp1 substrate, exhibiting <0.2% activity of AgtA-FL. This low activity was, however, measurable at prolonged times and greater AgtA-CAT concentration. Based on extrapolation of measurements conducted at different substrate concentrations, the activity of AgtA-CAT toward the Fuc-moiety was comparable irrespective of whether it was attached to Skp1 or pNP (data not shown). Removal of the His_{6} tag by TEV-protease did not affect activity, and a mutant version in which the DXD motif was modified by site-directed mutagenesis, His_{6}AgtA(D132A), was inactive (Fig. 2D). The importance of His-268, which is perfectly conserved in all CAZy GT77 family members to date, was examined by replacement with Ala. AgtA(H268A) exhibited 27% activity toward Skp1 but little activity toward F-pNP (Fig. 2D), and so is not absolutely required for catalysis.

The αGalT activity of a partially purified preparation of AgtA from Dictyostelium toward the disaccharide was previously shown to be sensitive to salt above 50 mM (11). Reinvestigation using purified recombinant proteins showed that intrinsic activity toward FGGn-Skp1 is progressively inhibited by increasing NaCl, reaching 60% at 450 mM NaCl, whereas F-pNP was unaffected (Fig. 2E). The difference in salt sensitivity toward small molecule substrates, between partially and fully purified AgtA, suggests the existence of a modifying factor in cells but was not investigated further. Because only the reaction with FGGn-Skp1 depends on the WD40 repeat domain, its function might be selectively sensitive to salt.
Non-enzymatic Binding Function of a Skp1 Glycosyltransferase

FIGURE 2. Domain dependence of AgtA α-GalT activity. A, His6-AgtA-FL was incubated with a series of FGGn-Skp1 concentrations in the presence of 40 μM UDP-[3H]Gal and 50 mM NaCl for 30 min. Incorporation of [3H]Gal was quantified by TCA precipitation. The curve was derived by non-linear regression according to the Michaelis-Menten model. B, His6-AgtA-FL or His6-AgtA-CAT was incubated in the presence of 10 μM UDP-[3H]Gal, 50 mM NaCl, and the indicated acceptor substrate for 30 min. Incorporation was quantitated by Sep-Pak C18 capture (pNP and octyl aglycons) or TCA precipitation (Skp1). His6-AgtA-CAT protein concentration was 10-fold higher than that of His6-AgtA-FL to yield similar activity toward F-pNP. Results are reported as the mean of triplicate reactions ± S.D. C, MALDI-TOF-MS of the reaction of His6-AgtA-FL or His6-AgtA-CAT with FG-octyl. Expected and experimental m/z values ([M + Na]+) for FG-octyl and the potential products GFG-octyl and GGFG-octyl are indicated. Other ions represent contaminants, which are common in the low mass range for MALDI-TOF-MS experiments. D, activities of AgtA (tag removed), His6-AgtA(D132A), and His6-AgtA(H268A), relative to full-length His6-AgtA, toward substrates F-pNP and FGGn-Skp1 were determined as in panel B. E, salt dependence of α-GalT activity of AgtA-FL toward F-pNP or FGGn-Skp1 was assayed as in B. Average values ± S.E. of triplicate reactions are reported. F, effect of non-substrate isoforms of Skp1 on AgtA activity, measured as transfer of [3H] from UDP-[3H]Gal to 0.25 μM FGGn-Skp1. Error bars represent ± S.D. for assays conducted in triplicate, which are representative of 3 independent trials. G, α-GalT reactions with 0.15 μM FGGn-Skp1 or 840 μM FG-octyl were supplemented with 0.31 μM His6-AgtA-WD40, and the results are reported as fold-activation relative to activity in its absence. Data are reported as representative mean ± S.D.
The WD40 repeat domain commonly mediates protein-protein interactions, and may recognize a second site on Skp1 to facilitate substrate recognition. This model predicts that unmodified Skp1, which lacks the trisaccharide required to interact with the active site, can compete with FGGn-Skp1 because of the second site interaction. Unmodified Skp1 reduced activity in a manner consistent with competitive inhibition, with 50% inhibition occurring at 25-fold excess over FGGn-Skp1 (Fig. 2E). Fully modified GGGFGn-Skp1, the product of the AgtA reaction, was also inhibitory but in a manner that plateaued near 30% inhibition at the same concentration. Thus the proposed second site that interacts with AgtA appears to be present in all Skp1 isoforms, but is less active in GGGFGn-Skp1, consistent with a need to separate from the enzyme to complete the catalytic cycle.

To more directly examine the importance of the WD40 domain, the effect of a recombinant preparation (Fig. 1C) was examined when introduced into AgtA reactions. AgtA-WD40 modestly stimulated the activity of AgtA-CAT toward FGGn-Skp1 under standard conditions (50 mM NaCl), but not of AgtA-CAT toward FG-octyl, or AgtA-FL toward FGGn-Skp1 (Fig. 2G). The finding that the unmodified WD40 repeat domain potentiates activity of the free catalytic domain for only the Skp1 substrate implies a direct role in Skp1 recognition, although more complex interpretations cannot be excluded. The WD40 domain may mediate the hypothesized salt-sensitive second-site interaction to facilitate Skp1 recognition in the cell.

**Skp1 Activation of AgtA Enzyme Activity**—The two-site hypothesis stipulates that the catalytic and WD40 repeat domains interact with separate sites on Skp1. To test for functional synergy of these proposed interactions, the effect of unmodified Skp1, lacking the acceptor Fuc that interacts with the active site, on catalytic modification of free F-pNP was examined. Remarkably, Skp1 exerted substantial concentration-dependent activation of F-pNP modification (Fig. 3A) such that, at saturation, activity of AgtA-FL toward F-pNP was stimulated 50-fold with activity approaching that toward the native substrate FGGn-Skp1 when analyzed at concentrations near their respective $K_m$ values. A comparison of acceptor substrates showed that 50 nM Skp1 activated the galactosylation of F-pNP, FG-octyl, and FGGn-pNP by 30-, 16-, and 17-fold, respectively (Fig. 3, A and B). A kinetic analysis of the dependence of the activated reaction on the F-pNP concentration showed a $K_m$ of 6 mM, similar to the previously reported value of 4 mM in the absence of Skp1 (11), indicating that Skp1 improved the reaction velocity rather than the affinity toward F-pNP. MALDI-TOF-MS analysis of a scaled-up activated reaction showed that still only a single Gal residue was added to F-pNP or FG-octyl (Fig. 2C). Skp1 activation exhibited hyperbolic saturation expected of a single binding event with an apparent $K_D$ of 60 nM. Consistent with the involvement of the WD40 repeat domain, Skp1 did not activate AgtA-CAT activity toward F-pNP (Fig. 3B). Furthermore, activation was suppressed at high salt (Fig. 3C), suggesting that activation was mediated by the same salt-sensitive, WD40-dependent second-site interaction that supports high activity toward FGGn-Skp1. HO-Skp1 and Gn-Skp1 were similarly effective at activating catalysis (Fig. 3A). In contrast, the product of the reaction, GGGFGn-Skp1, had no effect (Fig. 3B), suggesting that terminal galactosylation altered the second-site interaction, or the effect of the interaction on AgtA activation.

**AgtA Is a Novel Skp1-binding Protein**—Physical interactions between AgtA and Skp1 were investigated as a potential basis for the AgtA activation. Initial studies were conducted at pH 7.2 in a buffer supplemented with 50 mM NaCl, which correspond to optimal αGalT activity and to conditions reported for the Dictyostelium cytoplasm (22–24). Analysis of a 1:1 molar mixture of AgtA-FL and unmodified Skp1 by size exclusion chromatography revealed a novel peak that eluted earlier than AgtA-FL or Skp1 alone (Fig. 4A), at an elution position consistent with formation of a 1:1 complex at an estimated peak concentration of 2 μM. However, the assessment of stoichiometry is uncertain because AgtA-FL (Mr, 77,440) eluted after the Mr, 66,000 standard, which suggests retardation because of the interaction with resin, and Skp1 (Mr, 18,710) eluted before the

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**FIGURE 3. Skp1 activation of AgtA.** A, activity was measured based on the transfer of [3H] from UDP-[3H]Gal to 3.7 mM F-pNP by His6AgtA-FL, in the presence of the indicated concentrations of non-substrate Skp1 isoforms. The curve was fitted by non-linear regression modeled as a single site binding interaction. Data are reported as representative mean ± S.D. B, fold-activation of AgtA-FL and AgtA-CAT activity toward FG-octyl, FGGn-pNP, or FGGn-Skp1 in the presence of the indicated concentration of unmodified Skp1. C, reactions were prepared as in A with the indicated concentrations of unmodified Skp1 and NaCl. Total activation levels of triplicate reactions relative to AgtA-FL alone are plotted.


**Non-enzymatic Binding Function of a Skp1 Glycosyltransferase**

**A**, 0.1 nmol of His_{6}AgtA-FL, unmodified Skp1, or His_{6}AgtA-FL preincubated with unmodified Skp1 were chromatographed on a Superdex 200 column in 50 mM NaCl, and monitored based on A_{280}. Peak elution volumes are denoted. Elution positions for porcine thyroglobulin (M_{r} 670,000), alcohol dehydrogenase (150,000), bovine serum albumin (66,000), and carbonic anhydrase (29,000) are denoted. **B**, densitometric analysis of Western blots of Skp1 elution positions from A and an identical experiment in 150 mM NaCl. The elution positions of free AgtA-FL and the AgtA-Skp1 complex, which were not affected by the NaCl concentration, are denoted. C, gel filtration analysis of the interaction of AgtA-FL with fully modified Skp1 (GGFGGn-Skp1) as in A. Profiles shown in panels A–C are representative of 2–3 independent trials. **D**, parallel Western blots of cross-linking reactions performed on AgtA-FL (600 nm), unmodified Skp1 (600 nm), or a mixture with each at 600 nm. Samples were reacted with 0.25 mM DSS for 10 min at the indicated NaCl concentrations. Lower order multimers of Skp1 and AgtA are denoted with asterisks, and the position of a novel AgtA- and Skp1-dependent band is denoted with an arrow. Imaging of this region of the blot was reoptimized with respect to this band. The percentage of Skp1 and AgtA found in this band is plotted as a function of NaCl concentration above the Western blot lanes. Data shown are representative of 3 independent trials.

The nature of the complex was examined further by cross-linking studies with DSS, a bifunctional amine-reactive reagent incorporating an 8-carbon (11.4-Å) spacer. Using conditions optimized for DSS concentration and reaction time, SDS-PAGE/Western blotting showed that Skp1 and AgtA could be trapped as dimers and higher order multimers in 50 mM NaCl (Fig. 4D, lanes 2 and 14, asterisks). In comparison, the Skp1/AgtA mixture exhibited less oligomerization of Skp1, and a unique novel band with an apparent M_{r} of 108,000 was detected by both anti-Skp1 and anti-AgtA antibodies (arrow, in lanes 7 and 17). The intensity of this band was salt-sensitive (lanes 6–9 and 16–19, and histograms above), suggesting that it corresponds to the complex detected by gel filtration. The cross-linked complex appeared to be less salt-sensitive than was the α-galactosylation of FGGn-Skp1 (Fig. 2C), the activation of AgtA catalytic activity (Fig. 3), and the gel filtration complex (Fig. 4B), but this is likely because cross-linking is irreversible and thus cumulative over time. Although the fractions of total Skp1 and AgtA that could be trapped by cross-linking was low, it is likely that the majority of Skp1 was complexed with AgtA prior to denaturing SDS-PAGE. This is proposed based on the smaller fraction of Skp1 oligomers that could be cross-linked in the presence of AgtA because of competitive interaction.
Interestingly, interacting with Skp1 did not appear to impact AgtA oligomerization. The apparent $M_r$ of the novel band (108,000) lies between that of the 1:1 Skp1-AgtA complex (96,200) and a 2:1 complex (114,910). The observation that complex formation reduces the level of Skp1 dimer that can be detected (Fig. 2A), as revealed by disrupting $agtA$ and tested for its influence on AgtA-mediating enzymatic activity, suggests that not only does formation of the Skp1-Fbs1 complex block processing of Skp1 by AgtA, the interaction with Fbs1 is tight with high affinity, consistent with the previously reported $K_D$ value of 25 nM (20). In addition, Skp1 may exist as a dimer in vivo, as Fbs1 may act as a monomer as suggested by the substoichiometric nature of the interaction (20), but the possibility that the calculated extinction coefficient used to determine protein concentrations is inaccurate was not excluded.

The effect of Fbs1 on Skp1 association with AgtA modification of F-pNP is shown in Fig. 5C. Increasing concentrations strongly inhibited activation by either Skp1 or Gn-Skp1. As observed for inhibition of the AgtA reaction with FGGn-Skp1 (Fig. 5B), inhibition of activation by Fbs1 was strong and appeared to be substoichiometric. The similar effects of Fbs1 on both the substrate activity of Skp1 and activation by Skp1 of AgtA is consistent with a model in which they are both mediated by the same binding event, which may overlap with the site of interaction of Skp1 with the Fbs1 F-box.

A Nonenzymatic Role of AgtA in Dictyostelium Development—Based on previous studies of the $agtA$-disruption strain (6), AgtA contributes to the ability of cells to sense O$_2$ concentrations (8–10%) to allow culmination, i.e., the reorganization and differentiation of the migrating slug into a fruiting body consisting of spores supported on a cellular stalk. In the absence of AgtA, culmination can occur but requires a higher level of O$_2$ (18–21%), almost as much as that required by phyA$^-$ strains. Interestingly, partial functionality (culmination at ~12–15% O$_2$) is achieved by hydroxylation of Skp1 $\pm$ αGlcNAc addition, as revealed by disrupting gnt1 or $agtA$ (4). We reasoned that if AgtA contributes a function in addition to its role in α-galactosylation, it might be revealed in a genetic background in which the substrate for AgtA is not generated. Thus we created an $agtA^{−}$/gnt1$^{−}$ double knock-out, by disrupting gnt1 in an $agtA^{−}$ background. The double mutant lacked detectable Gnt1 activity in extracts, accumulated HO-Skp1, like gnt1$^{−}$ cells (Fig. 6C), and failed to culminate at even the highest O$_2$ level tested, 70%. An example at 21% O$_2$ is shown in Fig. 6A, where both gnt1$^{−}$ and $agtA^{−}$ cells formed normal appearing fruiting bodies (top row), as expected because O$_2$ was above their thresholds, but the double mutant (bottom left) remained in the slug-like stage. Failure to differentiate into spores is documented in Fig. 6B. Because the path to generating the double mutant involved dis-

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**FIGURE 5. Relationship between AgtA and other Skp1 interactors.** A, effect of AgtA on Skp1 PhyA and Gnt1 reactions. Reactions containing 0.1 μM unmodified Skp1 (left panel) or HO-Skp1 (right panel) were supplemented with 0, 0.1, or 0.5 μM His$_6$AgtA-FL or His$_6$AgtA(D132A) as indicated. B, effect of Fbs1 on AgtA-mediated α-galactosylation of Skp1. FGGn-Skp1 (0.5 μM) was preincubated with 0–50 μM His$_6$Fbs1, and its α-galactosylation was tested as described in the legend to Fig. 2A. C, effect of Fbs1 on Skp1 activation of AgtA activity. AgtA modification of 1 mM F-pNP was measured in the presence of 0–50 μM His$_6$AgtA, preincubated with 0–50 μM His$_6$Fbs1, and its α-galactosylation was tested as described in the legend to Fig. 2A.
rupturing \textit{gnt1} in a previously validated \textit{agtA} strain, specificity of the disruption was addressed by a rescue experiment in which \textit{gnt1} was restored by \textit{cotB}-promoter dependent expression of Gnt1 or catalytically inactive Gnt1 in prespore cells. As shown in Fig. 6, \textit{A} and \textit{B}, normal development was restored by active but not inactive Gnt1, demonstrating that the original change was specific to \textit{gnt1}, and confirming (4) that the \(\alpha\text{GlcNAcT}\) activity of Gnt1 is important for its function in culmination.

The results imply that AgtA is required for culmination when Skp1 is not glycosylated by a mechanism that does not involve its catalytic activity, which might involve the novel Skp1 binding activity documented \textit{in vitro} above.

In a second approach, the effect of semi-constitutive overexpression of native and catalytically inactive mutant AgtA was examined in \textit{agtA} and normal (Ax3) cells. As previously reported (6), AgtA-overexpression (AgtA\textsuperscript{oe}) rescued culmina-
Non-enzymatic Binding Function of a Skp1 Glycosyltransferase

Analysis of Skp1 modification in these strains showed that 

occurred at even 70% O2, were distinct from those seen in 

The cellular glycome is defined by the non-templated actions 

DISCUSSION

The cellular glycome is defined by the non-templated actions of GTs and glycosidases and, therefore, the varied mechanisms of their selectivity and regulation are of great interest. AgtA exhibits features that are, based on current knowledge, unusual among glycosyltransferases, including second-site recognition of the polypeptide carrier of the trisaccharide substrate, and catalytic activation via polypeptide recognition. Domain analysis indicates that these novel properties depend on an interplay between the two parts of AgtA, in which recognition of the Skp1 polypeptide by the C-terminal WD40 repeat domain activates the catalytic domain. Remarkably, polypeptide recognition includes all isoforms of Skp1, allowing for AgtA to influence Skp1 even before it becomes a glycosylation substrate. In turn, α-galactosylation of the Skp1 polypeptide alters recognition by AgtA, possibly allowing for termination of the non-enzymatic influence on Skp1 concomitant with acquisition of new func-

FIGURE 6. Non-enzymatic activity of AgtA in vivo. A. Representative images of development of wild-type (w/t) and mutant strains after 42 h at 21% O2. Normal and single knock-out strains are in the top row. The white oval encircles a typical fruiting body showing the spore-containing sorus linked to the stalk. The double agtA−/gnt1− mutant strain HWS20 and strains complemented with normal or enzymatically inactive Gnt1(D102A) expressed under control of the prespore cell specific cotB promoter are in the bottom row. B, quantification of culmination, based on spore number, as a function of O2 in the atmosphere. C, Western blot analysis for HO-Skp1 in S100 extracts from growing cells of the agtA−/gnt1− double mutant strain compared with other mutant strains. The dashed line shows where other lanes from the same blot were omitted. The lower panel shows a region of the blot after Coomassie Blue staining, as a loading control. D, representative images of development for 42 h at 21% O2 of strains constitutively expressing His6_AgtA or enzymatically inactive His6_AgtA(D132A) under control of the semi-constitutive dcsA promoter. E, a Western blot of growing cell S100 extracts from selected overexpression clones was probed with anti-AgtA, and with anti-Skp1 as a loading control. F, spore number as a function of the O2 concentration from strains shown in D. Data in B and D are representative of ≥3 independent trials where each strain was analyzed in parallel. G, examples of individual fruiting bodies from normal and AgtA(D132A)−/− strains from panel D, imaged using fluorescent Calcofluor staining for the cellulose cell walls of stalk and spore cells. Ax3/w/t shows a typical slender stalk and spores that disperse during preparation. The AgtA(D132A)−/− strain culminates to form a poorly organized bloated stalk, and the prespore cells remain contiguous because of failure to differentiate into spores. H, Western blots of S100 cell extracts from agtA−, His6_AgtA−, and His6_AgtA(D132A)−/− cells (in an Ax3 background) grown at 21% O2 were probed with anti-Skp1 antibodies that recognize total Skp1 (mAb 4E1), unmodified Skp1 (UOK85), or HO-Skp1 (UOK85). The slower migration of fully modified Skp1 relative to unmodified or hydroxylated Skp1 is denoted by the double bar.
A Role for the WD40 Repeat Domain in Catalytic Activation—The importance of the WD40 repeat domain in Skp1 recognition is indicated by its necessity for AgtA to efficiently galactosylate FG Gn-Skp1 at low, physiological concentrations (Fig. 2A). In fact, even the free WD40 repeat domain can promote activity of the catalytic domain toward FG Gn-Skp1 (Fig. 2G). Because the WD40 repeat domain is not required to modify the free glycan, we propose that it binds to Skp1 at a site separate from the interaction of the active site with the glycan acceptor (Fig. 7, step 4). The existence of the second site is supported by detection of a stable complex between AgtA-FL and Skp1 under native conditions by gel filtration chromatography in the absence or presence of the co-substrate UDP-Gal (Fig. 4, A and B), and a complex that could be trapped by DSS cross-linking (Fig. 4D). The binding interaction detected by gel filtration and cross-linking likely mediates Skp1 substrate recognition and AgtA activation because of their similar salt sensitivities. This might assist in substrate recognition in the cell, as proposed for the Dictyostelium myosin heavy chain kinases MhcA and MhcK8 whose WD40 repeat domains target their myosin substrate (26). Nevertheless, AgtA-FL also forms a stable complex with the reaction product GGFGGn-Skp1 (Fig. 4C), which raises the question of how the product dissociates to permit a new catalytic cycle.

Second-site recognition mechanisms that contribute to acceptor substrate specificity have been described in several membrane-bound GTs including βGalNAcT3 and -T4 (27), polysialyltransferase-I (18), polypeptide αGalNAc transfersases (16), and lysosomal enzyme GlcNAc-phosphotransferase (28). The mechanism of second-site recognition in these enzymes appears also to involve distinct domains, or subunits, and βGalNAcT3 and -T4 binding to their polypeptide substrates is sensitive to higher ionic strength (29) as for AgtA. The mechanism of product release for these enzymes is poorly understood and is usually assumed to involve a simple equilibrium process. The model that second-site recognition facilitates targeting of Skp1 seems sensible for an enzyme whose substrates are not concentrated within secretory pathway vesicles as for most glycosylation pathways, but a need for such as mechanism can be questioned considering that Skp1 is thought to be the only glycoprotein in the cytoplasmic compartment that bears the acceptor Fuc residue (2). Furthermore, the earlier GTs in the pathway, Gnt1 and PgtA, are also highly specific for Skp1 but lack obvious domains like that in AgtA to mediate this specificity (6, 19). Thus the WD40 repeat domain may have other functions.

A Role for the WD40 Repeat Domain in Substrate Recognition—The binding of AgtA-FL to substrate precursor isoforms of Skp1 (Fig. 4B) appears to be irrelevant to catalysis, because dissociation appears to be required for prior hydroxylation and αGlcNAcylation (Fig. 5A). This is reminiscent of the effect of the FBP FbxA, whose binding to Skp1 also inhibits its modification by Phya (7). This effect potentially occurs in vivo because overexpression of AgtA led to an accumulation of unmodified Skp1 in cells (Fig. 6H). Even higher levels accumulated using catalytically inactive AgtA(D132A), and HO-Skp1 was also detected. Thus AgtA is a novel Skp1-binding protein that may buffer against the actions of Phya and Gnt1 toward Skp1 (Fig. 7, step 2).

The binding affinity of Skp1 is sufficient to inhibit the reaction of AgtA with FG Gn-Skp1 (Figs. 2F and 7, step 2). The product GGFGGn-Skp1 was less inhibitory, corresponding to altered gel filtration behavior (Fig. 4C) and suggesting an altered binding mode that may be related to product release.

Competition between AgtA and an FBP was examined using guinea pig Fbs1, which forms a tight binding complex with Dictyostelium Skp1 based on gel filtration (25). Fbs1 efficiently blocked activation of AgtA by Skp1 (Fig. 5C), which could be explained by formation of a competing Fbs1-Skp1 complex. Indeed, Fbs1 has a lower KD for mammalian Skp1, 25 nM (20), than does AgtA (as measured indirectly) for Dictyostelium Skp1 (~60 nM). If AgtA and FBPs compete for binding Skp1 in cells, then they could dampen each other’s activities and jointly slow the hydroxylation and glycosylation of Skp1 in cells (Fig. 7, steps 1 and 3).

Importance of the Skp1 Binding Function of AgtA in Cells—Genetic studies suggest that the AgtA binding activities observed in vitro impact Skp1 in cells. Disruption of agtA has an anomalously severe effect on culmination in which O2 dependence rises to a level almost as high as that required for phya-null cells to culminate (6). This contrasts with the intermediate level of O2 required for gnt1- or pgtA-mutant cells, and the low level of O2 required for wild-type cells that can assemble the pentasaccharide. An even more severe effect was observed when agtA was deleted in a gnt1-null background (Fig. 6, A and 6B).
B), in which culmination could not be rescued at any O$_2$ level tested (up to 70%). Because the AgtA catalytic activity is irrelevant in the absence of a substrate that cannot be assembled without Gnt1, the failure to culminate suggests a need for the Skp1 binding activity of AgtA that is exacerbated by the inability to escape the effect of AgtA by completion of glycosylation. Indeed, the importance of glycosylation per se is emphasized by the ability of highly (23-fold) overexpressed AgtA-CAT to complement the absence of AgtA (5, 6). In complementary fashion, the binding activity of AgtA lessens the requirement for O$_2$, and may be a mediator of the O$_2$-sensing mechanism that rescues culmination of phyA$^-$ cells at high O$_2$ levels (19). Absence of the binding activity of AgtA may alter the kinetics of assembly of new SCF complexes containing different FBPs during developmental transitions (Fig. 7).

Overexpression of catalytically inactive AgtA(D132A) in a wild-type background was also deleterious with a selective and dramatic effect on late culmination. Proliferation and development up to the slug stage were morphologically normal and culmination was initiated at the normal time. However, the stalk that formed was flaccid with a larger volume (Fig. 6G) suggesting that a larger fraction of cells contributed to its formation, and the fewer cells that rose to the apex failed to sporulate (Fig. 6, D and F). This defect was not rescued at even the high O$_2$ levels that normally override Skp1 modification defects. The failure to sporulate under high O$_2$ is reminiscent of the inability of gnt1$^{-}$ or pgtA$^{-}$ cells to sporulate in O$_2$-limited submerged development (30). It is unlikely that the effect of AgtA(D132A) overexpression is due solely to hypomodified Skp1 that accumulates (Fig. 6H), because glycosylation mutants that accumulate these isoforms can form almost normal fruiting bodies at high O$_2$ (19, 4). The O$_2$-independent effect of AgtA(D132A) overexpression is most easily explained by the advantage that a high concentration affords the protein in competing with F-box interactions, which is expected to impede timely removal of critical regulatory proteins during culmination and terminal cell differentiation. Overexpression of active AgtA had, in contrast, little effect, suggesting that the ability to α-galactosylate Skp1 ultimately terminates the inhibition rendered by AgtA-binding. Because Skp1 is the only enzymatic substrate that has been detected in biochemical complementation studies (6) and agtA disruption suppresses disruption of one of the two Skp1 genes (8), the aberrant culmination likely involves Skp1 as a biochemical target despite the possibility that AgtA WD40 repeats bind other proteins as well.

Possible Impact of AgtA on SCF Complex Assembly—the importance of the Skp1 binding function of AgtA is consistent with previous evidence that AgtA and Skp1 are expressed at similar levels in vivo and to co-purify over DEAE- and phenyl- Sepharose chromatography columns (11), but future studies are needed to address the stability of AgtA binding to Skp1 in cells and potential interactions with other proteins. The optimal buffer for in vitro binding and catalysis contained 50 mM NaCl, and its osmolarity of 156 mosM lies within the physiological range of the Dictyostelium cytoplasm, which has been estimated at 85 to 200 mosM (23, 24). The dependence of binding on ionic strength raises the possibility that changes in osmolarity in response to varied environments may regulate the association of AgtA and Skp1.

Dictyostelium development involves global changes in protein profiles, which doubtless involve regulated protein turnover (e.g. Ref. 31). E$_3^{SCF}$-Ub-ligases (1) likely contribute to turnover as prescribed by developmentally programmed changes (32) in the expression of more than 50 predicted FBPs. Potentially, AgtA promotes equilibration of interactions between Skp1 and newly synthesized FBPs by serving as an E3 FBP exchange factor analogous to how Cand1 acts like an exchange factor to control Skp1/Cul1 interactions (33–35). Cand1, which binds to Cul1, helps dissociate the Skp1/FBP dimer from Cul1, but promotes SCF function in vivo, via a mechanism of mutually exclusive binding of Cand1 and Skp1 to Cul1. As evidenced by the effects of too much or too little binding competent AgtA in cells (Fig. 6), mutually exclusive binding of AgtA and an FBP to Skp1 may facilitate the switching and activation of FBPs during developmental transitions in a manner that is kinetically modulated by hydroxylation and glycosylation in response to environmental factors such as O$_2$ and internal metabolic factors such as sugar availability (Fig. 7). Structural studies to elucidate the molecular contacts that underlie these interactions are needed to further evaluate this model.

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