A Gene of the β3-Glycosyltransferase Family Encodes N-Acetylglucosaminyltransferase II Function in Trypanosoma brucei

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The bloodstream form of the human pathogen Trypanosoma brucei expresses oligomannose, paucimannose, and complex N-linked glycans, including some exceptionally large poly-N-acetyllactosamine-containing structures. Despite the presence of complex N-glycans in this organism, no homologues of the canonical N-acetylglucosaminyltransferase I or II genes can be found in the T. brucei genome. These genes encode the activities that initiate the elaboration of the Manα1–3 and Manα1–6 arms, respectively, of the conserved trimannosyl-N-acetylchitobiosyl core of N-linked glycans. Previously, we identified a highly divergent T. brucei N-acetylglucosaminyltransferase I (TbGnTII) among a set of putative T. brucei glycosyltransferase genes belonging to the β3-glycosyltransferase superfamily (Damerow, M., Rodrigues, J. A., Wu, D., Güther, M. L., Mehler, A., and Ferguson, M. A. (2014) J. Biol. Chem. 289, 9328–9339). Here, we demonstrate that TbGT15, another member of the same β3-glycosyltransferase family, encodes an equally divergent N-acetylglucosaminyltransferase II (TbGnTIII) activity. In contrast to multicellular organisms, where GntTII activity is essential, TbGnTII null mutants of T. brucei grow in culture and are still infectious to animals. Characterization of the large poly-N-acetyllactosamine containing N-glycans of the TbGnTII null mutants by methylation linkage analysis suggests that, in wild-type parasites, the Manα1–6 arm of the conserved trimannosyl core may carry predominantly linear poly-N-acetyllactosamine chains, whereas the Manα1–3 arm may carry predominantly branched poly-N-acetyllactosamine chains. These results provide further detail on the structure and biosynthesis of complex N-glycans in an important human pathogen and provide a second example of the adaptation by trypanosomes of β3-glycosyltransferase family members to catalyze β1–2 glycosidic linkages.

The African trypanosomes are protozoan parasites that cause human African sleeping sickness and Nagana in cattle. The parasite undergoes a complex life cycle between the mammalian host and the blood-feeding tsetse fly vector (Glossina sp.). Throughout this life cycle, Trypanosoma brucei is coated by glycosylphosphatidylinositol (GPI)-anchored proteins. The bloodstream form of the parasite in the mammalian host is covered by a coat of 5 × 10⁶ variant surface glycoprotein (VSGs) homodimers and evades the immune system by replacing one VSG coat by another, in a process known as antigenic variation (1–4). The VSG GPI anchors contain side chains of 0–6 Gal residues, depending on the VSG variant (5–7) and between 1 and 3 N-linked glycans. The latter can be of oligomannose, paucimannose, or complex types (6, 8, 9). T. brucei expresses numerous other GPI-anchored and transmembrane glycoproteins at the cell surface, in the flagellar pocket, and in the intracellular endosomal/lysosomal system, some of which are life cycle stage-specific or display life cycle stage-specific glycosylation differences. For example, the transmembrane invariant surface glycoproteins ISG65 and ISG75 (10) and the GPI-anchored flagellar pocket ESAG6/ESAG7 heterodimeric transferrin receptors (11–13) are specific to the bloodstream life cycle stage, whereas the major lysosomal glycoprotein p67 is common to bloodstream and procyclic stages but contains complex N-glycans only in the bloodstream stage (14). This control of stage-specific glycosylation resides primarily at the level of oligosaccharyltransferase expression (15). Thus, in the bloodstream form of T. brucei both the TbSTT3A and TbSTT3B genes are expressed, and it appears that TbSTT3A co-translationally scans for glycosylation sequons in relatively acidic local

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The abbreviations used are: GPI, glycosylphosphatidylinositol; VSG, variant surface glycoprotein; sVSG, soluble form VSG; poly-LacNAc, poly-N-acetylglucosamine; GT, glycosyltransferase; GnT, GlcNAc transferase; PAC, puromycin acetyltransferase; HYG, hygromycin phosphotransferase; Hex, hexose; HexNAc, N-acetylated hexosamine; HILIC, hydrophilic interaction liquid chromatography; PMAA, partially methylated alditol acetate; TLC, tosylsylchloromethyl ketone hydrochloride; CAZy, carbohydrate-active enzyme; bis-Tris, 2-[bis(2-hydroxyethyl)amino]-2-[hydroxyethyl]propane-1,3-diol.
Nevertheless, they identify the parasite genes as belonging to the (G/A), but not (G/A)'(V,F)XK motifs in the T. brucei GTs. All 21 putative acid sequences consistent with being UDP-sugar-dependent glycosyltransferase (GT) genes in the parasite. We previously reported a family of 21 genes with predicted amino acid sequences with being UDP-sugar-dependent glycosylation pathways are essential for the survival of the parasite in both the bloodstream and procyclic form life stages (20–22). Similarly, the creation of UDP-glucose 4'-epimerase (TbGalE) conditional null mutants showed that this gene, and hence UDP-Gal, is essential for the survival of the parasite in both the bloodstream and procyclic form life stages (20–22). Similarly, the creation of UDP-GlcNAc pyrophosphorylase (TbLAP) and glucosamine 6-phosphate N-acetyltransferase (TbGNA) conditional null mutants has shown that UDP-GlcNAc is essential for bloodstream form of T. brucei (23, 24). From these experiments, it is possible to conclude that one or more of the UDP-Gal- and UDP-GlcNAc-dependent glycosylation pathways are essential to the parasite. This has provided the impetus to identify and characterize the UDP-Gal- and UDP-GlcNAc-dependent glycosyltransferase (GT) genes in the parasite. We previously reported a family of 21 genes with predicted amino acid sequences consistent with being UDP-sugar-dependent GTs. All 21 putative T. brucei GT amino acid sequences are similar to those of the mammalian β3GT family (25). The mammalian β3GT family includes Gal, Glc, glucuronic acid, GlcNAc, and GalNAc β-3 transferases, and its members contain N-terminal transmembrane domains followed by three conserved motifs as follows: (I/L)RXXXWG, (F/Y)(V/L/M)XXXD, and (E/D)D(A/V)(Y/F)XGX(C/S). The comparable motifs in the T. brucei genes are slightly different, WG, Y(I,V,F)XXDXXD, and ED(A/V/I/L/M)(M/L)(G/A), but nevertheless, they identify the parasite genes as belonging to the β3GT superfamily (26). One of these genes (TbGT8) encodes a β1–3 GlcNAc transferase and another (TbGT3) a β1–3 Gal transferase that modifies the complex GPI anchor side chains of the procyclins (the major surface glycoproteins of the procyclic life cycle stage) (26–28). However, we recently reported that another gene (TbGT11) encodes a β1–2 GlcNAc transferase that performs a similar role to members of the N-acetylglucosaminyltransferase I family, in that it transfers GlcNAc in β1–2 linkage to the 6-arm of Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc (1).

Here, we report that another T. brucei β3GT superfamily gene member (TbGT15) encodes another β1–2 GlcNAc transferase that was already localized to the Golgi apparatus (29, 30). It performs a similar role to members of the N-acetylglucosaminyltransferase II family in that it transfers GlcNAc in β1–2 linkage to the 6-arm of Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc, emphasizing the highly divergent nature of the trypanosome genes involved in structurally conserved aspects of complex N-glycan biosynthesis.

**Experimental Procedures**

*Cultivation of Trypanosomes—* T. brucei brucei strain 427 bloodstream form parasites, expressing VSG variant 221 and transformed to stably express T7 polymerase and the tetracycline repressor protein under G418 antibiotic selection (31), were used in this study. This genetic background will be referred to as wild-type (WT). Cells were cultivated in HMI-9 medium containing 2.5 µg/ml G418 at 37 °C in a 5% CO2 incubator as described previously (31).

*DNA and RNA Isolation and Manipulation—* Plasmid DNA was purified from *Escherichia coli* (α-select chemically competent cells, Bioline, London, UK) using Qiagen Miniprep or Maxiprep kits, as appropriate. Gel extraction and reaction clean up was performed using QIAquick kits (Qiagen). Custom oligonucleotides were obtained from Eurofins MWG Operon or the Dundee University oligonucleotide facility. *T. brucei* genomic DNA was isolated from ∼2 × 10⁷ bloodstream form cells using DNAzol (Helena Biosciences, UK) by using standard methods. *T. brucei* mRNA was extracted from 1 × 10⁷ cells using RNeasy RNA extraction kit (Qiagen).

*Generation of Gene Replacement Constructs—* The 517-bp 5' and 454-bp 3' UTR sequences next to the Tb427.7.300 ORF were PCR-amplified from genomic DNA using Pfu DNA polymerase with primers 5'-cggttctgcaagtccgagcaaaagcgagc-3' and 5'-gttaactacgggacgtgtaaactttttactccacg-3' and 5'-gacgctcgaatttttaaacagggataggctc-3' and 5'-attcagatcttttactctttaatc-3' as forward and reverse primers, respectively. The two PCR products were used together in a further PCR to yield a product containing the 5'-UTR linked to the 3'-UTR by a short HindIII (underlined), Pmel (italics), and BamHI (underlined) cloning site and NotI and SacI restriction sites at each end (capital letters). The product was cloned between the NotI and SacI sites of the pGEM-5Zf(+) vector (Promega).

The hygromycin phosphotransferase (HYG) and puromycin acetyltransferase (PAC) drug-resistance genes were then introduced into the targeting vector via the HindIII and PacI cloning sites. For re-expression of Tb427.7.300, the ORF was PCR-amplified from genomic DNA with the primer pair 5'-agaaacttgggatgcttcatttcg3' and 5'-ctgcgcatgatgatgatgc-3' and cloned into pLEW100-Phleo (31).

For overexpression of full-length TbGT15 with a C-terminal 3× HA epitope tag, a plasmid was generated based on the trypanosome expression vector pLEW82 (31). *TbGT15 ORF* was amplified from *T. brucei* genomic DNA and the primers 5'-gacgctcgaatttttaaacagggataggctc-3' and 5'-ttcagatcttttactctttaatc-3' and cloned into pLEW20-Phleo (31).

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restriction sites under replacement of the GPIdeAc insert, but retention of the sequence encoding for one HA tag, resulting in the plasmid pLEW82-TbGT15-HA3. The identity of all constructs was confirmed by sequencing.

**Transformation of Bloodstream Form T. brucei**—Constructs for gene replacement and ectopic expression were purified, digested with NotI to linearize, precipitated, washed with 70% ethanol, and re-dissolved in sterile water. The linearized DNA was electroporated into *T. brucei* bloodstream form cells (strain 427, variant 221) that were stably transformed to express T7 RNA polymerase and the tetracycline repressor protein under the control of the *T. brucei* N-Acetylglucosaminyltransferase II promoter. Cell culture and transformation were carried out as described previously (31–33).

**Southern Blotting**—Aliquots of genomic DNA isolated from 100 ml of bloodstream form *T. brucei* cultures (~2 × 10⁸ cells) were digested with EcoRI, resolved on a 0.8% agarose gel and transferred onto a Hybond-N positively charged membrane (GE Healthcare, UK). Highly sensitive DNA probes labeled with digoxigenin-dUTP were generated using the PCR digoxigenin probe synthesis kit (Roche Applied Science) according to the manufacturer’s recommendations and hybridized overnight at 42 °C. Detection was performed using alkaline phosphatase-conjugated anti-digoxigenin Fab fragments and the chemiluminescent substrate CSPD (Roche Applied Science).

**Mouse Infectivity Studies**—Wild-type and *TbGT15* null mutant bloodstream form trypanosomes were grown in HMI-9T media, washed in media without antibiotics, and resuspended at 5 × 10⁸ cells/ml. Groups of five female BALB/c mice were used for each cell line, and 0.1 ml of the suspension above was injected intraperitoneally per animal. Infections were assessed 3 days post-infection by tail bleeding and cell counting using a Neubauer chamber in a phase contrast microscope.

**Semi-quantitative RT-PCR**—To assess the amount of Tb427.7.300 mRNA in the *TbGT15* conditional null mutant cells grown under permissive and non-permissive conditions, RT-PCRs were performed using AccessQuick RT-PCR System (Promega) according to the manufacturer’s recommendations. A *TbGT15* 350-bp fragment was amplified with the primer pair 5′-cactggtgctcggatagtgag-3′ and 5′-catctcccaagacccgggttaaattggg-3′. As a control to ensure similar RNA levels in both samples, primers 5′-aatgtgctggcactctgacacc-3′ and 5′-tagaaccctggccggttccta-3′ amplifying a 448-bp product of dolichyl phosphate mannose synthase (Tb10.70.2610) were used.

**Small Scale sVSG Isolation**—Soluble form sVSG (sVSG) was isolated from 100 ml of cultures containing ~2 × 10⁸ bloodstream form *T. brucei* by a modification of the method of Cross and co-workers (34, 35) as described previously (36). Briefly, cells were chilled on ice, centrifuged at 2500 × g for 10 min, and washed in an isotonic buffer. The pellet was resuspended in 300 μl of lysis buffer (10 mM NaH2PO4 buffer, pH 8.0, containing 0.1 mM tosyllysine chloromethyl ketone hydrochloride (TLCK), 1 μg/ml leupeptin, and 1 mg/ml aprotinin) and incubated for 5 min at 37 °C. The sample was centrifuged at 14,000 × g for 5 min, and the supernatant was applied to a 200-μl DE52 anion exchange column pre-equilibrated in 10 mM sodium phosphate buffer, pH 8.0. Elution was performed with 0.8 ml of 1 M sodium phosphate buffer, pH 8.0, and the eluate was concentrated and dialyzed with water on a YM-10 spin concentrator (Microcon). The final sample of 50–100 μg of sVSG221 was recovered in a volume of 100 μl of water.

**ES-MS Analysis of Intact sVSG**—50 μg of aliquots of sVSG preparations were diluted to ~0.05 μg/μl in 50% methanol, 1% formic acid and analyzed by positive ion ES-MS on a Q-Tof 6520 instrument (Agilent). Data were collected, averaged, and processed using the maximum entropy algorithm of the MassHunter software (Agilent).

**Lectin Blotting of Cell Extracts**—To analyze N-glycosylation of *T. brucei* bloodstream form cells, ~2 × 10⁸ cells were first depleted of VSG by hypotonic lysis (34, 35). For Western blot analysis, residual cell ghosts were solubilized in SDS sample buffer containing 8 M urea, boiled with DTT, separated by SDS-PAGE (~1 × 10⁵ cell eq/lane) on NuPAGE bis-Tris 4–12% gradient acrylamide gels (Invitrogen) and transferred to a nitrocellulose membrane (Invitrogen). Ponceau S staining confirmed equal loading and transfer. Glycoproteins were probed with 1.7 μg/ml biotin-conjugated ricin (RCA-120, Vector Laboratories, UK) in PBS before or after pre-incubation with 10 mg/ml d-galactose and 10 mg/ml α-lactose to confirm specific ricin binding. Detection was performed using IRDye 680LT-conjugated streptavidin and the LI-COR Odyssey infrared imaging system (LI-COR Biosciences, Lincoln, NE).

**Structural Analysis of the Large N-Glycan Fraction**—Bloodstream form cells of wild-type and *TbGT15* null mutant cells were isolated from infected rats and processed as described (19). Briefly, VSG-depleted cell ghosts of ~10⁸ cells eq were solubilized with SDS/urea buffer followed by lectin affinity chromatography using ricin-agarose (RCA-120, Vector Laboratories). N-Glycans from the ricin-binding glycoproteins were released with peptide:N-glycosidase F (Flavobacterium menin gosepticum, Roche Applied Science). N-Glycans were analyzed by GC-MS (Agilent) as trimethylsilylated derivatives. Monosaccharide composition analysis (37). Fractions that eluted in the void volume of the column (the total poly-LacNAc fraction, rich in Gal and GlcNAc) were pooled and used for methylation linkage analysis. After permethylation, acid hydrolysis, NaBD₄ reduction, and acetylation, the resulting partially methylated alditol acetates (PMAAs) were analyzed by GC-MS (Agilent) as described previously (38). Authentic glycans of Galβ1–4GlcNAcβ1–2Manα1–6(Galβ1–4GlcNAcβ1–2Manβ1–3GlcNAcβ1–4GlcNAc, lacto-N-neohexasa Galβ1–4GlcNAcβ1–6(Galβ1–4GlcNAcβ1–3)Galβ1–4Glc, lacto-neotetraose Galα1–4Glcβ1–3Galβ1–4GlcNAc, and Galβ1–6Gal (Dextra Laboratories, UK) were subjected to methylation linkage analysis alongside the experimental samples. Using the PMAA derivative derived from non-reducing terminal galactose residues, common to all of these structures (i.e. 1,5-di-O-acetyl-2,3,4,6-tetra-O-methyl-1-[3H]galactitol), we were able to inter-relate these data and determine the total ion current molar relative response factors for the PMAAs derived from terminal-Gal, 3-O-substituted Gal, 6-O-substituted Gal 3,6-di-O-substituted Gal, 2-O-substituted Man, 3,6-di-O-substituted Man, and 4-O-substituted GlcNAc (1.59, 0.99, 0.27, 0.90, 1.20, 1.00, and 0.23, respectively). These molar relative response fac-
tor values were used to correct the peak integrations of sample PMAA total ion current chromatograms and thus provide molar ratios of the PMAAs in the methylation linkage analyses of the wild-type and TbGT15 null glycan samples.

**GnTII in Vitro Activity Assay**—TbGT15 fused to a C-terminal triple HA tag was overexpressed in *T. brucei* bloodstream form cells. 1 × 10⁶ cells were lysed on ice in 25 mM Tris, pH 7.5, 100 mM NaCl, 1% Triton X-100 containing a mixture of protease inhibitors (CompleteMini, Roche Applied Science), and 0.1 mM TLCK. Expression was confirmed by SDSPAGE and Western blotting. Briefly, 5 × 10⁶ cell eq/lane were separated on NuPAGE bis-Tris 4–12% gradient acrylamide gels (Invitrogen) and transferred to nitrocellulose membrane (Invitrogen). Poncova S staining confirmed equal loading and transfer. Detection was performed using 0.5 μg/ml rabbit anti-HA antibody (QED Bioscience Inc., San Diego) and IRDye 800LT-conjugated donkey anti-rabbit IgG and the LI-COR Odyssey infrared imaging system (LI-COR Biosciences). For the *in vitro* activity assay, TbGT15-ΔHA₃ was immunoprecipitated using anti-HA magnetic beads (Pierce) and incubated with 1 μCi of UDP-³²P-GlcNAc (specific activity of 20–40 Ci/mmol, PerkinElmer Life Sciences), 1 mM cold UDP-GlcNAc (Sigma), and 5 μg of Man₁-6(Man₁-3)Manβ₁-4GlcNAcβ₁-4GlcNAc or 25 μg of α₁-3,α₁-6-mannosiose (both Dextera Laboratories, Reading, UK) in 50 mM Tris, pH 7.5, 10 mM MgCl₂, 10 mM MnCl₂ in a total volume of 50 μl. After overnight incubation under vigorous shaking at room temperature, samples were desalted via a mixed-bed ion exchange column of 100 μl of Chelex-100 (Na⁺) over 100 μl of AG50X12 (H⁺) over 200 μl of AG3X4 (OH⁻) over 100 μl of QAE-Sephadex 25 (OH⁻), all from Bio-Rad, UK, except QAE-Sephadex (Sigma). Finally, glycans were freeze-dried and re-dissolved in 20% 1-propanol and aliquots were spotted onto silica HPTLC plates (SI-60 HPTLC, Millipore) that were run twice in 1-propanol/acetic acid/H₂O (9:6:4). For product analysis, samples were treated with 128 units of α₁-2.3 mannosidase from *Xanthomonas manihotis* (New England Biolabs) or 0.2 units of β-N-acetylglucosaminidase from *Canavalia ensiformis* (Sigma) before TLC analysis. Plates were then dried, sprayed with EN³HANCE autofluorography enhancer (EN³HANCE, PerkinElmer Life Sciences), and exposed on x-ray film at −80°C for 1–2 days.

For mass spectrometric analysis of the reaction product, the assay was performed using 5 mM non-radioactive UDP-GlcNAc. Samples were analyzed by LC-MS using a HILIC column (Tosoh TSKgel Amide column, 1 mm × 10 cm) and a gradient of 80 to 5% acetonitrile in 0.1% formic acid at a flow rate of 50 μl/min using a TSQ Quantapeptide-triple-quadrupole mass spectrometer (Thermo Fisher Scientific). For methylation linkage analysis of the product, glycans were converted to constituent monosaccharides in the form of partially methylated alditol acetates and analyzed by GC-MS as described above.

**Scanning Electron Microscopy**—To analyze bloodstream form cells by scanning electron microscopy, cells were fixed in HMI-9 medium with 2.5% glutaraldehyde. They were further processed and examined in a Philips XL30 ESEM operating at an accelerating voltage of 15 kV by the Centre for High Resolution Imaging and Processing (CHIPS) at the University of Dundee.

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**Results**

**Analysis of the TbGT15 Gene Product**—We previously characterized the biological function of three members of a family of putative UDP-sugar-dependent GTs (1, 26, 27). In this study, Tb927.7.300 was selected for functional analysis. The gene encodes for a 367-amino acid protein with a theoretical molecular mass of 43.1 kDa. Stable isotope labeling with amino acids in cell culture-based quantitative proteomic data demonstrated that the protein expression level is 15 times higher in bloodstream form parasites compared with procyclic form parasites (40).

The *T. brucei* strain that was used in this study (Lister strain 427) differs from the one that was used for the reference genome sequencing project (TREU927). However, an alignment of Tb927.7.300 and its homologue Tb427.7.300 revealed a very high similarity with only three single nucleotide polymorphisms, none of them resulting in amino acid changes. The strain 427 gene and protein product will be referred to here as *TbGT15* and *TbGT15*, respectively.

The protein sequence contains several hallmarks of Golgi apparatus glycosyltransferases. First, a membrane protein topology prediction program based on a hidden Markov model (41) designates TbGT15 as a type II transmembrane protein. In addition, the sequence contains a DXXD motif, which is generally involved in catalytic activity of known GTs (42) as well as a dibasic motif, which functions as an endoplasmic reticulum exit signal (43). Indeed, a subcellular Golgi localization of TbGT15 was confirmed previously (29, 30).

**Creation of Bloodstream Form TbGT15 Null and Conditional Null Mutants**—As TbGT15 is predominantly expressed in bloodstream form parasites (40), we decided to investigate the protein function by creating null and conditional null mutants in this life cycle stage. BLAST search of the *T. brucei* genome suggested that TbGT15 is present as a single copy per haploid genome. Both alleles were sequentially replaced by homologous recombination using PAC and HYG drug resistance cassettes as summarized in Fig. 1A. After selection on the respective antibiotics, the generation of a *TbGT15* null mutant (∆*TbGT15::PAC/ΔTbGT15::HYG) was confirmed by Southern blot using probes for the *TbGT15* ORF and 3′ UTR (Fig. 1B). To allow for a tetracycline-inducible re-expression of the gene, an ectopic copy of *TbGT15* was introduced into the rRNA locus of the null mutant (*TbGT15* null mutant (Fig. 2A)). No morphological differences between the WT and *TbGT15* null mutant parasites could be ascertained by light microscopy or by scanning electron microscopy (Fig. 2A). Compared with WT cells, the *TbGT15* null mutant parasites exhibited slightly slower growth kinetics in vitro, and this mild growth phenotype was partially reversed in *TbGT15* conditional null cells grown under permissive conditions (Fig. 2B). In addition, no difference in its ability to infect mice could be detected for the *TbGT15* null mutant (Fig. 2C). From this we can conclude that *TbGT15* is a non-essential gene in *T. brucei* bloodstream form cells.
Characterization of VSG from WT and TbGT15 Null Mutant Parasites—VSG221 from WT cells is heterogeneously glycosylated, containing a highly galactosylated GPI anchor (5), one oligomannose N-glycan at Asn-428 (Man$_5$GlcNAc$_2$), as well as small biantennary structures ranging from Man$_3$GlcNAc$_2$ to Gal$_2$GlcNAc$_2$Man$_3$GlcNAc$_2$ at Asn-296 (8, 16). VSG can be isolated in its sVSG form by hypotonic lysis, which results in its release by endogenous GPI-specific phospholipase C (44).

To assess differences in the glycosylation phenotype, intact sVSGs from WT and TbGT15 null mutant parasites were analyzed by ES-MS in positive-ion mode (Fig. 3). VSG molecules containing a total of four or five GlcNAc residues were present at similar levels in both genotypes, but glycoforms with six GlcNAc residues were completely absent in the TbGT15 null mutant (see arrows in Fig. 3B and Table 1). Bearing in mind that four GlcNAc residues are necessary for the composition of the two N-glycan N-acetylchitobiose core structures, the lack of VSG glycoforms containing six GlcNAc residues strongly indi-
cates a deficiency in biantennary complex N-glycans at Asn-296. From this we can conclude that the mutant cells are unable to express complex N-glycans and that TbGT15 is involved in their biosynthesis.

In Vitro Functional Activity Assay—To verify that TbGT15 encodes a glycosyltransferase that is directly involved in the biosynthesis of hybrid or complex N-glycans, we performed an in vitro assay for enzymatic activity as described previously (1). Briefly, full-length TbGT15 fused to a C-terminal 3× HA epitope tag was expressed in T. brucei bloodstream form cells and immunoprecipitated using anti-HA magnetic beads. The protein was incubated with Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc as acceptor substrate and tritium-labeled UDP-[3H]GlcNAc as donor substrate. Following desalting and removal of excess UDP-[3H]GlcNAc donor by mixed-bed ion exchange, aliquots were separated by thin layer chromatography (TLC) and analyzed by fluorography.

Although a control immunoprecipitation with lysate from WT cells did not result in any [3H]GlcNAc incorporation (Fig. 4A, lane 2), the sample containing TbGT15-HA3 showed a strong signal of tritium-labeled reaction product (Fig. 4A, lane 1). This demonstrates that TbGT15 is able to transfer GlcNAc to biantennary Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc core structures. It is noteworthy that the shortened substrate Manα1–6(Manα1–3)Man lacking the chitobiose core was not used as an UDP-GlcNAc acceptor by TbG15 (Fig. 4A, lane 3).

To reveal whether GlcNAc is transferred to the Manα1–3 or Manα1–6 arm of Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc, the reaction product was treated with α1–2,3-mannosidase. As seen in the subsequent TLC/autofluorography analysis, mannosidase treatment caused an increased mobility of the reaction product (compare Rf values in Fig. 4B). This suggests that the 3-Man arm was not modified by TbGT15, leaving it susceptible to exoglycosidase cleavage, and allows us to conclude that the transferred GlcNAc residue is attached to the 6-Man arm of the Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc core structure. The anomeric configuration of the newly formed linkage was determined by β-N-acetylgalactosaminidase digestion of the reaction product. In the following TLC/autofluorography analysis, the band of tritium-labeled GlcNAcManα1GlcNAc disappeared, although the amount of free [3H]GlcNAc increased, demonstrating a β-configuration (Fig. 4C).

For further characterization of the reaction product, the assay was performed using non-radioactive UDP-GlcNAc. First, HILIC-MS was performed to identify the HexNAc3Hex3 reaction product (Fig. 5, A and B). A subsequent methylation linkage analysis on the reaction sample by GC-MS demonstrated the presence of 1,2,5-tri-O-acetyl-(1-deutero)-3,4,6-tri-O-methyl-mannitol, originating from 2-O-substituted mannose, which reveals that TbGT15 transfers GlcNAc in a 1–2 linkage to one of the non-reducing terminal mannose residues of the trimannosyl core (Fig. 5, C and D).

Taken together, these data show that TbGT15 is the glycosyltransferase responsible for the transfer of β1–2-linked GlcNAc to the α1–6-linked α-D-mannopyranosyl residue of Manα1–6(Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc and can therefore be termed an N-acetylgalactosaminyltransferase type II or TbGnTII.

N-Glycosylation Phenotype of Bloodstream Form TbGT15 Mutant Parasites—To investigate the effect of TbGT15 on the glycosylation of other proteins than VSG, total glycoproteins were extracted with SDS/urea from VSG-depleted trypanosome ghosts and analyzed by lectin blotting. As reported previously for WT T. brucei, ricin (RCA-120), a lectin that predominantly binds to non-reducing terminal β-galactose residues, showed strong binding to a series of glycoproteins running between 60 and 150 kDa (Fig. 6, 1st lane). Ricin binding to glycoproteins extracted from the TbGT15 null mutants was slightly reduced, and the apparent molecular mass of all signals was marginally smaller compared with WT signals (Fig. 6, 2nd lane). Albeit subtle, these changes in blotting pattern suggest an alteration in the synthesis of the large poly-LacNAc-containing glycans of the high molecular weight invariant glycoproteins (19).

To draw structural conclusions, we decided to analyze the ricin-binding glycoprotein fraction by methylation linkage as described previously (19). Briefly, WT and TbGT15 null mutant cells were isolated, depleted of VSGs, solubilized in SDS/urea, and glycoproteins were purified by ricin affinity chromatography. N-Linked glycans were released by peptide:N-glycosidase F and further fractionated by Bio-Gel P-4 gel filtration, resulting in two main fractions as follows: one containing the relatively small mannose-rich N-glycans, and a Gal/GlcNAc-rich high molecular mass fraction eluting at the void volume of the Bio-Gel P-4 column (the total poly-LacNAc fraction). Aliquots of these fractions were subjected to neutral monosaccharide composition analysis by GC-MS. The molar ratios of Gal/Man
in the total poly-LacNac fraction for WT (19) and TbGT15 null mutant parasites were found to be similar, 14.5:1 and 12.4:1, respectively.

Subsequent GC-MS methylation linkage analysis of the total poly-LacNac fraction revealed structural similarities but also some quantitative differences (Table 2). Thus, both WT and TbGT15 null total poly-LacNac glycans contain 2-O-substituted Man and 3,6-di-O-substituted Man, consistent with a conventional core structure of R-2Manα1→6(R′-2Manα1→3)Manβ1→4GlcNacβ1→4GlcNAC in all structures. Furthermore, both samples contained significant amounts of 4-O-substituted GlcNac, indicating the presence of multiple LacNac
repeats. However, although the numbers of terminal Gal residues were similar, there was a significant decrease in 6-O-substituted Gal and 3-O-substituted Gal residues and a concomitant increase in 3,6-di-O-substituted-Gal residues in the \( \text{TbGT15} \) null mutant glycans. Because \( \text{TbGT15} \) has a GnTII-type activity that initiates elaboration of the Man/Hex1–6 arm of Man/Hex1–6(Man/Hex1–3)Man/Hex1–4GlcNAc/Hex1–4GlcNAc core, these data are consistent with a model where linear poly-LacNAc chains predominate on the Man/Hex1–6 arm, and the Man/Hex1–3 arm is occupied by both linear and branched poly-LacNAc units (Fig. 7). The increase in the number of 3,6-di-O-substituted Gal residues in the \( \text{TbGT15} \) mutant glycans further suggests that deletion of the entire (predominantly linear) poly-LacNAc chain attached to the Man/Hex1–6 arm is, to some extent, compensated by further elaboration of the (linear and branched) poly-LacNAc chain attached to the Man/Hex1–6 arm.

Discussion

We have cloned the gene that encodes UDP-GlcNAc:α1-6-D-mannoside-β1–2-N-acetylglucosaminyltransferase II in \( \text{T. brucei} \), an enzyme that catalyzes an essential step on the route to complex N-glycans. In human patients with carbohydrate-deficient glycoprotein syndrome type II, congenital deficiency in GnTII activity is associated with severe psychomotor retardation and other multisystemic abnormalities (45, 46). In a mouse model with a homozygous null mutation in the gene encoding GnTII (\( \text{Mgat2}^{−/−} \)), 99% of
newborns die during the first postnatal week (47). These developmental defects highlight the importance of complex N-glycans in intercellular communication and signaling in multicellular organisms.

The significance of complex N-glycans in the unicellular protozoan T. brucei is less well understood. In the bloodstream form, the parasite expresses both conventional biantennary complex N-glycans and unique highly extended and branched poly-N-acetyllactosamine-containing complex N-glycans (8, 19, 48, 49). However, the T. brucei genome contains no obvious homologues of the canonical GnTI and GnTII genes that encode the β1-2GlcNAc transferases usually responsible for the initiation of complex N-glycans. In a previous study, we identified and characterized TbGnTI, the enzyme responsible for the transfer β1-2GlcNAc to the Manα1–3 arm of N-glycan core structures (1). TbGnTI showed unusual activity in that it acts on biantennary Manα1–3GlcNAc2 instead of triantennary core structures (1). TbGnTI showed unusual activity in that it acts on biantennary Manα1–3GlcNAc2 instead of triantennary core structures (1). TbGnTI showed unusual activity in that it acts on biantennary Manα1–3GlcNAc2 instead of triantennary core structures (1).

In this study, a reverse-genetics approach in T. brucei bloodstream form cells indicated that the deletion of TbGT15 (another trypanosome β3-glycosyltransferase superfamily member) is accompanied by the absence of complex N-glycans, as well as alterations in the biosynthesis of the giant poly-Lac-NAc-containing glycans. Using a direct enzymatic assay and comprehensive product analysis, we could show that purified TbGT15 catalyzes the conversion of Manα1–6(Manα1–3) Manβ1–4GlcNAcβ1–4GlcNAc to GlcNAcβ1–2Manα1–6 (Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc. We have therefore renamed TbGT15 to TbGnTII. The conversion of Manα3GlcNAc2 demonstrates that TbGnTII works indepen-

![FIGURE 6. Lectin blotting of VSG-depleted glycoproteins. Lysates of washed WT or TbGT15 null mutant (KO) trypanosome cell ghosts were subjected to SDS-PAGE and transferred to nitrocellulose membrane. The membrane was incubated with ricin (left panel) or, as a binding specificity control, with ricin that was pre-incubating with 10 mg/ml each of galactose and lactose (middle panel). Equal loading and transfer are demonstrated by Ponceau S staining (right panel).](image)

![FIGURE 7. Proposed scheme for poly-LacNAc-containing N-glycans of bloodstream form trypanosomes. The data presented here and in Ref. 1 are consistent with the model shown here whereby in the wild-type bloodstream form of T. brucei the majority of large complex N-glycans carry highly branched poly-LacNAc chains on the Manα1–3 arm and predominantly linear poly-LacNAc chains on the Manα1–6 arm. In the TbGT15 null mutant, all elaboration of the Manα1–6 arm is absent and partly compensated by an increase in the branched poly-LacNAc elaboration of the Manα1–3 arm. Note that the relative positions of branch points shown are arbitrary.](image)

### TABLE 2

Quantitative GC-MS methylation linkage analysis of the total poly-LacNAc fraction

The total poly-LacNAc fraction was permethylated, hydrolyzed, reduced, and acetylated for GC-MS analysis. The resulting PMAA derivatives were identified by retention time and electron impact mass spectra. Quantification was accomplished by integration of the total ion current using molar relative response factors deduced empirically from authentic standards, as described under “Experimental Procedures.”

<table>
<thead>
<tr>
<th>PMAA derivative</th>
<th>Origin</th>
<th>WT*</th>
<th>TbGT15 null mutant*</th>
</tr>
</thead>
<tbody>
<tr>
<td>2,4-Di-O-methyl-1,3,5,6-tetra-O-acetyl-1-[3H]mannitol</td>
<td>3,6-Di-O-substituted Man</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>3,4,6-Tri-O-methyl-1,2,5-tri-O-acetyl-1-[3H]mannitol</td>
<td>2-O-Substituted Man</td>
<td>2.3</td>
<td>2.3</td>
</tr>
<tr>
<td>2,3,4,6-Tetra-O-methyl-1,5-di-O-acetyl-1-[3H]galactitol</td>
<td>Terminal Gal</td>
<td>4.9</td>
<td>5.0</td>
</tr>
<tr>
<td>2,4,6-Tri-O-methyl-1,3,5-tri-O-acetyl-1-[3H]galactitol</td>
<td>3-O-Substituted Gal</td>
<td>2.2</td>
<td>1.7 (down 23%)</td>
</tr>
<tr>
<td>2,3,4-Tri-O-methyl-1,5,6-tri-O-acetyl-1-[3H]galactitol</td>
<td>6-O-Substituted Gal</td>
<td>23.4</td>
<td>15.9 (down 32%)</td>
</tr>
<tr>
<td>2,4-Di-O-methyl-1,3,5,6-tetra-O-acetyl-1-[3H]galactitol</td>
<td>3,6-Di-O-substituted Gal</td>
<td>3.6</td>
<td>5.0 (up 39%)</td>
</tr>
<tr>
<td>3,6-Di-O-methyl-1,4,5-tri-O-acetyl-2-N-methylacetamido-1-[3H]glucosaminol</td>
<td>4-O-substituted GlcNAc</td>
<td>24.6⁶</td>
<td>16.6⁶</td>
</tr>
</tbody>
</table>

* Molar quantities relative to 3,6-di-O-substituted Man (one per glycan) are shown.

⁶ Values for N-acetylglucosamine derivatives are less reliable than for hexoses.
dently from prior TbGnTI action, which is in contrast to canonical GnTIIIs that only use substrates after modification by GnTI, i.e. Manα1–6(GlcNAcβ1–2Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc (51, 52). This unusual acceptor specificity of the trypanosome enzyme was already indicated by previous data, which show the presence of “pseudohybrid” N-glycans in the absence of TbGnTI (1) and highlights the divergent nature of TbGnTII. A phylogram based on a multiple sequence alignment of TbGnTII and GnTIIIs of other species is shown in Fig. 8. The GnTIIIs of multicellular organisms belong the CAZy GT family 13 (39) and share only 9% identity with the human and mouse enzymes. In contrast, TbGnTII is a member of the CAZy GT31 family (26) and shares only 14% identity with the human sequence at the amino acid level. Interestingly, although TbGnTI is a member of the GnTII family, it is only weakly related to all closely related GnTIIIs, with nearly 90% identity between the human and mouse enzymes. Furthermore, the inability of TbGnTI to transfer 4GlcNAc (51, 52). This unusual acceptor specificity of the trypanosome enzymes, both of which work on the same acceptor substrate (Manα3GlcNAcα, although only the latter requires the N-acetylchitobiose core), whereas the canonical GnTII and GnTIII enzymes work on triantennary Manα2GlcNAcβ and Manα1–6(GlcNAcβ1–2Manα1–3)Manβ1–4GlcNAcβ1–4GlcNAc, respectively.

Methylation linkage analysis of the poly-LacNAc N-glycans of the TbGT15 null mutant showed a reduction in 6-O-substituted Gal and 3-O-substituted Gal but an increase in 3,6-O-substituted Gal. This allows us to augment our model of the parasites’ complex N-glycans and propose that the Manα1–6 arm is normally occupied by predominantly linear poly-LacNAc repeats and the Manα1–3 arm by branched as well as linear poly-LacNAc repeats (Fig. 7).

T. brucei has an unusual dual N-glycosylation mechanism with two paralogous oligosaccharyltransferases, TbSTT3A and TbSTT3B, that transfer biantennary Manα3GlcNAcα and triantennary Manα2GlcNAcβ, respectively, in a site-specific manner (15, 16). Because of the absence of Golgi α-mannosidase II in the parasite, triantennary structures cannot be processed to complex N-glycans, rendering biantennary Manα3GlcNAc transferred by STT3A the only route to paucimannose and complex structures. Furthermore, the inability of TbGnTI to act on triantennary Manα2GlcNAcβ (1) also means that biantennary Manα3GlcNAc transferred by STT3A is the only possible route to pseudohybrid N-glycans (i.e., those with only one arm of the trimannosyl-core modified by GlcNAc) as well as additional sug-

ars. RNAi knockdown of TbSTT3A showed that cells are viable in culture but not in mice (15). Interestingly, the deletion of TbGnTI (TbGT11) has no effect on in vitro growth rate, and the infectivity to mice was indistinguishable from wild type (1). This suggests that the presence of pseudohybrid N-glycans with glycan extensions to the 6-arm alone are sufficient to compensate for the loss of complex N-glycans. Here, the in vitro and in vivo viability of the TbGnTII null mutant shows that the reverse is true, in that the presence of hybrid structures with extensions to the 3-arm alone compensates for the loss of complex N-glycans. However, despite extensive attempts, a double knock-out lacking both TbGnTI and TbGnTII genes could not be generated in our hands, suggesting that extension of one or other of the arms of the N-glycan trimannosyl-core is essential for the growth and infectivity of bloodstream form of T. brucei.

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References
A Gene of the β3-Glycosyltransferase Family Encodes N-Acetylglucosaminyltransferase II Function in Trypanosoma brucei
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