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Deletion of GPI7, a Yeast Gene Required for Addition of a Side Chain to the Glycosylphosphatidylinositol (GPI) Core Structure, Affects GPI Protein Transport, Remodeling, and Cell Wall Integrity*

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Gpi7 was isolated by screening for mutants defective in the surface expression of glycosylphosphatidylinositol (GPI) proteins. Gpi7 mutants are deficient in YJL062w, herein named GPI7. GPI7 is not essential, but its deletion renders cells hypersensitive to Calcofluor White, indicating cell wall fragility. Several aspects of its deletion renders cells hypersensitive to Calcofluor White, indicating cell wall fragility. Several aspects of GPI biosynthesis are disturbed in Δgpi7. The extent of anchor remodeling, i.e. replacement of the primary lipid moiety of GPI anchors by ceramide, is significantly reduced, and the transport of GPI proteins to the Golgi is delayed. Gpi7p is a highly glycosylated integral membrane protein with 9–11 predicted transmembrane domains and a large, hydrophilic C-terminal part. Gpi7p contains significant homology with phosphodiesterases suggesting that Gpi7p itself is the transferase conferring on GPI-PLD, GPI-specific phospholipase D; Ins, inositol; Man, mannose; ORF, open reading frame; pC1 and pC2, protein-derived Ceramides 1 and 2; pG1 protein-derived Glycophospholipid 1; PI, phosphatidylinositol; PM, plasma membrane; ts, thermosensitive; wt, wild type; PAGE, polyacrylamide gel electrophoresis; PIPES, 1,4-piperazinediethanesulfonic acid; nt, nucleotide; PCR, polymerase chain reaction; kb, kilobase pair(s); HPLC, high pressure liquid chromatography.

Glycosylphosphatidylinositol (GPI)-anchored proteins represent a subclass of surface proteins found in virtually all eukaryotic organisms (1). The genome of Saccharomyces cerevisiae contains more than 70 open reading frames (ORFs) encoding for proteins that, as judged from the deduced primary sequence, can be predicted to be modified by the attachment of a GPI anchor (2, 3). In about 25 percent of them, the presence of an anchor has been confirmed biochemically. A majority of them lose part of the anchor and become covalently attached to the β1,6-glucans of the cell wall (4–6). A minority of GPI proteins retain the GPI anchor in an intact form and stay at the plasma membrane (PM).

For the biosynthesis of GPI anchors, phosphatidylinositol (PI) is modified by the stepwise addition of sugars and ethanolamine phosphate (EtN-P), thus forming a complete precursor lipid (CP) which subsequently is transferred en bloc to a transamidase onto newly synthesized proteins in the ER (7, 8). The identification of genes involved in the biosynthesis of the CP and its subsequent attachment to proteins has been possible through the complementation of mammalian and yeast gpi- mutants, i.e. mutants being deficient in GPI anchoring of membrane proteins (7, 9–20). In our laboratory, a series of recessive gpi- mutants (gpi4 to gpi10) has been obtained by screening for yeast mutants that are unable to display the GPI-anchored a-agglutinin (Sag1p) at the outer surface of the cell wall, although the synthesis and secretion of soluble proteins is normal (21, 22).

Here we report on the characterization of gpi7. Four independent gpi7 mutants accumulated M4, an abnormal GPI intermediate that is less hydrophilic than CP2, the precursor accumulating when the transfer of GPIs to proteins is interrupted (18, 19, 21, 23). Our preliminary characterization of M4 had shown that deacylation by NH3 followed by HF treatment, used to hydrolyze selectively the phosphodiester bonds (Fig. 1), yielded the same Man$_2$GlcN-inositol fragment as CP2, and we speculated that gpi7 mutants may be unable to add the EtN-P onto Man3 (Fig. 1) (21). Here we show that this speculation was wrong, that CP2 differs from M4 with regard to a previously unrecognized side chain attached to Man2 (Fig. 1), and that GPI7 is required for the attachment of this side chain.

**EXPERIMENTAL PROCEDURES**

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The abbreviations used are: GPI, glycosylphosphatidylinositol; ASAM, A. satoi α-mannosidase; CP, complete precursor; DAG, diacetyl-glycerol; DHS, dihydrosphingosine; ETN-P, ethanolamine phosphate; GPI-PLD, GPI-specific phospholipase D; Ins, inositol; JBM, jack bean α-mannosidase; Man, mannose; ORF, open reading frame; pC1 and pC2, protein-derived Ceramides 1 and 2; pG1 protein-derived Glycophospholipid 1; PI, phosphatidylinositol; PM, plasma membrane; ts, thermosensitive; wt, wild type; PAGE, polyacrylamide gel electrophoresis; PIPES, 1,4-piperazinediethanesulfonic acid; nt, nucleotide; PCR, polymerase chain reaction; kb, kilobase pair(s); HPLC, high pressure liquid chromatography.
FIG. 1. Presumed structure of the complete yeast precursor glycolipid GCP3. Relevant cleavage procedures are indicated. Man1, Man2, Man3, and Man4 designate the α1,4-linked, α1,6-linked, and α1,2-linked mannooses (Man). X indicates an HF-sensitive group that is not yet defined chemically. Alkyl, P, phosphate.

X2150-1A (MATaΔlys), FBY122 (MATa ade2-1 ura3-1 leu2-3,112 trpl-1 his3-11,15 gpi7-1) FBY182 (MATa ade2-1 ade2-1 ura3-1 leu2-3,112 his3-11,15 gpi7-7:KanMX4), HMSF176 (MATa sec18-1), FBY49 (MATa sec18-1 gpi7-KanMX4), C4 (MATa ura3-52 leu2-3,112 pir40), HMSF331 (MATa sec33-6), LB1234-3B (MATa mnn9), and YNS7-Aa (MATa ura3 his3-a11 och1::LEU2). Diploid strains were FBY118 (MATa a ade2-1 ade2-1 ura3-1 leu2-3,112, leu2-3,112 trpl-1 trpl-1 his3-11,15 his3-11,15 lys3-15 lys3-15), FBY40 (MATa a ade2-1 ade2-1 ura3-1 ura3-1 leu2-3,112/leu2-3,112 trpl-1 trpl-1 his3-11,15 his3-11,15 lys3-15 lys3-15 gpi7:GPI7-KanMX4), and FBY43 (MATa a ade2-1 ade2-1 ura3-1 ura3-1 leu2-3,112/leu2-3,112 trpl-1 trpl-1 his3-11,15 his3-11,15 lys3-15 lys3-15 gpi7:GPI7-KanMX4). 

Materials were obtained from the sources described recently (22). Cysteamine was from Sigma; [3H]dihydrosphingosine was synthesized as described (24); myriocin was a kind gift of Dr. N. Rao Movva (Novartis, Basel, Switzerland); antibodies to Och1p, alkaline phosphatase, Kex2p, and Wbp1p were kindly donated by Dr. Y. Jigami, National Institute of Bioscience and Human Technology, Ibaraki 305, Japan; Dr. S. Emr, Howard Hughes Medical Institute, University of California, San Diego; Dr. R. Fuller, University of Michigan Medical Center, Ann Arbor, MI; and Dr. M. Aebi, Mikrobiologisches Institut, ETH Zürich, Switzerland, respectively.

Cloning, Partial Sequencing, and Disruption of Gpi7p—The Gpi7p gene was cloned by complementation of the growth phenotype of the gpi7-1/gpi7-1 double mutant as described (19). The three plasmids complementing gpi7-1 contained a 4.3-kb common DNA restriction fragment that was partially sequenced by theideo sequencing method (25). The complementing insert Sppl/Ssp1 of 3.6 kb was cloned into the Sppl/Ssp1-digested Yep352 multicopy vector (26) or Yep243 single copy vector (27) to generate pBF41 (Fig. 3C) and pBF43, respectively.

One step disruption of Gpi7p was done as described (28). Briefly, the 1.5-kb long KanMX4 module was PCR-amplified by using pFA6a-KanMX4 as template and the following two adapter primers: GPI7-forwards (5′-CTTCCACAGGATGGAAAGACCTGACCGTCCTGCGTATGATCTCGTTC-3′) with 17 nucleotides (nt) of homology to the pFA6a-KanMX4 multiple cloning site (in lowercase) and 43 nt of homology to GPI7 (in uppercase) starting 18 nt upstream of the start codon (bold); GPI7-backwards (5′-ATCGAGACCCAGGAAAGGGGCA-AATTGGGATCCAGCCATTGAAGCTGAGCTCG-3′) with 18 nt of homology to the pFA6a-KanMX4 multiple cloning site (lowercase) and 44 nt of homology to the ORF of Gpi7p in the region immediately upstream of the stop codon. This PCR DNA fragment was used to transform the diploid strain FBY118, homozygous for Gpi7p, and FBY40, a heterozygous gpi7-1/GPI7 strain. The correct targeting of the PCR-made KanMX4 module into the Gpi7p locus in genetic-resistant clones was verified by PCR with whole yeast cells using primers GPI7-plus (5′-GTTTACATTCAACCGCGCACG-3′) starting 36 nt upstream of the ATG, GPI7-minus (5′-GACCGAAGTAATGGACCGGC-3′) starting 631 bp downstream of the stop codon, and the K2 primer of the KanMX4 module (5′-GGATTGATGTTGACG-3′).

Expression of Recombinant His-tagged Gpi7p and Antibody Production—Plasmid pBF41 (Fig. 3C) was digested with BstYI and EcoRV to generate a 633-base pair fragment of Gpi7p. This fragment was inserted into the multiple cloning site of the bacterial expression vector pQE-30 (Qiagen) digested with BsnHI/Smal thus generating the plasmid pBF402. This plasmid was used to transform the E. coli strain M15[pREP4]. Expression of the recombinant protein was induced with isopropyl-l-thio-β-D-galactoside and purified on a nickel-nitrotriacetic acid-agarose column (Qiagen) under denaturing conditions according to the manufacturer's instructions. A polyclonal antiserum was raised against this Gpi7p fragment by repeated intramuscular injections of 0.10 mg of recombinant protein. Ten mg of the recombinant protein were coupled to CNBr-activated Sepharose 4B (Amersham Pharmacia Biotech) according to the manufacturer's instructions, and antiserum was affinity purified as described (29).

Membrane Association, Protease Sensitivity, and Cellular Localization of Gpi7p—The nature of the association of Gpi7p with the membranes was studied by flotation.As a control, 1,000 µg of recombinant protein was solubilized. Ten mg of the recombinant protein were coupled to CNBr-activated Sepharose 4B (Amersham Pharmacia Biotech) according to the manufacturer's instructions, and antiserum was affinity purified as described (29). Briefly, 100 A260 of washed W303 cells were resuspended in 1 ml of lysis buffer (20 mM HEPES, pH 7.5, 500 mM sucrose, 3 mM magnesium acetate, 20 mM EDTA, 1 mM dithiothreitol) and were lysed by agitation with glass beads in 4 °C. The homogenate was centrifuged for 5 min at 600 × g to remove unbroken cells, and the supernatant was centrifuged for 20 min at 13,000 × g. The membrane pellet was resuspended in 240 µl of the same lysis buffer and split into 6 aliquots of equal size. Aliquots of microsome were incubated for 20 min on ice with or without 0.5% Triton X-100 and proteinase K. Digestions were stopped by addition of gelatin. The lipids were extracted by two-dimensional radioscanning (LB 2842; Berthold AG, Regensdorf, Switzerland). TLC plates were sprayed with EN3HANCE and exposed to X-ray film for 24 h at −70 °C to identify the lipids.

Labeling of Cells—Previously described procedures were used to label cells with [3H]dihydrosphingosine (21) or [3H]inositol (21). Lipids were extracted as described (24). Lipid extracts were analyzed by ascending TLC using 0.2-mm thick silica gel 60 plates with the solvent chloroform/methanol/water, 55:45:10, v/v as solvent 1 (chloroform/methanol, 0.25% KCl) for GPI-PLD treatment and solvent 2 (chloroform/methanol, 0.25% KCl) for GPI-AK. Lipids were treated with NaOH (34) before TLC analysis.

Western blotting—All the samples were denatured during 5 min at 95 °C in reducing sample buffer and run on a 6, 7.5, or 10% SDS-PAGE for detection of antibodies (32). Western blotting was carried out with anti-Wbp1p, anti-alkaline phosphatase, anti-Gas1p, or anti-Kex2p antiserum or with affinity purified anti-Gpi7p or anti-Och1p antibodies, always using the chemiluminescence ECL kit from Amersham Pharmacia Biotech, Buckinghamshire, UK.

Labeling of Cells—Previously described procedures were used to label cells with [3H]dihydrosphingosine (21) or [3H]inositol (21). Lipids were extracted as described (24). Lipid extracts were analyzed by ascending TLC using 0.2-mm thick silica gel 60 plates with the solvent chloroform/methanol/water, 55:45:10, v/v or solvent 2 (chloroform/methanol/water, 10: 10:3, v/v). Radioactivity was detected and quantitated by one- and two-dimensional radioscanning (LB 2842; Berthold AG, Regensdorf, Switzerland). TLC plates were sprayed with EN3HANCE and exposed to film (X-Omat; Eastman Kodak Co.) at −80 °C.

Analytical Methods—Lipid extracts were deacylated with NaOH (34) and treated with JAB (35) as described (38). For GPI-PLD treatment lipid extracts were dissolved in 20 mM Tris-HCl, pH 7.4, 0.1 mM CaCl2, 20% 1-propanol. Incubations were for 12 h at 37 °C. All treated lipid extracts were desalted by partitioning between n-butyl alcohol and an aqueous solution of 0.10 mM EDTA, 5.0 mM Tris-HCl, pH 7.5, and back extraction of the butanol phase with water before TLC (23). Lipids were treated with methanolic NH4 to remove the acyl group on Ins (36) and cleaved using nitrous acid (37) as described. Lipids were purified by preparative TLC on 0.2-mm thick Silica Gel 60 plates (Merck, Germany) in solvent 2. Radioactive spots were localized by radioscanning, scraped, and eluted in solvent 2. Thin-layer chromatography (TLC) was done to confirm the identity of the compounds by spraying with ninhydrin.
done as described (39). The generated fragments were analyzed by paper chromatography in methylethyl ketone/pyridine/H\textsubscript{2}O (20:12:11) as described (39). Before paper chromatography the products were N-acetylated and desalted over mixed-bed ion exchange resin AG-501-X8 (Bio-Rad) unless indicated otherwise (34). Acetolysis was done as described (39). Before paper chromatography the products were into view the faint bands of upper part of the fluorogram was scanned at higher sensitivity to bring contain residual amounts of free \([3H]\)Ins after extraction into butanol. The band migrating between CP2 and M4 in \(gpi8-1\) is a GPI-PLD-, mild base-, and JBAM-sensitive GPI intermediate containing the Man\(_4\)-GlcN-Ins core,\(^2\) but the structural differences between this species and CP2 or M4 have not been identified.

RESULTS

Cloning of GPI7—As reported before (19) and shown in Fig. 2, wild type (wt) cells do not contain polar GPIs (lane 1), \(gpi8-1\) accumulates CP2 as the most polar GPI lipid (lane 8), and \(gpi7-1\) and the \(gpi7-1/gpi8-1\) double mutant accumulate M4 (lanes 4 and 6), thus demonstrating that \(gpi7-1\) is epistatic to \(gpi8-1\) and suggesting that, during GPI biosynthesis, Gpi7p and Gpi8p function epistatically (Fig. 2, lanes 8 and 7). As expected, the accumulation of CP2 by \(gpi8-1\) was not abolished by the overexpression of \(gpi7-1\) (Fig. 2, lane 9). \(gpi7-1\) and the \(gpi7-1/gpi8-1\) double mutant were unable to sporulate (Fig. 2, lanes 6 and 8). All these clones harbored plasmids containing \(YJL062w\) as the only complete ORF. Transfection of a multicopy vector containing \(YJL062w\) under its own promoter (pBF41, Fig. 3C) into \(gpi7-1\) almost completely cured the accumulation of M4 (Fig. 2, lane 8). As expected, the accumulation of CP2 by \(gpi8-1\) was not abolished by the overexpression of \(YJL062w\) (Fig. 2, lanes 8 and 9). \(YJL062w\) predicts an 830-amino acid membrane protein with an N-terminal signal sequence for insertion into the ER, 5 potential N-glycosylation sites, \(^{3}\) and about 9–11 putative transmembrane domains (Fig. 3, A and B). \(YJL062w\) was deleted and replaced by the selectable marker KanMX4. On rich medium the deletants grew about as rapidly as wt cells at all temperatures. Thus, \(YJL062w\) is not an essential gene. We were unable to sporulate \(\Delta YJL062/\Delta YJL062\) diploids indicating that \(YJL062\) is required for sporulation. However, \(\Delta YJL062/\Delta YJL062\) heterozygotes sporulated readily and \(\Delta YJL062\) spores germinate normally. In accordance with previous results on \(gpi7\) mutants (21), growth of \(\Delta YJL062\) (\(=\Delta gpi7\), see below) on plates at 37 °C was severely inhibited by 0.5 mg/ml Calcofluor White. \(\Delta YJL062\) accumulated M4 at even higher levels than \(gpi7-1\), and this accumulation was almost completely suppressed by the transfection of pBF41 (Fig. 2, lanes 2 and 3). Residual accumulation of M4 may be due to

\(^2\)I. Flury, unpublished observations.
some cells that lost the complementing plasmid. Transfection of YJL062w under its own promoter on a single copy vector (plasmid pBF43) was sufficient to suppress the accumulation of M4 in a homozygous ∆YJL062/∆YJL062 diploid (Fig. 2, lanes 11 and 12). As can be seen in Fig. 2, gpi7-1, ∆YJL062, ∆YJL062/∆YJL062, and gpi8-1 mutants also show minor amounts of the GlcNα1,6(acyl→)Ins-P-DAG GPI intermediate M0, the accumulation of which is believed to reflect a build up of GPI intermediates throughout the biosynthetic pathway (Fig. 2, lanes 2, 4, 6, 8, and 11). (It should be noted that some intermediates of intermediate size are obscured on TLC by PI and inositol phosphoceramide (41).) As expected, expression of YJL062w abolishes the accumulation of M0 in gpi7-1 and ∆YJL062 (Fig. 2, lanes 3, 5, and 12) but not in gpi7-1/gpi8-1 nor gpi8-1 (lanes 7 and 9), since in the latter the GPI biosynthesis remains blocked. To evaluate if the mutation in gpi7-1 is genetically linked to YJL062w, YJL062w was disrupted in a heterozygous gpi7-1/GPI7 diploid. Correct replacement of one YJL062w locus was verified by PCR in two independent genetic-resistant transformants. The verified deletants were sporulated, and a total of 26 complete tetrads was labeled with [3H]Ins to analyze the accumulation of M4. In all 26 tetrads only two of the four segregants showed accumulation of M4, whereas the other two showed the lipid profile of wt cells. Geneticin resistance also segregated 2:2 and cosegregated with M4 accumulation in all cases. This demonstrates that the mutation of gpi7-1 is tightly linked to YJL062w which we henceforth call GPI7. Since a construct containing only 348 nucleotides 5’ of the initiation codon of GPI7 still retained significant complementing activity, we also can dismiss the possibility that the complementing activity of pBF41 is due to one of the two small ORFs located on the opposite strand in the 5’ upstream region of GPI7 and starting at −409 and −503 with regard to the start codon of GPI7.

Characterization of the GPI Intermediate M4—We found that M4, contrary to our initial expectation, contained an HF-sensitive group on Man3 (Fig. 1). Indeed, treatment of the lipid extracts of gpi7-1 with jack bean α-mannosidase (JBAM, an exomannosidase) shifted M4 to a slightly less hydrophilic position on TLC (Fig. 4A, lanes 1 and 2) but not to the position of M0. It seemed conceivable that JBAM did not remove more than one Man from M4 because it was sterically hindered by the detergent micelle in which M4 was embedded. To circumvent this problem, M4 was purified by preparative TLC, and its hydrophilic head group was liberated by GPI-PLD, O-deacylated by NH3, and then subjected to several treatments as indicated at the top of Fig. 5, A–D. The N-acetylated fragment comigrated with the Manα1-GlcNAc-Ins standard (Fig. 5A). When treated with JBAM before HF, the resulting N-acetylated fragment comigrated with the Manα1-GlcNAc-Ins standard, clearly indicating the presence of a blocking group on Man3 (Fig. 5B). The blocking group on Man3 was HF-sensitive, since JBAM done after HF produced a fragment comigrating with GlcNAc-Ins (Fig. 5C). Aspergillus satoi α-mannosidase (ASAM), a linkage-specific α1,2-exomannosidase, when used after HF treatment, produced Manα2-GlcNAc-Ins (Fig. 5D). The migration of the fragments shown in Fig. 5, A and B, was much slower when N-acetylation was omitted (not shown). This partial characterization of M4 is consistent with the presence of a classical Manα1,2/EtN-P→Manα1,2Manα1,6Manα1,4-GlcNα1,6Ins core structure.

Having recently discovered an additional EtN-P on Man1 of CP2 (22), we considered the possibility that M4 may be lacking EtN-P on Man1. We thus proceeded to compare the non-dephosphorylated head groups of M4 and CP2 by Dionex HPLC using a system in which the presence of negatively charged phosphodiester greatly retards the elution of oligosaccharides (42). The non-dephosphorylated head groups of M4 and CP2 eluted as sharp peaks at fractions 22 and 31, respectively (not shown). This wide separation suggested that the head group of M4 contains less negative charge than the one of CP2. To assay directly for a side chain on Man1 of M4, the head group of [3H]Ins-labeled W303 in the experiment described in Table I. M4 and anchor peptides were treated with HNO3 to liberate the acyl-1'HIns-P-lipid and [3H]Ins-P-lipid moieties, respectively (Fig. 1). Samples were incubated with (+) or without (−) methanolic NH3 to remove the acyl from the Ins, desalted and separated by TLC (solvent 1), and processed for fluorography. pG1, protein-derived glycosphospholipid 1; pC1 and pC2, protein-derived ceramides 1 and 2, see Sipos et al. (41). Other results of this same experiment were described before (Ref. 41, therein Fig. 2).
Fig. 5. Analysis of the head group of M4 using HF, JBAM, and ASAM. Δgpi7 was labeled with $^{[3]H}$Ins at 37°C; M4 was purified and used to prepare head groups. Head groups were subjected to HF and N-acetylation (A); JBAM, then HF, then N-acetylation (B); HF, then N-acetylation, then desalting, then JBAM (C); HF, then N-acetylation, then desalting, then ASAM (D). The thus generated fragments were separated by paper chromatography, and radioactivity contained in 1-cm wide strips was determined through scintillation counting. The position of standards run in parallel on the same paper are indicated: 2–4, Man$_1$-GlcN-Ins with $x = 2, 3, or 4, 0$, GlcN-Ins.

Fig. 6. M4 of Δgpi7 contains an HF-sensitive substituent on Man1 but lacks an HF-sensitive substituent on Man2. CP2 and M4 head groups were obtained from $^{[3]H}$Man labeled pmi40 and $^{[3]H}$Ins labeled Δgpi7, respectively. A and B, head groups of M4 were subjected to acetolysis and then either treated with JBAM (B) or left untreated (A). Finally all products were dephosphorylated with HF, N-acetylated, and analyzed by paper chromatography. C and D, head groups were treated for 12 h with HF, desalted, treated with JBAM, treated with HF for 60 h, N-acetylated, and finally analyzed by paper chromatography. Standards 0–4 are Man$_x$-GlcN-Ins (x = 0, 1, 2, 3, 4). Free $^{[3]H}$Man ran out of the paper shown in C.

HPLC (data presented above and in Refs. 22 and 23). We thus hypothesized that CP2 may contain either additional HF-sensitive groups on GlcN, Man1, Man2, or Man3 or may contain additional groups linked through the amino group of the EtN-P on M1. Of these several theoretical possibilities, the only ones that have been documented in other organisms are the Man1-P group on the GlcN in Paramecium aurelia (43) and the EtN-P group on Man2 in several mammalian GPI proteins, e.g., human erythrocyte acetylcholinesterase (44), CD52-II (45), and bovine liver 5′-nucleotidase (46). We used limiting HF treatment to test specifically if CP2 contains an HF-sensitive group on Man2. If we assume that during HF treatment the EtN-Ps are hydrolyzed in a random order, we may expect to find some reaction intermediates lacking the HF-sensitive group on Man3 while retaining EtN-P on Man2 or Man1. When such intermediates subsequently are treated with JBAM and then are dephosphorylated to completeness with HF, they should yield Man$_2$-GlcN-Ins and Man$_1$-GlcN-Ins fragments, respectively. For preliminary tests, CP2 head groups were first treated with HF for 0.5, 1, 3, 9, 12, 18, or 28 h, then with JBAM, and finally with HF for 60 h. These experiments showed that both Man$_2$-GlcN-Ins and Man$_1$-GlcN-Ins fragments became visible after 1 h of limiting HF treatment, peaked at 12 h, and remained detectable at all time points up to 28 h. In quantitative terms it appeared that Man$_2$-GlcN-Ins > Man$_2$-GlcN-Ins $\geq$ Man$_1$-GlcN-Ins at all time points. Importantly, treatment of the head group of CP2 with HF for 12 h yielded substantial amounts of Man$_2$-GlcN-Ins and Man$_1$-GlcN-Ins (Fig. 6C), whereas the identical treatment performed with the head group of M4 only yielded Man$_1$-GlcN-Ins but no Man$_2$-GlcN-Ins (Fig. 6D). This result is compatible with the idea that Δgpi7 cells are unable to add an HF-sensitive group onto Man2 of the GPI core (Fig. 1). It also confirms the presence of an HF-sensitive group on Man1 of both M4 and CP2.

The Lipid Moieties of GPI Intermediates in Δgpi7 Are Normal.—We looked for additional differences between M4 and CP2 by analyzing the lipid moiety of M4. M4 is sensitive to GPI-specific phospholipase D (GPI-PLD) and mild base treatment (Fig. 4A, lanes 3–6), suggesting that its lipid moiety consists of Ins-P-DAG. We previously reported that M4 is resistant to PI-specific phospholipase C (21). This finding, together with the GPI-PLD sensitivity, can be taken as an indication for the presence of an acyl moiety attached to the Ins of M4. We further released the (acyl)-Ins-P-DAG moiety of M4 with HNO$_2$, as described recently (41). As shown in Fig. 4B, the treatment of purified M4 by HNO$_2$ produced a very hydrophobic species, which migrates very closely to M0, i.e., the GlcN(acyl)-Ins-P-DAG accumulating in sec53 (Fig. 4B, lanes 7 and 9) (41). (As reported previously, the presence of GlcN on these early precursors does not significantly influence their migration in TLC, for discussion see Sipos et al. (41).) Partial deacylation of the M4-derived lipid moiety by NH$_3$ produced PI and lyso-PI (Fig. 4B, lane 10). This PI was compared with pG1, the PI species obtained by HNO$_2$ treatment of protein-bound GPI anchors from the corresponding wt strain (Fig. 4B, lane 11). The comparison shows that M4 contains a PI moiety that migrates clearly less than pG1, whereas a lyso-PI of M4 migrates slightly more than the lyso-PI species generated by methanolic NH$_3$ treatment of anchor peptides (Fig. 4B, lanes 10 and 12). Very similar results had been obtained previously when comparing protein-derived PI moieties with the PI moieties of M0 from sec53 and of CP2 from gpi8-1 (41). In addition we isolated from Δgpi7 the recently identified GPI intermediates that are obscured in TLC by PI and inositol phosphoceramides (Ref. 41, therein Fig. 6A), and we found that they are exactly the same as the corresponding intermediates from wt cells by all criteria (not shown). Thus, it seems that M4 and other GPI intermediates of Δgpi7 contain the same PI moiety as early and late GPI intermediates accumulating in other mutants or in wt cells, and we therefore conclude that the difference between CP2 and M4 is solely due the difference in their head groups.

Lack of Gpi7p Affects the in Vitro Biosynthesis of GPI Precursor Lipids.—When yeast microsomes are incubated in the presence of UDP-$^{[3]H}$GlCNac, ATP, coenzyme A, GDP-Man, and tunicamycin, they generate labeled GPI intermediates as the only kind of labeled lipids (22, 47). Wild type microsomes make GPI intermediates up to CP2. Although a large array of incomplete intermediates is also generated, the pattern of labeled intermediates is fairly reproducible. When we used Δgpi7
FIG. 7. Microsomes of Δgpi7 synthesize M4 and not CP2. Microsomes of W303 wild type or Δgpi7 were incubated with 6 μCi of UDP-[3H]GlcNAc, GDP-Man, tunicamycin, and ATP for 1 h at 37 °C as described (22). The glycolipid products were extracted and then run on TLC with solvent 2. The extract in lane 3 was first treated with JBAM. On the basis of the preceding analysis the band denoted with an asterisk can be presumed to be an M4 derivative in which Man4 has been removed.

microsomes, they reproducibly made all the normal intermediates down to a band comigrating with M4 of [3H]Ins-labeled microsomes, they reproducibly made all the normal intermediates, much in the same way as seen for [3H]Ins-labeled M4 glycans is not necessary for Gpi7p function since, when shifted be reproduced (Fig. 2, lanes 1 and 2). Thus, the M4 accumulation of Δgpi7 can be reproduced in vitro. This result implies that the Gpi7p present in wt microsomes is functional in vitro.

Characterization of Gpi7p—Gpi7p was characterized using affinity purified rabbit antibody made against the N-terminal, hydrophilic part of GPI7 (Fig. 3B). As shown in Fig. 8A, the antibody recognized a heterogeneously glycosylated 208-kDa protein, the estimated molecular mass of various glycoforms ranging, after heavy exposure, from about 130 to 230 kDa (Fig. 8A, lane 2). The predicted mass of the protein before and after removal of the signal sequence is 94,832 and 92,207 Da, respectively (Fig. 3B). When the labeled lipid extract was treated with JBAM, most of the band comigrating with M4 was shifted to a less hydrophilic position, much in the same way as seen for [3H]Ins-labeled M4 (Fig. 2, lanes 1 and 2). This, the M4 accumulation of Δgpi7 can be reproduced in vitro. This result implies that the Gpi7p present in wt microsomes is functional in vitro.

FIG. 8. Membrane association, orientation, and localization of Gpi7p. A, exponentially growing cells were broken with glass beads in TEP buffer as described (19); lysates were centrifuged at 10,000 × g for 15 min at 4 °C, and microsomal pellets were processed for SDS-PAGE. Cells in lanes 3 and 5 had been grown in 20 μg/ml tunicamycin (Tm) for 90 min. B–D, exponentially growing W303 cells were broken with glass beads using the buffers indicated under “Experimental Procedures,” and cell wall debris was removed by centrifugation at 600 × g. B, aliquots of cell lysate were incubated for 30 min at 0 °C with 0.5 mM NaCl, 0.8 M urea, 1% Triton X-100 (TX-100), 0.1 M Na2CO3, pH 11, or 1% SDS. Subsequently membranes were sedimented by ultracentrifugation to get supernatant (S) and pellet (P) fractions. C, cell lysate was sedimented at 13,000 × g for 15 min, and the membrane pellet was thoroughly resuspended and digested with 10 or 25 μg/ml proteinase K (prot K) at 0 °C for 20 min in the presence or absence of 0.5% Triton X-100. D, cell lysates were subjected to differential centrifugations at 8,000 and 100,000 × g. These centrifugations generated pellet P8 containing ER, PM, and vacuolar membranes and pellet P100 which contains Golgi membranes. The 8,000 × g supernatant was also precipitated with trichloroacetic acid (TCA). In all panels the lanes contain material derived from 1 A100 of cells except for lanes 1 and 7 of A which contain 0.3 A100.
Cells were labeled with \([^{3}H]_{\text{Ins}}\) for 75 min at 37 °C. Incorporation of \([^{3}H]_{\text{Ins}}\) into lipids ranged from 43 to 49% of the radioactivity added to cells. Proteins were extensively delipidated, and anchor peptides were prepared as described (41). Anchor peptides were treated with HNO2 to liberate the \([^{3}H]_{\text {Ins}-}\text {P-lipid moieties}; the products were desalted by partitioning between butanol and water, and the labeled lipid products were separated by TLC and quantitated by radioscanning. Results of the quantitation of anchor lipids \(pG1\), \(pC1\), and \(pC2\) are given as percentage of the total radioactivity of anchor peptides. The remaining 7–11% that are not accounted for stayed at the origin and represent residual anchor peptides that had not been cleaved by HNO2 treatment. An aliquot of W303 anchor lipids is shown in Fig. 4B, lane 11.

Deletion of GPI7 Alters GPI Protein Transport and Remodeling—We previously reported on the accumulation of the immature 105-kDa ER form of Gas1p in \(\text{gpi7}^{-}\) mutants (21). We therefore investigated GPI protein transport in \(\Delta\text{gpi7}\). Indeed, by pulse-chase experiments we found that the maturation of GPI proteins \(\text{Sag1p}\) and \(\text{Gas1p}\) was slowed 2–3-fold as compared with wt cells, whereas the maturation of carboxypeptidase Y proceeded with normal kinetics (not shown). This indicates that the transport of GPI proteins in \(\Delta\text{gpi7}\) is specifically retarded. Nevertheless, in rich media \(\Delta\text{gpi7}\) cells grow at roughly the same rate as wt cells. They also incorporate \([^{3}H]_{\text{Ins}}\) with the same efficiency as wt cells.

The lipid remodeling of GPI anchors is significantly altered in \(\Delta\text{gpi7}\). As seen in Table I, the proportion of ceramides (\(pC1\) and \(pC2\)) in anchor peptides from \(\Delta\text{gpi7}\) is drastically decreased, whereas the fraction of DAG-containing lipids (\(pG1\)) is correspondingly increased. It should be noted that at the time of analysis, \(i.e., 75\) min after addition of \([^{3}H]_{\text{Ins}}\), the relative amounts of mild base-sensitive and mild base-resistant anchors are no longer changing and represent the steady state proportion of these two anchor types (41, 52). It is important to realize that \(pG1\) also represents a remodeled form of the anchor lipid in which a long chain fatty acid has replaced the original fatty acid present in sn-2 of the glycerol of the CP (Fig. 1). It thus appears that the relative decrease of ceramide remodeling goes along with a compensatory increase in DAG remodeling. A relative reduction in ceramide remodeling was also observed when we compared the efficiency of \([^{3}H]_{\text{Ins}}\) and \([^{3}H]_{\text{DHS}}\) incorporation into GPI proteins. As can be seen in Fig. 10, the ratio of \([^{3}H]_{\text{DHS}}/[^{3}H]_{\text{Ins}}\) incorporation into proteins is much higher in wt than in \(\Delta\text{gpi7}\) (Fig. 10, lanes 1–4). The lack of incorporation of \([^{3}H]_{\text{DHS}}\) in \(\Delta\text{gpi7}\) cannot be explained by an increase of the endogenous production of DHS in \(\Delta\text{gpi7}\), since the difference between wt and \(\Delta\text{gpi7}\) persists, even when all endogenous DHS biosynthesis is blocked by myriocin (Fig. 10, lanes 5 and 6). The defect in

|     | \(pG1\) | \(pC1\) | \(pC2\) |
|-----|--------|--------|--------|
| \(\text{W303}\) | 22.2   | 63.6   | 7.4    |
| \(\text{X2180}\) | 24.2   | 61.2   | 4.8    |
| \(\text{gpi8–1}\) | 23.8   | 63.9   | 2.6    |
| \(\Delta\text{gpi7}\) | 68.8   | 19.3   | 1.1    |

TABLE I

Quantitation of GPI anchor lipids

Fig. 9. \(\text{Gpi7p}\) is localized at the cell surface. Cells in the early exponential phase growing at 30 °C in YPD were used. A, intact W303 cells were treated with cysteamine chloride and then treated with the indicated concentrations of proteinase K (\(\text{prot K}\)) exactly as described (65) except that the EDTA concentration was raised to 20 mm. In lane 6, Triton X-100 was added to 1%. B, W303 cells were treated with zymolyase 20T at the indicated concentrations. C, W303 or \(\Delta\text{gpi7}\) cells were either lysed directly or after having been incubated for 30 or 60 min at 10 \(\text{A}_{600}/\text{ml}\) in the presence of cycloheximide (200 \(\mu g/ml\)) or tunicamycin (\(\text{Tm}, 20 \mu g/ml\)). Cells were lysed by boiling in sample buffer and processed for SDS-PAGE and Western blotting with antibodies against \(\text{Gpi7p}\), \(\text{Gas1p}\), or \(\text{Wbp1p}\).

may be underestimated due to ongoing proteolytic degradation during the 100,000 × \(g\) spin. Therefore the supernatant of the 8,000 × \(g\) spin was split whereby proteins were immediately precipitated with trichloroacetic acid in one half, and the other half was pelleted at 100,000 × \(g\). \(\text{Gpi7p}\) was exclusively found in the 8,000 × \(g\) pellet (\(\text{P8}\)) and thus is associated with either an internal pool of \(\text{Gas1p}\) as proposed earlier (50). A tighter interaction of \(\text{Gas1p}\) with some cell wall components found to be partially resistant to proteinase K, indicating either a small amount of core-glycosylated material is found in the ER in transit to the cell surface. For the moment it is unclear why \(\text{Gpi7p}\) was completely protected in microsomes, since it has been claimed that PM does not form closed vesicles upon homogenization (51). It is conceivable that \(\text{Gpi7p}\) resides in special PM subdomains that form closed vesicles upon homogenization or that centrifugation of microsomes generated protease-resistant membrane aggregates (51). In contrast, \(\text{Wbp1p}\) and the 108-kDa ER form of \(\text{Gas1p}\) were completely resistant to this treatment unless membranes were permeabilized with Triton X-100 (Fig. 9A, lane 6). The mature 125-kDa form of \(\text{Gas1p}\) was found to be partially resistant to proteinase K, indicating either a tighter interaction of \(\text{Gas1p}\) with some cell wall components or the existence of an internal pool of \(\text{Gas1p}\) as proposed earlier (50). Crude zymolyase treatment of intact cells also removed all of the mature form of \(\text{Gpi7p}\) (Fig. 9B), whereas recombinant zymolyase left \(\text{Gpi7p}\) intact (not shown). Longer exposures showed the presence of several minor bands of smaller size all of which were also present in \(\Delta\text{gpi7}\) cells except for a 108-kDa form of \(\text{Gpi7p}\) (Fig. 9C, lanes 7 and 8). This material seems to be an ER form in transit to the surface since it was no more detectable if cells were preincubated with either cycloheximide or tunicamycin (Fig. 9C, lanes 1–6). These data also indicate that mature \(\text{Gpi7p}\) is relatively rapidly degraded or becomes resistant to extraction with SDS. Globally these data indicate that the bulk of \(\text{Gpi7p}\) is exposed at the cell surface but that a small amount of core-glycosylated material is found in the ER in transit to the cell surface. For the moment it is unclear why \(\text{Gpi7p}\) was completely protected in microsomes, since it has been claimed that PM does not form closed vesicles upon homogenization (51). It is conceivable that \(\text{Gpi7p}\) resides in special PM subdomains that form closed vesicles upon homogenization or that centrifugation of microsomes generated protease-resistant membrane aggregates (51).
remodeling seems to be affecting mostly the maturation processes in the Golgi and/or PM (Golgi/PM remodeling) since, as shown in Fig. 10, lanes 7–10, the ratio of [3H]DHS/[3H]Ins incorporation into proteins in the Δgpi7/sec18 double mutant was the same as in sec18. Also, when using stringent conditions under which one observes only ER or only Golgi/PM remodeling (24), remodeling in the ER appeared relatively normal, whereas remodeling in the Golgi/PM was reduced (Fig. 10, lanes 11–14, 11′, and 12′). The relatively low amount of pC2 in anchor lipids of Δgpi7 (Table I) may be a consequence of this relative deficiency of Golgi/PM remodeling, since pC2 type anchors are only generated by the Golgi/PM but not the ER remodelase (24, 41). The relationship between the specific retardation of GPI protein transport, reduced Golgi/PM remodeling, and increased remodeling toward pG1 is for the moment unclear.

**DISCUSSION**

Yeast and mammals contain the same GPI carbohydrate core structure. This suggests that the GPI anchoring pathway has been established early in evolution and has rigorously been conserved in widely diverging organisms. On the other hand, the side chains added to this core as well as the lipid moieties of the anchor tend to vary a lot between different species (1). The GPI anchors of *S. cerevisiae* contain two types of side chains as follows: one or two mannoses are linked to Man3 (53) and an EtN-P side chain is linked to Man1.3 Both side chains are already present on the precursor lipid CP2 (22, 23). These two side chains are also found in some vertebrates, including mammals, and possibly in *Dictyostelium discoideum* (1), suggesting that not only the GPI core structure but also certain kinds of side chains have been invented and conserved since early times of evolution. Here we present evidence for yet a further, possibly conserved HF-sensitive substituent on CP2 which is attached to Man2. So far, the only side chain attached to Man2 reported in the literature is EtN-P. EtN-P was found by mass spectrometry on 15% of anchors of human erythrocyte cholinesterase and 3% of bovine liver 5′-nucleotidase (44, 46). Partial acid hydrolysis has also indicated an HF-sensitive substituent on Man2 in 40% of CD52-II (45). Analysis of the ethanolamine/Ins ratio in GPI anchors of porcine renal membrane dipeptidase and of human placental alkaline phosphatase yielded values of 2.5 and 2.4, suggesting the presence of EtN-P.

Fig. 10. Ceramide remodeling of GPI anchors is reduced in Δgpi7. Cells growing exponentially in SDCUA were used. Cells were labeled by the addition of either 25 μCi of [3H]DHS (D) or [3H]Ins (I) to an aliquot of 2.5 A°_250 of cells exactly as described (24). Lanes 1–6, precultures and labelings were at 30 °C. Cells were preincubated with (+) or without (−) 40 μg/ml of myriocin (myr) for 20 min before addition of the radiotracers. Lanes 7–10, sec18 or sec18 Δgpi7 double mutants were preincubated at 24 °C and preincubated for 10 min at 37 °C before addition of the tracer. Lanes 11–14, cells were labeled in the presence of cycloheximide under conditions in which only the Golgi/PM remodelase (lanes 11 and 12) or only the ER remodelase (lanes 13 and 14) is probed (Ref. 24, therein Fig. 7 (Golgi/PM remodelase) and Fig. 8 (ER remodelase)). All samples were processed for SDS-PAGE under reducing conditions and fluorography. Lanes 11′ and 12′ were scanned at an increased sensitivity.

Δgpi7 cells are hypersensitive to Calcofluor White and hence have some difficulty in constructing their cell walls. Several reasons can be envisaged. (i) The side chain on Man2 may be important for the interaction of CPs with the transamidase complex and for their efficient transfer onto proteins. Recent data show that a small reduction of Gpi8p renders transamidase activity rate-limiting.3 The synthetic effect of gpi7 mutations with gpi8 mutations suggests that deletion of GPI may have a similar effect. A decreased transamidase activity may particularly affect the anchoring of certain GPI proteins that have a low affinity for the transamidase even though the global rate of GPI biosynthesis and [3H]Ins incorporation into proteins of Δgpi7 is not grossly reduced. Thus it is conceivable that some GPI proteins important for cell wall architecture are lacking in Δgpi7. (ii) The side chain on Man2 may serve as an attachment point for the covalent linkage of β1,6-glucans to the anchor moiety of cell wall proteins although a recent analysis of the linkage region between the GPI anchor remnant and β1,6-glucans rather showed a direct glycosidic linkage between Man1 and the β1,6-glucan (6). (iii) The side chain may serve as a recognition signal for enzymes or proteins that facilitate the packaging of GPI proteins into vesicles, for remodelases that

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3 I. Imhof, U. Meyer, A. Benachour, I. Flury, E. Canivenc-Gansel, C. Vionnet, and A. Conzelmann, manuscript in preparation.
exchange their lipid moieties, or for hydrolases or transglycosidases that remove parts of the GPI anchor of cell wall proteins and hook the GPI remnant onto β1,6-glucans (4, 5).

Our data further show that ceramide remodeling in the Golgi/PIM is significantly reduced in Δgpi7, whereas remodeling toward pG1 is increased whereby it is not clear if pG1 remodeling is increased because ceramide remodeling is decreased or if ceramide remodeling is decreased because pG1 remodeling is increased. Moreover, the relationship of the alteration of GPI remodeling with the other phenotypic changes of Δgpi7 can be explained in several ways. (i) Previous studies showed that remodeling toward pG1 occurs in the ER and that retention of GPI proteins in the ER in secretion mutants maintains a high pG1/pC1 ratio on these proteins (41). Thus, if the substituent on Man2 of GPI anchors is important for efficient packaging of GPI proteins into transport vesicles, the delay in export of GPI proteins out of the ER may give the ER remodelase generating pG1 prolonged access to the GPI proteins and may thus cause a relative increase of pG1. (ii) The side chain on Man2 may serve as a recognition signal for Golgi/PM remodelase. (iii) We also considered the possibility that Gpi7p itself may be a Golgi/PM remodelase. This latter hypothesis would not directly explain why Δgpi7 cells cannot attach the HF-sensitive substituent onto Man2 and would imply that the addition of this side chain somehow is directed by the prior attachment of a ceramide moiety. However, this is clearly not the case, since CP2 also contains the HF-sensitive side chain on Man2, although its lipid moiety consists of DAG (23). Thus, we believe that the reduced Golgi/PM remodeling of GPI proteins in Δgpi7 is secondary to the lack of a substituent on Man2.

Our previous data suggested that CP2 can be transferred to proteins (23), and our working hypothesis until recently assumed that CP2 represents the GPI lipid used for GPI anchoring also by normal cells that do not accumulate this lipid (“CP2 hypothesis”). By consequence we would have predicted that all the enzymes required for the elaboration of CP2 are localized in the ER. Paradoxically, the subcellular fractionation experiments and protease treatment of intact spheroplasts strongly suggest that the bulk of Gpi7p resides at the cell surface (Fig. 9, A and B). Moreover, although we recently succeeded in demonstrating the presence of an HF-sensitive group on Man1 of immature ER forms of GPI proteins, we presently lack the tools to look for such a group on Man2. Thus, the so far available data raise a doubt whether it is CP2 which is added to GPI proteins in the ER, and we therefore are presently considering the possibility that other GPI lipids than CP2 are the physiological substrate of the ER transamidase. In fact, neither CP2 nor M4 can be detected in wt cells. It therefore seems possible that under physiological conditions cells add M4 to GPI proteins (“M4 hypothesis”) and that CP2 is elaborated only in mutants in which M4 cannot be transferred to proteins, spills out of the ER, and reaches the PM. It is noteworthy that Δgpi7 incorporates [3H]Ins at a normal rate into proteins suggesting that the transamidase is perfectly able to transfer M4. Thus, the side chain on Man2 may normally not be added to GPI proteins or only be added after GPI proteins arrive at the surface. The M4 hypothesis, however, does not explain why M0 and M4 accumulate in Δgpi7, whereas M0, M4, and CP2 remain undetectable in wt cells (Fig. 2, lanes 1 and 2) or why gpi8-1, deficient in the transfer of GPIs onto proteins, accumulates CP2 (19, 21). It also fails to explain the delayed maturation of GPI proteins and the reduced rate of GPI remodeling observed in Δgpi7. To save the M4 hypothesis, the accumulation of GPI intermediates in Δgpi7 could be rationalized by assuming that the substituent on Man2 serves to mark supernumerary GPIs for degradation, but also this assumption does not explain the observed accumulation of CP2 in gpi8. Thus, although our results raised the possibility that M4 is the physiological GPI lipid for GPI anchoring, this M4 hypothesis leaves many results unexplained and the data are more easily explained by our original CP2 hypothesis. For one, the synchronous accumulation of M4 and CP2 in all our gpi8 mutants argues that M4 is not a better substrate for the transamidase than CP2. CP2 may physiologically be produced by the small amount of Gpi7p in the ER (Fig. 9C). Alternatively, it is conceivable that M4 is transported from the ER to the PM, is converted there to CP2, and is then transported back to the ER by some not yet elucidated mechanism. In this context it is noteworthy that the biosynthesis of GPIs by wt microsomes in vitro produces CP2 in good yield, i.e. the in vitro system adds the substituent on Man2. This in vitro system does not contain cytosol nor GTP and hence should not support vesicular transport from ER- to Golgi-derived microsomes (59). It is possible, however, that GPI lipids are transported between microsomes or membrane fragments by means of lipid transfer proteins or through direct contact between membranes. It also can be envisaged that juxtaposition of membranes allows enzymes present in one membrane to work on lipids in another membrane. The same mechanisms may also operate in intact cells. Clearly the identity of the physiological GPI lipid substrate of the transamidase will have to be established by further experiments.

Homology searches show that two ORFs of S. cerevisiae are related to GPI7, MCD4 (= YKL165c), and YLL031c. They belong to a novel gene family comprising for the moment the nine members shown in Table II which, based on the many predicted transmembrane domains, were previously classified as putative permeases (60). Pairwise alignment allows us to group

| Table II | GPI Anchor Side Chains |
|---------|-----------------------|
| 1       | YJL062w = GPI7, S.c.  |
| 2       | YA93, (1175452), S.p. |
| 3       | (1132507), C. elegans |
| 4       | YLL031c, S.c.         |
| 5       | (2984587) homo sapiens|
| 6       | (2257562) S.p.        |
| 7       | (2734088) C. elegans  |
| 8       | MCD4, YKL165c S.c.   |
| 9       | (2879870) S.p.        |
them into three subfamilies of more closely related ORFs. All nine ORFs predict membrane proteins of about 100 kDa having an N-terminal signal sequence, a hydrophilic N-terminal part, and multiple transmembrane domains in their C-terminal half. mcd4 mutants were obtained in a screen for cells deficient in the cell cycle controlled polarization of growth, a phenotype also generated by mutations in the exo cit or in N-glycosylation (61). The subfamilies typified by PIGF and YLL031c are more closely related to each other than to the MCD4 subfamily. All nine family members contain two conserved motifs at about the same position in the hydrophilic N-terminal domain, namely HXLGDXXXGH and DHGMXXGXGH. These motifs are also found in two EST clones from human cDNA that have high homology to MCD4 (NCBI PID 177947 and 1765215, ClustalW alignments giving aligned scores of 46 and 35). Very interestingly, by a reiterated Psi Blast search at the National Center of Biotechnology Information (NCBI) (62) one can find a highly significant homology of all three subfamilies with a large family of phosphodiesterases. The large majority of these homologous sequences encode mammalian cell-surface proteins classified as alkaline phosphodiesterase I, nucleotide pyrophosphatase, or alkaline phosphatase. The homology comprises a region of about 220–240 amino acids in the N-terminal hydrophilic part of PIGF, YLL031c, and MDC4. The homology of PIGF in this region with mammalian and plant phosphodiesterases amounts to 17–18% identity and 30–34% similarity and comprises a motif PTXXT, TGX, which is common to bacterial, viral, plant, and mammalian phosphodiesterase. This homology may suggest that Gpi7p itself is the transference adding the phosphodiester-linked substituent on Man2. In this context it is interesting to note that the Etn-P on Man3 of the GPI anchor has been shown to be transferred by transesterification using phosphatidylethanolamine as donor of Etn-P (63, 64). Mutants in YLL031c also accumulate abnormal GPI intermediates which on TLC have about the same mobility as M4 suggesting that YLL031c is similarly involved in adding Etn-P.2 Thus, it is conceivable that not only the PIGF subfamily but also other subfamilies are involved in the transfer of Etn-P onto the GPI core structure. However, the transesterification activity of Gpi7p will have to be shown directly before one can exclude that the primary function of Gpi7p is to generate some signal from the cell wall which regulates GPI protein transport and remodeling as well as side chain addition to structures.

It is interesting to note that side chain members belonging to different species are more closely related to each other than family members belonging to a single species. This can be seen when comparing the pairwise alignment scores among the three ORFs of S. cerevisiae or the three ORFs of Schizosaccharomyces pombe with the scores among subfamily members (Table II). In evolutionary terms this suggests that the divergence of these three subfamilies occurred earlier than the separation of the lineages leading to S. cerevisiae, S. pombe, and Caenorhabditis elegans. This implies that the HF-sensitive group on Man2 is of very ancient origin. PIGF bears no resemblance with PIG-F, a mammalian gene encoding for a highly hydrophilic membrane protein involved in the addition of Etn-P to Man3 (10). The exact role of PIG-F has not yet been elucidated.

It will be interesting to find the human homologues of PIGF. It may be that this gene, as in S. cerevisiae, plays a more dispensable role in GPI anchoring than the enzymes involved in the elaboration of the carbohydrate core structure such as for instance PIG-A/GPI3/CWH6/SPT14 (7). Thus, although deficiencies in PIG-A are only acquired by somatic cells, deficiencies in the human PIGF homologue may be transmittable through the germ line as well.

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Deletion of GPI7, a Yeast Gene Required for Addition of a Side Chain to the Glycosylphosphatidylinositol (GPI) Core Structure, Affects GPI Protein Transport, Remodeling, and Cell Wall Integrity

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