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TOWARDS A THERAPY FOR PHOSPHOMANNOMUTASE 2 DEFICIENCY, THE DEFECT IN CDG-Ia PATIENTS

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SUMMARY

Phosphomannomutase (PMM2, Mannose-6-P→ Mannose-1-P) deficiency is the most frequent glycosylation disorder affecting the N-glycosylation pathway. There is no therapy for the hundreds of patients who suffer from this disorder. This review describes previous attempts at therapeutic interventions and introduces perspectives emerging from the drawing boards. Two approaches aim to increase Mannose-1-P: small membrane permeable molecules that increase the availability or/and metabolic flux of precursors into the impaired glycosylation pathway; and, phosphomannomutase enhancement and/or replacement therapy. Glycosylation-deficient cell and animal models are needed to determine which individual or combined approaches improve glycosylation and may be suitable for preclinical evaluation.

Keywords

Glycosylation disorder; CDG; phosphomannomutase2; phosphomannose isomerase; therapy; high throughput screening

INTRODUCTION

Inherited deficiency of phosphomannomutase (PMM2) causes a human glycosylation disorder previously known as Congenital Disorder of Glycosylation Ia (CDG-Ia) [1]. Pediatric patients have a broad and variable clinical picture that affects nearly all systems leading to failure to thrive, hypotonia, variable developmental delay, ataxia, dysmorphism, skeletal abnormalities, coagulopathy with up to 20% mortality in the first 5 years due to organ failure and/or severe infections [2–5]. Later on they show mental retardation, stroke like episodes, and retinitis pigmentosa. Adults are usually wheelchair bound with peripheral neuropathy and stable mental retardation. The spectrum is broad and recently a considerable number of very mild patients have been reported [6–10]. There is no therapy for >600 identified patients; many more patients probably remain undetected. Scores of mutations have been identified in PMM2-deficient patients. Some of these patients have up to 25% normal enzymatic activity, while heterozygous parents with 50% activity are asymptomatic [9]. This surprisingly high threshold raises the question: how much enzymatic activity is required to prevent pathology? Since no precise genotype-phenotype correlation is seen [11], an ensemble of other unidentified genes must play important roles. However, patients with high residual activity tend to have milder phenotypes [11,12]. While developmental defects that occur *in utero* cannot be reversed, even

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a small improvement in the effective enzymatic activity might reduce the daily medical burden on patients and their families. This hope makes the pursuit of a therapy worthwhile.

The purpose of this article is to review previous attempts at therapy for PMM2- deficient patients and to present the rationale for the development of new ones.

METABOLIC PATHWAYS

PMM2 is an essential gene and its complete loss is lethal in yeast, mice and presumably humans [13,14]. PMM1 is a homolog of PMM2 [15], and catalyzes the same reaction, but it does not appear to substitute for loss of PMM2 in mice [16]. PMM1 ablation in mice produces no obvious pathology or phenotype [16]. Recent studies show that this enzyme, which is highly expressed in brain, is an IMP-stimulated glucose 1,6-P phosphatase [17]. PMM2 catalyzes the conversion of Mannose-6-P (Man-6-P) into Mannose-1-P (Man-1-P) and it is the first committed step in synthesis of the activated mannose donors GDP-mannose and dolichol-P-Mannose, as shown in Figure 1. Synthesis of N-glycans, O-mannose linked glycans, glycosylphosphatidylinositol anchors, and C-mannosylated proteins all require these donors. Cells transport exogenous mannose and convert it to Man-6-P using hexokinase. Alternatively, Man-6-P is formed from Fructose-6-P using phosphomannose isomerase (PMI or MPI). Both exogenous glucose and glycogen-derived glucose can contribute to this pathway [18,19]. The majority of mannose in N-glycans is derived from glucose in most cells (Freeze and Sharma, unpublished observations). Deficiencies in PMM2 and MPI reduce the amount of Man-1-P and Man-6-P [20,21], respectively, that are available for glycosylation, resulting in unoccupied N-linked glycosylation sites on a number of proteins. The absence of glycan chains is thought to cause the pathology seen in these patients. Excess Man-6-P derived from mannose is catabolized beginning with PMI. All of the biosynthetic reactions are thought to occur in the cytoplasm or on the cytoplasmic face of the endoplasmic reticulum.

SETTING THE STAGE FOR THERAPY: WHAT IS POSSIBLE AND PRACTICAL?

Cellular, genetic, and metabolic options should be considered to treat PMM2-deficient patients. Some are more practical than others and/or may be achievable within a reasonable timeframe; but all have shortcomings. See Table 1 and Figure 2 for an overview and summary. Applications of stem cell therapy are in the far future [22]. While they may hold promise for treating inherited liver disorders [23,24] or those disorders amenable to hematopoietic stem cell replacement [25,26] treating systemic glycosylation disorders would be considerably more difficult because of the ubiquitous and cell-autonomous nature of protein glycosylation. This approach will not be considered here in favor of near term options and better targets. The goal of therapy for PMM2-deficient patients is to increase the flux of metabolic precursors into the impoverished glycosylation pathways. Options include: increase the activity of PMM2, provide Man-1-P directly, increase the Man-6-P pool and/or redirect this substrate from glycolysis to glycosylation. Enzyme activity could be increased by gene therapy that included stem cells, but it is also unlikely to be useful in the near future, given the inherent risks and very small patient population [22]. Enzyme replacement therapy has not been tried and insulin is the only reported activator of PMM2 [27]. Man-1-P is not membrane permeable, but hydrophobic derivatives can enter cells and correct impaired glycosylation. However, these compounds tend to be unstable and toxic. Increasing exogenous mannose can improve [³H] mannose incorporation in PMM2-deficient cells, but mannose therapy was not effective in patients.

MANNOSE SUPPLEMENTS: SIMPLE BUT INEFFECTIVE

MPI-deficient patients (CDG-Ib) benefit from alimentary addition of 300–750 mg/kg/day Man [12,28] because it bypasses the defective step by allowing formation of Man-6-P via hexokinase (Figure 1). These children fail to thrive, coagulopathies, protein-losing enteropathy and liver fibrosis, and they show remarkable improvement when the therapy is initiated [12,29]. Fortunately, there are no neuronal pathologies in this disorder. Possibly mannose in mother's plasma provides an "endogenous therapy" during fetal life.

In contrast, mannose treatment in PMM2-deficient CDG-Ia patients was not successful. There was no measurable improvement in any clinical parameters in children during treatment [30, 31] nor was serum hypoglycosylation decreased. The original basis for suggesting the treatment was that PMM2-deficient fibroblasts synthesize a truncated LLO glycan, which becomes normal when cells are incubated with high concentrations of mannose [32]. The most likely explanation for the failure of mannose therapy in the patients was that Man-6-P resulting from increased mannose was not available because it was simply catabolized via MPI, since its activity remains normal in PMM2-deficient patients.

Another explanation for the failure of mannose therapy in these patients was suggested based on observations using streptolysin O-permeabilized cells [33,34]. These studies showed that addition of Man-6-P to cells preferentially reduced the amount of the major LLO species, $\text{Glc}_3\text{Man}_9\text{GlcNAc}_2\text{-PP-Dol}$ resulting in the liberation of the free glycan. Thus, increasing Man-6-P in either PMM or MPI deficient cells could potentially decrease the amount of available LLO and actually exacerbate deficient glycosylation. However, providing mannose to either PMM2- or MPI-deficient cells increases the deficient GDP-Mannose pools [35], showing that the metabolic flux increased, but it did not increase either the Mannose-6-P pool or decrease cellular glycosylation [36]. In contrast, providing mannose to intact *Mpi*-null mouse fibroblasts greatly increases Man-6-P pools and eventually depletes ATP stores, since there is no metabolic exit for Man-6-P after saturating the glycosylation pathway [37]. Increased Man-6-P in *Mpi*-null cells decreased the amount $\text{Glc}_3\text{Man}_9\text{GlcNAc}_2\text{-PP-Dol}$ LLO, but it did not appear to affect the amount of protein glycosylation [36]. This suggests that the amount of LLO is in excess. Changes in steady state level do not necessarily reflect the amount of flux through the pathway. It is important to distinguish between these.

BYPASS THE DEFECT: MEMBRANE PERMEABLE MANNOSE-1-P

Another potential approach is to provide Man-1-P directly to cells. However, Man-1-P does not diffuse, and is not transported across the cell membrane. To overcome this issue, several groups attempted to chemically synthesize cell permeable Man-1-P compounds that would be converted to free Man-1-P after entering the cell, i.e., prodrugs [38,39]. A series of compounds have been reported [38–40]. One set showed some biological activity [38]. Man-1-P can be made membrane permeable by covering phosphates with acetoxymethyl groups and protecting the OH-groups with ethylcarbonate or acetyl esters [38]. In cell culture, these compounds correct the LLO phenotype in several glycosylation-deficient fibroblasts at a lower concentration than Man, and also compete with radiolabeled Man in a glycosylation assay in cells [38]. The study also showed that the level of GDP-Man increased in response to the compounds. This validates the theoretical concept. However, all compounds synthesized to date are far too unstable to be clinically useful. The best compound has a half-life in serum of about 2.5 minutes. This is further confounded by the relatively high concentrations needed to get sufficient uptake through the cell membrane. The best compounds must be added to the medium at a high concentration of around 100 μM to be effective. A third issue is the toxicity of these compounds. The most effective compound synthesized so far shows negative effects on protein synthesis and cell viability at concentrations above 100 μM , i.e. in the same range

as its therapeutic dosage. In a more recent study [40], Man-1-P pro-drugs containing an additional phosphodiester-linked mannose residue along with a benzyl or phenyl group on the phosphate were reported to generate intracellular Man-1-P based on their ability to compete with ^3H -mannose for glycosylation. The compounds were also reported to have minimal toxicity after 16 hours based on release of lactate dehydrogenase (cell lysis). Additional studies are needed to confirm and extend these observations. Using less toxic blocking groups [41], or lipid carriers for delivery of free Man-1-P is another. Apart from the labor-intensive synthesis of making a stable, non-toxic and well-transported compound, no animal study system is currently available. All PMM2-deficient patients have some residual enzymatic activity, but complete elimination of *Pmm2* in mice is lethal in early embryogenesis [14]. This is also true for the *Mpi*-null mouse [37]. Model systems with quantifiable phenotypes will be needed to assess any therapy.

ENZYME REPLACEMENT THERAPY

Enzyme replacement therapy might be possible for PMM2-deficient patients. This approach has worked well for patients with lysosomal storage diseases. Gaucher's patients who are deficient in glucocerebrosidase are treated with Cerezyme a form of the expressed enzyme carrying Mannose-terminated N-glycan chains that targets the enzyme to macrophage in the liver spleen and skeleton the undegraded glucosecerebroside ($\text{Glc}\beta\text{-Cer}$) accumulates [42–44]. The enzyme gradually reduces the accumulation of the substrate since it is trafficked to its normal site of action in the lysosome [42–44]. Enzyme replacement therapy has also been approved for other lysosomal storage disorders including Pompe Disease (α -glucosidase deficiency) and MPS1 (α -iduronidase deficiency). Many of these patients have very low residual enzymatic activity (a few percent) and even slight increases of activity within the lysosome may be sufficient to improve patients' symptoms gradually as the accumulated substrate decreases. This may not true for PMM2-deficient patients where slight increases in activity might still be within the classical pathological range. Moreover, targeting PMM2 to the lysosome is not desirable, since it would likely be degraded; it needs to remain in the cytoplasm.

Many studies show that coupling arginine-rich peptides to high molecular mass cargos delivers them into cells, through an uptake system that probably involves multiple receptors, [45] and this has proven to work in rodents *in vivo* [46]. Whether sufficient PMM2 could enter cells and reside in the cytoplasm with an attached cationic sequence is not known. A slightly modified approach may offer another opportunity by binding cationic-derivitized molecules to heparan sulfate on the cell surface [47]. By converting all ammonium groups of the antibiotic neomycin to guanidinium groups, molecules >300 kDa could cross cell membranes at nanomolar transporter concentrations with delivery being entirely dependent on cell surface heparan sulfate proteoglycans. By conjugating the guanidinoneomycin to the ribosome-inactivating toxin, saporin, the study showed proteoglycan-dependent delivery of cargo into the cytoplasm. Coupling of typical arginine-rich peptides showed both heparan sulfate-dependent and -independent cellular uptake [47]. Since cell surface heparan sulfate is nearly ubiquitous, this presents an opportunity to deliver PMM2 to these cells.

The disadvantage of enzyme replacement therapy is the limited amount of enzyme that can be delivered to a cell/tissue. Another is that the enzymes are unlikely to cross the blood brain barrier providing limited benefit for brain and central nervous system pathologies.

ACTIVATE OR STABILIZE PMM2

The only reported activator of PMM2 is an insulin-dependent 2-fold increase in PMM2 activity seen in *Cos7* cells over expressing the enzyme. Transcription does not increase. Rather, it seems to involve a serum and glucocorticoid-regulated kinase (*Sgk1*), which regulates channels and

transporters including the renal epithelial Na(+) channel (ENaC) [48]. In vitro phosphorylation of PMM2 by this kinase completely eliminates activity both in the absence and in the presence of insulin. The interaction with PMM2 was observed in a yeast two-hybrid screen using Sgk1 as bait, and was confirmed by co-immunoprecipitation and Sgk1-dependent in vitro phosphorylation of PMM2. These data suggest that Sgk1 may affect glycosylation, but the physiological relationship is not clear.

There are no small molecule activators of PMM2, but the availability of large chemical libraries and robotic high throughput screening facilities offer a robust opportunity to explore this avenue. The simple coupled enzymatic assay using MPI, phosphoglucose isomerase and glucose-6-P dehydrogenase can be easily adapted for compound screening. Initial leads and structural optimization of selected chemical scaffolds can be combined with docking studies to further refine and optimize the structure activity relationships activators. The structure of PMM2 has not been solved, but the structure of its homolog, PMM1, was determined by X-ray crystallography [49]. The structure suggested a likely, but complex, reaction mechanism involving cap and core domains, which open to bind substrate and then close to provide a solvent-exclusive environment for catalysis. The substrate first binds to the cap and then is swept into the active site when the cap closes. Repulsion of positive charges at the interface of the cap and core domains stabilizes alpha-PMM1 the open conformation, but negatively charged substrate binds to the cap, thereby facilitating its closure over the core domain. A potential allosteric or conformational-dependent activator could be tested initially using normal PMM2, but its activity must also be confirmed on PMM2 carrying several common mutations to select the compounds with broadest application. A small molecule could potentially pass the blood brain barrier. A disadvantage of the approach is the possibility that effective inhibitors are more effective on some mutant forms than on others, which may limit their general application.

Enzyme enhancement therapy has shown some success in treating lysosomal storage disorders [42,43]. This approach exploits the well-known fact that substrates or their analogs stabilize enzymes. Competitive inhibitors bind at or near the active site of the mutated enzymes at neutral pH and are then released at low pH when the enzyme arrives in the lysosome. These molecules may only marginally increase the residual activity up to a few percent of normal, but this is sufficient to reduce the ongoing accumulation of non-degraded substrates. A competitive inhibitor is probably not useful for PMM2 because its synthesis and continued residence in the cytoplasm precludes a pH-dependant inhibitor dissociation to allow substrate binding. Identification of an allosteric activator, as described above, has a greater chance of success. Recently, a combination of protein specific pharmacologic chaperones and induction of the protein folding machinery synergized to allow mutated proteins to survive within the secretory pathway [50]. This approach may hold promise for ER or Golgi associated proteins that are mutated in other types of glycosylation disorders, but similar approach for PMM2 would require its stabilization and dampening of cytoplasmic protein degradation.

INCREASE MANNOSE-6-P FLUX INTO GLYCOSYLATION PATHWAYS

It should be possible to encourage the flux of Man-6-P toward the glycosylation pathways in PMM2-deficient cells. Metabolic labeling with [³H] Mannose showed these cells synthesize truncated LLO species and transfer them to proteins [32]. Addition of 250–500uM mannose to the culture medium, normalizes both the LLO pattern and the size of protein-bound glycans. Hyperphysiological (severe diabetic) concentrations of glucose (>10mM) also corrected the size, but clearly mannose was >20 fold more effective [32]. Since mannose is well tolerated at 4–5x the normal plasma concentration of 50uM, it would be reasonable to increase plasma mannose concentration. As mentioned above, mannose is not effective in PMM2-deficient patients, most likely because MPI is fully active and consumes the Man-6-P. The unfavorable

ratio of PMM:MPI activity does not favor glycosylation. Increasing mannose delivery to the cells either by increasing extracellular mannose concentration or by improving its delivery into the cells might improve this situation. The antidiabetic drug, metformin, which stimulates the AMP-activated protein kinase, was reported to provide a preferential 2-fold stimulus of mannose uptake into fibroblasts [51]. However, this stimulation is not seen at normal plasma glucose concentration (5mM), so it is unlikely to increase mannose delivery in normal conditions. Another way to increase Man-6-P is by increasing the flux of glucose into fructose-6-P via glycolysis. One study showed that 5-aminoimidazole-4-carboxamide riboside (AICAR), which activates glycogen phosphorylase and suppresses glucose-6-phosphatase transcription [52] can provide mannose for glycan synthesis [34,53]. Other studies also show that glycogen is a source of mannose in the plasma [19] AICAR was found to prevent fat accumulation in sedentary mice and was suggested as a way to increase endurance without exercise [54]. PMM2-deficient patients have normal plasma glucose and apparently normal glycogen stores. For unexplained reasons, they usually show a severe failure to thrive as infants, and as children, they struggle to gain weight, and are sometimes fed through gastrostomy tubes. Use of AICAR as a way to increase intracellular Man-6-P in PMM2-deficient patients would require caution.

An approach, which we have pursued, hinges on the observation that the amount of exogenous mannose directed toward glycan synthesis (PMM2-dependent) vs. catabolism (MPI-dependent) is determined by the ratio of PMM:MPI. Higher PMM2 activity, and lower MPI activity, drives more Man-6-P into the glycosylation pathway. Combining MPI inhibitors with hyper-physiological concentrations of mannose may increase Man-6-P flux in favor of glycosylation.

MPI is a zinc-dependent enzyme and so typical chelators such as 1,10-phenanthroline, EDTA and 2,2'-dipyridyl are effective inhibitors [55,56]. Competitive inhibitors include phosphorylated analogs such as arabinose-5-phosphate, erythrose-4-P, and ribose-5-P, but none of these are specific or membrane-permeable. Fructose-1-P was reported to inhibit MPI and this was offered as an explanation for why patients with uncontrolled hereditary fructose intolerance make hypoglycosylated proteins [57]. Previous screening of a combinatorial chemical library identified a membrane permeable compound, but its biological effects were uncharacterized [58]. The crystal structure of *Candida albicans* MPI is known [59], and software programs can model docking of substrate and/or potential competitors.

We used high throughput screening of chemical libraries to identify a first generation of MPI inhibitors [60]. The traditional MPI-coupled enzyme assay leading to NADPH⁺ formation was used, along with counter screens to eliminate effects on the coupling enzymes (phosphoglucose isomerase and G6PD) are required for production of NADPH⁺ [55]. After identifying possible inhibitors, dose-response, and IC₅₀ values for either competitive or (preferably) non-competitive inhibitors were done using the coupled assays. Direct in vitro confirmation is easy using only MPI, based on formation of ³HOH from [2-³H]Man-6-P (Figure 3) and expressed enzyme [28,37,61]. This simple assay can be directly applied to cells to assess the biological effectiveness of compounds by comparing the ratio of [2-³H]mannose incorporation into glycans (TCA precipitate) vs. ³HOH production. The inhibitor should increase the incorporation of label into glycans and decrease the amount of ³HOH produced. This method can be used to assess the effects of inhibitors in more complex models such as mice or zebrafish. Previous studies using mice and most cell lines show that the great majority of [2-³H] mannose is catabolized via MPI, with production of ³HOH ([61,62]; and unpublished observations). Increased organ specific incorporation of label into glycans would indicate that the MPI-specific inhibitors have been effective at that location. An inhibitor should decrease the amount or rate of ³HOH appearance in the plasma, but the organ/tissue that produced it could not be determined.

We screened ~200,000 compounds and identified a series of compounds with IC₅₀'s in the range of 1–5 μM. Tests on a multiple cell lines showed that, only a few of these increased [2-³H] mannose incorporation into proteins and lowered formation of ³HOH. The increased incorporation is consistent with about a 70% reduction MPI activity. Commercially available and custom-synthesized analogs are being interrogated in order to increase the efficiency of inhibition.

Such inhibitors have limitations. Besides the obvious caveats of off-target effects, toxicity, and short half life, inhibitors may have limited access to some tissues or only be effective in selected tissues where the normal ratio of PMM:MPI is favorable. For instance in the mouse, the MPI specific activity in liver was measured at about 15 nmole/min/mg protein, but it is nearly 60 nmole/min/mg in heart and ~120 nmole/min/mg in testis. It may be more challenging to reduce MPI activity sufficiently to drive the elevated plasma mannose into the glycosylation pathway in these tissues with high MPI activity.

FUTURE PERSPECTIVES AND CHALLENGES

Individual or combinations of PMM2 activators, MPI inhibitors+plasma mannose, and PMM2 replacement may offer hope to improve glycosylation in some PMM-deficient patients. Small molecules may traverse the blood brain barrier, but PMM2 will not. On the other hand, the half-life of PMM2 may be significantly longer than that of either inhibitors or activators. Differential accessibility of small molecules and of PMM2 are significant hurdles to overcome. Since PMM2- deficiency affects so many organ systems in acute, chronic, and progressive ways, it is worthwhile to consider simultaneous exploration of all these approaches. It is difficult to predict which aspects of PMM2-deficient patients' phenotype would respond to therapy.

The major limitation on development of these therapies is the lack of appropriate model systems, since the knockout mouse is lethal [14]. It is unlikely that organ- or temporal-specific knockout expression, e.g., floxed alleles, could be generally useful, but this would depend on the organ-to-organ variability in excision efficiency. Moreover, each cell is responsible for generation of its own Man-1-P and elimination of all PMM2 activity would preclude analysis of Pmm2 activators or Mpi inhibitors. Knock-in of human disease-causing PMM2 mutations may offer hope, although animals with observable phenotypes must be generated while avoiding embryonic or early post-natal lethality. A construct that allowed temporal or organ-specific elimination of the normal allele and simultaneous expression of the selected mutation might provide a useful model. Another approach might be to identify PMM2 shRNAi molecules that give 80–90% reduced enzymatic activity and stably integrate them into 2–4 cell embryos via lentiviral constructs [63]. This procedure could yield individuals for phenotypic analysis, but these viral constructs frequently integrate into multiple sites on different chromosomes [63], which would complicate the creation of a stable knockdown line. Nevertheless, such experiments may provide information on a range of phenotypic abnormalities in a series of individual mice. Zebrafish present another opportunity to test the therapeutic agents since the morpholino mutants produce hypomorphic alleles, similar to patients [64,65]. This system offers great promise. A non-genetic approach to generate a PMM2-deficient model system would be to identify a specific inhibitor of PMM2. This could easily be done in the same high throughput screen searching for a therapeutic PMM2 activator. A rather different approach would be to reduce the impact of hypoglycosylation of key proteins as suggested by Shang et al [66]. They propose that tailoring the rate of protein synthesis to impaired flux through the glycosylation pathway might be useful

CONCLUSIONS

In the last decade we have made great progress in identifying a few dozen human glycosylation disorders [2,4]. We have made almost no progress in providing the patients and families with potential therapies. The advent of high throughput compound screening offers a new avenue for development of therapies targeting the N-glycosylation pathway. The technology provides tools and reasonable approaches to assess the efficacy of potential lead compounds. Robust PMM2 deficiency models are needed to test these concepts and molecules before they will lead to useful products. International cooperation and collaboration among scientists are essential if we are to do “the right thing” and offer children and families evidence of our unified hopes for their future.

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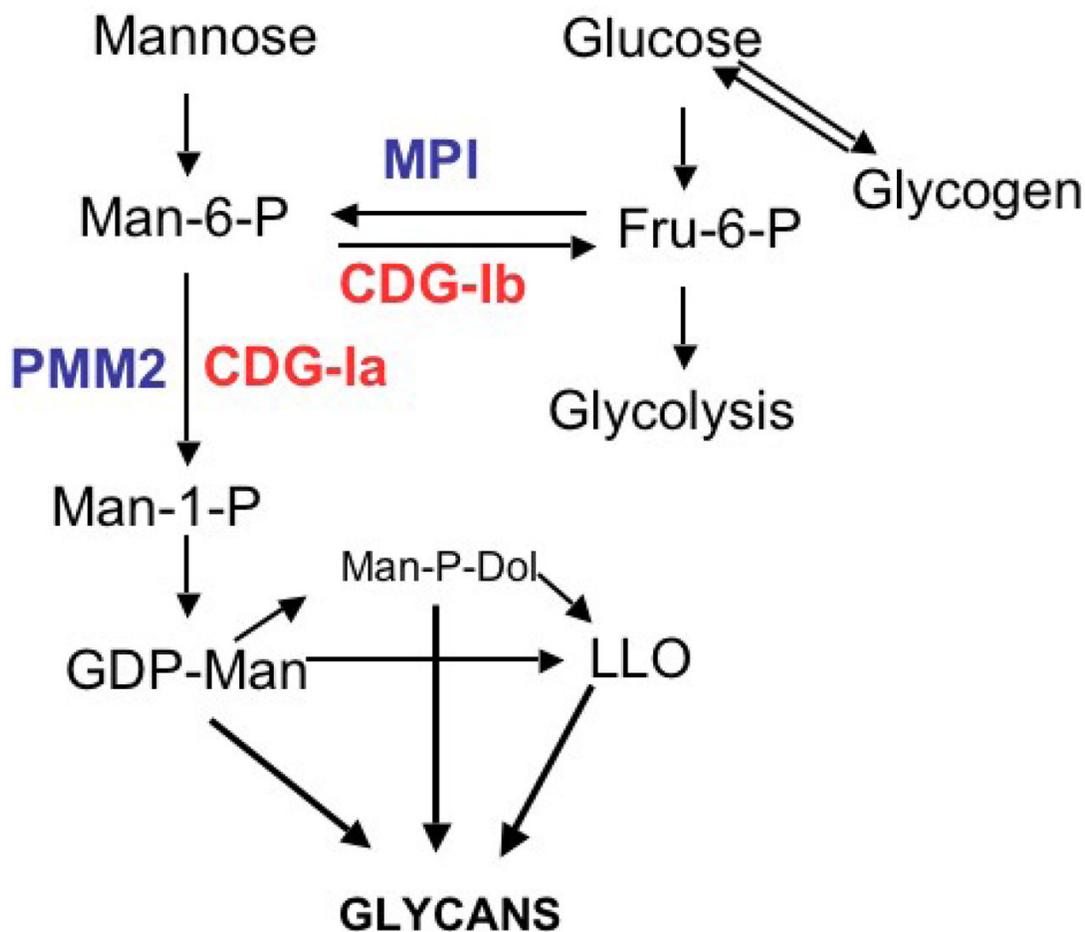


Figure 1. Mannose-related Metabolic Pathways in Glycan Synthesis

PMM2 deficiency, the cause of CDG-Ia, affects the first committed step required for the synthesis of mannosylated glycans. Reduced production of Man-1-P reduces its flux through the pathway and leads to reduced N-glycosylation. Whether other pathways are affected is not known. The goal of therapy is to increase the amount of Man-1-P available for glycosylation. This can be done by providing Man-1-P directly or by increasing the flux of Man-6-P to Man-1-P. Providing normal PMM2, stabilizing or activating the endogenous enzyme or driving more Man-6-P into the pathway may increase the amount of Man-1-P and improve impaired glycosylation. See text for a detailed discussion.

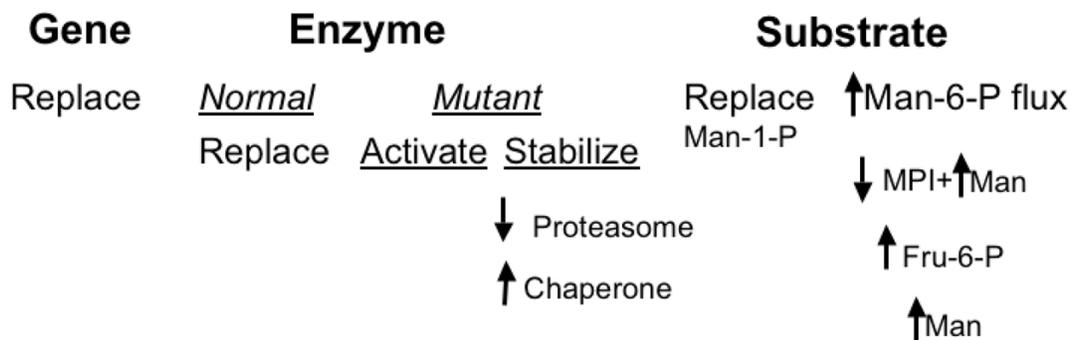


Figure 2. Summary of Potential Approaches for Treating PMM2-Deficient Patients

Therapeutic options include replacement of the gene, PMM2, supplementation with normal enzyme PMM2, or altering the stability or activity of the mutant enzyme. In addition, the substrate (Man-1-P) can be enhanced by direct replacement or alterations of the metabolic flux into the glycosylation pathways. These options are discussed in the text.

Formation of $^3\text{H}_2\text{O}$ from $[2\text{-}^3\text{H}]$ Mannose

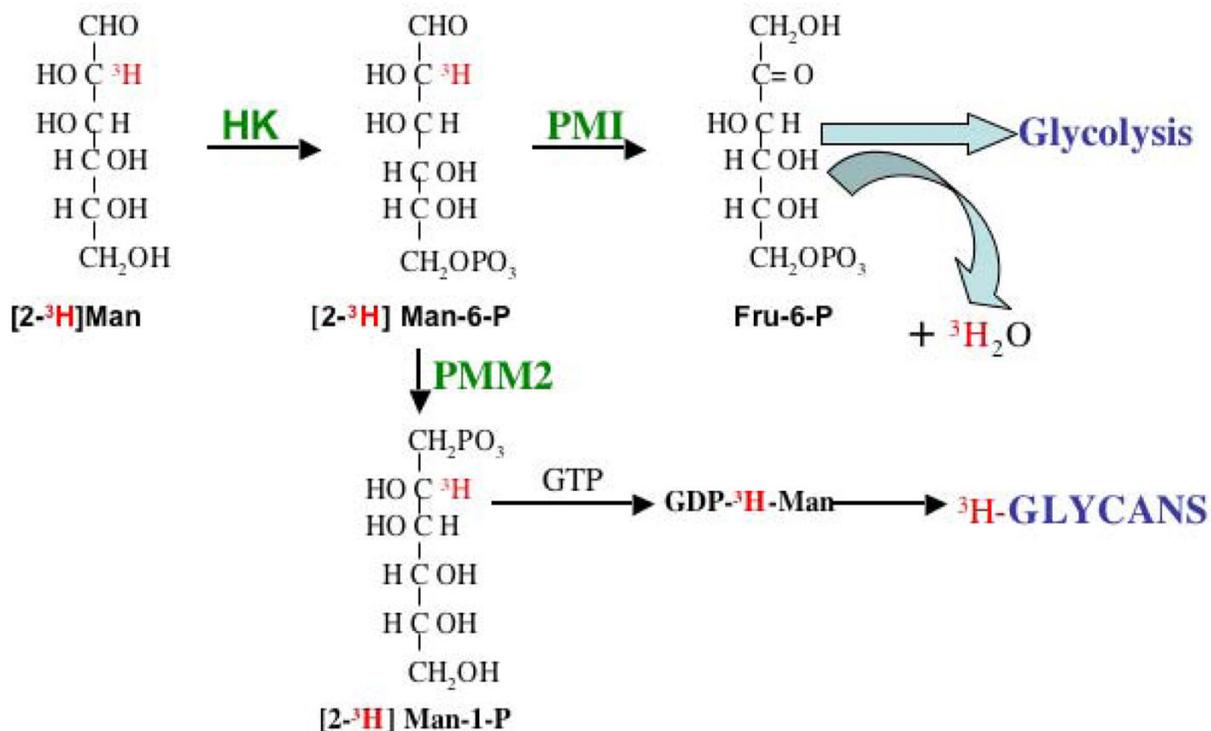


Figure 3. Metabolism of $[2\text{-}^3\text{H}]$ Mannose: the metabolic fate of exogenous mannose
 This versatile radiolabel is the best indicator of mannose flux into these pathways.

THERAPEUTIC APPROACHES FOR PMM2-DEFICIENT PATIENTS

Approach	Rationale	Advantages	Disadvantages
Enzyme Replacement	Provides normal active enzyme	Corrects defect in cells that take up and localize sufficient enzyme	Unequal accessibility to cells, especially CNS; would not cross blood brain barrier, requires cytoplasmic targeting; high cost
PMM2 Gene Therapy	Corrects defect by providing DNA encoding normal enzyme	Permanently corrects defect in cells that take up and integrate DNA	Variable accessibility to all cells, especially CNS, potential tumor formation, safety and regulatory hurdles, small number of patients
PMM2 Enzyme enhancement	Competitive inhibitor stabilizes mutated enzyme, but dissociates at lower pH to allow substrate binding	Small molecule may cross blood brain barrier; effective in some lysosomal storage disorders	pH does not change between site of synthesis and action; substrate competes with inhibitor
PMM2 activation	Non-competitive small molecule activates or stabilizes mutant enzyme increasing activity	Likely to cross BBB, may stabilize/activate in various ways	May not be useful for all mutant genotypes; will depend on whether specific mutation affects enzyme stability, Km, substrate binding or transcription
MPI Inhibition	Increase Man-6-P flux toward glycosylation by reducing MPI activity increasing PMM2:MPI	Small molecule likely to cross BBB activity; addition of mannose to the diet rescues even severe MPI deficiency	May not be effective in all tissues; likely to benefit those with higher residual activity.
AICAR	Activate glycogen phosphorylase, driving glucose→ fructose-6-P and increase Man-6-P	Orally active drug mimics effects of insulin and mobilizes glycogen	Studies only done on cell containing 0.5mM glucose. Application to physiological conditions unknown