Oculocutaneous albinism (OCA) is a group of genetic disorders characterized by hypopigmentation of the skin, hair, and eyes. Affected individuals experience reduced visual acuity and substantially increased skin cancer risk. There are four major types of OCA (OCA1–OCA4) that result from disruption in production of melanin from tyrosine. Current treatment options for individuals with OCA are limited to attempts to correct visual problems and counseling to promote use of sun protective measures. However, Onojafe et al., reporting in this issue of the JCI, provide hope for a new treatment approach for OCA, as they demonstrate that treating mice that model OCA-1b with nitisinone, which is FDA approved for treating hereditary tyrosinemia type 1, elevates plasma tyrosine levels, and increases eye and hair pigmentation.

Melanin synthesis and its disruption in OCA

Tyrosinase, a type I membrane protein, catalyzes the first and rate-limiting reaction during melanin synthesis. Unlike many membrane proteins that exit the ER rapidly, tyrosinase is retained for an unusually long period prior to transport to the Golgi, which suggests that it undergoes highly regulated/complex early processing. Maturation of tyrosinase begins during translation, when the peptide is inserted into the ER membrane by Sec61, then bound by the folding chaperone BiP (also known as HSPA5) (Figure 1). Additional chaperones — calnexin, then calreticulin — subsequently bind the peptide to facilitate glycosylation. The foldase ER protein 57 (ERp57) is also recruited to the complex to catalyze disulfide bond formation after signal sequence cleavage. The processed peptide is then subject to the ER quality control system that ensures that only correctly folded proteins exit the ER. Correctly folded tyrosinase is transported to the Golgi, where it undergoes additional glycosylation. Misfolded tyrosinase is retained in the ER, undergoing cycles of folding and unfolding, until it is eventually targeted for proteasomal degradation (10).

Many OCA1-causing TYR mutations result in a protein that fails to fold and is therefore unable to exit the ER (11). Tyrosinase is also retained in the ER, due to unknown mechanisms, in OCA2 (12) and OCA3 (13), whereas in OCA4, tyrosinase exits the ER but is not correctly transported to the melanosome (14). We have previously demonstrated that ER retention of tyrosinase activates the unfolded protein response (UPR) (15), a highly conserved cell signaling cascade that increases the capacity of a cell to process misfolded proteins but can lead to apoptosis if ER stress is sustained.

The precise functions of the remaining OCA-related proteins (excepting tyrosinase) remain unclear. We have proposed that OCA2 mediates glutathione transport into the ER to promote disulfide bond formation during tyrosinase maturation (16) and that TYRP1 stabilizes tyrosinase.
Costin et al. have proposed a role for the OCA4 protein, SLC45A2, in transport of tyrosinase from Golgi to melanosomes, given that mutations in the \textit{SLC45A2} locus result in trafficking of functional tyrosinase to the plasma membrane (14).

Improving tyrosinase folding and finding a treatment for OCA

Individuals with OCA1 can carry \textit{TYR} mutations that result in complete loss of tyrosinase activity (OCA-1A) or mutations that result in a protein with reduced activity at body temperature (OCA-1B). After postnatal molt, the hairs of mice carrying what is known as the Himalayan \textit{Tyr} mutation, which generates a mutant protein with reduced tyrosinase activity at body temperature, are darker in cooler, acral body areas (as in Siamese cats). In vitro studies have shown that folding of temperature-sensitive tyrosinase produced by human \textit{TYR} mutations can be improved by culturing cells at temperatures of approximately 31°C (11). In the early 1960s, long before the tyrosinase-encoding gene was identified, researchers found that hair bulbs from individuals with what we now call OCA-1A were incapable of making melanin when incubated at high concentrations of tyrosine, the substrate for tyrosinase, leading to OCA-1A (then termed tyrosinase-negative albinism). In contrast, incubation of hair bulbs from individuals with residual tyrosinase activity (which we now know include those with OCA-1B and OCA2 as well as other forms of OCA in which tyrosinase activity is diminished but not obliterated) in tyrosine causes pigment deposition, resulting in these forms being lumped together as tyrosinase-positive albinism. Topical application of tyrosine has not proven useful, presumably because of issues with permeation/penetration through skin, which acts as a barrier, to the melanocytes at the epidermal basal layer. Tyrosinase processing in cultured melanocytes can be greatly improved by tyrosine, DOPA (which is a cofactor for tyrosinase), and a variety of pH-altering agents, including monensin and bafilomycin (18).

Building on these observations, Onojafe et al. report in this issue of the \textit{JCI} on a potential treatment for OCA-1B (19). They were inspired by a side effect of sorts to make ingenious use of a drug, nitisinone, approved since 2002 to treat hereditary tyrosinemia type 1 (HT-1). Individuals with HT-1 have high levels of serum tyrosine as a result of deficiency of the enzyme fumarate-lactoacetate hydroxylase (FAH), the last enzyme in the tyrosine catabolism pathway. Somewhat counterintuitively, nitisinone raises serum tyrosine levels in individuals with HT-1 even further by blocking tyrosine catabolism one step upstream of FAH, acting as a competitive inhibitor of
Tyrosine catabolism. Five enzymes catalyze the cascade that leads to the degradation of tyrosine, and dysfunction of these enzymes leads to diseases of variable severity (reviewed in ref. 21). Tyrosine is converted to 4-hydroxyphenylpyruvate (HPP) by tyrosine aminotransferase (TAT). Mutations of this enzyme lead to HT-2. HPP is converted to homogentisic acid (HA) by hydroxyphenylpyruvate oxidase (HPD), the target of nitisinone. HPD mutations result in HT-3. HA is converted to 4-maleylacetoacetate (by homogentisate 1,2-dioxygenase [HGD], mutations of which cause alkaptonuria) then fumarylacetoacetate (by malelylacetoacetate isomerase [MAI]). These metabolites are toxic and accumulate, leading to the symptoms common in HT-1 (due to mutations of FAH), which converts fumaryl acetate to fumarate and acetooacetate.

4-hydroxyphenylpyruvate oxidase (Figure 2) and thus preventing accumulation of the toxic intermediates of tyrosine breakdown that cause the severe symptoms of HT-1. Inspired by this side effect (unlike typical drug repositioning), Onojafe and colleagues reasoned that administration of nitisinone to albino mice might result in elevated levels of tyrosine in tissues including RPE and epidermis (19), where it could then act (as in the hair bulb test) to rescue tyrosinase, presumably as a substrate chaperone. Substrate chaperones have been proposed for the treatment of genetic disorders including cystic fibrosis and Gaucher, Fabry, Tay-Sachs, and Sandhoff diseases.

Consistent with their hypothesis, Onojafe and colleagues found that administration of nitisinone at a dosage 2–4 times higher than used to treat HT-1 increased fur and eye pigmentation in OCA-1B mice (19). Treatment had no effect in OCA-1A mice, presumably because their misfolded tyrosinase is incapable of binding tyrosine. Based on these observations, one might expect that correction of tyrosinase misfolding in humans can be achieved, resulting in increased pigmentation of skin and hair. This would offer significant improvement for individuals with OCA, in that they will benefit from increased melanin protection from UV-induced skin cancers and amelioration of the significant psychosocial impact of hypopigmentation. Developmental defects due to lack of melanogenesis during embryogenesis are not expected to be correctable in an adult. Instead, correction would need to occur during the appropriate developmental window. To that end, Onojafe et al. gave nitisinone to pregnant OCA-1B mice and found that pigmentation was augmented in their offspring (19). While correction of developmental ocular defects remains to be proven, the augmented melanogenesis that they observed during gestation is highly encouraging. Since OCA-1B is a recessive disorder, parental carrier status is usually unknown until a first affected child is born. It is therefore unlikely that we will be able to treat women early enough during pregnancy to affect optic tract development, which is complete by 54–56 days. Nitisinone treatment may thus be an option in second pregnancies or in cases where family history suggests a high chance of an affected pregnancy.

Challenges for the future
The challenge remains to develop similar approaches for other OCAs. We have previously demonstrated that tyrosine can improve tyrosinase folding in OCA2 melanocytes (20), which suggests that nitisinone deserves exploration in OCA2 models. This must be tempered with the knowledge that a proportion of tyrosinase in OCA2 is trapped in vesicles and fails to reach the melanosome. Furthermore, excess tyrosine increases tyrosinase expression and may exacerbate ER stress as a result of increased retention of the enzyme (also a possible complication in the treatment of other forms of OCA, including OCA-1B). While our prior studies showed that UPR activation caused by tyrosinase misfolding does not lead to apoptosis (15), a further tipping of the balance may lead to melanocyte death, although this awaits experimental testing.

The work of Onojafe et al. (19) represents a substantial leap forward toward the possible treatment of all forms of OCA. However, it also reminds us that in addition to rational crystallographic design and screening of large compound libraries, informed reasoning can identify promising therapeutics.

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Viruses and human brain tumors: cytomegalovirus enters the fray

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Medulloblastoma is the most common malignant brain tumor in children. Overall survival rates have improved in recent years as a result of risk-stratified treatment regimens. However, medulloblastoma remains associated with substantial mortality, and survivors often experience debilitating neurological, endocrinological, and social sequelae as a result of treatment. Targeted and less toxic therapeutic strategies are therefore needed. In this issue of the JCI, Baryawno et al. report their findings that a large percentage of primary medulloblastomas and medulloblastoma cell lines are infected with human cytomegalovirus (HCMV) and suggest that targeting this virus could provide a new way to treat individuals with medulloblastoma.

Brain tumors account for 20% of all neoplasms in children and are the largest group of solid tumors that develop in childhood (1). Medulloblastoma is the most common malignant pediatric brain tumor, constituting 20%–25% of pediatric central nervous system neoplasms (2, 3). Its incidence is estimated at 2–6 cases per million children per year, with approximately 540 new cases diagnosed annually in the United States (2, 4). Medulloblastoma typically arises in the midline cerebellum, in the region of the mid- and inferior vermis. Current treatment includes surgery, craniospinal irradiation, and chemotherapy. Overall survival for all medulloblastoma patients is roughly 50%–60% in population-based studies (5). However, as we have gained a better understanding of clinical risk factors, the incorporation of patient stratification in larger, multi-institutional studies has resulted in improved survival. Currently, there are three major treatment strategies for medulloblastoma patients based on their clinical status. For patients younger than 3–5 years, treatments are aimed at maximizing survival while avoiding radiation (6, 7). Older patients are stratified by the extent of resection and metastatic status. Patients with metastatic disease or a less-than-optimal resection are classified as high-risk and are treated with high doses of craniospinal irradiation (36–39 Gy) and aggressive chemotherapy. Patients with totally, or near-totally, resected, non-disseminated disease are designated average-risk. This is the most prevalent group, and these patients are treated with a combination of lower-dose irradiation and chemotherapy. This clinical stratification has resulted in higher cure rates for all groups, with 80% of average-risk patients reaching progression-free survival at 5 years (8, 9). Despite the improved survival, there still remains a substantial amount of mortality associated with medulloblastoma, and survivors often experience neurological, endocrinological, and social sequelae as a result of treatment. Thus, many researchers are seeking to develop new, more targeted and less toxic therapeutic strategies. In this issue of the JCI, Baryawno et al. describe a potential novel therapeutic strategy for medulloblastoma.