Phenylketonuria (PKU) is an inborn error of metabolism caused by loss of function of the liver-expressed enzyme phenylalanine hydroxylase and is characterized by elevated systemic phenylalanine levels that are neurotoxic. Current therapies do not address the underlying genetic disease or restore the natural metabolic pathway resulting in the conversion of phenylalanine to tyrosine. A family of hepatotropic clade F adeno-associated viruses (AAVs) was isolated from human CD34+ hematopoietic stem cells (HSCs) and one (AAVHSC15) was utilized to deliver a vector to correct the phenylketonuria phenotype in Pahenu2 mice. The AAVHSC15 vector containing a codon-optimized form of the human phenylalanine hydroxylase cDNA was administered as a single intravenous dose to Pahenu2 mice maintained on a phenylalanine-containing normal chow diet. Optimization of the transgene resulted in a vector that produced a sustained reduction in serum phenylalanine and normalized tyrosine levels for the lifespan of Pahenu2 mice. Brain levels of phenylalanine and the downstream serotonin metabolite 5-hydroxyindoleacetic acid were restored. In addition, the coat color of treated mice darkened following treatment, indicating restoration of the phenylalanine metabolic pathway. Taken together, these data support the potential of an AAVHSC15-based gene therapy as an investigational therapeutic for phenylketonuria patients.

INTRODUCTION

Phenylketonuria (PKU) is a genetic disorder of liver metabolism that arises primarily from loss of function of the hepatic enzyme phenylalanine (Phe) hydroxylase (PAH), which is caused by mutations in the PAH gene. In the presence of a co-factor, tetrahydrobiopterin (BH4), PAH hydroxylates Phe to produce tyrosine (Tyr) and subsequently dopamine, adrenaline, and melanin. Lack of functional PAH results in hyperphenylalaninemia (HPA) after dietary intake of protein, causing elevated systemic Phe levels. HPA also causes a neurometabolic phenotype accompanied by cognitive impairment and behavior problems in patients on uncontrolled diets. PAH deficiency manifests as a continuum of HPA phenotypes from mild HPA (blood Phe from 120 to 360 μmol/L) to severe, termed as classical PKU, which is the most common form and clinically defined as blood Phe levels exceeding 1,200 μmol/L. If left untreated, classical PKU results in progressive, irreversible neurological impairment during infancy and early childhood. Classical PKU patients have neurocognitive issues even when they are on a controlled, Phe-restricted diet after adolescence. The disease shows multiple characteristics such as hypopigmentation in skin, hair, and eyes; eczema; growth retardation; and a mousy odor caused by accumulation of phenylacetate and phenyllactate. There are more than 950 known mutations in the PAH gene, with the vast majority being missense mutations. Although recessively inherited mutations in the PAH gene underlie most PKU cases, about 2% of the PKU patient population have a deficiency of the cofactor BH4 that can be corrected by oral supplementation of the missing cofactor. The cornerstone of current PKU therapy is restriction of Phe where dietary protein is replaced with a Phe-free liquid formula. The onerous diet makes it difficult to balance nutrients and eventually leads to lack of compliance, with a serious impact on neurocognitive health. Additionally, the reduction of dietary Phe intake does not restore Tyr levels and subsequent neurotransmitter synthesis. Introduction of functional PAH by gene therapy has the potential to correct the deficiency in the metabolic pathway by addressing the underlying genetic defect and provide a lasting effect following a single dose.

The most widely used murine model of PKU, the homozygous Pahenu2 mouse, has a point mutation in exon 7 (c.835T>C; p.F263S) resulting in approximately 1% of wild-type (WT) hepatic PAH activity. Blood Phe levels are increased 15- to 20-fold, and hypopigmentation is observed due to reduced melanin production, which manifests as lighter color fur. These features recapitulate classical PKU in humans, making this a valid model to evaluate the
therapeutic potential of gene therapy approaches for the treatment of PKU.

Early research has shown that liver-mediated gene therapy using an adenovirus vector transiently reverses HPA in Pahen2 mice, but efficacy was lost due to an immune response to the adenoviral capsid.9

Alternatively, adeno-associated viruses (AAVs) are non-pathogenic viruses of lower immunogenicity that can mediate long-term episomal transgene expression in non-dividing or slowly dividing cells.10 PAH gene therapy using AAVs has been attempted in mice by a number of groups using different routes of administration, AAV serotypes, or vector designs.5,11–13 More recently, Yagi et al.14 have reported that the intraperitoneal administration of a self-complementary AAV8 expressing murine PAH under the control of a liver-specific promoter to Pahen2 mice led to a reduction of serum Phe to normal or near-normal levels up to a year following administration. No published report has demonstrated sustained correction in a murine model of PKU following a single intravenous (i.v.) administration of human PAH at low doses of viral vectors.

In the studies described herein several vectors were screened and packaged in recombinant AAVHSC15, a clade F AAV isolated from human hematopoietic stem cells.15 Two of the vectors tested reduced blood Phe within a week of administration and darkened coat color in Pahen2 mice, a sign of restoration of melanin production. The administration of an AAVHSC15 containing a ubiquitous promoter (AAVHSC15-chicken β-actin [CBA]-PAH) rescued the high blood Phe phenotype within a week at a dose approximately a log lower than previously reported.5,11–13 But the reduction of serum Phe was transient. Further optimization including the use of a liver-specific promoter and a codon-optimized human PAH cDNA resulted in a vector (AAVHSC15-PAH) designated HMI-102. A single i.v. dose of HMI-102 restored the underlying loss of function of PAH activity caused by mutations in the PAH gene, achieved a sustained reduction of serum Phe, and normalized serum Tyr levels for the lifespan of Pahen2 mice at a 10-fold lower dose than AAVHSC15-CBA-PAH. In addition, brain levels of Phe and 5-hydroxyindoleacetic acid (5-HIAA) were restored. Taken together, these data support further development of AAVHSC15-PAH as an investigational gene therapy candidate for PKU.

RESULTS

Characterization of the Pahen2 Mouse

The Pahen2 mouse model has a homozygous missense mutation (F263S in exon 7) in the active site of PAH, resulting in 1% of WT activity.8 Mutations in exon 7 are common in PKU patients in a compound heterozygous context.16 Impaired Phe metabolism reduces production of Ty and eumelanin and is responsible for hypopigmentation in Pahen2 mice. Mutant Pahen2 males are smaller and weigh less than WT males (29.5 ± 4.5 g compared to 38.3 ± 2.2 g; p < 0.001, Figure S1A).

The murine model has several features consistent with the classical PKU phenotype, notably high blood Phe levels compared to the parent strain (1,458 ± 378 μM as opposed to 92.6 ± 37.6 μM; p < 0.0001) when maintained on a Phe-containing normal chow diet. Tyr levels show an inverse trend with lower levels in mutant mice (21.3 ± 5.7 μM as opposed to 83.5 ± 34.6 μM; p < 0.0001). The average lifespan of male Pahen2 mice was 235.5 days and that of female Pahen2 mice was 157 days on a normal chow diet (Figure S1B). This is consistent with literature reports indicating that the background strain (BTBR) for the Pahen2 mouse has a reduced lifespan compared to other mouse strains such as C57BL/6J.17

AAVHSC15-CBA-PAH Administration Corrected the PKU Phenotype In Vivo

The vector containing the human PAH cDNA flanked by the CBA promoter and the SV40 polyadenylation sequence was packaged into AAVHSC15. The expression cassette was designated AAVHSC15-CBA-PAH. This capsid was chosen based on its hepatic tropism previously observed.5,11 To assess whether AAVHSC15-CBA-PAH could restore Phe metabolism in vivo, groups of four male Pahen2 mice were injected with a single i.v. dose of AAVHSC15-CBA-PAH at various dose levels, that is, 1.6 × 1011, 5 × 1011, 1.6 × 1012, and 5 × 1012 vector genomes (VG)/mouse. Phe and Tyr levels were measured in blood collected immediately prior to dosing and then at weekly intervals for 4 weeks.

At all doses, injected mice showed a dose-dependent reduction in Phe levels and a concomitant increase in Tyr levels in Phe levels in the blood within 1 week of vector administration (Figure 1A). Phe levels rebounded and Tyr levels declined in mice treated with the two lower doses of AAVHSC15-CBA-PAH (Figures 1A and 1B). Similar findings were observed in female Pahen2 mice (data not shown).

A second study was performed using a group of six male mice dosed at 2 × 1012 VG/mouse with AAVHSC15-CBA-PAH to examine hepatic transduction, human PAH expression, and PAH enzyme activity. All mice treated with AAVHSC15-CBA-PAH showed darkened coat color (Figure 1C) and a reduction in serum Phe levels within 1 week of administration, which was sustained to the end of the 4-week study (Figure 1D). At study termination, VG levels were 1.2 × 106 VG/μg of DNA in the liver of treated mice (Figure 1E), indicating the high liver tropism of the AAVHSC15 capsid. The presence of human PAH VGs in the liver was associated with increased expression of human PAH mRNA (Figure 1F) and restoration of PAH enzyme activity to approximately 20% of the levels observed in WT mice (Figure 1G). These data are consistent with liver repopulation studies conducted in PKU-deficient mice by Hamman et al.25 using hepatocytes from WT or heterozygous Pah mice demonstrating that blood Phe levels within the normal range were observed when approximately 10% of the hepatocytes expressed PAH.

Vector Optimization to Increase the Efficacy of AAVHSC15-Based Gene Therapy

Durability of expression and phenotypic correction are key criteria for a successful gene therapy approach. To evaluate whether expression of PAH was sustained, groups of five male Pahen2 mice were dosed
with AAVHSC15-CBA-PAH at $1.6 \times 10^{12}$ and $5 \times 10^{12}$ VG/mouse, the two higher doses tested previously. Both doses showed a reduction of Phe to normal levels ($81.1 \pm 32.5$ μM; p < 0.0001) within 1 week (Figure 2A). The highest dose tested led to a sustained reduction in serum Phe for 12 weeks, the duration of the study (Figure 2A).

Injected mice showed a dose-dependent accumulation of VGs and mRNA levels (Figures S2A and S2B). The absence of sustained reduction of serum Phe at the low doses tested prompted modifications in the construct design to increase the efficacy of the vector. Several new constructs, including changes in promoters and codon optimization, were made (Figure 2B). Codon optimization was used to identify a cDNA sequence for the human PAH gene that would exhibit elevated and durable levels of human PAH mRNA in mouse liver.

Two methods were used to optimize transgene expression; variation in the number of cytosine-guanosine (CpG) dinucleotides in the PAH cDNA and the codon adaptation index (CAI). The CAI indicates usage of highly expressing codons; that is, higher CAI values indicate higher estimated protein expression. Nevertheless, the higher CpG dinucleotide count within a construct increases the potential likelihood of silencing mediated by CpG methylation,21 degradation of the VG mediated by TLR9 activation,22 or degradation of RNA mediated by the zinc finger antiviral protein (ZAP).23 Predicted CAI and CpG dinucleotide counts for each sequence obtained from the COOL program are shown in Figure 2C. An increase along the x axis indicates higher predicted protein expression. Constructs that had a lower CpG content but higher potential for durable expression compared to the unoptimized PAH sequence (Figure 2C) were selected for testing. Tested cDNAs were compared for relative identity and clustered together. The results are superimposed on a heatmap indicating relative identity between the algorithm-generated sequences (Figure 2D). Sequences selected for testing in vivo in Pahen(u2) mice are indicated by closed diamonds and compared to the WT human PAH sequence (closed squares).
hypothesis, a subset of vectors (namely constructs 5, 6, 7, and 8 containing 22, 10, 7, and 23 CpG counts, respectively) optimized for CpG counts in the human PAH cDNA sequence under the same promoter (CBA) were injected into the Pahenu2 mice (n = 4) and assayed for their ability to reduce serum Phe during 24 weeks post-injection (Figure 2E). No correlation between CpG content and efficacy was
observed. Construct 5 reduced and maintained serum Phe levels for the duration of the study while a loss of efficacy was observed with the unoptimized WT cDNA sequence vector (construct 8) despite having similar CpG counts of 22 and 23 CpG for constructs 5 and 8, respectively. Similarly, construct 7 (Figure 2E, closed circles) showed a rebound in blood Phe levels in spite of having three times lower CpG counts than construct 5. Construct 6 showed more sustained efficacy than construct 7 despite having more CpG counts of 10 and 7 CpG for constructs 6 and 7, respectively. Taken together, these findings indicate that the number of CpG counts in the PAH cDNA is not sufficient to explain the differences in efficacy observed in Pahenu2 mice. Similar hepatic VG levels were observed in all groups at the termination of the study (Figure 2F), indicating comparable hepatic transduction by all designs tested. The ratios of mRNA to VG were similar across different codon-optimized designs, further supporting the hypothesis that the CpG content within the human PAH cDNA does not affect the mRNA levels or efficacy of the tested constructs (Figure 2G).

We hypothesized that enhanced expression of the vector in the liver at lower doses may require a combination of a liver-specific promoter and a hepatotropic capsid. Thus, we developed chimeric promoters from combinations of the human apolipoprotein E/C-I hepatic control region enhancer (HCR1) and genes that express a large number of transcripts such as the alpha1 antitrypsin gene (SERPINA1) and the elongation factor 1 alpha (EF1a) to make three constructs: constructs 3, 4, and 9. A computationally derived chimeric promoter containing the hepatocyte-specific cis-regulatory module (HS-CRM8) was combined with the transthyretin (TTR) promoter to make construct 2.26 Construct 1 contained CBA as a ubiquitous promoter and was used as a reference. All designs were generated by using the same codon-optimized version of the human PAH containing 22 CpG counts, packaged in AAVHSC15, and tested at a dose of $2 \times 10^{12}$ VG/mouse. All vectors with liver-specific promoters normalized blood Phe levels in Pahenu2 mice within the first week of administration and sustained the phenotypic correction out to 48 weeks. In contrast, construct 1, containing the ubiquitous CBA promoter, showed a loss of efficacy over time (Figure 3A). In this study, constructs 2, 3, 4, and 9 containing liver-specific promoters were administered at a log-lower dose than the dose tested for construct 1. Liver concentrations of VGs did not differ significantly among all liver-specific promoter designs when compared to construct 1 (Figure 3B). Hepatic PAH mRNA levels were significantly higher (Figure 3C, $p < 0.01$) in Pahenu2 mice treated with AAVHSC15 vector containing liver-specific promoters (constructs 2, 3, 4, and 9) when compared to the AAVHSC15 vector design containing a ubiquitous CBA promoter (construct 1). Based on these findings, construct 9 was selected for further experiments.

Optimized Vector Efficiently Corrected PKU in Pahenu2 Mice

To assess the efficacy of the optimized vector AAVHSC15-PAH (construct 9), Pahenu2 male mice were dosed at $4 \times 10^{11}$ VG/mouse (n = 4 per group), a dose 10-fold lower than that used for the non-optimized AAVHSC15-CBA-PAH vector. Treated mice showed darkened coat color (data not shown) and a reduction in serum Phe within 1 week of administration (Figure 4A) that persisted until the end of the study at 12 weeks. All cohorts of mice sacrificed at earlier time points showed the same kinetics on Phe lowering. Vector
genomes in livers declined from week 1 to week 2 after the dose and remained stable from 2 to 12 weeks (Figure 4B).

While this study was underway, a long-term single i.v. dose efficacy study with AAVHSC15-PAH was initiated in male and female Pahenu2 mice fed ad libitum a normal chow diet for the duration of the study. Serum Phe and Tyr levels were assessed for 48 weeks. The 48-week time point was chosen for terminal sacrifice, as it is consistent with the lifespan of the model. A single-dose i.v. administration of AAVHSC15-PAH at 4 × 10^{11} VG/mouse in males (Figure 4C) and 9 × 10^{11} VG/mouse in females (Figure 4E) reduced blood Phe concentrations to normal levels (55 ± 24 μM in males and 54 ± 4 μM in females). Mean serum Phe concentrations remained below 120 μM and consistent with normal levels during the course of the 48-week study in both male (mean, 101 ± 38 μM) and female (mean, 94 ± 38 μM) mice. A corresponding increase from baseline in Tyr concentration was also observed (Figures 4D and 4F; p < 0.05) in male and female Pahenu2 mice, indicating restoration of the Phe/Tyr metabolic pathway. Consistent with the reduced levels in serum Phe, vector genomes were detected in liver tissues at 48 weeks (2.1 × 10^6 ± 4 × 10^4 VG/μg of DNA and 3.4 × 10^6 ± 7.9 × 10^5 VG/μg of DNA for males and females, respectively).

The effect of AAVHSC15-PAH in the metabolism of Phe and related metabolites in the brain was also examined. The serotonin metabolite 5-HIAA (Figure 5A), a pharmacodynamic marker for PKU patients,27 is significantly decreased in Pahenu2 mice when compared to WT parental mice (Figure 5B, p < 0.001), suggesting that the Pahenu2 mice reflect neurotransmitter deficits observed in PKU patients.28 Treatment with AAVHSC15-PAH was associated with increases in levels of 5-HIAA in brain, reaching levels comparable to those found in the parental strain at 4 weeks post-dose (Figure 5B). In addition, Phe levels in the brains of Pahenu2 mice were decreased to levels observed in WT mice 4 weeks following the administration of AAVHSC15-PAH (Figure 5C). Restauration of normal brain Phe levels and concomitant increases in brain 5-HIAA were also seen in female Pahenu2 mice (data not shown). These findings support the potential of a liver-mediated PAH gene therapy to address the elevated Phe levels observed in the brain tissue of PKU patients.29

AAVHSC15-PAH Restored Phe Levels in the PKU Mouse Model in a Dose-Dependent Manner

In order to determine the pharmacologically effective dose range of AAVHSC15-PAH, we conducted a dose-ranging efficacy study in...
Pahenu2 mice. In this study, AAVHSC15-PAH was administered by a single i.v. injection via the tail vein at one of five doses over approximately a 1.5-log range. Mice were sacrificed 4 weeks after dosing. A dose-responsive decrease in blood Phe was observed over the course of the study, in both male and female mice. As shown in Figures 6A and 6B, after treatment with 2 × 10¹¹ VG/mouse AAVHSC15-PAH, serum Phe levels in male mice were 127 ± 106 μM on day 8, as were those in female mice treated with 4 × 10¹¹ VG/mouse (150 ± 144 μM). The group mean Phe concentrations for male mice treated with 4 × 10¹¹, 8 × 10¹¹, and 1.6 × 10¹² VG/mouse and female mice treated with 8 × 10¹¹ and 1.6 × 10¹² VG/mouse were below 120 μM at all intervals and maintained for the duration of the study. A corresponding increase in serum Tyr was observed relative to baseline and vehicle control in all mice at all doses evaluated (Figures 6C and 6D). As shown in Figure 6B, female Pahenu2 mice had an approximately 34% higher level of serum Phe at baseline than that in males (1,400 ± 95 μM and 1,048 ± 108 μM, respectively. At lower doses (1 × 10¹¹ and 2 × 10¹¹ VG/mouse), female mice showed higher levels of blood Phe following treatment with AAVHSC15-PAH when compared to male Pahenu2 mice.

Analysis of liver samples was also performed to assess the level of vector genomes 28 days following administration of AAVHSC15-PAH (Figures 7A and 7B). All mice treated with AAVHSC15-PAH had detectable levels of vector genomes. The levels increased in both male and female mice with each increasing dose level. Notable differences between male and female mice in the levels of AAVHSC15-PAH vector genomes were observed. The levels of vector genomes in the male mice were 1.7- to 3.2-fold higher when compared to female mice, consistent with the sex difference in AAV transduction previously reported in mice.30 As with the vector genome results, the levels of expression increased with each dose level, and differences were noted between male and female mice (1.5- to 2.8-fold higher in males compared to females; data not shown).

Liver samples were also collected for PAH enzyme activity, and results are shown in Figures 7C and 7D. A dose-dependent increase in PAH activity was observed. At the highest dose tested (1.6 × 10¹² VG/mouse), PAH activity was 98% ± 67% and 37% ± 11% of WT levels in males and females, respectively. Mean PAH activity was above 10% of WT levels at doses up to and above 2 × 10¹¹ VG/mouse (16% ± 8%) in males and 4 × 10¹¹ VG/mouse (12% ± 5%) in females 28 days following AAVHSC15-PAH administration and at sufficient levels to restore serum Phe to levels below 120 μM.
**DISCUSSION**

PKU is an inborn error of Phe metabolism primarily caused by mutations in the *PAH* gene. At present, a strict low-Phe diet is the standard of care for most PKU patients. Compliance, however, is poor and such diets are also low in long-chain polyunsaturated fatty acids (LCPUFAs) and docosahexaenoic acid (DHA), both of which are important in neurological development. Dietary Phe restriction does not address the underlying genetic deficiency of *PAH*, leading to the deficiency of Tyr and related downstream metabolites observed in patients with PKU. PKU patients have persistent white matter lesions, subtle neurologic changes, mood and behavior disturbances, and quality of life concerns even when their Phe intake is well controlled.

Initial enzyme replacement studies with functional PAH did not prove effective for PKU. Subcutaneous administration of an alternative enzyme that deaminates Phe to ammonia and *trans*-cinnamic acid, PAL, reduces Phe within the blood but does not restore the affected biological pathway, including the synthesis of Tyr, a well-known neurotransmitter precursor. Current treatment strategies also include dietary restriction of Phe with or without augmentation of residual hepatic PAH activity by delivery of the BH₄ cofactor necessary to metabolize Phe.

In the present study, we showed that gene therapy has the potential to restore the underlying deficient biochemical pathway. Providing copies of the human *PAH* gene to produce functional enzyme would enable metabolism of Phe to its normal by-products and decrease its toxic accretion.

AAVs are potentially suitable gene therapy delivery vehicles given their tissue tropism and limited immunogenicity. Previous studies indicate that the presence of anti-AAV neutralizing antibodies, even at low titers, can block the transduction and compromise the effectiveness of AAV-based therapy in both preclinical and clinical studies. Two of the AAVHSCs (AAVHSC15 and AAVHSC17) were tested against a panel of human sera samples and were shown to have a low seroprevalence of neutralizing antibodies. Successful correction of PKU in the established PKU mouse model using recombinant AAVs (rAAVs) carrying *PAH* has been shown by several laboratories, although the therapy was less effective in female mice. Similar sex-specific differences were observed in our studies consistent with literature reports. Interestingly, enzyme substitution therapy using pegylated PAL, which acts by a different mechanism than *PAH* gene replacement also showed lower efficacy in female Pah<sup>em2</sup> mice. Sex-based differences in transduction by AAVHSCs in non-human primates or even sex-based differences in basal Phe levels of human patients have not been reported, suggesting that the sex-based difference appears to be a characteristic specific to the PKU mouse model. Other than the sex-specific response to therapy, Pah<sup>em2</sup> mice recapitulate several symptoms of human patients with PKU and represent a well-established model to test the efficacy of investigational therapies for PKU.

**Figure 6. Dose Range Efficacy Study in Male and Female Pah<sup>em2</sup> Mice following Administration of AAVHSC15-PAH**

(A–D) Serum Phe (A and B) and Tyr levels in male (A and C) and female (B and D) Pah<sup>em2</sup> mice following single tail vein injection of a range of doses of AAVHSC15-PAH (n = 5 mice/group) as described in Materials and Methods. Data in (A–D) are represented as mean ± SD.
Effectiveness of a gene therapy vector is a function of delivery to the target tissue, efficacy of the expression cassette, and durability of effect. We pursued a multipronged approach to develop an AAV-based therapy for PKU by combining a hepatotropic capsid with the human PAH gene and liver-restricted expression. This combination allowed for lowering the vector dose while achieving long-term and sustained efficacy in the murine model.

Construct 1, the vector containing the ubiquitous promoter (AAVHSC15-CBA-PAH), restored clinically relevant levels of PAH enzyme activity that were accompanied by the normalized blood Phe levels in Pahenu2 mice within the first week of administration. Normalization of blood Phe levels was transient despite the presence of high levels of vector genomes in the liver, suggesting that the loss of efficacy was not due to poor transduction but rather low levels of PAH mRNA and, consequently, insufficient levels of PAH activity. Optimization of CpG dinucleotide repeats within the PAH sequence to address potential immune-related loss of transduced cells and efficacy was unsuccessful in restoring long-term efficacy.

As demonstrated in studies on mouse models for Pompe disease and GSD1b, promoter specificity can determine the efficacy of a given vector design by restricting the transgene expression to the intended tissue, avoiding cytotoxic T lymphocyte (CTL)-mediated toxicity and resulting in the clearance of the transduced cells and loss in therapeutic effects. We tested several liver-specific promoters on a side-by-side comparison between AAVHSC15 vectors containing ubiquitous or liver-specific promoters and demonstrated a significant difference in hepatic mRNA levels consistent with the hypothesis that promoter specificity improves the efficacy of the therapeutic cassette.

The PKU mice developed by Shedlovsky et al. (Pahenu2/BTBR) survive for approximately 60 weeks in our laboratory; thus, all long-term studies were carried out until 48 weeks post-injection since mice were injected as adults at 7 weeks of age. In the studies described herein, several vectors demonstrated potential in delivering the human PAH gene at levels sufficient to restore PAH enzyme activity for the lifespan of the Pahenu2 mice even in the presence of a consistent and normal influx of Phe from the diet.

In conclusion, this work describes the identification of an investigational gene therapy for PKU designated HMI-102 based on a novel optimized AAVHSC15 that was isolated from human CD34+ hematopoietic stem cells and injected i.v. to deliver a corrected copy of human PAH. Taken together, the findings from these studies demonstrate that liver-restricted expression of a codon-optimized version of human PAH resulted in the persistence of VGs in the liver accompanied by durable hepatic expression of human PAH mRNA and PAH enzyme activity levels sufficient to restore the natural metabolic pathway throughout the lifespan of the Pahenu2 mouse model, as shown by the normalization of serum and brain Phe, Tyr, and 5-HIAA levels. Collectively, these data support the potential of a gene therapy approach for PKU that could reduce Phe to normal levels and substantially restore the natural biological pathway.
MATERIALS AND METHODS

Vector Design

ssAAV (single-stranded AAV)/CAG-hPAH (human PAH) plasmid was used to build the vector construct 1 with the CBA promoter: CMV immediate early enhancer and codon-optimized human PAH cDNA, followed by the SV40 early polyadenylation signal. Construct 1 was used to generate AAVHSC15-CBA-hPAH by cloning into a plasmid containing AAV2 inverted terminal repeats (ITRs). All constructs were fully sequenced and verified. The self-complementary AAV vector AAVHSC15-PAH (construct 9) was developed with a 2,093-bp synthetic double-stranded DNA (dsDNA) gBlock fragment (IDT) that was subsequently PCR amplified and cloned into an AAV2 ITR-containing plasmid by Gibson Assembly. It included a hybrid liver-specific promoter, with a codon-optimized form of the PAH gene (NCBI GenBank: NM_000277.2), and an SV40 late polyadenylation signal. The promoter contained a hepatic control region (HCR1), the SERPINA1 (AAT) promoter, and 5’ UTR. The total construct size was 2,356 bp. Construct 2 contained the HS-CRM8-TTR promoter as previously described (PMID: 30700722). Construct 3 contained HCR1, the SERPINA1 promoter, and modified 5’ UTR from the construct described in Picancão-Castro et al. (2018). Construct 4 contained the HCR1 enhancer combined with human EF1α promoter with a 104-bp truncation, derived from the eEF1α-P-mCherry-HPos plasmid. All constructs had the same PAH cDNA sequence. Promoters for constructs 2 and 3 were synthesized as 424- and 1,191-bp dsDNA gBlocks, respectively. The promoter for construct 4 was synthesized as an 1,832-bp fragment (Genscript), which was subsequently PCR amplified and cloned into the same hPAH/SV40 pA plasmid as described above by Gibson Assembly. All constructs were fully sequenced and verified. Vector genomes were packaged into AAVHSC15, a clade F AAV isolated from human CD34+ human peripheral blood stem cells.19

The WT PAH cDNA was modified based on individual codon usage, codon context, codon adaption index, and low CpG count relative to the human genome by the COOL codon optimization tool. COOL generated 56 codon-optimized cDNAs. A subset of five cDNAs was selected for screening. The percentage identity was determined in a pairwise manner for cDNA sequences by the Geneious program. cDNA heatmaps and clustering were performed with the heatmap package in R. Additional codon optimization was performed with Genscript’s OptimumGene tool (US patent no. 8,326,547).

Production of AAV Vectors

AAVHSC15 vector stocks were produced by transient triple transfection of HEK293 cells and CaCl2 gradient sedimentation as previously described, and formulated in a buffer consisting of 5% sorbitol and 0.001% pluronic F-68 in PBS (pH 7.6). Vector preparations were titrated by droplet digital PCR (ddPCR), purity of vectors was assessed by 4%–12% sodium dodecyl sulfate (SDS)-acylamide gel electrophoresis and silver staining (Invitrogen, Carlsbad, CA) and morphological integrity of virions was assessed by transmission electron microscopy of negative-stained rAAV virions at Electron Microscopy Core (UMass Medical School, Worcester, MA). Vector genome titers were determined by ddPCR analysis using the following SV40 primers and probe—forward, 5’-CCACACATGATAAGATACATTGAGTT-3’; reverse, 5’-AG CAATACGCATCAAATTTCAAAA-3’; probe, 5’-ACGATTTTTT TCACGTGATTCTGTTGTTGTGTC-3’—or the following set for the human PAH sequences: forward, 5’-GTATCATTGTTGCG CGAGTCTTT-3’; reverse, 5’-GTATATTCCAGCACACTCCTGATCC-3’; probe, 5’-ACGATGCGAAAGGAGGTTGAGAA-3’.

Animal Procedures

Colonies of PKU mice, BTBR-Pahnu2, and their parental strain, BTBR (obtained from Jackson Laboratory, Bar Harbor, ME, USA), were maintained in the animal facility at Homology Medicines, Inc. under standard laboratory conditions (temperature, 20°C–24°C; relative humidity, 50%–60%; 12-h light/12-h dark cycle). Animals were supplied with standard chow (PicoLab 5058, providing approximately 25% of energy as protein and containing 0.97% Phe) and sterile water ad libitum. For experiments, 7-week-old adult males were injected with AAV vectors into tail veins using a 1-mL syringe with a 28-gauge needle. For Phe monitoring, blood was collected from the facial vein, and serum or plasma was extracted and evaluated for Phe and Tyr levels. All procedures were approved by the Institutional Animal Care and Use Committee at Homology Medicines, Inc. No changes in survival were observed in mice treated with AAVHSC15 when compared to control untreated mice.

Determination of Phe and Tyr by RapidFire-MS

Blood samples were taken as described above and analyzed using a RapidFire 300/Sciex API 4000 QqQ mass spectrometer (MS). Standard curve and internal standards were 2 mmol/L 15C6-Phe and 0.2 mmol/L 13C6-Tyr in water. The RapidFire-MS protocol, as developed, provided a linear signal up to 20 μmol/L with a lower limit of quantitation approximately 10 nmol/L. Prior to analysis, samples were diluted 250-fold in high-pressure liquid chromatography (HPLC) water. Following analysis on the RapidFire-MS, the calculated peak areas of the eight 13C6-Tyr and 13C6-Phe concentrations were plotted versus the internal standard concentrations. The linear fit for the eight data points (including the blank where only background signal should be detected) had an R² correlation value of 0.97 or higher. The average peak area of the eight injections for each sample for both Phe and Tyr fell within the linear range of the internal standard injections and had a CV of 15% or better, or was retested. The stable isotope standard curves for 15C6-Tyr and 13C6-Phe were used to calculate the unknown concentrations of Phe and Tyr in each specimen.

Determination of Vector Genome Copy Numbers in Liver Samples

rAAV genomes were quantified in 1-μg samples of genomic DNA extracted from mouse livers using the QIAamp DNA mini kit for quantitative detection of vector genome copies by TaqMan probes.
(Applied Biosystems, Foster City, CA, USA) with a single-copy endogenous glyceraldehyde 3-phosphate dehydrogenase (GAPDH) gene as the diploid cell number reference. A TaqMan real-time PCR kit was run with primer sets that amplified regions of the hPAH or SV40. Briefly, isolated DNA was tested for purity and concentration by spectrophotometry. PCR reactions were performed in 20 μL of final volume using the 2× Universal TaqMan master mix (Applied Biosystems, Foster City, CA, USA) supplemented with 100 nM each of sense and antisense primer, and 2 μL of diluted template DNA (either plasmid standard or extracted sample DNA according to the manufacturer’s instructions). All samples and controls were run in three replicates of 20 μL of reactions in alternate rows of a 384-well optical plate. The PCR profile contained an initial denaturation step at 95°C for 10 min followed by 40 cycles of denaturation at 95°C for 15 s and annealing or extension at 60°C for 1 min. Data analysis was performed using Applied Biosystems QuantiStudio software v1.3. The sensitivity of the assay was 10 copies/150 ng of cellular DNA. Samples were evaluated for vector biodistribution using the following primers and probes: SV40, forward, 5'-AGCAATAGCATCACAAATTTCACAA-3', reverse, 5'-AGCATTTTTTTTCACCTGATTCATGTTGTGTTGTGT-3'; human PAH, forward, 5'-GTACTATGTGGCCGAGTCTTT-3', reverse, 5'-GACATTTTTTTTCACCTGATTCATGTTGTGTTGT-3'; human hypoxanthine guanine phosphoribosyl transferase (HPRT), were determined by Applied Biosystems QuantStudio software v1.3. The sensitivity of the assay was 10 copies/150 ng of cellular DNA. RNA concentrations were determined by NanoDrop and subsequently treated with DNase. Samples with a spectrophotometric ratio of absorbance at 260 and 280 nm between 1.8 and 2.0 were considered acceptable for use, diluted to 100 ng/μL, and reverse transcribed to cDNA.

mRNA levels were measured by qRT-PCR using an Applied Biosystems QuantiStudio 6 Flex system (Thermo Fisher Scientific). Briefly, mRNA levels for PAH and the reference gene, hypoxanthine guanine phosphoribosyl transferase (HPRT), were determined by qPCR using specific sets of primers and probes as follows: human PAH, forward, 5'-GTACTATGTGGCCGAGTCTTT-3', reverse, 5'-GTATTATCCAGCACCTGATCC-3', probe, 5'-ACGATGCCAAGGAGAACTG-3'; human PAH, forward, 5'-GTACTATGTGGCCGAGTCTTT-3', reverse, 5'-GTATTATCCAGCACCTGATCC-3', probe, 5'-ACGATGCCAAGGAGAACTG-3'. Data were collected in triplicate as Ct (threshold cycle) values, and the ΔΔCt method was applied to calculate PAH mRNA levels relative to the levels of the reference gene, HPRT. Results were expressed as ratios of the gene-specific signal to the internal reference signal, yielding a corrected relative value for the gene-specific product in each sample. Samples with Ct values higher than 32 or equivalent to water controls Ct were considered outside the optimal ranges.

**Determination of mRNA levels by qRT-PCR**

A QIAGEN RNeasy Plus mini kit was used to extract mRNA from mouse livers following the manufacturer’s protocol. RNA concentrations were determined by NanoDrop and subsequently treated with DNase. Samples with a spectrophotometric ratio of absorbance at 260 and 280 nm between 1.8 and 2.0 were considered acceptable for use, diluted to 100 ng/μL, and reverse transcribed to cDNA.

**Determination of PAH Activity in Liver Samples**

The liver PAH enzyme activity analysis was conducted using a research-grade non-good laboratory practice (GLP) method developed at PureHoney Technologies (Billericia, MA, USA). Briefly, approximately 50 mg of liver tissue was homogenized using a Precellys homogenizer. Protein concentration was assessed (Pierce BCA [bicinchoninic acid] protein assay kit) and the homogenate diluted with phosphate-buffered saline to a 1 mg/mL concentration for use in the PAH enzyme assay. The 2× substrate solution (final assay concentration of 200 mM potassium phosphate/150 mM potassium chloride, 400 μg/mL catalase, 300 mM ammonium iron (II) sulfate, 2 mM dithiothreitol, 200 mM [13C6]-Phe, and 200 μM tetrahydrobiopterin in aqueous solution) and 2× enzyme solution (final assay concentration of 200 mM potassium phosphate/150 mM potassium chloride, 400 μg/mL catalase, 300 mM ammonium iron (II) sulfate, 2 mM dithiothreitol, and 400 μg/mL liver extract) were added to a 96-well microplate and incubated at room temperature for a specified time period (during this time period, any PAH present in the liver sample converts Phe to Tyr through oxidation). The enzyme reaction was quenched by the addition of 10 μL of 10% formic acid containing 5 μM [13C9]-Tyr. Samples were then analyzed by the RapidFire liquid chromatography-tandem mass spectrometry (LC-MS/MS) method.

The slope was determined for each individual sample within the linear range, either from 0 to 10 min or from 0 to 15 min. The background for the assay was determined by taking the average of the vehicle control and subtracted from the slope of each individual animal as well as from the control samples.

**Determination of 5-HIAA**

Mouse brains were homogenized in PBS and vortexed for 15 min. Lysates were spun down at maximum speed for 5 min to clarify the debris, and the supernatant was extracted with a 50:50 mixture of methanol and acetonitrile. Extract was clarified by vortexing at high speed and centrifugation at 16,000 rpm for 15 min. Supernatant was transferred to an amber liquid chromatography vial and diluted with deionized water to prepare for LC-MS/MS injection. 5-HIAA-d5 was used as an internal standard.

**SUPPLEMENTAL INFORMATION**

Supplemental Information can be found online at [https://doi.org/10.1016/j.omtm.2020.03.009](https://doi.org/10.1016/j.omtm.2020.03.009).

**AUTHOR CONTRIBUTIONS**

S.S.A., A.S., J.L.E., and O.L.F. designed research; S.S.A., H.R., M.W., D.F., A.S., S.N.D., N.A., and J.L. performed research; S.S.A., M.L., and L.A.-S. analyzed data; D.L. and T.W. had scientific input; and S.S.A. and O.L.F. wrote the manuscript.

**CONFLICTS OF INTEREST**

All authors are employees or former employees of Homology Medicines, Inc. and hold equity in the company.
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