Safety and efficacy of an engineered hepatotropic AAV gene therapy for ornithine transcarbamylase deficiency in cynomolgus monkeys

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INTRODUCTION

X-linked inherited ornithine transcarbamylase deficiency (OTCD) is the most common disorder affecting the liver-based urea cycle, a pathway enabling detoxification of nitrogen waste and endogenous arginine biosynthesis. Patients develop acute hyperammonemia leading to neurological sequelae or death despite the best-accepted therapy based on ammonia scavengers and protein-restricted diet. Liver transplantation is curative but associated with procedure-related complications and lifelong immunosuppression. Adeno-associated viral (AAV) vectors have demonstrated safety and clinical benefits in a rapidly growing number of clinical trials for inherited metabolic liver diseases. Engineered AAV capsids have shown promising enhanced liver tropism. Here, we conducted a good-laboratory practice-compliant investigational new drug-enabling study to assess the safety of intravenous liver-tropic AAVLK03 gene transfer of a human codon-optimized OTC gene. Juvenile cynomolgus monkeys received vehicle and a low and high dose of vector (2 × 10^{12} and 2 × 10^{13} vector genome (vg)/kg, respectively) and were monitored for 26 weeks for in-life safety with sequential liver biopsies at 1 and 13 weeks post-vector administration. Upon completion of monitoring, animals were euthanized to study vector biodistribution, immune responses, and histopathology. The product was well tolerated with no adverse clinical events, predominant hepatic biodistribution, and sustained supra-physiological OTC overexpression. This study supports the clinical deployment of intravenous AAVLK03 for severe OTCD.

X-linked inherited ornithine transcarbamylase deficiency (OTCD) has been recognized for many years as an appealing disease candidate for gene therapy. A historical clinical trial with an adeno-viral vector encoding the OTC gene caused the death of one late-onset OTCD patient receiving a high-dose (HD) vector due to a severe exacerbation of liver disease. Here, we have shown that AAVLK03 is associated with significant morbidity in severe OTCD.
immune response syndrome triggered by innate immunity against the vector capsid.\textsuperscript{5,6} This fatal outcome somewhat compromised the development of gene therapy for diseases such as OTCD until non-pathogenic adeno-associated viruses (AAVs) were successfully delivered for other liver monogenic disorders such as hemophilia A and B.\textsuperscript{7–9} AAV vectors have demonstrated long-lasting transgene expression and disease-modifying efficacy in hemophilia B.\textsuperscript{10} The safety profile is satisfactory, predominantly triggering a CD8 T cell-mediated immune response causing asymptomatic transaminitis well controlled by an oral course of corticosteroids.\textsuperscript{10} However, very HDs (\(>5 \times 10^{13}\) vector genome [vg]/kg) have induced a severe immune response syndrome with complement activation in large animal models and in clinical trials with fatal outcomes highlighting AAV dose-limited toxicity.\textsuperscript{11–13}

Using a wild-type AAV8 capsid, a phase I/II trial recruiting late-onset adult OTCD patients sponsored by Ultragenyx has shown partial or complete response in 6 out of 9 patients at doses of \(2 \times 10^{12}\) to \(2 \times 10^{11}\) vg/kg.\textsuperscript{14} To improve hepatocyte transduction, engineered AAV capsids with enhanced liver tropism have been generated.\textsuperscript{15,16} AAVLK03 is an engineered capsid that shares 97.7% homology of the cap sequence and 98.9% homology of the amino acid sequence with the wild-type hepatotropic AAV serotype AAV3B.\textsuperscript{15} AAVLK03 transduces human hepatocytes 1 log better than AAV8 in chimera mouse-human livers and is more resistant to neutralizing antibodies (Nabs) compared to wild-type capsids.\textsuperscript{15,17,18} Interestingly, the prevalence of anti-AAVLK03 Nabs is particularly low in pediatric populations.\textsuperscript{19} This capsid is currently in clinical development for adults with hemophilia A sponsored by Spark Therapeutics (ClinicalTrials.gov: NCT03003533).\textsuperscript{20} Moreover, one patient was injected with an AAVLK03 vector containing the MMUT gene in a phase I/II open-label clinical trial for pediatric patients with methylmalonic academia (SUNRISE study; ClinicalTrials.gov: NCT04581785).\textsuperscript{21}

Treating pediatric OTCD patients is clinically relevant, as these patients have a more severe phenotype and are at risk of developing hyperammonemic decompensation with fatal outcome or causing severe lifelong neurological sequelae. AAV vectors deliver a transgene cassette, which persists mainly as a non-integrating nuclear episome that is not effectively transmitted during mitosis. The liver growth in young patients represents a challenge, as transgene loss and dilution may limit the long-term efficacy of AAV gene therapy. To advance AAV gene therapy for pediatric OTCD patients, we have designed an AAVLK03 vector encoding the human codon-optimized OTC gene. We performed a good laboratory practice-compliant toxicology study using juvenile cynomolgus macaques to assess vector safety, biodistribution, and immune responses.

RESULTS

Study design

Three gender-matched groups of six cynomolgus macaques, aged 18 months, seronegative for Nabs against AAVLK03 received a single peripheral intravenous injection of vehicle, low dose (LD; \(2 \times 10^{12}\) vg/kg) or HD (\(2 \times 10^{11}\) vg/kg) of AAVLK03.human OTC (hOTC) vector encoding the hOTC transgene (Figure 1A). Over a 26-week monitoring period, biofluids were collected at regular intervals, and needle liver biopsies were performed at 1 and 13 weeks post-administration. The animals were sacrificed after 26 weeks, and organs were collected for analysis. No immunosuppression was given (Figure 1B).

For dose selection, we considered the efficacy outcome of a former clinical trial targeting adult hemophilia B patients with a single intravenous injection at \(2 \times 10^{12}\) vg/kg with an AAV8-derived vector.\textsuperscript{10} For safety reason, no liver biopsy was performed in this trial. We assumed that a mild increase of plasma factor IX from <1% to 6% should correspond to a low percentage (likely <5%) of hepatocyte transduction. This conclusion is in accordance with the percentage of human hepatocytes transduced by AAV8 vectors in Fah\(^{-/-}\)/Rag2\(^{-/-}\)/Il2rg\(^{-/-}\) (FRG) mice.\textsuperscript{15} As the AAV-LK03 capsid is expected to have at least 10 times better transduction efficacy in human hepatocytes compared to AAV8 based on \textit{in vivo} studies in FRG mice,\textsuperscript{15,22} a LD of \(6 \times 10^{11}\) vg/kg AAV-LK03 vector would be equivalent to \(6 \times 10^{12}\) vg/kg AAV8 vector and transduce around 10%–15% hepatocytes, which is deemed sufficient to observe already an improvement of the phenotype according to experiments in the sparse fur/abnormal skin and hair (Sp\(^{m1y}\)) mice and estimated enzymatic threshold expected to normalize the phenotype in urea-cycle defects in humans.\textsuperscript{23}

AAV vector quality control (QC) testing

The AAVLK03.hOTC appeared in a clear, colorless solution with no visible particles by visual inspection. A bioburden test by direct plating did not identify any bacterial growth (<2 colony formation units/mL), and bacterial endotoxin assessed by Limulus amebocyte lysate (LAL) kinetic chromogenic method was not detected (<0.05 endotoxin units (EU)/mL). The empty-to-full AAV capsid ratio was assessed by optical density by A260/A280 ratio spectrophotometry and found to be at 1.5, in line with acceptable criterion for the manufacturer (>1.3). The SDS-PAGE/silver stain test assessing purity was comparable to the reference. The titer measured by qPCR was \(2.66 \times 10^{13}\) vg/mL.

In-life safety parameters

All animals survived until their scheduled necropsy at 26 weeks. The vector-treated animals had no treatment-related clinical changes identified on general observations, temperature, and heart-rate measurements. Food consumption was adjusted according to body weight, and all animals maintained their weight gain during the study. Mild bruising was noted at the injection site lasting 4 days or less after vector administration. Ophthalmological evaluations were satisfactory. Electrocardiogram was unremarkable in all animals.

Laboratory parameters

Clinical laboratory parameters, including hematology, coagulation, and biochemistry, did not differ between vehicle- and vector-treated animals at 4, 13, and 26 weeks post-administration (Tables S1–S4). Urine analysis revealed significantly higher relative urine gravity in
HD compared to LD, the control group, and pretest values. This difference was considered as not clinically significant.

**Histopathology**
An extensive tissue collection listed in Table 1 was performed at necropsy 26 weeks after treatment. All tissue samples had a normal macroscopic appearance. There was no difference in organ weight between AAVLK03.hOTC-treated and control animals. Histologic evaluation did not reveal any noticeable pathological lesions in the AAVLK03.hOTC-treated or control monkeys. Specifically, no ischemic, apoptotic, inflammatory, or mitotic changes were seen. The microscopic examination of necropsy tissues also did not reveal treatment-related changes.

**Immune response**

**Humoral immunity**
There were no detectable anti-AAVLK03 Nabs in any vector-treated animals before administration. Control animals were either seronegative or seropositive with a low antibody titer (1/5 dilution). No relevant changes in the Nab titers against AAVLK03 were recorded in control animals during the 26-week observation period. Overall, all AAVLK03.hOTC-dosed animals showed a peak of Nab titers in plasma 2 weeks after vector administration, except for female 20 and male 7 from the HD group, for which the peak occurred at weeks 4 and 13 after the administration, respectively.

In the HD group, the Nab peak plateaued from weeks 2 to 13 before decreasing by week 26. In the LD group, the peak was evident at 2 weeks, followed by a fall in the titers from 4 weeks onward, which was quicker than for the animals in the HD group (Figure 2A). 3/6 animals in the LD group were seronegative at 26 weeks (Figure 2A). Different animals in both LD and HD cohorts mounted variable intensity of humoral immune response (Table S5). There were no marked gender differences in humoral immune responses between the groups at the end of the study (Figures 2B and 2C). In the LD group, females showed a lower production of Nabs compared to males (Figure 2C). The assay at weeks 13 and 26 did not work for an unknown reason for animal 18F and could not be repeated.

**Cellular immunity**
The cellular immune response was assessed prior to vector administration and at weeks 2, 4, 13, and 26 in peripheral blood mononuclear cells (PBMCs). The PBMCs were incubated with 15-mer overlapping peptide libraries for viral protein VP1 of the AAVLK03 capsid and the hOTC protein, and the percentage of helper (CD4+) and cytotoxic (CD8+) T cells secreting interleukin 2 (IL-2) and interferon gamma (IFN-γ) was detected by flow cytometry. The results showed no cellular response against the viral protein VP1 of the AAVLK03 capsid, suggesting the absence of capsid-specific T cells (Figures S1A–S1C). There was also no response against the hOTC protein. The percentage of T cell-producing cytokines was negligible and very similar to that observed in PBMCs treated with the control Roswell Park Memorial Institute (RPMI) cell culture medium buffer. All PBMC samples at all time points demonstrated capacity to respond to an appropriate stimulus by secreting pro-inflammatory cytokines.
after incubation with leukocyte activation cocktail (LAC; a polyclonal stimulator) (Figures S1A–S1C).

All samples in all groups yielded results below or near the assay range for IL-1β, IL-2, IL-4, IL-5, IL-6, tumor necrosis factor α (TNF-α), granulocyte colony-stimulating factor (G-CSF), and IFN-γ concentrations with the exception of IL-8. IL-8 concentrations were seen to be variable across all time points and groups, including controls. IL-8 concentrations were lower with the lower doses of AAVLK03.hOTC compared to the HD cohort at all time points (Figure 3A). In saliva, the AAVLK03.hOTC vector was quantifiable from day 1 up to day 15 from the LD cohort and from day 1 to week 26 in samples from the HD cohort, with a peak at day 1 (Figure 3A). In the LD group, vector DNA was quantifiable in the plasma from day 1 up to day 15. vg copies were at least 1 log lower in the LD cohort compared to the HD cohort at all time points (Figure 3A). In saliva, the AAVLK03.hOTC viral vector was quantifiable in all swab samples from the LD and HD groups from day 4 and day 8 (Figure 3B). In the LD group, viral vector was detected up to week 4 (only one positive animal), and in the HD group, viral vector was detected up to week 13 (only one positive animal). No vector copies were detected in saliva samples at week 26. In feces samples, the AAVLK03.hOTC vector was detected in LD up to 4 weeks and in HD up to 13 weeks. In urine samples, the vector was detected up to day 8 for the LD cohort and up to week 4 for the HD cohort (Figures 3C and 3D). No vector copies were detected in urine and feces samples from week 13 and week 26.

**Biodistribution**

Vector biodistribution analysis was performed by qPCR on all tissues listed in Table 1, collected at sacrifice 26 weeks after vector administration (Figure 4). The highest number of genome copies (GCs) was found in the liver with an average of \(1.6 \times 10^7\) GC/µg DNA in the HD group. In contrast, the average GC per microgram in the liver was 31-fold lower in the LD group at an average of \(5 \times 10^5\) GC/µg DNA. Not surprisingly, gallbladder biodistribution was also high. Most other organs displayed a GC number <1 \(\times 10^3\) GC/µg DNA (Figure 4) except for the spleen and aorta (circa \(10^5\) GC/µg DNA), lymph nodes, and adrenals (circa \(10^4\) GC/µg DNA). GC number in the central nervous system and genitals was particularly low (undetectable for most samples and <1 \(\times 10^2\) GC/µg DNA for all samples). No vector copies were detected in samples from the control group. No gender difference was observed.

**Supraphysiological liver OTC enzyme activity was sustained in all AAVLK03.hOTC-treated animals**

We sequentially measured liver OTC enzyme activity in biopsy samples obtained at 1 and 13 weeks and then at necropsy at 6 months post-administration of AAVLK03.hOTC in treated and control animals. Increased OTC activity was observed in the livers from all treated animals at all time points. In addition, a trend for dose response was observed at 1 and 13 weeks. OTC activity was 2 and 11 times higher in the HD compared with the LD group at 1 and 13 weeks post-administration, respectively. The increase of the OTC activity was similar in both treatment groups at week 26 (Figure 5A; Table S6). GC was almost 1 log lower in the LD group compared to the HD group at all time points (Figure 5B), which was consistent with the log lower dose administered in LD versus HD groups. No gender difference was observed in liver OTC activity either at LD (Figure 5C) or at HD (Figure 5D).

**DISCUSSION**

In this good laboratory practice-compliant study, we investigated the safety of AAVLK03.hOTC gene transfer administered intravenously to cynomolgus macaques at doses intended for the first-in-human clinical trial. To our knowledge, this is the first detailed report of a
Figure 2. Humoral immune response against AAVLK03 capsid

(A) Overall neutralization titers. Gender difference of neutralization titers between low-dose (B) and high-dose (C) groups. Neutralization titers are expressed as 1 in X dilution of serum. Horizontal lines display the mean ± SEM.
safety study using a vector with an engineered human liver-tropic AAV capsid in non-human primates. There were no adverse events or abnormal histological lesions observed at the doses intended for clinical use, supporting safety of this vector in juvenile animals. It is expected that as for wild-type AAV capsids, the testing and dose extrapolation of engineered capsids in animal models for human use will be reliable.\(^2\)

The biodistribution of AAVLK03.hOTC confirmed a remarkable liver tropism with partial biodistribution to the spleen. There was no transduction of the central nervous system and no or very low transduction of the genital organs. AAV8 has been historically the first serotype demonstrating long-lasting efficacy of AAV liver-directed gene therapy.\(^5,10\) AAVLK03 was shown to selectively transduce primary human hepatocytes in vivo, which was 12-fold better than AAV8 in chimeric mouse-human livers.\(^15\) However, AAVLK03 was not found to be superior to AAV8 in non-human primates \(s\) in terms of liver tropism.\(^24\)

The AAVLK03.hOTC vector was quantifiable from day 1 up to day 15 from the LD cohort and from day 1 to week 26 in plasma samples from the HD cohort, with a peak at day 1 (Figure 3A). In comparison, peripheral vein administration of the \(2 \times 10^{12} \text{ vg/kg}\) resulted in an average of \(3.6 \pm 0.9 \times 10^9 \text{ vg/mL}\) of self-complementary (sc)AAV2/8-LP1-hFIXco DNA in the plasma on day 1 following vector administration, which gradually declined to undetectable levels by day 10 in non-human primates.\(^25\) In a preclinical study assessing the biodistribution of AAVLK03.hOTC, the highest number of VGCNs in the liver and the gallbladder as expected, followed by spleen, aorta, lymph nodes, and adrenals. In a preclinical study using an AAV8 vector at a dose of \(2 \times 10^{11} \text{ vg/kg}\), liver was also the main transduced organ followed by adrenals, spleen, aorta, and heart. Similarly to our observation with AAVLK03.hOTC, transduction levels were lowest in the central nervous system and the genital organs using AAV8.\(^25\)

Immune responses to AAV can be classified as innate and adaptative. Adaptive immunity entails humoral and cell-mediated reactions. Several studies have shown the seroprevalence of Nabs directed against various AAV serotypes among the general population.\(^27,28\) This can be caused by multiple infections from various wild-type AAV serotypes.\(^27,29,30\) Another explanation can be the wide cross-reactivity of Nabs between AAV serotypes caused by homology of the amino acid sequence.\(^28\) In this study, after single administration of AAVLK03.hOTC, the humoral immune response had a maximal titer at week 2 post-administration and was similar in both genders. These titers decreased rapidly over the 26-week observation period with undetectable titers in some animals and a significant decrease in most animals especially in the LD group. The transient and limited immune response in our study supports the belief that engineered AAV capsids can evade Nabs more easily or generate Nabs, which rapidly decrease overtime.\(^15,16\) This might explain why anti-AAVLK03 Nabs are particularly low in childhood.\(^19\)

AAV vectors can also trigger the cellular immune response.\(^31\) Immune-mediated destruction of transduced hepatocytes is associated with CD8^+ T cell responses to AAV capsid antigens, which are processed and presented by major histocompatibility complex (MHC) class I.\(^32\) AAV vectors with a single-stranded (ss)DNA genome are able to stimulate the innate immune system through the Toll-like receptor (TLR)9/myeloid differentiation primary response gene 88 (MyD88) and type I IFN cascades.\(^33\) However, only a mild and highly transient immune response has been observed for ssAAVs regardless of capsid sequence, whereas changing the genome to scAAV increases innate immunity in a TLR9-dependent manner.\(^34\) In our study, the single administration of AAVLK03.hOTC did not show any cell-mediated immune response.

**Figure 3. Vector shedding profile of AAVLK03.hOTC**

Vector shedding profile of AAVLK03.hOTC in (A) plasma, (B) saliva, (C) feces, and (D) urine. Horizontal lines display the mean ± SEM. VGCN, vector genome copy number.
against either AAV capsid or OTC protein from isolated PBMCs. Similarly, a T cell response was not detected in non-human primates transduced with $2 \times 10^{12}$ vg/kg of AAV8 in a hemophilia B study. However, increased AAV8 capsid-reactive T cells were seen within 3 months after the delivery in a human clinical trial in patients injected with both $6 \times 10^{11}$ and $2 \times 10^{12}$ vg/kg doses. Long-term monitoring showed capsid-reactive T cells detected in peripheral blood for the first year following injection. Thus, a cellular immune response may be present in human trials despite an absence of its detection in preclinical studies.

Asymptomatic elevation in liver function tests is one of the adverse events associated with liver-targeted AAV gene therapy. Recombinant AAV encoding the hFIX gene caused transient elevations in transaminases in 2 out of 5 non-human primates in another preclinical study. In contrast, in a long-term safety and efficacy study for hemophilia B, macaques injected with $2 \times 10^{12}$ vg/kg of AAV8 vector had normal liver function tests for the 5 years of follow-up. However, the subsequent clinical trial revealed an asymptomatic transient increase of transaminases between week 7 and week 10 in 4 out of 6 patients injected with the same vector at the same dose. A phase I/II clinical trial in hemophilia A with AAVLK03 also revealed that three participants had elevation of transaminases. Therefore, despite the absence of increased transaminases in our preclinical study, careful monitoring is still warranted in future human clinical trials.

It has been previously shown that small differences in OTC activity can be responsible for the variability of the clinical phenotype in humans, which ranges between mild and severe. This study was performed in juvenile animals mimicking the growth of a pediatric liver. Interestingly, the increase in OTC activity following vector administration was sustained throughout the 26-week study period with a trend for dose response at early time points. This trend was surprisingly not observed at the end of the study, although there was larger variability in values at this time point compared to others, affecting all experimental groups including controls, which might have masked this effect. A sampling effect and the uneven OTC expression throughout the liver due to metabolic zonation could partially explain this variability. This contrasts with a significant dose response of VGNC at all time points. The GC levels in the growing liver of juvenile macaques also remained high over the 26 weeks with only a mild decline as expected, suggesting only a partial loss of episomal transgene copies in the growing liver. Cynomolgus macaques can live as long as 40 years (average 27 years). Female and male cynomolgus macaques reach sexual maturity at 4 and 7 years old, respectively. The animals used in this study were 14 to 18 months at the time of injection and considered as juvenile animals with ongoing liver growth, as seen in young pediatric patients. Altogether, these findings support the potential benefit of AAVLK03.hOTC in pediatric OTCD patients.

Overall, this study of intravenous AAVLK03.hOTC in juvenile cynomolgus macaques demonstrated safety of the tested vector with no toxicity, an excellent liver tropism with limited off-target biodistribution, a limited humoral immune response, and sustained increase in the enzyme activity in vector-treated animals. This study supports the clinical development of AAVlk03.hOTC for a first-in-human clinical trial.
MATERIALS AND METHODS

Vector production and study design

The codon-optimized hOTC (hcoOTC) transgene was generated by GenScript (Piscataway, NJ, USA) using the OptimumGene algorithm. The hcoOTC sequence is available as Supplemental information.

AAV (AAV-LK03) with the OTC transgene was produced by the Clinical Vector Core at the Children’s Hospital of Philadelphia. The process and materials were identical to those used in good manufacturing practice (GMP) manufacturing following Food and Drug Administration recommendations that materials designated for pharmacology/toxicology safety studies are GMP-process comparable. Briefly, the AAVLK03.hOTC vector was generated by triple calcium phosphate transient transfection of adherent human embryonic kidney (HEK) epithelial cells (HEK293) expanded in tissue-culture flasks and roller bottles prior to transfection. Cells were transfected using three plasmids that included the transgene plasmid (pAAV2-LPhOTC), Adenoviral-helper plasmid (pCCVC-AD2HPv2), and capsid plasmid (LK03). Post-transfection, the media were replaced with serum-free medium, and at the optimal time point post-transfection, cells and media were harvested. Crude harvest, including cells and media, was concentrated by hollow fiber tangential flow filtration (TFF), and cells were lysed by microfluidization (MF), treated with Benzonase, and clarified by 0.2 \( \mu \)lteration. AAV vector in the post-MF-filtered material was purified using ion exchange chromatography. Transgenes containing particles were fully separated from empty particles using cesium-chloride gradient centrifugation and collected. Viral vector derived from multiple batches was pooled, diafiltered into final formulation buffer (180 mM NaCl, 10 mM Na phosphate, 0.001% poloxamer 188) by TFF, sterile 0.2 \( \mu \)lter, aseptically filled in 1.5 mL cryovials at 1.0 and 0.3 mL per vial, and stored at \(-80^\circ\)C pending completion of QC testing and quality analysis (QA) release.

The AAVLK03.hOTC was administered via a single intravenous injection. The administration was performed in the left cephalic vein in both LD and HD groups and in the left saphenous vein in controls. After the injection, 1 mL of diluent provided by the manufacturer was flushed through the catheter. The amount of the test item to be administered was calculated according to the body weight recorded on the administration day. Control animals were administered with diluent following the same administration pattern as that followed for animals treated with the test item. Administration volume was set as 0.8 mL/kg (PBS), 0.36 mL/kg (2 \( \times \) 10^{12} vg/kg), and 0.8 mL/kg (2 \( \times \) 10^{13} vg/kg) in controls and LD and HD groups, retrospectively. Duration of monitoring post-injection was 26 weeks.

Animal procedures

All animal procedures were approved by the Ethical Institutional Committee at Envigo and were compliant with Animal Research: Reporting of In Vivo Experiments (ARRIVE) guidelines. Naive cynomolgus monkeys (Macaca fascicularis; supplier, KHI Bioservices, Hong Kong; breeder, Nafovanvn, Vietnam) were pre-screened to confirm absence of Nabs. The females were nulliparous and nonpregnant. 10 males and 10 females (9 males and 9 females were allocated to treatment groups) were between the age of 14 and 18 months at the beginning of treatment. In the interim period, their body-weight...
ranges were 1.5–2.2 kg and 1.6–2.3 kg in males and females, respectively. Animals were seronegative against AAV-LK03 and were allocated at random to the two treatment groups.

Safety and toxicity assessments

In-life analysis

The observations listed below were recorded. Viability/mortality/cage-side observations were done twice a day. Detailed clinical signs, including revision of the injection area, were recorded on the day of administration and weekly after dosing. Food consumption was evaluated by visual assessment and described as none, normal, and low. Daily check was done from the pre-test period to 26 weeks following the day of administration. Body weight was assessed weekly from pre-test, for 26 weeks after administration, and before sacrifice (fasted).

Signs are reported in four categories: detailed clinical signs, veterinary inspection, dose-site observations, and pen-side observations. Clinical observations are presented for all animals showing signs, providing detailed information of each sign, day or week of occurrence, and information on the duration of the sign.

All animals had an indirect ophthalmoscopic evaluation (Welch Allyn; model number [no.] 12500) at the pre-test period and in week 26. Observation areas were cornea, lens, conjunctiva, sclera, iris, and fundus.

All animals had electrocardiograms at the pre-test period and in week 26. Electrocardiograms (single snapshots) were obtained using Eindhoven (I, II, and III) and Goldberger (aVR, aVL, and aVF) leads. The heart rate, P-wave duration and amplitude, PQ interval, QRS interval, and QT interval were measured using a representative section of the electrocardiogram from lead II. Correction of the QT interval for heart rate (Fridericia) was also calculated.

Clinical laboratory investigations

The samples were collected early in the working day to reduce biological variation caused by circadian rhythms. The monkeys were fasted for blood collections. Blood samples were drawn from the femoral vein. Blood and urine samples were collected at the pre-test period and weeks 3–4, 12–13, and 25–26. The assay was performed at Envigo CRS, S.A.U., under internal laboratory QC conditions to assure reliable test results.

Hematology. Samples were collected in 0.5 mL tri-potassium-EDTA tubes. The following parameters were determined using an ADVIA 120 hematology analyzer: erythrocyte count (RBC), hematocrit (Hct), hemoglobin (Hb), mean corpuscular Hb (MCH), mean corpuscular Hb concentration (MCHC), mean corpuscular volume (MCV), platelet (thrombocyte) count (Plt), reticulocyte count (absolute and relative; Ret), total leucocyte count (WBC), differential leucocyte count (neutrophils [N], lymphocytes [L], monocytes [M], eosinophils [E], and basophils [B]), and large unstained cells (LUCs).

Coagulation. Samples were collected in 0.5 mL 3.2% sodium citrate (1 part anticoagulant to 9 parts blood) tubes in order to obtain the plasma. The following parameters were determined: prothrombin time (SPT) and activated partial thromboplastin time (SAPT).

Clinical biochemistry. Samples were collected in 0.8 mL lithium heparin tubes in order to obtain the plasma. The resultant plasma was analyzed using the Cobas 6000 analyzer for the following parameters: alanine aminotransferase (ALT); creatinine (Creat); alkaline phosphatase (ALP); albumin (Alb); Alb/globulin ratio (A/G); aspartate ALT (AST); bilirubin, total (Bili); calcium (Ca); chloride (Cl); cholesterol, total (Chol); creatine kinase (CK); gamma-glutamyltransferase (gGT); globulin*; glucose (Gluc); phosphorus, inorganic (Phos); potassium (K); protein, total (Total Prot); sodium (Na); triglycerides (Trig); and urea.

Urine analysis. The following parameters were determined: relative gravity (SG1), color (COL), appearance (App), pH, nitrite (NITE), protein (PROT), Gluc, ketone (Keto), urobinogen (Urob), Bili, erythrocytes (UBld), and leukocytes (UWBC).

Efficacy assessments

OTC enzyme activity was measured by a colorimetric assay, which detects the formation of L-citrulline. Liver tissue was prepared as a 1% (w/v) homogenate in deionized water, and fetal liver pellet was prepared as a 3%–5% (w/v) suspension in deionized water using a tissue homogenizer. Samples were stored at −70°C until the assay. Citrulline is produced by the OTC reaction, similar to the reaction in vivo, and was quantitated using a phenazone diacetyl reagent producing a strong yellow color read on a spectrophotometer set at 464 nm. CPS was measured as a control enzyme.

OTC + ornithine

Carbamoyl phosphate → citrulline

Citrulline + phenazone/diacetyl reagent → yellow product (absorbance [Abs.] 464 nm)

600 µL of working phenazone diacetyl reagent was added to 100 µL of the assay supernatants in 5 mL stopped glass tubes and for the standards. They were heated at 100°C for 10 min in the dark. Then they were cooled in ice water and then allowed to come to room temperature before reading. The color was stable at room temperature. The absorbance was measured at 464 nm using a glass microcuvette. Then the cuvette was rinsed with phenazone diacetyl reagent and the sensitivity of the spectrophotometer adjusted using the standard containing no citrulline. Results were expressed in micromoles of citrulline/hour/milligram of proteins.

Viral shedding and biodistribution

DNA was extracted from tissues issuing the QIAamp Fast DNA Tissue Kit (ref. 51404; QIAGEN). Assays results were reported by Real-time PCR Equipment Mx3005P Stratagene (Agilent Technologies). For tissues, results were adjusted to double-stranded vector copies/
serum samples were incubated for 1 h at 37°C from 1 in 5 and then in 2-fold serial dilutions to 1:640. The diluted medium (DMEM) (Gibco, Invitrogen, Grand Island, NY, USA) without antibiotics at 15,000 rpm for 10 min. On the day of transduction, dilution was incubated at the same concentration to reach a predetermined final multiplicity of infection (MOI) into a 100-μL final volume for transduction. The optimal transduction for the assay was reached at an MOI of 1,000 vg/cell for AAVLK03. Purified immunoglobulins (Octagam 10% [v/v]; Octapharma, Manchester, UK) were used as positive control. Each assay was run in duplicate. The next day, wells were complemented with DMEM and 10% (v/v) FCS. Fluorescence-activated cell sorting BD FACSVerse (BD Biosciences, San Jose, CA, USA) analysis of the GFP signal was performed 72 h post-transduction. Samples were considered positive when a 1:5 dilution of serum reduced the vector transduction by 50% or more. The Nab titer was determined as the highest positive serum dilution.

**Necropsy**

Animals were euthanized with intravenous pentobarbital overdose and necropsied and tissues harvested for comprehensive histopathologic examination. Table 1 lists tissues that were collected and examined histologically. Organ weights were presented both as absolute/ unadjusted and adjusted for terminal body weight using the weight recorded on the day of necropsy.

**Histology**

All organ and tissue samples to be examined by the study pathologist were fixed and sent to ProPath for processing. The samples were processed, embedded, cut at a nominal thickness of 4–5 μm, and stained with hematoxylin and eosin. After evaluation, any remaining tissues and organs were stored for archiving. The bone marrow smears were stained using the May-Grunwald Giemsa method and stored for possible further investigation.

**Pathology**

Slides of all organs and tissues collected at necropsies of all animals were examined by a veterinary pathologist at Envigo CRS, S.A.U. Where possible, the microscopic findings were correlated with the gross observations. A peer review of microscopic findings was performed in accordance with Envigo standard operating procedures.

**Immunological assessments**

**Humoral**

Blood samples (nominally, 500 μL) were collected from the femoral vein from all animals on the following occasions: pretreatment and weeks 2, 4, 13, and 25–26. The blood samples were maintained at room temperature until centrifugation. Each blood sample was centrifuged at 1,500 × g during 15 min at room temperature (11 to 68 min from collection) after coagulation in order to obtain around 200 μL of serum. The resulting serum was transferred to a polypropylene vial and then was immediately frozen in dry ice and stored below −80°C ± 10°C until dispatch.

Human hepatocellular carcinoma HuH-7 cells were seeded at 5 × 10³ cells per well on a 48-well plate and left for at least 7 h before transduction. Before use, sera were heat inactivated at 56°C for 30 min. Precipitate resulting from heat inactivation was removed by centrifugation at 15,000 × g for 10 min. On the day of transduction, dilution of serum sample was performed in Dulbecco’s modified Eagle’s medium (DMEM) (Gibco, Invitrogen, Grand Island, NY, USA) without fetal calf serum (FCS) (JRH Biosciences, Lenexa, KS, USA) starting from 1 in 5 and then in 2-fold serial dilutions to 1:640. The diluted serum samples were incubated for 1 h at 37°C with AAVLK03.hAAT.GFP diluted in an equal volume of DMEM. AAVLK03 was incubated at the same concentration to reach a predetermined final multiplicity of infection (MOI) into a 100-μL final volume for transduction. The optimal transduction for the assay was reached at an MOI of 1,000 vg/cell for AAVLK03. Purified immunoglobulins (Octagam 10% [v/v]; Octapharma, Manchester, UK) were used as positive control. Each assay was run in duplicate. The next day, wells were complemented with DMEM and 10% (v/v) FCS. Fluorescence-activated cell sorting BD FACSVerse (BD Biosciences, San Jose, CA, USA) analysis of the GFP signal was performed 72 h post-transduction. Samples were considered positive when a 1:5 dilution of serum reduced the vector transduction by 50% or more. The Nab titer was determined as the highest positive serum dilution.

**Cytokine determination**

Totally, 90 blood samples (nominally, 250 μL) were collected in tubes with EDTA K₂ from the femoral vein from all animals on the following occasions: day 1 (predose, 4, 8, and 24 h after administration) and weeks 25–26. The blood samples were maintained at room temperature until centrifugation. Each blood sample was centrifuged at 2,000 × g during 10 min at 4°C in order to obtain plasma. The resulting plasma was transferred to a polypropylene vial and then was immediately frozen on dry ice and stored below −80°C ± 10°C until dispatch. Plasma levels of IL-1β, IL-2, IL-4, IL-5, IL-6, IL-8, TNF-α, G-CSF, and IFN-γ were analyzed at Envigo CRS (UK). This assay is based on the Luminex xMAP technology. Data are presented as concentrations in whole matrix expressed in picograms per milliliter.

**Statistical analysis**

Statistical analysis has been performed; however, due to the small sample size, there may not be adequate power to identify biologically relevant differences as being statistically significant. All statistical analyses were carried out separately for males and females. For all parameters, the analyses were carried out using the individual animal as the basic experimental unit. The following data types were analyzed at each time point separately: body weight, electrocardiography, hematology, blood chemistry, urine analysis, macropathology, and organ weights (absolute and adjusted for terminal body weight). Comparison of continuous variables between two experimental groups was performed using the Student’s two-tailed t test. p values < 0.05 were considered statistically significant.
were considered statistically significant. Standard error of the mean (SEM) was used as the indicator of dispersion.

SUPPLEMENTAL INFORMATION
Supplemental information can be found online at https://doi.org/10.1016/j.omtm.2021.09.005.

ACKNOWLEDGMENTS
The views expressed are those of the author(s) and not necessarily those of the NHS, NIHR, or Department of Health. We are grateful to the team of the Clinical Vector Core, The Raymond G. Perelman Center for Cellular and Molecular Therapeutics, and The Children’s Hospital of Philadelphia (Philadelphia, PA, USA) for their assistance in vector production and characterization, especially Dr. Johannes Van Der Loo, Olga Zelenaia, and Stacey Piecyk. This work was supported by funding from the United Kingdom Medical Research Council grants MR/N019075/1 and MR/R026084/1 to P.G., Great Ormond Street Hospital Charity Clinical Research Starter grant 531808 to P.G., and NIHR Great Ormond Street Hospital Biomedical Research Centre (to P.G. and J.B.). J.B. is currently supported by an MRC Clinician Scientist Fellowship MR/T008024/1. B.S.Y. is supported by the Turkish Scientific and Technological Research Council (TUBITAK), grant 2219/2018-2.

AUTHOR CONTRIBUTIONS
J.B. and P.G. designed and supervised the study. J.B. and B.S.Y. wrote the manuscript. D.P.P. performed OTC enzymatic assays and immunnoassays. S.C.C., A.J.T., L.L., and I.E.A. offered scientific and technical assistance. D.B. provided assistance in the OTC enzyme assay. All authors reviewed and approved the manuscript.

DECLARATION OF INTERESTS
I.E.A. and L.L. are inventors on patents related to the AAVLk03 capsid or the transgene cassette used in AAVLk03.hOTC. The other authors have no competing interests.

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