RESEARCH ARTICLE

Oral tetrahydouridine and decitabine for non-cytotoxic epigenetic gene regulation in sickle cell disease: A randomized phase 1 study

Robert Molokie, Donald Lavelle, Michel Gowhari, Michael Pacini, Lani Krauz, Johara Hassan, Vinzon Ibanez, Maria A. Ruiz, Kwok Peng Ng, Philip Woost, Tomas Radivoyevitch, Daisy Pacelli, Sherry Fada, Matthew Rump, Matthew Hsieh, John F. Tisdale, James Jacobberger, Mitch Phelps, James Douglas Engel, Santhosh Saraf, Lewis L. Hsu, Victor Gordeuk, Joseph DeSimone, Yogen Saunthararajah

1 Department of Medicine, University of Illinois Hospital and Health Sciences System, Chicago, Illinois, United States of America, 2 Jesse Brown VA Medical Center, Chicago, Illinois, United States of America, 3 Translational Hematology and Oncology Research, Taussig Cancer Institute, Cleveland Clinic, Cleveland, Ohio, United States of America, 4 Case Comprehensive Cancer Center, Case Western Reserve University, Cleveland, Ohio, United States of America, 5 Department of Quantitative Health Sciences, Cleveland Clinic, Cleveland, Ohio, United States of America, 6 Department of Hematology and Oncology, Taussig Cancer Institute, Cleveland Clinic, Cleveland, Ohio, United States of America, 7 Molecular and Clinical Hematology Section, National Institutes of Health, Bethesda, Maryland, United States of America, 8 College of Pharmacy, The Ohio State University, Columbus, Ohio, United States of America, 9 Cell and Developmental Biology, University of Michigan, Ann Arbor, Michigan, United States of America

* saunthy@ccf.org

Abstract

Background

Sickle cell disease (SCD), a congenital hemolytic anemia that exacts terrible global morbidity and mortality, is driven by polymerization of mutated sickle hemoglobin (HbS) in red blood cells (RBCs). Fetal hemoglobin (HbF) interferes with this polymerization, but HbF is epigenetically silenced from infancy onward by DNA methyltransferase 1 (DNMT1).

Methods and findings

To pharmacologically re-induce HbF by DNMT1 inhibition, this first-in-human clinical trial (NCT01685515) combined 2 small molecules—decitabine to deplete DNMT1 and tetrahydouridine (THU) to inhibit cytidine deaminase (CDA), the enzyme that otherwise rapidly deaminates/inactivates decitabine, severely limiting its half-life, tissue distribution, and oral bioavailability. Oral decitabine doses, administered after oral THU 10 mg/kg, were escalated from a very low starting level (0.01, 0.02, 0.04, 0.08, or 0.16 mg/kg) to identify minimal doses active in depleting DNMT1 without cytotoxicity. Patients were SCD adults at risk of early death despite standard-of-care, randomized 3:2 to THU–decitabine versus placebo in 5 cohorts of 5 patients treated 2X/week for 8 weeks, with 4 weeks of follow-up. The primary endpoint was ≥ grade 3 non-hematologic toxicity. This endpoint was not triggered, and
adverse events (AEs) were not significantly different in THU-decitabine—versus placebo-treated patients. At the decitabine 0.16 mg/kg dose, plasma concentrations peaked at approximately 50 nM (C<sub>max</sub>) and remained elevated for several hours. This dose decreased DNMT1 protein in peripheral blood mononuclear cells by >75% and repetitive element CpG methylation by approximately 10%, and increased HbF by 4%–9% (P < 0.001), doubling fetal hemoglobin-enriched red blood cells (F-cells) up to approximately 80% of total RBCs. Total hemoglobin increased by 1.2–1.9 g/dL (P = 0.01) as reticulocytes simultaneously decreased; that is, better quality and efficiency of HbF-enriched erythropoiesis elevated hemoglobin using fewer reticulocytes. Also indicating better RBC quality, biomarkers of hemolysis, thrombophilia, and inflammation (LDH, bilirubin, D-dimer, C-reactive protein [CRP]) improved. As expected with non-cytotoxic DNMT1-depletion, platelets increased and neutrophils concurrently decreased, but not to an extent requiring treatment holds. As an early phase study, limitations include small patient numbers at each dose level and narrow capacity to evaluate clinical benefits.

Conclusion

Administration of oral THU-decitabine to patients with SCD was safe in this study and, by targeting DNMT1, upregulated HbF in RBCs. Further studies should investigate clinical benefits and potential harms not identified to date.

Trial registration

ClinicalTrials.gov, NCT01685515

Author summary

Why was this study done?

- Sickle cell disease, one of the most frequent inherited diseases in humans, is driven by aggregation and precipitation of less soluble sickle cell hemoglobin in red blood cells—this destroys red blood cells and blocks blood vessels, causing organ damage, suffering, and early death.

- Fetal hemoglobin, expressed in red blood cells until infancy, mixes with sickle cell hemoglobin and promotes its dissolution. Individuals who naturally express high levels of fetal hemoglobin beyond infancy thus receive some protection from sickle complications.

- To mimic this natural state using drugs, one relevant observation was that fetal hemoglobin is increased during recovery of bone marrow from extreme stress. This led to evaluation and approval of the cytotoxic (cell killing) drug hydroxyurea, the only drug approved to treat sickle cell disease. This approach to fetal hemoglobin induction, however, is limited in potency and sustainability.

- Our goal was to increase fetal hemoglobin by inhibiting an enzyme, DNA methyltransferase 1 (DNMT1), involved in shutting off the fetal hemoglobin gene from infancy onward.
What did the researchers do and find?

- We used the drug decitabine to inhibit DNMT1. However, decitabine is very rapidly inactivated in the body by another enzyme, cytidine deaminase (CDA). We therefore combined it with a CDA inhibitor, tetrahydrouridine (THU). Oral THU and decitabine were given to patients with severe, symptomatic sickle cell disease who had not derived benefit from the standard treatment of hydroxyurea.

- Oral THU-decitabine was safe and well-tolerated by the patients.

- Measurement of decitabine levels in the blood confirmed that THU enabled oral absorption of very small doses of oral decitabine. Measurements of DNMT1 protein levels and DNA methylation confirmed that this decitabine exposure was sufficient to deplete DNMT1 from cells.

- DNMT1 depletion by decitabine produced large increases in fetal hemoglobin and increased numbers of healthy red blood cells.

What do these findings mean?

- Oral THU-decitabine to deplete DNMT1 safely increased fetal hemoglobin and numbers of healthy red blood cells.

- Since this was a first-in-human study, only a small number of patients were treated and for a fairly short time. Further clinical studies are therefore needed.

Introduction

Sickle cell disease (SCD) is a congenital hemolytic anemia caused by polymerization and precipitation of mutated sickle hemoglobin (HbS, $\alpha_2\beta^S_2$) in red blood cells (RBCs). This decreases RBC life span by $>90\%$, causing severe anemia. The unhealthy RBCs also adhere to and occlude blood vessels. Ensuing tissue hypoxia damages all organs, triggers severe pain, and compromises immunity. In low-income countries, most children with SCD do not survive to adulthood. Even in high-income countries, morbidity can be severe and life spans are reduced by several decades—the median life expectancy for people with SCD in the United States is approximately 45 years [1,2].

RBCs at the fetal stage of life contain fetal hemoglobin (HbF, $\alpha_2\gamma_2$). Normal adult hemoglobin (HbA, $\alpha_2\beta_2$) polymerizes with HbS, while HbF intercalates with but does not polymerize with HbS [3–5]. HbF thus interrupts SCD pathophysiology at its inception [3,4], and higher HbF correlates with fewer vaso-occlusive pain crises, less renal damage, less pulmonary hypertension, fewer strokes, and longer survival [1,6–12] (reviewed in [5]). That is, any increase in HbF provides some benefit. Those few SCD patients who inherit HbF at 20%–30% of total hemoglobin (hereditary persistence of fetal hemoglobin, HPFH) have essentially normal life spans [13–15]. Decades of research has hence been directed towards pharmacologic recapitulation of this naturally selected protective state [5]. A key observation directing early efforts was that HbF is enriched during bone marrow recovery from extreme stress [16–21]. Cytotoxic (cell killing) drugs can create such stress, leading to evaluation in SCD of the oral ribonucleotide reductase inhibitor hydroxyurea [20–22]. In a pivotal trial, hydroxyurea (15–35 mg/kg) increased HbF for
2 years in approximately 50% of the adult SCD patients treated [22,23]. HbF induction correlated strongly with increased RBC half-life [24,25], fewer pain crises [23], and better quality of life [26]. Trial patients with HbF levels >0.5 g/dL survived longer [8]. Average HbF increases at 2 years, however, were modest (3.6%) [20–23,27]. Moreover, HbF increases were particularly unlikely in patients with the lowest baseline HbF levels and thus at highest risk of morbidity and mortality [23,25,28,29], and even patients with excellent initial HbF inductions demonstrated diminishing inductions over time [23,30]. Lower and less durable HbF increases also correlated with fewer reticulocytes (<300,000 × 10^9/L) and neutrophils (<7.5 × 10^9/L) at baseline. This correlation underscores that HbF induction by cytotoxicity requires reserves of hematopoietic precursors sufficient to repeatedly mount recoveries from bone marrow stress that destroys their counterparts [21,23]. Cumulative attrition of these reserves occurs via vaso-occlusion in the marrow and to the kidneys [23,25,28,29]. This is a problem even separate from considerations of sustainable HbF induction via cytotoxicity: in SCD, erythropoiesis has to operate at >10-fold the normal rate to barely sustain hemoglobin levels compatible with life, and dwindling compensatory capacity is a major cause of early death [8,23,31,32]. Therefore, new, non-cytotoxic, durable, and more potent methods of inducing HbF are needed.

The chromatin-modifying enzyme DNA methyltransferase 1 (DNMT1) maintains methylation marks on DNA through cell division. DNMT1 is also a co-repressor—it executes the biochemical work of silencing genes for sequence-specific DNA-binding factors, e.g., BCL11A and TR2/TR4, that direct epigenetic silencing of the HbF gene (γ-globin, HBG) [33–45]. The deoxycytidine analog decitabine can deplete DNMT1—a nitrogen substituted for a carbon in the decitabine pyrimidine ring covalently binds to DNMT1 and causes its degradation [46]. Importantly, the decitabine deoxyribose moiety is unmodified; thus, it can incorporate into the elongating DNA strand during S-phase without terminating chain extension or causing cytotoxicity [47,48]. High concentrations of decitabine, however, do produce off-target antimitabolite effects and cytotoxicity, in significant part via the uridine moiety degradation products that can misincorporate into DNA or inhibit thymidylate synthase [49,50]. Decitabine regimens approved by the US Food and Drug Administration (FDA) for the treatment of myeloid cancer, having evolved out of the traditional cytotoxic intent of oncology, utilize such cytotoxic doses and require pulse-cycled administration to recover from cytotoxic side effects. We therefore redesigned decitabine application for non-cytotoxic, molecularly-targeted therapy of non-malignant and malignant diseases [43,51–54]. Specifically, we selected demonstrably non-cytotoxic yet DNMT1-depleting doses, used a subcutaneous route of administration to blunt C_max, and administered these doses frequently and in distributed fashion (2–3X/week) to increase the fraction of target cells exposed in S-phase, as DNMT1-depletion by decitabine is S-phase–dependent [43,51–54]. This approach increased HbF by >10% in SCD patients who had no HbF response to hydroxyurea in the pivotal clinical trial (i.e., a mean change in HbF after 2 years of treatment with hydroxyurea of 0.3%) [42,43,55]. Hindering wide clinical application of decitabine in this way, however, are a number of limitations with its pharmacology: it has a very brief plasma half-life of only minutes [56,57] (a problem since DNMT1-depletion is S-phase, and hence exposure-time, dependent); has negligible solid tissue distribution (a problem when target cells reside in solid tissues, e.g., erythroid precursors in the spleen in β-thalassemia); and hence also has negligible oral bioavailability (a problem for worldwide and long-term therapy). These pharmacology problems have a common cause: rapid deamination/inactivation of decitabine to uridine degradation products by the pyrimidine metabolism enzyme cytidine deaminase (CDA) that is highly expressed in tissues such as the intestines and liver [58–62].

Thus, we combined a CDA inhibitor, oral tetrahydrouridine (THU), with oral decitabine: THU is a uridine analog competitive inhibitor of CDA that, although not FDA-approved for
any indication (new chemical entity), has not caused toxicity to animals or to humans in several clinical trials, including a trial with oral administration exceeding a year [40,59–61,63–69]. Here, building on encouraging results from pre-clinical in vivo studies [70], we report the first-in-human clinical translation of oral THU–decitabine for HbF induction in SCD. Specific goals were to evaluate safety of oral THU–decitabine and recommend a phase 2 dose for DNMT1-depletion and HbF induction in SCD [70].

### Materials and methods

#### Ethics statement

All research involving human participants was approved by the Cleveland Clinic and University of Illinois at Chicago Institutional Review Boards (IRBs), and all clinical investigation was conducted according to the principles expressed in the Declaration of Helsinki. Written informed consent was obtained from the participants on the IRB approved protocol. The protocol number was Case 10z11. The United States FDA Investigational New Drug number is 112,914.

#### Study design

This was a single-blind dose-escalating phase 1 clinical trial with a maximum of 5 decitabine dose levels. Five patients were enrolled at each dose level, randomized with 3:2 odds to study drugs versus placebo (5 cohorts of 5 randomized patients).

This design was not altered during the course of the study.

#### Patient population

Written informed consent was obtained prior to treatment in all patients. The treatment population was adult (>18 years of age) SCD (SS or S-β-thalassemia) patients who despite standard-of-care hydroxyurea for ≥6 months (or being intolerant or unwilling to take hydroxyurea) were at risk of early death as defined by published criteria [8]. These criteria were at least 1 of the following: (i) HbF <0.5 g/dL, (ii) 3 or more pain episodes per year requiring parenteral narcotics, (iii) 1 or more acute chest syndrome episodes, and (iv) total hemoglobin <9 g/dL and absolute reticulocyte count ≤250,000 × 10⁹/L.

#### Interventions

Decitabine and THU were synthesized by Ash Stevens (Detroit, MI). Drugs were stored in glass bottles at −20˚C. Bottles were opened after equilibration to room temperature. The appropriate amount of drug was weighed out and reconstituted with water for consumption by patients in the clinic within 30 minutes of drug reconstitution. Placebo was an equivalent amount of water without study drug.

Fixed-dose oral THU 10 mg/kg was administered 60 minutes before oral decitabine at 0.01, 0.02, 0.04, 0.08, or 0.16 mg/kg (Figs 1, 2A and 2B). Repeat dose administration 2X/week for 8 weeks, instead of single dose administration, was used to assess safety and efficacy—to increase the likelihood that the dose identified for further studies would be safe and efficacious for the intended application of chronic disease modification (Fig 2A and 2B).

Patients at each dose level were monitored weekly to determine if next treatments should be withheld based on laboratory endpoints; in previous clinical trials, the most sensitive indices of decitabine biologic activity were an increase in the platelet count and a decrease in the absolute neutrophil count (ANC). The plan was to use threshold values of these parameters not associated with clinical toxic events, being within the range observed in patients with SCD during their routine care with hydroxyurea and/or splenectomy, to trigger dose modification, and
Fig 1. Fetal hemoglobin (HbF) blocks polymerization of deoxy sickle hemoglobin (HbS), the root cause of sickle cell disease (SCD) pathophysiology, and is the most powerful known disease modifier. (A) Polymerization of deoxy HbS drives all SCD pathophysiology; in contrast to HbF, normal adult hemoglobin (HbA, ß-chains) can participate in polymerization. (B) The gene for HbF (HBG) is silenced by DNA methyltransferase 1 (DNMT1). Although DNA-binding factors, e.g., BCL11A, direct this silencing, the biochemical work of epigenetic repression is executed by chromatin-modifying enzymes, amongst which DNMT1 is central. Decitabine depletes DNMT1 and can do so without cytotoxicity because in contrast to other cytidine analogues (e.g., cytarabine) the deoxyribose moiety (green dotted circle) is natural, although higher concentrations do cause anti-metabolite effects and DNA damage, in part by degradation into uridine counterparts that misincorporate into DNA. (C) Several pharmacologic limitations of decitabine hinder safe, effective, practical clinical translation. The limitations have a common cause, the enzyme cytidine deaminase (CDA). Tetrahydrodouridine (THU) inhibits CDA. No toxicities have been found for THU in animals or humans.

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Fig 2. Design rationale, schema, flow, and patient characteristics. (A) Decitabine regimens approved by the US Food and Drug Administration (FDA) to treat myeloid malignancy utilize cytotoxic decitabine doses (red zone), requiring pulse-cycled administration to recover from cytotoxic side effects. This clinical trial instead escalated oral decitabine doses from almost zero (0.01 mg/kg) gradually upward to find the minimum doses required to deplete DNA methyltransferase 1 (DNMT1) without cytotoxicity (green zone) when administered after oral tetrahydrouridine (THU) 10 mg/kg. Because such doses are non-cytotoxic, they can be administered frequently in distributed fashion to increase the fraction of target cells subject to S-phase-dependent DNMT1 depletion. (B) The study schema. (C) Flow of patients through the trial. (D) Patient clinical characteristics at baseline. Abbreviations: ACS, acute chest syndrome; AVN, acute vascular necrosis; CVA, cerebral vascular accident; ED/Hosp, emergency department/hospital; PE/DVT, pulmonary embolus/deep vein thrombosis. (E) Patient demographics and baseline laboratory variables. Median and range showed for continuous variables. P values Wilcoxon ranked sums 2-tailed or Fisher exact test.

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thereby maintain safety [43,51]. These thresholds were ANC < 1.5 × 10⁹/L and platelets > 1,200 × 10⁹/L. Non-hematologic toxicity ≥ grade 3 attributed to study drug was also to trigger a dose modification, as was decitabine C<sub>max</sub> > 0.2 μM. Drug was to be held until recovery below these thresholds, when drug was to be restarted with a 25% decrease in dose.

Concurrent hydroxyurea therapy was explicitly disallowed, with a requirement for a 28-day washout period from the last hydroxyurea dose to initiation of study drug or placebo.

**Outcomes**

The primary endpoint was ≥ grade 3 non-hematologic toxicity. In addition, the a priori study design required dose modification for platelets >1,200 × 10⁹/L, or neutrophils < 1.5 × 10⁹/L. Our goal was to provide evidence in favor of a null hypothesis that patients in treated groups (oral THU-decitabine 2X/week over 8 weeks; n = 15) do not experience treatment-related events requiring dose modification more than patients in the placebo group (n = 10). Secondary endpoints included (i) sickle cell crisis frequency (efficacy), (ii) coagulation (D-dimer) and inflammatory (C-reactive protein [CRP]) pathway activity (efficacy), (iii) HbF levels measured by high-performance liquid chromatography (HPLC) (efficacy and pharmacodynamics of decitabine), (iv) DNA methylation levels at repetitive elements in buffy coat DNA (pharmacodynamics of decitabine), and (v) DNMT1 levels in buffy coat cells (pharmacodynamics of decitabine). A priori secondary endpoints not measured for cost and technical reasons were CDA genotype and CDA functional activity in serum.

**Sample size**

The study employed a randomized design to identify an oral dose of decitabine that can be administered twice a week in combination with oral THU over an 8-week period without requiring dose modification. We used a standard 3+3 dose escalation design with at most 5 dose levels. There would thus have been 6 in the maximum dose group if a dose-limiting toxicity had been detected. Such a 6-patient group would have detected at least 1 toxicity with a probability of 0.88 if it occurred in 30% of the population; this value is calculated as 1 minus the probability of zero toxicities, i.e., 1 − 0.7<sup>6</sup>. Ten patients were treated with placebos, 2 at each dose level, so the maximum sample size was 15 + 10 = 25. If a dose modification was required because of ANC < 1.5 × 10⁹/L or platelets > 1200 × 10⁹/L or ≥ grade 3 non-hematologic toxicity attributed to the study drug, in 1 or more of the 3 patients who received the study drug at a dose-level cohort, the plan was to accrue another cohort of 5 new patients (3:2 study-drug:placebo) with the treatment dose based on the cumulative dose administered to patients receiving the study drug in the preceding cohort. The Safety and Intent-To-Treat (ITT) populations included all enrolled patients receiving at least one dose of decitabine.

**Randomization procedures**

Randomization was at the University of Illinois at Chicago by Dr. Michael Pacini using a randomization table created on www.randomization.com. Patients were randomly allocated to study drug treatment versus placebo.

**Blinding**

Patients, but not investigators, were blinded as to the assigned treatment. The experimental treatment was highly diluted in water. Therefore, the method of patient blinding was administration of water placebo with similar appearance, volume, and taste as the study drug.
Pharmacokinetics

Blood was collected for pharmacokinetic analysis at 0, 2, 4, and 24 hours after the first decitabine dose (week 1). Blood samples were drawn over a period of less than 1 minute into tubes pre-loaded with heparin and THU 10 μg/ml (to prevent in vitro metabolism by CDA) and immediately transferred onto ice. Samples were then centrifuged as soon as possible at 600 g for 5 minutes at 4°C. After separation, plasma was transferred in 0.2 ml aliquots into pre-frozen vials and stored frozen at −80°C until shipment to Ohio State University for analysis (shipment on dry ice) by a liquid chromatography tandem mass spectrometry (LCMS/MS) method that has been previously described in detail for determination of decitabine in human, baboon, mouse, and rat plasma [56,57,70]. Pharmacokinetic data were analyzed using non-compartmental methods or a 2-compartment model with instantaneous/intravenous input using the R package PKLMfit. The model-fitting method allowed estimation of terminal half-lives for some data sets. The AUC_{last} (the area under the curve from the time of dosing to the last measurable concentration) was calculated by the linear trapezoidal method.

DNMT1 protein measurement by flow cytometry of peripheral blood mononuclear cells (pharmacodynamic analyses)

Phlebotomized whole blood was layered over Histopaque. The interface was collected and washed with phosphate-buffered saline (PBS). Cell suspensions of approximately 200,000 cells were fixed with 4% paraformaldehyde for 30 minutes on ice. Cells were spun down, washed in PBS, and suspended in 70% ethanol and stored at −20°C for subsequent batched analyses. For analyses, patient samples were thawed and centrifuged. All procedures were performed on ice and all centrifugations were done at 400 g for 5 minutes at 4°C. The pellet was hydrated overnight in 1 ml sterile, distilled water to partially reverse the alcohol fixation. After incubation, samples were centrifuged and then resuspended in 1 ml PBS/2% bovine serum albumin (BSA) and incubated for 30 minutes to block nonspecific antibody binding. Following centrifugation, unlabeled anti-DNMT1 antibody [EPR 3522] (0.0625 μg/test; Abcam; catalog no. ab92314) was added in a final volume of 100 μl and incubated for 1 hour. Samples were washed 3 times: each wash was a 10-minute incubation with 1 ml PBS/2% BSA followed by centrifugation.

After the third wash, CD64-Alexa Fluor 488 (5 μl/test; BioLegend; catalog no. 305010), Cyclin A2-PE (4μl/test; Beckman Coulter; catalog no. B15092), CD33-APC/Cy7 (5 μl/test; BioLegend; catalog no. 366614), and F(ab')2-goat anti-rabbit IgG (H+L) Alexa Fluor 647 (0.0938 μg/test; Life Technologies; catalog no. A21246) were added in a final volume of 100 μl and incubated for 1 hour. After the incubation and without washing, 3.5 ml PBS containing 0.5 μg/ml DAPI was added to each sample. All samples were analyzed on an Attune Nxt Acoustic Focusing Cytometer (Life Technologies) at a flow rate of 500 μl per minute. Compensation was performed with CompBeads Set Anti-Mouse Ig, κ (BD Biosciences; catalog no. 552843) and Flow Cytometry Protein G Antibody Binding Beads (Bangs Laboratories, Inc.; catalog no. 554/11863).

Data analysis. Data were normalized to the instrument using 8-peak SPHERO Rainbow Calibration Particles (Spherotech; catalog no. RCP-30-5A). WinList 3D v8.0 (Verity Software House) was used for post-acquisition analysis. Doublet discrimination was performed using the DAPI area and peak signals. Gated singlet events were displayed in a bivariate plot of cyclin A2 versus DNA content to identify S + G2 + M phase cells as both cyclin A2 positive and >2C DNA. These color-evented cells were used as a guide to set the boundary between DNMT1 positive and negative events. The median DNMT1-negative value was subtracted from the median DNMT1-positive value. This net value was further processed by normalization using linear regression of 8 peak bead sets between runs performed on different days.
Methylation level of LINE-1 repetitive element CpG by pyrosequencing

DNA was purified from peripheral blood mononuclear cells (isolated by Ficoll-Hypaque density centrifugation) using the Wizard Genomic DNA Purification Kit (Promega). DNA was bisulfite-converted using EZ DNA Lightning Methylation kit (Zymo Research). The PCR primers were 10 pmol of 5'-TTTGTAGTTAGTGGG-3' and 10 pmol of biotinylated-5'-TCTCAAAAAATACCAAC-3'. PCR cycling conditions were: cycle temperature of 94˚C for 30 seconds, 60˚C for 30 seconds, and 72˚C for 30 seconds, for 45 cycles to consume all the biotinylated primers. The biotinylated strand was captured on streptavidin sepharose beads (Amersham Biosciences, Uppsala, Sweden) and annealed with the sequencing primer 5'-GGGTGGGAGTGAT-3'. The methylation degree of long interspersed nuclear elements (LINE-1) was computed at 3 CpG sites pyrosequenced with the Pyromark Q24 Pyrosequencer (Qiagen) using the dispensation order GTCGATTAGTGCTAGTGTATC.

Clinical pathology tests

Blood counts, blood chemistries, total bilirubin, HbF, LDH, and D-dimer levels were standard clinical pathology tests through the CLIA-certified Clinical Pathology Laboratory at the University of Illinois at Chicago.

Measurement of HbF levels

Analysis of globin chains was performed on a TSP Spectra HPLC system using a LiChristopher 100 RP-8 column and a gradient of acetonitrile-methanol-NaCl as per the standard CLIA-certified Clinical Pathology Laboratory methods at University of Illinois at Chicago. For quantification of F-cells by flow cytometry, peripheral blood samples were fixed and stained with phycoerythrin-conjugated anti-HbF (Caltag) per manufacturer’s instructions. A normal adult negative control and cord blood positive control were run with each Becton-Dickinson Facs- Calibur (Sunnyvale, CA) analysis.

Statistics

For statistical comparisons of adverse events (AEs) between placebo and each dose group, we performed exact tests of the null hypothesis that the rate parameters of Poisson distributed AEs are equal. Relative risks are given as the ratio of such rate parameters. Thus, differences are insignificant if relative risks confidence intervals include 1. The R function poisson.test(), an exact test, was used. Pain, the most frequent AE, was analyzed similarly.

For time course analyses of HbF and total hemoglobin endpoints in placebo versus each dose group, linear functions of time were fitted to each patient’s time course of measurements on-treatment. The slopes of such fits versus the dose that the patient received were fitted to linear dose-response to generate P values of such slopes of slopes, i.e., lines in dose-response plots. This approach controls for inter-patient differences in initial values as per the planned mixed-effects modeling.

Results

Patient flow and characteristics

The trial protocol is provided as S1 Text, the CONSORT statement as S2 Text, and the patient flow is shown in Fig 2C. Twenty-seven patients were screened and 25 eligible patients enrolled at the University of Illinois at Chicago with the first patient starting the study drug on Sept 5, 2012 and the last patient receiving the last dose of the study drug on Jan 6, 2016. Patients met inclusion criteria for high-risk disease despite standard of care, and hydroxyurea treatment
was discontinued at least 28 days prior to initiation of the study drug or placebo [8] (Fig 2D). Placebo-treated patients were older than study drug–treated patients, a difference produced randomly. Median age of the 4 male and 21 female patients was 34 years (range 23–56) (Fig 2E). Baseline laboratory values reflected hemolytic anemia and were similar in drug- and placebo-treated patients (Fig 2E).

**AEs**

The primary endpoint of ≥ grade 3 non-hematologic toxicities was not triggered, nor did any patients trigger the platelets >1,200 × 10^9/L, or neutrophils <1.5 × 10^9/L thresholds requiring dose modification. There were no grade 4 AEs (Fig 3A).

This study was not designed or powered to demonstrate clinical benefit (e.g., reduction in grade 3 AEs from SCD). Such grade 3 AEs unrelated to the study drug occurring in study drug–treated patients were sickle cell vaso-occlusive pain crises (X14) and a pulmonary embolus (Fig 3B). Of these 15 events in study drug–treated patients, 2 events occurred prior to study drug administration (between study enrollment and initiation of drug), and 6 events occurred after discontinuation of study drug, including 3 events in week 3 or 4 of post-drug follow-up (Fig 3B). The pulmonary embolus occurred in a patient with prior history of this complication and without baseline or on-treatment platelet count elevations (a cohort 2 patient). All patients except for one had required emergency room or hospital admission for sickle cell vaso-occlusive pain crises between 1 and 18 times in the 12 months prior to study enrollment (Fig 3B). By including the event occurring prior to study drug administration, the rate of grade 3 sickle cell vaso-occlusive pain crises was statistically significantly increased in cohort 4 versus placebo patients: cohort 4 patients had a rate of this complication in the 12 months preceding study enrollment that was more than 2-fold greater than the rate in placebo-treated patients (Fig 3B).

The most frequent AE of any grade in placebo- and study drug–treated patients was pain from vaso-occlusive sickle cell crisis (Fig 3A). The rates of vaso-occlusive pain crisis AEs of all grades appeared lower in THU-decitabine dose level versus placebo-treated patients except in cohort 4, as noted above (Fig 4). The overall pattern of AEs was similar in placebo- and study drug–treated patients (Fig 4).

No patients discontinued the study drug or placebo because of AEs.

**Decitabine pharmacokinetics**

Samples for decitabine pharmacokinetic measurements by LCMS/MS were obtained immediately prior to and at 2, 4, and 24 hours after decitabine administration in 12 of 15 study drug–treated patients (dictated by venous access). Decitabine was detected in the plasma even at the lowest decitabine dose level of 0.01 mg/kg, and a dose-dependent increase was observed (Fig 5A). The highest dose of oral decitabine, 0.16 mg/kg (cohort 5), produced decitabine plasma concentrations at 2 hours from 39–54 nM (C_max). The dose level below this, 0.08 mg/kg (cohort 4), produced C_max of 9–21 nM (Fig 5A).

**Molecular pharmacodynamics**

The intended molecular pharmacodynamic effect with oral THU-decitabine therapy is DNMT1 depletion. DNMT1 protein levels were measured by flow cytometric analysis of peripheral blood mononuclear cells obtained at baseline, 3, 6, and 8 weeks after initiation of treatment. DNMT1 protein levels decreased by >75% from baseline in 2 of the 3 decitabine 0.16 mg/kg–treated patients (cohort 5), and by approximately 50% in all 0.08 mg/kg–treated patients (cohort 4), but not in patients with lower decitabine doses or in placebo-treated
**Fig 3. Adverse events (AEs).** (A) AEs with possible relatedness to the study drug (AEs other than “unrelated”) per judgement of the treating clinical teams. (B) All grade 3 AEs occurring in study drug—and placebo-treated patients. These grade 3 AEs were sickle cell complications judged to be unrelated to the study drug by the treating clinical teams. (C) Statistical comparison of grade 3 AEs between placebo and decitabine dose-level patients. R function poisson.test(), an exact test, was used. Abbreviation: ED/Hosp, emergency department/hospital.

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patients (Fig 5B). An expected consequence of DNMT1 depletion is reduction in DNA methylation at LINE-1 repetitive element CpGs, a measurement that has been used in other clinical trials of DNMT1-depleting drugs [71]. LINE-1 CpG methylation was measured in peripheral blood mononuclear cells obtained at baseline, 4, 8, and 10 weeks after initiation of therapy.
LINE-1 CpG methylation decreased consistently by approximately 10% with decitabine at 0.16 mg/kg (cohort 5), but not with lower decitabine doses or in placebo-treated patients (Fig 5C).

**Induction of HbF**

The primary efficacy objective with non-cytotoxic DNMT1 depletion in patients with SCD is to increase HbF expression in erythroid precursors, and thereby to decrease HbS polymerization.
and stop the SCD pathophysiological cascade at its inception (Fig 1). HbF was measured by HPLC in the Clinical Pathology Laboratory and also by flow cytometric quantification of HbF content in individual RBCs. HbF percentage (HbF%) increases by HPLC were observed with decitabine 0.08 mg/kg and 0.16 mg/kg (cohorts 4 and 5, respectively), but not with other decitabine dose levels or placebo (Fig 6A–6D, S1 Fig, S1 Data). Rates of increase in HbF% (time-slope estimates) increased with dose ($P < 0.001$) (Fig 6B). The largest HbF increases were in cohort 5 with increases of 4%, 9%, and 9%, corresponding to absolute HbF increases of 0.4, 0.85, and 1.1 g/dL from baseline (Fig 6A–6D). Upon study drug discontinuation, HbF plateaued for 2 weeks and then began to decline (Fig 6A and 6C). By flow cytometry, major increases in RBC enriched for HbF (F-cells) were observed in cohorts 4 and 5, with the largest F-cell increases in cohort 5 patients, up 2.1-, 2.1- and 1.4-fold, thereby reaching levels up to approximately 40%, 65%, and 80% of total RBCs, respectively (Fig 6E and 6F, S2 Fig). F-cells did not increase with placebo (Fig 6E and 6F). F-cells plateaued in the 2 weeks after study drug discontinuation (Fig 6D).

**Total hemoglobin and other hematology/efficacy parameters**

Non-cytotoxic DNMT1 depletion by decitabine in vitro and in vivo is known to bias commitment decisions of multi-potent hematopoietic precursors towards erythroid and megakaryocyte lineage-fate and away from granulocyte-monocyte fate [43,51,53,72]. That is, non-cytotoxic DNMT1 depletion is expected to increase hemoglobin and platelets and concurrently decrease neutrophils. Hemoglobin levels increased in all cohort 4 and 5 (0.08 and 0.16 mg/kg, respectively) patients (in one cohort 4 patient, an isolated large increase in hemoglobin and ANC coincided with a pneumonia diagnosis and dehydration) (Fig 7A, S1 Fig, S1 Data). The total hemoglobin increases from baseline to maximum during 8 weeks of treatment at 0.16 mg/kg dose (cohort 5) were 1.2, 1.8, and 1.9 g/dL (Fig 7A). Rates of increase in total hemoglobin (time-slope estimates) increased with dose ($P = 0.01$) (Fig 7B). Total hemoglobin increased even as reticulocyte counts decreased; that is, better quality and efficiency of HbF-enriched erythropoiesis permitted increases in hemoglobin with fewer reticulocytes (Fig 7C). Also indicating better RBC quality were biomarkers of hemolysis, thrombophilia, and inflammation, namely serum LDH, total bilirubin, D-dimer, and CRP levels, all of which improved in cohorts 4 and 5 (Fig 7D–7G, S1 Fig, S1 Data).

In parallel, as expected with non-cytotoxic DNMT1 depletion, platelets increased concurrently with neutrophil decreases (Fig 7H and 7I). Platelet and neutrophil counts on therapy did not reach a priori–defined triggers for dose modification: neither a platelet count of $>1,200 \times 10^9/L$, nor an ANC $<1.5 \times 10^9/L$, were reached, as the highest platelet count on therapy was $1,122 \times 10^9/L$ in a patient with a pretreatment count of $733 \times 10^9/L$, and the lowest ANC on therapy was $1.6 \times 10^9/L$ in a patient with a pretreatment count of $1.8 \times 10^9/L$ (Fig 7H and 7I). In contrast to the clear trends in myeloid lineages, there was no clear pattern of change in absolute lymphocyte counts on therapy (S1 Fig).

All blood counts reverted towards baseline 2 weeks after discontinuation of study drug (Fig 7).

**Discussion**

This first-in-human clinical trial evaluated safety of oral THU-decitabine, and identified an appropriate dose, for DNMT1 depletion and HbF induction in SCD [70]. Oral THU-decitabine was safe and well-tolerated in this study and produced a wide decitabine concentration-time profile (low $C_{max}$, long $T_{max}$) ideal for non-cytotoxic DNMT1 depletion, since DNMT1 depletion can occur at low nanomolar concentrations but depends on exposure timing [46,73–75]. Increases in fetal and total hemoglobin expected to be clinically significant were produced...
at the highest decitabine dose administered, 0.16 mg/kg, accompanied by improvements in laboratory biomarkers of hemolysis, coagulation, and inflammation. The side effects were a concurrent increase in platelets and decrease in neutrophils, expected with non-cytotoxic DNMT1 depletion.

Fig 6. Fetal hemoglobin (HbF) induction in cohort 4 and 5 patients. (A) HbF percentage (HbF%) over time in cohorts 4 and 5 (decitabine 0.08 mg/kg and 0.16 mg/kg, respectively) patients. (B) Statistical analysis of the rates of change in HbF% with increasing doses of decitabine (0 = placebo); P value determined by linear regression. (C) Change in HbF% from pretreatment to week 8 or 10. (D) Absolute HbF levels. (E) Proportion of red blood cells (RBCs) expressing high levels of HbF (F-cells) in cohort 4 and 5 patients. (F) Raw F-cell flow cytometry data for cohort 5 tetrahydro尿idine (THU)-decitabine—and placebo-treated patients. Abbreviation: FS, forward scatter.

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The main limitation was narrow ability to document clinical benefits, because this was a first-in-human study with a single-blind design, with only a small number of patients treated at each decitabine dose level, and for only 8 weeks. Non-blinding of investigators could produce over- or under-reporting of AEs. The major rationale for not blinding investigators to

Fig 7. Blood counts and other parameters in cohort 4 (decitabine 0.08 mg/kg) and 5 (decitabine 0.16 mg/kg) tetrahydrodouridine (THU)-decitabine–treated patients. (A) Total hemoglobin. (B) Statistical analysis of the rates of change in total hemoglobin (Hbg) with increasing doses of decitabine (0 = placebo); P value determined by linear regression. (C) Absolute reticulocyte counts (ARC). (D, E) LDH and total bilirubin. These are biomarkers of hemolysis. (F) D-dimer. A biomarker of coagulation activation. (G) C-reactive protein (CRP). A biomarker of inflammation. (H) Platelets. (I) Absolute neutrophil counts (ANC).

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treatment assignment was the intent to use intra-patient dose modification for decitabine $C_{\text{max}} > 0.2 \mu M$, study drug–related $\geq$ grade 3 toxicity, platelets $> 1,200 \times 10^9/L$, or neutrophils $< 1.5 \times 10^9/L$. In the study, intra-patient dose modification was not triggered.

Doses of oral decitabine administered after oral THU were escalated from a very low starting level (0.01 mg/kg), to find the minimal decitabine doses active in depleting DNMT1 without cytotoxicity. Our specific pharmacokinetic objectives were to (i) distribute decitabine through the most CDA-enriched tissues, intestines and liver, which need to be surmounted for oral bioavailability; (ii) extend decitabine half-life/T$_{\text{max}}$ to hours, instead of minutes, to increase S-phase–dependent DNMT1 depletion; (iii) avoid off-target cytotoxicity by maintaining $C_{\text{max}}$ in the range of $>5 \text{nM}$ and $<200 \text{nM}$ (nucleotide pool imbalance, DNA-damage, and cytotoxicity correlates with $C_{\text{max}} > 500 \text{nM}$ (reviewed in [76])). These pharmacokinetic objectives were met: mini-doses of oral decitabine administered 60 minutes after THU 10 mg/kg traversed the intestines and liver to produce systemic exposure with a wide decitabine concentration-time profile ($C_{\text{max}}$ approximately 50 nM at 0.16 mg/kg decitabine, T$_{\text{max}}$ of hours instead of minutes). By contrast, intravenous administration of decitabine produces a very high $C_{\text{max}}$ and very brief half-life of minutes. Although continuous intravenous or subcutaneous infusion can in principle lower $C_{\text{max}}$ and extend T$_{\text{max}}$, such approaches are impractical, expensive, and do not solve the problem of negligible distribution into CDA-rich solid tissues —target cells residing in such tissues remain unexposed to treatment [62,77,78]. Increasing intravenous or subcutaneous dose is not a solution for uneven tissue distribution, since toxic exposures are produced in sensitive CDA-poor tissues (e.g., bone marrow) while negligible distribution into CDA-rich tissues persists [62,77]. Similarly, attempting to overcome the intestinal/liver CDA first-pass barrier with high oral doses risks luminal drug concentrations toxic to intestinal enterocytes while systemic exposures remain suboptimal [70,79]. Subcutaneous administration blunts $C_{\text{max}}$ but does not resolve short half-life and uneven tissue distribution problems [70].

The molecular pharmacodynamic objective of DNMT1 depletion and its corollary, hypomethylation of repetitive element LINE-1 CpG, were produced in peripheral blood mononuclear cells most consistently by oral decitabine 0.16 mg/kg (approximately 5 mg/m$^2$). The degree of LINE-1 CpG hypomethylation generated was comparable to that produced by intravenous decitabine regimens FDA-approved to treat myeloid malignancies, that infuse an approximately four-fold higher intravenous daily dose of 20 mg/m$^2$/day for 5 days. However, instead of being pulse-cycled, the hypomethylation with oral THU-decitabine is sustained week to week [80,81]. Also contrasting with intravenous infusion, decitabine exposure was throughout CDA-rich solid tissues, demonstrated by its distribution through intestine and liver into plasma where it was measured. This is clinically pertinent beyond oral bioavailability, since target cells may reside in CDA-rich tissues, e.g., erythroid precursors in the spleen in β-thalassemia [62,77,80].

We previously compared head-to-head in the same non-human primates ($n = 14$) HbF induction by hydroxyurea or cytarabine versus 5-azacytidine (a prodrug of decitabine); 5-azacytidine produced 2 to 20-fold greater increases in HbF than cytarabine or hydroxyurea, without the cytotoxic side effects of cytarabine and hydroxyurea and with consistent effects in older animals less able to respond to cytarabine or hydroxyurea [21]. From this, we repositioned parenterally administered decitabine for non-cytotoxic DNMT1 depletion in SCD patients who had not benefitted from hydroxyurea. In these trials, HbF increases plateaued at $>10\%$ from baseline after 12 weeks of treatment, including in patients with essentially 0% HbF increases after 2 years of monitored treatment with hydroxyurea in the pivotal clinical trial [42,43]. The observations here are in line with these previous observations: 8 weeks of treatment with oral THU-decitabine 0.16 mg/kg 2X/week increased HbF 4%–9% from baseline,
again in patients who had not benefitted from hydroxyurea. F-cells increased by >2-fold, up to approximately 80% of the total RBC population. Total hemoglobin increased 1.2–1.9 g/dL even as reticulocyte counts decreased, reflecting the better efficiency and quality of HbF-enriched erythropoiesis, also shown by improvements in biomarkers of hemolysis, thrombophilia, and inflammation (LDH, bilirubin, D-dimer, CRP). By way of comparison, transfusion of 1 unit of blood increases hemoglobin by approximately 1 g/dL, and hydroxyurea in the pivotal trial increased total hemoglobin by an average of 0.6 g/dL. Previous parenteral decitabine trials evaluated RBC adhesion to thrombospondin and laminin, RBC hemoglobin concentration, and RBC phosphatidylserine exposure, documenting improvements in these parameters as well [43,51].

In parallel, as expected with non-cytotoxic DNMT1 depletion, platelet counts increased while neutrophil counts concurrently decreased [43,51–53], although remaining within ranges observed in SCD or β-thalassemia patients receiving standard-of-care interventions such as splenectomy or hydroxyurea. Platelet increases and neutrophil decreases are dose/frequency-limiting parameters. In a previous clinical trial in which subcutaneous decitabine was administered up to 3X/week, the more frequent administration, expected to produce DNMT1 depletion in a greater fraction of the erythroid precursor population, did produce even larger increases in HbF and total hemoglobin; however, there were correspondingly greater platelet increases and neutrophil decreases [43]. Platelet increases in SCD, already a thrombophilia, are a concern. The trade-off with better RBC quality, however, is worth closer examination. Several groups have found that platelet activation in β-hemoglobinopathies is by diseased RBC and the endothelial damage they cause [82–94]. In thrombophilic myeloproliferative neoplasms also, thrombotic risk has been found to be a function of qualitative defects in RBC and platelets and not higher platelet counts (reviewed in [95]). Similarly, no link between higher platelets and thrombosis was observed in series of post-splenectomy β-thalassemia patients [91,92], nor during 12–15 years of follow-up of >5,000 normal individuals [96]. The improvement in RBC quality produced by HbF induction thus explains improvements in thrombophilia biomarkers (D-dimer, Von Willebrand factor [vWF] propeptide, RBC adhesion to thrombospondin and laminin, SVCA1 [endothelial damage biomarker]) in this and previous studies of decitabine, despite concurrent platelet count increases to >800 × 10^9/L [43]. In short, an association between platelet count and thrombosis risk, though intuitive and possible, is not clear cut in ex and in vivo studies, while qualitative RBC defects, improved by HbF induction, are linked (reviewed in [83]). Future trials of non-cytotoxic DNMT1 depletion will need to continue to carefully monitor the risks/benefits of better RBC quality and higher platelets.

Viscosity that increases with hematocrit also contributes to thrombophilia [97]. That non-cytotoxic DNMT1 depletion produces large increases in total hemoglobin could thus mean that this approach is less suited to SCD subtypes with relatively high hematocrits at baseline, e.g., S-β’-thalassemia or S-C disease, even if RBC quality is improved concurrently by HbF induction. This question will need evaluation in future clinical trials.

Another risk is that high concentrations of decitabine can be DNA damaging and mutagenic and hence potentially pro-oncogenic. DNA damage can also be cytotoxic; this is a concern independent of potential oncogenicity because of the extraordinary demands on SCD bone marrow to compensate for severe hemolytic anemia—dwindling of such compensation by vaso-occlusive damage and age contributes to early death [8,23,31,32,98]. A major rationale for oral THU-decitabine, therefore, is creating decitabine pharmacology that reduces off-target anti-metabolite effects, DNA-damage, and cytotoxicity. First, combination with THU decreases formation of uridine degradation products that contribute significantly to off-target DNA damage and cytotoxicity [49,50]. Second, oral THU-decitabine produces a low C_{max}–
extended T_max concentration-time profile that is conducive to DNMT1 depletion without measurable DNA damage or cytotoxicity, shown extensively in vitro (reviewed in [76]), in non-human primates [21,70], and in clinical trials in which very low doses of decitabine were administered subcutaneously by metronomic schedules [43,51,52]; the clinical trials documented non-cytotoxic DNMT1 depletion by a number of assays including bone marrow evaluation of DNMT1 protein levels, cellularity, γH2AX, and sub-G1 fraction, and peripheral blood evaluation of erythrocyte micronucleus and VDJ recombination assays [43,51,52].

Also regarding concerns of oncogenicity, DNMT1 is highly validated as a molecular target to prevent and treat cancer, offering a p53-independent mechanism of action distinct from conventional anti-metabolite therapy and preserving normal dividing cells (excellent therapeutic index) [43,51–54,76,99–103] (reviewed in [104]). These properties of DNMT1 as a molecular target for oncotherapy likely explain why decitabine and the related 5-azacytidine are the only drugs FDA-approved to treat all subtypes of myelodysplastic syndromes, a type of myeloid malignancy usually seen in the elderly in which better outcomes depend on improving blood counts [52,76,99,100,105,106].

Oral THU-decitabine, like standard-of-care hydroxyurea, could be teratogenic and should be restricted accordingly.

Many patients with myeloid malignancies have been treated chronically for years with metronomic very-low-dose decitabine without significant side effects [52]. Although THU is a new chemical entity that is not FDA-approved for any indication, there have been no side effects observed in clinical trials, some with treatment durations up to and exceeding a year (reviewed in [70,107]). The absence of side effects with THU likely reflects the adaptive network structure of pyrimidine metabolism that is robust to inhibition of CDA; THU has been shown to have no effect on nucleotide pool sizes [108]. Also supporting that its inhibition can be non-detrimental to normal physiology is large natural variation in CDA levels between species [109]. Mostly, THU has been used to try to increase anti-metabolite, DNA-damaging effects of coadministered cytotoxic cytidine analogues, an approach that increases systemic toxicity. Here, by contrast, the intent was to create lower Cmax, extended T_max, more equitable tissue distribution and decrease uridine degradation products of decitabine for non-cytotoxic (non-toxic), molecularly targeted therapy goals (reviewed in [76,99]).

Fetal hemoglobin induction involves chromatin remodeling of the HbF gene locus (HBG) [110]. Cytotoxicity generates this remodeling indirectly, via bone marrow stress [21,110,111] (S3 Fig). The inefficiency of this approach versus directly inhibiting epigenetic enzymes is underscored by greater HbF increases produced by <1/1,000th the molar amount of decitabine versus hydroxyurea in the same patients (decitabine approximately 0.2 mg/kg 2X/week versus hydroxyurea approximately 20 mg/kg daily) [43,112] (S3 Fig). Several other drug development efforts are thus also directed towards direct inhibition of epigenetic enzymes that silence HbF (HBG). Besides DNMT1, enzymes targeted include histone deacetylases (HDAC), KDM1A/LSD1, and PRMT5. DNMT1-depleting and HDAC-inhibiting drugs are already FDA-approved for other clinical indications. Potential application of marketed HDAC inhibitors for HbF induction is limited by pleiotropic roles of HDAC in cells outside of chromatin, rendering it difficult to separate anti-metabolite/cytotoxicity from epigenetic effects [113–117]. The limitations of marketed parenteral decitabine also apply to 5-azacytidine, a decitabine prodrug. By depleting DNMT1 protein, decitabine and 5-azacytidine disrupt its scaffolding functions for other epigenetic enzymes such as LSD1/KDM1A [118,119]. This potent epigenetic effect, and its known safety profile, justify efforts to improve decitabine pharmacology and accessibility.

HbF is the most powerful known natural modulator of the root cause of SCD pathophysiology. This clinical trial affirms that substantial HbF and total hemoglobin increases can be
produced by inhibiting an epigenetic enzyme that mediates HbF silencing and by rationally avoiding cytotoxicity. The oral route of administration and safety profile of the agent used for this purpose, oral THU-decitabine, could have global health implications. Further clinical development and evaluation are thus warranted.

Supporting information

S1 Data. All patient laboratory data.

S1 Fig. Changes in hematologic, hemolysis, thrombophilia, and inflammation biomarkers on therapy. Plots of median values in decitabine dose level and placebo cohorts. (A) HbF%. (B) Total hemoglobin. (C) Absolute reticulocyte counts (ARC). (D) Serum lactate dehydrogenase levels (LDH). (E) Total bilirubin levels. (F) D-dimer levels. (G) Serum C-reactive protein levels (CRP). (H) Platelet counts. (I) Absolute neutrophil counts (ANC). (J) Absolute lymphocyte counts (ALC). (K) Total white blood cell counts (WBC).

S2 Fig. F-cells in cohort 5—Original flow cytometry data. Peripheral blood samples were fixed and stained with phycoerythrin-conjugated anti-HbF (Caltag) per manufacturer’s instructions. Analysis on a Becton-Dickinson FacsCalibur (Sunnyvale, CA).

S3 Fig. Fetal hemoglobin induction requires chromatin remodeling, including DNA hypomethylation, of the HbF gene locus (γ-globin, HBG). Bone marrow stress, e.g., from cytotoxic drugs such as hydroxyurea, can create such remodeling during the recovery phase by surviving erythroid precursors [21,110,111]. An alternative approach is to remodel the HbF locus directly, e.g., by directly inhibiting DNMT1 using decitabine. The relative efficiencies of these approaches are illustrated by the greater HbF increases produced in the same non-human primates or human patients by decitabine approximately 0.2 mg/kg 2X/week, versus hydroxyurea approximately 20 mg/kg daily. That is, the molar amount of decitabine administered per week is <1/1,000th the amount of hydroxyurea administered per week [21,43].

S1 Text. Clinical protocol document.

S2 Text. CONSORT statement.

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Author Contributions

Conceptualization: Joseph DeSimone, Yogen Saunthararajah.

Data curation: Lani Krauz, Sherry Fada, Matthew Rump.
Formal analysis: Donald Lavelle, Vinzon Ibanez, Maria A. Ruiz, Kwok Peng Ng, Philip Woost, Tomas Radivoyevitch, James Jacobberger, Mitch Phelps, Yogen Saunthararajah.

Funding acquisition: Joseph DeSimone, Yogen Saunthararajah.

Investigation: Robert Molokie, Donald Lavelle, Michel Gowhari, Michael Pacini, Johara Hassan, Vinzon Ibanez, Maria A. Ruiz, Kwok Peng Ng, Philip Woost, Matthew Hsieh, John F. Tisdale, James Douglas Engel, Santhosh Saraf, Lewis L. Hsu, Victor Gordeuk, Yogen Saunthararajah.

Methodology: Robert Molokie, Donald Lavelle, Michael Pacini, Lani Krauz, James Jacobberger, Mitch Phelps, Joseph DeSimone, Yogen Saunthararajah.

Project administration: Robert Molokie, Donald Lavelle, Michel Gowhari, Michael Pacini, Lani Krauz, Johara Hassan, Daisy Pacelli, Sherry Fada, Joseph DeSimone, Yogen Saunthararajah.

Resources: Yogen Saunthararajah.

Software: Matthew Rump.

Supervision: Robert Molokie, Donald Lavelle, Michel Gowhari, Michael Pacini, Lani Krauz, Johara Hassan, Daisy Pacelli, Santhosh Saraf.

Validation: Robert Molokie, Donald Lavelle, Philip Woost, Daisy Pacelli.

Writing – original draft: Yogen Saunthararajah.

Writing – review & editing: Robert Molokie, Donald Lavelle, Michel Gowhari, Michael Pacini, Lani Krauz, Johara Hassan, Vinzon Ibanez, Maria A. Ruiz, Kwok Peng Ng, Philip Woost, Tomas Radivoyevitch, Daisy Pacelli, Sherry Fada, Matthew Rump, Matthew Hsieh, John F. Tisdale, James Jacobberger, Mitch Phelps, James Douglas Engel, Santhosh Saraf, Lewis L. Hsu, Victor Gordeuk, Joseph DeSimone, Yogen Saunthararajah.

References

1. Platt OS, Brambilla DJ, Rosse WF, Milner PF, Castro O, Steinberg MH, et al. (1994) Mortality in sickle cell disease. Life expectancy and risk factors for early death. N Engl J Med 330: 1639–1644. https://doi.org/10.1056/NEJM199406093302303 PMID: 7993409

2. Lanzkron S, Carroll CP, Haywood C Jr. (2013) Mortality rates and age at death from sickle cell disease: U.S., 1979–2005. Public Health Reports 128: 110–116. https://doi.org/10.1177/003335491312800206 PMID: 23450875

3. Goldberg MA, Husson MA, Bunn HF (1977) Participation of hemoglobins A and F in polymerization of sickle hemoglobin. J Biol Chem 252: 3414–3421. PMID: 16902

4. Nagel RL, Bookchin RM, Johnson J, Labie D, Wajcman H, Isaac-Sodeye WA, et al. (1979) Structural bases of the inhibitory effects of hemoglobin F and hemoglobin A2 on the polymerization of hemoglobin S. Proc Natl Acad Sci U S A 76: 670–672. PMID: 284392

5. Akinsheye I, Alsultan A, Solovieff N, Ngo D, Baldwin CT, Sebastiani P, et al. (2011) Fetal hemoglobin in sickle cell anemia. Blood 118: 19–27. https://doi.org/10.1182/blood-2011-03-325258 PMID: 21490337

6. Platt OS, Thorington BD, Brambilla DJ, Milner PF, Rosse WF, Vichinsky E, et al. (1991) Pain in sickle cell disease. Rates and risk factors. N Engl J Med 325: 11–16. https://doi.org/10.1056/NEJM199107043250103 PMID: 1710777

7. Powars DR, Weiss JN, Chan LS, Schroeder WA (1984) Is there a threshold level of fetal hemoglobin that ameliorates morbidity in sickle cell anemia? Blood 63: 921–926. PMID: 6200161

8. Steinberg MH, Barton F, Castro O, Pegelow CH, Ballas SK, Kullar A, et al. (2003) Effect of hydroxyurea on mortality and morbidity in adult sickle cell anemia: risks and benefits up to 9 years of treatment. JAMA 289: 1645–1651. https://doi.org/10.1001/jama.289.13.1645 PMID: 12672732

9. Atweh GF, Schechter AN (2001) Pharmacologic induction of fetal hemoglobin: raising the therapeutic bar in sickle cell disease. Curr Opin Hematol 8: 123–130. PMID: 11224687
10. Powars DR, Elliott-Mills DD, Chan L, Niland J, Hiti AL, Opas LM, et al. (1991) Chronic renal failure in sickle cell disease: risk factors, clinical course, and mortality. Ann Intern Med 115: 614–620. PMID: 1892333

11. Lebensburger J, Johnson SM, Askenazi DJ, Rozario NL, Howard TH, Hilliard LM (2011) Protective role of hemoglobin and fetal hemoglobin in early kidney disease for children with sickle cell anemia. Am J Hematol 86: 430–432. https://doi.org/10.1002/ajh.21994 PMID: 21523807

12. Aban I, Baddam S, Hilliard LM, Howard TH, Feig DI, Lebensburger JD (2017) Severe anemia early in life as a risk factor for sickle-cell kidney disease. Blood 129: 385–387. https://doi.org/10.1182/blood-2016-09-738104 PMID: 27919909

13. Conley CL, Weatherall DJ, Richardson SN, Shepard MK, Charache S (1963) Hereditary persistence of fetal hemoglobin: a study of 79 affected persons in 15 Negro families in Baltimore. Blood 21: 261–281. PMID: 14022587

14. Perrine RP, Pembrey ME, John P, Perrine S, Shoup F (1978) Natural history of sickle cell anemia in Saudi Arabs. A study of 270 subjects. Ann Intern Med 88: 1–6. PMID: 619731

15. Ngo DA, Aygun B, Akinsheyeye I, Hankins JS, Bhan I, Luo HY, et al. (2012) Fetal hemoglobin levels and haematological characteristics of compound heterozygotes for haemoglobin S and deletional hereditary persistence of fetal haemoglobin. Br J Haematol 156: 259–264. https://doi.org/10.1111/j.1365-2457.2011.08916.x PMID: 22017641

16. DeSimone J, Biel SI, Heller P (1978) Stimulation of fetal hemoglobin synthesis in baboons by hemolytic and hypoxia. Proc Natl Acad Sci U S A 75: 2937–2940. PMID: 96444

17. DeSimone J, Heller P, Adams JG (1979) Hemopoietic stress and fetal hemoglobin synthesis: comparative studies in vivo and in vitro. Blood 54: 1176–1181. PMID: 115514

18. DeSimone J, Heller P, Amsel J, Usman M (1980) Magnitude of the fetal hemoglobin response to acute hemolytic anemia in baboons is controlled by genetic factors. J Clin Invest 65: 224–226. https://doi.org/10.1172/JCI109654 PMID: 6765958

19. DeSimone J, Heller P, Biel M, Zwiers D (1981) Genetic relationship between fetal Hb levels in normal and erythropoietically stressed baboons. Br J Haematol 49: 175–183. PMID: 6170305

20. DeSimone J, Biel M, Heller P (1982) Maintenance of fetal hemoglobin (HbF) elevations in the baboon by prolonged erythropoietic stress. Blood 60: 519–523. PMID: 6178456

21. Lavelle D, DeSimone J, Heller P, Zwiers D, Hall L (1986) On the mechanism of Hb F elevations in the baboon by erythropoietic stress and pharmacologic manipulation. Blood 67: 1083–1089. PMID: 2420395

22. Charache S, Terrin ML, Moore RD, Dover GJ, Barton FB, Eckert SV, et al. (1995) Effect of hydroxyurea on the frequency of painful crises in sickle cell anemia. Investigators of the Multicenter Study of Hydroxyurea in Sickle Cell Anemia. N Engl J Med 332: 1317–1322. https://doi.org/10.1056/NEJM19950518332201 PMID: 7715639

23. Steinberg MH, Lu ZH, Barton FB, Terrin ML, Charache S, Dover GJ (1997) Fetal hemoglobin in sickle cell anemia: determinants of response to hydroxyurea. Multicenter Study of Hydroxyurea. Blood 89: 1078–1086. PMID: 9028341

24. Ballas SK, Marcolina MJ, Dover GJ, Barton FB (1999) Erythropoietic activity in patients with sickle cell anaemia before and after treatment with hydroxyurea. Br J Haematol 105: 491–496. PMID: 10233426

25. Charache S, Dover GJ, Moore RD, Eckert S, Ballas SK, Koshy M, et al. (1992) Hydroxyurea: effects on hemoglobin F production in patients with sickle cell anemia. Blood 79: 2555–2565. PMID: 1375104

26. Ballas SK, Barton FB, Waclawiw MA, Swerdlow P, Eckman JR, Pegelow CH, et al. (2006) Hydroxyurea and sickle cell anemia: effect on quality of life. Health and Quality of Life Outcomes 4: 59. https://doi.org/10.1186/1477-7525-4-59 PMID: 16942629

27. Segal JB, Strouse JJ, Beach MC, Haywood C, Witkop C, Park H, et al. (2008) Hydroxyurea for the treatment of sickle cell disease. Evidence Report/technology Assessment: 1–95.

28. Machado RF, Anthi A, Steinberg MH, Bonds D, Sachdev V, Kato GJ, et al. (2006) N-terminal pro-brain natriuretic peptide levels and risk of death in sickle cell disease. JAMA 296: 310–318. https://doi.org/10.1001/jama.296.3.310 PMID: 16849664

29. Ware RE, Eggleton B, Redding-Lallinger R, Wang WC, Smith-Whitley K, Daeschner C, et al. (2002) Predictors of fetal hemoglobin response in children with sickle cell anemia receiving hydroxyurea therapy. Blood 99: 10–14. PMID: 11756146

30. Green NS, Manwani D, Qureshi M, Ireland K, Sinha A, Smaldone AM (2016) Decreased fetal hemoglobin over time among youth with sickle cell disease on hydroxyurea is associated with higher urgent hospital use. Pediatric Blood & Cancer 63: 2146–2153.
31. West MS, Wethers D, Smith J, Steinberg M (1992) Laboratory profile of sickle cell disease: a cross-sectional analysis. The Cooperative Study of Sickle Cell Disease. J Clin Epidemiol 45: 893–909. PMID: 1624972

32. Saraf S, Farooqui M, Infusino G, Oza B, Sidhwani S, Gowhari M, et al. (2011) Standard clinical practice underestimates the role and significance of erythropoietin deficiency in sickle cell disease. Br J Haematol 153: 386–392. https://doi.org/10.1111/j.1365-2141.2010.08479.x PMID: 21418176

33. van der Ploeg LH, Flavell RA (1980) DNA methylation in the human gamma delta beta-globin locus in erythroid and nonerythroid tissues. Cell 19: 947–958. PMID: 6247075

34. Mavilio F, Giampaolo A, Care A, Migliaccio G, Calandrini M, Russo G, et al. (1983) Molecular mechanisms of human hemoglobin switching: selective undermethylation and expression of globin genes in embryonic, fetal, and adult erythroblasts. Proc Natl Acad Sci U S A 80: 6907–6911. PMID: 6316333

35. Tagle DA, Koop BF, Goodman M, Slightom JL, Hess DL, Jones RT (1988) Embryonic epsilon and gamma globin genes of a prosimian primate (Galago crassicaudatus). Nucleotide and amino acid sequences, developmental regulation and phylogenetic footprints. J Mol Biol 203: 439–455. PMID: 3199442

36. DeSimone J, Heller P, Hall L, Zwiers D (1982) 5-Azacytidine stimulates fetal hemoglobin synthesis in anemic baboons. Proc Natl Acad Sci U S A 79: 4428–4431. PMID: 6181507

37. Ley TJ, DeSimone J, Anagnou NP, Keller GH, Humphries RK, Turner PH, et al. (1982) 5-azacytidine selectively increases gamma-globin synthesis in a patient with beta+ thalassemia. N Engl J Med 307: 1469–1475. https://doi.org/10.1056/NEJM198212093072401 PMID: 6183586

38. Ley TJ, DeSimone J, Noguchi CT, Turner PH, Schechter AN, Heller P, et al. (1983) 5-Azacytidine increases gamma-globin synthesis and reduces the proportion of dense cells in patients with sickle cell anemia. Blood 62: 370–380. PMID: 6191799

39. Charache S, Dover G, Smith K, Talbot CC Jr., Moyer M, Boyer S (1983) Treatment of sickle cell anemia with 5-azacytidine results in increased fetal hemoglobin production and is associated with nonrandom hypomethylation of DNA around the gamma-delta-beta-globin gene complex. Proc Natl Acad Sci U S A 80: 4842–4846. PMID: 6192443

40. Dover GJ, Charache S, Boyer SH, Vogelsang G, Moyer M (1985) 5-Azacytidine increases HbF production and reduces anemia in sickle cell disease: dose-response analysis of subcutaneous and oral dosage regimens. Blood 66: 527–532. PMID: 2411310

41. Lowrey CH, Nienhuis AW (1993) Brief report: treatment with azacitidine of patients with end-stage beta-thalassemia. N Engl J Med 329: 845–848. https://doi.org/10.1056/NEJM199309163291205 PMID: 7689171

42. Koshy M, DeSimone J, Molokie R, Dorn L, van der Galien T, Bressler L (1998) Augmentation of fetal hemoglobin (HbF) levels by low-dose short-duration 5'-aza-2'-deoxycytidine (decitabine) administration in sickle cell anemia patients who had no HbF elevation following hydroxyurea therapy. Blood 92: 30b–30b.

43. Saunthararajah Y, Hillery CA, Lavelle D, Molokie R, Dorn L, Bressler L, et al. (2003) Effects of 5-aza-2'-deoxycytidine on fetal hemoglobin levels, red cell adhesion, and hematopoietic differentiation in patients with sickle cell disease. Blood 102: 3865–3870. https://doi.org/10.1182/blood-2003-05-1738 PMID: 12907443

44. Xu J, Bauer DE, Kerenyi MA, Vo TD, Hou S, Hsu YJ, et al. (2013) Corepressor-dependent silencing of fetal hemoglobin expression by BCL11A. Proc Natl Acad Sci U S A 110: 6518–6523. https://doi.org/10.1073/pnas.1303976110 PMID: 23576758

45. Cui S, Kolodzije K, Obara N, Amaral-Psarris A, Demmers J, Shi L, et al. (2011) Nuclear receptors TR2 and TR4 recruit multiple epigenetic transcriptional corepressors that associate specifically with the embryonic beta-type globin promoters in differentiated adult erythroid cells. Mol Cell Biol 31: 3298–3311. https://doi.org/10.1128/MCB.05310-11 PMID: 21670149

46. Santi DV, Garrett CE, Barr PJ (1983) On the mechanism of inhibition of DNA-cytosine methyltransferases by cytotoxic analogs. Cell 33: 9–10. PMID: 6205762

47. Covey JM, D’Incalci M, Titchen EH, Zaharko DS, Kohn KW (1986) Differences in DNA damage produced by incorporation of 5-aza-2'-deoxycytidine or 5,6-dihydro-5-azacytidine into DNA of mammalian cells. Cancer Res 46: 5511–5517. PMID: 2428479

48. Schemelhiah L, Haemmer A, Spada F, Rosing N, Meilingr D, Rothbauer U, et al. (2007) Dynamics of Dnmt1 interaction with the replication machinery and its role in postreplicative maintenance of DNA methylation. Nucleic Acids Res 35: 4301–4312. https://doi.org/10.1093/nar/gkm432 PMID: 17576694

49. Zauri M, Berridge G, Thezenas ML, Pugh KM, Goldin R, Kessler BM, et al. (2015) CDA directs metabolism of epigenetic nucleosides revealing a therapeutic window in cancer. Nature 524: 114–118. https://doi.org/10.1038/nature14948 PMID: 26200337
50. Almqvist H, Axelsson H, Jafari R, Dan C, Mateus A, Haraldsson M, et al. (2016) CETSA screening identifies known and novel thymidylate synthase inhibitors and slow intracellular activation of 5-fluoro-uracil. Nat Commun 7: 11040. https://doi.org/10.1038/ncomms11040 PMID: 27010513

51. Olivieri NF, Saunthararajah Y, Thayalasuthan V, Kwatkiowski J, Ware RE, Kuypers FA, et al. (2011) A pilot study of subcutaneous decitabine in beta-thalassemia intermedia. Blood 118: 2708–2711. https://doi.org/10.1182/blood-2011-03-341909 PMID: 21700776

52. Saunthararajah Y, Sekeres M, Advani A, Mahfouz R, Durkin L, Radivoyevitch T, et al. (2015) Evaluation of noncytotoxic DNMT1-depleting therapy in patients with myelodysplastic syndromes. J Clin Invest 125: 1043–1055. https://doi.org/10.1172/JCI78789 PMID: 25621498

53. Sauntharajah Y, Sekeres M, Mahfouz R, Durkin L, Radivoyevitch T, et al. (2015) Evaluation of noncytotoxic DNMT1-depleting therapy in patients with myelodysplastic syndromes. J Clin Invest 125: 1043–1055. https://doi.org/10.1172/JCI78789 PMID: 25621498

54. Milhem M, Mahmud N, Lavelle D, Araki H, DeSimone J, Sauntharajah Y, et al. (2004) Modification of hematopoietic stem cell fate by 5aza-2’-deoxycytidine and trichostatin A. Blood 103: 4102–4110. https://doi.org/10.1182/blood-2003-07-2431 PMID: 14976039

55. Hu Z, Negrotto S, Gu X, Mahfouz R, Ng KP, Ebrahem Q, et al. (2010) Decitabine maintains hematopoietic progenitor self-renewal by preventing repression of stem cell genes by a differentiation-inducing stimulus. Mol Cancer Ther 9: 1536–1543. https://doi.org/10.1158/1535-7163.MCT-10-0191 PMID: 20501800

56. DeSimone J, Koshy M, Dorn L, Lavelle D, Bressier L, Molokie R, et al. (2002) Maintenance of elevated fetal hemoglobin levels by decitabine during dose interval treatment of sickle cell anemia. Blood 99: 3905–3908. PMID: 12010787

57. Liu Z, Marcucci G, Byrd JC, Grever M, Xiao J, Chan KK (2006) Characterization of decomposition products and preclinical and low dose clinical pharmacokinetics of decitabine (5-aza-2’-deoxycytidine) by a new liquid chromatography/tandem mass spectrometry quantification method. Rapid Commun Mass Spectrom 20: 1117–1126. https://doi.org/10.1002/rcm.2423 PMID: 16523529

58. Li, Marcucci G, Byrd JC, Grever M, Xiao J, Chan KK (2006) Characterization of decomposition products and preclinical and low dose clinical pharmacokinetics of decitabine (5-aza-2’-deoxycytidine) by a new liquid chromatography/tandem mass spectrometry quantification method. Rapid Commun Mass Spectrom 20: 1117–1126. https://doi.org/10.1002/rcm.2423 PMID: 16523529

59. Neil GL, Moxley TE, Kuentzel SL, Manak RC, Hanka LJ (1975) Enhancement by tetrahydrouridine (NSC-112907) of the oral activity of 5-azacytidine (NSC-102816) in L1210 leukemic mice. Cancer Chemother Rep 59: 459–465. PMID: 54211

60. DeSimone J, Heller P, Molokie RE, Hall L, Zwiers D (1985) Tetrahydrouridine, cytidine analogues, and hemoglobin F. Am J Hematol 18: 283–288. PMID: 2579548

61. Beumer JH, Eiseman JL, Parise RA, Joseph E, Covey JM, Egorin MJ (1982) Influence of tetrahydrouridine on the pharmaco kinetics of intrathecally administered 1-beta-D-arabinofuranosylcytosine. Cancer Res 42: 1736–1739. PMID: 6896013

62. Kreis W, Budman DR, Chan K, Allen SL, Schulman P, Lichtman S, et al. (1991) Therapy of refractory/relapsed acute leukemia with cytosine arabinoside plus tetrahydrouridine (an inhibitor of cytidine deaminase)—a pilot study. Leukemia 5: 991–998. PMID: 1961042

63. Wong PP, Currie VE, Mackey RW, Krakoff IH, Tan CT, Burchenal JH, et al. (1979) Phase I evaluation of tetrahydrouridine combined with cytosine arabinoside. Cancer Treat Rep 63: 1245–1249. PMID: 383291

64. Ho DH, Bodey GP, Hall SW, Benjamin RS, Brown NS, Freireich EJ, et al. (1978) Clinica, pharmacology of tetrahydrouridine. J Clin Pharmacol 18: 259–265. PMID: 641214

65. Kreis W, Woodcock TM, Gordon CS, Krakoff IH (1977) Tetrahydrouridine: Physiologic disposition and effect upon deamination of cytosine arabinoside in man. Cancer Treat Rep 61: 1347–1353. PMID: 589600

66. Marsh JH, Kreis W, Barile B, Akerman S, Schulman P, Allen SL, et al. (1993) Therapy of refractory/relapsed acute myeloid leukemia and blast crisis of chronic myeloid leukemia with the combination of cytosine arabinoside, tetrahydrouridine, and carboplatin. Cancer Chemother Pharmacol 31: 481–484. PMID: 8453689
69. Goldenthal EI, Cookson KM, Geil RG, Wazeter FX (1974) Preclinical toxicologic evaluation of tetrahydouridine (NSC-112907) in beagle dogs and rhesus monkeys. Cancer Chemother Rep 5: 15–16.

70. Lavelle D, Vaitkus K, Ling Y, Ruiz MA, Mahfouz R, Ng KP, et al. (2012) Effects of tetrahydouridine on pharmacokinetics and pharmacodynamics of oral decitabine. Blood 119: 1240–1247. https://doi.org/10.1182/blood-2011-08-371690 PMID: 22160381

71. Aparicio A, North B, Barske L, Wang X, Bollati V, Weisenberger D, et al. (2009) LINE-1 methylation in plasma DNA as a biomarker of activity of DNA methylation inhibitors in patients with solid tumors. Epigenetics 4: 176–184. PMID: 19421002

72. Saunthararajah Y, Sekeres M, Advani A, Mahfouz R, Durklin L, Radiyovitch T, et al. (2015) Evaluation of noncytotoxic DNMT1-depleting therapy in patients with myelodysplastic syndromes. J Clin Invest. 2; 125(3):1043–55. https://doi.org/10.1172/JCI78789 PMID: 25621498

73. Santi DV, Norment A, Garrett CE (1984) Covalent bond formation between a DNA-cytosine methyltransferase and DNA containing 5-azacytosine. Proc Natl Acad Sci U S A 81: 6993–6997. PMID: 6209710

74. Patel K, Dickson J, Din S, Macleod K, Jodrell D, Ramsahoye B (2010) Targeting of 5-aza-2'-deoxycytidine residues by chromatin-associated DNMT1 induces proteasomal degradation of the free enzyme. Nucleic Acids Res 38: 4313–4324. https://doi.org/10.1093/nar/gkq187 PMID: 20348135

75. Creusot F, Acs G, Christman JK (1982) Inhibition of DNA methyltransferase and induction of Friend erythroleukemia cell differentiation by 5-azacytidine and 5-aza-2'-deoxycytidine. J Biol Chem 257: 2041–2048. PMID: 6173384

76. Saunthararajah Y (2013) Key clinical observations after 5-azacytidine and decitabine treatment of myelodysplastic syndromes suggest practical solutions for better outcomes. Hematol Am Soc Hematol Educ Program 2013: 511–521. https://doi.org/10.1182/asheducation-2013.1.511 PMID: 24319226

77. Schrump DS, Fischette MR, Nguyen DM, Zhao M, Li X, Kunst TF, et al. (2006) Phase I study of decitabine-mediated gene expression in patients with cancers involving the lungs, esophagus, or pleura. Clin Cancer Res 12: 5777–5785. https://doi.org/10.1158/1078-0432.CCR-06-0669 PMID: 17020984

78. Jansen RS, Rosing H, Wijermans PW, Keizer RJ, Schellens JH, Beijnen JH (2012) Decitabine triphosphate levels in peripheral blood mononuclear cells from patients receiving prolonged low-dose decitabine administration: a pilot study. Cancer Chemotherapy and Pharmacol 69: 1457–1466.

79. Garcia-Manero G, Gore SD, Cogle C, Ward R, Shi T, Macbeth KJ, et al. (2011) Phase I study of oral azacitidine in myelodysplastic syndromes, chronic myelomonocytic leukemia, and acute myeloid leukemia. J Clin Oncol 29: 2521–2527. https://doi.org/10.1200/JCO.2010.34.4226 PMID: 21576646

80. Stewart DJ, Issa JP, Kurzrock R, Nunez MI, Jelinek J, Hong D, et al. (2009) Decitabine effect on tumor global DNA methylation and other parameters in a phase I trial in refractory solid tumors and lymphomas. Clin Cancer Res 15: 3881–3888. https://doi.org/10.1158/1078-0432.CCR-08-2196 PMID: 19470736

81. Fang F, Balch C, Schilder J, Breen T, Zhang S, Shen C, et al. (2010) A phase 1 and pharmacodynamic study of decitabine in combination with carboplatin in patients with recurrent, platinum-resistant, epithelial ovarian cancer. Cancer 116: 4043–4053. https://doi.org/10.1002/cncr.25204 PMID: 20564122

82. Goldschmidt N, Spectre G, Brill A, Zelig O, Goldfarb A, Rachmilewitz E, et al. (2008) Increased platelet adhesion under flow conditions is induced by both thalassemic platelets and red blood cells. Thrombosis and Haemostasis 100: 864–870. PMID: 18989531

83. Zhang D, Xu C, Manwani D, Frenette PS (2016) Neutrophils, platelets, and inflammatory pathways at the nexus of sickle cell disease pathophysiology. Blood 127: 801–809. https://doi.org/10.1182/blood-2015-09-618538 PMID: 26758915

84. Habib A, Kunzelmann C, Shamseddien W, Zobairi F, Freyssinet JM, Taher A (2008) Elevated levels of circulating procoagulant microparticles in patients with beta-thalassemia intermedia. Haematologica 93: 941–942. https://doi.org/10.3324/haematol.12460 PMID: 18460647

85. Gladwin MT, Kato GJ (2006) Hemolysis-associated hypercoagulability in sickle cell disease: the plot (and blood) thickens! Haematologica 93: 1–3. https://doi.org/10.3324/haematol.12318 PMID: 18166776

86. Ataga KI, Moore CG, Hillery CA, Jones S, Whinna HC, Strayhorn D, et al. (2008) Coagulation activation and inflammation in sickle cell disease-associated pulmonary hypertension. Haematologica 93: 20–26. https://doi.org/10.3324/haematol.11763 PMID: 18166781

87. Wheelihan MF, Zachary V, Orfeo T, Mann KG (2012) Prothrombin activation in blood coagulation: the erythrocyte contribution to thrombin generation. Blood 120: 3837–3845. https://doi.org/10.1182/blood-2012-05-427856 PMID: 22966460
88. Ruf A, Pick M, Deutsch V, Patschke H, Goldfarb A, Rachmilewitz EA, et al. (1997) In-vivo platelet activation correlates with red cell anionic phospholipid exposure in patients with beta-thalassaemia major. Br J Haematol 98: 51–56. PMID: 9233563

89. Tripodi A, Cappellini MD, Chantarangkul V, Padovan L, Fasulo MR, Marcon A, et al. (2009) Hypercoaguibility in splenectomized thalassemic patients detected by whole-blood thromboelastometry, but not by thrombin generation in platelet-poor plasma. Haematologica: 94: 1520–1527. https://doi.org/10.3324/haematol.2009.010546 PMID: 19648162

90. Whelihan MF, Mooberry MJ, Zachary V, Bradford RL, Ataga KI, Mann KG, et al. (2013) In-vivo platelet aggregation by decitabine of melanoma differentiation in vitro and in vivo. Int J Cancer 131: 18–29. https://doi.org/10.1002/ijc.29124 PMID: 23798841

91. Chunchanuenee S, Archararin N, Ungkanont A, Jootar S, Angchaissrisi P, Bunyaratavej A, et al. (2000) Etiology and incidence of thrombotic and hemorrhagic disorders in Thai patients with extreme thrombocytosis. J Med Assoc Thai 83 Suppl 1: S95–100.

92. Hathirat P, Mahaphan W, Chuansumrit A, Pintadit P, Sasanakul W, Isarangkura P (1993) Platelet counts in thalassemic children before and after splenectomy. Southeast Asian J Trop Med Public Health 24 Suppl 1: 213–215.

93. Setty BN, Kulkarni S, Rao AK, Stuart MJ (2000) Fetal hemoglobin in sickle cell disease: relationship to erythrocyte phosphatidylserine exposure and coagulation activation. Blood 96: 1119–1124. PMID: 10910931

94. Setty BN, Kulkarni S, Dampier CD, Stuart MJ (2001) Fetal hemoglobin in sickle cell anemia: relationship to erythrocyte adhesion markers and adhesion. Blood 97: 2568–2573. PMID: 11313243

95. Tefferi A (2008) Platelet count in essential thrombocythemia: the more the better? Blood 112: 3526–3527. https://doi.org/10.1182/blood-2006-06-036088 PMID: 1840727

96. van der Bom JG, Heckbert SR, Lumley T, Holmes CE, Cushman M, Folsom AR, et al. (2009) Platelet counts and the risk for thrombosis and death in the elderly. J Thromb Haemost 7: 399–405. https://doi.org/10.1111/j.1538-7836.2008.03267.x PMID: 19143922

97. Sverdlow PS (2006) Red cell exchange in sickle cell disease. Hematology Am Soc Hematol Educ Program: 48–53. https://doi.org/10.1182/asheducation-2006.1.48 PMID: 17124039

98. de Haan G, Nijhof W, Van Zant G (1997) Mouse strain-dependent changes in frequency and proliferation of hematopoietic stem cells during aging: correlation between lifespan and cycling activity. Blood 89: 1543–1550. PMID: 9057635

99. Saunthararajah Y, Triozzi P, Rini B, Singh A, Radivoyevitch T, Sekeres M, et al. (2012) p53-independent, normal stem cell sparing epigenetic differentiation therapy for myeloid and other malignancies. Semin Oncol 39: 97–108. https://doi.org/10.1053/j.seminoncol.2011.11.011 PMID: 22829496

100. Ng KP, Ebrahim Q, Negrotto S, Mahfouz RZ, Link KA, Hu Z, et al. (2011) p53 independent epigenetic-differentiation treatment in xenotransplant models of acute myeloid leukemia. Leukemia 25: 1739–1750. https://doi.org/10.1038/leu.2011.159 PMID: 21701495

101. Negrotto S, Ng KP, Jankowska AM, Bodo J, Gopalan B, Guinta K, et al. (2012) CpG methylation patterns and decitabine treatment response in acute myeloid leukemia cells and normal hematopoietic precursors. Leukemia 26: 244–254. https://doi.org/10.1038/leu.2011.207 PMID: 21836612

102. Tsai HC, Li H, Van Neste L, Cai Y, Robert C, Rassool FV, et al. (2012) Transient Low Doses of DNAdamethylating Agents Exert Durable Antitumor Effects on Hematological and Epithelial Tumor Cells. Cancer Cell 21: 430–446. https://doi.org/10.1016/j.ccr.2012.02.039 PMID: 22439938

103. Momparler RL, Cote S, Momparler LF (2013) Epigenetic action of decitabine (5-aza-2'-deoxycytidine) is more effective against acute myeloid leukemia than cytotoxic action of cytarabine (ARA-C). Leuk Res. 37: 980–984. https://doi.org/10.1016/j.leukres.2013.04.019 PMID: 23660386

104. Velcheli V, Radivoyevitch T, Saunthararajah Y (2017) Higher-Level Pathway Objectives of Epigenetic Therapy: A Solution to the p53 Problem in Cancer. Am Soc Clin Oncol Educ Book 37: 812–824. https://doi.org/10.14694/EDBK_174175 PMID: 28561650

105. Negrotto S, Hu Z, Alcazor O, Ng KP, Triozzi P, Lindner D, et al. (2011) Non-cytotoxic differentiation treatment of renal cell cancer. Cancer Res 71: 1431–1441. https://doi.org/10.1158/0008-5472.CAN-10-2422 PMID: 21303982

106. Alcazor O, Achberger S, Aldrich W, Hu Z, Negrotto S, Saunthararajah Y, et al. (2012) Epigenetic regulation by decitabine of melanoma differentiation in vitro and in vivo. Int J Cancer 131: 18–29. https://doi.org/10.1002/ijc.26320 PMID: 21796622

107. Terse P, Engelke K, Chan K, Ling Y, Sharpnack D, Saunthararajah Y, et al. (2014) Subchronic oral toxicity study of decitabine in combination with tetrahydrouridine in CD-1 mice. Int J of Toxicol 33: 75–85.
108. Heinemann V, Plunkett W (1989) Modulation of deoxynucleotide metabolism by the deoxycytidylic deaminase inhibitor 3,4,5,6-tetrahydrodeoxyuridine. Biochem Pharmacol 38: 4115–4121. PMID: 2688654

109. Dedrick RL, Forrester DD, Cannon JN, el-Dareer SM, Mellett LB (1973) Pharmacokinetics of 1-beta-D-arabinofuranosylcytosine (ARA-C) deamination in several species. Biochem Pharmacol 22: 2405–2417. PMID: 4200888

110. Goren A, Simchen G, Fibach E, Szabo PE, Tanimoto K, Chakalova L, et al. (2006) Fine tuning of globin gene expression by DNA methylation. PLoS ONE 1(1): e46. https://doi.org/10.1371/journal.pone.0000046 PMID: 17183675

111. Platt OS, Orkin SH, Dover G, Beardsley GP, Miller B, Nathan DG (1984) Hydroxyurea enhances fetal hemoglobin production in sickle cell anemia. J Clin Invest 74: 652–656. https://doi.org/10.1172/JCI111464 PMID: 6205021

112. Saunthararajah Y, Lavelle D, DeSimone J (2004) DNA hypo-methylating agents and sickle cell disease. Brit J Haematol 126: 629–636.

113. Scuto A, Kirschbaum M, Kowolik C, Kretzner L, Juhasz A, Atadja P, et al. (2008) The novel histone deacetylase inhibitor, LBH589, induces expression of DNA damage response genes and apoptosis in Ph- acute lymphoblastic leukemia cells. Blood 111: 5093–5100. https://doi.org/10.1182/blood-2007-10-117762 PMID: 18349321

114. Lee JH, Choy ML, Ngo L, Foster SS, Marks PA (2010) Histone deacetylase inhibitor induces DNA damage, which normal but not transformed cells can repair. Proc Natl Acad Sci U S A 107: 14639–14644. https://doi.org/10.1073/pnas.1008522107 PMID: 20679231

115. Conti C, Leo E, Eichler GS, Sordet O, Martin MM, Fan A, et al. (2010) Inhibition of histone deacetylase in cancer cells slows down replication forks, activates dormant origins, and induces DNA damage. Cancer Res 70: 4470–4480. https://doi.org/10.1158/0008-5472.CAN-09-3028 PMID: 20460513

116. Gaymes TJ, Padua RA, Pla M, Orr S, Omidvar N, Chomienne C, et al. (2006) Histone deacetylase inhibitors (HDI) cause DNA damage in leukemia cells: a mechanism for leukemia-specific HDI-dependent apoptosis? Mol Cancer Res 4: 563–573. https://doi.org/10.1158/1541-7786.MCR-06-0111 PMID: 16877702

117. Minucci S, Pelicci PG (2006) Histone deacetylase inhibitors and the promise of epigenetic (and more) treatments for cancer. Nat Rev Cancer 6: 38–51. https://doi.org/10.1038/nrc1779 PMID: 16397526

118. Clements EG, Mohammad HP, Leadem BR, Easwaran H, Cai Y, Van Neste L, et al. (2012) DNMT1 modulates gene expression without its catalytic activity partially through its interactions with histone-modifying enzymes. Nucleic Acids Res 40: 4334–4346. https://doi.org/10.1093/nar/gks031 PMID: 22278882

119. Brenner C, Luciani J, Bize M, Ndlovu M, Josseaux E, Dedeuvaerder S, et al. (2016) The interplay between the lysine demethylase KDM1A and DNA methyltransferases in cancer cells is cell cycle dependent. Oncotarget 7: 58939–58952. https://doi.org/10.18632/oncotarget.10624 PMID: 27449299