

1-1-2018

Mortality in children, adolescents and adults with sickle cell anemia in Rio de Janeiro, Brazil.

Clarisse Lopes de Castro Lobo

Instituto de Hematologia Arthur de Siqueira Cavalcanti (HEMORIO)

Emilia Matos do Nascimento

Instituto de Hematologia Arthur de Siqueira Cavalcanti (HEMORIO); Centro Universitário Estadual da Zona Oeste (UEZO)

Leonardo José Carvalho de Jesus

Instituto de Hematologia Arthur de Siqueira Cavalcanti (HEMORIO)

Thiago Gotelip de Freitas

Instituto de Hematologia Arthur de Siqueira Cavalcanti (HEMORIO)

Jocemir Ronaldo Lugon

Follow this and additional works at: https://jdc.jefferson.edu/cardeza_foundation
Universidade Federal Fluminense (UFF)



Part of the [Hematology Commons](#)

[Let us know how access to this document benefits you](#)

See next page for additional authors

Recommended Citation

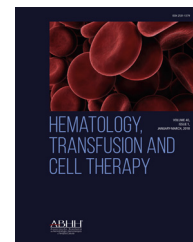
Lobo, Clarisse Lopes de Castro; Nascimento, Emilia Matos do; Jesus, Leonardo José Carvalho de; Freitas, Thiago Gotelip de; Lugon, Jocemir Ronaldo; and Ballas, Samir K., "Mortality in children, adolescents and adults with sickle cell anemia in Rio de Janeiro, Brazil." (2018). *Cardeza Foundation for Hematologic Research*. Paper 46.

https://jdc.jefferson.edu/cardeza_foundation/46

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's [Center for Teaching and Learning \(CTL\)](#). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Cardeza Foundation for Hematologic Research by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.

Authors

Clarisse Lopes de Castro Lobo, Emilia Matos do Nascimento, Leonardo José Carvalho de Jesus, Thiago Gotelip de Freitas, Jocemir Ronaldo Lugon, and Samir K. Ballas



Original article

Mortality in children, adolescents and adults with sickle cell anemia in Rio de Janeiro, Brazil



Clarisse Lopes de Castro Lobo^a, Emilia Matos do Nascimento^{a,b}, Leonardo José Carvalho de Jesus^a, Thiago Gotelip de Freitas^a, Jocemir Ronaldo Lugon^c, Samir K. Ballas^{a,d,*}

^a Instituto de Hematologia Arthur de Siqueira Cavalcanti (HEMORIO), Rio de Janeiro, RJ, Brazil

^b Centro Universitário Estadual da Zona Oeste (UEZO), Rio de Janeiro, RJ, Brazil

^c Universidade Federal Fluminense (UFF), Niteroi, RJ, Brazil

^d Cardeza Foundation, Sidney Kimmel Medical College, Thomas Jefferson University, Philadelphia, United States

ARTICLE INFO

Article history:

Received 8 July 2017

Accepted 22 September 2017

Available online 6 December 2017

Keywords:

Sickle cell anemia
Vaso-occlusive crisis
Mortality
Newborn screening
Survival

ABSTRACT

Objective: To determine the mortality rate of children, adolescents and adults with sickle cell anemia in Rio de Janeiro, Brazil.

Methods: The number of deaths, the mortality rate and the causes of deaths in patients with sickle cell anemia who were treated and followed up at our institution for 15 years were determined and compared to data available for the Brazilian population.

Results: The overall number of deaths was 281 patients with a mortality rate of 16.77%. Survival probability was significantly higher in females. The number of deaths and the mortality rate were age-specific with a significant increase in the 19- to 29-year-old age group. The remaining life expectancy of the patients with sickle cell anemia was less than that of Brazilians at large. The gap between the two was about 20 years for ages between one and five years with this gap decreasing to ten years after the age of 65 years. The most common causes of death were infection, acute chest syndrome, overt stroke, organ damage and sudden death during painful crises.

Conclusion: To the best of our knowledge, this is the first Brazilian study in a single institution in Rio de Janeiro; the mortality rate was 18.87% among adult patients with sickle cell anemia. The mortality rates in children and adults are higher than those reported in developed countries of the northern hemisphere.

© 2017 Associação Brasileira de Hematologia, Hemoterapia e Terapia Celular. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

* Corresponding author at: Cardeza Foundation, Department of Medicine, Sidney Kimmel Medical College, Thomas Jefferson University, 1020 Locust Street, Philadelphia, PA 19107, United States.

E-mail address: samir.ballas@jefferson.edu (S.K. Ballas).

<https://doi.org/10.1016/j.bjhh.2017.09.006>

2531-1379/© 2017 Associação Brasileira de Hematologia, Hemoterapia e Terapia Celular. Published by Elsevier Editora Ltda. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Introduction

The mortality rate among infants, children and young adults with sickle cell disease (SCD) in general and sickle cell anemia (SCA) in particular is high. The overall death rates in under 20-year-old children reported by the Cooperative Study of Sickle Cell Disease (CSSCD) about 30 years ago were 2.6% for patients with all types of SCD and 3.3% for patients with SCA.¹ Sickle cell disease-related mortality decreased significantly from 1999 to 2009 in the USA in all pediatric age groups up to age 19 years.² This decline in mortality was the highest for children younger than one year where the decrease was 61% less than the mortality rate during 1979–1998.² The mortality in young adults (ages 20–24 years), however, increased sharply compared to the mortality in children ages 15–19 years from 0.6/100,000 to 1.4/100,000.² This increase is attributed to problems associated with the transition from pediatric to adult care.

Unlike the USA, the mortality rate in developing countries continues to be high especially in Africa.^{3,4} In the state of Minas Gerais, Brazil the mortality rate among children with SCD born during 2009–2011 was not significantly different from that for children born during 1999–2001 (6.2% vs. 5.8%).⁵ This suggests that newborn screening alone is not enough to reduce the mortality rate of children with SCA. Regular follow-ups, compliance to treatment, family support and the style of life are important factors to maintain the decreases in both morbidity and mortality rates. The purpose of this study was to determine the mortality rates and the cause of death in children, adolescents and adults with SCA followed-up at a single institution in Rio de Janeiro and compare these rates to Brazilian children and adults at large.

Methods

Patients

Data of patients with SCA including patients with sickle/ β^0 -thalassemia (S/ β^0 thal), who were seen and followed at HEMORIO for 15 years from January 1, 1998 through December 31, 2012 were retrospectively collected and analyzed. Children (<18 years old) and adults (\geq 18 years) were included in the study. Age and sex related mortality were also determined and compared to available data of the Brazilian population at large.

The diagnosis of SCA including S/ β^0 thal was confirmed by high performance liquid chromatography (HPLC) as previously described.⁶ The date and cause of deaths were confirmed on patients' charts if death occurred at HEMORIO. Death outside HEMORIO, suspected when patients failed to show up for a follow-up, was confirmed by interviews with the patient's families and from death certificates. Accordingly, the death of 69 patients occurred outside HEMORIO. The family brought the death certificates of 62 patients. Of these 13 deaths were due to accidents unrelated to SCA and in 49 patients, death was due to stroke, acute chest syndrome (ACS) or infection. The cause of death in the remaining seven patients was not clear or could not be confirmed and hence, their cause of death was included in the unknown category.

The study was approved by the Institutional Review Board of HEMORIO.

Statistical analysis

Statistical analysis was performed using the R software⁷ and the Kaplan–Meier estimator to assess the overall survival of patients with SCA. Comparison of survival curves was made using the Tarone–Ware test.⁸ Exclusions from statistical analyses included patients whose deaths were unrelated to SCD and those whose cause of death was unknown. Comparisons of frequencies between nominal variables were made using the chi-squared test without continuity correction or the Fisher exact test. Determination of the average remaining life expectancy was determined by the empirical likelihood (Emplik) ratio test package⁹ of the R software. Statistical significance was based on a *p*-value <0.05.

Results

The total number of patients enrolled in this study was 1676. Age (both at enrollment and at death) and sex are summarized in Table 1. A total of 281 patients died during the study period (Table 1). The mortality rate is summarized in Table 2.

A Kaplan Meier analysis⁸ of the survival probability of all enrolled patients showed that survival of females was significantly higher than males (*p*-value = 0.007). The median age of survival was 53.3 years for males and 56.5 years for females (Figure 1). The median age at death was 28.98 years for men and 34.02 years for women (Table 1).

Table 1 – Characteristics of patients with sickle cell anemia.

	Female	Male	Total
Patients – n (%)	889 (53)	787 (47)	1676 (100)
Deaths – n (%)	139 (49)	142 (51)	281 (100)
Age at enrollment (years) – mean \pm SD	12.52 \pm 11.06	10.99 \pm 10.46	11.80 \pm 10.81
Median (range)	8.89 (0.02–66.99)	7.30 (0.14–62.66)	8.12 (0.02–66.99)
Age at death (years) – mean \pm SD	33.85 \pm 14.80	32.71 \pm 14.25	33.27 \pm 14.51
Median (range)	34.02 (2.74–74.79)	28.98 (0.56–68.03)	31.98 (0.56–74.79)

SD: standard deviation.

Table 2 – Mortality rate of the enrolled patients.

	Patients n	Deaths n	Mortality rate %
Children (<18 years old)	420	44	10.48
Adults (>18 years old)	1256	237	18.87
All patients	1676	281	16.77

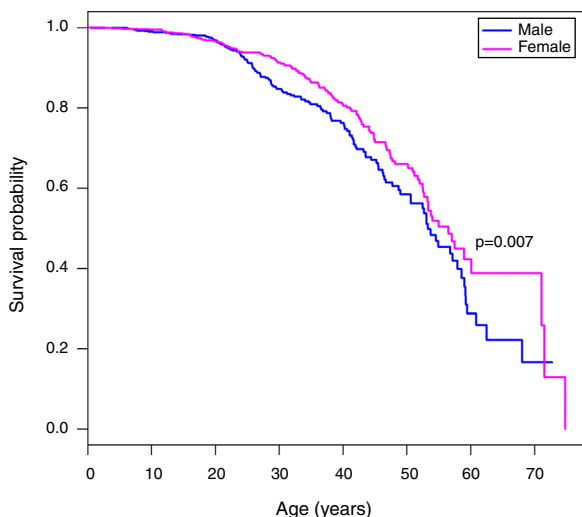


Figure 1 – Kaplan–Meier curves of patients with sickle cell anemia stratified by sex. The median age of survival of males is 53.3 years, while for females the median survival is 56.5 years. The Tarone-Ware test indicates a statistically significant difference between the curves (p -value = 0.007).

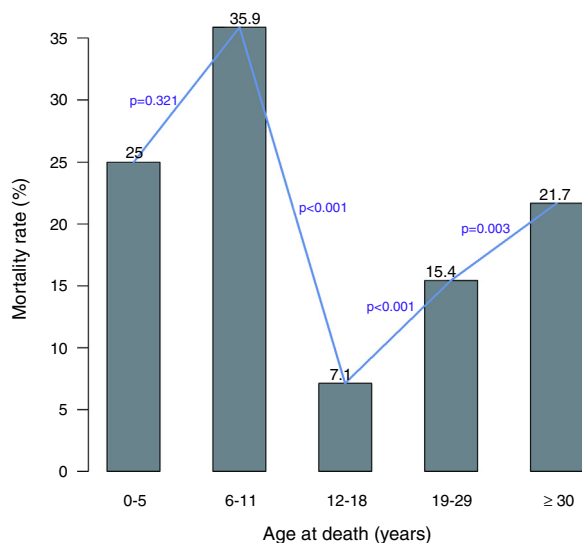


Figure 2 – The mortality rate varies considerably between the age groups. The mortality rate decreases significantly after the age of five years and increases significantly after the age of 18 years.

The mortality among patients varied significantly with age. As shown in Figure 2 the mortality rate in under 5-year-old patients was lower than the mortality rate in the 6–11 age group. However, there was a significant difference in the mortality rates between the 6–11 and 12–18 age groups. After the age of 18 years, there was a significant and sudden increase in the mortality in the 19–29 age group. The mortality continued to increase significantly after the age of 30 years.

The following assumptions were made in order to compare the life expectancy of patients with SCA in Rio de Janeiro with the life expectancy of Brazilians at large. The prevalence of

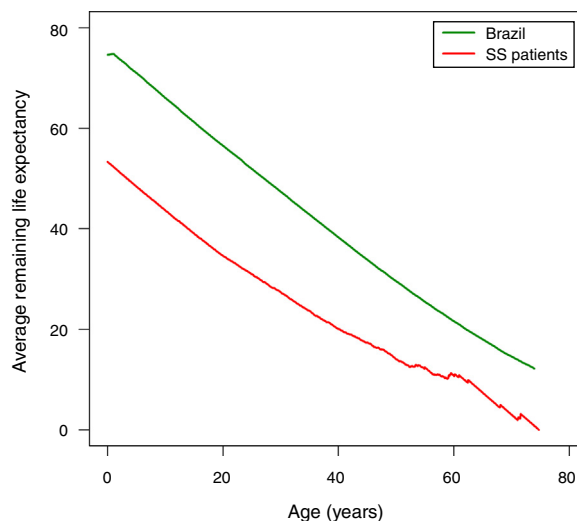


Figure 3 – The average remaining life expectancy for Brazilians and for the patients with sickle cell anemia enrolled in this study.

SCD in the state of Rio de Janeiro is the second highest in Brazil after the state of Minas Gerais. In 2016, about 4000 patients with SCD were treated at HEMORIO representing about 80% of the patients with SCD in the state of Rio de Janeiro. Thus, by extrapolation we assumed that the life expectancy of patients with SCA in Rio de Janeiro may represent the life expectancy of patients with SCA in Brazil. We next compared the life expectancy of patients with SCA described in this study with the life expectancy in Brazilians at large (Figure 3). The latter was based on the 2013 life expectancy report of Brazilians published by the *Instituto Brasileiro de Geografia e Estatística* (IBGE).¹⁰ It is important to note that life tables are used to measure the mortality, survival, and remaining lifetime of a population at various ages as previously described.^{11,12}

Accordingly, the life expectancy of Brazilian males is 71.0 years and for females it is 78.3 years and the estimated life expectancy for both sexes at birth is 74.6 years (Table 3). The remaining life expectancy at birth of the patients with SCA is 53.3 years, that is, 21.3 years less than the life expectancy of Brazilians at large. Between the ages of one and five years, the remaining life expectancy of the patients in this study is about 22 years less than that of Brazilians at large. This gap decreased to about ten years after the age of 60 as shown in Table 3.

The causes of death of the 281 patients who died are listed in Table 4. The most common causes of death were infection, mostly due to sepsis, ACS, overt stroke, sudden death during painful crises and organ damage due to hepatic or renal failure. Thirteen patients died of unrelated causes mostly due to trauma.

Discussion

Although previous studies determined the mortality rate in Brazilian children with SCA, to the best of our knowledge this is the first study that determined the mortality rate in Rio de Janeiro and probably, by extrapolation, in Brazilian adults with

Table 3 – Average remaining life expectancy of sickle cell anemia (SCA) patients compared to that of Brazilians at large.

Age	Life expectancy Brazilians	Life expectancy SCA patients	Age	Life expectancy Brazilians	Life expectancy SCA patients
0	74.6	53.3	38	40.1	21.4
1	74.8	52.3	39	39.2	20.9
2	73.8	51.3	40	38.3	20.0
3	72.9	50.4	41	37.4	19.4
4	71.9	49.4	42	36.5	19.0
5	71.0	48.4	43	35.6	18.5
6	70.0	47.4	44	34.7	17.8
7	69.0	46.5	45	33.9	17.3
8	68.0	45.6	46	33.0	16.7
9	67.0	44.6	47	32.1	16.3
10	66.0	43.7	48	31.3	15.8
11	65.1	42.7	49	30.4	15.1
12	64.1	41.8	50	29.6	14.1
13	63.1	40.9	51	28.8	13.5
14	62.1	39.9	52	27.9	12.9
15	61.2	39.0	53	27.1	12.5
16	60.2	38.1	54	26.3	12.7
17	59.3	37.3	55	25.5	12.4
18	58.3	36.4	56	24.7	11.4
19	57.4	35.5	57	23.9	10.9
20	56.5	34.7	58	23.2	10.8
21	55.6	33.9	59	22.4	10.3
22	54.7	33.2	60	21.6	10.8
23	53.8	32.4	61	20.9	10.8
24	52.8	31.8	62	20.1	9.8
25	51.9	31.0	63	19.4	9.4
26	51.0	30.3	64	18.7	8.4
27	50.1	29.4	65	18.0	7.4
28	49.2	28.7	66	17.3	6.4
29	48.3	28.1	67	16.6	5.4
30	47.4	27.4	68	15.9	4.4
31	46.5	26.7	69	15.2	4.1
32	45.5	25.8	70	14.6	3.1
33	44.6	25.0	71	14.0	2.1
34	43.7	24.3	72	13.4	2.8
35	42.8	23.7	73	12.8	1.8
36	41.9	22.8	74	12.2	0.8
37	41.0	22.1			

Table 4 – Causes of death of the patients with sickle cell anemia.

Cause	n	%
Acute chest syndrome	71	25.27
Cardiac causes	13	4.63
Hemolytic crisis	4	1.42
Hepatic crisis	1	0.36
Infection	82	29.18
Organ damage	30	10.68
Overt stroke	33	11.74
Splenic sequestration	1	0.36
Sudden during painful crisis	26	9.25
Unknown	7	2.49
Unrelated	13	4.63
Total	281	100

SCA. The overall 10.48% mortality rate of all children in this study is higher than that reported in the USA and Europe^{1,2,13} but similar to other reports on Brazilian children.^{5,14} Moreover, the Minas Gerais study reported no significant difference in the mortality rate in 1999–2001 after the initiation of their

newborn screening program in 1998 compared to that in 2009–2011.⁵

On comparing age groups, after the significant decrease in the mortality rate in children aged 6–11 years and in adolescents, there was a sudden and sharp increase in the mortality rate in patients aged 19–29 years similar to that reported in the USA. This surge in mortality after adolescence is not well understood and usually it is assumed to be due to a number of issues related to the psychosocial domain.^{2,15} However, this sudden increase in mortality in patients with SCD aged >18 years of age was not present in a Belgian study that evaluated the survival of patients with SCD recorded in the SCD registry.¹³

As reported in previous studies, the mortality rate in children is similar between males and females.^{1,2,16,17} The divergence in the mortality rate between the sexes in favor of females becomes apparent in adults as shown in this report as well as in previous studies.^{1,18} The reason why adult women with SCA live longer than males is not known. One possibility is that Hb F may be higher in females due to the partial control of Hb F by an X-linked gene located at Xp22.2.^{19–21} Another possibility is the relatively lower

blood viscosity due to lower Hb and hematocrit levels in women.

In order to determine the significance of the mortality rate of patients with SCA, the data of this study were compared with what is known about mortality in children and the life expectancy in the Brazilian population. The last mortality rate for all children <5 years of age in Brazil reported by the United Nations Children's Fund (UNICEF)²² was 1.6% for males and 1.3% for females. These rates are 14.3% for males and 33.3% for females with SCA in the current study. Thus, the mortality rates for under 5-year-old patients with SCA in Rio de Janeiro are about 8.9 times and 25.6 times higher for males and females, respectively. Moreover, the mean difference in average remaining life expectancy for a Brazilian patient with SCA in Rio de Janeiro is 17.6 years (range: 10.0–22.6 years) less than that in the general Brazilian population.^{7,9,10} The average life expectancy for Brazilians at birth is 74.6 years. The mean survival of Brazilian patients with SCA is 53.3 years, 21.3 years less than the general population.

Among the known causes of death due to SCA, infection was the most common cause of death in both children and adults followed by ACS. The most common causes of death in the Brazilian population at large are heart problems, stroke, pneumonia, chronic obstructive pulmonary disease and diabetes mellitus.²³

In conclusion, to the best of our knowledge this is the first Brazilian study that reports the mortality rate among adult patients with SCA in Rio de Janeiro. Unfortunately, this mortality rate is higher than that reported in developed countries in the northern hemisphere. Similarly, the mortality rate among Brazilian children with SCA is higher than that in developed countries. We hope that the mortality rate in Brazilian children and adults will decrease with time as the experience of providers and logistic issues of the newborn screening program improve. Currently, newborn screening seems to be necessary but not sufficient to reduce the mortality rate of children with SCA. In addition, other factors such as regular follow-ups, compliance to treatment, family support and the style of life are important factors to decrease both morbidity and mortality.

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgement

Supported in part by the Office of the Secretary of Health for the State of Rio de Janeiro and the Brazilian Ministry of Health.

REFERENCES

1. Leikin SL, Gallagher D, Kinney TR, Sloane D, Klug P, Rida W. Mortality in children and adolescents with sickle cell disease. Cooperative Study of Sickle Cell Disease. *Pediatrics*. 1989;84(3):500–8.
2. Hamideh D, Alvarez O. Sickle cell disease related mortality in the United States (1999–2009). *Pediatr Blood Cancer*. 2013;60(9):1482–6.
3. McGann PT. Sickle cell anemia: an underappreciated and unaddressed contributor to global childhood mortality. *J Pediatr*. 2014;165(1):18–22.
4. Grosse SD, Odame I, Atrash HK, Amendah DD, Piel FB, Williams TN. Sickle cell disease in Africa: a neglected cause of early childhood mortality. *Am J Prev Med*. 2011;41 6 Suppl. 4:S398–405.
5. Sabarense AP, Lima GO, Silva LM, Viana MB. Survival of children with sickle cell disease in the comprehensive newborn screening programme in Minas Gerais Brazil. *Paediatr Int Child Health*. 2015.
6. Lobo CL, Ballas SK, Domingos AC, Moura PG, do Nascimento EM, Cardoso GP, et al. Newborn screening program for hemoglobinopathies in Rio de Janeiro, Brazil. *Pediatr Blood Cancer*. 2014;61(1):34–9.
7. R Core Team. R: A language and environment for statistical computing. Vienna, Austria: R Foundation for Statistical Computing; 2015.
8. Tarone R, Ware J. On distribution-free tests for equality of survival distributions. *Biometrika*. 1977;64:1561–660.
9. Zhou M, Jeong JH. Empirical likelihood ratio test for median and mean residual lifetime. *Stat Med*. 2011;30(2):152–9.
10. IBGE. Tábuas Completas de Mortalidade por Sexo e Idade Brasil 2012: Breve análise da mortalidade no período 2011–2012. Rio de Janeiro: IBGE; 2013.
11. Barclay GW. The study of mortality. Techniques of population analysis. New York: John Wiley and Sons; 1958. p. 123–34.
12. Raymondo JC. Survival rates: census and life table methods. Population estimation and projection. New York: Quorum Books; 1992. p. 43–60.
13. Le PQ, Gulbis B, Dedeken L, Dupont S, Vanderfaeillie A, Heijmans C, et al. Survival among children and adults with sickle cell disease in Belgium: benefit from hydroxyurea treatment. *Pediatr Blood Cancer*. 2015;62(11):1956–61.
14. Sabarense AP, Lima GO, Silva LM, Viana MB. Characterization of mortality in children with sickle cell disease diagnosed through the Newborn Screening Program. *J Pediatr (Rio J)*. 2015;91(3):242–7.
15. Quinn CT, Rogers ZR, McCavit TL, Buchanan GR. Improved survival of children and adolescents with sickle cell disease. *Blood*. 2010;115(17):3447–52.
16. Quinn CT, Rogers ZR, Buchanan GR. Survival of children with sickle cell disease. *Blood*. 2004;103(11):4023–7.
17. Rogers DW, Clarke JM, Cupidore L, Ramlal AM, Sparke BR, Serjeant GR. Early deaths in Jamaican children with sickle cell disease. *Br Med J*. 1978;1(6126):1515–6.
18. Wierenga KJ, Hambleton IR, Lewis NA. Survival estimates for patients with homozygous sickle-cell disease in Jamaica: a clinic-based population study. *Lancet*. 2001;357(9257):680–3.
19. Dover GJ, Smith KD, Chang YC, Purvis S, Mays A, Meyers DA, et al. Fetal hemoglobin levels in sickle cell disease and normal individuals are partially controlled by an X-linked gene located at Xp22.2. *Blood*. 1992;80(3):816–24.
20. Chang YC, Smith KD, Moore RD, Serjeant GR, Dover GJ. An analysis of fetal hemoglobin variation in sickle cell disease: the relative contributions of the X-linked factor, beta-globin haplotypes, alpha-globin gene number, gender, and age. *Blood*. 1995;85(4):1111–7.
21. Chang YP, Maier-Redelsperger M, Smith KD, Contu L, Ducrocq R, de Montalembert M, et al. The relative importance of the X-linked FCP locus and beta-globin haplotypes in determining haemoglobin F levels: a study of SS patients homozygous for beta S haplotypes. *Br J Haematol*. 1997;96(4):806–14.

22. You D, Bastian P, Wu J, Wardlaw T. Levels and trends in child mortality. Report 2013. Fund UNCs ed; 2013.
23. GBD 2013 Mortality and Causes of Death Collaborators. Global, regional, national age-sex specific all-cause, cause-specific mortality for 240 causes of death, 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet*. 2015;385(9963):117–71.