



**FACULTY OF HEALTH SCIENCES
DEPARTMENT OF COMMUNITY & PUBLIC HEALTH**

FINAL YEAR PROJECT REPORT

**PREVALENCE OF AND FACTORS ASSOCIATED WITH COMBINED USE OF
HERBAL AND CONVENTIONAL MEDICINE BY CARETAKERS OF CHILDREN
WITH SICKLE CELL DISEASE IN JINJA REGIONAL REFERRAL HOSPITAL: A
CROSS-SECTIONAL STUDY**

BY:

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**A RESEARCH REPORT SUBMITTED TO THE FACULTY OF HEALTH SCIENCES
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ABSTRACT

Background: Concurrent use of Herbal (HM) and Conventional Medicine (CM) poses a huge risk of drug-herbal interactions that can result in therapeutic failure or toxicity. However, little is known about the prevalence of combined use of HM and CM and the associated factors by caretakers of children with sickle cell disease (SCD). This study aimed to determine the prevalence of and factors associated with the combined use of HM and CM by caretakers of children with SCD attending the Sickle Cell Clinic (SCC) at Jinja Regional Referral Hospital (JRRH), East-Central Uganda.

Methods: A mixed-method, explanatory design was used. Data were collected between January and March 2022. Quantitatively, a sample size of 394 caretakers of children with SCD aged 1-18 years was targeted. Structured questionnaires were used to collect data on socio-demographic characteristics, caretakers' take on SCD treatment, perceptions of and intentions of the caretakers to use HM, CM, or both, and community and health-related factors. The primary outcome was the combined use of HM and CM. Quantitative data analysis was performed using Stata version 15. A logistic regression model was fitted to determine the factors associated with the use of combined therapy and all factors with a p-value less than 0.05 were considered significantly associated with the primary outcome. After completion of the structured interviews, qualitative data were collected from 26 purposively selected caretakers who reported use of both HM and CM and had participated in the quantitative interviews. Four focus group discussions (FGDs) were conducted to collect data on caretakers' take on treatment outcomes; cultural, family, friends or relatives' influence, and caretakers' considerations when choosing both remedies. Qualitative data were analyzed following a thematic framework approach, using NVIVO software.

Results: A total of 372 (94.4% of the sample) caretakers were interviewed; 341 (91.7%) of whom were females. Respondents' average age was 34.3 (SD: ± 9.8) years. More than half (55.1%, n=205) of the caretakers were aged 18-34 years while 40.9% (n=152) had primary education. Fifty-eight percent (n=217) of the caretakers reported use of only CM, 37.1%, (n=138) used combined therapy while 4.6% (n=17) reported use of HM only. In the multivariate analysis, caretakers aged 60+ years (adjusted odds ratio [AOR] = 11.8; 95% Confidence Interval [95%CI]: 1.2, 115.2), those with lower secondary education (AOR=6.2; 95% CI: 1.5, 26.0), those who believed that HM is beneficial (AOR=3.3; 95% CI: 1.5, 7.6) and those who believed that the use of both CM and HM is safe (AOR=7.7; 95% CI: 3.5, 17.0) were significantly more likely to use combined therapy than their counterparts. Caretakers who intended to use only CM (AOR=0.1; 95% CI: 0.1, 0.3) were significantly less likely to use combined HM and CM. From the qualitative findings, lack of trust in the use of either CM alone or HM alone and recommendations from relatives and friends were the main factors that influenced the use of combined therapy.

Conclusion: Slightly more than half of the caretakers used CM only while more than a third used both. Caretakers' age and level of education, compliance with recommendations from significant others, and lack of trust in either therapy alone were significantly associated with the use of combined therapy. We recommend that further research be done targeting the understanding of the use of combined medicines.

DECLARATION

I, Apolot Consiliate, certify that this study is my original work and it has never been published or submitted to any university or institution of higher learning for any academic award.

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APPROVAL

This dissertation has been with our approval as the academic supervisor(s)

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DEDICATION

This research is dedicated to Jehovah Jireh, the Alpha, and Omega, for without Him, I would not have started this journey.

To you, I give all the Glory and the Honor.

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ACRONYMS AND ABBREVIATIONS

ADR: Adverse Drug Reaction

CAM: Complementary and Alternative Medicines

CM: Conventional Medicine

HM: Herbal Medicine

HU: Hydroxyurea

MOH: Ministry of Health

PHC: Primary Health Care

SCD: Sickle Cell Disease

SSA: Sub Saharan Africa

TPB: Theory of Planned Behaviour

WHO: World Health Organization

OPERATIONAL DEFINITIONS

Key Concepts	Definitions
Conventional Medicine	These are medicines formulated using known formulas and quantifications, produced under approval by Food and Drug Administration (FDA). In this study, I referred to conventional medicines as approved medicines that caretakers received from the sickle cell clinic of JRRH and administered to their children with sickle cell disease following the physician’s prescription such as folic acid, and penicillins, and those provided during hospital admissions like blood transfusions, and other intravenous treatments. In addition, any other approved medicines that a caretaker bought from over – the counter in pharmacies or drug stores for disease management were also considered conventional.
Herbal Medicine	Herbal medicines in this study were referred to as medications prepared from one or more herbs or plant parts (roots, stem bark, seeds, and/or fruits and those medicines bought from the herbalists that caretakers of children with SCD may use to treat a child with sickle cell disease. Unlike prescription or over-the-counter drugs that may contain purified ingredients from plants, herbal medicines contain entire plants or plant parts, and the

	<p>medication is usually not measured in any known units and the amount administered at any one given time largely depends on the caregiver's perception of the severity of disease or herbal drug efficacy.</p>
Combined use/ Therapy	<p>In this study, combined therapy/use referred to the use of both HM and CM concurrently by the caretaker to treat a child with sickle cell disease.</p>
Child	<p>In this study, a child was referred to as one who is between 1-10 years, diagnosed with Sickle Cell Disease (SCD) attending the sickle cell clinic of JRRH and under the care of an adult (caretaker).</p>
Caretaker	<p>A caretaker was referred to as someone who offered emotional, psychological, and physical support to the child with SCD, attending the Sickle Cell Clinic at JRRH.</p>
Significant other	<p>These referred to the people that caretakers believed are important to them, hence their say in the treatment choices also regarded important. This include friends, family, health professionals, neighbors.</p>

CHAPTER ONE: INTRODUCTION

1.1 Background

Globally, it is estimated that about 300,000 to 400,000 babies are born annually with Sickle Cell Disease (SCD), with 90% of the cases occurring in sub-Saharan Africa (McGann, 2016). The prevalence of SCD ranges between 1% and 2% in North Africa to less than 1% in Southern Africa and in countries such as Cameroon, the Republic of Congo, Gabon, Ghana, Nigeria, and Uganda the prevalence of the SCD trait is between 20% and 30%. A survey done on SCD in Uganda revealed that the prevalence of SCD trait in Uganda is estimated to be 13.3%, and the disease at 0.7% (Ndeezi et al, 2016), across various regions. Another study further revealed that of the 900,000 thousand children born annually in Uganda (UDHS, 2006), approximately 2.8% have sickle cell disease (Serjeant & Ndugwa, 2003).

SCD disease management involves palliative care, comprising preventive, symptomatic, and supportive approaches (Sinha et al., 2019). Preventive therapy most commonly consists of hydroxyurea, routine vaccinations, blood transfusions, prophylactic antibiotics, and antimalarials coupled with iron supplements and pain management which are the mainstay treatments as the disease persists (Johnso et al., 2015) throughout life.

In recent decades, however, there has been a global surge in popularity of the use of Complementary and Alternative Modalities/Medicines (CAM) for disease management in both developing and developed countries with over 80% accounting for those in Sub-Saharan Africa (Sinha et al., 2019). Herbal medicines (HM) account for more than half of the various alternative medicines used in Africa (Mahomoodally, 2013; Ndeezi et al., 2016; Oshikoya et al., 2013; WHO, 2013). HM may be used alone as a substitute or concurrently with Conventional Medicine (CM).

The use of HM has been widely reported in the Caribbean, Trinidad, Uganda, South Africa, and Nigeria with a prevalence of 20-80% in both young and old patients with chronic health conditions such as SCD, diabetes, asthma, epilepsy, hypertension, HIV infection and cancer (Kamatenesi-Mugisha & Oryem-Origa, 2005; Langlois-Klassen et al., 2007; Nsibirwa et al.,

References

- Adewoyin, A. S., Oghuvwu, O. S., & Awodu, O. A. (2017). Hydroxyurea therapy in adult Nigerian sickle cell disease: A monocentric survey on the pattern of use, clinical effects and patient's compliance. *African Health Sciences*, *17*(1), 255-261. <https://doi.org/10.4314/ahs.v17i1.31>
- Ahmadi, M., Ilkhani, M., Beiranvand, S., Poormansouri, S., & Sedighie, L. (2017). Massage for Pain Management in Patients with Sickle Cell Disease: A Review Study. *Jundishapur Journal of Chronic Disease Care*, *7*(1), 1–8. <https://doi.org/10.5812/jjcdc.62315>
- Ameade, Evans P K, Amalba, A., Helegbe, G. K., & Mohammed, B. S. (2015). Herbal medicine: a survey on the knowledge and attitude of medical students in Tamale, Ghana. *Peak Journal of Medicinal Plant Research*, *3*(1), 1–8. www.peakjournals.org/sub-journals-PJMPPR.html
- Ameade, Evans Paul Kwame, Ibrahim, M., Ibrahim, H.-S., Habib, R. H., & Gbedema, S. Y. (2018). Concurrent Use of Herbal and Orthodox Medicines by Residents of Tamale, Northern Ghana, Who Patronize Hospitals and Herbal Clinics. *Evidence-Based Complementary and Alternative Medicine*, *2018*(2002), 1–8. <https://doi.org/10.1155/2018/1289125>
- Ameh, S. J., Tarfa, F. D., & Ebeshi, B. U. (2012). Traditional herbal management of sickle cell anemia: Lessons from Nigeria. *Anemia*, *2012*. <https://doi.org/10.1155/2012/607436>
- Aziato, L., & Antwi, H. O. (2016). Facilitators and barriers to herbal medicine use in Accra, Ghana: An inductive exploratory study. *BMC Complementary and Alternative Medicine*, *16*(1), 1–9. <https://doi.org/10.1186/s12906-016-1124-y>

- Ballas, S. K. (2017). The Use of Cannabis by Patients with Sickle Cell Disease Increased the Frequency of Hospitalization due to Vaso-Occlusive Crises. *Cannabis and Cannabinoid Research*, 2(1), 197–201. <https://doi.org/10.1089/can.2017.0011>
- Bodeker, G., & Kronenberg, F. (2002). A public health agenda for traditional, complementary, and alternative medicine. *American Journal of Public Health*, 92(10), 1582–1591.
<https://doi.org/10.2105/AJPH.92.10.1582>
- Brumatti, L. V., Marcuzzi, A., Tricarico, P. M., Zanin, V., Girardelli, M., & Bianco, A. M. (2014). Curcumin and inflammatory bowel disease: Potential and limits of innovative treatments. *Molecules*, 19(12), 21127–21153.
<https://doi.org/10.3390/molecules191221127>
- Busari, A. A., & Mufutau, M. A. (2017). High prevalence of complementary and alternative medicine use by patients with sickle cell disease in a tertiary hospital in Lagos, South West, Nigeria. *BMC Complementary and Alternative Medicine*, 17(1), 1–8.
<https://doi.org/10.1186/s12906-017-1812-2>
- Clement, Y. N., Morton-Gittens, J., Basdeo, L., Blades, A., Francis, M. J., Gomes, N., Janjua, M., & Singh, A. (2007). Perceived efficacy of herbal remedies by users accessing primary healthcare in Trinidad. *BMC Complementary and Alternative Medicine*, 7 (May 2014). <https://doi.org/10.1186/1472-6882-7-4>
- Floden, A, Combs, C. (2012). NIH Public Access. *Bone*, 23(1), 1–7.
<https://doi.org/10.1097/01.qai.0000437619.23031.83.Reduced>
- Gardner, R. V. (2018). Sickle cell disease: Advances in treatment. *Ochsner Journal*, 18(4), 377– 389. <https://doi.org/10.31486/toj.18.0076>
- Gasmi, A., Chirumbolo, S., Peana, M., Noor, S., Menzel, A., Dadar, M., & Bjørklund, G. (2021).

The Role of Diet and Supplementation of Natural Products in COVID-19 Prevention. *Biological Trace Element Research*, 10–13. <https://doi.org/10.1007/s12011-021-02623-3>

Guest, G., Bunce, A., & Johnson, L. (2006). How Many Interviews Are Enough?: An Experiment with Data Saturation and Variability. *Field Methods*, 18(1), 59–82. <https://doi.org/10.1177/1525822X05279903>

Harvard, T. (2020). *Barriers to Adherence to Hydroxyurea in Rural Global Low Resource Settings : Lessons From Rural Central India and Beyond*.

James, P. B., Kaikai, A. I., Bah, A. J., Steel, A., & Wardle, J. (2019). Herbal medicine use during breastfeeding: A cross-sectional study by mothers visiting public health facilities in the Western area of Sierra Leone. *BMC Complementary and Alternative Medicine*, 19(1), 1–11. <https://doi.org/10.1186/s12906-019-2479-7>

James, P. B., Wardle, J., Steel, A., & Adams, J. (2018). Traditional, complementary and alternative medicine use in Sub-Saharan Africa: A systematic review. *BMJ Global Health*, 3(5). <https://doi.org/10.1136/bmjgh-2018-000895>

Johnso, P. L., Lombardo, F. A., & Moore, C. D. (2015). Options in the management of sickle cell disease. *U.S. Pharmacist*, 40(7), 4–7.

Jose, J., Elsadek, R. A., Jimmy, B., & George, P. (2019). Hydroxyurea: Pattern of use, patient adherence, and safety profile in patients with sickle cell disease in Oman. *Oman Medical Journal*, 34(4), 327–335. <https://doi.org/10.5001/omj.2019.64>

Kamatenesi-Mugisha, M., & Oryem-Origa, H. (2005). Traditional herbal remedies used in the management of sexual impotence and erectile dysfunction in western Uganda. *African Health Sciences*, 5(1), 40–49. <https://doi.org/10.4314/ahs.v5i1.6896>

- Langlois-Klassen, D., Kipp, W., Jhangri, G. S., & Rubaale, T. (2007). Use of traditional herbal medicine by AIDS patients in Kabarole District, western Uganda. *The American Journal of Tropical Medicine and Hygiene*, 77(4), 757–763.
- Mahomoodally, M. F. (2013). Traditional medicines in Africa: An appraisal of ten potent African medicinal plants. *Evidence-Based Complementary and Alternative Medicine*, 2013. <https://doi.org/10.1155/2013/617459>
- McGann, P. T. (2016). Time to invest in sickle cell anemia as a global health priority. *Pediatrics*, 137(6). <https://doi.org/10.1542/peds.2016-0348>
- Medicine, T. T., & Products, N. (2016). *The Traditional Medicine and Modern Medicine from Natural Products*. <https://doi.org/10.3390/molecules21050559>
- Mpiana, P. T. (2014). *The Potential Effectiveness of Medicinal Plants used for the Treatment of Sickle Cell Disease in the Democratic Republic of Congo Folk Medicine : A Review. January 2012.*
- Mpiana1, P. T., Ngbolua, K. N., Mudogo, V., Tshibangu, D. S. T., Atibu, E. K., Mbala, B. M., Kahumba, B., Bokota, M. T., & Makelele, L. T. (2012). The Potential effectiveness of medicinal plants used for the treatment of Sickle Cell Disease in the Democratic Republic of Congo Folk Medicine : A review. *Progress in Traditional and Folk Herbal Medicine*, 1(January 2012), 1–11.
- Mulumba, L. L., & Wilson, L. (2020). *International Journal of Africa Nursing Sciences Sickle cell disease by children in Africa : An integrative literature review and global recommendations*. 3(2015), 56–64. <https://doi.org/10.1016/j.ijans.2015.08.002>
- Ndao-Brumblay, S. K., & Green, C. R. (2010). Predictors of complementary and alternative medicine use in chronic pain patients. *Pain Medicine*, 11(1), 16–24.

<https://doi.org/10.1111/j.1526-4637.2009.00767.x>

Ndeezi, G., Kiyaga, C., Hernandez, A. G., Munube, D., Howard, T. A., Ssewanyana, I., Nsungwa, J., Kiguli, S., Ndugwa, C. M., Ware, R. E., & Aceng, J. R. (2016). The burden of sickle cell trait and disease in the Uganda Sickle Surveillance Study (US3): A cross-sectional study. *The Lancet Global Health*, 4(3), e195–e200. [https://doi.org/10.1016/S2214109X\(15\)00288-0](https://doi.org/10.1016/S2214109X(15)00288-0)

Nsibirwa, S., Anguzu, G., Kamukama, S., Ocamo, P., & Nankya-Mutyoba, J. (2020). Herbal medicine use by patients with viral and non-viral hepatitis in Uganda: Prevalence, patterns and related factors. *BMC Complementary Medicine and Therapies*, 20(1), 1–11. <https://doi.org/10.1186/s12906-020-02959-8>

Okoh, M. P., Alli, L. A., Tolvanen, M. E. E., & Nwegbu, M. M. (2018). Herbal Drug use in Sickle Cell Disease Management; Trends and Perspectives in Sub-Saharan Africa - A Systematic Review. *Current Drug Discovery Technologies*, 16(4), 372–385. <https://doi.org/10.2174/1570163815666181002101611>

Okoronkwo, I., Onyia-Pat, J. L., Okpala, P., Agbo, M. A., & Ndu, A. (2014). Patterns of complementary and alternative medicine use, perceived benefits, and adverse effects by adult users in Enugu Urban, Southeast Nigeria. *Evidence-Based Complementary and Alternative Medicine*, 2014(October). <https://doi.org/10.1155/2014/239372>

Opoka, R. O., Ndugwa, C. M., Latham, T. S., Lane, A., Hume, H. A., Kasirye, P., Hodges, J. S., Ware, R. E., & John, C. C. (2017). Novel use Of Hydroxyurea in an African Region with Malaria (NOHARM): a trial for children with sickle cell anemia. *Blood*, 130(24), 2585–2593. <https://doi.org/10.1182/blood-2017-06-788935>

Oreagba, I. A., Oshikoya, K. A., & Amachree, M. (2011). Herbal medicine use by urban residents in Lagos, Nigeria. *BMC Complementary and Alternative Medicine*, 11(1), 117.

<https://doi.org/10.1186/1472-6882-11-117>

Oshikoya, K. A., Oreagba, I. A., Ogunleye, O. O., Oluwa, R., Senbanjo, I. O., & Olayemi, S. O.

(2013). Herbal medicines supplied by community pharmacies in Lagos, Nigeria: pharmacists' knowledge. *Pharmacy Practice (Internet)*, *11*(4), 219–227.

<https://doi.org/10.4321/s1886-36552013000400007>

Oshikoya, K. A., Senbanjo, I. O., Njokanma, O. F., & Soipe, A. (2008). Use of complementary and alternative medicines for children with chronic health conditions in Lagos, Nigeria. *BMC Complementary and Alternative Medicine*, *8*, 1–8.

<https://doi.org/10.1186/1472-6882-8-66>

Rashrash, M., Schommer, J. C., & Brown, L. M. (2017). Prevalence and Predictors of Herbal Medicine Use By Adults in the United States. *Journal of Patient Experience*, *4*(3),

108–113. <https://doi.org/10.1177/2374373517706612>

Rossignol, M., Bégaud, B., Avouac, B., Lert, F., Rouillon, F., Bénichou, J., Massol, J., Duru, G., Magnier, A. M., Guillemot, D., Grimaldi-Bensouda, L., & Abenhaim, L. (2011). Who seeks primary care for musculoskeletal disorders (MSDs) with physicians prescribing homeopathic and other complementary medicine? Results from the EPI3-LASER survey in France. *BMC Musculoskeletal Disorders*, *12*, 1–6.

<https://doi.org/10.1186/1471-2474-12-21>

Ryan, N., Dike, L., Ojo, T., Vieira, D., Nnodu, O., Gyamfi, J., & Peprah, E. (2020).

Implementation of the therapeutic use of hydroxyurea for sickle cell disease management in resource-constrained settings: A systematic review of adoption, cost, and acceptability. *BMJ Open*, *10*(11), 1–9. <https://doi.org/10.1136/bmjopen-2020-038685>

- Sanchez, H. C., Karlson, C. W., Hsu, J. H., Ostrenga, A., & Gordon, C. (2015). Complementary and Alternative Medicine Use in Pediatric Hematology/Oncology Patients at the University of Mississippi Medical Center. *Journal of Alternative and Complementary Medicine*, 21(11), 660–666. <https://doi.org/10.1089/acm.2014.0371>
- Saraf, S. L., & Rondelli, D. (2019). Allogeneic Hematopoietic Stem Cell Transplantation for Adults with Sickle Cell Disease. *Journal of Clinical Medicine*, 8(10), 1565. <https://doi.org/10.3390/jcm8101565>
- Segal, J. B., Strouse, J. J., Beach, M. C., Haywood, C., Witkop, C., Park, H., Wilson, R. F., Bass, E. B., & Lanzkron, S. (2008). Hydroxyurea for the treatment of sickle cell disease. In *Evidence report/technology assessment*. <https://doi.org/10.1056/NEJMct0708272>
- Sinha, C. B., Bakshi, N., Ross, D., & Krishnamurti, L. (2019). Management of chronic pain in adults living with sickle cell disease in the era of the opioid epidemic a qualitative study. *JAMA Network Open*, 2(5), 1–9. <https://doi.org/10.1001/jamanetworkopen.2019.4410>
- Skovgaard, L., Nicolajsen, P. H., Pedersen, E., Kant, M., Fredrikson, S., Verhoef, M., & Meyrowitsch, D. W. (2012). Use of complementary and alternative medicine by people with multiple sclerosis in the nordic countries. *Autoimmune Diseases*, 1(1). <https://doi.org/10.1155/2012/841085>
- Stratton, T. D., & McGivern-Snofsky, J. L. (2008). Toward a sociological understanding of complementary and alternative medicine use. *Journal of Alternative and Complementary Medicine*, 14(6), 777–783. <https://doi.org/10.1089/acm.2007.7006>
- Sun, C., Desai, G. J., Pucci, D. S., & Jew, S. (2004). Musculoskeletal Disorders: Does the Osteopathic Medical Profession Demonstrate Its Unique and Distinctive Characteristics?

Journal of the American Osteopathic Association, 104(4), 149–155.
<https://doi.org/10.7556/jaoa.2004.104.4.149>

Tabuti, J. R. S., Lye, K. A., & Dhillion, S. S. (2003). Traditional herbal drugs of Bulamogi, Uganda: Plants, use and administration. *Journal of Ethnopharmacology*.
[https://doi.org/10.1016/S0378-8741\(03\)00161-2](https://doi.org/10.1016/S0378-8741(03)00161-2)

Tesfamariam, S., Tesfai, F., Hussien, L., Ateshim, Y., Yemane, D., Russom, M., Ahmed, H., Bahta, I., Kidane, S. N., Namboze, J., & Kasilo, O. M. J. (2021). Traditional medicine by the community of Gash-Barka region, Eritrea: attitude, societal dependence, and pattern of use. *BMC Complementary Medicine and Therapies*, 21(1), 1–9.
<https://doi.org/10.1186/s12906-021-03247-9>

Thompson, W. E., & Eriator, I. (2014). Pain control in sickle cell disease patients: Use of complementary and alternative medicine. *Pain Medicine (United States)*, 15(2), 241–246.
<https://doi.org/10.1111/pme.12292>

Toklu, H. Z. (2013). Herbal medicine use by diabetes mellitus patients in Northern Cyprus. *Journal of Medicinal Plants Research*, 7(22), 1652–1664.
<https://doi.org/10.5897/JMPR12.1207>

Tugume, P., & Nyakoojo, C. (2019). Ethno-pharmacological survey of herbal remedies used in the treatment of pediatric diseases in Buhunga parish, Rukungiri District, Uganda. *BMC Complementary and Alternative Medicine*, 19(1), 1–10.
<https://doi.org/10.1186/s12906-0192763-6>

Wambebe, C., Khamofu, H., Momoh, J. A., Ekpeyong, M., Audu, B. S., Njoku, O. S., Bamgboye, E. A., Nasipuri, R. N., Kunle, O. O., Okogun, J. I., Enwerem, M. N., Audam, J. G., Gamaniel, K. S., Obodozie, O. O., Samuel, B., Fojule, G., & Ogunyale, O. (2001). Double-blind, placebo-controlled, randomized cross-over clinical trial of NIPRISAN®

in patients with Sickle Cell Disorder. *Phytomedicine*. <https://doi.org/10.1078/0944-711300040>

WHO_EDM_TRM_2001.2_rus.pdf. (n.d.).

World Health Organization (WHO). (2013). WHO Traditional Medicine Strategy 2014-2023.

World Health Organization (WHO), 1–76. <https://doi.org/2013>

Yoon SL, Black S. Comprehensive, integrative management of pain for patients with sickle-cell disease. *J Altern Complement Med*. 2006 Dec;12(10):995-1001. DOI:10.1089/acm.2006.12.995. PMID: 17212571.