

Mechanisms and Therapeutic Potential of *Gongronema latifolium* in Sickle Cell Disease

Nnodim Johnkennedy *, Uzodima Udoha Augustine, Nwanguma Eberechi Doris, Nwaneri Faith Ekpere Kelechi

Department of Medical Laboratory Science, Imo State University, Owerri.

*Corresponding Author: Nnodim Johnkennedy, Department of Medical Laboratory Science, Imo State University, Owerri.

Received date: 24 December 2025 | Accepted: 07 January 2026 | Published: 14 January 2026

Citation: Nnodim Johnkennedy, Uzodima U. Augustine, Nwanguma E. Doris, Ekpere Kelechi NF, (2026), Mechanisms and Therapeutic Potential of *Gongronema latifolium* in Sickle Cell Disease, *Clinical Endocrinology and Metabolism*, 5(1); Doi:10.31579/2834-8761/102

Copyright: © 2026, Nnodim Johnkennedy. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Abnormal haemoglobin polymerisation, red blood cell deformation, chronic haemolysis, vaso-occlusion, and systemic inflammation are the hallmarks of sickle cell disease (SCD), a genetic hemoglobinopathy. Many patients in developing nations rely on medicinal plants as supplemental therapies despite advancements in management. The tropical medicinal plant *Gongronema latifolium* Benth., which is commonly used in West Africa, has drawn notice for its haematopoietic, antioxidant, anti-inflammatory, and antisickling qualities. This article examines *G. latifolium*'s pharmacodynamics in relation to sickle cell disease, emphasising its bioactive phytochemicals and modes of action. The plant contains flavonoids, alkaloids, saponins, tannins, and phenolic compounds that can reduce inflammation, improve erythrocyte membrane stability, modulate oxidative stress, and inhibit haemoglobin S polymerisation, according to experimental research. These pharmacodynamic effects imply that *G. latifolium* might be helpful in sickle cell disease treatment. To confirm its effectiveness, find the best dosage, and create safety profiles for long-term therapeutic use, more clinical trials are needed.

Keywords: oxidative stress; *gongronema latifolium*; sickle cell disease; pharmacodynamics; antisickling agents; phytochemicals

Introduction

One of the most common genetic diseases in the world, sickle cell disease (SCD) is especially common in sub-Saharan Africa. Haemoglobin S (HbS) is produced when a point mutation in the β -globin gene causes valine to replace glutamic acid at position six of the haemoglobin molecule. Low oxygen tension causes HbS to polymerise, giving erythrocytes a sickled shape and stiffness. These aberrant cells cause organ damage, haemolytic anaemia, vaso-occlusion, and frequent, excruciating crises [1]. Hydroxyurea, blood transfusions, and bone marrow transplants are examples of pharmacological treatments that have improved patient outcomes, but they are costly and difficult to obtain in many low-income nations. As a result, medicinal plants continue to be a significant complementary or alternative treatment for sickle cell disease [2]. *Gongronema latifolium*, a climbing shrub in the Asclepiadaceae (Apocynaceae) family, is one plant that has garnered growing scientific attention. The plant is widely found throughout tropical Africa, and in Nigeria, it is frequently used as a vegetable and medicinal herb. It has historically been used to treat sickle cell disease, diabetes, hypertension, malaria, and gastrointestinal issues [3]. *G. latifolium* has been shown to have antioxidant, anti-inflammatory, hepatoprotective, antimicrobial, and antisickling properties in a number of pharmacological studies. Its rich phytochemical composition and capacity to alter several biochemical pathways linked to disease processes are primarily responsible for these therapeutic qualities. This review discusses the pharmacodynamic

mechanisms through which *G. latifolium* may exert beneficial effects in patients with sickle cell disease [4].

Gongronema latifolium's Phytochemical Composition

The bioactive phytochemical components of medicinal plants play a major role in their pharmacological actions. Several classes of secondary metabolites have been found in *G. latifolium*, according to phytochemical analyses, including: Flavonoids, Alkaloids, Saponins, Tannins, Terpenoids, Phenolic substances and Glycosides [5]. The leaves of *G. latifolium* are especially rich in phenolic compounds and flavonoids. These substances have potent antioxidant qualities and are essential for preventing oxidative damage to erythrocytes. Additionally, molecular docking studies have predicted that some of the plant's phytochemicals will interact with sickle haemoglobin, indicating potential therapeutic activity against SCD [6].

Sickle Cell Disease Pharmacodynamic Mechanisms

The antisickling activity of *G. latifolium* is one of its most significant pharmacodynamic characteristics. By preventing haemoglobin polymerisation, antisickling agents stop or reverse red blood cell deformation.

The plant's phytochemicals may interact with haemoglobin molecules to stabilise the oxygenated form of haemoglobin and stop HbS from polymerising in hypoxic environments. This effect enhances blood flow in microcirculation and lessens erythrocyte sickling [7]. *G. latifolium* is frequently used in polyherbal antisickling formulations used in traditional medicine to treat sickle cell anaemia. Such plant-based formulations have been shown in experiments to affect haematological parameters like haemoglobin levels and red blood cell counts [8].

A key factor in the pathophysiology of sickle cell disease is oxidative stress. Excessive reactive oxygen species (ROS) produced by sickled erythrocytes harm cell membranes and encourage haemolysis. Antioxidant enzyme activities have been demonstrated to be increased by *G. latifolium* extracts, including: SOD, or superoxide dismutase GPx, or glutathione peroxidase and Reductase of glutathione (GR). In addition, the plant protects against oxidative damage by lowering lipid peroxidation markers like malondialdehyde (MDA). *G. latifolium* may improve erythrocyte survival and lessen haemolysis in sickle cell patients through these antioxidant mechanisms [9]. Another important component of sickle cell disease is chronic inflammation. Increased expression of adhesion molecules and inflammatory cytokines is linked to vaso-occlusive crises. *G. latifolium* contains bioactive substances with strong anti-inflammatory properties, especially flavonoids and saponins. These substances block inflammatory mediators like: TNF- α , or tumour necrosis factor-alpha, IL-6, or interleukin-6, NF- κ B, or nuclear factor kappa B. *G. latifolium* may lessen vascular inflammation and the frequency of vaso-occlusive episodes by inhibiting inflammatory pathways [10]. The development of sickle cell disease is significantly influenced by the immune system. Research has shown that *G. latifolium* extracts improve humoral and cellular immune responses. According to experimental data, the plant extract boosts phagocytic activity, increases neutrophil counts, and induces the production of immunoglobulins in animal models. People with sickle cell disease, who are frequently prone to infections, may benefit from this immunomodulatory action by having better host defence mechanisms [11]. The delicate membranes of sickle erythrocytes are easily ruined during circulation. It is thought that some of the phytochemicals in *G. latifolium* stabilise erythrocyte membranes, which lowers haemolysis. Membrane stabilisation happens via a number of processes, such as: Lipid peroxidation inhibition, improvement of the structural integrity of the membrane defence of membrane proteins [12]. Patients with sickle cell disease may benefit from these effects in terms of better haematological outcomes and red blood cell survival.

Bioavailability and Pharmacokinetics

The information that is currently available indicates that *G. latifolium*'s active compounds are absorbed through the gastrointestinal tract after oral administration, despite the fact that research on its pharmacokinetics is still scarce. Following absorption, these substances have systemic effects via metabolic, anti-inflammatory, and antioxidant pathways [13]. The absorption, distribution, metabolism, and excretion (ADME) profiles of the plant's main bioactive components require additional research.

Implications for Clinical Practice

G. latifolium's pharmacodynamic characteristics imply that the plant may be helpful as an adjunctive treatment for sickle cell disease. Among the possible clinical advantages are: decrease in erythrocyte sickling, Reduced oxidative stress, Enhanced immunity, Decreased inflammation and Increased survival of erythrocytes. Medicinal plants like *G. latifolium* may offer sickle cell disease patients an inexpensive supportive treatment in areas where access to traditional therapies is restricted. Nonetheless, in vitro research and animal models provide the majority of the evidence currently available. To establish

standardised dosage regimens and validate its therapeutic potential, rigorous clinical trials involving human subjects are required [14]. *G. latifolium* extracts are generally safe at moderate doses, according to preclinical toxicological research. However, because strong phytochemicals are present, excessive consumption may have negative effects. Therefore, before the plant is suggested for regular clinical use in sickle cell patients, standardisation of extracts and controlled clinical trials are crucial [15].

Conclusion

With several pharmacodynamic actions pertinent to the treatment of sickle cell disease, *Gongronema latifolium* is a promising medicinal plant. Its antisickling, antioxidant, anti-inflammatory, immunomodulatory, and membrane-stabilizing qualities imply that it may lessen the consequences of illness and enhance the lives of those who are impacted. More clinical research is required to validate its therapeutic efficacy and create standardised treatment protocols, despite promising experimental evidence. Combining contemporary pharmacological research with traditional medicinal plants like *G. latifolium* may help create new, reasonably priced sickle cell disease treatments.

References

1. Nnodim J., Meludu S.C., Dioka C.E., Martin I., Ukaibe N., Ihim A. (2015). Cytokine expression in homozygous sickle cell anaemia. *JKIMSU Journal of Krishna Institute of Medical Sciences University*; 4(1): 34–37.
2. Nnodim J., Udujih B.U., Nwaokoro J.C., Uche U., Onah C. (2015). Alterations of homocysteine in sickle cell anaemia. *Journal of Medical and Biological Science Research*; 1(4): 44–46.
3. Morebise O., Fafunso M.A., Makinde J.M., Olajide O.A., Awe E.O. (2002). Antiinflammatory property of the leaves of *Gongronema latifolium*. *Phytotherapy Research*; 16: 75–77.
4. Nwinyi O.C., Chinedu N.S., Ajani O.O. (2008). Evaluation of antibacterial activity of *Psidium guajava* and *Gongronema latifolium*. *Journal of Medicinal Plants Research*; 2: 189–192.
5. Nnodim J., Chinedu-Madu J.U., Nwaokoro J.C., Bede-Ojimadu O.O. (2021). Effects of *Gongronema latifolium* leaf extract on malondialdehyde concentration of Wistar rats administered with a toxic dose of ibuprofen. *Sumerianz Journal of Biotechnology*; 4(1): 1–3.
6. Ebo N.E.C., Okolie N.J.C., Nnodim J. (2024). Histomorphological changes of heart and kidney of alloxan-induced Wistar rats treated with *Gongronema latifolium* leaf extract. *Medical Case Chronicles and Case Investigations*; 1(1): 4–7.
7. Owu D.U., Nwokocho C.R., Obembe A.O., Essien A.D., Ikpi D.E., Osim E.E. (2012). Effect of *Gongronema latifolium* ethanol leaf extract on gastric acid secretion and cytoprotection in streptozotocin-induced diabetic rats. *West Indian Medical Journal*; 61: 853–860.
8. Omodamiro O.D., Ekeleme C.M. (2013). Comparative study of in vitro antioxidant and antimicrobial activities of *Piper guineense*, *Curcuma longa*, *Gongronema latifolium*, *Allium sativum* and *Ocimum gratissimum*. *World Journal of Medical and Medical Science*; 1: 51–69.
9. Egba S.I., Omeoga H.C., Njoku O.U. (2014). Oral administration of methanol extract of *Gongronema latifolium* (Utazi) upregulates cytokine expression and influences the immune

- system in Wistar albino rats. *World Applied Sciences Journal*; 31: 745–750.
10. Okpala J.C., Sani I., Abdullahi R., Ifedilichukwu H.N., Igwe J.C. (2014). Effects of n-butanol fraction of *Gongronema latifolium* leaf extract on some biochemical parameters in CCl₄-induced oxidative damage in Wistar albino rats. *African Journal of Biochemistry Research*; 8: 52–64.
 11. Ezekwe C.I., Nwodo O.F.C., Ezea S.C. (2014). Chemical and phytochemical components of *Gongronema latifolium* (Asclepiadaceae). *Research Journal of Pharmaceutical, Biological and Chemical Sciences*; 5: 857–866.
 12. Al-Hindi B., Yusoff N.A., Atangwho I.J., Ahmad M., Asmawi M.Z., Yam M.F. (2016). A Soxhlet extract of *Gongronema latifolium* retains moderate blood glucose lowering effect and produces structural recovery in the pancreas of STZ-induced diabetic rats. *Medical Sciences*; 4.
 13. Okon I.A., Ufot U.F., Onoyeraye U.G., Nwachukwu E.O., Owu D.U. (2019). Effect of *Gongronema latifolium* on lipid profile, oral glucose tolerance test and some hematological parameters in fructose-induced hyperglycemia in Wistar rats. *Pharmaceutical and Biomedical Research*; 5: 25–31.
 14. Nnodim J., Meludu S., Dioka C.E., Onah C.E., Ihim A., Atuegbu C. (2014). Trace elements deficiency in patients with homozygous sickle cell disease. *Journal of Advances in Medicine and Medical Research*; 4(21): 3878–3883.
 15. Nnodim J., Udujih B.U., Udujih H.I., Nwaokoro J.C., Obeagu E.I., Aloy-Amadi O. (2015). Hepcidin and erythropoietin level in sickle cell disease. *Journal of Advances in Medicine and Medical Research*; 8(3): 261–265.

Ready to submit your research? Choose ClinicSearch and benefit from:

- fast, convenient online submission
- rigorous peer review by experienced research in your field
- rapid publication on acceptance
- authors retain copyrights
- unique DOI for all articles
- immediate, unrestricted online access

At ClinicSearch, research is always in progress.

Learn more <https://clinicsearchonline.org/journals/clinical-endocrinology-and-metabolism>



© The Author(s) 2026. **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>. The Creative Commons Public Domain Dedication waiver (<http://creativecommons.org/publicdomain/zero/1.0/>) applies to the data made available in this article, unless otherwise stated in a credit line to the data.