Peer-to-Peer Learning Networks: Sickle Cell Disease Education Among Adolescents

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Abstract

Sickle Cell Disease (SCD) presents unique challenges for adolescents, impacting their health, social interactions, and overall well-being. Peer-to-peer learning networks emerge as promising avenues for SCD education among adolescents, offering opportunities for knowledge exchange, support, and empowerment. This review explores the role of peer-to-peer learning networks in SCD education, assessing their effectiveness, benefits, and challenges. By synthesizing existing literature and best practices, this article highlights the significance of peer-led education in raising awareness, enhancing self-management skills, and fostering a sense of community among adolescents living with SCD. Through peer-to-peer learning networks, adolescents can gain valuable insights, connect with peers facing similar experiences, and develop the skills and resilience necessary to navigate life with SCD effectively.

Keywords: Peer-to-Peer Learning, Sickle Cell Disease, Adolescents, Education, Awareness, Support

Introduction

Sickle Cell Disease (SCD) represents a chronic health condition that significantly impacts the lives of adolescents, influencing their physical health, emotional well-being, and social interactions. Adolescents with SCD face unique challenges related to managing their condition, including coping with chronic pain, navigating complex healthcare systems, and maintaining social connections. Peer-to-peer learning networks, characterized by interactions and knowledge

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exchange among individuals of similar age and experiences, offer a promising avenue for addressing these challenges and enhancing SCD education among adolescents. This introduction sets the stage for exploring the role of peer-to-peer learning networks in SCD education, focusing on their potential to empower adolescents, promote self-management skills, and foster a sense of community and support. Adolescents living with SCD often encounter difficulties in understanding and managing their condition, as well as in accessing appropriate support and resources. Peer-to-peer learning networks provide a platform for adolescents to share their experiences, insights, and coping strategies with peers facing similar challenges. By connecting with others who understand their experiences firsthand, adolescents can gain valuable knowledge and support that may not be readily available through traditional educational channels.¹⁻¹⁰

Moreover, peer-to-peer learning networks offer adolescents with SCD a sense of belonging and community, reducing feelings of isolation and loneliness commonly experienced by individuals with chronic health conditions. Through peer interactions, adolescents can forge meaningful connections, build supportive relationships, and develop a sense of empowerment and resilience in managing their condition. Additionally, peer networks serve as a safe space for adolescents to express their thoughts, concerns, and emotions openly, fostering mutual understanding and empathy among participants. Despite their potential benefits, peer-to-peer learning networks in SCD education also face challenges and considerations. Maintaining active participation and engagement among adolescents, especially in virtual or online settings, can be challenging due to competing priorities, limited time, and fluctuating health status. Furthermore, ensuring the accuracy and reliability of information shared within peer networks is essential to prevent the spread of misinformation and misconceptions about SCD. Addressing these challenges requires careful planning, facilitation, and ongoing support from educators, healthcare providers, and community leaders. 11-20

Benefits of Peer-to-Peer Learning Networks

Peer-to-peer learning networks provide adolescents with Sickle Cell Disease (SCD) the opportunity to connect with others who share similar experiences and challenges. By interacting with peers who understand their condition firsthand, adolescents can exchange insights, coping strategies, and practical tips for managing their SCD effectively. This shared experience fosters empathy, understanding, and solidarity among participants, creating a supportive environment where individuals feel valued, validated, and understood. Peer-to-peer learning networks offer a valuable source of peer support and empowerment for adolescents with SCD. Through peer interactions, adolescents can receive encouragement, validation, and emotional support from individuals who are facing similar struggles. Peer support helps adolescents build resilience, confidence, and self-efficacy in managing their condition, empowering them to take an active role in their healthcare and well-being. Additionally, peer networks provide a platform for adolescents to share their accomplishments, celebrate successes, and inspire others within the community. ²¹⁻²⁵

Peer-to-peer learning networks facilitate peer-to-peer education, allowing adolescents to learn from each other's experiences, knowledge, and expertise. Participants can exchange information

about SCD, its symptoms, treatment options, and self-management strategies, thereby enhancing their understanding of the condition and improving their ability to cope with its challenges. Moreover, peer networks provide opportunities for skill-building, such as communication skills, problem-solving abilities, and advocacy techniques, which are essential for effectively managing SCD and navigating healthcare systems. Adolescents with chronic health conditions, such as SCD, often experience feelings of isolation and loneliness due to the unique challenges they face. Peerto-peer learning networks offer a sense of belonging and community, connecting adolescents with others who understand their experiences and can offer support and companionship. By fostering social connections and meaningful relationships, peer networks alleviate feelings of isolation and loneliness, promoting mental well-being and overall quality of life for adolescents with SCD. Peerto-peer learning networks empower adolescents to advocate for themselves and their peers, raising awareness about SCD and advocating for improved support and resources. Through collective action, participants can amplify their voices, share their stories, and advocate for policy changes and initiatives that address the needs of individuals living with SCD. By mobilizing their collective strength and solidarity, peer networks contribute to broader efforts to improve outcomes and enhance the quality of life for adolescents with SCD.²⁶⁻³⁰

Challenges and Considerations

One of the primary challenges facing peer-to-peer learning networks for adolescents with Sickle Cell Disease (SCD) is ensuring accessibility and active participation among participants. Adolescents with SCD may face barriers such as limited mobility, fluctuating health status, and competing priorities, which can impact their ability to engage consistently in peer network activities. Addressing these challenges requires flexibility in scheduling, accommodating individual needs, and providing alternative participation options, such as virtual meetings or asynchronous communication. Sustaining engagement and interest among participants in peer-topeer learning networks can be challenging over time. Adolescents may experience fluctuations in motivation, interest, and willingness to participate in network activities, particularly as they navigate the demands of school, work, and social life. Maintaining a dynamic and engaging environment, incorporating diverse activities and topics, and providing opportunities for active participation and leadership can help sustain engagement and foster a sense of ownership and belonging among participants. Peer-to-peer learning networks rely on the exchange of information and experiences among participants, which may pose challenges in ensuring the accuracy and reliability of information shared. Adolescents with SCD may have varying levels of knowledge and understanding about their condition, leading to the spread of misinformation or misconceptions within peer networks. Facilitators must provide guidance, oversight, and factchecking mechanisms to ensure that information shared within peer networks is accurate, evidence-based, and reliable. 31-35

Peer-to-peer learning networks must strive to be inclusive and representative of the diverse experiences and backgrounds of adolescents with SCD. Individuals from different cultural, socioeconomic, and geographic backgrounds may have unique perspectives, beliefs, and experiences related to their condition, requiring sensitivity and inclusivity in peer network

activities and discussions. Creating a safe and supportive environment where all voices are heard, valued, and respected is essential for fostering meaningful peer interactions and collaboration. Adolescents participating in peer-to-peer learning networks may have concerns about confidentiality and privacy, particularly when sharing personal experiences and sensitive information about their health condition. Maintaining confidentiality and privacy protocols, establishing clear guidelines for sharing personal information, and ensuring that peer network activities are conducted in a safe and respectful manner are essential for building trust and fostering open communication among participants. Effective facilitation and support from trained mentors or facilitators are critical for the success of peer-to-peer learning networks. Facilitators play a key role in guiding discussions, providing information and resources, resolving conflicts, and ensuring that peer network activities align with the goals and objectives of the program. Investing in facilitator training, mentorship, and ongoing support is essential for maintaining the quality and integrity of peer network initiatives.³⁶⁻⁴⁰

Best Practices and Recommendations

Clearly define the goals and objectives of the peer-to-peer learning network, outlining specific outcomes and desired impacts. Establishing clear goals helps focus activities, guide discussions, and measure the effectiveness of the network in meeting its objectives. Offer comprehensive training and support for both peer mentors/facilitators and participants to ensure that they have the knowledge, skills, and resources needed to effectively engage in peer network activities. Training should cover topics such as effective communication, active listening, conflict resolution, confidentiality, and cultural sensitivity. Create a safe and inclusive environment where all participants feel welcome, respected, and valued. Establish ground rules for communication and behavior, and actively promote diversity, equity, and inclusion within the peer network. Encourage open dialogue, mutual respect, and empathy among participants. Offer a variety of activities and topics within the peer network to cater to the diverse interests, preferences, and needs of participants. Incorporate interactive discussions, educational sessions, skill-building workshops, social events, and peer support groups to provide a well-rounded experience for participants.

Encourage active participation and leadership among participants by providing opportunities for them to take on roles and responsibilities within the peer network. Empower participants to lead discussions, organize events, and contribute their ideas and expertise to the network. Recognize and celebrate the contributions of participants to foster a sense of ownership and belonging. Utilize technology and social media platforms to enhance accessibility, engagement, and communication within the peer network. Create online forums, chat groups, or virtual meetings where participants can connect, share information, and engage in discussions remotely. Leverage social media channels to promote peer network activities, share resources, and connect with a broader audience. Regularly evaluate the effectiveness of the peer-to-peer learning network through participant feedback, surveys, and assessments. Use evaluation data to identify strengths, areas for improvement, and opportunities for adaptation. Be willing to adjust strategies, activities, and approaches based on feedback and evolving needs to ensure that the peer network remains relevant and impactful. 41-45

Conclusion

Peer-to-peer learning networks represent a valuable and effective approach for supporting adolescents living with Sickle Cell Disease (SCD). By fostering a supportive environment for learning, sharing experiences, and building connections, these networks empower adolescents to navigate the challenges of their condition with resilience and confidence. Through comprehensive training, inclusive practices, and diverse activities, peer networks provide participants with the knowledge, skills, and support needed to effectively manage their SCD and improve their overall well-being. Furthermore, peer-to-peer learning networks offer numerous benefits, including shared experiences, peer support, enhanced learning, reduced isolation, and opportunities for advocacy and leadership. By leveraging technology and social media platforms, peer networks can reach a broader audience and provide accessible and engaging experiences for participants. Moreover, ongoing evaluation and adaptation ensure that peer networks remain responsive to the evolving needs and preferences of participants, maximizing their impact and effectiveness.

References

- 1. Ata F, Rahhal A, Malkawi L, Iqbal P, Khamees I, Alhiyari M, Yousaf Z, Qasim H, Alshurafa A, Sardar S, Javed S. Genotypic and phenotypic composition of sickle cell disease in the Arab population-a systematic review. Pharmacogenomics and Personalized Medicine. 2023:133-144.
- 2. Piel FB, Williams TN. Sickle cell anemia: history and epidemiology. Sickle Cell Anemia: From Basic Science to Clinical Practice. 2016:23-47.
- 3. Aneke JC, Okocha CE. Sickle cell disease genetic counseling and testing: A review. Archives of Medicine and Health Sciences. 2016;4(1):50-57.
- 4. Bitoungui VJ, Pule GD, Hanchard N, Ngogang J, Wonkam A. Beta-globin gene haplotypes among cameroonians and review of the global distribution: is there a case for a single sickle mutation origin in Africa? Omics: a journal of integrative biology. 2015;19(3):171-179.
- 5. Obeagu EI, Ochei KC, Nwachukwu BN, Nchuma BO. Sickle cell anaemia: a review. Scholars Journal of Applied Medical Sciences. 2015;3(6B):224422-52.
- 6. Obeagu EI. Erythropoeitin in Sickle Cell Anaemia: A Review. International Journal of Research Studies in Medical and Health Sciences. 2020;5(2):22-28.
- 7. Obeagu EI. Sickle Cell Anaemia: Haemolysis and Anemia. Int. J. Curr. Res. Chem. Pharm. Sci. 2018;5(10):20-21.
- 8. Obeagu EI, Muhimbura E, Kagenderezo BP, Uwakwe OS, Nakyeyune S, Obeagu GU. An Update on Interferon Gamma and C Reactive Proteins in Sickle Cell Anaemia Crisis. J Biomed Sci. 2022;11(10):84.
- 9. Obeagu EI, Bunu UO, Obeagu GU, Habimana JB. Antioxidants in the management of sickle cell anaemia: an area to be exploited for the wellbeing of the patients. International Research in Medical and Health Sciences. 2023;6(4):12-17.

- 10. Obeagu EI, Ogunnaya FU, Obeagu GU, Ndidi AC. Sickle cell anaemia: a gestational enigma. European Journal of Biomedical and Pharmaceutical Sciences. 2023;10((9): 72-75
- 11. Obeagu EI. An update on micro RNA in sickle cell disease. Int J Adv Res Biol Sci. 2018; 5:157-8.
- 12. Obeagu EI, Babar Q. Covid-19 and Sickle Cell Anemia: Susceptibility and Severity. J. Clinical and Laboratory Research. 2021;3(5):2768-0487.
- 13. Obeagu EI. Depression in Sickle Cell Anemia: An Overlooked Battle. Int. J. Curr. Res. Chem. Pharm. Sci. 2023;10(10):41-.
- 14. Obeagu EI, Obeagu GU. Evaluation of Hematological Parameters of Sickle Cell Anemia Patients with Osteomyelitis in A Tertiary Hospital in Enugu, Nigeria. Journal of Clinical and Laboratory Research. 2023;6(1):2768-0487.
- 15. Obeagu EI, Dahir FS, Francisca U, Vandu C, Obeagu GU. Hyperthyroidism in sickle cell anaemia. Int. J. Adv. Res. Biol. Sci. 2023;10(3):81-89.
- 16. Njar VE, Ogunnaya FU, Obeagu EI. Knowledge And Prevalence of The Sickle Cell Trait Among Undergraduate Students Of The University Of Calabar. Prevalence.;5(100):0-5.
- 17. Swem CA, Ukaejiofo EO, Obeagu EI, Eluke B. Expression of micro RNA 144 in sickle cell disease. Int. J. Curr. Res. Med. Sci. 2018;4(3):26-32.
- 18. Obeagu EI. Sickle cell anaemia: Historical perspective, Pathophysiology and Clinical manifestations. Int. J. Curr. Res. Chem. Pharm. Sci. 2018;5(11):13-15.
- 19. Obeagu EI, Obeagu GU. Sickle Cell Anaemia in Pregnancy: A Review. International Research in Medical and Health Sciences. 2023 Jun 10;6(2):10-13.
- 20. Obeagu EI, Mohamod AH. An update on Iron deficiency anaemia among children with congenital heart disease. Int. J. Curr. Res. Chem. Pharm. Sci. 2023;10(4):45-48.
- 21. Edward U, Osuorji VC, Nnodim J, Obeagu EI. Evaluation Trace Elements in Sickle Cell Anaemia Patients Attending Imo State Specialist Hospital, Owerri. Madonna University journal of Medicine and Health Sciences. 2022;2(1):218-234.
- 22. Umar MI, Aliyu F, Abdullahi MI, Aliyu MN, Isyaku I, Aisha BB, Sadiq RU, Shariff MI, Obeagu EI. Assessment Of Factors Precipitating Sickle Cell Crises Among Under 5-Years Children Attending Sickle Cell Clinic of Murtala Muhammad Specialist Hospital, Kano. blood.;11:16.
- 23. Obeagu EI. Vaso-occlusion and adhesion molecules in sickle cells disease. Int J Curr Res Med Sci. 2018;4(11):33-35.
- 24. Ifeanyi OE, Stella EI, Favour AA. Antioxidants in the Management of Sickle Cell Anaemia. Int J Hematol Blood Disord: https://symbiosisonlinepublishing.com/hematology/hematology/5. php. 2018.
- 25. Inusa BP, Stewart CE, Mathurin-Charles S, Porter J, Hsu LL, Atoyebi W, De Montalembert M, Diaku-Akinwumi I, Akinola NO, Andemariam B, Abboud MR. Paediatric to adult transition care for patients with sickle cell disease: a global perspective. The Lancet Haematology. 2020;7(4): e329-341.
- 26. Buhari HA, Ahmad AS, Obeagu EI. Current Advances in the Diagnosis and Treatment of Sickle Cell Anaemia. APPLIED SCIENCES (NIJBAS). 2023;4(1).

- 27. Nnodim J, Uche U, Ifeoma U, Chidozie N, Ifeanyi O, Oluchi AA. Hepcidin and erythropoietin level in sickle cell disease. British Journal of Medicine and Medical Research. 2015;8(3):261-5.
- 28. Obeagu EI. BURDEN OF CHRONIC OSTEOMYLITIS: REVIEW OF ASSOCIATIED FACTORS. Madonna University journal of Medicine and Health Sciences. 2023;3(1):1-6.
- 29. Aloh GS, Obeagu EI, Okoroiwu IL, Odo CE, Chibunna OM, Kanu SN, Elemchukwu Q, Okpara KE, Ugwu GU. Antioxidant-Mediated Heinz Bodies Levels of Sickle Erythrocytes under Drug-Induced Oxidative Stress. European Journal of Biomedical and Pharmaceutical sciences. 2015;2(1):502-507.
- 30. Obeagu EI, Bot YS, Opoku D, Obeagu GU, Hassan AO. Sickle Cell Anaemia: Current Burden in Africa. International Journal of Innovative and Applied Research. 2023;11(2):12-14.
- 31. Obeagu EI, Obeagu GU. Sickle Cell Anaemia in Pregnancy: A Review. International Research in Medical and Health Sciences. 2023; 6 (2): 10-13.
- 32. Obeagu EI, Ogbuabor BN, Ikechukwu OA, Chude CN. Haematological parameters among sickle cell anemia patients' state and haemoglobin genotype AA individuals at Michael Okpara University of Agriculture, Umudike, Abia State, Nigeria. International Journal of Current Microbiology and Applied Sciences. 2014;3(3):1000-1005.
- 33. Ifeanyi OE, Nwakaego OB, Angela IO, Nwakaego CC. Haematological parameters among sickle cell anaemia... Emmanuel Ifeanyi1, et al. pdf• Obeagu. Int. J. Curr. Microbiol. App. Sci. 2014;3(3):1000-1005.
- 34. Obeagu EI, Abdirahman BF, Bunu UO, Obeagu GU. Obsterics characteristics that effect the newborn outcomes. Int. J. Adv. Res. Biol. Sci. 2023;10(3):134-143.
- 35. Obeagu EI, Opoku D, Obeagu GU. Burden of nutritional anaemia in Africa: A Review. Int. J. Adv. Res. Biol. Sci. 2023;10(2):160-163.
- 36. Ifeanyi E. Erythropoietin (Epo) Level in Sickle Cell Anaemia (HbSS) With Falciparum Malaria Infection in University Health Services, Michael Okpara University of Agriculture, Umudike, Abia State, Nigeria. Paripex Indian Journal of Research, 2015; 4(6): 258-259
- 37. Ifeanyi OE, Nwakaego OB, Angela IO, Nwakaego CC. Haematological parameters among sickle cell anaemia patients in steady state and haemoglobin genotype AA individuals at Michael Okpara, University of Agriculture, Umudike, Abia State, Nigeria. Int. J. Curr. Microbiol. App. Sci. 2014;3(3):1000-1005.
- 38. Ifeanyi OE, Stanley MC, Nwakaego OB. Comparative analysis of some haematological parameters in sickle cell patients in steady and crisis state at Michael Okpara University of agriculture, Umudike, Abia state, Nigeria. Int. J. Curr. Microbiol. App. Sci. 2014;3(3):1046-1050.
- 39. Ifeanyi EO, Uzoma GO. Malaria and The Sickle Cell Trait: Conferring Selective Protective Advantage to Malaria. J Clin Med Res. 2020; 2:1-4.
- 40. Isa H, Okocha E, Adegoke SA, Nnebe-Agumadu U, Kuliya-Gwarzo A, Sopekan A, Ofakunrin AO, Ugwu N, Hassan AA, Ohiaeri C, Madu A. Strategies to improve healthcare services for patients with sickle cell disease in Nigeria: The perspectives of stakeholders. Frontiers in Genetics. 2023; 14:1052444.

- 41. Martinez RM, Osei-Anto HA, McCormick M, National Academies of Sciences, Engineering, and Medicine. Community Engagement and Patient Advocacy. In Addressing Sickle Cell Disease: A Strategic Plan and Blueprint for Action 2020. National Academies Press (US).
- 42. Issom DZ. Digital Health Interventions to Empower People with Sickle Cell Disease: Toward Patient-Led Design. In Sickle Cell Disease 2022. Intech Open.
- 43. Obeagu EI, Obeagu GU. From Classroom to Home: Strengthening the Continuum of Sickle Cell Disease Knowledge. Elite Journal of Health Science, 2023; 1(1):23-29
- 44. Obeagu EI, Obeagu GU. Incorporating Sickle Cell Disease Curriculum in Schools: An Effective Approach. Elite Journal of Health Science, 2023; 1(1):30-36
- 45. Obeagu EI, Obeagu GU. Community Leaders as Educators: Mobilizing for Sickle Cell Disease Reduction. Elite Journal of Health Science, 2023; 1(1):37-43