Sickle Cell Anaemia- A Synopsis of the Inherited Ailment

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Abstract

Sickle Cell Anaemia (SCA) is one of the most prevalent monogenic disorders. The formation of polymerised haemoglobin leading to erythrocyte rigidity and appearance of characteristic sickle-shaped Red blood Cells (RBCs) resulting in vascular occlusion and haemolysis is central to the molecular pathogenesis of the disease. A major drawback of the disease in children is the development of cerebrovascular disease, hypoxia, and neuro-cognitive impairment. The recurrent episodes of vascular occlusion and inflammation of the vessels lead to progressive organ damage which becomes apparent with age. In addition to hydroxyurea and butyrate treatments, novel methods such as gene therapy and others to prevent complications from SCA are being evaluated. This article reviews the diagnostic approaches, therapies and advancements in SCA treatment with particular emphasis on the future perspective of research towards a cure for SCA.

Keywords: Sickle cell anaemia; Haemoglobin S; Hydroxyurea; Gene Therapy; Genetics

Introduction

Sickle Cell Anaemia (SCA), the most common type of Sickle Cell Disease (SCD) is believed to be a multisystem, monogenic heritable blood disorder, often associated with occurrences of acute illness and progressive organ damage [1,2]. The disease is characterised by abnormally shaped Red Blood Cells (RBCs), which are removed from circulation and destroyed at increased rates, leading to anaemia. In addition to the aforementioned clinical features, the distorted RBCs also cause vascular occlusion, leading to haemolysis and tissue infarction. The first published account of SCD was described in a dental student in 1910 who was presented with pulmonary symptoms [3]. Henrick was the first to coin the term “sickle-shaped” to describe the characteristic appearance of the erythrocytes of the patient (Figure 1) [3].

Figure 1: A peripheral blood smear demonstrating irreversibly sickled cells, and red blood cells (RBCs).

Studies by Pauling and colleagues [4] later determined that a point mutation in the gene coding the β-chain of the haemoglobin molecule resulted in a single amino acid substitution, thereby resulting in sickle cell haemoglobin or HbS. The structure of haemoglobin contains four protein subunits, of which two are β-globin and the remaining two are α-globin subunits. Specifically, β-globin is a larger protein subunit of haemoglobin compared to α-globin. Haemoglobin S (HbS) replaces both the β-globin subunits, and sickle cell anaemia occurs when the patient possesses two copies of the sickle cell genes (HbSS). Later, in 1956 Ingram and colleagues identified the anomaly in the amino acid sequence and described the replacement of amino acid- glutamic acid by valine at 6th position in β-globin protein subunit [5,6]. The substitution causes the abnormal haemoglobin S subunits to form aggregates with other haemoglobin molecules within the RBC. These molecules then adhere together and develop into long and rigid chains which distort the RBC into a characteristic sickle shape, thereby leading to blockage of blood vessels and finally impair blood flow through vessels [7]. This paves the way for a cascade of complications like anaemia, hypoxia, fatigue, delayed growth...
etc. Additionally, the increased breakdown of RBC can cause yellowing of eyes and skin, showing signs of jaundice and the lack of oxygen rich blood results in organ damage of the spleen, liver, lungs, kidney, especially the brain which can also result in stroke [8].

SCA is a prevalent single gene disorder among black Americans affecting nearly 0.26% Black Americans in the United States. Additionally, an approximate of 0.15% of the black population in the United States is afflicted with SCA. The disease is prevalent among the Senegal, Benin, Bantu and Cameroonian tribes of Arica and Arabian-Indian tribes of the Asian haplotype [9]. More than 305,000 infants are conceived every year with SCD for the most part in low and middle income nations, and around 75% of these births occur in sub-Saharan Africa, representing an expanding well-being burden and adding to early adolescence mortality [10]. Every year, 275,000 youngsters with sickle cell weakness are conceived on the planet, primarily in Africa, where around 1.5% of babies have the sickness. In corroboration of this, Nigeria has the largest population of people with SCD in the world with over 150,000 babies born with this serious condition every year [11].

This review discusses the pathophysiology and genetics of the disease in addition to illustrating the various manifestations. Furthermore, a widespread account of the advancements in therapy and future prospective of treatment has been discussed.

Genetics of SCA

Pioneering studies leading to the discovery of Haemoglobin S (HbS) by Linus Pauling and colleagues in 1949 demonstrated for the first time that the production of an abnormal protein could be the cause of a genetic disorder [4]. It was during the same year, that studies revealed an autosomal recessive model of inheritance of the disease [12]. The iron-containing metalloprotein, Haemoglobin (Hb), encoded by the HBB gene, in the mammalian red blood cells (RBCs), responsible for transport of oxygen to the tissues is known to change structure during human development [13]. Broadly classifying, two components exist in adults, namely: major haemoglobin (HbA), and minor haemoglobin A2 (HbA2). The bulk of the haemoglobin during advanced foetal life is foetal haemoglobin (HbF). Each of these haemoglobins is composed of two pairs of dissimilar globin chains. The adult haemoglobins and foetal haemoglobin have α chains combined with β (HbA, α2β2), δ (HbA2, α2δ2), or γ chains (HbF, α2γ2) [13]. Studies have revealed that genetic changes encircling mutations in the haemoglobin gene (HBB) which codes for the protein haemoglobin appears to be the causative rationale for the widespread occurrence of SCA [14]. Additionally, the HBB gene is known to provide instructions for creating the beta-globin (β-globin), the various versions of which are a result of diverse mutations in the HBB gene [14].

A normal molecule of haemoglobin A (HbA) predominates as the adult human haemoglobin tetramer consisting of four globin subunits, typically two α chains and two β chains [15]. Two normal globin chains of α and β constitute the normal structural composition of HbA, while anomalies in the structural components of the β-globin chain often leads to structural malformations thereby inducing the production of defective haemoglobin [15,16]. Briefly, certain mutations in the HBB gene produce an abnormal version of β-globin known as Haemoglobin S (HbS), while other mutations result in additional anomalous forms, namely Haemoglobin C (Hbc) and Haemoglobin E (Hbe) [14]. In individuals’ diagnosed with SCD, at least one of the β-globin subunits is replaced with HbS. However, in SCA, known to be the most common form of SCD, both the β-globin subunits of haemoglobin are replaced by two copies of the variant haemoglobin S (HbSS) resulting in the production of sickled haemoglobin in the red blood cells (RBCs) [6,14]. In other variants of SCD, individuals are often heterozygotes wherein, they possess a copy of the HbS variant and another β-globin namely, Hbc or haemoglobin β-thalassemia (H β-thal) [6].

Pattern of inheritance of SCA

SCA is an autosomal recessive gene disorder [17]. The parents of an individual with an autosomal recessive condition carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition. The health of the parent is rarely affected, but they have one mutated gene (recessive gene) and one normal gene (dominant gene) which leads to the defective condition in the offspring. If the recessive genes from both the parents fuse together, the symptoms of SCD are observed in the offspring (Figure 2) [17]. It follows an autosomal recessive pattern of inheritance, generating a phenotypic ratio of 1:2:1 (Unaffected child: Unaffected carrier child: Affected child). The heterozygous mutated gene carrying individual is the carrier for SCA.

Figure 2: Pictorial representation of the pattern of inheritance of SCA.

Molecular Pathophysiology

In the course of the process of deoxygenation which follows the passage of RBCs in the microcirculation, the Hb molecule undergoes a conformational change which requires the packaging of high concentration of Hb molecules in RBCs to be extraordinarily soluble [18,19]. SCA occurs due to a single
nucleotide polymorphism (SNP) in the short arm of chromosome 11 [20]. In SCA, the presence of HbS results in replacement of the hydrophilic glutamic acid with the hydrophobic amino acid valine at position 6 of the short arm of chromosome 11 of the β-globin chain. This results in a hydrophobic interaction between the haemoglobin molecules, thereby triggering the formation of a large polymer [15,18,19]. The process of polymerization of deoxygenated HbS forms the essential and primary event in the molecular pathogenesis of the disease. This further induces the formation of helical, twisted, rope-like fibres which when grouped together stiffen and distort the shape of RBC into a variety of elongated shapes, decreasing the deformability and conferring it a characteristic “sickle shape” [15,18,21]. The process of polymerization triggers a cascade of several cellular and molecular events, ultimately leading to vascular obstruction and haemolysis which participate in the overall molecular pathophysiological mechanism of the disease (Figure 3) [19]. The series of events following polymerization results in a non-selective increase in membrane cation permeability to sodium, potassium, calcium and magnesium, which upon accumulating in the RBCs, activate several cell membrane transport systems resulting in a cumulative discharge of water causing the RBCs to dehydrate and become dense. Consequently, the concentration of haemoglobin in the cell increases, in turn accelerating the process of polymerization which leads to added dehydration. Such alternate episodes of reversible polymerization and irreversible dehydration forms a dense, irreversible sickled RBC which appears distorted in shape and contributes to vaso-occlusion and haemolysis [21-23].

The elucidation of the molecular mechanisms of SCD remains an unparalleled model, and marks a remarkable achievement in itself. However, factors and processes such as vaso-occlusion and haemolysis require further illustration.

**Diseases Associated with SCA**

Paleness is a typical condition in principally sick patients and results in a high prerequisite for blood transfusions related with poor results [24]. Anaemia is the primary symptom which is associated with SCA. The adaptability of the erythrocytes is crumbled, its life expectancy is abbreviated, and the capillaries in the body are most likely going to impede. This implies that there are insufficient solid red platelets to transport oxygen all through the body. When this happens, a person might present with tiredness, pale skin colour, and delayed puberty. Pain is the most widely recognized entanglement of SCD, and typically most common reason that the patients with SCD go through the crisis. In the sequential effect of SCA, RBCs of most of the patient loose affinity for oxygen and the patient suffers with hypoxia. The patient becomes fatigued easily and experiences shortness of breath.

Acute chest syndrome (ACS), described by fever and new opacification of the lungs on chest radiograph, is the second most common reason for hospitalization in patients with sickle cell ailment and causes 25% of deaths. Rehashed scenes of ACS are related with advancement of incessant lung sickness, with dynamic hypoxemia and diminished survival [25]. About 30% of adult patients with SCA suffer from pulmonary hypertension, and the complication is a strong negative prognostic factor. Recent examinations have uncovered that adenosine concentrations are fundamentally lifted in SCD and add to infection pathology by initiating adenosine receptors on red platelets. Apart from adenosine, hypoxia additionally causes haemoglobin discharge through haemolysis [26]

SCA influences vast majority of tissues in the body. SCA brings about both anatomical adjustments and aggravations in renal capacity. Majority of the sickle cell iron deficient patients have enormous kidneys, which may be attributed to expanded renal blood volume [27]. The imperfect in concentrating urine capacity in sickle cell trait is accepted to be the result of intracellular polymerization of HbS in erythrocytes that causes impediment of tiny vessels in the vasa recta of the renal medulla. The seriousness of the deformity in concentration of urine may be connected to the level of sickling process of erythrocytes [28]. During later stages of this disorder, renal dysfunction results in sickle nephropathy.

In a severe situation, large number of sickle cells gets trapped in the spleen and makes it enlarged. Symptoms shown by the individual are extreme thirst, left abdominal pain, weakness, pale lips, fast heartbeat and breathing. This event can be life threatening, in most cases. Medically, this discomfort is known as splenic sequestration [29].

In the worst case scenario, major organ failure is the critical situation which arises. Initially, clinically apparent as autosplenectomy, followed by cerebral infarction during childhood and finally in young adulthood as end stage renal failure (glomerulosclerosis), disabling leg ulcers, generalized osteonecrosis, intracranial haemorrhage, retinopathy and sickle chronic lung disease [30].
Diagnosis of SCA

Prenatal screening

Currently it is feasible for specialists to analyse sickle cell ailment before birth. This can be accomplished by utilizing a specimen of amniotic fluid (liquid in the sac encompassing a developing foetus) or tissue taken from the placenta. This test should be possible in the initial couple of long stretches of pregnancy. Pre-birth conclusion of sickle cell iron deficiency is conceivable utilizing amniocentesis or chorionic villus sampling. The specimen received is then tried for DNA investigation of the foetal cells. Testing before birth should be possible as early as first trimester of pregnancy. This testing examines for the sickle haemoglobin gene instead of the abnormal haemoglobin [31].

New-born screening

In the United States, the District of Columbia, and the U.S. domains require that each child is tried for SCD as a component of an infant screening program. It is essential to analyse SCD right on time to avoid entanglements. In infant screening programs, blood from a foot rear area prick is gathered in "spots" on a unique paper. The haemoglobin from this blood is then dissected in exceptional labs. Results will then be discussed with the parents. If the test shows sickle haemoglobin, a second blood test is done to confirm the diagnosis [32].

It is noteworthy that, when a child has sickle cell trait or SCD, a future sibling, or the child’s own future may be at risk. These possibilities should be discussed with the primary care doctor, haematologist, and/or genetic counsellor [33].

For young individuals displaying symptoms of SCA, blood smear is observed under microscope to check the abnormal sickle-shaped cells. Smearing of blood using a special low-oxygen preparation is done for testing and this is referred to as sickle prep. Other prep tests can likewise be utilized to distinguish the strange HbS. Preparatory tests also include dissolvability tests performed on the blood samples. The infection can be affirmed by particularly evaluating the kinds of haemoglobin, utilizing a haemoglobin electrophoresis test [34]. This electrophoresis test recognizes the haemoglobin in the blood by isolating them. The partition of the diverse haemoglobins is conceivable due to the unique electrical charges they have on their protein surfaces, causing them each to move distinctively in an electric field [35].

Treatment Procedures

Blood transfusions are utilized to treat extreme anaemia. A sudden exacerbation of weakness caused due to disease or augmentation of the spleen is a typical purpose behind a transfusion. Nonetheless, excess blood transfusions might cause medical issues as a result of the excess iron pumping in the blood. This over-burden, called haemosiderosis, can harm liver, heart, pancreas and different organs, prompting maladies, for example, diabetes mellitus. Press chelation treatment ought to be begun in patients with SCD accepting general blood transfusions to lessen abundant press levels.

SCD is a genetic disorder endemic in malaria prone areas. Therefore, the need for prompt initiation of efficacious antimalarial therapy in patients with SCD is of utmost importance. In a randomized controlled trial, 60 children with SCD and acute uncomplicated malaria were treated with artesunate-amodiaquine (AA) also known as artemether-lumefantrine (AL). It was observed that AA and AL showed similar clinical and parasitological effects in the SCD and non-SCD groups. This could be attributed to alterations in WBC and platelet counts [36].

The goals of treating SCD are to prevent infections, relieve pain, prevent organ damage, strokes, treat anaemia and control complications [23].

Treating pain

The most difficult sign of SCA appears to be the pain. Mild pain is often treated with heating pads and counter medicines. In order to suppress the pain, fluids and pain-controlling medicines are administered. The body tends to become dehydrated due to pain, thus fluids are conferred either orally or intravenously. Severe pain may need to be treated in a hospital [37].

Common medicines used to treat pain crises include acetaminophen, narcotics and non-steroidal anti-inflammatory drugs (NSAIDs). Mild to moderate pain are usually treated with NSAIDs or acetaminophen. Moderate to severe pain is often treated with narcotics. The narcotic may be used alone or in combination with NSAIDs or acetaminophen. Certain counter pain medications available are aspin and ibuprofen. In case of severe pain opioid (i.e. morphine) medications is given on a daily basis, along with additional pain medication [31].

Several parallel treatments have been used as combination or primary therapy for acute episodes of sickle cell pain. Nonsteroidal antiinflammatory agents such as ketorolac, piroxicam, and ibuprofen have been assessed for monotherapy and in combination with opiates for vasoocclusive crises associated with SCA [38]. A Nigerian trial found that piroxicam was as effective as aspirin for the treatment of vasoocclusive crisis and had fewer side effects [39]. Furthermore, in a blinded, crossover trial, ketorolac provided superior pain control with fewer side effects than meperidine [40-42].

Prevention

HbF is a variant of Haemoglobin, which is found to impede the advance of HbS quality in an individual. In an examination, it was demonstrated that large amounts of HbF in children may diminish the seriousness of the sickness. This could be possible because of the hindrance of polymerization of HbS by HbF [43].

Hydroxyurea treatment might be useful in decreasing emergencies and the requirement for transfusions. Hydroxyurea does not cure sickle cell infection. It can incredibly lessen a portion of the complexities of the illness. It doesn't work effectively, if not taken as recommended. For the major part, it takes a while before any positive outcomes are observable. It is vital to recollect that hydroxyurea must be taken each day for it

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to function effectively. Those with extreme sickle cell iron deficiency may profit by daily consumption of hydroxyurea. Hydroxyurea is utilized to anticipate difficult pain but not to treat them [44].

Hydroxyurea treatment enables the red platelets to remain round and adaptable. This gives them a chance to move effortlessly through venules. To some extent, this happens because hydroxyurea expands the measure of foetal haemoglobin (HbF) in red platelets. As mentioned above, infants have HbF when they are born. HbF shields them from sickle cell complexities (medical issues) amid the first couple of months of life. With an expanded measure of HbF, red platelets are less inclined to change into sickle or banana shape. In most of us, the measure of HbF diminishes after the initial couple of long stretches of life. A few people have more HbF than others. Individuals with sickle cell disease who have more elevated amounts of HbF for the most part have less difficulties of the ailment [45]. The exact mechanism of activity of hydroxyurea isn’t known in spite of the fact that the enlistment of HbF amalgamation and macrocytosis appear to be the possible mechanism [46]. Hydroxyurea was endorsed by the Food and Drug Administration (FDA) for the treatment of grown-ups with direct to extreme SCA and sickle-β-thalassemia [47].

The side effect of hydroxyurea includes myelotoxicity, which is reversible upon discontinuation of the medication. Opposite reactions that have been portrayed in animal studies considers its teratogenic and cancer-causing potential. Also, hydroxyurea has idiosyncratic reactions that happen in a few people and this might be a hereditary or epigenetic phenomenon. Studies report three idiosyncratic side effects of hydroxyurea in four patients with sickle cell iron deficiency (SS). Desquamating rash, diminished charisma and incomplete complex seizures have been already portrayed in patients who are hydroxyurea responders. In spite of the fact that these responses are uncommon, they can be not serious in few patients [48].

Preventing infections associated with SCA

Bacterial contaminations can be a noteworthy complexity of SCA, yet regularly they can be forestalled or treated. Children who have sickle cell illness hints at early onset of infections and diseases, for example, fever, trouble breathing, or restricted bone agony.

To avert diseases in infants and youthful kids, medications include:

- Daily dosages of penicillin: Treatment may start ahead of schedule at 2 months of age and proceed until the point when the kid is no less than 5 years of age.
- All routine immunizations (counting a yearly influenza shot), in addition to vaccination(s) against Streptococcus pneumonia.

Adults who have sickle cell ailment ought to likewise get influenza shots consistently and get inoculated against pneumococcal diseases. The two grown-ups and youngsters are in danger for an assortment of diseases, for example, pneumonia and bone contaminations. They ought to be inspected at whatever point they encounter fever since early finding and treatment result in better results [49].

Approaches and advancements in the therapy of SCA

The current understanding of the molecular pathophysiology of SCA has led to the development of several independent approaches and advancements in the therapy. A notable few of these theories, namely, pharmacologic induction of HbF, chemical inhibition of Hb polymerization and reduction of the intracellular haemoglobin concentration have been subjected to thorough laboratory and clinical investigations.

Pharmacological induction of HbF: Several biochemical, clinical, epidemiological and laboratory investigations have converged to support the notion that HbF is a very potent inhibitor of the deoxygenation of HbS in vitro [50]. Patients homozygous for SCA and belonging to the Bedouin area of Saudi Arabian Peninsula and central provinces of India, typically have relatively high amounts of HbF associated with relatively mild clinical manifestations of the disease [43,51-53]. Based on these compelling biochemical and clinical evidence, it was proposed that the pharmacological inhibition of HbF production may act as an effective therapeutic strategy to benefit patients with SCA. Additionally, with studies elucidating the cloning of different globin genes and their mechanisms responsible for regulation gained importance in the late 1970s and 1980s [54-56]. It became very evident that epigenetic factors such as DNA methylation and histone acetylation were influential towards developmental regulation and the expression of globin genes [57-59]. This led to the theory that drugs which alter the epigenetic configuration of the γ-globin genes may furnish a viable therapeutic approach to the pharmacological induction of HbF, thereby influencing the inception of the first pharmaceutical agent, namely 5-azacytidine [53].

5-Azacytidine, an antineoplastic drug was the first agent used to induce HbF expression via epigenetic silencing of the γ-globin genes in adult life, the rationale for which was based on the discovery that the actively transcribed adult β-globin genes are hypomethylated and the non-transcribed foetal γ-globin genes are hypermethylated. Contrastingly, in foetal life, the adult β-globin genes are hypermethylated and the nontranscribed foetal γ-globin genes are hypomethylated [53,58,59]. Furthermore, in addition to inhibiting the methylation of DNA, it was observed to induce a marked increase in the levels of HbF in anaemic baboons [60]. Inspite of these promising results, limited clinical trials testing demonstrated the concerns regarding potential carcinogenicity of this drug in patients with SCA. Thus, focus was shifted towards other anti-tumour drugs, particularly hydroxyurea which showed increase in the production of HbF in non-human primates and patients with SCA [61-63].

Chemical inhibition of Hb polymerization: The controversy over the mechanism of inhibition of HbF by 5-Azacytidine laid the groundwork for a study in which hydroxyurea, a relatively non-toxic, 5-phase-specific chemotherapeutic agent with readily reversible myelosuppressive effects which does inhibit the
enzyme DNA methyltransferase was shown to result in a marked increase in the levels of HbF in baboons [64].

Hydroxyurea, an inhibitor of ribonucleotide reductase that was in use for several years for the treatment of myeloproliferative disorders is an orally available drug, relatively well tolerated and easy to use. After demonstration of its ability to induce the production of increased levels of HbF in baboons, a number of clinical trials were carried out in patients with SCA who were administered doses sufficient to cause mild myelosuppression. These patients showed marked increase in the number of HbF–containing red cells and the percentage of HbF in addition to the level of HbF per F cell [62,65]. In a separate national multicentre study comprising 299 adults with moderate to severe SCA, it was observed that the drug was relatively nontoxic and showed a marked decrease in the frequency and severity of painful crises. In addition to this, considerable reduction in the incidence of acute chest syndromes, the need for transfusions and hospitalizations were also observed [66,67]. Additional studies in children with SCA demonstrated the clinical efficacy and short-term safety of hydroxyurea resulting in a substantial reduction of hospitalization [68-70]. Albeit, similarities were observed in the HbF inducing activity of hydroxyurea and 5-azacytidine among anaemic baboon and patients with SCA. However, its molecular targets and mechanisms of action require further elucidation. The original theory suggested that hydroxyurea may elevate HbF levels by accelerating the differentiation of erythroid formation in the bone-marrow, leading to the appearance of ‘foetal-like’ cells in the peripheral blood [71]. More conventional theories have shown that hydroxyurea generates NO in vivo resulting in the activation of the NO/cGMP signalling pathway leading to the up-regulation of γ-globin gene expression in patients with SCA [72]. Additionally, the hydroxyurea has other effects that may be beneficial to patients with SCA. For example, hydroxyurea has shown to reduce the number of reticulocytes and young, low-density sickled red cells to the endothelium and also decrease the expression level of soluble VCAM-1 [73,74].

Butyrate: Due to the concerns about the long-term administration of an antitumor drug and its potential serious side effects, there has been considerable interest in identifying safe and effective inducers for the production of HbF. Initially, studies on recombinant human erythropoietin suggested promising results in stimulating HbF production in patients with SCA [75]. However, subsequent studies on human erythropoietin administration unaided or in combination with hydroxyurea have yielded conflicting results [76-78]. This led to a considerable interest in the physiologic and pharmacologic roles of butyrate in the regulation of HbF production.

A short-chain fatty acid that inhibits HDAC was shown to stimulate embryonic or foetal gene expressions in sheep, mice, chicken and baboons [79-81]. Furthermore, the switch from HbF to HbA in infants with diabetic mothers is delayed due to the increased serum concentration of amino-n-butyric acid [82]. Additionally, studies involving intermittent administration of arginine butyrate in patients with SCA resulted in sustained induction in the production of HbF [83]. While there exists considerable evidence of this agent in the treatment of SCA, there is an added need to assess the efficacy and safety of the administration of this drug.

The butyrates are perhaps the first class of drugs designed to transcriptionally activate specific genes—in this particular case, to reactivate the developmentally silenced foetal globin genes. In recent times, research focusing on the conventional use of sodium butyrate is used for foetal haemoglobin induction is being studied. It is suggested that sodium butyrate treatment can be replaced by safer therapeutic tools like microRNAs, small non-coding RNAs that control number of epigenetic mechanisms. In a study by Tayabi and colleagues, it was demonstrated that gamma chain was 5.9-fold up-regulated in the treated group [84].

It can be assumed that sodium butyrate can up-regulate gamma chain gene, and change the expression of micro RNAs (miRNAs). These results can be profitable in future studies to discover therapeutics suitable for such disorders.

Bone marrow transplantation: The concept of replacing the source of defective sickled RBCs, the bone marrow, with a healthy bone marrow which produces normal RBC is an insightful therapeutic approach in the treatment of SCA. However, for several years the possibility of such an approach was considered uncertain for a non-malignant disorder like SCA, due to the high chances of mortality. Nonetheless, recent advancements in immunosuppressive therapy and supportive care resulted in reduction in the mortality rates following bone marrow transplantation (BMT). Patients with β-thalassemia, after treatment with BMT showed a considerable 90% increase in survival rates [85]. Promising results such as those described above led to improved interest in the use of BMT as a therapy for treatment of SCA and clinical trials were conducted in Europe and USA to exploit the full potential of this therapy [86-88]. However, the foremost drawback in using BMT as a therapeutic approach lies in the fact that it is almost impossible to find a matched sibling donor as suitable candidate for transplantation [89]. Nevertheless, efforts and clinical trials are underway to increase the availability of sources of haematopoietic stem cells (HSCs) for transplantation to identify procedures to reduce the mortality and morbidity rates in adults [90]. Currently, BMT appears to be the only effective curative theory and the main challenge that exists is to make it commonly available to patients.

Future Prospects

Gene therapy

In order to cure or prevent SCD, researchers could settle patient’s undifferentiated cells of hemoglobin by utilizing quality altering property and afterward transplant the changed cells back into the patient. In the event that specialists could alter a patient’s hematopoietic undifferentiated cells, either to settle the β-globin transformation or to restart the generation of fetal hemoglobin. Thereafter, utilize them to repopulate the bone marrow. As we know that, there is rare compelling reason to locate a good bone-marrow contributor.
Repairing or replacing the β-globin gene is not a new idea, but researchers have found fresh promise in the relative ease of the Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR) technique, achieving early success in mouse models and genome editing system in the treatment of cancer [91]. The materials required for CRISPR are easily accessible. Researchers need only an enzyme that cuts DNA, such as the endonuclease Cas9, and a guide RNA molecule that directs the enzyme to cut the gene of interest. Instead of leaving the cell to make its own error-prone repairs to the CRISPR-induced cuts, researchers supply further guidance by delivering a strand of DNA that contains the corrected snippet of the haemoglobin sequence—a kind of ‘repeat after me’. This approach is called homology-directed repair [84].

Repairing or supplanting the β-globin quality isn’t another thought, yet researchers have discovered new guarantee in the relative simplicity of the CRISPR method, making early progress in mouse models. CRISPR-Cas9 was adjusted from a normally infectious’ system in the treatment of cancer [91]. The materials required for CRISPR are easily accessible. Researchers need only an enzyme that cuts DNA, such as the endonuclease Cas9, and a guide RNA molecule that directs the enzyme to cut the gene of interest. Instead of leaving the cell to make its own error-prone repairs to the CRISPR-induced cuts, researchers supply further guidance by delivering a strand of DNA that contains the corrected snippet of the haemoglobin sequence—a kind of ‘repeat after me’. This approach is called homology-directed repair [84].

Red blood cells produced from the edited hematopoietic stem cells of patients with SCD produced 25% more fetal haemoglobin than unedited cells, and the edited cells successfully engrafted in mice. After those cells are transplanted into mice with smothered invulnerable frameworks, their numbers tumble off: 10% of the human cells that were joined, or engrafted, into the bone marrow of the mice created solid haemoglobin. Future research can be steered in this direction, where alteration in various factors can attain the results [93]. Another study performed in patients with SCD who received gene therapy with the use of lentiviral gene addition of an anti-sickling β-globin variant, showed that normal blood cell counts were attained after the transduced stem cells were engrafted. While adverse events related to the BB305-transduced cells were not observed, and the pattern of vector integration remained polyclonal without clonal dominance, insertional oncogenesis was reported in clinical gene-transfer studies with gamma retroviral vectors [94,95].

This successful experiment suggests that SCD can be cured by using gene therapy technique; however, it is not implemented in regular patients of SCA due to the fact that it has to be tested on a large scale. In this regard, it opens the door to further research by considering a large population to see its success.

Conclusion

International health care providers and health care agencies are faced with multiple challenges to overcome the burden of SCA among children in backward countries. In order to spearhead the process, emphasis should be laid on diagnostic procedures such as new-born screening, and anti-microbial prophylaxis. While these are affordable resources available in low income countries, efforts should also be made to improve morbidity as well as quality of life. Many advances have taken place in the past decade to improve treatment options for SCA. However, most of these treatments are expensive and dependent on medical infrastructure and only a few patients with SCA are able to afford treatments such as gene therapy and hematopoietic stem cell transplantation. There is a need for active measures and collaborations to be deployed on the part of the government to effectively implement population screening. Additionally, counseling programs and epidemiological studies in SCA stricken countries are necessary to assess current and future health needs.

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