Sickle Cell Disease: 2022 Reading List

This brief bibliography contains a variety of resources focused on sickle cell disease. The list contains three sections: factsheets and toolkits for consumers, reports and recommendations for professionals, and peer-reviewed journal articles. All the resources in this list are available to read and download for free. For more resources on sickle cell disease, contact the OMH Knowledge Center or browse the online catalog.

Consumer Materials

Do You Use the Emergency Department for Care of Sickle Cell Disease? What to Know Before You Go. Centers for Disease Control and Prevention (CDC), 2017.
- English version: Click here to read
- Spanish version: Click here to read

Toolkit for Living Well with Sickle Cell Disease. Centers for Disease Control and Prevention (CDC), 2014. Click here to read

Sickle Cell Disease: A Family Guide. (4th edition) Savage B., et al. New Jersey Department of Health; Division of Family Services; Newborn Screening and Genetic Services, 2017. Click here to read

Understanding Sickle Cell Disease. American Society of Hematology (ASH), 2019. Click here to read

Hydroxyurea for Sickle Cell Disease: Treatment Information from the American Society of Hematology. American Society of Hematology (ASH), 2020. Click here to read

A Century of Progress – Sickle Cell Disease: Milestones in Research and Clinical Progress. National Heart Lung and Blood Institute (NHLBI), 2018. Click here to read

- English version: Click here to read
- Spanish version: Click here to read

- English version: Click here to read
- Spanish version: Click here to read

Reports

Addressing Sickle Cell Disease: A Strategic Plan and Blueprint for Action. National Academies of Sciences, Engineering, and Medicine, 2020. Click here to read


Strategies from the Field: Data Collection. National Center on Birth Defects and Developmental Disabilities, Division of Blood Disorders, 2018. Click here to read

Strategies from the Field: Health Promotion. National Center on Birth Defects and Developmental Disabilities, Division of Blood Disorders, 2018. Click here to read

Journal Articles


Barriers to Care and Quality of Primary Care Services in Children with Sickle Cell Disease. Jacob E., et al. Journal of Advanced Nursing, v. 72, #6 (June), p. 1417-1429, 2016. Click here to read


Lifespan Care in SCD: Whom to Transition, the Patients or the Health Care System? Minniti C.P., et al. American Journal of Hematology, v. 92, #6 (June), p. 487-489, 2017. Click here to read


The OMH Knowledge Center contains a collection of over 70,000 documents, reports, books, consumer materials, and journal articles related to the health status of racial and ethnic minority populations. Knowledge Center staff can respond to requests from the public by providing selective bibliographies of articles and books in its collection.

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