

DAFTAR PUSTAKA

- Adeel Rehman, Kumayl Abbas Meghji, Haji Abdullah Memon, Ahmed Mustafa Burney, Wasi Ur Rehman Siyal, & Muhammad Jawad. (2024). Correlation between hepatomegaly and transfusion frequency in thalassemia patients: A cross-sectional study. *The Professional Medical Journal*, 31(02), 277–282. <https://doi.org/10.29309/tpmj/2024.31.02.7729>
- Ahmed, M. H., Ghatge, M. S., & Safo, M. K. (2020). Hemoglobin: Structure, Function and Allostery. In *Subcellular Biochemistry* (Vol. 94, pp. 345–382). Springer. https://doi.org/10.1007/978-3-030-41769-7_14
- Andika, O., & Puspitasari, A. ; (2019). *Buku Ajar Mata Kuliah Hematologi Diterbitkan oleh UMSIDA PRESS*.
- Anggoro, A. (2017). *Talasemia pada Kehamilan* (Vol. 1).
- Ansari-Moghaddam, A., Adineh, H. A., Zareban, I., Mohammadi, M., & Maghsoodlu, M. (2018). The survival rate of patients with beta-thalassemia major and intermedia and its trends in recent years in Iran. *Epidemiology and Health*, 40. <https://doi.org/10.4178/epih.e2018048>
- Arfie, N. G., Zulkarnaen, B. S., & Sudarmanto, S. (2022). Efektivitas Deferasirox pada Pasien Talasemia Mayor: Artikel Review. *Jurnal Sains Dan Kesehatan*, 4(3), 354–362. <https://doi.org/10.25026/jsk.v4i3.1159>
- Armina, A., & Pebriyanti, D. K. (2021). Hubungan Kepatuhan Transfusi Darah dan Kelasi Besi dengan Kualitas Hidup Anak Talasemia. *Jurnal Akademika Baiturrahim Jambi*, 10(2), 306. <https://doi.org/10.36565/jab.v10i2.336>
- Asadov, C., Alimirzoeva, Z., Mammadova, T., Aliyeva, G., Gafarova, S., & Mammadov, J. (2018). β -Thalassemia intermedia: a comprehensive overview and novel approaches. *International Journal of Hematology*, 108(1), 5–21. <https://doi.org/10.1007/s12185-018-2411-9>
- Babker, A. M. A. (2022). An overview on thalassemia and challenges during COVID-19. *International Journal of Health Sciences*, 3207–3220. <https://doi.org/10.53730/ijhs.v6ns1.5446>
- Bodge, M., & Cumpston, A. (2019). Pharmacology of Drugs Used in Hematopoietic Cell Transplantation. In *Hematopoietic Cell Transplantation for Malignant Conditions* (pp. 19–35). Elsevier. <https://doi.org/10.1016/B978-0-323-56802-9.00002-X>
- Bouguila, J., Besbes, G., & Khochtali, H. (2015). Skeletal facial deformity in patients with β thalassemia major: Report of one Tunisian case and a review of the literature. *International Journal of Pediatric Otorhinolaryngology*, 79(11), 1955–1958. <https://doi.org/10.1016/j.ijporl.2015.08.037>
- Calderon Moreno, R., Navas-Acien, A., Escolar, E., Nathan, D. M., Newman, J., Schmedtje, J. F., Diaz, D., Lamas, G. A., & Fonseca, V. (2019). Potential Role of Metal Chelation to Prevent the Cardiovascular Complications of Diabetes. In *Journal of Clinical Endocrinology and Metabolism* (Vol. 104, Issue 7, pp. 2931–2941). Endocrine Society. <https://doi.org/10.1210/jc.2018-01484>
- Chiabrando, D., Mercurio, S., & Tolosano, E. (2014). Heme and erythropoiesis: More than a structural role. In *Haematologica* (Vol. 99, Issue 6, pp. 973–983). Ferrata Storti Foundation. <https://doi.org/10.3324/haematol.2013.091991>
- Clénin, G. E., Cordes, M., Huber, A., Schumacher, Y., Noack, P., Scales, J., & Kriemler, S. (2016). Iron deficiency in sports - definition, influence on performance and

- therapy. *Schweizerische Zeitschrift Fur Sportmedizin Und Sporttraumatologie*, 64(1), 6–18. <https://doi.org/10.4414/smw.2015.14196>
- Dorgaleleh, S., Barahouie, A., Naghipoor, K., Dastaviz, F., Ghodsalavi, Z., & Oladnabi, M. (2020). Transfusion Related Adverse Effects on Beta-Thalassemia Major and New Therapeutic Approaches: A Review Study. *Int J Pediatr*, 8(79). <https://doi.org/10.22038/ijp.2020.46749.3794>
- Edi, I. G. M. S. (2017). *FAKTOR-FAKTOR YANG MEMPENGARUHI KEPATUHAN PASIEN PADA PENGOBATAN: TELAAH SISTEMATIK (FACTORS AFFECTING THE PATIENT ADHERENCE TO MEDICAL TREATMENT: A SYSTEMATIC REVIEW)*.
- Elalfy, M. S., Hamdy, M., Adly, A., Ebeid, F. S. E., Temin, N. T., Rozova, A., Lee, D., Fradette, C., & Tricta, F. (2023). Efficacy and safety of early-start deferiprone in infants and young children with transfusion-dependent beta thalassemia: Evidence for iron shuttling to transferrin in a randomized, double-blind, placebo-controlled, clinical trial (START). *American Journal of Hematology*, 98(9), 1415–1424. <https://doi.org/10.1002/ajh.27010>
- Eleftheriou, A., & Angastiniotis, M. (2023). *GLOBAL EPIDEMIOLOGY OF B-THALASSAEMIA*. www.thalassaemia.org.cy
- Entezari, S., Haghi, S. M., Norouzkhani, N., Sahebazar, B., Vosoughian, F., Akbarzadeh, D., Islampanah, M., Naghsh, N., Abbasalizadeh, M., & Deravi, N. (2022a). Iron Chelators in Treatment of Iron Overload. In *Journal of Toxicology* (Vol. 2022). Hindawi Limited. <https://doi.org/10.1155/2022/4911205>
- Entezari, S., Haghi, S. M., Norouzkhani, N., Sahebazar, B., Vosoughian, F., Akbarzadeh, D., Islampanah, M., Naghsh, N., Abbasalizadeh, M., & Deravi, N. (2022b). Iron Chelators in Treatment of Iron Overload. In *Journal of Toxicology* (Vol. 2022). Hindawi Limited. <https://doi.org/10.1155/2022/4911205>
- Farashi, S., & Hartevelde, C. L. (2018). Molecular basis of α -thalassemia. In *Blood Cells, Molecules, and Diseases* (Vol. 70, pp. 43–53). Academic Press Inc. <https://doi.org/10.1016/j.bcmd.2017.09.004>
- Farmakis, D., Porter, J., Taher, A., Cappellini, M. D., Angastiniotis, M., Eleftheriou, A., Alassaf, A., Angastiniotis, M., Angelucci, E., Aydinok, Y., Bou-Fakhredin Rayan, R., Brunetta, L., Cappellini, M. D., Constantinou, G., Daar, S., De Sanctis, V., Dusheiko, G., Elbard, R., Eleftheriou, A., ... Yardumian, A. (2022a). 2021 Thalassaemia International Federation Guidelines for the Management of Transfusion-dependent Thalassaemia. *HemaSphere*, 6(8). <https://doi.org/10.1097/HS9.0000000000000732>
- Farmakis, D., Porter, J., Taher, A., Cappellini, M. D., Angastiniotis, M., Eleftheriou, A., Alassaf, A., Angastiniotis, M., Angelucci, E., Aydinok, Y., Bou-Fakhredin Rayan, R., Brunetta, L., Cappellini, M. D., Constantinou, G., Daar, S., De Sanctis, V., Dusheiko, G., Elbard, R., Eleftheriou, A., ... Yardumian, A. (2022b). 2021 Thalassaemia International Federation Guidelines for the Management of Transfusion-dependent Thalassaemia. *HemaSphere*, 6(8). <https://doi.org/10.1097/HS9.0000000000000732>
- Farmakis, D., Triposkiadis, F., Lekakis, J., & Parissis, J. (2017). Heart failure in haemoglobinopathies: pathophysiology, clinical phenotypes, and management. In *European Journal of Heart Failure* (Vol. 19, Issue 4, pp. 479–489). John Wiley and Sons Ltd. <https://doi.org/10.1002/ejhf.708>

- Fibach, E., & Rachmilewitz, E. A. (2017). Pathophysiology and treatment of patients with beta-thalassemia - an update. In *F1000Research* (Vol. 6). Faculty of 1000 Ltd. <https://doi.org/10.12688/f1000research.12688.1>
- Firdaus, H. F. F., Karnita, R., & Isnaini, W. (2023). *Perancangan Website Sebagai Media Informasi Mengenai Thalassemia Untuk Remaja TUGAS AKHIR*.
- Gell, D. A. (2018). Structure and function of haemoglobins. *Blood Cells, Molecules, and Diseases*, 70, 13–42. <https://doi.org/10.1016/J.BCMD.2017.10.006>
- Gumilang, L., Dewi Judistiani, T., Aryuti Nirmala, S., & Wibowo, A. (2021). *Korelasi Asupan Zat Besi dan Protein dengan Kadar Ferritin Serum Ibu Hamil di Kabupaten Waled dan Sukabumi*. <https://doi.org/10.15294/higeia/v2i3/44805>
- Hartina, M., Politeknik, U., Palembang, K., & Yuliany, E. H. (2017). *KORELASI TRANSFUSI DARAH DENGAN KADAR FERITIN PASIEN THALASSEMIA CORRELATION BLOOD TRANSFUSION AND FERRITIN LEVELS IN PATIENTS WITH THALASSEMIA DISEASE: A LITERATURE REVIEW*. <https://www.researchgate.net/publication/340501921>
- KEMENKES. 2018. PEDOMAN NASIONAL PELAYANAN KEDOKTERAN TATA LAKSANA THALASEMIA. KEPUTUSAN MENTERI KESEHATAN REPUBLIK INDONESIA NOMOR HK.01.07/MENKES/1/2018. Kementerian Kesehatan Republik Indonesia. Jakarta.
- Kontoghiorghes, G. J., Kleanthous, M., & Kontoghiorghes, C. N. (2020). THE HISTORY OF DEFERIPRONE (L1) AND THE COMPLETE TREATMENT OF IRON OVERLOAD IN THALASSAEMIA. *Mediterranean Journal of Hematology and Infectious Diseases*, 12(1), e2020011. <https://doi.org/10.4084/mjhid.2020.011>
- Kowalak, J. P., Welsh, W., & Mayer, B. (2016). *Buku Ajar Patofisiologi*.
- Lee, J.-S., Rhee, T.-M., Jeon, K., Cho, Y., Lee, S.-W., Han, K.-D., Seong, M.-W., Park, S.-S., & Lee, Y. K. (2022). Epidemiologic Trends of Thalassemia, 2006–2018: A Nationwide Population-Based Study. *Journal of Clinical Medicine*, 11(9), 2289. <https://doi.org/10.3390/jcm11092289>
- Lee, W. J., Mohd Tahir, N. A., Chun, G. Y., & Li, S. C. (2024). The impact of chelation compliance in health outcome and health related quality of life in thalassaemia patients: a systematic review. In *Health and Quality of Life Outcomes* (Vol. 22, Issue 1). BioMed Central Ltd. <https://doi.org/10.1186/s12955-023-02221-y>
- Lewandowski, J., Komur, A. A., & Sobańska, D. (2021). Structural and functional relationship of mammalian and nematode ferritins. In *Biotechnologia* (Vol. 102, Issue 4, pp. 457–471). Termedia Publishing House Ltd. <https://doi.org/10.5114/BTA.2021.111110>
- Mahroum, N., Alghory, A., Kiyak, Z., Alwani, A., Seida, R., Alrais, M., & Shoenfeld, Y. (2022). Ferritin – from iron, through inflammation and autoimmunity, to COVID-19. In *Journal of Autoimmunity* (Vol. 126). Academic Press. <https://doi.org/10.1016/j.jaut.2021.102778>
- Mattioli, F., Puntoni, M., Marini, V., Fucile, C., Milano, G., Robbiano, L., Perrotta, S., Pinto, V., Martelli, A., & Forni, G. L. (2015). Determination of deferasirox plasma concentrations: do gender, physical and genetic differences affect chelation efficacy? *European Journal of Haematology*, 94(4), 310–317. <https://doi.org/10.1111/ejh.12419>
- Mazzola, M., Crippa, J., Bertoglio, C. L., Andreani, S., Morini, L., Sfondrini, S., & Ferrari, G. (2021). Postoperative risk of pancreatic fistula after distal

- pancreatectomy with or without spleen preservation. *Tumori*, 107(2), 160–165. <https://doi.org/10.1177/0300891620936744>
- Mobarra, N., Shanaki, M., Ehteram, H., Nasiri, H., Sahmani, M., Saeidi, M., Goudarzi, M., Pourkarim, H., & Azad, M. (2016). A Review on Iron Chelators in Treatment of Iron Overload Syndromes. In *International Journal of Hematology-Oncology and Stem Cell Research 5 Review Article IJHOSCR* (Vol. 10, Issue 4).
- Nasar, I. M., & Cornain, S. (2015). *Buku Ajar Patologi Robbins* (9th ed.). Elsevier.
- Nini, C. (2019). *PENGARUH KADAR FERRITIN SERUM TRANSFUSI DARAH BERULANG TERHADAP GANGGUAN PERTUMBUHAN PADA ANAK THALASSEMIA β MAYOR*.
- Nurbahiyah, E., & Maulina, D. (2023). PROFIL PENGGUNAAN OBAT KELASI BESI PADAPASIEEN TALASEMIA POLI ANAK DI RUMAH SAKIT X JATINEGARA. In *Indonesian Journal of Health Science* (Vol. 3, Issue 2).
- Origa, R. (2017a). β -Thalassemia. In *Genetics in Medicine* (Vol. 19, Issue 6, pp. 609–619). Nature Publishing Group. <https://doi.org/10.1038/gim.2016.173>
- Origa, R. (2017b). β -Thalassemia. *Genetics in Medicine*, 19(6), 609–619. <https://doi.org/10.1038/GIM.2016.173>
- Panigrahi, M., Swain, T., Jena, R., Panigrahi, A., & Debta, N. (2020). Effectiveness of Deferasirox in Pediatric Thalassemia Patients: Experience from a Tertiary Care Hospital of Odisha. *Indian Journal of Pharmacology*, 52(3), 172. https://doi.org/10.4103/ijp.IJP_68_18
- Peng, Y. Y., & Uprichard, J. (2017). Ferritin and iron studies in anaemia and chronic disease. In *Annals of Clinical Biochemistry* (Vol. 54, Issue 1, pp. 43–48). SAGE Publications Ltd. <https://doi.org/10.1177/0004563216675185>
- Piolatto, A., Berchialla, P., Allegra, S., De Francia, S., Ferrero, G. B., Piga, A., & Longo, F. (2021). Pharmacological and clinical evaluation of deferasirox formulations for treatment tailoring. *Scientific Reports*, 11(1), 12581. <https://doi.org/10.1038/s41598-021-91983-w>
- Plays, M., Müller, S., & Rodriguez, R. (2021). Chemistry and biology of ferritin. In *Metallomics* (Vol. 13, Issue 5). Oxford University Press. <https://doi.org/10.1093/mtomcs/mfab021>
- Porter, D., & Taher, J. (2021). *2021 GUIDELINES FOR THE MANAGEMENT OF TRANSFUSION DEPENDENT THALASSAEMIA (TDT) PUBLISHERS THALASSAEMIA INTERNATIONAL FEDERATION 4 TH EDITION*.
- Porter, J. B., & Garbowski, M. (2014). The Pathophysiology of Transfusional Iron Overload. *Hematology/Oncology Clinics of North America*, 28(4), 683–701. <https://doi.org/10.1016/J.HOC.2014.04.003>
- Reddy, P. S., Locke, M., & Badawy, S. M. (2022). A systematic review of adherence to iron chelation therapy among children and adolescents with thalassemia. In *Annals of Medicine* (Vol. 54, Issue 1, pp. 326–342). Taylor and Francis Ltd. <https://doi.org/10.1080/07853890.2022.2028894>
- Rediyanto, D. K. (2023). *Analisis Polimorfisme Gen CYP pada Metabolisme Obat Deteksi Dini Thalassemia* (Vol. 8, Issue 1).
- Roemhild, K., Von Maltzahn, F., Weiskirchen, R., Knüchel, R., Von Stillfried, S., & Lammers, T. (2021). Iron metabolism: Pathophysiology and Pharmacology. *Trends Pharmacol Sci*. <https://doi.org/10.1016/j.tips.2021.05.001>
- Rujito, L. (2019a). *Buku Referensi Talasemia : Genetik Dasar dan Pengelolaan Terkini*.
- Rujito, L. (2019b). *Buku Referensi Talasemia : Genetik Dasar dan Pengelolaan Terkini*.

- Salem, A., Desai, P., & Elgebaly, A. (2023). Efficacy and Safety of Combined Deferiprone and Deferasirox in Iron-Overloaded Patients: A Systematic Review. *Cureus*. <https://doi.org/10.7759/cureus.48276>
- Sanchez-Villalobos, M., Blanquer, M., Moraleda, J. M., Salido, E. J., & Perez-Oliva, A. B. (2022). New Insights Into Pathophysiology of β -Thalassemia. In *Frontiers in Medicine* (Vol. 9). Frontiers Media S.A. <https://doi.org/10.3389/fmed.2022.880752>
- Sawitri, H., & Asmaul Husna, C. (2018). KARAKTERISTIK PASIEN THALASEMIA MAYOR DI BLUD RSU CUT MEUTIA ACEH UTARA TAHUN 2018. In *Jurnal Averrous* (Vol. 4, Issue 2).
- Shafique, F., Ali, S., Almansouri, T., Van Eeden, F., Shafi, N., Khalid, M., Khawaja, S., Andleeb, S., & Ul Hassan, M. (2023). Thalassemia, a human blood disorder. *Brazilian Journal of Biology*, 83. <https://doi.org/10.1590/1519-6984.246062>
- Shah, F. T., Sayani, F., Trompeter, S., Drasar, E., & Piga, A. (2019). Challenges of blood transfusions in β -thalassemia. In *Blood Reviews* (Vol. 37). Churchill Livingstone. <https://doi.org/10.1016/j.blre.2019.100588>
- Sharma, A., Arora, E., & Singh, H. (2015). Hypersensitivity reaction with deferasirox. *Journal of Pharmacology and Pharmacotherapeutics*, 6(2), 105–106. <https://doi.org/10.4103/0976-500X.155491>
- Shukla, S. K., Shrivastava, A., & Mishra, P. C. (2019). Effect of deferasirox on serum ferritin level in children with thalassemia major: impact of transfusional iron load. *International Journal of Contemporary Pediatrics*, 6(5), 2081. <https://doi.org/10.18203/2349-3291.ijcp20193729>
- Sikorska, K., Bernat, A., & Wróblewska, A. (2016). Molecular pathogenesis and clinical consequences of iron overload in liver cirrhosis. *Hepatobiliary & Pancreatic Diseases International*, 15(5), 461–479. [https://doi.org/10.1016/S1499-3872\(16\)60135-2](https://doi.org/10.1016/S1499-3872(16)60135-2)
- Suparyanto, T., Anggraeni, M., & Yolandia, R. A. (2023). HUBUNGAN PEMBERIAN TABLET ZINK, GAYA HIDUP, DAN ASUPAN PROTEIN TERHADAP KADAR FERITIN PADA IBU HAMIL TRIMESTER III DI RUMAH SAKIT KRAKATAU MEDIKA CILEGON TAHUN 2022. *SENTRI: Jurnal Riset Ilmiah*, 2(4).
- Supriatna, C., Indriani, B. K., & Akbari, R. (2020). EVALUASI PENGGUNAAN OBAT KELASI BESI DALAM MENURUNKAN KADAR FERITIN PADA PASIEN THALASEMIA ANAK DI RSUD 45 KUNINGAN. *Jurnal Ilmiah Indonesia*, 5.
- Supriyanti, E. S. I., & Mariana, M. R. (2019). *Faktor-Faktor Yang Berhubungan Dengan Kepatuhan Transfusi Pada Pasien Thalassemia*.
- Susilowati, R. P., Pratanu, L., Ayu, D., & Putri, A. (2024). Case overview of children's thalassemia at RSAB Harapan Kita Jakarta. *Biogenesis: Jurnal Ilmiah Biologi*, 12(1), 74–80. <https://doi.org/10.24252/bio.v12i1.41218>
- Taher, A. T., Origa, R., Perrotta, S., Kouraklis, A., Ruffo, G. B., Kattamis, A., Goh, A. S., Huang, V., Zia, A., Herranz, R. M., & Porter, J. B. (2018). Patient-reported outcomes from a randomized phase II study of the deferasirox film-coated tablet in patients with transfusion-dependent anemias. *Health and Quality of Life Outcomes*, 16(1). <https://doi.org/10.1186/s12955-018-1041-5>
- Taher, A. T., & Saliba, A. N. (2017). | *EMERGING ISSUES IN CLINICAL CARE IN THALASSEMIA* | Iron overload in thalassemia: different organs at different rates.

- Tathe, P. A., Pramod Gosavi, P., & Sanap, G. (2023). *Thalassemia (Beta-Thalassemia)*. www.ijfmr.com
- Thein, S. L. (2018). Molecular basis of β thalassemia and potential therapeutic targets. *Blood Cells, Molecules, and Diseases*, 70, 54–65. <https://doi.org/10.1016/J.BCMD.2017.06.001>
- Tripathy, I., Panja, A., Dolai, T. K., & Mallick, A. K. (2021). Comparative Efficacy and Safety Between Deferiprone and Deferasirox with Special Reference to Serum Ferritin Level and Cardiac Function in Bengali β -Thalassemia Major Children. *Hemoglobin*, 45(5), 296–302. <https://doi.org/10.1080/03630269.2021.1999258>
- Tuo, Y., Li, Y., Li, Y., Ma, J., Yang, X., Wu, S., Jin, J., & He, Z. (2024). Global, regional, and national burden of thalassemia, 1990–2021: a systematic analysis for the global burden of disease study 2021. *EClinicalMedicine*, 72. <https://doi.org/10.1016/j.eclinm.2024.102619>
- Viprakashit, V., & Ekwattanakit, S. (2018). Clinical Classification, Screening and Diagnosis for Thalassemia. In *Hematology/Oncology Clinics of North America* (Vol. 32, Issue 2, pp. 193–211). W.B. Saunders. <https://doi.org/10.1016/j.hoc.2017.11.006>
- Vlachaki, E., Bakır Koyuncu, M., Salido, E. J., Perez-Oliva, A. B., Sanchez-Villalobos, M., Blanquer, M., & Moraleda, J. M. (2022). New Insights Into Pathophysiology of β -Thalassemia. *Frontiers in Medicine | Wwww.Frontiersin.Org*, 1, 880752. <https://doi.org/10.3389/fmed.2022.880752>
- Yampayon, K., Anantachoti, P., Chongmelaxme, B., & Yodsurang, V. (2023). Genetic polymorphisms influencing deferasirox pharmacokinetics, efficacy, and adverse drug reactions: a systematic review and meta-analysis. *Frontiers in Pharmacology*, 14. <https://doi.org/10.3389/fphar.2023.1069854>
- Yiannikourides, A., & Latunde-Dada, G. (2019). A Short Review of Iron Metabolism and Pathophysiology of Iron Disorders. *Medicines*, 6(3), 85. <https://doi.org/10.3390/medicines6030085>
- Yusuf, S., Herdata, H. N., Edward, E. D., & Khairunnisa, K. (2023). Comparison of oral iron chelators in the management of transfusion-dependent β -thalassemia major based on serum ferritin and liver enzymes. *F1000Research*, 12, 154. <https://doi.org/10.12688/f1000research.128810.1>