

Obstetrics and Gynecology III

Editor

Süleyman Cansun DEMİR



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PREFACE

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Chapter 1

DIAGNOSIS AND MANAGEMENT OF HYPERANDROGENISM

Senem ALKAN AKALIN¹

Hyperandrogenism is characterized by excessive androgen synthesis in the ovaries and/or adrenal glands. The most common clinical manifestation of hyperandrogenism in women is hirsutism, excessive terminal hair growth in androgen-dependent body areas.

Other clinical findings of hyperandrogenism;

- Acne vulgaris
- Weight gain
- Menstrual irregularities
- PCOS
- Acanthosis nigricans

The underlying cause of androgen increase can often be determined by a good history, including age of onset, duration and severity of symptoms, and examination of the skin, breasts, pelvis and abdomen. Specific laboratory studies (Serum total/free testosterone, DHEA-S) are requested if no cause is found in the history and FM. Most causes of hyperandrogenism are benign. However, the sudden onset and progressive worsening of symptoms should suggest malignancy.

EVALUATION OF HYPERANDROGENISM

Hirsutism, excessive terminal hair growth in androgen-dependent areas, is the most common finding of hyperandrogenism and is present in approximately 60-80% of patients. (1.Nikolaou D, Gilling-Smith C. Hirsutism. Curr Obstet Gynaecol 2005;15:174-82.)

Evaluation of Hyperandrogenism;

Hirsutism; It should not be confused with hypertrichosis, which occurs with diffuse growth of vellus hair. Hirsutism is seen in 5-10% of the general female

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- Assess for ovarian tumors

Lipid profile

-Lipids elevated in Hyperandrogenism

Source

REFERENCES

1. Nikolaou D, Gilling-Smith C. Hirsutism. *Curr Obstet Gynaecol* 2005;15:174–82
2. Rosenfield RL. Clinical practice. Hirsutism *N Engl J Med* 2005;353:2578–88
3. Hahn S, Kuehnel W, Tan S, et al. Diagnostic value of calculated testosterone indices in the assessment of polycystic ovary syndrome. *Clin Chem Lab Med* 2007;45:202–7.
4. Azziz R. The evaluation and management of hirsutism. *Obstet Gynecol* 2003;101(5 Pt 1):995–1007.
5. Sheehan MT. Polycystic ovarian syndrome: diagnosis and management. *ClinMed Res* 2004;2:13–27.
6. Chang RJ. A practical approach to the diagnosis of polycystic ovary syndrome. *Am J Obstet Gynecol* 2004;191:713–7
7. Barbieri RL, Makris A, Ryan KJ. Insulin stimulates androgen accumulation in incubations of human ovarian stroma and theca. *Obstet Gynecol* 1984;64(3 Suppl):73S–80S
8. Practice Committee of the American Society for Reproductive Medicine. The evaluation and treatment of androgen excess. *Fertil Steril* 2006;86(5 Suppl): S241–7.
9. Moran C. Nonclassic adrenal hyperplasia. *Fertil Steril* 2006;86 Suppl 1:S3
10. Ross EJ, Linch DC. Cushing's syndrome—killing disease: discriminatory value of signs and symptoms aiding early diagnosis. *Lancet* 1982;2:646–9.
11. Kaltsas GA, Korbonits M, Isidori AM, et al. How common are polycystic ovaries and the polycystic ovarian syndrome in women with Cushing's syndrome? *Clin Endocrinol* 2000;53:493–500.)
12. Gilling-Smith C. Hirsutism. *Curr Obstet Gynaecol* 2002;12:144–9.
13. Danilowics K, Albiger N, Vanegas M, et al. Androgen-secreting adrenal adenomas. *Obstet Gynecol* 2002;100(5 Pt 2):1099–102
14. Escobar-Morreale HF. Macroprolactinemia in women presenting with hyperandrogenic symptoms: implications for the management of polycystic ovary syndrome. *Fertil Steril* 2004;82:1697–9.

Chapter 2

GYNECOLOGY AND OBSTETRICS PSYCHIATRY

Ömer Furkan YILMAZ¹

INTRODUCTION

Gynecology and Obstetrics Psychiatry is a sub-specialty of psychiatry that focuses on the mental health of women throughout pregnancy and postpartum period. This field of study is becoming increasingly important as research has shown that up to 20% of women through some form of mental health issue during pregnancy or postpartum (1).

Common conditions in this field include postpartum depression and anxiety, prenatal depression, bipolar disorder, and psychosis. These conditions can have significant consequences for both of the mother and the child, and early identification and intervention is crucial (2).

Obstetricians and gynecologists play an significant role in identifying and managing these conditions, as they are commonly the first point of contact for pregnant women. They can screen for these conditions using standardized questionnaires and refer to an obstetric psychiatrist or other mental health professional as needed (3).

Treatment options for obstetric psychiatric conditions include psychotherapy, medication, and support groups. Antidepressant medication is commonly used, but it is vital to note that these medications can have potential risks and advantage for both of the mother and the baby, and should be attentively considered (4).

It is critical to note that perinatal mental health is a complex and multifaceted field and it is essential that obstetric and gynecological care providers work in collaboration with mental health professionals to provide the finest care possible for women experiencing mental health issues throughout pregnancy and postpartum (5).

In conclusion, Gynecology and Obstetrics Psychiatry is an important field of study that focuses on the mental health of women while pregnancy and postpartum

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CONCLUSION: THE IMPORTANCE OF OBSTETRIC PSYCHIATRY IN WOMEN'S HEALTH CARE

Obstetric psychiatry is a vital field of study that concentrates on the mental well-being of women during pregnancy and postpartum. Research has shown that as much as 20% of women encounter some form of mental health issue during this period, with the most prevalent conditions being postpartum depression, prenatal depression, anxiety, and bipolar disorder (26).

Early identification and intervention is crucial to ensure the finest outcome for both of the mother and the child. Obstetricians and gynecologists play an critical role in identifying and managing these conditions, and a variety of treatment options are available, including psychotherapy, medication, and support groups (26).

It's important to note that perinatal mental health is a complex and multifaceted field and it is essential that obstetric and gynecological care providers work in collaboration with mental health professionals to provide the optimal care possible for women experiencing mental health issues throughout pregnancy and postpartum (5).

The field of Obstetric Psychiatry is also important in addressing the mental health requirements of women, regardless of their life stage, and it is essential to evaluate the potential long-term impacts of psychiatric disorders on the mother's overall well-being and on her future pregnancies (27).

In summary, Obstetric Psychiatry is a significant field of study that concentrates on the mental well-being of women during pregnancy and postpartum. Early identification and intervention are crucial to guarantee the optimal outcome for both of the mother and the child. Obstetricians and gynecologists have an essential role in identifying and treating these conditions, and various treatment options are accessible. It is crucial that healthcare providers collaborate with mental health professionals to provide the optimal care possible for women and to address their mental health requirements in the long-term (27).

REFERENCES

1. Tsakiridis, I., Bousi, V., Dagklis, T. et al. Epidemiology of antenatal depression among women with high-risk pregnancies due to obstetric complications: a scoping review. *Archives of Gynecology and Obstetrics* 300, 849–859 (2019). <https://doi.org/10.1007/s00404-019-05270-1>.
2. Collardeau F, Corbyn B, Abramowitz J, et al. Maternal unwanted and intrusive thoughts of infant-related harm, obsessive-compulsive disorder and depression in the

- perinatal period: study protocol. *BMC Psychiatry*. 2019;19(1):94. Published 2019 Mar 21. doi:10.1186/s12888-019-2067-x.
3. Jin, Y., Bi, Q., Song, G. et al. Psychological coherence, inclusive leadership and implicit absenteeism in obstetrics and gynecology nurses: a multi-site survey. *BMC Psychiatry* 22, 525 (2022). <https://doi.org/10.1186/s12888-022-04137-1>.
4. Garbarino Abigail AB; Kohn Jaden BS; Coverdale John MD, et al. Current Trends in Psychiatry Education Among Obstetrics & Gynecology Residency Programs: A Cross-sectional Survey of Program Directors. *Obstetrics & Gynecology* 132():p 38S, October 2018. | DOI: 10.1097/01.AOG.0000546607.11541.ce.
5. Vannuccini S., Lazzeri L., Orlandini C., et al (2018) Mental health, pain symptoms and systemic comorbidities in women with endometriosis: a cross-sectional study, *Journal of Psychosomatic Obstetrics Gynecology*, 39:4, 315-320, DOI: 10.1080/0167482X.2017.1386171.
6. Yan H, Ding Y, Guo W. Mental Health of Pregnant and Postpartum Women During the Coronavirus Disease 2019 Pandemic: A Systematic Review and Meta-Analysis. *Frontiers in Psychology*. 2020;11:617001. Published 2020 Nov 25. doi:10.3389/fpsyg.2020.617001.
7. Silva BPD, Matijasevich A, Malta MB, et al. Common mental disorders in pregnancy and postnatal depressive symptoms in the MINA-Brazil study: occurrence and associated factors. *Revista de saude publica*. 2022;56:83. Published 2022 Sep 26. doi:10.11606/s1518-8787.2022056004028.
8. Mangla K., Hoffman M. C., Trumpff C., et al (2019), Maternal self-harm deaths: an unrecognized and preventable outcome, *American Journal of Obstetrics and Gynecology* 221 (4), 295-303. <https://doi.org/10.1016/j.ajog.2019.02.056>.
9. Kingston D.E., McDonald S., Austin MP. et al. The Public's views of mental health in pregnant and postpartum women: a population-based study. *BMC Pregnancy Child-birth* 14, 84 (2014). <https://doi.org/10.1186/1471-2393-14-84>.
10. Bener A., Gerber L. M, Sheikh J. (2012) Prevalence of psychiatric disorders and associated risk factors in women during their postpartum period: a major public health problem and global comparison, *International Journal of Women's Health*, 4:, 191-200, DOI: 10.2147/IJWH.S29380.
11. Hamidia, A, Kheirkhah, F, Chehrazi, M, et al. Screening of psychiatric disorders in women with high-risk pregnancy: Accuracy of three psychological tools. *Health Science Reports*. 2022; 5:e518. doi:10.1002/hsr2.518.
12. Cox JL, Chapman G, Murray D, Jones P. Validation of the Edinburgh Postnatal Depression Scale (EPDS) in non-postnatal women. *Journal of affective disorders*. 1996;39(3):185-189. doi:10.1016/0165-0327(96)00008-0.
13. Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity measure. *Journal of general internal medicine*. 2001;16(9):606-613. doi:10.1046/j.1525-1497.2001.016009606.x.
14. Hoffman MC, Wisner KL. Psychiatry and Obstetrics: An Imperative for Collaboration. *The American journal of psychiatry*. 2017;174(3):205-207. doi:10.1176/appi.ajp.2016.16111233.
15. Bharadwaj B, Endumathi R, Parial S, Chandra PS. Management of Psychiatric Disorders during the Perinatal Period. *Indian journal of psychiatry*. 2022;64(Suppl 2):S414-S428. doi:10.4103/indianjpsychiatry.indianjpsychiatry_12_22.

16. Molyneaux E, Poston L, Ashurst-Williams S, Howard LM. Obesity and mental disorders during pregnancy and postpartum: a systematic review and meta-analysis. *Obstetrics and gynecology*. 2014;123(4):857-867. doi:10.1097/AOG.000000000000170.
17. Glasheen C., Colpe L., Hoffman V. et al. Prevalence of Serious Psychological Distress and Mental Health Treatment in a National Sample of Pregnant and Postpartum Women. *Maternal and Child Health Journal* 19, 204–216 (2015). <https://doi.org/10.1007/s10995-014-1511-2>.
18. Byatt N, Cox L, Moore Simas TA, et al. How obstetric settings can help address gaps in psychiatric care for pregnant and postpartum women with bipolar disorder. *Archives of women's mental health*. 2018;21(5):543-551. doi:10.1007/s00737-018-0825-2.
19. Rodriguez-Cabezas L, Clark C. Psychiatric Emergencies in Pregnancy and Postpartum. *Clinical obstetrics and gynecology*. 2018;61(3):615-627. doi:10.1097/GRF.0000000000000377.
20. Bhat A, Reed SD, Ünützer J. The Obstetrician-Gynecologist's Role in Detecting, Preventing, and Treating Depression. *Obstetrics and gynecology*. 2017;129(1):157-163. doi:10.1097/AOG.0000000000001809.
21. Nagle-Yang, S., Sachdeva J., Zhao L.X. et al. Trauma-Informed Care for Obstetric and Gynecologic Settings. *Maternal and Child Health Journal* 26, 2362–2369 (2022). <https://doi.org/10.1007/s10995-022-03518-y>.
22. Dathe K, Schaefer C. The Use of Medication in Pregnancy. *Deutsches Arzteblatt international*. 2019;116(46):783-790. doi:10.3238/arztebl.2019.0783.
23. Lynch MM, Squiers LB, Kosa KM, et al. Making Decisions About Medication Use During Pregnancy: Implications for Communication Strategies. *Maternal and child health journal*. 2018;22(1):92-100. doi:10.1007/s10995-017-2358-0.
24. Smith Erin K. MD; Gopalan Priya MD; Glance Jody B. MD; Azzam, Pierre N. MD. Postpartum Depression Screening: A Review for Psychiatrists. *Harvard Review of Psychiatry* 24(3):p 173-187, May/June 2016. | DOI: 10.1097/HRP.0000000000000103.
25. Meltzer-Brody S., Maegbaek M., Medland S., et al. (2017). Obstetrical, pregnancy and socio-economic predictors for new-onset severe postpartum psychiatric disorders in primiparous women. *Psychological Medicine*, 47(8), 1427-1441. doi:10.1017/S0033291716003020.
26. Ghahremani T, Magann EF, Phillips A, Ray-Griffith SL, Coker JL, Stowe ZN. Women's Mental Health Services and Pregnancy: A Review. *Obstetrical & gynecological survey*. 2022;77(2):122-129. doi:10.1097/OGX.0000000000000994.
27. Raiff EM, D'Antonio KM, Mai C, Monk C. Mental Health in Obstetric Patients and Providers During the COVID-19. *Clinical obstetrics and gynecology*. 2022;65(1):203-215. doi:10.1097/GRF.0000000000000668.

Chapter 3

INTRAHEPATIC CHOLESTASIS OF PREGNANCY

Gülay BALKAŞ¹

Intrahepatic cholestasis of pregnancy (ICP), characterised by pruritus and elevated serum bile acid concentrations, typically develops in the late second and/or third trimester and is associated with increased rates of adverse fetal outcomes. The etiology of cholestasis is poorly understood, and management is difficult due to limited data on diagnosis, treatment, and associated adverse outcomes. It is thought to result from the cholestatic effects of reproductive hormones and their metabolites in genetically susceptible women and resolves rapidly after delivery. The mechanisms by which fetal complications occur are also unclear.

INTRODUCTION

Cholestasis, translated from the Greek, describes the “stoppage of bile”. This disturbance of bile flow can occur at any site in the biliary system and can be caused by extrahepatic mechanical obstruction, pathology of the intrahepatic biliary tree, or dysfunction of individual hepatocytes. Usually, the disorder does not result in a complete interruption of bile flow, and often only certain components of bile become pathologic. The clinical and biochemical manifestations of cholestasis may therefore vary. Conventionally, cholestatic liver diseases are first divided into those with extrahepatic and those with intrahepatic etiology. Cholestasis in pregnancy is an intrahepatic disorder that occurs only during pregnancy and regresses rapidly after delivery (1). ICP was originally described by Ahlfeld in 1883 as recurrent jaundice in pregnancy which resolved after delivery. Pruritus was not mentioned in this report, but subsequent case reports published in the 1950s reported severe pruritus with or without jaundice associated with the condition, in addition to complete resolution after delivery and high recurrence rates in subsequent pregnancies (2). It is the most common pregnancy-specific liver disorder. The etiology of intrahepatic cholestasis in pregnancy (ICP) is poorly understood. It is likely to be a multifactorial disease, with genetic, environmental,

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- With peak bile acids 40 to 99 micromol/L and no other risk factors, the risk of stillbirth is similar to the background risk up to 38 to 39 weeks' gestation. Consider planned delivery at 38 to 39 weeks' gestation.
- For peak bile acids ≥ 100 micromol/L, the risk of stillbirth is higher than the background risk. Consider planned delivery at 35 to 36 weeks' gestation.
- The American College of Obstetricians and Gynecologist (ACOG)/Society for Maternal-Fetal Medicine (SMFM) recommends (40):
- For patients with total bile acid levels < 100 micromol/L, delivery is recommended at 36/7 to 39/7 weeks' gestation or at diagnosis if diagnosed at $> 39/7$ weeks.
- For patients with total bile acid levels ≥ 100 micromol/L, delivery is recommended at 36/7 weeks or at diagnosis if diagnosed later.

There are no special considerations for labour in patients with ICP. Continuous fetal monitoring during labour is indicated because of the increased incidence of fetal death and non-fatal asphyxia. Induction of labour does not necessarily increase the risk of caesarean section compared with expectant management. The risk of postpartum hemorrhage is not increased when ICP is treated with UDCA. Therefore, do not routinely assess coagulation parameters or prescribe vitamin K before delivery. In rare refractory cases, prothrombin time can be checked and vitamin K administered if it is prolonged (41, 42).

REFERENCES

1. Kenyon AP, Piercy CN, Girling J, Williamson C, Tribe RM, Shennan AH. Obstetric cholestasis, outcome with active management: a series of 70 cases. *Journal of Obstetrics and Gynaecology*. 2002; 109(3): 282-8.
2. Svanborg A. A study of recurrent jaundice in pregnancy. *Acta Obstetrica et Gynecologica Scandinavica*. 1954; 33(4): 434-44.
3. Williamson C, Geenes V. Intrahepatic cholestasis of pregnancy. *Obstetrics and Gynecology*. 2014; 124(1): 120-33.
4. Manzotti C, Casazza G, Stimac T, et al. Total serum bile acids or serum bile acid profile, or both, for the diagnosis of intrahepatic cholestasis of pregnancy. *Cochrane Database of Systematic Reviews*. 2019; 7: pCD012546
5. Geenes V, Williamson C. Intrahepatic cholestasis of pregnancy. *World Journal of Gastroenterology*. 2009; 15(17): 2049-66.
6. Gao XX, Ye MY, Liu Y, et al. Prevalence and risk factors of intrahepatic cholestasis of pregnancy in a Chinese population. *Scientific Reports*. 2020; 10(1): 16307.
7. Jacquemin E, De Vree JM, Cresteil D, et al. The wide spectrum of multidrug resistance 3 deficiency: from neonatal cholestasis to cirrhosis of adulthood. *Gastroenterology*. 2001; 120(6): 1448-58.

8. Jacquemin E, Cresteil D, Manouvrier S, et al. Heterozygous non-sense mutation of the MDR3 gene in familial intrahepatic cholestasis of pregnancy. *Lancet*. 1999; 353(9148): 210-1.
9. Bacq Y, Gendrot C, Perrotin F, Lefrou L, Chretien S, Vie-Buret V, et al. ABCB4 gene mutations and single-nucleotide polymorphisms in women with intrahepatic cholestasis of pregnancy. *Journal of Medical Genetics*. 2009; 46(10): 711-5.
10. Keitel V, Vogt C, Haussinger D, et al. Combined mutations of canalicular transporter proteins cause severe intrahepatic cholestasis of pregnancy. *Gastroenterology*. 2006;131(2):624-9.
11. Mutlu MF, Aslan K, Guler I, Mutlu I, Erdem M, Bozkurt N, et al. Two cases of first onset intrahepatic cholestasis of pregnancy associated with moderate ovarian hyperstimulation syndrome after IVF treatment and review of the literature. *Obstetrics and Gynaecology*. 2017; 37(5): 547-9.
12. Abu-Hayyeh S, Ovadia C, Lieu T, et al. Prognostic and mechanistic potential of progesterone sulfates in intrahepatic cholestasis of pregnancy and pruritus gravidarum. *Hepatology*. 2016; 63(4): 1287-98.
13. Floreani A, Gervasi MT. New Insights on Intrahepatic Cholestasis of Pregnancy. *Clinics in Liver Disease*. 2016; 20(1): 177-89.
14. Ropponen A, Sund R, Riikonen S, Ylikorkala O, Aittomäki KJH. Intrahepatic cholestasis of pregnancy as an indicator of liver and biliary diseases: a population-based study. *Hepatology*. 2006; 43(4): 723-8.
15. Piechota J, Jelski W. Intrahepatic Cholestasis in Pregnancy: Review of the Literature. *Clinical Medicine*. 2020; 9(5): 1361.
16. Sahni A, Jogdand SD. Effects of Intrahepatic Cholestasis on the Foetus During Pregnancy. *Cureus*. 2022; 14(10): e30657.
17. Gorelik J, Shevchuk A, De Swiet M, Lab M, Korchev Y, Williamson CJBAlJoO, et al. Comparison of the arrhythmogenic effects of tauro- and glycoconjugates of cholic acid in an in vitro study of rat cardiomyocytes. *Journal of Obstetrics and Gynaecology*. 2004; 111(8): 867-70.
18. Israel EJ, Guzman ML, Campos GA. Maximal response to oxytocin of the isolated myometrium from pregnant patients with intrahepatic cholestasis. *Acta Obstetrica et Gynecologica Scandinavica*. 1986; 65(6): 581-2.
19. Sepulveda WH, Gonzalez C, Cruz MA, et al. Vasoconstrictive effect of bile acids on isolated human placental chorionic veins. *European Journal of Obstetrics & Gynecology and Reproductive Biology*. 1991; 42(3): 211-5.
20. Geenes V, Lövgren-Sandblom A, Benthin L, et al. The reversed feto-maternal bile acid gradient in intrahepatic cholestasis of pregnancy is corrected by ursodeoxycholic acid. *PloS one*. 2014; 9(1): e83828.
21. Ovadia C, Seed PT, Sklavounos A, Geenes V, Di Ilio C, Chambers J, et al. Association of adverse perinatal outcomes of intrahepatic cholestasis of pregnancy with biochemical markers: results of aggregate and individual patient data meta-analyses. *Lancet*. 2019; 393(10174): 899-909.
22. Di Mascio D, Quist-Nelson J, Riegel M, et al. Perinatal death by bile acid levels in intrahepatic cholestasis of pregnancy: a systematic review. *Maternal Fetal Neonatal Medicine*. 2021; 34(21): 3614-22.

23. Wood AM, Livingston EG, Hughes BL, et al. Intrahepatic Cholestasis of Pregnancy: A Review of Diagnosis and Management. *Obstetrical & Gynecological Survey*. 2018; 73(2): 103-9.
24. Geenes V, Chappell LC, Seed PT, et al. Association of severe intrahepatic cholestasis of pregnancy with adverse pregnancy outcomes: a prospective population-based case-control study. *Hepatology*. 2014; 59(4): 1482-91.
25. Williamson C, Miragoli M, Sheikh Abdul Kadir S, et al. Bile acid signaling in fetal tissues: implications for intrahepatic cholestasis of pregnancy. *Digestive Disease*. 2011; 29(1): 58-61.
26. Williamson C, Hems LM, Goulis DG, et al. Clinical outcome in a series of cases of obstetric cholestasis identified via a patient support group. *Obstetrics and Gynaecology*. 2004; 111(7): 676-81.
27. Wensink MJ. The risk of infant and fetal death by each additional week of expectant management in intrahepatic cholestasis of pregnancy by gestational age: various objections. *American Journal of Obstetrics & Gynecology*. 2016; 215(6): 807-8.
28. Saleh MM, Abdo KR. Intrahepatic cholestasis of pregnancy: review of the literature and evaluation of current evidence. *Womens Health (Larchmont)*. 2007; 16(6): 833-41.
29. Manna L, Ovadia C, Lövgren-Sandblom A, et al. Enzymatic quantification of total serum bile acids as a monitoring strategy for women with intrahepatic cholestasis of pregnancy receiving ursodeoxycholic acid treatment: a cohort study. *Obstetrics & Gynaecology*. 2019; 126(13): 1633-40.
30. Zhang Y, Lu L, Victor DW, et al. Ursodeoxycholic acid and S-adenosylmethionine for the treatment of intrahepatic cholestasis of pregnancy: a meta-analysis. *Hepatitis Monthly*. 2016; 16(8): e38558
31. Coltorti M, Bortolini M, Di Padova C. A review of the studies on the clinical use of S-adenosylmethionine (SAMe) for the symptomatic treatment of intrahepatic cholestasis. *Methods and Findings in Experimental Clinical and Pharmacology*. 1990; 12(1): 69-78.
32. Mela M, Mancuso A, Burroughs AK. Review article: pruritus in cholestatic and other liver diseases. *Aliment Pharmacol Ther*. 2003; 17(7): 857-70.
33. Liu J, Murray AM, Mankus EB, et al. Adjuvant Use of Rifampin for Refractory Intrahepatic Cholestasis of Pregnancy. *Obstetrics and Gynecology*. 2018; 132(3): 678-81.
34. Walker KE, Chappell LC, Hague WM, et al. Pharmacological interventions for treating intrahepatic cholestasis of pregnancy. *Cochrane Database System Revision*. 2020; 7(7): CD000493.
35. Cappell MS. Hepatic disorders mildly to moderately affected by pregnancy: medical and obstetric management. *Medical Clinics of North America*. 2008; 92(4): 717-737.
36. Lee RH, Incerpi MH, Miller DA, et al. Sudden fetal death in intrahepatic cholestasis of pregnancy. *Obstetrics and Gynecology*. 2009; 113(2 Pt 2): 528-531.
37. Sentilhes L, Verspyck E, Pia P, et al. Fetal death in a patient with intrahepatic cholestasis of pregnancy. *Obstetrics and Gynecology*. 2006; 107(2 Pt 2): 458-60.
38. Heinonen S, Kirkinen P. Pregnancy outcome with intrahepatic cholestasis. *Obstetrics and Gynecology*. 1999; 94(2): 189-93.
39. Girling J, Knight CL, Chappell L, et al. Intrahepatic cholestasis of pregnancy: Green-top Guideline No. 43 June 2022. *Journal of Obstetrics and Gynaecology*. 2022; 129(13): e95-e114.

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40. Late-Preterm MI, Deliveries E-T. *Obstetrics and Gynecology*. ACOG Committee Opinion, Number 818. 2021;138(1): e35-9.
41. Furrer R, Winter K, Schaffer L, et al. Postpartum Blood Loss in Women Treated for Intrahepatic Cholestasis of Pregnancy. *Obstetrics and Gynecology*. 2016; 128(5): 1048-52.
42. Maldonado M, Alhousseini A, Awadalla M, Idler J, Welch R, Puder K, et al. Intrahepatic Cholestasis of Pregnancy Leading to Severe Vitamin K Deficiency and Coagulopathy. *Case Reports in Obstetrics and Gynecology*. 2017; 2017: 5646247.

Chapter 4

MORPHOLOGY OF THE OVARY

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INTRODUCTION

A crucial element of female reproductive capacity lies in a cyclic nature, which is prominently exemplified by the growth and development of dominant follicles. To gain insight into the menstrual cycle, it becomes imperative to comprehend the cycle of the dominant follicle and the mechanisms that govern it.

FOLLICULOGENESIS

Folliculogenesis commences with the selection of a primordial follicle to join the cohort of developing follicles, culminating in either ovulation or cessation through atresia.

OVULATION

Histologically, ovulation witnesses various tissue changes, especially during the transformation and release of the Graafian follicle. The connections between the granulosa cells weaken and the stigma is formed.

LUTEINIZATION

The corpus luteum, during the luteal phase of the menstrual cycle, functions as a prominent endocrine gland, generating substantial quantities of progesterone and estradiol.

CONCLUSION

Knowledge of the interaction between the morphological structure and function is of great importance in a number of medical and research applications such as reproductive health, infertility treatments and hormonal regulation.

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the corpus luteum functions for approximately eleven to fourteen days, but in cases of luteal phase defects, its activity diminishes in less than nine days. This deficiency is commonly attributed to inadequate progesterone levels, which are responsible for sustaining the corpus luteum. Clinically, luteal phase deficiency can contribute to recurrent early pregnancy loss. In the past, the gold standard for diagnosis involved a luteal phase biopsy; however, due to its recognized imprecision, it is no longer considered clinically relevant. Treatment options may include progesterone supplementation (17).

The visualization of the corpus albicans is a rare occurrence in clinical imaging. However, in post-menopausal women, alterations in the appearance of the corpus albicans may arise as a consequence of hormonal changes related to menopause. Reduced estrogen levels and decreased immune-mediated phagocytic and fibroblastic activity can result in inadequate development and regression of the corpus albicans. Consequently, this can lead to the ultrasound detection of these structures, potentially causing confusion with ovarian neoplasms. The presence of hemosiderin and calcium deposits within the corpus albicans itself can enhance its ultrasound visibility, manifesting as small echogenic foci devoid of distal acoustic shadowing. In addition to ultrasound, these calcifications may also be evident in plain film radiographs of the pelvis (18).

REFERENCES

1. Williams CJ, Erickson GF. Morphology and Physiology of the Ovary. In: Feingold KR, Anawalt B, Blackman MR, et al., eds. *Endotext*. South Dartmouth (MA): MDText.com, Inc.; January 30, 2012.
2. Baker Tg. A Quantitative And Cytological Study Of Germ Cells In Human Ovaries. *Proc R Soc Lond B Biol Sci*. 1963; 158:417-433. doi:10.1098/rspb.1963.0055.
3. MacLennan M, Crichton JH, Playfoot CJ, Adams IR. Oocyte development, meiosis and aneuploidy. *Semin Cell Dev Biol*. 2015; 45:68-76. doi: 10.1016/j.semcdb.2015.10.005
4. Richards JS, Russell DL, Robker RL, Dajee M, Alliston TN. Molecular mechanisms of ovulation and luteinization. *Mol Cell Endocrinol*. 1998;145(1-2):47-54. doi:10.1016/s0303-7207(98)00168-3
5. Johnson J, Canning J, Kaneko T, Pru JK, Tilly JL 2004 Germline stem cells and follicular renewal in the postnatal mammalian ovary. *Nature* 428:145-150
6. Schwartz D, Mayaux MJ 1982 Female fecundity as a function of age: results of artificial insemination in 2193 nulliparous women with azoospermic husbands. *Federation CECOS. N Engl J Med* 306:404-406
7. Oktay K, Briggs D, Gosden RG 1997 Ontogeny of follicle-stimulating hormone receptor gene expression in isolated human ovarian follicles. *J Clin Endocrinol Metab* 82:3748-3751

8. Albertini DE, Combelles CM, Benecchi E, Carabatsos MJ. Cellular basis for paracrine regulation of ovarian follicle development. *Reproduction*. 2001;121(5):647-653. doi:10.1530/rep.0.1210647
9. Makabe S, Naguro T, Stallone T. Oocyte-follicle cell interactions during ovarian follicle development, as seen by high resolution scanning and transmission electron microscopy in humans. *Microsc Res Tech*. 2006;69(6):436-449. doi:10.1002/jemt.20303
10. Simon AM, Goodenough DA, Li E, Paul DL. Female infertility in mice lacking connexin 37. *Nature*. 1997;385(6616):525-529. doi:10.1038/385525a0
11. Erickson GF, Magoffin DA, Dyer CA, Hofeditz C. The ovarian androgen producing cells: a review of structure/function relationships. *Endocr Rev*. 1985;6(3):371-399. doi:10.1210/edrv-6-3-371
12. Richards JS, Pangas SA. The ovary: basic biology and clinical implications. *J Clin Invest*. 2010;120(4):963-972. doi:10.1172/JCI41350
13. Erickson GF, Shimasaki S. The role of the oocyte in folliculogenesis. *Trends Endocrinol Metab*. 2000;11(5):193-198. doi:10.1016/s1043-2760(00)00249-6
14. Eppig JJ. Oocyte control of ovarian follicular development and function in mammals. *Reproduction*. 2001;122(6):829-838. doi:10.1530/rep.0.1220829
15. Russell DL, Robker RL. Molecular mechanisms of ovulation: co-ordination through the cumulus complex. *Hum Reprod Update*. 2007;13(3):289-312. doi:10.1093/humupd/dml062
16. Crisp TM, Dessouky DA, Denys FR. The fine structure of the human corpus luteum of early pregnancy and during the progestational phase of the menstrual cycle. *Am J Anat*. 1970;127(1):37-69. doi:10.1002/aja.1001270105
17. Mesen TB, Young SL. Progesterone and the luteal phase: a requisite to reproduction. *Obstet Gynecol Clin North Am*. 2015;42(1):135-151. doi: 10.1016/j.ogc.2014.10.003
18. Millet J, Much M, Gunabushanam G, Buza N, Schwartz PE, Scoutt LM. Large ovarian calcifications from an unresorbed corpus albicans. *J Ultrasound Med*. 2012;31(9):1465-1468. doi:10.7863/jum.2012.31.9.1465

Chapter 5

OSTEOPOROSIS

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INTRODUCTION

Osteoporosis (OP) is the most common disease affecting the metabolism of bones. The first definitive definition of OP was made in 1820 by a French pathologist named Jean Georges Lobstein as “porous bone”, then in 1947 by Albright as “too little bone in bone”. Osteoporosis is a condition characterized by decreased bone density, resulting in fragile and porous bones. It is defined as a reduction in bone mass per unit volume. (1). The World Health Organization (WHO) defines OP as “a systemic skeletal disease characterized by low bone mass and an increase in bone fragility and fractures as a result of impairment of the microarchitectural structure of bone tissue”. This definition is an international consensus set at the 1990 World Health Organization Osteoporosis Conference (2). With the consensus of WHO in 1994, the definition of OP was revised, so it was adopted that the definition of OP should be made according to the values obtained using DXA (dual x-ray absorptiometry) and the presence of fracture (WHO Study Group 1994). In fact, the purpose of this definition is both to prevent conceptual confusion and to clarify the prevalence of OP over certain numerical values (3).

According to this;

Normal: Bone mineral density (BMD) and bone mineral content below 1 standard deviation (SD) compared to a young adult,

Osteopenia: BMD is between -1 and -2.5 SD compared to young adults,

Osteoporosis: BMD greater than -2.5 SD compared to young adults,

Established OP: BMD greater than -2.5 SD compared to a young adult and the presence of one or more additional fractures.

According to this last definition, fracture is not essential for my definition of osteoporosis. If DEXA is not used, the old definition will apply.

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is recommended in patients with acute vertebral fractures or chronic pain from multiple vertebral fractures. However, patients should be trained to perform isometric exercises when using back braces, as long-term brace application can lead to muscle atrophy and muscle weakness. walking aids; It consists of conventional canes, broad-based supporting canes and walkers. Most hip fractures occur as a result of falls. Therefore, in addition to the above measures, specially designed trochanteric cushions can be used to provide external protection to the hip joint during falls in elderly men and women (39,40). In the prevention of osteoporosis, pain due to fractures, prevention of falls and improving the quality of life, physical therapy agents as well as exercise and physical activity are useful. its importance will gain more importance with increasing age as it is today (39,40,44).

REFERENCES

1. Melikoğlu M. Osteoporoz tanımlaması ve sınıflaması. Türkiye Klinikleri J PM&R-Special Topics 2012;5(3):1-5.
2. Sweet MG, Sweet J, Jeremiah MP, Galazka SS. Diagnosis and treatment of osteoporosis. Am Fam Physician 2009;79(3):193-200.
3. Edwards MH, Jameson K, Denison H, Harvey NC, Sayer AA, Dennison EM, et al. Clinical risk factors, bone density and fall history in the prediction of incident fracture among men and women. Bone 2013;52(2):541-7.
4. Şen N, Tuncer T. Osteoporoz patofizyolojisi. Türkiye Klinikleri J PM&R-Special Topics 2012;5(3):11-6.
5. Bonjour JP, Chevalley T, Ferrari S, Rizzoli R. The importance and relevance of peak bone mass in the prevalence of osteoporosis. Salud Publica Mex 2009;51(Suppl 1):5-17.
6. Wong PK, Christie JJ, Wark JD. The effects of smoking on bone health. Clin Sci (Lond.) 2007;113(5): 233-41.
7. Türkiye İstatistik Kurumu [internet]. Küresel yetişkin tütün araştırması, 2012 [cited 2015 Jul 9]. Available from: <http://www.tuik.gov.tr/PreHaberBultenleri.do?id=13142>.
8. Araslı T. Osteoporoz epidemiyolojisi ve Türkiye verileri. Türkiye Klinikleri J PM&R-Special Topics 2012;5(3):6-10.
9. Tuzun S, Eskiurt N, Akarirmak U, Saridogan M, Senocak M, Johansson H, et al. Incidence of hip fracture and prevalence of osteoporosis in Turkey: the FRACTURK study. Osteoporos Int 2012;23(3):949-55.
10. Uğur M. Osteoporozda risk faktörleri. Türkiye Klinikleri J PM&R-Special Topics, 2012;5(3):17-22.
11. Erdem HR. Osteoporozda tanı yöntemleri. Türkiye Klinikleri J PM&R-Special Topics 2012;5(3):34-42.
12. Civitelli R, Armamento-Villareal R, Napoli N. Bone turnover markers: understanding their value in clinical trials and clinical practice. Osteoporos Int 2009;20(6):843-51.
13. National Osteoporosis Foundation [Internet]. Clinician's guide to prevention and treatment of osteoporosis 2010 [cited 2015 Oct 10]. Available from: <http://nof.org/files/nof/public/content/file/344/upload/159.pdf>

14. Wheeler G, Elshahaly M, Tuck SP, Datta HK, van Laar JM. The clinical utility of bone marker measurements in osteoporosis. *J Transl Med* 2013;(11):201.
15. Tang BM, Eslick GD, Nowson C, Smith C, Bensoussan A. Use of calcium or calcium in combination with vitamin D supplementation to prevent fractures and bone loss in people aged 50 years and older: a meta-analysis. *Lancet* 2007;370(9588):657-66.
16. Yamauchi H, Suzuki H, Orimo H. Calcitonin for the treatment of osteoporosis: dosage and dosing interval in Japan. *J Bone Miner Metab* 2003;21(4):198-204.
17. Akesson K. New approaches to pharmacological treatment of osteoporosis. *Bull World Health Organ* 2003;81(9):657-64.
18. McClung M, Harris ST, Miller PD, Bauer DC, Davison KS, Dian L, et al. Bisphosphonate therapy for osteoporosis: benefits, risks, and drug holiday. *Am J Med* 2013;126:13-20.
19. Roelofs AJ, Thompson K, Gordon S, Rogers MJ. Molecular mechanisms of action of bisphosphonates: current status. *Clin Cancer Res* 2006;12:6222-30.
20. Xu XL, Gou WL, Wang AY, Wang Y, Guo QY, Lu Q, et al. Basic research and clinical applications of bisphosphonates in bone disease: what have we learned over the last 40 years? *J Transl Med* 2013;11:303.
21. National Osteoporosis Foundation. Clinician's Guide to Prevention and Treatment of Osteoporosis. Washington, DC: National Osteoporosis Foundation, 2013.
22. Rizzoli R, Adachi JD, Cooper C, Dere W, Devogelaer JP, Diez-Perez A, et al. Management of glucocorticoid-induced osteoporosis. *Calcif Tissue Int* 2012;91:225-43.
23. Herrera A, Lobo-Escolar A, Mateo J, Gil J, Ibarz E, Gracia L. Male osteoporosis: A review. *World J Orthop* 2012;3:223-34.
24. Lippuner K. The future of osteoporosis treatment - a research update. *Swiss Med Wkly* 2012;142:13624.
25. Diab DL, Watts NB. Denosumab in osteoporosis. *Expert Opin Drug Saf* 2013.
26. Cummings SR, San Martin J, McClung MR, Siris ES, Eastell R, Reid IR, et al. Denosumab for prevention of fractures in postmenopausal women with osteoporosis. *N Engl J Med* 2009;361:756-65.
27. Seeman E, Delmas PD, Hanley DA, Sellmeyer D, Cheung AM, Shane E, et al. Microarchitectural deterioration of cortical and trabecular bone: differing effects of denosumab and alendronate. *J Bone Miner Res* 2010;25:1886-94.
28. Bone HG, Chapurlat R, Brandi ML, Brown JP, Czerwinski E, Krieg MA, et al. The effect of three or six years of denosumab exposure in women with postmenopausal osteoporosis: results from the FREEDOM extension. *J Clin Endocrinol Metab* 2013;98:4483-92.
29. Grey A, Bolland M. Web of industry, advocacy, and academia in the management of osteoporosis. *BMJ* 2015;(351):h3170. doi: 10.1136/bmj.h3170.
30. Eskiuyurt N. Osteoporozdan korunma; genel önlemler (kalsiyum, d vitamini ve fiziksel aktivite). *Türkiye Klinikleri J PM&R-Special Topics* 2012;5(3):98-103.
31. Das S, Crockett JC. Osteoporosis - a current view of pharmacological prevention and treatment. *Drug Des Devel Ther* 2013;7:435-48.
32. Hurtel-Lemaire AS, Mentaverri R, Caudrillier A, Cournarie F, Wattel A, Kamel S, et al. The calcium-sensing receptor is involved in strontium ranelate-induced osteoclast apoptosis. New insights into the associated signaling pathways. *J Biol Chem* 2009;284:575-84.

33. Cianferotti L, D'Asta F, Brandi ML. A review on strontium ranelate long-term antifracture efficacy in the treatment of postmenopausal osteoporosis. *Ther Adv Musculoskelet Dis* 2013;5:127-39.
34. Reginster JY, Kaufman JM, Goemaere S, Devogelaer JP, Benhamou CL, Felsenberg D, et al. Maintenance of antifracture efficacy over 10 years with strontium ranelate in postmenopausal osteoporosis. *Osteoporos Int* 2012;23:1115-22.
35. Rizzoli R, Reginster JY. Adverse drug reactions to osteoporosis treatments. *Expert Rev Clin Pharmacol* 2011;4:593-604.
36. <http://www.ema.europa.eu/ema/index>.
37. Whitaker M, Guo J, Kehoe T, Benson G. Bisphosphonates for osteoporosis--where do we go from here? *N Engl J Med* 2012;366:2048-51.
38. Diab DL, Watts NB. Bisphosphonate drug holiday: who, when and how long. *Ther Adv Musculoskelet Dis* 2013;5:107-11.
39. Sindel D. Osteoporozda rehabilitasyon. *Osteoporoz Kitabı*. İstanbul: Epsilon Matbaası; 2002. s.122-34.
40. Oral A, Küçükdeveci AA, Varela E, Ilieva EM, Valero R, Berteanu M, et al. Osteoporosis. The role of physical and rehabilitation medicine physicians. The European perspective based on the best evidence. A paper by the UEMS-PRM Section Professional Practice Committee. *Eur J Phys Rehabil Med* 2013;49:565-77.
41. Howe TE, Shea B, Dawson LJ, Downie F, Murray A, Ross C, et al. Exercise for preventing and treating osteoporosis in postmenopausal women. *Cochrane Database Syst Rev* 2011;CD000333.
42. Slatkovska L, Alibhai SM, Beyene J, Cheung AM. Effect of wholebody vibration on BMD: a systematic review and meta-analysis. *Osteoporos Int* 2010;21:1969-80.
43. Lau RW, Liao LR, Yu F, Teo T, Chung RC, Pang MY. The effects of whole body vibration therapy on bone mineral density and leg muscle strength in older adults: a systematic review and meta-analysis. *Clin Rehabil* 2011;25:975-88.
44. Sindel D, Dilşen G, Kubat A. Postmenopozal osteoporozda rehabilitasyon açısından yaşam kalitesi sonuçları. *Romatol Tıp Rehab* 1995;6:144-8.

Chapter 6

THE APPROACH TO MALE FACTOR INFERTILITY IN IVF

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INTRODUCTION

A) Definition of male factor infertility

Male factor infertility refers to a condition in which a man's reproductive system has difficulties or abnormalities that contribute to the couple's inability to conceive a child. This can result from various factors such as low sperm count, poor sperm motility (movement), abnormal sperm shape, or other issues affecting sperm production or function. Male factor infertility can be a significant factor in a couple's fertility problems and may require medical evaluation and intervention to address.

B) Prevalence and significance

The prevalence and significance of male factor infertility can vary depending on geographic location, lifestyle factors, and individual health. Here are some general points:

1. **Prevalence:** Male factor infertility is a common issue, accounting for approximately 40-50% of infertility cases. However, it's important to note that male infertility can often be a contributing factor alongside female infertility issues.
2. **Significance:** Male factor infertility is significant because it can impact a couple's ability to conceive naturally. It can lead to emotional and psychological stress for both partners and can strain relationships. Additionally, male infertility may be indicative of underlying health conditions that need attention.
3. **Potential Causes:** Several factors can contribute to male factor infertility, including medical conditions, lifestyle choices (such as smoking or excessive alcohol consumption), exposure to environmental toxins, genetic factors, and hormonal imbalances.

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Hope for Couples Facing Male Factor Infertility:

Couples facing male factor infertility should remain hopeful and proactive. With the advancements in reproductive medicine, personalized treatment plans, and ongoing research, there are numerous options and strategies available to help individuals and couples achieve their dream of parenthood. It's crucial to seek the guidance of fertility specialists who can provide tailored solutions and support throughout the journey. While the path to parenthood may have its challenges, many couples ultimately find success and fulfillment in building their families.

REFERENCES

1. Agarwal, A., & Esteves, S. C. (2016). Sperm Retrieval Techniques in Azoospermic Men: A Systematic Review. *International Journal of Urology*, 23(2), 174-188.
2. Palermo, G. D., & Devroey, P. (2014). What next in the toolbox of assisted reproductive techniques? The search for the best sperm. *Fertility and Sterility*, 101(3), 631-632.
3. Jungwirth, A., et al. (2018). Guidelines on Male Infertility. *European Association of Urology*. Retrieved from <https://uroweb.org/guideline/male-infertility/>
4. Practice Committee of the American Society for Reproductive Medicine. (2018). Diagnostic Evaluation of the Infertile Male: A Committee Opinion. *Fertility and Sterility*, 110(7), 1007-1013.
5. Esteves, S. C., & Agarwal, A. (2013). Novel Concepts in Male Infertility. *International Journal of Urology*, 20(1), 22-35.
6. American Society for Reproductive Medicine (ASRM): The ASRM website (<https://www.asrm.org/>) offers a wealth of information on infertility, including male infertility. They provide educational materials, guidelines, and resources for individuals and couples seeking help with fertility issues.
7. The American Urological Association (AUA): AUA (<https://www.auanet.org/>) provides guidelines and information on male reproductive health, including male infertility, urological treatments, and specialists.
8. Resolve: The National Infertility Association: Resolve (<https://resolve.org/>) is a non-profit organization dedicated to providing support, education, and advocacy for individuals and couples dealing with infertility. They offer a variety of resources, including support groups and educational materials.
9. Mayo Clinic: Mayo Clinic's website (<https://www.mayoclinic.org/>) contains informative articles on male infertility, its causes, diagnosis, and treatment options. It's a trusted source for medical information.
10. National Institutes of Health (NIH): The NIH's MedlinePlus (<https://medlineplus.gov/>) provides comprehensive information on male infertility, including articles, videos, and links to clinical trials.
11. Books: There are many books available on male infertility that provide in-depth information and personal stories. Some recommended titles include "The Male Biological Clock" by Harry Fisch and "Sperm Wars" by Robin Baker.
12. Support Groups: Consider joining online or in-person support groups for individuals and couples dealing with male infertility. Websites like Fertility Network (<https://fertilitynetworkuk.org/>) and Fertility Friends (<https://www.fertilityfriends.co.uk/>)

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provide forums and communities where you can connect with others facing similar challenges.

13. Consulting a Specialist: If you or your partner are experiencing male infertility, it's essential to consult a reproductive specialist or urologist. They can prov