

International Journal of Medical Science and Innovative Research (IJMSIR)

IJMSIR: A Medical Publication Hub Available Online at: www.ijmsir.com

Volume - 7, Issue - 2, April - 2022, Page No.: 441 - 450

Association of vitamin d supplementation with clinical and biochemical parameters in vitamin d deficient pcos patients-A prospective interventional study

¹Suman S, ²Nayan D, ³Singh A, ⁴Banerjee BD, ⁵Grover C

Corresponding Author: Suman S, VCMS and GTB Hospital, New Delhi

Citation this Article: Suman S, Nayan D, Singh A, Banerjee BD, Grover C, "Association of vitamin d supplementation with clinical and biochemical parameters in vitamin d deficient pcos patients-A prospective interventional study", IJMSIR-

April - 2022, Vol - 7, Issue - 2, P. No. 441 - 450.

Type of Publication: Original Research Article

Conflicts of Interest: Nil

Abstract

Aim: To study the effect of vitamin D therapy on various clinical and biochemical parameters in vitamin D deficient PCOS patients.

Material and method: The present study was conducted in Department of Obstetrics and Gynaecology in collaboration with Department of Biochemistry at University College of Medical Sciences (UCMS) and Guru Teg Bahadur Hospital (GTBH), Delhi from November 2017 to April 2019 (including enrollment of subjects, data analysis and thesis writing). 30 vitamin D deficient PCOS patients were recruited. They received vitamin D supplementation for 8 weeks and their clinical and biochemical parameters were compared before and after the therapy.

Result: Significant increase in serum vitamin D levels post supplementation was observed from pre supplementation 9.524±4.41 to post supplementation 32.07±12.40; p-value <0.001. We observed improvement in clinical and biochemical Parameters of the subjects.

Conclusion: Vitamin D supplementation in PCOS patients who are vitamin D deficient, leads to significant fall in angiogenesis and neo-proliferation in ovaries,

which is manifested as improvement in their clinical and biochemical parameters.

Keywords: PCOS, Vitamin D deficiency, Vitamin D supplementation, angiogenesis, polycystic ovaries

Introduction

PCOS is the most common endocrinopathy affecting reproductive age women with prevalence of 6-10% 2. It's a heterogeneous disorder characterised by obesity, insulin resistance, adverse lipid profile and hyper androgenism3. Important reproductive aspects associated with PCOS are polycystic ovaries, anovulation leading to oligomenorrhea or amenorrhea. Metabolic disorders include hyper insulinemia, insulin resistance impaired pancreatic cell insulin secretion, increased risk of type 2 diabetes mellitus and hypertension.4 It's also frequently associated with increased risk of depression, anxiety, endometrial carcinoma5.

Aims and objectives

Aim: To study the effect of vitamin D therapy on various clinical and biochemical parameters in vitamin D deficient PCOS patients.

Objective

- a) Estimation and comparison of serum Vitamin D levels before and after vitamin D supplementation in vitamin D deficient PCOS patients.
- b)To study the effect of vitamin D therapy on various clinical parameters like hirsutism, acne status, BP, and intermenstrual interval.
- c)To study the changes in insulin resistance, hyperandrogenism markers and lipid profile of study subjects after vitamin D supplementation.

Materials and methods

Study setting: The present study was conducted in Department of Obstetrics and Gynaecology in collaboration with Department of Biochemistry at University College of Medical Sciences (UCMS) and Guru Teg Bahadur Hospital (GTBH), Delhi.

Duration: This study was conducted from November 2017 to April 2019 (including enrollment of subjects, data analysis and thesis writing).

Study design: Prospective interventional study. Ethical clearance and consent Clearance from institutional ethical committee was obtained and a written informed consent was taken from all the participants.

Inclusion criteria: All the women between 16-45 years diagnosed as Vitamin D deficient PCOS.

Exclusion criteria

Comparison before and after vitamin d supplementation

Table 1: PCOS related clinical features

Clinical features	Pre Vit-D therapy No. (%)	Post Vit D therapy No. (%)	p-value
Menstrual history			
Infrequent cycles	16(61.53%)	12(46.1%)	
Amenorrhea	7(26.92%)	4(15.3%)	
Regular cycles	3(11.53%)	10(38.4%)	
Hyperandrogenic features			
Acne	7(26.925)	4(15.38%)	

- Women on any hormonal therapy in last 3 months.
- Women on vitamin D supplementation
- Women with known hypersensitivity to vitamin D.
- Patients on any chronic medication like metformin or lipid lowering drugs etc.
- Patients with chronic medical illness like tuberculosis, diabetes mellitus, hypertension etc.

Statistical analysis

Whole data analysis was done by SPSS 20.0 software. At the end of the study data was compiled and outcome parameters were measured as follows:

- 1. Comparison of changes in various clinical and biochemical parameters in study subjects after vitamin D supplementation.
- Comparison of change in the serum levels of vitamin
 in study subjects pre and post vitamin
 supplementation.

Comparison of pre and post vitamin D supplementation of clinical parameters, biochemical parameters, serum vitamin D levels was obtained by Paired t-test. P value of <0.05 was considered significant.

Pillsbury shelly	kligman 0.38±0.68	0.307±.61	0.004
scoring(mean±SD)			
Hirsutism	11(42.30%)	8(30.7%)	
FG score (mean±SD)	9.88±5.218	8.65±3.417	0.001
Insulin resistance			
Acanthosis	6(23.07%)	6(23.07%)	
BMI (kg/m ²)			
18.5-22.9	4(15.38%)	4(15.38%)	
23-24.9	1(3.8%)	9(34.61%)	
≥25	21(80.7%)	13(50%)	
mean±SD	26.884±3.289	25.56±2.838	0.060
Blood pressure (mm/Hg)			
SBP (mean±SD)	122.07±7.494	120.23±5.225	0.97
DBP (mean±SD)	79.096±9.644	77.62±5.216	0.96

Table 2: Effect of vitamin D supplementation on biochemical parameters in vitamin D deficient PCOS patients

Biochemical parameters	Pre vit D mean ±SD	Post vit D mean±SD	p-value
Fasting blood sugar(mg/dl)	81.42±7.66	81.34±7.66	0.45
Postprandial blood sugar (mg/dl)	106.53±12.70	101.73±9.31	0.003
Serum fasting insulin (µIU/ml)	13.97±11.68	12.48±9.81	0.99
QUICKI	0.3455±.045	0.347±.044	0.8224
HOMA-IR	2.91±.5351	2.58±.460	0.070
Serum total testosterone (mIU/ml)	0.77±0.38	0.729±.338	0.001
Serum DHEAS(µg/dl)	309.176±71.8586	302.23±68.87	0.002
Lipid profile			
Total cholesterol (mg/dl)	188.23±41.139	174.96±37.43	1.00
HDL-C (mg/dl)	46.46±8.026	50.54±8.60	0.998
VLDL-C (mg/dl)	23.15±11.540	19.35±12.27	0.70
TG (mg/dl)	139.85±50.454	125.62±43.12	0.010
Dained 4 4est			

Paired t test

Results and discussion

PCOS is the most common endocrinopathy affecting reproductive age women. Increased ovarian mass

supported with new blood vessel proliferation in stroma and theca is a key feature in PCOS pathophysiology.^{6,7} Various angiogenic factors have a role in

neovascularisation in ovarian stroma, chief among them being VEGF.⁸

Studies have suggested role of Vitamin D in pathology of PCOS.⁹ Hypovitaminosis D may induce higher inflammatory response thereby leading to higher angiogenesis. Vitamin D deficiency related increased angiogenesis in PCOS is believed to be mediated by vascular endothelial growth factor (VEGF) dysregulation.¹ It has been shown that vitamin D supplementation decreases VEGF production by human lumbar annulus cells and cancer cells.¹

Most of the information available in literature are from foreign studies. Taking all of these in consideration, we have hypothesized that vitamin D supplementation will lead to decrease in inflammation in ovaries, consequently decrease in ovarian neo angiogenesis and proliferation which would be manifested by improvement in clinical and biochemical parameters.

In the present study, 80 women with clinical diagnosis of PCOS like acne, obesity, oligomenorrhea, hirsutism etc. were screened from November 2017 to October 2018. Out of them 40 women who satisfied selection criteria were subjected to serum vitamin D levels estimation. 30 women out of these 40 were found to be Vitamin D deficient and were enrolled in the study.

Detailed history taking, general examination was done and blood sample was taken for biochemical parameter assessment.

All women received vitamin D supplementation in form of cholecalciferol granules in sachet of 60,000 IU for 8 weeks followed by maintenance therapy. Post vitamin D supplementation, blood sample was collected for biochemical parameter assessment.

Out of 30 participants, 1 woman did not complete the study, 1 spontaneously conceived, 1 took alternative

treatment like metformin during the study and 1 did not come for follow up.

Total 26 women completed the study who were evaluated after 8 weeks for biochemical and at fifth month for clinical parameters.

Serum vitamin D levels along with various clinical and biochemical parameters were compared before and after vitamin D supplementation.

30 out of 40 women i.e., 75% were found to be vitamin D deficient. 26 out of 30 subjects completed the present study. Out of 26 subjects, 15 (57.69%) had severe VDD while 11 (42.30%) had mild VDD. This favours that vitamin D deficiency is 67-85% prevalent in PCOS women as observed in a study by Keshavarz et al¹⁰ conducted in 2013 in Iran, wherein out of 73 recruited women 64 (79%) of them had serum vitamin D level<20 ng/ml. A study done on Indian population by Garg et al¹¹ in 2015, hypovitaminosis D was observed in 93.8% of all subjects with mean serum vitamin D level of 7.30±4.45 ng/ml. Among 32 subjects who completed the study 13(40.3%) had severe vitamin D deficiency, while 16(50%) had mild vitamin D deficiency.

Pre supplementation mean BMI in current study was 26.884 kg/m^2 , (table 1) which lies in overweight category, correlating with the fact that obesity is one of the clinical features in PCOS. Similar results were also reported by Irani et al¹ in a 2017 study done on 53 subjects where the mean BMI of study population was $28\pm1.6 \text{ kg/m}^2$.

Pre vitamin D supplementation, 16 subjects had irregular cycles, while post supplementation 12 subjects had irregular cycles which showed 15% improvement in cyclicity of menses. (Table 1)

Similarly we observed an improvement in cycle regularisation in 10 subjects post supplementation in

comparison to 3 subjects pre supplementation which is 27% improvement. (Table 1)

Improvement in intermenstrual intervals after vitamin D supplementation was also observed by Irani et al 1 in their study which was a randomized placebo controlled trial on 53 subjects (80 \pm 9 to 60 \pm 6 days; p=0.04) .Similar results were observed by Tehrani et al 12 improvement in menstrual cycles in 13 out of 65 in metformin + calcium +vitamin D supplementation group and improvement in 4 subjects out of 20 in calcium+vitamin D supplementation group

We observed a significant improvement in acne status of subjects. Mean Pillsbury shelly kligman scoring change was from. 0.38 ± 0.68 to 0.307 ± 0.61 supplementation. This difference in mean was statistically significant, p value 0.004.(Table 1). Tehrani et al12 conducted a study on 80 PCOS women improvement in acne was seen following vitamin D supplementation 50% improvement, in metformin + calcium +vitamin D supplementation group and 25% improvement in calcium+vitamin D supplementation group.

Post vitamin D therapy, significant improvement in hirsutism was observed. Mean FG score pre supplementation 9.88±5.218 to post supplementation 8.65±3.417, with p-value 0.001 which is also statistically significant (Table 1) our study results were consistent with the study conducted by Irani et al¹ in 2017, following vitamin D supplementation, significant decrease in Ferriman Gallwey hirsutism score (9.8±1.5 to 8.1±1.5 p<0.001) was seen. Improvement in hirsutism post vitamin D supplementation was also observed by Tehrani et al^{12.}

In present study, mean BMI post vitamin D supplementation was 25.56±2.838 which was lower than

pre vitamin D mean of 26.884±3.289 and p-value was 0.01. But it was not statistically significant. These results are similar to that observed in study by Garg et al¹¹ where mean BMI changed from 26.8±4.57 to 25.4±5.65 with p-value 0.1, post vitamin D supplementation.

Post vitamin D supplementation mean Systolic BP and mean Diastolic BP values were lower, however these changes are not significant in our study. This is consistent with results obtained by Raja khan et al¹³, change in mean Systolic BP (117.46 ± 10.00 to 118.53 ± 6.97) and mean Diastolic BP change was (79.08 ± 8.28 to 78.97 ± 5.27) which was not significant, however vitamin D supplementation was seen to have protective effect on diastolic blood pressure, similar to our study. In a study by Pal L berry ¹⁴ et al, significant lowering in BP parameters was seen in participants with baseline BP \geq 120/80 mmHg (n = 8) and in those with baseline serum 250HD \leq 20 ng/ml (n = 9).

The change in mean FBS post vitamin D supplementation was from 81.42±7.66 to 81.34±7.66 mg/dl; p-value =0.45 which was not statistically significant. We observed a significant fall in PPBS post supplementation, 106.53±12.70 to 101.73±9.31 mg/dl; pvalue 0.003 which was statistically significant. (Table 2) We observed a fall in parameters of insulin resistance. Fasting serum insulin mean pre supplementation 13.97±11.68 to 12.48±9.81 μIU/ml post supplementation with p-value=0.99. Mean QUICKI pre supplementation was 0.3455±0.045 to 0.347±0.44 post supplementation p value being 0.822 and mean HOMA IR values pre vitamin D supplementation 2.91±0.535 to 2.58±0.460 post supplementation with p-value =0.07, but these changes were not statistically significant which may be due to small sample size. (Table 2)

Considering HOMA IR > 2.5 as measure of insulin resistance, the post vitamin D HOMA IR mean was 2.58 which is although an improvement from pre vitamin D HOMA IR of 2.91 but it is not statistically significant.

Similar results were obtained by Garg et al¹¹, vitamin D supplementation did not have any significant effect on glycemic picture of the study subjects. Mean fasting plasma glucose change from pre to post vitamin D supplementation (87.9 \pm 7.22 to 90 \pm 7 mg/dl; p=0.43), mean fasting insulin (17.3 \pm 15.28 to 10.3 \pm 5.92; p<0.01), mean HOMA IR (3.8 \pm 3.40 to 2.3 \pm 1.32; p=0.03).

Similar to this in study by Raja khan¹³ et al using a high dose of vitamin-D for 12 weeks, there was no significant differences in QUICKI and other measures of insulin sensitivity but trends towards lower two-hour insulin and lower 2- hr glucose was observed which is consistent with results of present study. In this study, mean fasting plasma glucose change from pre to post vitamin D supplementation was (84.92±9.46 to 83.82±8.02 mg/dl; p=0.7), post prandial plasma glucose change was (122.08±36.29 to 110.73±24.84 mg/dl; p=11.66) mean fasting insulin (26.31±9.60 to 38.09±37.60 μIU/ml; p=13.04), mean HOMA IR (5.47±1.82 to 7.79±7.37; p=2.57), mean QUICKI (0.302±0.014 to 0.296±0.022; p=0.008)

Contrary to our results, Pal L Berry¹⁴ et al observed parameters of glucose homeostasis and IR remained unchanged (p > 0.05).

A RCT carried out by Bonakdaran et al¹⁵ among 51 PCOS patients, did report a decrease in HOMA IR after 3 months of vitamin D supplementation, but this decrease was not significant due to small sample size

In the only study, by Seligmolu et al¹⁶ where vitamin D supplementation resulted in decrease in IR, the results were of borderline significance. In this study they gave

single oral dose of 300,000IU of vitamin D to 11 PCOS women 3 weeks after administration HOMA IR decreased significantly (p=0.043). However, it would be difficult to compare results of our study with these studies as the dose and duration of vitamin D supplementation is quite variable among the different studies.

We observed an improvement in lipid profile of subjects post vitamin D therapy. Changes in mean values pre to post vitamin D supplementation were, of total cholesterol from 188.23±41.139 to 174.96±37.43 mg/dl; p-value=1.00, of mean HDL-C from 46.46±8.026 to 50.54±8.60 mg/dl; p-value 0.998 of mean VLDL-C from 23.15±11.54 to 19.35±12.27 mg/dl; p-value 0.70 but they were not statistically significant. A significant fall was observed in serum triglycerides level from pre vitamin D supplementation 139.85±50.454 to 125.62±43.12 mg/dl post supplementation with p value-0.01 (Table 2)

Similar observations were made by Irani et al¹. In the study conducted by Garg et al¹¹, mean value of total cholesterol (172±31 to 158±20 mg/dl; p-value=0.08), mean HDL-C from 42±3 to 43±6 mg/dl; p-value 0.36 mean serum triglyceride from 116±102 to 81±42 mg/dl; p-value< 0.01 which although was an improvement but they were not statistically significant.

In study by Raja Khan¹³ et al, where high dose vitamin D supplementation was given, the results were, mean value changes from pre supplementation to post supplementation, of total cholesterol from 172±42.70 to 177.18±37.17 mg/dl; p-value=1.69), of mean HDL-C from 45.54±17.60 to 45.73±18.40 mg/dl ;p-value 0.7 of mean serum triglyceride from 139.08±71.61 to 127.73±58.09 mg/dl; p-value< 2.21)which is an overall improvement in lipid profile same as our current study.

In the present study, biochemical parameters of hyperandrogenism like serum total testosterone and serum DHEAS were reduced post vitamin D therapy with mean values of 0.72 ± 0.338 and 302.23 ± 68.875 respectively. Out of 26 subjects 4 had serum total testosterone ≥ 9 mIU/ml which is 50% reduction than pre supplementation value of 8 subjects, similarly serum DHEAS of 9 subjects were>333 µg/dl instead of 15 subjects in the pre vitamin D therapy group. This change in serum testosterone and serum DHEAS values post vitamin D supplementation was statistically significant (p value 0.001 and 0.002 respectively). (Table 2)

This is similar to the, study by Pal L Berry et al¹⁴ in 2012, which demonstrated positive effects of vitamin D and calcium supplementation on metabolic and hormonal milieu in PCOS patients. It showed improved serum 25OHD (p<0.001) and reductions in total testosterone (p

= 0.036) and androstenedione (p=0.090) levels were noted following 3-month supplementation, compared to baseline. In the study conducted by Garg et al¹¹ although there was no statistically significant difference in serum testosterone after vitamin D supple mentation, but a significant fall in serum testosterone was observed from pre supple mentation 0.47±0.19 to post supple mentation 0.35±0.12 ng/ml with p-value=0.03, similarly fall in serum DHEAS was also observed from pre supple mentation 236.5±85.57 to post supple mentation238.3±126 with p-value=0.93. A direct effect of vitamin D on steroidogenesis pathway has been proposed to explain the observed fall in circulating androgens.

Table 3: Comparison of vitamin D supple mentation effect on serum vitamin D levels pre and post vitamin D therapy

	Serum vitamin D level Pre therapy	Serum vitamin D level Post
	(ng/ml)	therapy (ng/ml)
mean ± SD of study population (n=26)	9.524±4.41	32.07±12.40
p value of study population (n=26)	<0.001	
Severe vit D def (mean±SD)	6.233±1.957	24.158±5.17
Range	0.200_000	
(n=15)	(1.362 - 9.413)	(14.431 - 31.68)
Mild vit D def (mean±SD)	14.01±2.23	42.86±11.20
Range		
(n=11)	(10.25 - 17.094)	(32.78 - 66.85)

Paired t-test

We observed a significant rise in serum vitamin D levels in study subjects after vitamin D supplementation. The mean serum vitamin D values raised from 9.524±4.41 pre supplementation to 32.07±12.40 ng/ml post supplementation, the p-value being <0.001 which was statistically significant. (Table 3; Fig 1).

Irani et al¹ also reported significant change in serum vitamin D levels post supplementation. They observed pre supplementation mean was 16.3±0.9 to post supplementation mean 43.2±2.4 ng/ml with p value <0.01.

Similar results were observed in study by Raja khan et al¹³ post supplementation $(19.95\pm9.47 \text{ to } 67.36\pm28.62)$

ng/ml; p<0.001) and Garg et al¹¹ from pre vitamin D therapy 7.7±6.05 to post vitamin D serum vitamin D mean 31.5±13.88 ng/ml; with p<0.001, where there was significant rise in serum vitamin D post supplementation After supplementation 23 out of 26 subjects achieved normal serum vitamin D levels and 3 subjects in severe vitamin D deficient category, remained mild vitamin D deficient who were kept on maintenance therapy.

We did not observe any symptoms or signs of vitamin D toxicity like nausea, gastrointestinal disturbances or skin manifestations in any of our subjects after vitamin D supplementation.

Conclusion

PCOS is a heterogeneous disorder characterised by obesity, insulin resistance, adverse lipid profile, hyperandrogenism. Important reproductive aspect associated with PCOS are polycystic ovaries, anovulation leading to oligo or amenorrhea. One of the important aspects of PCOS pathophysiology is inflammation and neo angiogenesis in the PCOS ovaries. Vitamin D is anti-inflammatory. It reduces angiogenesis and excessive proliferation in ovaries of PCOS women.

- Vitamin D supplementation improves clinical parameters in PCOS women manifested by, improvement in acne, hirsutism, improvement in menstrual cycle regularity. Trends towards lower BMI, lower SBP and lower DBP was observed
- We observed improvement in biochemical parameters like significant fall in PPBS, serum testosterone and serum DHEAS level.
- Improvement in parameters of insulin resistance like HOMA IR, QUICKI and serum fasting insulin post vitamin D supplementation. Improved lipid profile after vitamin D therapy. However the values were not significant due to small sample size.

- Improvement in lipid profile post vitamin D supplementation, with a significant reduction in hypertriglyceridemia was observed.
- Significant increase in serum vitamin D levels post supplementation was observed from pre supplementation 9.524±4.41 to post supplementation 32.07±12.40; p-value <0.001.

So, we propose that vitamin D supplementation is beneficial in PCOS women as it decreases angiogenesis and inflammation in ovaries as evident by improvement in clinical and biochemical picture.

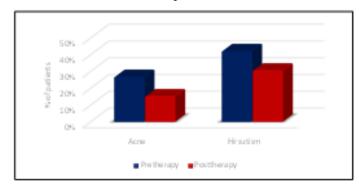


Fig. 1 Distribution of acre and hirsutism in study population.

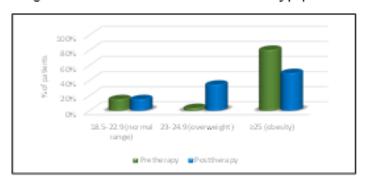


Fig. 2: Distribution of BMI in study population

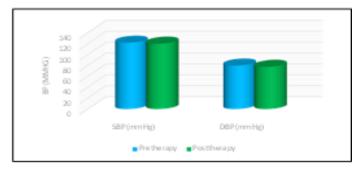


Fig. 3: Distribution of SBP and DBP in study population

References

- 1. Irani M, Seifer DB, Grazi RV, Irani S, Rosenwaks Z, Tal R. Vitamin D Decreases Serum VEGF Correlating with Clinical Improvement in Vitamin D-Deficient Women with PCOS: A Randomized Placebo-Controlled Trial. Nutrients. 2017;28;9(4).
- 2. Bozdag G, Mumusoglu S, Zengin D, Karabulut E, Yildiz BO. Atlas of Science. The worldwide prevalence and phenotypic features of polycystic ovary syndrome. Available from: https://atlasofscience.org/the-worldwide-prevalence-and-phenotypic-features-of-polycystic-ovary-syndrome/. Accessed on 10th April 2019.
- 3. Setji TL, Brown AJ. Polycystic ovary syndrome: update on diagnosis and treatment. Am J Med. 2014;127(10):912-9.
- 4. Hull MJR. Epidemiology of infertility and polycystic ovarian disease: endocrinological and demographic studies. Gynaecol Endocrinol 1987; 1: 233-245.
- 5. Rotstein A. Polycystic ovarian syndrome (PCOS). Available from: http://www.pathophys.org/pcos/.
- 6. Zaidi J, Campbell S, Pittrof R, Kyei-Mensah A, Shaker A, Jacobs HS, Tan SL. Ovarian stromal blood flow in women with polycystic ovaries--a possible new marker for diagnosis? Hum Reprod. 1995;10(8):1992-6.
- 7. Pan HA, Wu MH, Cheng YC, Li CH, Chang FM. Quantification of Doppler signal in polycystic ovary syndrome using three-dimensional power Doppler ultrasonography: a possible new marker for diagnosis. Hum Reprod. 2002;17(1):201-6.
- 8. Peitsidis P, Agrawal R. Role of vascular endothelial growth factor in women with PCO and PCOS: a systematic review. Reprod Biomed Online. 2010 Apr;20(4):444-52.

- 9. Thomson RL, Spedding S, Buckley JD. Vitamin D in the aetiology and management of polycystic ovary syndrome. Clin Endocrinol (Oxf). 2012 Sep;77(3):343-50 10. Keshavarz MA, Moradi S, Emami Z, Rohani F. Association between serum 25(OH) vitamin D and metabolic disturbances in polycystic ovary syndrome. Neth J Med. 2017;75(5):190-195.
- 11. Garg G, Kachhawa G, Ramot R, Khadgawat R, Tandon N, Sreevivas V, et al. Effect of vitamin D supplementation on insulin kinetics and cardiovascular risk factors in polycystic ovarian syndrome: a pilot study. Endocrine Conn 2015;4:108-16.
- 12. Tehrani HG, Mostajeran F, Shahsavari S. The effect of calcium and vitamin D supplementation on menstrual cycle, body mass index and hyperandrogenism state of women with poly cystic ovarian syndrome. J Res Med Sci. 2014;19(9):875-80.
- 13. High-dose Vitamin D Supplementation and Measures of Insulin Sensitivity in Polycystic Ovary Syndrome: a Randomized Controlled Pilot Trial Nazia Raja-Khan, Julie Shah, Christy M. Stetter, Mary E.J. Lott, Allen R. Kunselman, William C. Dodson, Richard S. Legro Fertil Steril. Author manuscript; available in PMC 2015 Aug 14.Published in final edited form as: Fertil Steril. 2014 Jun; 101(6): 1740–1746. Published online 2014 Mar 14. doi: 10.1016 /j. fertnstert. 2014. 02.021 PMCID: PMC 4537163
- 14. Pal L Berry, A Coraluzzi ,L Kustan,E Danton, C Shaw, J Taylor H. Therapeutic implications of vitamin D and calcium in overweight women with polycystic ovary syndrome. Gynecological Endocrinology201228965–968. (doi:10.3109/09513590.2012.696753).
- 15. Bonakdaran S, Khorasani ZM, Davachi B, Khorasani JM. The effects of calcitriol on improvement of insulin resistance, ovulation and comparison with

metformin therapy in PCOS patients: a randomized placebo-controlled clinical trial. Iranian Journal of ReproductiveMedicine201210465–472.

- 16. Selimoglu H, Duran C, Kiyici S, Ersoy C, Guclu M, Ozkaya G, et al. The effect of vitamin D replacement therapy on insulin resistance and androgen levels in women with polycystic ovary syndrome. J Endocrinol Invest. 2010; 33:234–8
- 17. Irani M, Seifer DB, Grazi RV, Julka N, Bhatt D, Kalgi B,Irani S, Tal O, Lambert-Messerlian G, Tal R. Vitamin D Supplementation Decreases TGF- β 1 Bioavailability in PCOS: A Randomized Placebo-Controlled Trial. J Clin Endocrinol Me tab. 2015 Nov;100(11):4307-14