Polycystic ovarian syndrome: pathogenesis and management

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Received: 08 December 2020
Accepted: 13 January 2021

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ABSTRACT
Polycystic ovarian syndrome (PCOS) is an endocrine disorder that commonly affects women of reproductive age. Its prevalence varies quite significantly and its clinical presentation includes: reproductive, metabolic and psychological issues. PCOS develops due to a variety of factors some of which are not yet fully understood. Due to the complexity in the development and presentation of this condition, the management of PCOS in most instances is complicated and involves the combination of pharmacological, non-pharmacological and surgical interventions. This review discusses the pathogenesis and management of PCOS as means of providing information that would enhance understanding and control of this disease.

Keywords: Hyperandrogenemia, Insulin resistance, Polycystic ovarian syndrome, Type-2 diabetes mellitus

INTRODUCTION
PCOS is considered one of the most common endocrine disorders affecting women of reproductive age.1,2 Although the aetiology of this condition is unknown, the prevalence of PCOS has been proven to vary quite widely ranging from as low as 2.2% to as high as 26%.3 The clinical presentations of this disease are quite variable and include: reproductive problems such as infertility, metabolic manifestations such as type II diabetes mellitus (T2DM) and psychological issues such as depression.4

Research into the pathophysiology of this condition has led to significant discoveries regarding PCOS, some of which have brought to light the importance of insulin resistance and obesity as part of the presentation of the syndrome.5,6 This knowledge is important as it directs clinicians to not only focus on managing the common manifestations of PCOS, such as infertility and hyperandrogenemia, but to also take into consideration the metabolic abnormalities that may present as part of this disease. As it relates to the pathogenesis of PCOS, different theories have been proposed for its development. PCOS is considered multifactorial and evidence suggests that it potentially develops due to various factors which may be genetic or environmental in nature.7,8

Despite the wealth of knowledge already acquired on this condition, much research is still required into its pathogenesis and aetiology. In addition, with the general effects that PCOS has on the body (reproductive, metabolic and psychological) physicians may often struggle to successfully control the clinical manifestations and associated features of the syndrome.

Therefore, this review serves to further investigate the pathogenesis and management of PCOS as means of providing clinicians with the necessary information needed to not only understand this disease but to also effectively manage and control its presentation.
**TYPES OF PCOS**

Takai et al described PCOS based on androgenic function. Their study identified three types of PCOS.⁹

*Type 1:* Characterized by an absence of features of hirsutism and no increase in androgen levels (androstenedione and or testosterone)

*Type 2:* Characterized by an absence of features of hirsutism with elevated levels of androgens.

*Type 3:* Characterized by both features of hirsutism and elevated blood levels of androgens.

**DIAGNOSTIC CRITERIA FOR PCOS**

A series of criteria has been developed for use in the diagnosis of PCOS. In 1990, at a conference on PCOS sponsored by The National Institutes of Health/National Institute of Child Health and Human Development (NIH/NICHD), the following criteria, with features presented in order of importance, were developed for the diagnosis of PCOS: a) high levels of androgens manifested clinically or biochemically, b) prolonged anovulation and c) the elimination of other similar conditions such as thyroid diseases or hyperprolactinemia.¹⁰,¹¹

In 2003, another conference, which was sponsored by two bodies, the American Society for Reproductive Medicine and the European Society for Human reproduction and Embryology, was held in Rotterdam, Netherlands. During this conference, the following three features were identified as part of the diagnostic criteria for PCOS: a) oligoovulation or anovulation, b) increased levels of androgens manifested clinically or biochemically and c) polycystic ovaries (PCOs) detected upon ultrasound examination. After the successful exclusion of other similar disorders, the Rotterdam criteria require the presence of at least two out of these three features for the accurate diagnosis of PCOS.¹² The Rotterdam criteria added two diagnostic features to the NIH criteria: a) the presence of PCOs with ovulatory problems and b) the presence of PCOs with ovulatory problems in the absence of an increased production of androgens.¹³

In 2006 further reviews led to the amendment of the diagnostic criteria for PCOS by a board of experts from The Androgen Excess Society (AES). The AES recommended that the presence of excess androgens be given great attention as a critical feature for diagnosing PCOS. According to the AES criteria, in order for a diagnosis of PCOS to be given, the following features are required: a) hyperandrogenism detected clinically or biochemically, combined with b) irregular ovulation and/or the presence of PCOs.¹⁴

Table 1: Summary of diagnostic criteria for polycystic ovarian syndrome.

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Year of development</th>
<th>Features</th>
</tr>
</thead>
<tbody>
<tr>
<td>NIH/NICHD</td>
<td>1990</td>
<td>Excess androgen production and chronic anovulation</td>
</tr>
<tr>
<td>Rotterdam</td>
<td>2003</td>
<td>Excess androgen production, chronic anovulation and Polycystic ovaries on ultrasound examination NB: Requires the presence of at least two of these features for diagnosis</td>
</tr>
<tr>
<td>AES</td>
<td>2006</td>
<td>Excess androgen production, irregular ovulation and/or polycystic ovaries on ultrasound examination</td>
</tr>
</tbody>
</table>

NIH/NICHD: The National Institutes of Health/ National Institute of Child Health and Human Development, AES: The Androgen Excess Society

**Pathogenesis**

PCOS is considered a multifactorial condition which may develop due to different genetic or environment causes. The following identifies hormonal, neuroendocrine, genetic and developmental factors that may contribute to the development of this syndrome.

**Hormonal factors**

The role of Anti-Mullerian hormone (AMH) in the development of PCOS has been investigated extensively. This hormone is normally produced in the male foetus where it supports the degeneration of the Mullerian duct. In females, it is produced by ovarian granulosa cells.¹⁵

In the ovaries, AMH renders the developing follicles less sensitive to follicle stimulating hormone (FSH). It also interferes with the normal functioning of the ovaries and the enzyme aromatase; therefore, inhibiting the initiation of the primordial follicle. In a study carried out by Durlinger et al to investigate the effects of AMH on the recruitment of primordial follicles, ovaries from 2 day old mice were harvested and cultured. After culturing these ovaries for 2 to 4 days, it was observed that there was a decrease in the number of growing follicles in ovaries that were exposed to AMH as compared to the control.¹⁶ Therefore, this demonstrated that AMH may have the ability to directly affect developing primordial follicles.

Another study was carried out by Durlinger et al to examine the effects that AMH has on the ovaries. In this study the entire ovarian population of three categories of mice were examined: 1) AMH (+/+)(wild-type) 2) AMH
Neuroendocrine factors

Studies into the relationship between AMH and the development of PCOS show that PCOS patients exhibit a significantly higher level of AMH as compared to control groups. An in-vivo study was carried out to demonstrate the effects of AMH on LH secretion through recruitment of the hypothalamic neurons that release gonadotropin releasing hormone (GnRh). It was demonstrated that AMH increases LH levels by binding to AMH receptors found on GnRh neurons in the hypothalamus. In patients with PCOS, AMH inhibits FSH release while it induces an increase in LH secretion. LH levels are therefore significantly increased with a typical increase in the LH: FSH ratio, a finding which is a classical feature of PCOS.

Genetic factors

Research evidence suggests a genetic arm to the development of PCOS. Twin studies demonstrate that this condition is highly hereditable with a stronger correlation of the condition being observed amongst monozygotic twins as compared to dizygotic twins. Furthermore, studies have also attempted to assign autosomal dominant and x-linked modes of inheritance for this condition; however, due to limitations in study designs, selection methods and the categorization of the features of the syndrome, results obtained in this regard are majorly inconclusive.

A procedure called the transmission disequilibrium test (TDT) has been applied with success in identifying potential genetic defects that may be associated with certain features of PCOS. TDT is used to determine and link the transmission of susceptible genes from parents to offsprings. With the use of this test, a defect linked to the reproductive phenotype (hyperandrogenemia) observed in PCOS patients was identified on Chromosome 19p 13.2 (dinucleotide repeat D19S884). A case-control study which investigated 85 Caucasian females with PCOS showed a relationship between the D19S884 genetic defect and PCOS. However, two larger case control studies, which investigated this association in Caucasian females of European decent, both failed to further demonstrate an association between this identified genetic defect and PCOS.

The fibrillin-3 gene has also been identified for the role that it may play in the pathogenesis of PCOS. Observation of both polycystic and non-polycystic ovaries shows that PCO demonstrate a marked reduction in their expression of fibrillin-3 in stromal cells immediately surrounding follicles maturing from primordial to primary follicles; a phase of follicular development which is usually affected in PCOS.

Developmental factors

Fetal programming is another factor that has been
postulated as a major contributor to the pathogenesis of PCOS. However, further research is required to fully understand this effect in humans as research has yielded inconclusive or opposing results in this regard.

**Prenatal exposure to excess androgens**

Studies on rhesus monkeys demonstrated that female off-springs, exposed to excess androgens prenatally, exhibit reproductive and metabolic features of PCOS during adolescent periods of their development. In humans, it is theorized that females exposed to elevated levels of androgens at any point starting from the beginning of fetal ovarian development to the commencement of puberty are predisposed to developing reproductive and metabolic features of PCOS. However, further research is required to fully understand the role that prenatal exposure to excess androgens may play in the development of PCOS in humans as previous evidence shows contrary results.

**Restricted intrauterine growth**

It has been proposed that restricted intrauterine growth resulting in low birth weight predisposes girls to developing features of PCOS during puberty.

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**Table 2: Highlighting the targets and the benefits of different treatment methods used in PCOS.**

<table>
<thead>
<tr>
<th>Target</th>
<th>Benefits</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Non-pharmacological</strong></td>
<td>Weight loss</td>
</tr>
<tr>
<td>Hyperinsulinemia</td>
<td></td>
</tr>
<tr>
<td>Hyperandrogenemia</td>
<td></td>
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<tr>
<td>Hyperglycemia</td>
<td></td>
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<tr>
<td>Infertility</td>
<td></td>
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<tr>
<td>Obesity</td>
<td></td>
</tr>
<tr>
<td><strong>Pharmacological</strong></td>
<td>Improves ovulation and menstruation</td>
</tr>
<tr>
<td>Hirsutism related to hyperandrogenemia</td>
<td></td>
</tr>
<tr>
<td>Anovulation/ menstural dysfunction and infertility</td>
<td></td>
</tr>
<tr>
<td><strong>Surgical</strong></td>
<td>Improves acne and hirsutism</td>
</tr>
<tr>
<td>Blood androgens and testosterone levels</td>
<td></td>
</tr>
</tbody>
</table>

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**NON-PHARMACOLOGICAL**

Lifestyle modifications such as diet and exercises are natural methods that play pivotal roles in the management of PCOS. It is observed that lifestyle interventions have the potential to improve infertility issues and metabolic disturbances. It is also implied that lifestyle modifications can possibly reduce future risks for developing diabetes and cardiovascular diseases. All these benefits are derived from a general improvement in the state of insulin resistance that is usually observed with the application of lifestyle changes. Lifestyle modifications are considered as part of the first-line therapy for PCOS. Since the prevalence of obesity is quite alarming in women suffering from PCOS, weight reduction is given high priority during treatment. Weight loss in PCOS patients has been associated with observed improvements in metabolic and reproduction issues. Evidence reveals that weight reduction by as little as 5%, potentially improves ovulation cycles and restores normal menstruation.

The use of exercise regimens such as Hiit and steady training are shown to improve insulin sensitivity. Physical activities may also have some benefits on psychological manifestations as it helps to improve states...
of depression and anxiety. Weight loss resulting from exercises also positively impacts perceptions of body image which in turn helps to improve self-esteem. 53,54

Blood glucose levels stimulate pancreatic beta-cells to produce insulin and are normally affected by carbohydrate consumption. Since impaired blood glucose regulation and insulin resistance are commonly seen in PCOS patients, diet control is also an important factor to consider in the management of the disease. PCOS patients should consume diets low in carbohydrates. Examples of low carbohydrate diets (LCD) include: 1) ketogenic diet (daily carbohydrate intake is less than 30g, or a total of 5% of daily energy intake, and 2) mediterranean diet (involves the consumption of a variety of fruits and vegetables including nuts and seeds; also requires minimal intake of dairy products and red meat). 55 LCD diets have been shown to improve reproductive and metabolic problems in PCOS patients. In addition, they improve the symptoms that are related to hyperandrogenemia. They also aid in reducing blood glucose and insulin levels in addition to increasing FSH and SHBG. Increased circulation of FSH and SHBG reduce testosterone levels and also help to reverse the hypothalamic-hypophyseal dysfunction that is normally responsible for the fertility issues seen in PCOS. 56

Furthermore, the use of supplements has also been recommended for treating PCOS. Since different nutrients potentially regulate insulin signalling pathways and the synthesis of androgens, their deficiency may be linked to the development of some common complications of PCOS. 57 Therefore, the use of supplements in the form of vitamins or minerals may be used to complement the treatment of PCOS. 52 Vitamins such as vitamins A,B,D,E or inositol has antioxidant properties which have several potential benefits including: 1) help in oocyte maturation, 2) decrease cardiovascular risk factors, 3) improve menstruation, 4) improve fertility, 5) improve endometrial lining thickness, and 6) reduce androgen levels. Minerals such as calcium, chromium and zinc, may also contribute to improvements in ovulatory cycles, menstruation, and metabolic profiles in patients diagnosed with PCOS. 57

Another method that is observed for its effectiveness in the management of PCOS is fasting or intermittent fasting. During periods of fasting, blood glucose levels are low which in turn lowers blood insulin levels. Fasting combined with dietary and other lifestyle modifications have proven quite effective in lowering blood insulin levels, promoting weight loss, improving skin conditions and increasing ovulation. 58

Table 3: Summarizing the treatment methods available for treating polycystic ovarian syndrome.

<table>
<thead>
<tr>
<th>Non-pharmacological</th>
<th>Pharmacological</th>
<th>Surgical</th>
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</thead>
<tbody>
<tr>
<td><strong>Lifestyle modification</strong></td>
<td><strong>Hirsutism and hyperandrogenemia</strong></td>
<td></td>
</tr>
<tr>
<td>Diet</td>
<td>1) Estro-progestins pills</td>
<td></td>
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<tr>
<td>Exercise</td>
<td>Oral contraceptives combining oestrogen and progestin</td>
<td></td>
</tr>
<tr>
<td><strong>Dietary supplements</strong></td>
<td>2) Anti-androgens</td>
<td></td>
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<tr>
<td>Vitamins, A, B, D, E or inositol</td>
<td>Androgen receptor blockers</td>
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<tr>
<td>Minerals: calcium, chromium, zinc</td>
<td>Spironolactone</td>
<td>LOD</td>
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<td></td>
<td>Flutamide</td>
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<td></td>
<td>Alpha reductase inhibitors</td>
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<td></td>
<td>Finasteride</td>
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<td></td>
<td>3) Insulin-sensitizing agents</td>
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<tr>
<td></td>
<td>Biguanides-metformin</td>
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<tr>
<td></td>
<td>TZDs</td>
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<tr>
<td></td>
<td>Ovarian/menstrual dysfunction and infertility</td>
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<tr>
<td></td>
<td>MPA</td>
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<tr>
<td></td>
<td>Metformin</td>
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<tr>
<td></td>
<td>GLP-1 receptor agonists</td>
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<td></td>
<td>CC</td>
<td></td>
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<tr>
<td></td>
<td>Gonadotropins</td>
<td></td>
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<tr>
<td></td>
<td>Letrozole</td>
<td></td>
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<tr>
<td>GLP-1- Glucagon-like peptide-1, CC- Clomiphene Citrate, TZDs- Thiazolidinedones, LOD- Laparoscopic Drilling, MPA- Medroxyprogesterone acetate</td>
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</table>

**PHARMACOLOGICAL APPROACHES**

When weight loss seems difficult or in instances where PCOS patients are of normal body weight or are underweight, the use of various pharmacological agents may prove helpful in treating the disease. Pharmacological agents mainly target the symptoms and complications of PCOS such as hirsutism related to
hyperandrogenemia and anovulation\menstrual dysfunction and infertility.

**Hirsutism and hyperandrogenemia**

In cases where hirsutism is mild or localized, simple cosmetic measures may be sufficient for management and control. However, in instances where hirsutism is moderate or severe, pharmacological intervention becomes imperative. Treatment options usually preferentially target hyperandrogenemia, since excess androgen production is the main influencing factor for the development of hirsutism. In the presence of hyperandrogenemia, estrogen-progestins or anti-androgens are used as mono-therapy or in combination with ethinylestradiol. If required, insulin sensitizers such as metformin are also included as part of the treatment regimen.59

Oral contraceptive pills (OCPs) that combine oestrogen with progesterin are the primary option employed when managing hirsutism and acne in PCOS.60 Progesterin potentially suppresses LH levels, which results in decreased androgen production from the ovaries. Oestrogen, on the other hand, increases SHBG, thus reducing the levels of free androgens in the blood.61 These pharmacological options are excellent choices for controlling hirsutism in PCOS as they are inexpensive, safe and are very effective in controlling acne and hyperandrogenemia.60 Anti-androgens can also be successfully used for managing hirsutism and hyperandrogenemia when oestrogen-progestin combinations seem ineffective. Drugs that are currently used to counteract the effects of androgens in the blood include androgen receptor blockers such as spironolactone and flutamide, and 5α-reductase inhibitors such as finasteride.59 These classes of pharmacological agents work by reducing the effects or the levels of androgen in the blood and they have the added benefit of regulating blood lipid levels.60 Flutamide, though effective, potentially impairs liver function and should be used with caution or administered at lower doses when used for the management of PCOS in teenagers and adults. In general, anti-androgens can cause abnormal development in male foetuses; therefore, with their use, contraception is recommended.60 The combination of anti-androgens with oestrogen-progestin pills is proven more effective in successfully treating hirsutism as compared to the use of oestrogen-progestin pills alone.61

Insulin-sensitizing agents may also be used as part of the treatment regimen; however, their use is still questionable as evidence shows that their effects on hyperandrogenemia and hirsutism are only moderate.61 Medications that may be considered as insulin-sensitizers include biguanides (metformin) and thiazolidinediones (TZDs). Metformin can potentially control hyperinsulinemia and therefore, decrease insulin-stimulated androgen production by the ovaries. The use of TZDs in the treatment of hyperandrogenemia and hirsutism in PCOS may be limited as its use potentially supports weight gain.51

**Management of ovarian\menstrual dysfunction and infertility**

Ovarian dysfunction and chronic anovulation are common in PCOS patients and are due to hyperinsulinemia, hyperandrogenemia and obesity that are usually present as part of the syndrome. Anovulation is primarily responsible for observed infertility and reproductive issues; therefore, if ovulation is successfully restored fertility issues would also be resolved. With this in mind, the main aim when treating fertility problems in PCOS women is to achieve mono-follicular development.52,61

In PCOS patients there is hormonal dysregulation in the hypothalamic-pituitary-gonadal (HPG) axis, causing increased blood levels of androgens and oestrogen in addition to increased LH and decreased FSH levels. These effects lead to abnormal menstruation and amenorrhea (observed in 75% to 85% of PCOS patients).59 The use of medication in the management of ovarian dysfunction in PCOS should only be employed after the patient is fully educated on the risks and benefits that may accompany this method of treatment. Traditionally, OCPs are used in the long-term management of PCOS. OCPs prevent overgrowth of the endometrium and therefore, protect patients against cancers. OCPs are often used as they help to regulate menstrual cycles and improve co-morbidities by decreasing plasma levels of LH thus reducing hyperandrogenemia.59 It is important to note that the use of OCPs is not recommended in patients who smoke, has a history of breast cancer or in patients with a significant family history for heart diseases.64

The following medications may prove helpful in the management of ovarian\menstrual dysfunction and infertility in PCOS.

**Medroxyprogesterone acetate**

Medroxyprogesterone acetate (MPA) is a progestin or female hormone that potentially regulates ovulation and menstruation. This pharmacological agent can be applied to the management of various conditions which may include: amenorrhea or dysfunctional uterine bleeding. It can also be used to prevent endometrial overgrowth and therefore, reduces the risk for developing uterine cancers. MPA also has the added benefits of promoting insulin sensitivity and improving blood lipid profiles in PCOS patients.60 It can also be used in PCOS women to avoid pregnancies.

**Metformin**

Women who are diagnosed with PCOS face an increased risk for developing diabetes mellitus (DM) and usually
have elevated insulin levels. Insulin resistance is quite common in PCOS patients and it contributes significantly to the ovulatory abnormalities that are often present. Metformin can be used in PCOS women to reduce the risk of diabetes or to treat diabetes if it has already developed. Metformin is an insulin sensitizing agent and it therefore, promotes insulin sensitivity, thus decreasing blood insulin levels. This in turn suppresses insulin stimulated androgen production from the ovaries, potentially supporting menstruation. It is often used in women who have difficulties achieving weight loss. It is also used as an add-on therapy in women who experience menstrual irregularities or women who are unable to use estro-progestinics. This medication may also be prescribed particularly in instances where there are impaired glucose tolerance (IGT).

The use of metformin can also contribute to improvement in fertility in PCOS women. Metformin regulates the levels of reproductive hormones in the blood and therefore helps in restoring ovulation. In addition, metformin may also induce weight loss in PCOS women in whom infertility drugs are used to achieve ovulation. Metformin helps to improve fertility rates in PCOS women who experience fertility issues. This is achieved through an induced reduction in blood insulin levels, thus causing normal release of GnRH and gonadotropins.

Evidence suggests that prescribing metformin prior to the introduction of infertility drugs or the attempt of in-vitro fertilization, increases the rates of ovulation and successful pregnancies. Despite the promising effects of the use of metformin on fertility, its clinical effects may become evident following weeks of treatment as the onset of this drug is gradual and slow.

The use of metformin is associated with a myriad of side effects (nausea, diarrhoea, abdominal pain, bloating); therefore, its use is often substituted by other insulin sensitizers. Evidence suggests that treatment with TZDs is effective in PCOS patients and TZDs are prescribed as second-line management options in PCOS patients who are overweight or exhibit insulin resistance.

**Glucagon-like peptide-1 (GLP-1) receptor agonists**

Glucagon-like peptide-1 (GLP-1) receptor agonists is a novel class of drug proven effective in the management of T2DM. In addition to lowering blood glucose levels, GLP-1 receptor agonists also reduces body weight and promotes insulin sensitivity. Additionally, in PCOS patients who are obese or overweight, this drug class improves menstruation and suppresses blood levels of androgen. Combination therapies of GLP-1 receptor agonists and metformin are potentially more effective than their mono-therapy in the management of metabolic and reproductive abnormalities of PCOS; however, further studies are pending to determine the full effects of combination therapy with these agents.

**Clomiphene citrate**

Clomiphene Citrate (CC) provides a simple and cost effective way of promoting ovulation in PCOS women. It is described as a weak oestrogen-like hormone. This drug exhibits its effects on endocrine glands (hypothalamus, pituitary and ovaries), thus increasing the plasma levels of LH and FSH. This is achieved through the potential of CC to prevent the binding of estradiol to receptors in these major glands therefore, eliminating the negative feedback loop for endogenous oestrogens and estradiol. By inhibiting the negative feedback loop between the ovaries, pituitary gland and the hypothalamus, CC directly induces ovulation and thus directly target infertility issues that are observed in PCOS. Evidence reveals that the use of CC is associated with ovulation rates of about 60%-85% and pregnancy rates of about (30%-50%) following six ovulatory cycles.

CC is considered the first-line treatment option for the complication of infertility in PCOS. Increased FSH levels increases the chance for ovum maturation and ovulation. CC exhibits the potential to induce mono-follicular development and carries a low possibility for causing multiple pregnancies (2 to 13%).

The use of this drug is associated with side effects such as ovarian hyper-stimulation syndrome (OHSS), bloating, ovarian enlargement, hot flashes and multiple pregnancies. If following six months of treatment women are non-responsive to CC, then CC resistance is present. In instances like these, a second pharmacological agent is added or second-line therapy is required. Metformin is the most recommended add on therapy when resistance occurs and the combination of metformin with CC in the presence of CC resistance increases the rates of ovulation and successful pregnancies.

**Gonadotropins**

In PCOS patients who are not responsive to metformin or clomiphene, Human menopausal gonadotropin (HMG) daily injections serve excellent alternatives. Gonadotropins (LH and FSH) are hormones naturally produced by the pituitary gland. These hormones promote follicle maturation and ovulation. The use of gonadotropins in the treatment of ovulatory dysfunctions in PCOS may yield greater effects than the use of clomiphene. Despite this fact, CC is favoured as first-line therapy in the management of infertility issues in PCOS as it is easier to administer and is more cost effective as compared to the use of gonadotropins. Since FSH promotes follicular growth, its use, as part of therapy, may result in multiple pregnancies or OHSS due to an over response of the ovaries to the hormone. These risks factors are controlled through the co-administration of gonadotropins with letrozole as the use of letrozole reduces the length of stimulation and amount of gonadotropins required for a desired effect.
Letrozole

Letrozole prevents the conversion of androgens to oestrogen. As a result, when utilized in the management of PCOS, letrozole directly addresses fertility complications through its potential to inhibit the negative feedback loop between the ovaries, hypothalamus and pituitary gland. The mechanism of action of this drug promotes a rise in the plasma levels of FSH which in turn induces follicle development, maturation and ovulation. Letrozole is more effective than CC in addressing fertility abnormalities of PCOS as it is associated with improved gestational outcomes and greater events of live births. Even though the World Health Organization (WHO) supports the use of letrozole as first-line treatment for ovulation induction and some guidelines propose letrozole as an alternative to clomiphene, more research on safety and efficacy is required due to concerns risen regarding congenital malfunctions associated with the use of letrozole.71

SURGICAL APPROACH

The use of surgical interventions aids to decrease the plasma levels of LH and testosterone. This in turn promotes the successful restoration of normal, regular menstruation in addition to controlling symptoms of acne and hirsutism. Surgical methods directly destroy parts of the ovaries (follicles and surrounding stroma), therefore, reducing the amount of androgens that are available for conversion through aromatization. This induces an increased secretion of FSH, thus supporting follicular development, maturation and ovulation.72

Laparoscopic drilling (LOD) is the surgical method that is now preferred over laparotomy and ovarian wedge surgeries. This procedure can be utilized in patients who are resistant to CC. It carries many advantages which include: 1) helps to decrease potential risks for developing OHS, 2) promotes mono-ovulation, 3) supports higher pregnancy rates, 4) reduces length of time to pregnancy and 5) decreases the need for ovulation promoting drugs.72 Studies demonstrate that the performance of LOD on one ovary or both ovaries shows no major differences in outcomes. However, unilateral interventions are deemed more appropriate in reducing the occurrence of adverse effects. As with all surgical procedures, LOD may be associated with intra and post-operative complications. Bleeding and infections may occur. In addition, during LOD, if the ovaries are extensively damaged, the woman may enter menopause prematurely. It is also important to note that this procedure is not permanent and with time menstrual irregularities may re-occur.73

Table 4: Highlighting recommendation for the treatment of infertility in polycystic ovarian syndrome.74

<table>
<thead>
<tr>
<th>Treatment option</th>
<th>Recommendations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lifestyle modifications</td>
<td>In cases where the patient is overweight or obese, lifestyle changes are required as the first-line treatment for PCOS. It provides tremendous benefits in improving mental states, reproductive issues and metabolic abnormalities.</td>
</tr>
<tr>
<td>Pharmacological approaches</td>
<td></td>
</tr>
<tr>
<td>Letrozole</td>
<td>Recommended as part of the first-line treatment for inducing ovulation in PCOS.</td>
</tr>
<tr>
<td>CC</td>
<td>Recommended as part of the first-line treatment for PCOS; however, it is slightly less effective than letrozole.</td>
</tr>
<tr>
<td>Metformin</td>
<td>Can be used as a mono-therapy as part of the first-line treatment for PCOS; however, it is slightly less effective as compared to clomiphene citrate and letrozole.</td>
</tr>
<tr>
<td>Combination therapy of clomiphene citrate and metformin</td>
<td>This combination therapy can be considered for its use as a first-line therapy in the treatment of PCOS. Clomiphene citrate combined with metformin provides more benefits for obese patients as compared to the single use of metformin. In addition, in all women particularly those that are resistant to clomiphene citrate, the use of this combination is more effective as compared to the single use of clomiphene citrate. Despite these benefits that are potentially derived from this treatment combination, it is not recommended over letrozole as the first-line treatment in PCOS.</td>
</tr>
<tr>
<td>Gonadotropins</td>
<td>Can be considered as part of the second-line treatment regimen for PCOS in instances where first-line treatments are proven ineffective. The use of gonadotropins is more effective in women who are resistant to clomiphene citrate as compared to the use of the combination therapy of clomiphene citrate and metformin. In addition, in women that are resistant to clomiphene citrate, gonadotropins can be combined with metformin. Gonadotropins can be considered for use as first-line therapy and is seen as more effective when combined with ultrasound monitoring. In this instance, it should only be considered after the patient is educated on its cost and possible risk of multiple pregnancies.</td>
</tr>
<tr>
<td>LOD</td>
<td>Recommended as part of the second-line treatment regimen when patient is resistance to clomiphene citrate.</td>
</tr>
</tbody>
</table>

CC- Clomiphene citrate, LOD- Laparoscopic ovarian drilling, PCOS- Polycystic Ovarian Syndrome Laparoscopic ovarian drilling
CONCLUSION

In conclusion, PCOS is a multifactorial disorder which may develop secondary to hormonal, neuroendocrine, genetic and developmental factors. Its management involves the combination of non-pharmacological, pharmacological and surgical interventions. PCOS remains a complex condition and more research is required to further understand the role that genetics and excess androgen exposure during development may play in its development. In addition, more exploration is needed into the side effect profile of the drugs that are currently used for the treatment of this condition and probably into the production of drugs that are capable of targeting the multiple complications that accompanies PCOS.

Funding: No funding sources
Conflict of interest: None declared
Ethical approval: Not required

REFERENCES

23.  Laven JS, Mulders AG, Visser JA, Themmen AP, De Jong FH, Fauser BC. Anti-Mullerian hormone serum concentrations in normoovulatory and anovulatory


59. Pasquali R. Contemporary approaches to the management of polycystic ovary syndrome. Ther Adv Endocrinol Metab. 2018;9:123-34.


