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SEXUAL RECEPTIVITY, REPRODUCTIVE HORMONES AND PERFORMANCE TRAITS OF AGING FEMALE RABBITS AS AFFECTED BY DIFFERENT SOURCES OF UNSATURATED FATTY ACIDS

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Sixty V-lines doe rabbits, 24 months of age weighing 3244 ± 58.9 g, which had low conception rate and repeated refuse of mating were used. Rabbits were randomly divided into five equal treatment groups and orally administered with 3 ml of different sources of oil/doe/day, for eleven consecutive days prior to insemination as follows: group 1 sterilized water and served as control (C), group 2 sesame oil (Ses), group 3 linseed oil (Lin), group 4 sunflower oil (Sun), group 5 mixture oil (Mix) containing equal portions of Ses, Lin and Sun oils. Estradiol (E217- β), progesterone (P4) and prostaglandin (PGF_{2a}), sexual receptivity of does, fertility rate, litter size and bunny weight at birth were measured. **Results revealed that:**

1- All treated groups represented significant ($P \le 0.01$) increase of hormonal profile (E217- β , P4 and PGF_{2a}), sexual receptivity, fertility rate, litter size and bunny weight at birth compared to the control group.

2- Aged doe rabbits supplemented with Ses and Sun oils recorded highest concentrations of $E2_{17\beta}$ and $PGF_{2\alpha}$ and decrease of P_4 than Lin and Mix oils or the control groups.

3-Sexual receptivity and fertility rate were improved ($P \le 0.01$) for groups supplemented with Ses and Sun oils compared with other groups.

4- Litter size and bunny weight at birth were significantly ($P \le 0.01$) increased for groups supplemented with Lin and Mix oils compared with other groups.

Conclusively, supplementing the aged doe rabbits with Ses, lin and Sun oils or their mixture for 11 consecutive days perior

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insemination could be beneficial for improving estradiol and prostaglandin hormone with subsequent improvement of does receptivity, litter size and bunny weight combined with progesterone reduction

Key words: Aging doe rabbits, reproductive hormones, fertility rate, sexual receptivity.

Reproductive rabbits capacity is starting at puberty age, increases for a short time and eventually decreases at a rate characteristic of the species (*Talbert, 1968*). Falling in fertility with age suppose due to an attrition of the oocyte population (*Ingram et al., 1958*), increase in the incidence of defective oocytes (*Henderson and Edwards, 1968*) and reduce in the potency of the uterus to maintain pregnancy (*Adams, 1970*). Furthermore, ovaries can not produce steroid hormones in aging animals (*Deng, 2012*) and reduce the capacity of the hypothalamic-pituitary-gonadal axis in reproductive system (*Walker et al., 2013*).

In context, aging women oocytes may contain abnormal chromosome division, decreased mitochondrial quality, including the accumulation of mutations in the mitochondrial DNA and low ATP production, increased oxidative stress, and decreased antioxidant levels, also, decreases drastically the quality of fertility with the qualitative deterioration of oocytes (Meldrum et al., 2016). In addition, old age mares is associated with increased accumulation of inflammatory cells within the endometrium, reduced pregnancy rate, and increased chance of pregnancy loss when compared with younger mares (Carnevale and Ginther, 1992). Also, Simmen et al., (2015) reported that aging of mice uteri leads to the down regulation of several genes associated with cell proliferation, indicating the presence of senescent cells with impaired proliferation owing to uterine aging. On the other hand, aging female rabbits able to still produce an adequate number of fertilizable oocytes which are capable of normal development provided but uterus incapable to maintained pregnancy (Maurer and Foote, 1971) which lead to decreased litter size this is not due to low ovulation rate (Thorneycroft and Soderwall, 1969). Furthermore, decrease in progesterone flow by the ovaries has been implicated failure pregnancy in aging rabbits (Maurer and Foote, 1972). In the context, Larson et al., (1973) who reported that aging female rabbits were given supplemental exogenous progesterone and/or oestradiol benzoate in an effort to increase reproductive efficiency. Also, elevated basal FSH and LH can induce reproductive and endocrinological signs of aging women (Ahmed et al., 1994).

Several studies on lipids metabolism have shown that alpha linolenic acid (ALA) (18:3 n-3) and linoleic acid (LA) (18:2 n-6) can not be synthesized in the most mammals' species including rabbits due to the lack of appropriate fatty acid desaturase enzymes and consequently they must be supplied in the diet (*Vrablik and Watts, 2013*).

An important function of PUFAs is the components of phospholipids in cell membrane. Different types of cells, membranes and phospholipids have different compositions of PUFAs and this fatty acid composition may affective by metabolism, state of cell activation, hormone and genetic factors (*Calder*, 2015). The physical features of cell membrane that include fluidity and permeability are influenced by PUFAs acids composition of cell membrane. Also, the PUFAs as precursors are related to PG family synthesis and steroidogenesis (*Stocco et al.*, 2005).

Polyunsaturated fatty acids (PUFAs) influence oocyte maturation, fertilization and subsequent development through altering the biochemical composition of follicular fluid (*Bilby et al., 2006*). Another studies by *Scott et al., (1995*) showed that cows fed with supplements containing long chain PUFAs present stronger estrous had more active ovaries and less derived PGF_{2a} was requiring for estrous induction. Another study by *Fouladi-Nashta et al.,* (2009) reported that fatty acid composition in granulosa cells were changed by supplementation and it had influence to oocyte maturation and subsequent development *in vivo*. Otherwise, *Wathes et al., (2007)* reported that omega-3 a one of PUFAs regulates different types of reproductive function in mammals.

Cholesterol is described to be the precursor for the synthesis of steroid hormones, P4 and E2, while AA is the precursor for $PGF_{2\alpha}$ (*Santos et al., 2008*), $PGF_{2\alpha}$, P4 and E2 are essential hormones induct in ovulation, oestrous cycle and maintenance of pregnancy in dairy cows (*Field and Taylor, 2008*).

Therefore, the objective of this study was to investigate the effect of oral administration of polyunsaturated fatty acids (different sources of omega-3 and omega-6) on some reproductive hormones, sexual receptivity prior insemination, fertility and litter size and bunny weight at birth for aged doe rabbits.

MATERIALS AND METHODS

The present work was carried out at a private rabbit farm at Qaluobia Governorate from October 2017 to April 2018 (temperature ranged from 19 to 27°C while, humidity was 43 to 54% and light period 16 hr light : 8 hr dark).

Experimental design:

Sixty aged V-line does 24 month age, with average body weight of 3244 ± 58.9 g, had low conception rate and repeated refuse of mating were used for the present study.

Rabbits were randomly divided into five equal treatment groups and orally administered with 3 ml of different sources of oil/doe/day, for eleven consecutive days prior insemination as follows:

Group 1: 3 ml sterilized water was given orally and served as control group (C),

Group 2: Orally 3 ml of Sesame oil* (Ses), Group 3: Orally 3 ml of Linseed oil* (Lin), Group 4: Orally 3 ml of Sunflower oil* (Sun), Group5: Orally 3 ml of Mixture oils** (Mix)

*Mixture oils. Contains equal portion of Ses, Lin and Sun oils.

Housing and management:

The rabbits were housed in a naturally ventilated building and kept in individual wire galvanized cages ($60 \times 55 \times 40$ cm). Batteries were accommodated with feeders for pelleted rations and automatic drinkers. Animals were kept under similar management, hygienic conditions, healthy and clinically free of external and internal parasites.

Diet nutrient profiles:

Does were allowed to a standard pelleted diet (Table 1). Feed and water were available *ad libitum*.

Polyunsaturated fatty acids composition of Ses, Lin and Sun oils are presented in Table 2.

Does were inseminated using a heterospermic pool diluted 3 times (1 Semen : 3 extender). A pool of semen was collected from bucks of proven fertility for artificial insemination (AI), which performed by depositing 0.5 ml of fresh diluting semen deeply in the upper of vagina by sterile catheter.

Semen was diluted with extender stored at room temperature (20° C) and used within 4 hr of collection (semen ejaculates were individually evaluated microscopically and the ejaculates, which showed active progressive motility percentages ($\geq 70\%$) were pooled and extended to Tris-buffer extender.

Ingredients	<u>useu</u> %
Vellow corn	6.22
Sovbean meal 44%	22 33
Wheat bran	22.33
Barley	15.00
Alfalfa hav	30.12
Ground limestone	1.00
Di-calcium phosphate	1 20
Common salt	0.50
Vit + Min premix [*]	0.30
Total/kg	100
Calculated analysis ^{**}	100
Crude protein. %	18.8
Crude fiber. %	13.0
Ether extract. %	3.0
Digestible energy (kcal/kg diet)	2680
n-6 PUFAs%	0.3
n-3 PUFAs%	1.03
n-6 / n-3 ratio	0.3:1
Determined analysis (g/kg)	
Dry matter	902.4
Organic matter	912.1
Crude protein	172.4
Crude fiber	138.5
Ether extract	26.2
Nitrogen-free extract	575.0
Ash	87.9
Cell wall constituents (g/kg)	
Neutral detergent fiber (NDF)	331.1
Acid detergent fiber (ADF)	160.9
Hemicellulose	170.2

Table1. Composition and determined analysis (on a dry matter basis) of the experimental commercial diet used

Each 3kg of premix contains: Vit. A: 12,000,000 IU; Vit. D3: 3,000,000 IU; Vit. E:10.0 mg; Vit. K3: 3.0 mg; Vit. B1: 200 mg: Vit. B2: 5.0 mg Vit. B6: 3.0 mg: Vit. B12: 15.0 mg; Biotin: 50.0 mg; Folic acid: 1.0 mg; Nicotinic acid: 35.0 mg: Pantothenic acid: 10.0 mg; Mn: 80 g; Cu: 8.8 g; Zn: 70 g; Fe: 35 g; I: 1 g; Co: 0.15g and Se: 0.3g.

	Mean Fatty acids %								
Type of oil	Palmatic	Stearic	Olic	Linoleic	Linolenic	n-6/n3			
	16:0	18:0	18:1	(n-6)18:2	(n-3)18:3	Ratio			
Sesame oil (Ses) ¹	8.47	5.53	41.63	42.77	0.42	101:1			
Linseed oil (Lin) ²	5.51	3.8	13.5	14.8	62.1	0.24:1			
Sunflower (Sun) ³	6.65	4.97	52.83	33.15	0.51	65:1			

 Table 2. Polyunsaturated fatty acid analysis of sesame, linseed and sunflower oils

Polyunsaturated fatty acid composition of Sesame, linseed and sunflower oils are based on references by Hassan $(2012)^1$, El-Prollosy, $(2012)^2$ and Ismaill and Arafat $(2014)^3$, respectively.

*Mix oil, the ratio of n-6/n3 ratio calculated according to previous analysis recorded 1.4:1

The final concentration rate was $80-100 \times^{10}$ motile sperm/ml. The insemination was immediately followed by the administration of 0.8 µg of buserelin IM (0.2 ml Receptal; Hoechst, Frankfurt, Germany) to induce ovulation (Lopez and Alvariño, 2000).

Blood biochemical constituents:

Blood samples (3 ml) were withdrawn at morning from marginal ear veins for each treatment group before access feed and water.

Blood samples were centrifuged at 3000 r.p.m for 15 min to obtain clear serum samples and stored at -20 °C until analysis the following hormones.

Determination hormones:

Serum samples were used for determining the concentrations of E217- β (the Estradiol ELISA Test Kit has a sensitivity of 6.5 pg/ml) and P4 (the Progesterone ELISA Test Kit has a sensitivity of 0.105 ng/ml) by Immunoassay Technique Elisa Kits (Fortress Diagnostics Ltd, Antrim, UK), and PG concentrations of prostaglandin (PGF_{2a}) were measured using a PG ELISA kit (the Sensitivity: 50% B/B0:52 pg/ml) (Cayman Chemicals, Ann Arbor, MI, USA).

Reproductive performance:

Sexual receptivity of does was tested in the presence of a vasectomies buck as described by the International Rabbit Reproduction Group (2005). Fertility rate (number of parturitions/number of inseminations×100), litter size (total number of born) and bunny body weight at birth were recorded.

Statistical analysis:

All data were subjected to analysis of variance as described in SAS (program) (SAS, 2002).

The significant means differences among groups were separated by Duncan's multiple rang test (Duncan, 1955).

RESULTS AND DISCUSSION

Reproductive hormones concentration:

Table 3 shows the effect of different PUFAs oils administration as sources of n-3 and n-6 for aging does on E217 β , P4 and PGF_{2 α} hormonal concentrations. The obtained data revealed that E217 β and PGF_{2 α} hormone concentrations for all groups were higher (P \leq 0.01) compared to those for control group. However the rabbits of Ses and Sun oils groups recorded a highest E217 β , PGF_{2 α} concentration and low progesterone value than those for the Lin and Mix oils or control groups. Results show that there were no statistical differences between Lin and Mix oils groups with respect to E217 β , P4 and PGF_{2 α} hormones. On the other hand, control group exhibited significantly the lowest concentration of E217 β and PGF_{2 α} besides highest level of progesterone hormone.

Essential reproductive hormones are E2, P4 and PGF_{2α}. Estradaiol17 β is involved with the preparation of the reproductive genital tracts for ova fertilization and the initiation of pulsative surge of LH (Ball and Peters, 2004). Progesterone is one of the majority prominent fertility hormones safe for carrying out pregnancy to period, whereas PGF_{2α} acts against the service of P4 on the CL after failed fertilization (Funston, 2004). Cows fed sources of omega-6 PUFAs are known to produce more cholesterol. Omega-3 PUFAs enables P4 to be synthesized by negatively affecting PGF_{2α} synthesis (Mattos *et al.*, 2004). The cholesterol can be used in the presence of steroidogenic acute regulatory hormone to synthesize P4 (Piccinato *et al.*, 2010).

Different type of cells, membranes and phospholipids have various compositions of PUFAs and these fatty acids composition may be affected by metabolism, state of cell activation, hormone, and genetic factors (Calder, 2015). The physical features of cell membrane that include fluidity and permeability are influenced by unsaturated fatty acids composition of cell membrane, furthermore, the unsaturated fatty acids as precursors are

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Parameters	Control	Ses	Lin	Sun	Mix	Sig test
E217-	22.23 ^c	47.65 ^a	33.98 ^b	45.21 ^a	35.22 ^b	**
B,(pg/mL)	±0.84	±0.91	±0.41	±0.91	±0.54	
P4	4.39 ^a	1.88 ^c	2.45 ^b	1.89 ^c	2.37 ^b	*
(ng/ml)	±0.04	±0.07	± 0.08	±0.06	±0.12	
PGF2a	450.97 ^c	612.34 ^a	580.25 ^b	605.12 ^a	598.22 ^b	**
(ng/ml)	±12.23	±17.4	±13.5	±13.6	±12.23	

Table 3. Effect of unsaturated fatty acids on serum estradaiol17- β , progesterone and prostaglandin F_{2a} for aged V-line doe rabbits

a,b,c Means within the same row for each effect with different superscripts are significantly different (P \leq 0.05).

Sesame oil = Ses, Linseed oil = Lin, Sunflower oil = Sun, Estradiol 17- β , =E217- β pg/mL, Progesterone =P4 ng/ml, Prostaglandin=, PGF_{2a} ng/ml

* P≤0.05 ** P≤0.01

related to prostaglandins (PGs) synthesis and steroidogenesis (Stocco *et al.*, 2005).

Current results indicated that increasing concentration of $PGF_{2\alpha}$ by treatment with different oils are in agreement with those reported by Petit et al., (2004) who revealed that the linoliec acids as Ses and Sun oils (C18:2,n-6) are converted into arachidonic acid (C20:4, n-6), which is the precursor of the dienoic (2-series) PGs, such as PGF_{2α}, while, omega-3 in Lin oil (α-linolenic acid; C18:3n-3) is converted into eicosapentaenoic acid (C20:5n- 3), which is the precursor of the trienoic (3- series) PGs, such as PGF_{3α}.

Moreover, Mattos *et al.*, (2004) showed that feeding a diet rich in linoleic acid could contribute to increasing secretion of PGF2 α , compared with feeding a diet rich in linolenic acid. Harmony with the context, Lands (1992) reported that the proportion of different PUFAs in the diet alters cell membrane phospholipid composition and this becomes quantitatively significant because the precursors of each group of PGs compete for the same enzyme systems for metabolism. This in turn will have considerable effects on the types of PG synthesized and hence, physiological responses (Abayasekara and Wathes, 1999). Also, Cheng *et al.*, (2004) recommended that the supplementation of linoleic acids to endometrial cells in vitro isolated from late pregnant ewes caused a significant decrease in production of 2-series PGs.

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Diminishing P4 concentration due to using unsaturated fatty acids in this excremental is in harmony with results reported by Rebollar et al., (1992) who stated that the PGF_{2 α} posses luteolytic properties in rabbits and induce a fall in P4 concentration. Same results obtained by Robinson et al., (2002) who demonstrated that feeding cows with a Linseed rich diet (linoleic acid, C: 18:3, n-3) leads to significant reduce of plasma P4 concentrations. Likewise, Acosta et al., (2002) and Beatrice and Ulrich, (2012) reported that administration of $PGF_{2\alpha}$ significantly decreased the volume of the CL and the blood P4 concentration of cyclic cows. On the other hand, Tran et al., (2016) demonstrate that the DHA increase GnRH enhancer-derived RNA levels. Overall, these provide evidence that GnRH neurons directly sense FFAs. This will advance our understanding of the mechanisms underlying FFA sensing in the brain and provides insight into the links between nutrition and reproductive function. Furthermore, Thatcher *et al.*, (2006) revealed that injected of $PGF_{2\alpha}$ (0.5 mg dinoprost, a synthetic analogue of $PGF_{2\alpha}$) impacts on increasing blood PG profile and decreasing blood P4 and this effect provided the opportunity to increased secretion of gonadotropin hormones from the pituitary and secretion estrogen from the ovary through inhibiting the effect of P4 on gonadotropin release. Whereas, increased biological effect of $PGF_{2\alpha}$ from Sun led to a decrease in blood P4 and an increase in blood 17- β E2 level, which is necessary for the growth and maturation of oocytes (Thatcher et al., 2006). Besides, E2 can increase the sensitivity of the CL to $PGF_{2\alpha}$, thus causing a more complete regression of the CL (Howard et al., 1990). On the other hand, $PGF_{2\alpha}$ reduced the production of P4 because PUFAs converted to $PGF_{2\alpha}$ that caused an increase PGF concentration in the plasma with a reduction of the P4 synthesis (Thatcher et al., 2006). According to Smits (2010), the follicular phase begins after luteolysis and ends at ovulation. Gonadotropins (GnRH) released from hypothalamus, FSH and LH, released from the anterior pituitary, stimulate follicles to produce E2 and the follicular phase is dominated by E2, which produced by developing follicles (Adams, 1999).

Present data on Ses and Sun oils revealed that the $PGF_{2\alpha}$, which produced in vivo from n-6 (PG precursor) more effective on reproductive functions of rabbit does than that of the synthetic $PGF_{2\alpha}$. Also, current results revealed that Ses and Sun oils as a precursor of $PGF_{2\alpha}$ had a biological activity more than the Lin and Mix oils as a precursor of $PGF_{3\alpha}$. This finding was consistent with the finding of Lauderdale (2002) who found that in domestic animals, the most important and practical utility of PG appears to be $PGF_{2\alpha}$ and the actions of $PGF_{2\alpha}$ are mediated by the PGF receptor (Sugimoto *et al.*, 1994). Finally, Elkomy and Elspeiy (2015) reported that treated aged rabbit does by oral administration with Sun oil improved reproductive hormone synthesis and secretion, this demonstrated that treated aged does with PUFAs is a good tool for improving the reproductive traits.

Reproductive performance:

Date of Table 4 showed that all groups of rabbit does received different oil sources either separated or combined represented significant (P \leq 0.01) increase of sexual receptivity, fertility rate, litter size and body weight at birth in treated groups compared with control. However, rabbits of Ses and Sun oils groups significantly (P \leq 0.01) recorded highest sexual receptivity and fertility percentages besides low litter size and body weight at birth than those for Lin and Mix oils groups. Also, results showed that there were no statistical differences between Ses and Sun oils groups with respect to previous mentioned parameters. Interestingly, doe rabbits of Lin and Mix oils groups represented highest (P \leq 0.01) records than other groups with respect to litter size and body weight at birth, while control group recorded low values for all studied parameters.

Table	4	Effe	ct of	unsatu	irated	fatty	acids	on	sex	ual r	recept	ivity,	fertility
		rate,	litter	size	and	bunny	/ wei	ght	at	birtl	h of	aged	V-line
		multi	parous	s does									

Parameters	Control	Ses	Lin	Sun	Mix	Sig test
Sexual receptivity,	58.43 ^c	82.67 ^a	75.34 ^b	84.87 ^a	77.35 ^b	**
%	±2.89	± 1.78	± 1.42	± 1.84	±1.44	
Fertility rate,	51.19 ^c	80.45 ^a	71.23 ^b	82.22 ^a	73.42 ^b	**
%	±1.56	±1.54	±124	±1.31	±1089	
L.S.at birth	4.98 ^c	8.11 ^b	8.54 ^a	8.08 ^b	7.75 ^b	**
(n)	±0.73	±1.45	±0.61	± 0.56	±0.67	
B.W.at birth	41.21 ^c	44.11 ^b	47.25 ^a	43.65 ^b	45.82 ^a	**
(g)	±0.61	±0.59	±0.34	±0.78	±0.62	

a,b,c Means within the same row for each effect with different superscripts are significantly different ($P \le 0.05s$).

Sesame oil = Ses, Linseed oil = Lin, Sunflower oil = Sun Litter size at birth (n)=L.S.at birth (n), Bunny weight at birth (g)=B.W.at birth (g)

** P≤0.01

Current results for that increasing concentration of $PGF_{2\alpha}$ by treatment with different oils and sexual receptivity improvement are in agreement with those reported by Dragan *et al.*, (1996) who mentioned that sexual receptivity of rabbit does by rise level of $PGF_{2\alpha}$ analogues is improved than the control. Also, Hassanein (2000) found that the increased percentage of mating acceptance was 100% when does were treated with lutalyase (PGF_{2a} analogues) 48 h before natural mating.

The results of this research proved that amendment in dietary PUFAs significantly increases fertility rate due to raise of $PGF_{2\alpha}$. This result is keeping with those previously reported by Facchin et al., (1992) who stated that injected $PGF_{2\alpha}$ before insemination for rise PG level to induce oestrus synchronization of multiparous doe rabbits. Moreover, Alvarino et al., (1995) observed that $PGF_{2\alpha}$ improved the fertility rate in nulliparous and multiparous inseminated does. Similarly, results of Gogol (2009) showed that simultaneous treatment with $PGF_{2\alpha}$ to induce oestrus synchronization can increase reproductive performance in postpartum rabbit does, whereas, fertility rate and litter size were significantly increased in multiparous does treated with unsaturated fatty acids. On the other hand, Adamiak et al., (2005) showed that PUFAs content in follicular fluid is highly correlated to that of the diet and it is generally accepted that alterations in dietary fatty acid intake cause a similar shift in the fatty acid profile of the follicular fluid. In addition, Zachut et al., (2008) mentioned that supplemented diet with lipid increases the size of the pre-ovulatory follicle and its production of E2. Otherwise, the lipids stored within the oocyte and early embryo represents an important source of energy for the early embryo (Mc Keegan and Sturmey, 2012). Moreover, Kowalska (2008) found that the female rabbits fed a complete diet fish oil (n-3) supplemented had a quantitavely and qualititately better milk fat content, higher fertility prolificacy values, and higher body weight of young rabbits at birth. The observed effects in this study regarding the increase of litter size, fertility and higher body weight at birth support the conclusion of Nemeth et al., (2017) who recorded that PUFAs supplementation may contribute to an adjustment of energy accumulations to the total litter mass, which may both favor a mother's body condition at parturition and perhaps increase the offspring survival at birth.

In conclusion, supplementing the aged doe rabbits with sesame, linseed, sunflower and their mixture could be a useful tool for improving reproductive hormones, sexual receptivity, fertility rate, litter size and weight at birth.

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تم إستخدام عد ٦٠ أنثى من خط V-Line، عمر ٢٤ شهر بمتوسط وزن جسم ٣٢٤٤ ± ٩.٨٩م منخفضة فى معدل الحمل وتكرر رفضها للتزاوج حيث تم توزيعها على ٥ مجاميع تجريبية عشوائيا وتغنت جميعها على العليقة الأساسية بينما تم تجريع الأرانب بالفم ٣ مل من الماء المقطر (كنترول) أو زيت السمسم أو زيت الكتان أو زيت عبد الشمس أو خليط بنسب متساوية من كل من زيت السمسم والكتان وعباد الشمس وذلك لمدة ١١ يوم متصلة قبل التلقيح : وقد أوضحت النتائج ما يلى:

- ارتفع مستوى هرمون الأستراديول فى سيرم الدم بدرجة عالية فى المجاميع المعامله بزيت السمسم أو زيت عباد الشمس مقارنة بالمجاميع المعاملة بزيت الكتان أو خليط الزيوت أوالكنترول. EL-SAWY et al.

- إنخفض معنويا على مستوى ١% مستوى هرمون البروجستيرون فى كل المجاميع المعاملة بزيت السمسم أو زيت الكتان أو زيت عباد الشمس أو خليط الزيوت مقارنة بمجموعة الكنترول. - لوحظت زياده معنوية على مستوى ١% فى قابلية الإناث للتزاوج ومعدل الخصوبة لكل المجاميع المعامله بالزيوت الثلاثة أو خليطهم مقارنة بالكنترول. - حدثت زيادة معنوية على مستوى ١% فى عد ووزن الخلفة عند الميلاد للمجاميع المعاملة بالزيوت الثلاثة أو خليطهم مقارنة بالكنترول. - أظهرت الإناث المعاملة بزيت السمسم أو عباد الشمس أو خليط الزيوت أعلى عد وزن الخلفة عند الميلاد مقارنة بمجموعة الكنترول. - حدثت تحسن فى قابلية الإناث المعاملة بزيت السمسم أو عباد الشمس أو خليط الزيوت أعلى عد وزن الخلفة عند الميلاد مقارنة بمجموعة الكنترول. - حدث تحسن فى قابلية الإناث التزاوج بعد تجريعها كل من زيت السمسم أو زيت الكتان أو زيت عباد الشمس أو خليطهم مقارنة بمجموعة الكنترول. التوصية: تجريع إناث الأرانب المتقدمة فى العمر والمنخفضة الخصوبة ٣ مل بالفم من زيت السمسم أو زيت الكتان أو زيت عباد الشمس او خليطهم لمدة ١١ يوم متصلة قبل التلقيح يؤدى إلى تحسن مستوى هرمونات التناسل، القابلية التزاوج ،عدد الجيسم مد الميلاد قبل التلقيح يؤدى والخصائص التناسلية.