Primates: Hylobatidae



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Contraceptive methods:	GnRH agonist (implant)	GnRH agonist (injection)	Progestagen (implants)	Progestagen (implants)	Progestagen (injection)	Combination Birth-Control Pills	Progestagen only Birth-Control Pills	Surgical/ Permanent
Contraceptive Product:	Deslorelin acetate	Leuprolide acetate	Etonogestrel 68 mg	Levonorgestrel 2x 75mg	medroxyprogesterone acetate	Combinations of a synthetic progestagen and oestrogen at various doses are available	Oral synthetic progestagens without any oestrogen component	N/A
Commercial Name:	Suprelorin ®	Lupron °	Implanon® Nexplanon®	Jadelle*	Depo-Provera®, Depo-Progevera®,	Several commercial oral combination pills are available in the market for human use.	Several commertial oral progestagen pills are available in the market for human use.	Vasectomy
Product Availbility:	4.7mg ('Suprelorin 6') and 9.4 mg ('Suprelorin 12') widely available through veterinary drug distributors in the EU. 9.4 mg ('Suprelorin 12') is also available through Peptech Animal Health, Australia.	Leuprolide acetate licenced for human use	Manufactured by Organon. Available through human drug distributors	Manufactured by Organon. Available through human drug distributors	Manufactured by Pfizer. Widely avilable throughout Europe through human drug distributors.	Widely availabe in pharmacies for human use	Widely availabe in pharmacies for human use	N/A
Restrictions and/or permit required by Importing Country:	EGZAC recommends: always check with your local licencing authority	Data deficient	EGZAC recommends: always check with your local licencing authority	EGZAC recommends: always check with your local licencing authority	EGZAC recommends: always check with your local licencing authority	N/A	N/A	N/A
Mechanism of action:	GnRH agonist suppress the reproductive endocrine system, preventing production of pituitary and gonadal hormones. As an agonist of the GnRH initially stimulates the reporductive system -which can result in oestrus and ovulation in females or temporary enhancement of testosterone and spermatogenesis in males - therefore additional contraception needed during this time. Please see below and refer to Desiorelin datasheet for detailed information.	GnRH agonist suppress the reproductive endocrine system, preventing production of pituitary and gonadal hormones	Interference with fertilization by thickening cervical mucus, interrupting gamete transport, disruption of implantation, inhibition of LH surge necessary for ovulation	Interference with fertilization by thickening cervical mucus, interrupting gamete transport, disruption of implantation, inhibition of LH surge necessary for ovulation	Anti-estrogenic activity. Interference with fertilization by thickening cervical mucus, interrupting gamete transport, disruption of implantation, inhibition of LH surge necessary for ovulation	Inhibit follicular development and LH surge preventing ovulation. Progestagen part also blocks fertilisation and/or implantation.	Interference with fertilization by thickening cervical mucus, interrupting gamete transport. Disruption of implantation. Inhibition of the LH surge necessary for ovulation. These mechanisms are dose dependant, typically higher dose of synthetic progestagens are required to block ovulation than to block fertilisatio and/or implantation.	Surgical procedure in which the ductus deferens are cut, tied, cauterized, or otherwise interurrupted

Insertion/Placement:	Sub-cutaneous, in a place where it can be easily detected or seen for removal at a later date (I.e. upper inner arm); refer Suprelorin fact sheet for effective method of implant (tunnelisation) Data deficient	Injectable Data deficient	Intramuscular or subcutaneous. EGZAC recommends sub-cutaneous, upper inner arm for visibility (aid for later removal)	Intramuscular or subcutaneous, EGZAC recommends sub-cutaneous, upper inner arm for visibility (aid for later removal)	Injectable intramuscular	Oral Data deficient	Oral	Surgical
Females	Data deficient	Data deficient				Data deficient		
Dose	Dosages and duration of efficacy have not been well established as a guide for hylobatidae: 1 x 4.7 mg; 1 x 9.4 mg	Dosing information is not available; extrapolation from human literature is likely the best place to start	Doses not well established. Recommended 1/2 to 1 implant, depending on species and weight	Recommended 1 rod. Doses not well established .	2.5-5 mg/kg body weight every 45-90 respectively days has been effective in most NHP secies	Dosage has not been well established; whole and half pills have been given. There are some reports that commercially available birth control pills are effective.	Dosage has not been well established; whole and half pills have been given. There are some reports that commercially available birth control pills are effective.	N/A
Latency to effectiveness:	3 weeks average as GnRH agonist initially stimulates the reporductive system-please refer to Deslorelin datasheet for detailed information - additional contraception needed during this time (see product data sheet. "Zmg/Kg Megestrol acetate pills daily 7 days before and 8 days after has been used)	Same as desiorelin with an initial stimulation phase and suppression should then occur 3-4 weeks later (please refer to desiorelin and lupron datasheet for more details)	In general inhibition of ovulation after 1 day when inserted on day 1-5 of cycle or when replacing oral progestogen. As the right stage during menstrual cycle is often unknown, it is advised to use other contraceptive methods for at least 7-14 days after insertion of the implant depending on administration route (Im or SC)	In general inhibition of ovulation after 1 day when inserted on day 1-5 of cycle or when replacing oral progestogen. As the right stage during menstrual cycle is often unknown, it is advised to use other contraceptive methods for at least 7-14 days after insertion of the implant depending on administration route (Im or SC)	1-3 days post injection. However, if the cycle stage is not known then extra time must be allowed; therefore, separation of the sexes or alternative contraception should be used for at least 1 week.	1 to 2 weeks but can take up to one month if treatment starts near the time of ovulation (refer to product insert for exact information on this)	1 to 2 weeks, although this varies depending on the brand. Please read the packet insert. The packet will outline when to start and how long to use secondary protection and/or how long the individual may need to be separated.	N/A
Oestrus cycles during contraceptive treatment:	Initial oestrus and ovulation (during the 3 weeks of stimulation) then no oestrus cycle. To supress the initial osetrus and ovulation you can follow the megestrol acetate protocol mentioned above.	Same as deslorelin.	Oestrus is inhibited. Menstruation in non- human primates are more or less present with regular cyclicity. This is an individual and dose-dependent response. Some will swell during treatment and some will not.	Oestrus is inhibited. Menstruation in non- human primates are more or less present with regular cyclicity. This is an individual and dose-dependent response. Some will swell during treatment and some will not.	Oestrus behaviour may be observed. Ovulation and cycling can occur in adequately contracepted individuals (but is unlikely and the degree of suppression is dose dependent).	Sings of oestrus can occur druing the placebo week if treatment not administer continuously (placebo week not necessary)	Oestrus behaviour may be observed. Ovulation and cycling can occur in adequately contracepted individuals (but is unlikely and the degree of suppression is dose dependent). Be aware that progestagen-only pills are not being as effective at suppression oestrus as the combination pills.	N/A
Use during pregnancy:	Not recommended	Not recommended	In non-human primates progestagens normally do not interfere with parturition. However in other species progestagens are not recommended for use in pregnant animals because of the risk of prolonged gestation, stillbirth or abortion.	In non-human primates progestagens normally do not interfere with parturition. However in other species progestagens are not recommended for use in pregnant animals because of the risk of prolonged gestation, stillbirth or abortion.	Progestagens are not recommended in pregnant animals because of the possibility of prolonged gestation, stillbirth, abortion, etc. in some species, although the effect may depend on dose. Progestagens in late pregnancy seem not to interfere with parturition in primates, but this is a taxon-specific phenomenon.	Not recommended - risk to foetus unknown	Progestagens are not recommended in pregnant animals because of the possibility of prolonged gestation, stillbirth, abortion, etc. in some species, although the effect may depend on dose. Progestagens in late pregnancy seem not to interfere with parturition in primates, but this is a taxon-specific phenomenon.	N/A
Use during lactation:	No contraindications once lactation established	No contraindications once lactation established	Concidered safe for nursing; Does not affect lactation, but etonogestrel is excreted in milk.	Considered safe for nursing infant.	Considered safe for nursing infant.	Not recommended - may interfere with milk production and affect the developing infant. Progestin-only birth control pills can be used instead.	Considered safe for the nursing infant	N/A
Use in prepubertals or juveniles:	Data deficient in this group, see product information sheet	Data deficient in this group, see product information sheet	The use of synthetic progestagens in pre- pubertals or juveniles has not been fully assessed. Possible long-term effects on fertility are not known.	The use of synthetic progestagens in pre- pubertals or juveniles has not been fully assessed. Possible long-term effects on fertility are not known.	The use of synthetic progestagens in pre- pubertals or juveniles has not been fully assessed. Possible long-term effects on fertility are not known.	Not recommended - data deficient and potential long-term effects in fertility	The use of synthetic progestagens in pre-pubertals or juveniles has not been fully assessed. Possible long-term effects on fertility are not known.	N/A
Use in seasonal breeders:	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A

Duration	Duration of efficacy has not been well established as a guide: 4.7 mg implants will suppress for a minimum of 6 months; 9.4mg will be effective for a minimum of 12 months	Not well established, duration of effect being likely related to the dose. Higher doser result in longer duration of effect. This is extremely data deficient	2-3 years in various primates	2-3 years in various primates	Dose dependant: 45-90 days in general. However, effects could last 1-2 years in some individuals.	More than 1 day as effective during the placebo week in human. Duration for other species not fully established.	Not more than one day. Pills need to be administered daily (follow packet insert instructions if one day is missed).	N/A
Reversibilty	Considered reversible but every species has not been tested. duration to reversibility extremely variable. Removal of implant to aid reversibility is recommended.	Considered reversible but every species has not been tested. duration to reversibility extremely variable. Removal of implant to aid reversibility is recommended.	Designed to be fully reversible but individual variations can occurTo increase potential for full reversibility implants must be removed. A reversal has been reported in a female agile gibbon who became pregnant 2 years after removal of the implant.	Designed to be fully reversible but individual variations can occur. To increase potential for full reversibility implants must be removed.	Designed to be fully reversible but individual variations can occur	Reversibility is unknown but presumably would occur after cessation of treatment, although return to cycling can vary per individual. Even in humans, it may take several months (cycles) before normal ovulation returns.	It should be reversible after cessation of treatment, although return to cycling can vary per individual. Even in humans, it may take several months (cycles) before normal ovulation returns.	N/A
Effects on Behaviour	None observed except lack of libido. There are anecdotal reports of change of hierarchy with the behavoirual implications that this may have.	Same as deslorelin	Effects on behaviour have not been studied, every individual may react differently. Because progestagens can suppress ovulation it can be expected that courtship and mating behaviour will be affected in some way. Further research in the subject is necessary.	Effects on behaviour have not been studied, every individual may react differently. Because progestagens can suppress ovulation it can be expected that courtship and mating behaviour will be affected in some way. At high doses can have masculinising effect. Further research in the subject is necessary.	Effects on behaviour have not been studied, every individual may react differently. Because it binds readily to androgen receptors and is antiestrogenic, females may experience male-like qualities (increased aggression , development of male secondary sex characteristics, etc.) Further research in the subject is necessary.	Effects on behaviour have not been studied, every individual may react differently. Further research in the subject is necessary.	Effects on behaviour have not been studied, every individual may react differently. Further research in the subject is necessary.	N/A
Effects on sexual physical characteristics	Some dichromatic species may change colour. Loss of muscle mass may also be seen.	Some dichromatic species may change colour.	There might be some degree of sexual swelling and menstruation might occur. Ovulation may also occur even though pregnancy does not ensue.	There might be some degree of sexual swelling and mensturation might occur. Ovulation may also occur even though pregnancy does not ensue.	See above	Data deficient	Data deficient	N/A
Males	Data deficient	Data deficient see comment for deslorelin	Not Recommended	Not Recommended	Not recommended	N/A	N/A	Reported
Dose	Usually a higher dose than in females are required in males. Data deficient	Usually a higher dose than in females are required in males. Data deficient	N/A	N/A	N/A	N/A	N/A	N/A
Latency to effectiveness:	Depending on the species there may be fertile sperm present in vas deferens for 6-8 weeks post treatment. Testosterone decreases after 3-4 weeks but sperm can stay fertile for many weeks after. Additional contraception needed during this time or separtion of the sexes.	Depending on the species there may be fertile sperm present in vas deferens for 6-8 weeks post treatment. Testosterone decreases after 3-4 weeks but sperm can stay fertile for many weeks after. Additional contraception needed during this time or separtion of the sexes.	N/A	N/A	N/A	N/A	N/A	Depending on species and individual, perhaps as long as 2 months or more
Use in prepubertals or juveniles:	Data deficient in this group, see product information sheet	Data deficient in this group, see product information sheet	N/A	N/A	N/A	N/A	N/A	Data deficient
Use in seasonal breeders:	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
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Effects on Behaviour	Testosterone related aggression is likely to decrease. Data deficient in this group, see product information sheet.	Testosterone related aggression is likely to decrease. Data deficient in this group, see product information sheet.	N/A	N/A	N/A	N/A	N/A	Vasectomy will not affect androgen- dependant behaviours
Effects on sexual physical characteristics	Some dichromatic species may change colour if testosterone ralated. Decrease in body size, decrease testicular size, feminisation of males.	Some dichromatic species may change colour if testosterone ralated. Decrease in body size, feminisation of males.	N/A	N/A	N/A	N/A	N/A	None observed in non-human primates
General:								
Side effects	Similar to gonadectomy; especially weight gain. Some dichromatic species may change colour.	Similar to gonadectomy; especially weight gain. Some dichromatic species may change colour.	Possible weight gain, possible increased or decreased frequency of bleeding during menstruation. EGZAC recommends always reading the manufacturer's data sheet	Possible weight gain, possible increased or decreased frequency of bleeding during menstruation. At high doses can have masculinising effect. EGZAC recommends always reading the manufacturer's data sheet	Progestagens are likely to cause weight gain in all species. Possible deleterious effects on uterine and mammary tissues vary greatly by species; (see taxon sheets). In the human literature, Depo-Provera® has been linked to mood changes. Because it binds readily to androgen receptors and is anti-estrogenic, females may experience male-like qualities (increased aggression, development of male secondary sex characteristics, etc.) EGZAC recommends always reading the manufacturer's data sheet	Weight gain is less likely than with the progestagen only pills. Mood changes might occur.	Progestagens likely cause weight gain in all species. Possible deleterious effects on uterine and mammars tissues vary greatly by species. To date, few studies have shown link between synthetic progestagen treatment and serious health risk in non-human primates.	N/A
Warnings	Causes initial gonadal stimulation; correct administration essential - see product information sheet	Causes initial gonadal stimulation	Interaction with other drugs are known to occur and may influence protection against pregnancy. In some diabetic animals progestagens has led to an increased insulin requirement, it is advised that the product be used with caution in diabetic animals and that urine glucose levels are carefully monitored during the month after dosing. EGZAC recommends always reading the manufacturer's data sheet.	Interaction with other drugs are known to occur and may influence protection against pregnancy. In some diabetic animals progestagens has led to an increased insulin requirement, it is advised that the product be used with caution in diabetic animals and that urine glucose levels are carefully monitored during the month after dosing. EGZAC recommends always reading the manufacturer's data sheet.	Interaction with other drugs are known to occur and may influence protection against pregnancy in some diabetic animals progestagens has led to an increased insulin requirement, it is advised that the product be used with caution in diabetic animals and that urine glucose levels are carefully monitored during the month after dosing. EGZAC recommends always reading the manufacturer's data sheet.	Interaction with other drugs are known to occur and may influence protection against pregnancy. In some diabetic animals progestagens has led to an increased insulin requirement, it is advised that the product be used with caution in diabetic animals and that urine glucose levels are carefully monitored during the month after dosing. EGZAC recommends always reading the manufacturer's data sheet.	Progestagen only contraceptive pills can fail in obese animals. Be aware that progestagen-only pills are not being as effective at suppression oestrus as the combination pills. In some diabetic animals progestagens has led to an increased insulin requirement, it is advised that the product be used with caution in diabetic animals and that urine glucose levels are carefully monitored during the month after dosing.	Infection of the surgical wound might occur. Intradermal closure of the skin is advised together with prophilactic antibiotic treatment and NSAID

Reporting Requirements: In order to increase our knowledge of the efficacy of contraception methods in the Hylobatidae family it is recommended that all individuals on contraception be reported to EGZAC

References:

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Disclaimer: EGZAC endeavours to provide correct and current information on contraception from various sources. As these are prescription only medicines it is the responsibility of the veterinarian to determine the dosage and best treatment for an individual