A quantitative investigation of gonadal feminization by diethylstilboestrol of genetically male embryos of the quail Coturnix coturnix japonica

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The effect of diethylstilboestrol on gonad development in quail embryos has been quantitatively analysed. Quail embryos at 4 days of incubation were treated with diethylstilboestrol (DES), using the egg dipping method. At 10 days of incubation, embryos were removed and killed by decapitation. Tissues were prepared for chromosome analysis, and the parts of the abdomen containing the gonads were prepared for serial sectioning and quantitative assessment. Left gonads of DES-treated male embryos resembled ovaries histologically, while their right gonads were markedly reduced in size. Right gonads of DES-treated female embryos were also further reduced by treatment with DES. There was no statistically significant effect by DES treatment on the size of left gonads, although the ratio of left compared with right gonadal volumes was highly significant. Since, in birds, the left embryonic gonad has bisexual potential, while the potential of the right gonad is exclusively masculine, these results exemplify the adverse effect exerted by oestrogen on male sexual development in vertebrates.

Introduction

Studies on the genetics of sex differentiation in birds lag behind those of mammals and of reptiles. Yet the avian embryo offers advantages in both accessibility and availability and is thus a potentially useful candidate for studying the effects of exogenous substances on the development of the gonads.

A characteristic feature of sexual development in birds is the asymmetry of their gonads. In most species, ovaries are confined to the left side, while the right gonad of females remains rudimentary, but, in certain circumstances, has the potential to develop into a testicular structure (Lillie, 1952). Moreover, in the early embryo, the left gonad in both sexes has the potential to develop ovarian tissue, while the only potential of the right gonad is for testicular development. Evidently, the chromosomal sex-determining mechanism is more open to environmental modification than is that of mammals.

Wolff and Ginlinger (1935) reported that injecting diethylstilboestrol (DES) into incubating chick eggs produces a feminization of male chicks. A simplified method is described by Seltzer (1956; cited by Pincus, 1958), which, in place of injection, substitutes dipping the eggs into the solutions to be tested. Dipping quail (Coturnix coturnix) eggs that have been incubated for 4 days into an alcoholic solution of 40 mg per 100 ml ethanolic DES achieves a modification of male gonads (Haffen, 1965). Instead of appearing symmetrical, the left gonad of treated embryos assumes the appearance of an ovary, while the right gonad is reduced. The feminization of quail embryos appears to be more pronounced than in chicks; whereas feminized chicks revert to male sex characteristics soon after hatching (Wolff and Haffen, 1961), sex reversal in quails seems more stable. Haffen (1965) reports that an 8-week-old male had an ovotestis containing follicles.

Here, we report a detailed quantitative investigation of DES-treated and control gonads in 10-day-old quail embryos, the genetic sex of which has been verified by chromosome analysis. As in other birds, the sex chromosomes of males consist of two equal Z chromosomes, while females have one Z chromosome and a smaller W chromosome.

Materials and Methods

Fertilized quail eggs were obtained from a breeding colony maintained by J. Bee at the Royal Veterinary College, London. Each batch of eggs was divided into three groups and placed in randomized positions in an incubator kept at 37.5°C. After 4 days of incubation, one group of eggs was dipped into a solution of 40 mg DES in 100 ml of 95% (v/v) ethanol, while the second group was dipped into 95% ethanol, and the third group was left untreated. Dipping was carried out by placing each egg, with its pointed end downwards, into a net and immersing it along two-thirds of its length for 10 s. The eggs were then returned to the incubator and left for another 6 days. At the end of this period, the embryos were removed, and their crown-rump lengths were measured, using digital calipers. The heads of the embryos were set up for chromosome analysis, using the method of Tuinen and Valentine (1983), while the parts of the bodies containing the gonads were prepared...
for serial transverse sectioning at 7 µm and staining in haematoxylin and eosin.

Volumes of left and right gonads were computed from section areas as described by Baker et al. (1993), using a Leitz camera lucida and a digitizing tablet connected to an Opus computer. Areas of between 30 and 40 sections spanning the entire gonad were measured and the sum of the areas was converted into volumes by multiplying by the effective section thickness (for example, if every eighth section was measured, the effective section area would be 56 µm²); this method is unaffected by the angle of sectioning. The results obtained for DES-treated gonads and the two types of controls were compared by Student’s t test.

Results

Mean crown–rump lengths varied between 18.1 and 19.8 mm in different groups but the differences were not significant; these data are not shown.

Male embryos

The histological appearance of DES-treated and control gonads in genetically male embryos is illustrated in Fig. 1.

Whereas in untreated and ethanol-treated embryos, left and right gonads appeared to be symmetrical (Fig. 1a), after DES treatment, the right gonad was markedly reduced in size, while the comparatively much larger left gonad assumed the general appearance of an ovary (Fig. 1b). It had a more rugose outline than did the control gonads, and signs of an incipient ovarian cortex could be made out along its lower edge.

The difference in size between left and right DES-treated gonads was highly significant, whereas there were no significant differences between left and right untreated or ethanol-treated gonads (Table 1). Neither treatment exerted a significant effect on the size of the left gonad, and there was no significant difference in size between untreated and ethanol-treated right gonads. By contrast, treatment with DES caused a highly significant reduction in the size of the right gonad.

Female embryos

All genetically female embryos had a left ovary and a reduced right gonad, and there was no obvious effect produced by any of the treatments (Table 2). The difference in size between left and right gonads is highly significant in all groups, and the effect of DES in further reducing the size of the right gonad is also significant.

Comparison between male and female embryos

Figure 2 illustrates the difference in gonadal volume obtained when the mean volumes of female gonads are subtracted from those of male gonads in any one group. In untreated and ethanol-treated embryos, the results are negative for left gonads and positive for right gonads, but this effect is not evident in DES-treated embryos. In control embryos, females have a larger left gonad and a smaller right gonad than do males. This relationship is no longer seen in embryos treated with DES, in which neither difference is significantly different from zero.

Discussion

Our results confirm the reports by Haffen (1965) and Scheib et al. (1981) that the dipping of quail eggs into an alcoholic solution of DES results in the partial feminization of embryos and hatched birds. We have shown that 10-day-old male embryos exposed to DES show a marked reduction in the size of the right gonad, and that female embryos show a somewhat smaller, but still significant, reduction in the size of the right gonad. By contrast, the hormone did not reduce the size of the left gonad, the morphology of which was similar in some ways to an ovary. This gonad was a little larger than in the controls, but the difference was not significant. In female embryos treated with DES, volumes of both gonads were reduced, but the difference was significant only for the right gonad. The DES-induced changes in the growth patterns of the four types of gonad abolished the normal relationship seen in the controls, in which the left ovary is larger than the testis, and the right rudimentary gonad of the female is very much smaller than a testis.
The difference in gonadal volumes between untreated and ethanol-treated embryos was not significant in any of the four types of gonad. However, ethanol-treated gonads were reduced in size in all four groups, suggesting that the alcohol may have a slight inhibitory effect on gonadal development; this possibility needs to be tested with a larger number of samples. The only increase in gonadal volume, albeit statistically not significant, was found in the left gonads of DES-treated male embryos, which became feminized as a result of the treatment.

In young chick embryos it can be seen clearly in both sexes that the left gonad has an incipient ovarian cortex and is thus potentially hermaphroditic. The cortex normally regresses in male embryos and is altogether absent in the right gonad of either sex, the only potential of which is to develop into a testis (Mittwoch, 1973, 1986). The feminization by DES of the left gonad of male embryos illustrates its fundamentally hermaphroditic nature. The marked reduction of the right gonad of male embryos exemplifies the deleterious nature of oestrogen on the developing testis and, to a lesser extent, this is also seen in the reduction in size of the right female gonad, which has the potential to develop into a testis.

Treating pregnant rats and mice with oestrogen is known to give rise to cryptorchidism (Hadžiselimović et al., 1980) and there is evidence in human males that the administration of oestrogen to mothers during pregnancy also results in an increased incidence of cryptorchidism and inguinal hernia (Depue, 1984). A rise in exogenous oestrogen acting on the male fetus has been suggested as a contributory factor for a possible secular fall in the number of spermatozoa and rises in disorders of the testis and the male reproductive tract, including testicular cancer, hypospadias and cryptorchidism (Giwersman et al., 1993). Since oestrogens are normally present in the uterine environment during pregnancy, the fast development of the male embryo and the early secretion of testosterone by the fetal testis, before high oestrogen concentrations are established, may be a necessary adaptation in the development of eutherian males (Mittwoch, 1993).

Failure of male sex differentiation in the presence of oestrogen appears to be widespread among vertebrates. In the red-eared slider, *Trachemys scripta*, a reptile with temperature-dependent sex determination, Wibbels et al. (1993) reported that eggs incubated at the male-producing temperature of 26°C and treated with oestradiol develop as phenotypic females. In another turtle, *Caretta caretta*, with a similar temperature-dependent system of sex determination, Harry and Williams (1991) reported different growth patterns for the male and female urogenital system during the sex-determining period.

Quantitative studies of the effect of oestrogen on the developing urogenital system of the quail offer considerable...
Fig. 2. Differences in size obtained after subtracting mean volumes of female gonads from those of male gonads in control and diethylstilboestrol (DES)-treated quail embryos (means ± SEM). L: left gonad; R: right gonad. *0.02 < P < 0.05; **0.001 < P < 0.01; ***P < 0.001.

potential for a better understanding of the different mechanisms operating in the process of sex determination in vertebrates (Mittwoch and Burgess, 1991). An important theoretical question that needs to be addressed is whether the asymmetric response to DES by the two gonads is caused by an initial difference in the presence of receptors, and whether the primary effect of oestrogen is the ovarian transformation of the left gonad or the inhibition of the right gonad.

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