Luteinized Ovarian Cysts in Mediterranean Striped Dolphins

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ABSTRACT: The morbillivirus epizootic during 1990 to 1992 in Mediterranean striped dolphins (Stenella coeruleoalba) off the Mediterranean coast of Spain diminished these populations directly through mortalities, and indirectly through loss of normal fecundity. High levels of polychlorinated biphenyls (PCB's) also were detected in stranded animals. In addition to high numbers of abortions during the epidemic, unusual cystic structures were noted in the ovaries of several morbillivirus-infected dolphins with high PCB levels. These structures were identified as multiple luteinized cysts from their gross and histomorphologic characteristics. No morbillivirus antigens were detected in the lesions by immunohistochemistry. Because luteinized cysts occur when ovulation is impeded, either an effect of morbillivirus or PCB's on hypothalamic/pituitary function or an effect of PCB's on ovarian responsiveness are proposed as pathogenic mechanisms. These cysts may impede population recovery from the epidemic if similar cysts occurred on surviving dolphins.

Key words: Luteinized cyst, Mediterranean striped dolphin, morbillivirus, ovary, polychlorinated biphenyl, Stenella coeruleoalba.

The morbillivirus epidemic of 1990 to 1992 resulted in the stranding and death of over 1,000 Mediterranean striped dolphins (Stenella coeruleoalba) off the Spanish Mediterranean coast (Aguilar and Raga, 1993; Calzada et al., 1994). Most of the stranded dolphins also had high tissue concentrations of polychlorinated biphenyls (PCB), and the role of these environmental contaminants on the health of the dolphins remains unclear (Aguilar and Borrell, 1994). The epidemic occurred during the height of the breeding season (September and October), and the impact on reproductive functions has not been fully assessed. During the epidemic, an abnormally high percentage of females had young corpora albicans and uterine distention suggestive of recent abortion, and characteristically high numbers of dead fetuses were found (Calzada et al., 1996). Ovaries from several other females examined during necropsy had unusual cystic structures not previously observed in the ovaries of any healthy related Stenella spp. The purpose of this study was to characterize these ovarian structures and determine if these lesions were caused by morbillivirus infection of the ovary.

Ovaries of 56 sexually-mature female striped dolphins were collected in summer of 1990 and 1992. All dolphins in the study group had stranded during the epidemic and had high PCB levels (Aguilar and Borrell, 1994). The ages of dolphins were determined by counting growth layer groups in the dentine of decalcified and longitudinally-sectioned teeth. Each growth layer group was assumed to represent 1 yr (Calzada et al., 1994). Sexual maturity was determined by identifying corpora lutea and corpora albicans in serial sectioned fixed ovaries. From the population of 56 sexually mature females, the ovaries of four females had unusual cystic structures that were examined by histopathology. Dolphins with cysts were 14-, 14-, 18- and 19-yr-old. They were confirmed to have morbillivirus infection (Domingo et al., 1992), and were not pregnant. The stranding sites of these four dolphins were Castelldefels (41.17°N, 1.58°E), Salou (41.04°N, 1.08°E), Tarragona (41.07°N, 1.15°E) and Cartagena (37.36°N, 0.59°W). The ovaries of two healthy pregnant females also were included as examples of normal pregnancy corpora lutea.

For histopathology, formalin-fixed ovaries were deparaffinized in xylene, embedded in paraffin, sectioned at 7 µm, and stained with hematoxylin and eosin, Mas-
son's trichrome, and periodic acid Schiff's reagent methods (Luna, 1992). The presence of morbillivirus antigen in tissues was assessed by immunohistochemistry using a previously published method (Kennedy et al., 1991), and a mouse monoclonal antibody (clone 1.3) against glycosylated hemagglutinin protein of phocine distemper virus as the primary antibody (Trudgett et al., 1991).

The unusual ovarian structures were large thick-walled cysts that ranged from 16 to 23 mm in diameter (Fig. 1). The 23 mm diameter cyst had an 18 mm diameter central cavity, in contrast to the corpora lutea from normal pregnant dolphins that were solid structures 24 to 26 mm in diameter. One dolphin had a single 23 mm diameter cyst, two dolphins had two cysts and one dolphin had four cysts. The cysts were located in the cortical parenchyma and were covered by ovarian stroma without an ovulation stigma. The cyst lumina were round, lined with mottled gray-yellow soft tissue, and filled with grey gelatious material. Histologically, the cysts were lined by a thick, disorganized, multicellular layer composed of elongate granulosa/theca cells with minimal luteinization (Fig. 2). The luteinized cells were arranged haphazardly and were divided into irregular lobules by dense fibrous septa with minimal vascularization. The cyst lumina were filled with fibrillar proteinaceous fluid surrounded by a thick rim of mature fibrous connective tissue. In comparison, corpora lutea of normal dolphins had plump polygonal luteal cells with prominent intercellular vascularization surrounding a core of solid fibrous tissue. Ovaries with the cysts also had developing follicles, multiple corpora albica and no normal corpora lutea. Morbillivirus antigen was not detected in any ovary from the females with cysts.

The multiple cystic structures on the ovaries of the stranded dolphins were most compatible with luteinized follicular cysts (McEntee, 1990; Kennedy and Miller, 1993). The absence of an ovulation stigma and presence of more than one struc-

**FIGURE 1.** Ovary from a striped dolphin with two luteinized cysts. Bar = 3 mm.

**FIGURE 2.** Histologic appearance of a luteinized cyst in the ovary of a striped dolphin. The cyst is lined by spindle-shaped granulosa/thecal cells with minimal luteinization that are separated from the lumen (L) by a mature fibrous band (arrow). Masson's trichrome stain. Bar = 30 μm.
ture in three of four females from a species that is monotocous (Marshall, 1984) indicates that these structures were abnormal. Morphologic characteristics that suggest they are luteinized cysts and not cystic corpora lutea (CL) were the large expansive central cavity with a thick mature fibrous lining and the small, angular shape of the luteal cells. (Kennedy and Miller, 1993). The lack of concordance between the immaturity of the spindle-shaped luteal cells and maturity of the fibrous tissue lining of the central cavity indicated that these structures were not developing CLs. These cysts did not have features of cystic CLs which are characterized by a larger overall size, an irregular central cavity containing organizing loose fibrous tissue, and plump luteal cells (Van Lennep, 1950; Harrison and McBrearty, 1977; Kennedy, 1993). Although the dolphin cysts were similar to accessory corpora lutea (CL) that rarely have been observed in the ovaries of pregnant Odontocetes (Brodie, 1972) and Delphinids (Harrison and McBrearty, 1977; Marsh and Kasuya, 1984), none of the affected dolphins were pregnant.

In monotocous cetacean species, a single Graafian follicle ovulates resulting in a corpus luteum that may have a central cavity which usually becomes obliterated as it matures (Harrison and McBrearty, 1977; Harrison, 1981). In land mammals, if one or more Graafian follicles fail to ovulate, follicular cysts develop and luteinization of granulosa or theca cells can ensue. Ovulation failure in land mammals is thought to be due to inadequate levels of luteinizing hormone (LH) or inappropriate timing of LH secretion from the pituitary (Youngquist, 1986).

The pathogenesis of the luteinized cysts in the dolphins also may be due to inadequate or untimely release of LH or lack of LH responsiveness in the follicle. LH release may have been affected by morbillivirus infection of the central nervous system, PCB toxicity, stranding stress, or poor nutritional status. All dolphins with cysts were infected with morbillivirus, including viral infection of the central nervous system (Domingo et al., 1992; Duignan et al., 1992). The absence of viral antigens in the ovary of affected dolphins does not exclude the possibility that morbillivirus indirectly affected ovulation through infection of the hypothalamus or pituitary, thereby impeding gonadotrophin-releasing hormone (GnRH) or LH synthesis or release.

Another possible cause of the cysts in these dolphins was high systemic PCB levels (Aguilar and Borrell, 1994). Polychlorinated biphenyls and related xenobiotics act as estrogen agonists/antagonists (Gorospe and Reinhard, 1995) that could impair follicular development and positive feedback of estrogens on pituitary LH release (Mahony and Hodgen, 1995). Polychlorinated biphenyls affect fecundity in other species, including marine mammals (Reijnders, 1980; Golub et al., 1991; Gorospe and Reinhard, 1995), and reduced fertility has been proven experimentally in monkeys (Arnold et al., 1995), rodents (Marks et al., 1989; Golub et al., 1991), and rabbits (Lindenauf et al., 1994). Unique isomers of PCBs accumulate in the follicular fluid of the ovary, resulting in reduced ovulation in rabbits (Lindenauf et al., 1994). In teleosts fed PCBs, gonadal hormone levels were reduced and pituitary secretion was diminished (Thomas, 1990). Polychlorinated biphenyls therefore could contribute to the formation of luteinized follicles by altering follicular responsiveness directly or reducing pituitary secretion of LH. Polychlorinated biphenyls have been linked with declining reproductive function in seals at 701 µg/g fat (range 87 to 1,447 µg/g fat) (Reijnders, 1980), and mean levels (±SD) of PCBs were even greater in dolphins with luteinized cysts (1,081.3 ± 684.4 µg/g liver; 1,017.3 ± 665.0 µg/g fat) than in the population as a whole (median, 778 µg/g lipid basis) (Aguilar and Borrell, 1994). The poor nutritional status of the infected dolphins also could potentially in-
terfere with normal pituitary LH release, leading to luteinized cyst formation. Poor nutritional status compromises normal hypothalamic and pituitary function (Dubey et al., 1986), and the morbillivirus-infected dolphins had lipid reserves that were estimated at 60% below normal (Aguilar and Raga, 1993). Stress from stranding also may have contributed to abnormal reproductive function, because experimentally-induced stress results in increased endogenous corticosteroids and decreased LH release in other species (Moberg, 1985). Luteinized and unluteinized follicular cysts have been reported on ovaries from other stranded and captured delphinids (Harrison and McBrearty, 1977; Harrison, 1981), supporting a possible role of stress.

The final outcome of the reproductive disorders that occurred in the Mediterranean dolphin population during the morbillivirus stranding has yet to be assessed. While these ovarian cysts are of no life-threatening consequence to individual dolphins, their significance to the population may be more portentous. Mediterranean striped dolphins are seasonal breeders with most breeding occurring during September and October (Forcada et al., 1994), precisely when the morbillivirus epidemic occurred (Forcada et al., 1994; Calzada et al., 1996). High numbers of abortions also were suspected (Calzada et al., 1996), and school size and structure which determine reproductive behavior were markedly altered (Forcada et al., 1994). If luteal cysts also occurred in surviving females, these females would be unlikely to produce offspring until the next breeding season, augmenting losses in the population. If morbillivirus caused these reproductive changes, population recovery might occur during the subsequent period of population immunity (Appel, 1987) if reproductive functions are restored. However, if PCB toxicity is the basis for the reproductive disorders observed during the Mediterranean dolphin strandings, continued reproductive problems are likely, threatening long-term population recovery.

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