Unexpected Perinatal Loss versus Sids - a Common Neuropathologic Entity

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Abstract: Objective: To evaluate the involvement of alterations of the central autonomic nervous system, particularly of the brainstem and cerebellum, in a wide set of victims of sudden and unexplained perinatal and infant death.

Material and Methods: The study population consisted of 63 stillbirths, 28 neonatal deaths and 140 suspected SIDS. The victims were subjected to in-depth anatomopathological examination following appropriate guidelines. The protocol included, in particular, the histological evaluation on serial sections of the cardiorespiratory autonomic nervous system.

Results: A diagnosis of “unexplained death” was established for 217 of the 231 victims (59 stillbirths, 28 newborns and 130 SIDS). In a very high percentage of these deaths (84%) we observed one or more anomalies of the nuclei and/or structures of the brainstem and cerebellum related to vital functions.

Conclusion: Unexpected perinatal loss should not be regarded as a separate entity from SIDS, given the common neuropathological substrates.

Keywords: Stillbirth, SIDS, Central Autonomic Nervous System, Brainstem, Neuropathology, Developmental Alterations.

INTRODUCTION

Sudden perinatal death (that includes “ante-partum death”, from 22 completed gestational weeks, “intra-partum death” and “early neonatal death” occurring during the first seven completed days of life) and sudden infant death syndrome (SIDS) represent facets of a multifactorial problem that has not yet found a univocal approach on the clinical plane [1-6]. Despite the efforts of many investigators, the true causes of unexpected stillbirth and SIDS remain unknown. This is due to a lack of appropriately detailed post-mortem studies.

A fundamental component of these deaths is neuropathology. In fact, pathogenesis of unexplained perinatal death and SIDS seems to privilege, in most cases, some nervous dysfunctions. Nevertheless, only a few studies in this field have adequately examined the neurological substrates, although even subtle abnormalities of structures of the autonomic nervous system can determine dysfunctions in the control of vital functions, leading to unexpected perinatal loss as well as infant death [7-10].

The present study will focus upon the multifaceted involvement of the central autonomic nervous system, particularly of the brainstem and cerebellum, in the disruption of neuronal pathways observed in a large population of sudden and unexplained perinatal and infant death victims.

MATERIAL AND METHODS

Our study was focused on 231 victims of sudden and unexplained perinatal and infant death, aged from 22 gestational weeks to 10 postnatal months. This was a selected set of cases, sent to our Center over a 6-year period (2000-2006) on the basis of a specific decree passed by the Lombardy Region [11]. This decree specifies that all cases of unexplained perinatal death and/or of suspected SIDS must be sent to the L. Rossi Center of Milan University, being the Regional Reference Center, for in-depth anatomopathological examination, in accordance with specific guidelines [12-14]. This protocol includes, in particular, the histological evaluation on serial sections of the cardiorespiratory autonomic nervous system, besides the examination of the placental disk, umbilical cord and membranes in fetuses.

For every case, a complete clinical history, particularly referred to the mother in perinatal deaths, and including the death scene examination in infant deaths, was collected. Cases were number-coded and identifying data (name, address, etc.) were removed prior to analysis.

The study population consisted of 63 stillbirths, 28 neonatal deaths and 140 suspected SIDS.

The 63 fetuses, 36 males and 27 females, were aged from 22 to 41 gestational weeks, with a peak incidence at 36 to 38 weeks (62%). The weights of victims ranged from 352 to 3500 grams, being below the corresponding reference value in most cases.

The 28 newborns, 16 males and 12 females, died intra-partum or in the first week after birth.
Among the infant victims, 52 females and 88 males, sudden death occurred between 1 and 10 months of life, with the greatest incidence around the third month.

After an in-depth autopsy and histological examination, a diagnosis of “unexplained death” was established for 217 of the 231 victims, namely 59 stillbirths, 28 newborns and 130 infants, all clearly SIDS victims.

The remaining 14 cases (4 fetal and 10 infant deaths) (6%), were classified as “border-line”, since it was difficult to establish whether the pathological findings were sufficiently severe to have caused the death. In the infant deaths we observed: neoplastic lesions of the cardiac conduction system, i.e. Purkinje cell tumor and cardiac fibroma in 3 cases; encephalitic features (probably viral in etiology) in nuclei of the brainstem and moderate myocarditis in 4 cases; a case with focal necrosis of the tractus solitarii nucleus, a case with capillary hemangioendothelioma in the area postrema and a case with pneumonia associated to hypoglossal nucleus hypoplasia. In four stillbirths, abnormalities of the annexa were diagnosed (umbilical cord around the neck, head band of the umbilical cord, multifocal placental infarcts). Table 1 summarizes the case profiles of the study.

None of the mothers had any significant pathology. With reference to smoking habit, 81 mothers (35%) declared that they were active smokers (all of more than 3 cigarettes/day and all already before becoming pregnant) while 129 (56%) were non smokers. In 21 cases no information about smoking habit was available.

The focus of this study is the in-depth histological examination of the central autonomic nervous system. Hereafter, we will briefly describe the study protocol. More detailed information is available on the previous publications [12-14].

The brainstem and cerebellum, where the main vital centers are located, were processed and embedded in paraffin. Transverse serial sections were made at intervals of 30 µm. For each level, twelve 5 µm sections were obtained, two of which were routinely stained for histological examination using alternately hematoxylin-eosin and Klüver-Barrera stains. Two additional sections at each level were subjected to immunohistochemistry for the study of neurotransmitters (somatostatin and tyrosine hydroxylase (TH), indicative of catecholamine synthesis) and apoptosis. The remaining sections were saved and stained as deemed necessary for further investigations.

The main nuclei were analyzed in serial sections of the brainstem, namely the parafacial nucleus, the locus coeruleus and the parabrachial/Kölliker-Fuse complex in the pons and mesencephalon; the hypoglossus, the dorsal motor vagal, the tractus solitarius, the ambiguous, the pre-Bötzinger, the inferior olivary and the arcuate nuclei in the medulla oblongata. Fig. (1) shows the sampling of the brainstem (three specimens are obtained: the first, ponto-mesencephalic, includes the upper third of the pons and the adjacent portion of mesencephalon; the second extends from the rostral limit of the medulla oblongata to the adjacent caudal pons, and the third specimen takes as reference point, the obex and extends 2-3 mm above it and below it). Fig. (2) shows the histological sections corresponding to the three brainstem samples of Fig. (1) indicating the main nuclei and structures to be examined.

Table 1. Case Profiles of the Study

| Victims    | Age (range) | Sex (n.) | Unexplained deaths (n.217) | Border line deaths (n.14) |
|------------|-------------|----------|--------------------------|-------------------------|
| Fetuses (n.63) | 22-41 gw    | 36       | 27 | unexplained stillbirth (n.59) | Annexa anomalies (n.4) |
| Newborns (n.28) | 1-7 pd      | 16       | 12 | unexplained neonatal death (n.28) | _ |
| Infants (n.140) | 1-10 m     | 88       | 52 | SIDS (n.130) | Neoplastic lesions of the cardiac conduction system (n.3) Brainstem encephalitis + moderate myocarditis (n.4) Tractus solitarii nucleus necrosis (n.1) Area postrema hemangioendothelioma (n.1) Pneumonia + hypoglossal nucleus hypoplasia (n.1) |

gw = gestational week; pd = postnatal day; m = month; SIDS = Sudden Infant Death Syndrome.

Fig. (1). Sampling of the brainstem: ventral (left) and dorsal (right) surface.

In the cerebellum, the cortex layers (external granular layer, molecular layer, Purkinje cell layer and internal granular layer) and the medullary deep nuclei (the dentate nucleus, the fastigial nucleus, the globose nucleus and the emboliform nucleus) were examined.
Immunohistochemical Methods

In order to analyze the immuno-expression of neurotransmitters, particularly of somatostatin and catecholamines, we used specific primary antibodies after applying the avidin-biotin-peroxidase technique, in conformity with conventional immunohistochemical procedures. Immunohistochemical visualization of apoptotic cells was obtained by the TUNEL method (TdT-mediated dUTP-biotin nick end labeling). A detailed description of the immunohistochemical procedures that we applied, including the immunopositivity evaluation, is available in our previous works [15-17].

Statistical Analysis

The statistical significance of direct comparison between groups of victims (unexplained fetal, neonatal and infant deaths, border-line cases) was determined using one way analysis of variance (ANOVA) and Student’s paired t-test. The selected threshold level for statistical significance was p<0.05.

RESULTS

The in-depth anatomo-pathological examination of the central autonomic nervous system performed in the 217 victims (59 stillbirths, 28 newborns and 130 SIDS) of unexplained perinatal and infant deaths, revealed one or more anomalies of the nuclei and/or structures of the brainstem and cerebellum related to vital functions, mainly to cardio-respiratory control, in 182 cases (84%).

Main Morphological Findings in Brainstem and Cerebellum

Medulla Oblongata

Hypoplasia of the arcuate nucleus emerged in 56% of the sudden fetal deaths (33 cases) and in 50% of both unexplained neonatal deaths (14 cases) and in SIDS victims (65 cases). Different degrees of hypodevelopment were observed, ranging from a normal structure with neuronal immaturity to total agenesis of the nucleus, observed in 8 stillbirths (Fig. 3).

Fig. (3). Transversal section of medulla oblongata with severe hypoplasia of the arcuate nucleus (magnification: 10x)
Abbreviations: ARcn, arcuate nucleus; POn, principal inferior olivary nucleus; PYR, pyramid.

Hypoplasia of the pre-Bötzinger nucleus, with a decreased number of neuronal bodies and/or dendritic fibers, was diagnosed in 25% of stillbirths (15 cases) and in 8% of SIDS (10 cases).

Pons-Mesencephalon

In transverse sections of rostral pons and caudal mesencephalon of 14 unexplained stillbirths (24%) and of 8 unex-
plained neonatal deaths (30%), prevalently intra-partial deaths, a very few immature neurons were observed in both the medial and lateral parabrachial nuclei and in the Kölliker-Fuse area; altogether, these findings led to a diagnosis of hypoplasia of the parabrachial-Kölliker-Fuse complex.

Hypoplasia of the parafacial nucleus, with a decreased neuronal density and transverse area, was diagnosed in the caudal pons of 41 stillbirths (70% of unexplained fetal deaths) but in no case of post-natal death.

**Cerebellum**

Histological examination revealed morphological alterations of the cortex development. In fact, the cerebellar cortex showed an immature structure, uniformly made up of small round cells without the usual four-layered shape, in 10% of the unexplained stillbirths (6 cases, all after the 36th gestational week), in 6 neonatal deaths (20%) and in 26% of the SIDS victims (34 cases) (Fig. 4).

![Fig. (4). Transversal section of cerebellar cortex with immature structure, uniformly made up of small round cells without the usual four-layered shape in a case of sudden neonatal death (magnification: 20x).](image)

Table 2 shows the morphological alterations of the central autonomic nervous system observed in all the unexplained sudden death victims.

**Main Functional Findings in Brainstem and Cerebellum**

Using specific immunohistochemical methods, altered expression of somatostatin, a neurotransmitter that is widely present in brainstem nuclei during fetal life but scarcely expressed after birth, was highlighted in the hypoglossus nucleus. Despite a normal structure, this nucleus was in fact somatostatin-immunonegative in 27% of the unexplained fetal deaths (17 cases) and, on the contrary, immunopositive in 44% of the SIDS victims (57 cases).

Negative expression of tyrosine-hydroxylase (TH), an essential enzyme in catecholamine biosynthesis, was detectable in a well-structured locus coeruleus (the major brainstem producer of noradrenaline) in 57% of the perinatal and infant sudden deaths (124 of the 217 cases).

We observed unusual apoptosis of the Purkinje cells and of the internal granular layer in the cerebellar cortex of 23 SIDS victims (18%) and high apoptotic index in the dentate nucleus of 44 SIDS cases (34%). A more in-depth description of these results can be found in our previous immunohistochemical works [15-17].

Frequently, two or more morphological and/or physiological alterations were observable in the same case, particularly in fetal deaths. In seven victims, we found the simultaneous presence of hypoplasia of different nuclei (the arcuate, pre-Bötzinger and parafacial nuclei) and of negative somatostatin expression in the hypoglossus nucleus.

None of the above reported alterations was found in the border-line population.

**DISCUSSION**

This study summarizes the results of our in-depth neuropathological researches performed in a 6-year period on a very wide, selected set of unexpected and unexplained perinatal deaths and SIDS, investigated following specific guidelines in accordance with the decree of the Lombardy Region [11]. The results hereafter indicated are, in part, reported in previous works [15-24].

The accurate examination of the central autonomic nervous system, allowed us to identify even minute alterations of the nuclei and/or structures overseeing the vital functions in these pathologies.

In particular, in 84% of all the unexplained deaths, we highlighted structural and/or functional alterations, particu-

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**Table 2. Morphological Alterations of the Brainstem and Cerebellum in Unexpected Death Victims**

| Unexpected death victims | Arcuate nucleus hypoplasia | Pre-Bötzinger nucleus hypoplasia | Parabrachial Kölliker-Fuse complex hypoplasia | Parafacial nucleus hypoplasia | Cerebellar cortex immature structure |
|--------------------------|---------------------------|---------------------------------|---------------------------------------------|-------------------------------|-----------------------------------|
| stillbirths n/tot(%)      | 33/59 (56)                | 15/59 (25) *                    | 14/59 (24) *                                | 41/59 (70) *                  | 6/59 (10)                        |
| newborns n/tot(%)         | 14/28 (50)                | 0 *                             | 8/28 (29) *                                 | 0 *                          | 6/28 (20)                        |
| SIDS n/tot(%)             | 65/130 (50)               | 10/130 (8) *                    | 0 * *                                       | 0 *                          | 34/130 (26)                      |
| statistics                | Not significant           | # SB>NB p=0.0030                | * SB>SIDS p=0.0008                           | * SB>SIDS p<0.0001            | # SB>NB p<0.0001                 |

SIDS=sudden infant death syndrome.

One way ANOVA and Student’s paired t-test were performed to compare the incidence of hypoplasia in the different districts of central autonomic nervous system.
larly of the brainstem and cerebellum. These alterations represent a common morphological substrate in both unexplained perinatal death and SIDS and, therefore, can be considered congenital in nature.

A common finding was hypoplasia of the arcuate nucleus, a chemoreceptor component of the ventral surface of the medulla oblongata, which was present in over 50% of all the victims, albeit with different degrees of extension and severity. In perinatal deaths, in addition to hypoplasia of the arcuate nucleus, we observed hypodevelopment of the dendritic fibers with a decreased neuronal density in the pre-Bötzinger area, besides hypoplasia of the parabrachial/Kölliker-Fuse complex. The pre-Bötzinger nucleus has been physiologically defined as a group of neurons of the ventrolateral medulla, that are essential for generating the respiratory rhythm, as well as for modulating eupneic breathing [24,25]. Instead, the Kölliker-Fuse nucleus has an important function during intrauterine life, inhibiting the response of the respiratory center, able to coordinate the pulmonary motor response to a variety of mechanisms and, above all, to maternal cigarette smoking in pregnancy, as we have previously suggested [16,28,29].

In conclusion, the neuropathologic substrates common to both fetal and neonatal deaths can be ascribed to a process of unnatural selection caused by exogenous pathogenic noxae. The effects of these are particularly important, in view of their frequency and the variety of alterations observed, during fetal life. Therefore, unexpected perinatal loss should not be regarded as a separate entity from SIDS, since both are an integral part of autonomic nervous system abnormalities.

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