Gasping and autoresuscitation in the developing rat: effect of antecedent intermittent hypoxia

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Gozal, David, Evelyne Gozal, Stephen R. Reeves, and Andrew J. Lipton. Gasping and autoresuscitation in the developing rat: effect of antecedent intermittent hypoxia. J Appl Physiol 92: 1141–1144, 2002. First published November 2, 2001; 10.1152/japplphysiol.00972.2001.—Gasping is a critically important mechanism for autoresuscitation and survival during extreme tissue hypoxia. Evidence of antecedent hypoxia in sudden infant death syndrome suggests that intermittently occurring hypoxic episodes may modify gasping and autoresuscitation. To examine this issue, an intermittent hypoxia (IH) profile consisting of alternating room air and 10% O2-balance N2 every 90 s was applied to pregnant Sprague-Dawley rats (IHRA; n = 50) and to pups after a normal pregnancy (RAIH; n = 50) as well as to control pups (RARA; n = 50). At postnatal day 5, pups were exposed to 95% N2-5% CO2, and gasping and the ability to autoresuscitate were assessed. Compared with RARA, IHRA- and RAIH-exposed pups had a reduced number of gasps, decreased overall gasp duration, and were less likely to autoresuscitate on introduction of room air to the breathing mixture during the last phase of gasping (P < 0.001 vs. RARA). We conclude that both prenatal and early postnatal IH adversely affect gasping and related survival mechanisms.

anoxic tolerance; respiration; sudden infant death syndrome; apnea

IN THE LAST DECADE, THE INCIDENCE of sudden infant death syndrome (SIDS) has decreased primarily as a result of public campaigns aiming to reduce prone sleeping position in babies (27). However, SIDS still remains the major cause of death in apparently healthy infants. Although the precise mechanisms underlying SIDS remain undefined, one of the leading hypotheses posits that recurrent hypoxemia may precede the fatal event and affect gasping and autoresuscitation mechanisms (15, 16, 22).

When exposed to asphyxic conditions, most mammalian species will develop a transient hyperpnea, then apnea, and, within a variable period of time (gassing latency), gasping respiratory activity will emerge and continue until terminal apnea and death (2). Our laboratory has previously shown that, in developing Sprague-Dawley rats, a triphasic gasping response occurs in response to anoxia and that such response becomes monophasic at ~20 days postnataally (14). Using a similar approach, Fewell et al. (4, 5) showed that prenatal exposure to nicotine modifies anoxia-induced gasping in an age-dependent fashion and that autoresuscitation is also adversely affected. However, whether intermittent fetal or postnatal hypoxia modifies gasping activity in the young postnatal rat and has detrimental effects on its ability to autoresuscitate.

METHODS

Time-pregnant Sprague-Dawley rats were purchased from Charles River and used for all experiments. The experimental protocols were approved by the Institutional Animal Use and Care Committee and are in close agreement with the National Institutes of Health Guide for the Care and Use of Laboratory Animals. All efforts were made to minimize animal suffering and to reduce the number of animals used. For prenatal intermittent hypoxia exposures, pregnant rats were placed from day 5 of pregnancy until delivery in alternating room air and 10% O2-balance N2 every 90 s (IHRA; n = 6 litters). For postnatal exposures, litters were exposed within 12 h from delivery to the intermittent hypoxia profile (RAIH; n = 6 litters). As control, both pregnancy and postnatal development occurred in normoxic conditions (RARA; n = 6 litters).

Intermittent hypoxia protocol. Animals were placed in four identical commercially designed chambers (30 × 20 × 20 in.; Oxycycler model A44XO, Reming Bioinstruments, Redfield, NY), which were operated under a 12:12-h light-dark cycle (6:00 AM to 6:00 PM). Gas was circulated around each of the chambers, attached tubing, and other units at 60 l/min (i.e., 1 complete change/10 s). The O2 concentration was continuously measured by an O2 analyzer and was changed by a computerized system controlling the gas valve outlets, such that the moment-to-moment desired O2 concentration of the chamber was programmed and adjusted automatically. Deviations from the desired concentration were met by addition of N2 or O2 through solenoid valves. Ambient CO2 in the chamber was periodically monitored and maintained at <0.01% by adjusting overall chamber basal ventilation. The

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gas was also circulated through a molecular sieve (type 3A, Fisons, Cheshire, UK) so as to remove ammonia. Humidity was measured and maintained at 40–50% by circulating the gas through a freezer and silica gel. Ambient temperature was kept at 22–24°C. Control animals were exposed to circulating normoxic gas in one of the chambers.

**Respiratory measurements.** Respiratory signals elicited by displacement of the chest were acquired in the restrained animals placed in a size-appropriate head-out plethysmographic chamber (Buxco Electronics, Troy, NJ) by using the barometric method (1, 21), such that gas mixtures could be rapidly changed as needed. For all experiments, environmental temperature was maintained constant, slightly below the thermoneutral range for rat pups (30°C). Pressure changes in the chamber due to the inspiratory and expiratory displacements were measured by using a high-gain differential pressure transducer (model MP45-1, Validyne), displayed on the computer screen, and continuously digitized and stored onto a MacIntosh Personal Computer System at 125-Hz sampling frequency as previously described (14). Differences between the various treatment groups were compared by two-way analysis of variance (ANOVA) and the Newman-Keuls multiple-range test for multiple comparisons. For comparisons of autosuscitation rates, one-way analyses of variance and χ² analyses were conducted followed by Fisher exact test as appropriate. A P value of <0.05 was considered to achieve statistical significance.

**RESULTS**

IHRA pups were smaller at birth (5.4 ± 0.4 g) compared with RARA (7.2 ± 0.3 g; P < 0.001) and RAIH pups (7.0 ± 0.3 g; P < 0.001 vs. IHRA; P = not significant vs. RARA). At postnatal day 5, IHRA pups were still smaller (8.6 ± 0.5 g), and RAIH pups showed significant decreases in weight gain (7.7 ± 0.5 g) compared with RARA pups (10.7 ± 0.5 g; P < 0.001 vs. IHRA and RAIH). However, brain weights were similar for all three groups at all postnatal ages (P = not significant).

The triphasic gasping pattern occurred in all experimental groups (Fig. 1). Mean gasp latencies were similar in RAIH (32 ± 6 s), IHRA (34 ± 6 s), and RARA pups (30 ± 7 s; n = 20/group; P = not significant). However, the mean total gasp duration was significantly reduced in both IHRA and RAIH compared with RARA pups (18.6 ± 3.1, 15.6 ± 2.9, and 24.7 ± 4.6 min, respectively; Fig. 2; P < 0.001). Similarly, the overall number of gasps was diminished in IHRA and RAIH pups (Fig. 2; P < 0.001). Indeed, the mean total number of gasps was 31.5 ± 5.2 in IHRA, 26.2 ± 5.1 in RAIH, and 44.2 ± 7.3 in RARA pups. The mean gasp frequency was not modified during each of the gasping phases (Fig. 2).

The ability to autosuscitate from gasping during phase 3 was assessed in thirty 5-day-old pups from the various litters corresponding to each experimental group. Significantly fewer IHRA (21 of 30) and RAIH (19 of 30) pups successfully autosuscitated compared with RARA pups (28 of 30; Fig. 3; P < 0.03 RARA vs. IHRA and P < 0.01 RARA vs. RAIH respectively).

**DISCUSSION**

In this study, we show that intermittent hypoxia occurring either prenatally or during the initial days of life adversely affects gasp generating mechanisms and that the ability to autosuscitate during the late phase of gasping is markedly curtailed.

The neural substrate for gasp generation has been assigned to a discrete region within the caudal brain stem, the lateral tegmental field of the medulla (8, 9, 25, 26). This concept has been further expanded to incorporate the possibility that the critical regions for respiratory rhythmicity may dynamically reconfigure to generate gasping (19, 25, 26). Studies from our laboratory have previously shown that N-methyl-D-aspartate glutamate receptors are critically involved in
particular components of gasp maturation (11), specifically in the early phases of respiratory activity that follow primary apnea. In addition, our laboratory has shown that increased neuronal nitric oxide synthase expression and activity occur with increasing postnatal age within the neural sites responsible for gasp generation and underlie some of the characteristic developmental changes in gasp activity (12, 13). The overall changes in N-methyl-D-aspartate receptor and neuronal nitric oxide synthase expression associated with intermittent hypoxia during fetal and early postnatal life are currently unknown.

The overall reduction in gasping duration and total gasp number as well as the increased failure rates during autoresuscitation attempts could reflect deficient glycogen stores. IHRA pups were smaller at birth, and, although they demonstrated some catch-up growth after birth, they remained smaller than controls. Similarly, RAIH pups showed lesser weight increments than RARA, suggesting that fuel tissue reserves may be compromised in both experimental groups. Relative depletion of cardiac glycogen has been proposed as a potential mechanism underlying significant differences in the autoresuscitation of SWR compared with BALB/c weanling mice (3). Similarly, depletion of cardiac glycogen with recurrent autoresuscitations was advanced as one of the mechanisms potentially leading to autoresuscitation failure in the usually successful autoresuscitating BALB/c mice (3). However, external fuel delivery as brought about by glucose supplementation before the asphyxial exposure exerts divergent effects on gasping and autoresuscitation (28). Indeed, Yuan and colleagues (28) have shown that hyperglycemia will prolong the overall duration of gasping but reduce the frequency of autoresuscitation. Yuan et al. (29) also examined the role of adrenergic receptors, and found that adrenalectomy shortened gasping duration in both 1-day-old and 8-day-old rats, whereas the nonselective α-receptor antagonist phentolamine reduced the duration of gasping in 1-day-old rats but prolonged this duration in 8-day-old rats, with similar effects on gasping by the nonselective β-receptor antagonist propranolol. Thus these investigators concluded that gasping requires intact adrenal function and primarily involves α-adrenergic receptors. However, no significant differences were found in α-adrenergic receptor expression within the putative gasping centers of infants who succumbed to SIDS (20).

An interesting analogy to intermittent hypoxia can be drawn with prenatal nicotine exposures. Continuous administration of nicotine via an osmotic pump to pregnant rats from day 6 of pregnancy was associated with decreased gasp duration and with reduced ability to autoresuscitate until postnatal age 5–6 days but not thereafter (4, 5, 21). The adverse effect of perinatal nicotine was further shown by its acute administration to neonatal piglets at doses leading to serum levels similar to those found in infants of smoking mothers (7). In these experiments, laryngeal stimulation not...
only induced more severe central apneas in nicotine-treated piglets but also was associated with reduced ventilatory compensation after induced apnea (7). Thus both prenatal and early postnatal exposure to nicotine lead to impaired protective responses of rat pups that may sustain life during exposure to asphyxia. It should be emphasized, however, that the potentially deleterious effects of prenatal nicotine on gasping have not been consistently found (23). Whether these discrepant findings reflect the differences in dosages used or in the susceptibility of particular rat substrains remains unclear.

Although environmental temperature can modify gasping duration and the degree of successful autoresuscitation (24), we cannot a priori ascribe a role to this particular factor by virtue of our strict standardization of environmental temperature during all of our experiments. Similarly, all animals were studied at the same postnatal age, such that it is unlikely that there were marked differences in their developmental stages so as to induce the relatively robust changes in gasping and autoresuscitation characteristics associated with increasing postnatal age (6, 14, 18).

In summary, perinatal intermittent hypoxic exposures are associated with altered defense mechanisms against asphyxia, manifesting both as decreased total survival and as a relative inability to autoresuscitate. Although the implications of such increased susceptibility are unknown, we postulate that it may play a role in conditions such as SIDS.

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