

基础研究

孕期酒精暴露对新生大鼠延髓基本节律性呼吸放电的作用

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摘要:目的 探讨孕期酒精暴露对子代新生大鼠延髓脑片基本节律性呼吸放电(rhythmic respiratory discharge activity, RRDA)的作用。方法 制作包含面神经后核内侧区(the medial region of the nucleus retrofacialis, mNRF)并保留舌下神经根的新生大鼠离体延髓脑片标本,给予灌流改良的Kreb's液(modified Kreb's solution, MKS),使用吸附电极记录延髓脑片腹侧面舌下神经根RRDA,分析RRDA的变化。实验分为5组:对照组、4%酒精暴露组、6%酒精暴露组、8%酒精暴露组和10%酒精暴露组,研究孕期不同浓度酒精暴露对RRDA的作用。结果 对照组、4%酒精暴露组、6%酒精暴露组、8%酒精暴露组各组组内RRDA在50 min内吸气时程(TI)、呼吸频率(RF)、放电积分幅度(IA)无统计学差异,10%酒精暴露组RRDA节律不规整;与对照组相比,4%~10%酒精暴露组的RRDA随酒精浓度增加逐渐降低,TI缩短、RF下降、IA减弱。结论 孕期酒精暴露抑制子代新生大鼠延髓脑片舌下神经根RRDA,这可能是孕期酒精暴露抑制子代呼吸功能的基础。

关键词: 孕期酒精暴露;基本节律性呼吸放电;延髓脑片;新生大鼠

Effect of prenatal alcohol exposure on rhythmic respiratory discharge activity in medullary slices of neonatal rats

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Abstract: Objective To investigate the effects of prenatal alcohol exposure on rhythmic respiratory discharge activity (RRDA) in the medullary slices of neonatal rats. **Methods** Ten pregnant female SD rats were exposed to 0, 4%, 6%, 8%, and 10% alcohol in drinking water from 1 week before till 3 days after delivery. The medullary slices of the neonatal rats containing the medial region of the nucleus retrofacialis (mNRF) with the hypoglossal nerve rootlets were prepared and perfused with modified Kreb's solution to record RRDA from the hypoglossal nerve rootlets using suction electrodes. **Results** No significant difference was found in RRDA in 50 min among the neonatal rats with prenatal exposure to 0, 4%, 6%, and 8% alcohol, but the RRDA in 10% alcohol exposure group became irregular. Prenatal exposure to increased alcohol concentrations caused attenuated RRDA attenuated in the neonatal rats, shown by shortened inspiratory time (TI), decreased respiratory frequency (RF), and reduced integral amplitude (IA) as compared with those in the control group. **Conclusion** Prenatal alcohol exposure inhibits RRDA in medullary slices of neonatal rats, which might be a mechanism by which maternal alcohol exposure causes suppressed offspring respiratory functions.

Key words: prenatal alcohol exposure; rhythmic respiratory discharge activity; medullary slices; neonatal rats

呼吸是生命存在的前提和基本条件,呼吸运动是呼吸肌在呼吸中枢支配下节律性舒缩完成的。延髓是呼吸放电的产生部位,在本实验中所使用的新生大鼠延髓脑片包括面神经后核内侧区(the medial region of the nucleus retrofacialis, mNRF),而该区域是延髓内基本节律性呼吸放电产生和调节的精确部位^[1-2]。

女性孕期摄入酒精抑制胎儿各系统、器官的发育,

尤其对中枢神经系统发育的影响更严重^[3]。有研究表明,孕期母体饮酒能够降低幼鼠的每分通气量,损伤呼吸系统网络在低氧情况下的反应性^[4]。但孕期摄入酒精对新生动物呼吸中枢作用的报导尚不多见:Duncan等研究发现孕期饮酒降低了婴儿猝死综合征死亡的婴儿脑干呼吸循环相关核团神经元(3)H-nicotinic受体浓度,并随着饮酒量的增加呈剂量依赖性降低^[5];Conrad等发现孕期饮酒增加1月龄婴儿的呼吸性窦性心律不齐发生率^[6]。而孕期摄入酒精对新生动物延髓呼吸中枢基本节律性呼吸放电的作用未见报导。为此,我们通过记录分析孕期酒精暴露对新生大鼠延髓脑片基本节律性呼吸放电(RRDA)来研究孕期酒精暴露对新生大鼠延髓呼吸中枢的作用。

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1 材料与方法

1.1 实验动物和分组

30只SPF级Sprague-Dawley成年大鼠,雌雄比例2:1,购自郑州大学实验动物中心(许可证号SCXK豫2010-002),雌鼠和雄鼠随机分为数量相等的5组对照组、4%酒精暴露组、6%酒精暴露组、8%酒精暴露组和10%酒精暴露组(酒精浓度为体积/体积),按照雌雄大鼠2:1合笼交配。对照组大鼠自由饮用纯净水,不同浓度酒精暴露组大鼠自合笼前1周至产后3d以相应浓度酒精水溶液作为唯一饮用水(自由饮用)。实验使用各组2d龄新生大鼠(每组6只),雌雄不拘。

1.2 试剂

无水乙醇,天津市德恩化学试剂有限公司生产;改良Kreb's液(modified Kreb's solution, MKS) in mmol/L: NaCl 124、KCl 5、CaCl₂ 2.4、MgSO₄ 1.3、NaHCO₃ 26、Glucose 30所用药品购自Sigma;95% O₂+5% CO₂混合气购自河南源正科技发展有限公司。

1.3 脑片制备及电生理记录

2d龄5组新生大鼠各6只被乙醚深度麻醉后在C4~C5之间断头,将鼠头置于盛有95% O₂+5% CO₂混合气充分饱和的冰MKS的解剖盒内。眼科剪剪去皮肤和肌肉,从颈椎处剪开椎管并向吻端延伸,完整剪开颅骨、去除颅骨、暴露脑组织,手术刀片割断颅底脑神经、游离出完整的脑组织。夹去小脑、在C2~C3间切下脊髓,将脑组织腹侧面向上,在门前切下厚约900~1100 μm的延髓脑片,在操作过程中持续通95% O₂+5% CO₂的混合气。脑片切好后转移至脑片灌流槽内,使用混合气充分饱和并持续通气的MKS以5~7 ml/min的速度灌流,保持灌流液pH7.35~7.45、温度25~28℃^[7-8]。在体视显微镜下辨认脑片腹侧面舌下神经根并用内含银-氯化银电极的吸附电极(内径180~220 μm)在舌下神经根处加以负压吸附,记录RRDA。RRDA经直流前置放大器放大后输入BL-420F生物信号采集和处理系统在计算机显示器上显示并存储,以备实验后数据统计分析^[8]。

1.4 数据处理

为直观简洁对比实验结果,分别以对照组RRDA的吸气时程(inspiratory time, TI)、呼吸频率(respiratory frequency, RF)、放电积分幅度(integral amplitude, IA)等观测指标为100%,对各组10、20、30、40、50 min时间点RRDA进行标准化处理后再进行统计分析。计算公式:各浓度组原始数据/对照组原始数据×100%。

1.5 统计学方法

实验结果以均数±标准差的形式表示,使用SPSS 13.0统计软件重复测量方差分析和单因素方差分析检验,多重比较采用LSD法进行分析, $P<0.05$ 被认为有显

著性差异。

2 结果

2.1 对照组RRDA

经统计学分析,与10 min相比20、30、40、50 min时间点RRDA的TI、IA、RF无统计学差异,说明RRDA在实验过程中无衰减,证明本实验模型稳定可靠($P>0.05$) (表1,图1)。

表1 对照组不同时间点RRDA

Tab.1 RRDA in the control group at different time points

Timing	TI	IA	RF
10 min	100	100	100
20 min	101.24±2.18	100.84±1.39	101.63±1.75
30 min	99.67±2.12	101.67±1.82	99.20±2.04
40 min	100.36±2.55	98.43±2.28	100.96±1.86
50 min	99.01±2.74	99.08±2.77	98.67±2.31
<i>F</i>	0.263	0.328	0.438
<i>P</i>	0.849	0.764	0.599

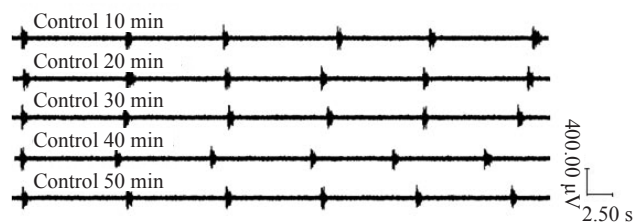


图1 对照组在50 min内不同时间点RRDA

Fig.1 RRDA in 50 min in the control group.

2.2 不同浓度酒精孕期暴露组

4%酒精暴露组、6%酒精暴露组、8%酒精暴露组各组内新生大鼠脑片RRDA各指标在不同时间点无统计学差异($P>0.05$,表2~4)成年大鼠饮水中酒精浓度增加,新生鼠RRDA逐渐降低,TI缩短,RF下降、IA减弱。10%酒精暴露组RRDA节律不规整,并且弱于8%酒精暴露组。因此,我们选择8%酒精作为孕期酒精暴露作用的最适浓度(图2)。

2.3 对照组和不同浓度酒精暴露组30 min时新生大鼠延髓脑片RRDA比较

孕期酒精暴露抑制剂量依赖性的新生大鼠延髓脑片RRDA,浓度为4%的孕期酒精暴露对RRDA的抑制作用已有统计学差异,RF低于对照组,6%酒精暴露组TI、RF、IA低于对照组和4%酒精暴露组;8%酒精暴露组TI、RF、IA均低于对照组、4%酒精暴露组和6%酒精暴露组(图3)。与对照组相比,孕期酒精暴露组新生大鼠延髓脑片RRDA的TI缩短,RF下降,IA减弱,具有统计学差异($P<0.05$,表5,图4)。

表2 4%酒精暴露组不同时间点RRDA
Tab.2 RRDA in 4% alcohol exposure group at different time points

Timing	TI	IA	RF
10 min	97.64±2.89	98.17±2.52	95.45±2.96
20 min	96.60±2.58	96.59±3.57	94.30±3.29
30 min	98.42±3.21	96.33±3.42	95.02±3.88
40 min	97.67±3.02	95.67±3.03	94.47±4.11
50 min	95.73±4.08	95.29±3.38	93.72±4.21
<i>F</i>	0.501	0.204	0.216
<i>P</i>	0.571	0.948	0.877

表3 6%酒精暴露组不同时间点RRDA
Tab.3 RRDA in 6% alcohol exposure group at different time points

Timing	TI	IA	RF
10 min	92.67±3.57	92.56±3.24	90.39±3.17
20 min	94.83±3.22	90.24±3.50	89.23±3.69
30 min	92.47±2.81	93.56±2.46	90.33±4.12
40 min	92.25±3.52	91.61±3.13	88.33±4.35
50 min	91.37±4.07	92.22±3.04	90.80±3.51
<i>F</i>	0.215	0.238	0.302
<i>P</i>	0.884	0.865	0.794

表4 8%酒精暴露组不同时间点RRDA
Tab.4 RRDA in 8% alcohol exposure group at different time points

Timing	TI	IA	RF
10 min	86.65±3.87	87.22±4.34	83.45±3.56
20 min	87.28±3.65	85.17±3.86	84.08±4.70
30 min	84.85±4.28	86.46±4.24	82.58±5.05
40 min	85.34±3.75	85.28±4.39	84.53±4.57
50 min	86.17±3.11	84.86±5.04	83.76±5.23
<i>F</i>	0.312	0.207	0.266
<i>P</i>	0.773	0.939	0.801

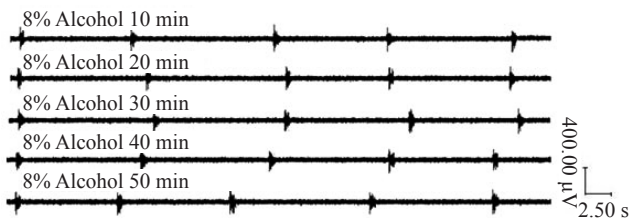


图2 8%酒精暴露组在50 min内不同时间RRDA
Fig.2 RRDA in 8% alcohol exposure group at different time points in 50 min.

3 讨论

近年来,关于酒精暴露对神经系统的影响已成为国内外研究的热点。在神经系统发育早期,即神经元突触

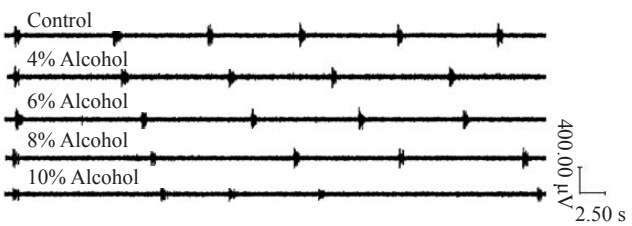


图3 对照组和不同浓度酒精暴露组RRDA的比较
Fig.3 Comparison of RRDA in neonatal rats among the groups. Prenatal exposure to increased alcohol concentrations attenuated RRDA in the medullary slices of the neonatal rats with shortened TI and decreased IA and RF.

表5 各组30 min时RRDA数据比较
Tab.5 Comparison of RRDA among the groups at 30 min

Groups	TI	IA	RF
Control group	99.67±2.12	101.67±1.82	99.20±2.04
4% Alcohol exposure group	98.42±3.21	96.33±3.42	95.02±3.88
6% Alcohol exposure group	92.47±2.81	93.56±2.46	90.33±4.12
8% Alcohol exposure group	86.85±4.28	86.46±4.24	82.58±5.05
<i>F</i>	35.938	24.776	29.621
<i>P</i>	0.000	0.000	0.000

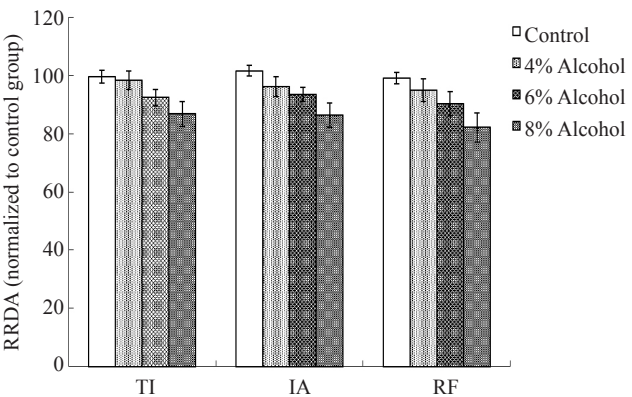


图4 不同浓度酒精暴露对新生大鼠RRDA的影响
Fig.4 Influence of prenatal exposure to different alcohol concentrations on RRDA of neonatal rats. RRDA in the alcohol exposure groups were lower than those in the control group.

形成时期,短暂接触酒精就能引起神经元的大量死亡。慢性酒精暴露能够影响神经元的分化和迁移,影响突触和轴突的生成,并可通过影响Na⁺/K⁺-ATP酶、腺苷酸环化酶的活性以及Ca²⁺通道开放,诱导神经元凋亡^[9],引起形态结构发育障碍,进而引起胎儿发育不良。而婴儿发育不良的程度与母亲孕期的饮酒量、饮酒时间和次数有关^[10-11]。

本实验发现,与对照组相比,4%~8%酒精暴露组的RRDA随酒精浓度增加而逐渐减弱,TI缩短、RF下降、IA降低。提示孕期酒精暴露对RRDA具有浓度依赖性

抑制作用。孕期酒精暴露抑制RRDA的机制可能与以下几方面相关:首先,孕期酒精暴露改变子代中枢神经系统神经元细胞膜受体表达,通过增加抑制性受体表达减少兴奋性受体来降低神经元兴奋性;胎儿在母体内接触酒精降低兴奋性受体NMDA-2B受体^[12]和5-HT_{2A}受体表达^[13],孕期酒精暴露增加GABA受体活性并增加其表达^[14-15]。以上各受体均参与延髓呼吸的发生和调节^[16-18],孕期酒精暴露通过抑制呼吸兴奋性受体,激活抑制性受体,从而抑制延髓呼吸中枢功能,我们的实验结果也与上述文献一致。其次,酒精及其代谢产物还能通过离子通道来发挥作用,如降低Na⁺/K⁺-ATP酶^[19]和Ca²⁺通道^[20]的活性,而Na⁺内流和K⁺外流是动作电位形成的基础,钙离子是重要的生物信号传导分子,调控细胞的理化活动。再次,孕期酒精暴露可降低脑源性神经生长因子BDNF浓度,BDNF在神经元的存活和突触形成中起重要作用,其浓度的下降影响了神经元活性,降低神经网络功能^[21-22]。另外,孕期酒精暴露使胎儿脑内线粒体发育迟缓,功能受损,ATP供给不足,从而影响了神经元活性及功能^[23]。有研究表明,孕期酒精暴露能降低子代大鼠的每分钟通气量^[4],而这极可能与孕期酒精暴露抑制新生大鼠延髓基本节律性呼吸放电有关。

综上所述,本研究结果表明,孕期酒精暴露抑制新生大鼠延髓脑片基本节律性呼吸放电活动,其机制可能是酒精通过诱导呼吸相关性受体、离子通道、神经生长因子及线粒体等发生改变,进而促使呼吸相关性神经元的活性下降和功能紊乱。孕期酒精暴露抑制新生大鼠延髓脑片基本节律性呼吸放电,这可能是孕期酒精暴露抑制子代整体水平呼吸功能的基础。

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