Push-in Head Restraining Apparatus for Intracranial Self Stimulation Tasks in Rats

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Head restraining is an experimental technique that firmly secures the animal's head to a fixation apparatus for the precise control and sensing of behaviors. However, procedural and surgical difficulties and limitations have been obstructing the use of the technique in neurophysiological and behavioral experiments. Here, we propose a novel design of the head-restraining apparatus which is easy to develop and convenient for practical use. Head restraining procedure can be completed by sliding the head mounter, which is molded by dental cement during implantation surgery, into the port, which serves as matching guide rails for the mounter, of the fixation bar. So neither skull-attached plates nor screws for fixation are needed. We performed intracranial self stimulation experiment in rats using the newly designed device. Rats were habituated to acclimatize the head-restraint environment and trained to discriminate two spatially distinguished cues using a customized push-pull lever as an operandum. Direct electrical stimulation into the medial forebrain bundle served as reward. We confirmed that head restraining was stable throughout experiments and rats were able to learn to manipulate the lever after successful habituation. Our experimental framework might help precise control or sensing of behavior under head fixed rats using direct electrical brain stimulation as a reward.

Key Words: Apparatus, Head fixation, Intra-cranial self-stimulation, Operant conditioning, Rat

INTRODUCTION

Head restraining (or head fixation) is an experimental technique that firmly secures the animal's head (or skull) to a fixing apparatus not to move freely. This technique has been used for experimentation on precise control and sensing of behavior [1-4] and brain imaging [5] in awake behaving animals. The experimental procedure incorporating head restraining basically begins with exposing the animals repeatedly to head restraining situation. Throughout several sessions of habituation, animals are getting accustomed to the unfamiliar and stressful environment. Skillful handling and not to make them annoying seem to be essential for the successful head restraining. More details about head restraining, particularly in rodents, are well documented by Schwarz [6].

Most of the fixation has attained by anchoring plates that is attached on the animal's skull to the fixation apparatus [7,8]. Under such designs, additional plates or pieces should be surgically glued on top of the animal's skull [1,5,9,10] and the plates are fixed to the fixation apparatus by screws. If the electrode implantation for recording and/or stimulation needs to be concurrent with the head restraining, for example, to examine neuronal activity while whisker movements and intracranial self stimulation (ICSS) task under the head restraining, the glued plates could spatially restrict the implantation of electrodes. Another minor problem is the perturbation of animals' natural behavior when they are not engaged in experiments, for example, when they are in the home cage. Moreover, inconvenience is originated from screw-tightening. During the habituation period of the head restraint, animals show struggling behavior at the moment of which the head is about to be fixed. This indicates that experimenters who have little experience of handling animals could be somewhat uncomfortable with tightening screws.

To overcome the incompatibility of the fixation apparatus and electrodes implantation and its inconvenience, a novel design for head fixation apparatus is suggested and deve-
METHODS

Design of head restraining apparatus

The drawing and implementation of the fixation apparatus are shown in Fig. 1A and B. The apparatus is composed of the fixation bar and box, both made of acrylic panel (6 mm thickness). The bar is the key component for a convenient fixation procedure and can be installed into the box by tightening screws through the two elongated holes at each side. Those holes allowed the bar to be installed with a tilt. A port is prepared in the middle of the bar (Fig. 1A, shaded area) and it fits the overhead mounter molded in head-restrained rats [6]. Therefore, we decided to make a small lever to serve as an operandum for head-restrained rats. The push-pull lever was basically designed to be small enough to be installed in the fixation box and could indicate at least two states using a single arm operation of the animal. The lever was assembled with a stick and two micro-switches disjoined from the computer mouse, and cased with acrylic plate of 1 mm thickness. Those two micro-switches were oppositely placed adjacent to the lever stick as shown in Fig. 1B and E. Operational principles are simple. If the lever stick was pushed, it triggered the micro-switch of the pull position and vice versa. The on and off signal was then delivered to the digital IO device (National Instruments) and detected by computer programs. Since about 70 to 80% of rats are right-pawed [7,8], the lever was attached at the right side of the floor where rat’s forepaw was naturally placed upon the lever stick when head was fixed (Fig. 1F).

Animals and surgery

Male Sprague-Dawley rats (n=4) weighing 300~350 g were housed in home cages with ad libitum access to food and water. The light cycle begins 7 AM and lasts for 12 hours and room temperature was maintained as 20 to 25 °C. After surgery, rats were housed individually in separate cages. All experiments were performed according to Guide for the Care and Use of Laboratory Animals (National Institute of Health, 1996).

Rats were anesthetized with the cocktail mixture of ketamine 10 ml (50 mg/ml), xylazine hydrochloride 1.5 ml (mg/ml), and saline 2.5 ml. Anesthesia was confirmed by no responses to tail and toe pinches. Animals were then fixed in a stereotaxic apparatus (David Kopf Instruments, USA). The position of rat skull was set to be flattened. After 2% lidocaine s.c. injection and midline incision, the periosteum was completely scraped off the skull surface. Eight holes were drilled and screws (1mm diameter) were implanted into the holes of the skull (frontal AP5.0, ML±2.0 and AP0, ML±5.0, parietal AP-9.0, ML±5.0, occipital AP-3.0 from Lambda, ML±2.0) according to Paxinos and Watson coordinates [13]. Then super bond (Super-Bond C&B, Sun medical, Japan) was carefully applied around the screws over the skull. Two holes for stimulation electrodes (insect pins – stainless steel, No. 00 – coated with formvar except the tip) were drilled after the super bond became hard. The stimulation electrodes were then implanted into the bilateral MFB area of coordinate AP-2.3 mm, ML±1.8 mm,
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DV-8.2 (from the dura). Dental cement was applied to fix the electrodes firmly.

After the implantation surgery, dental cement was applied around the edge of the port of the fixation bar (Fig. 1A) which is wrapped with Parafilm® (Pechiney Plastic Packaging, Chicago II, USA) (Fig. 1C). Care was taken not to cover the bare acrylic part of the fixation bar with dental cement. When it hardened enough, the bar was fixed on the stereotaxic frame by aligning the port of the bar with the center of implantation cement (Fig. 1C, left). Additional application of dental cement into the gap between the port and cement applied during surgery adjoins dental cement of both permanently (Fig. 1C, center). When cement hardened, the mounter and fixation bar was separated by pulling the overhead mounting out of the port and this naturally formed the overhead mounter (Fig. 1C, right). Parafilm® covering the port enables this procedure to be simply achieved. After surgery, it is natural that the head mounter serves guide rails for head restraint as shown in Fig. 1D.

**Behavioral tasks**

After recovery, rats were first tested whether they respond to the brain stimulation reward by introducing them to a general operant chamber containing a lever. Whenever rats press the lever, they could receive an electrical stimulation directly into their MFB. During the initial shaping stage, stimulation intensity was gradually increased. Reached at a specific intensity, rats kept on pressing the lever and this was regarded as a successful demonstration of ICSS. For that case, it deemed that stimulation electrode was precisely located at the MFB area.

Next, rats showing operant behavior were introduced to initial head restraining training using the devised head fixation apparatus. The first step of head fixation was simply pushing the overhead mounter into the port. Then the overhead mounter was tightly secured to the bar using a general rubber band (the wrist rings of latex gloves were pre-

![Table](Table.png)

**Fig. 2.** Procedures of habituation and experiment. (A) Habituation procedures for desensitizing rats to head-restraint constitutes repetitive and longer exposure to head-restraining condition. HR: head-restraining habituation only. HR+ICSS: ICSS was allowed (B) The basic structure of the rat conditioning.

Previously suggested protocols [1,9] were selectively modified and combined for the habituation of head restraints. The detailed protocol for head restraining used in this study is described in Fig. 2A. Basic idea is that rats became exposed to head restraining condition with short time duration and this duration is getting longer across sessions (from 5 min to 60 min across 5 sessions). Rats that did not show any struggling and/or awkward behavior at least 1 hour of head restraining at the final session proceeded to the ICSS tasks under the head restraining (ICSS-HR) condition. For ICSS-HR training, the push-pull lever was installed on the box floor.

In general, reward stimulation into the MFB elicits hyperactivity including exploratory and sniffing actions in the freely moving condition. In the head restraining condition, such hyperactivity could result in the separation of the overhead mounter from the rat’s skull. To prevent this tragic accident, stimulation intensity was lowered about 10 to 50 μA from the previous phase at the initial stage of ICSS-HR phase.

![Graph](Graph.png)

**Fig. 3.** Behavioral data for the initial ICSS shaping phases. The number of lever-pulls (triangle) and consequent rewards (circle) are depicted for the rat RS007 (A), RS008 (B), and RS009 (C) for consecutive three sessions (Day1 to 3). Numbers in the parenthesis indicates predetermined inter-trial intervals and stimulation current intensity. When the parameters are modified, it is denoted by the dotted vertical line. The number of rewards that is larger than the number of lever-pulls indicates the delivery of free-reward by the experimenter.
The experimental design of the initial ICSS-HR training was simple as shown in Fig. 2B. In a dark room, the house light and white noise were turned on, indicating the beginning of training. After three sec of delay, the light cue was turned on, indicating that rats could be rewarded when they pull the lever. If rats pull the lever by their right fore-limb, the light cue immediately turned off, and after 100 msec delays, contingent reward stimulation (biphasic, train duration 300 msec, pulse duration 0.2 msec, pulse width 4 msec) was delivered. The light cue was turned on again following a break period (0.2 to 9.6 sec, pre-defined) and next trial was begun. This procedure lasts for about an hour for each of the individual rat.

Next, the design was slightly modified to see whether rats could learn the association between the cue and lever-pull. The duration of the light cue was fixed to 3 sec, and the break period after reward was randomly set for each trial (1.6 to 4.6 sec). Correct lever-pulls resulted in immediate cue-off and reward delivery. If rats pull the lever during the break, the period was reset from that moment indicating a longer break period.

Lastly, rats were introduced to spatial cue discrimination tasks after several sessions of left cue and lever-pull training. In this task, rats were rewarded by pushing or pulling the lever when the left or right cue had been presented, respectively. The left or right cue was randomly presented at each trial. The duration of both cues was fixed to 5 sec but correct lever manipulation shortened the duration (that is, cue-off). The break between trials was set to 2.6 sec.

All training and task was under control of customized software implemented by Labview 8.5 (National Instruments) and behavioral data was collected using data acquisition device (National Instruments). Data was analyzed using Matlab software (Mathworks) off-line. The number of lever-pull or push was converted “events per min” on a 5 min bin basis. The performance of lever-pull training is shown in Fig. 3. In the first session (Day0), no rat manipulated the lever. In the second session (Day1), the rat (RS007) merely pulled the lever until 50 min (Fig. 3A). However, the rat began to pull the lever from 45 min and the frequency tended to increase until the end of the session. The more number of rewards than lever-pulls in the first bin indicates that free reward was delivered by the experimenter for shaping purposes. At Day2, the rat pulled the lever from the beginning of the session. The higher number of lever-pulls than rewards implies that the rat pulled the lever during the break period. The slight reduction in stimulation intensity (100 to 90 μA; separated by a dotted bar, Day2 in Fig. 3A) resulted in less lever-pulls (30 to 55 min). The longer inter-trial-interval was associated with the decrease in the number of lever-pulls in the later part (2 to 10 sec; separated by a dotted bar, Day3 in Fig. 3A). Another rat (RS008) showed an abrupt increase in lever pulls around 30 min of Day1 (Fig. 3B). The number of lever-pulls increased in Day2. Increased inter-trial interval resulted in the decreased number of rewards (Day3). The other rat (RS009) showed no interest in pulling the lever even though extensive free reward in Day1 (Fig. 3C). The intermittent lever-pulls were highly associated with struggling behavior.

The performance of lever-pull training is shown in Fig. 4. The rat RS007 showed overall 87.94% of success rate and the rat RS008 showed overall 74.94% of success rate. The performance of rats during the final 10 to 15 min of training was nearly perfect in both of the rats, indicating that rats learned how to behave (pull the lever) to receive reward. The rat RS009 did not show any capacity of task learning.

RESULTS

During the initial habituation in the head restraining apparatus, rats showed struggling behavior, usually running in place and squeaked when touched their back or trunk by an experimenter accidentally. However, those stressful responses were diminishing across the habituation session. The overhead mounter inserted into the port was firmly secured by general rubber band and rats could never fall off by their muscle force. The rats sit in the rear part of the box but nestled down somewhat forward when they were fully habituated. Respiration rate slowed down across sessions was observed by visual inspection. Food reward which was given to the rats after daily habituation was consumed by both rat RS007 and RS008 but the rat RS009 was not.

In the ICSS check phase, three rats showed repeated lever presses at the stimulation intensity of 90 to 150 μA, thus those rats underwent subsequent ICSS-HR training. Rats readily pulled the lever even though their head was fixed. Their locomotion activity was increased by stimulation reward at first but decreased across session. Shaping of two rats (lever-pull task only) was successful in the ICSS-HR paradigm as shown in Fig. 3. The rat RS007 rarely pulled the lever until 50 min (Fig. 3A). While the rat RS008 showed an abrupt increase in lever pulls around 30 min of Day1 (Fig. 3B). The number of lever-pulls increased in Day2. Increased inter-trial interval resulted in the decreased number of rewards (Day3). The other rat (RS009) showed no interest in pulling the lever even though extensive free reward in Day1 (Fig. 3C). The intermittent lever-pulls were highly associated with struggling behavior.

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so it was excluded in the following cue discrimination task. Lastly, the result of spatial cue discrimination in the ICSS-HR condition is shown in Fig. 5. During the consecutive four sessions of training, the rat RS007 showed growing performance (Fig. 5A), while the other rat RS008 showed nebulous performance (Fig. 5B). Those results are summarized in Fig. 6. Overall performance tends to increase (Fig. 6A) and the response latency tends to decrease (Fig. 6B) across session in the RS007 rat.

DISCUSSION

In this study, a novel head-restraining apparatus was developed and tested for its feasibility of instrumental conditioning using brain stimulation reward.

Head-restraining using the devised apparatus had been stable enough throughout experiments. The overhead moun- ter guaranteed a firm fixation. It was designed to use no additional attachment such as plates and pieces and be compatible with brain recording and stimulation electrodes by providing sufficient space above the skull. The exper- imenter can easily and firmly fix a rat’s head to the appara- tus without any screw-like fastener.

The design of the push-pull lever was also adequate to manipulate by rats. The stick of the lever could be grabbed by the right forepaw and manipulated within the moving range of the lever, indicating ergonomically suitable for rats. The lever position that should be placed beneath the right forepaw might be important for successful behavioral tasks in head-restrained rats.

In the following behavioral tasks, successful operant conditioning was not observed for all rats. Only the rat, RS007, operantly conditioned and performed discrimination tasks successfully under head-restraining using the push-pull lever. The behavioral performance was better than other rats from the initial shaping phases. Another rat, RS008, showed an imperfect task performance. Though it showed good performance during initial shaping phases which used lever-pull task, but it did not learn to manipulate the push-pull lever during the cue discrimination task. It seems that the rat was not exposed enough to precedent push training to learn the association between the left cues and push behavior. Another possibility might be an improper conditioning. The rat showed excessive running behavior during shaping. This resulted in unintended lever-pulls and reward delivery. Admitting the individual differences, RS007 showed better leaning capability compared with RS008. The other rat, RS009, showed no interest in manipulating the lever. The rat did not show intake of food reward after head-restraining habituation, which might be unable to overcome the stress. During the initial shaping phases, the rat received many free rewards without any voluntary or even unintended lever manipulation, indicating an improper conditioning. So it is possible that the rat was highly stressful during the habituation and was not properly habituated and trained.

Overall, rats showing a poor performance were not trained with additional sessions, so if they had been habi- tuated and trained more sessions they might have learned and performed the task better under minimized stress and aversion. While other study had spent two or more weeks [6,10], only minimum time (five days) was spent for habituating head-restrained rats. For a better behavioral performance under head-restraint condition, sufficient habituation period seems to be crucial.

This study demonstrated a successful operant conditioning in head-restrained rats using brain stimulation reward for the first time. ICSS paradigm was compatible with head-restraining condition, thus, our experimental frame-
work might contribute to examine not only the precise control or sensing of behavior but also behavioral operant conditioning in head fixed rats. Though this study focused on the feasibility of operant conditioning using brain stimulation reward (ICSS) under head restraining, the apparatus can be used in combination with electrophysiological techniques that are extensively employed such as single or multi unit recording and \textit{in vivo} patch clamping.

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