Social Isolation in the Rat Produces Developmentally Specific Deficits in Prepulse Inhibition of the Acoustic Startle Response Without Disrupting Latent Inhibition

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A series of experiments examined the effects of 8 weeks of social isolation on spontaneous locomotor activity, prepulse inhibition (PPI) of the acoustic startle response, latent inhibition (LI) in a conditioned suppression paradigm, and basal and d-amphetamine stimulated dopamine (DA) release in the ventral striatum, as measured by in vivo microdialysis. Both isolation-reared animals (those isolated from the weaning age) and isolation-housed animals (those isolated as adults) were hyperactive when placed in a novel environment. Social isolation also led to deficits in PPI of the acoustic startle

response that were specific to isolation-reared animals. Isolation rearing was without effect on the expression of LI but did lead to an enhanced response to systemic d-amphetamine in terms of striatal DA release. The data are discussed with respect to the involvement of ventral striatal DA mechanisms in the expression of PPI and LI, differences in the impact of social isolation in young and adult animals, and the utility of social isolation model as a nonlesion, nonpharmacologic means of perturbing ventral striatal DA function.

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Variations in early social experience have been shown to have profound effects on the adult behavior, brain morphology, and neurochemistry of a variety of species (Sachser 1986; Lewis et al. 1990; Martin et al. 1991; Jones et al. 1992). For example, rearing rats in social isolation from the weaning age produces behavioral abnormalities that include increases in spontaneous and

conditioned locomotor activity (Gentsch et al. 1981; Sahakian et al., 1982; Jones et al., 1990), altered responses to environmental novelty (Sahakian et al. 1977; Gentsch et al. 1982a, 1982b, 1983), impairments in discrimination learning (Jones et al. 1991), and resistance to extinction (Einon et al. 1975; Morgan et al. 1977). Rats isolated in this way also show an increased sensitivity to psychomotor stimulant drugs such as *d*-amphetamine and cocaine, both in terms of the locomotor enhancing properties of these agents and their ability to induce stereotyped behaviors (Sahakian et al. 1975; Jones et al. 1990, 1992), responses that depend on the mesolimbic and mesostriatal dopamine (DA) projections, respectively (Kelly et al. 1975). Consistent with these behavioral data are studies utilizing in vivo microdialysis techniques, which have provided evidence that isolation-reared rats exhibit greater elevations in extracellular DA in both dorsal and ventral striatum in response to systemic *d*-amphetamine (Jones et al. 1992), thereby

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substantiating the hypothesis that early social isolation produces enduring changes in central dopaminergic function.

Recently, there has been considerable interest in the modeling of aspects of the attentional deficits in schizophrenia by paradigms that can be used in both humans and experimental animals. In this regard, schizophrenic patients have been reported to show deficits in both prepulse inhibition (PPI) of the startle response, in which a weak lead stimulus markedly reduces the response to strong startle-eliciting stimuli (Ison and Hoffman 1983; Braff et al. 1992), and latent inhibition (LI), a procedure in which nonreinforced preexposure to a stimulus retards subsequent conditioning to that stimulus (Lubow 1973; Baruch et al. 1988). Both PPI and LI have been shown to be affected by treatments that alter central dopaminergic transmission. For example, PPI is disrupted by systemic administration of *d*-amphetamine or the direct DA agonist apomorphine, effects that can be blocked by neuroleptic drugs such as raclopride (Davis et al. 1990; Mansbach et al. 1988; Swerdlow et al. 1991). Similarly, the LI effect is abolished by systemic *d*-amphetamine, whereas neuroleptic drugs enhance the LI effect (Weiner et al. 1988; Weiner and Feldon, 1987). In the case of PPI, there is strong evidence that some of the effects of dopaminergic manipulations involve the ventral striatum and its associated neural circuitry. Thus, infusions of DA directly into the nucleus accumbens (NAC) have been shown to disrupt PPI (Swerdlow et al. 1990a): an effect that can be reversed by infusion of the gamma-aminobutyric acid (GABA)-ergic agonist muscimol into the ventral pallidum, a major target area of the NAC (Swerdlow et al. 1990b). The situation with LI is less clear, in that recent evidence has indicated that the disruption of LI induced by systemic *d*-amphetamine is unlikely to be due to DA-dependent mechanisms of the ventral striatum (Killcross and Robbins 1993). Furthermore, discrete lesions of the ventral subiculum, the source of a major connection between the hippocampus and the ventral striatum (Groenewegen et al. 1991) are without effect on LI (Clarke et al. 1992). It would seem that, although the LI effect is sensitive to treatments that perturb DA transmission, the precise neuroanatomic substrate for these effects remains unknown.

The changes in central DA function brought about by social isolation suggest that it may be of interest to compare the effects of this manipulation on PPI and LI. Given the basic finding that treatments that enhance dopaminergic transmission in the striatum reduce PPI, one main prediction would be that isolates should show similar deficits in PPI. In fact, previous experiments have indicated that early social isolation does indeed impair PPI (Geyer et al. 1993). Nevertheless, these data leave several questions unresolved: in particular, the

developmental specificity of the effects of isolation. That is to say, does a period of isolation as an adult give rise to the same changes in PPI as that seen in animals isolated at the weaning age. This issue is important because both manipulations might be hypothesized to be forms of chronic mild stress, which is known to affect the functioning of the mesolimbocortical dopaminergic systems (LeMoal and Simon 1991). It has rarely been considered, however, that stressors might affect these systems differentially depending on the age of the animal. Therefore, the first experiment compared the effects of isolation rearing (i.e., isolating weanlings) and isolation housing (i.e., isolating adults) on PPI of the acoustic startle response. The second experiment examined the effects of isolation rearing on the development of LI in a within-subject, conditioned-suppression paradigm. Here, any predictions must be less firm in that, as mentioned previously, the precise role of striatal DA mechanisms in the expression of LI is not settled. Accordingly, it could be predicted that social isolates would show no deficits in LI, with the caveat that isolation may modify LI by producing changes in DA function at sites other than the ventral striatum. In view of the particular uncertainty regarding the effects on LI of elevating DA transmission in ventral striatum, these animals were also subsequently assessed for levels of extracellular DA in the NAC utilizing *in vivo* microdialysis, a technique that provides a direct measure of presynaptic transmitter dynamics (Ungerstedt 1984).

MATERIALS AND METHODS

Subjects and Housing Conditions

The experiments used a total of 64 male, Lister Hooded rats, 40 weanlings (21 days postnatal) and 24 adults (weighing approximately 300 g). All rats were obtained from Harlan Olac Ltd (Bicester, U.K.). On arrival at our laboratory, the animals were divided into two rearing conditions counterbalanced by weight. Animals were housed either in isolation or in social groups for the duration of the experiment. Social groups were housed in fours in cages 56 × 38 × 18 cm, and isolated animals were housed singly in cages 45 × 20 × 20 cm. All cages were of plastic construction with steel grid floors and underhanging sawdust trays (North Kent Plastic Cages Ltd, Dartford, U.K.). The animals were kept in a colony room maintained at 21°C on a 12-hour light/12-hour dark cycle (light on 0700) with free access to food and water. All rats could see, hear, and smell the other rats in the colony.

Experimental Groups

Testing began after 8 weeks of the various housing conditions. All animals were first scored for spontaneous

activity in a novel environment. Thereafter, rats entered one of two experimental pathways. In the first experiment, a total of 48 animals were assessed for acoustic startle and PPI of startle. This group contained rats that had been isolated from weaning age as well as rats isolated as adults together with their respective social controls ($n = 12$ for each group). The former group was defined as isolation-reared animals and the latter as isolation-housed animals to distinguish between the two isolation conditions. In the second experimental pathway, 16 animals (eight isolation reared and eight social controls) were tested for the expression of LI in an on-baseline, within-subjects, conditioned-suppression paradigm followed by assessment of DA levels in the NAC using in vivo microdialysis techniques.

Assessment of Spontaneous Activity

Apparatus. Spontaneous locomotor activity was monitored in 16 individual wire-mesh photocell cages ($40 \times 25 \times 18$ cm). Each cage was fitted with two parallel horizontal infrared beams, 1 cm above the floor, spaced equally along the long axis of the cage. Interruption of either beam resulted in an incremental count for that cage. Beam breaks were the dependent measure and were recorded on-line in an adjacent room by a BBC Master Series microcomputer equipped with a SPIDER extension (Paul Fray Ltd, U.K.). Data from the 2-hour session were collected in 12, 10-min time bins.

Behavioral Procedures. All animals were naive to the activity boxes and were tested in batches of 16 between 1600 and 1800 hours, that is, towards the end of their light period.

Assessment of Startle and Prepulse Inhibition of Startle

Apparatus. A SR-LAB (San Diego Instruments, San Diego, CA) test station was used to detect startle responses. The illuminated, ventilated chamber contained a stabilimeter composed of an 8.2-cm diameter Plexiglas cylinder mounted on a Plexiglas base as described previously (Swerdlow et al. 1991). A speaker mounted 24 cm above the animal provided the background noise, prepulse stimuli, and startle stimuli, which were controlled by the SR-LAB. Startle responses were transduced by a piezoelectric accelerometer mounted below the cylinder, digitized (0-4095), rectified, and recorded as 100 1-millisecond readings, starting at the onset of each startle stimulus. The average of these 100 readings was used as the dependent measure.

Behavioral Procedures. For testing, rats were placed into the startle apparatus and allowed a 5-minute acclimation period during which a 65-dB [A] (all sound levels refer to the A scale) background noise was ad-

ministered. The startle session began with three successive 120-dB startle stimuli of 30-millisecond duration, followed by five blocks of trials, each comprising two 120-dB startle stimuli alone, four prepulse stimuli + startle stimuli, and two no-stimuli trials. Within each block, individual trials were distributed in a pseudo-random fashion. Four different intensities of acoustic prepulse stimuli (2, 4, 8, and 16 dB above background) were used, each prepulse being 30 millisecond in duration. The time from onset of the prepulse to onset of the startle stimulus was always 100 milliseconds. In all experiments, the variable interval between trials averaged 10 seconds; hence, the session lasted approximately 15 minutes.

Assessment of Latent Inhibition in a Within-Subjects Conditioned-Suppression Paradigm

Apparatus. Four standard operant chambers ($26.5 \times 22 \times 20$ cm; Campden Instruments) housed in light- and sound-attenuating boxes were used for the procedures. The chambers were illuminated by a diffuse light source located above the translucent plastic ceiling. The floor of the chambers consisted of 16, 5-mm diameter, steel rods spaced 5 mm apart. These grids were each connected to a shock generator (Model 512/C; Campden Instruments), and a shock scrambler (Model 512/S; Campden Instruments), set at 0.3 mA. Two discrete auditory stimuli were available in each chamber: one was a 3-kHz tone, produced by a Sonalert module (Model SC 628) and delivered through a wall-mounted speaker located opposite to the food magazine, and the other derived from a heavy-duty relay mounted behind the back wall providing a 10-Hz clicker. Each chamber contained a single lever, 3.8-cm wide, located to the left-hand side of a central, recessed magazine that provided access, via a hinged Plexiglas panel, to a dipper delivering 20% sucrose solution. Lever presses were the dependent measure (see below). A BBC Master-128 microcomputer, equipped with a SPIDER extension for on-line control (Paul Fray Ltd., UK), controlled the equipment and recorded the data.

Behavioral Procedures. Two days before the beginning of the test procedures, the animals were placed on a 22.5-hour schedule of food deprivation and maintained on this schedule throughout the rest of the experiment. Water was available ad libitum. All animals received two initial 30-minute magazine-training sessions with the levers removed from the chambers; sucrose was delivered according to a variable time 60-second (VT60) schedule. The levers were then replaced and two 30-minute sessions given on a continuous reinforcement schedule in which every lever press was rewarded, up to a maximum of 60 rewards per session. For the next two sessions, a variable interval 30-second

(VI30) schedule was employed: the first session lasting 30 minutes and the second 40 minutes. Thereafter 40-minute sessions on VI60 were used. After three such 40-minute VI60 sessions had been given, the preexposure phase of the experiment began.

Isolated and social animals were divided randomly into two groups. One group in each of the two conditions was assigned to preexposure to the 3-kHz tone and the other to preexposure to the 10-Hz clicker ($n = 4$ in all cases). Preexposure consisted of three 40-minute on-baseline VI60 lever-press sessions during which the to-be-conditioned stimulus was presented 12 times according to a pseudo-random schedule. Each presentation lasted 30 seconds, and the average ISI was approximately 3 minutes. The preexposure sessions and all subsequent sessions were conducted on alternate days. On the intervening days, animals were given one, 30-minute VI60 lever-press session to maintain baseline responding. Following the preexposure sessions, an assessment of unconditioned suppression to the non-preexposed stimulus was made. This session served to remove the initial disruption of lever pressing observed following presentation of any novel stimulus (Baker and Mercier 1982) and also provided a demonstration that the animals could, in fact, distinguish between the two auditory stimuli. During a VI60 lever-press session, both auditory stimuli were presented twice to every animal, in a counterbalanced order, and at equal intervals throughout the session. Finally, conditioning to both of the auditory stimuli occurred over four subsequent conditioning sessions. In each of these sessions, each of the auditory stimuli was presented once, again in a counterbalanced order, but in this case, the final 0.5 seconds of the stimulus presentation was accompanied by a 0.3-mA electric shock, delivered through the grid floor of the cage. The first conditioned stimulus was presented 10 minutes after the start of the session, and the second conditioned stimulus came 23 minutes later. This interval provided a sufficiently long period between the two shock presentations to allow for recovery from any suppression of baseline responding following shock presentation.

Measurement of Extracellular Dopamine in Nucleus Accumbens

Apparatus. Basal and *d*-amphetamine stimulated levels of extracellular DA in the nucleus accumbens were assessed using in vivo microdialysis. Animals were anesthetized with halothane (0.9% in O₂) and placed in a stereotactic frame with the incisor bar set at +5 mm. The skull was exposed, a burr hole made, and a 2-mm dialysis probe lowered into the brain so that the tip of the probe was at the following stereotactic coordinates, AP: +3.4 mm, L: -1.5 mm (from bregma), V: -7.5 mm (from dura). The dialysis probe was of a concentric de-

sign, made of 23-gauge steel tubing coupled to Hospal dialysis membrane (nominal 5000 molecular weight cut off). The probe was perfused with ACSF at a flow rate of 2 μ l/min. The composition of the ACSF was as follows: NaCl 120 mmol/L, NaHCO₃ 27 mmol/L, KCl 2.5 mmol/L, NaH₂PO₄ 0.6 mmol/L, Na₂HPO₄ 1.27 mmol/L, Na₂SO₄ 1 mmol/L, MgCl₂ 0.47 mmol/L, CaCl₂ 1.5 mmol/L, pH 7.4. At room temperature, the in vitro recovery of the probe for DA was 12% to 14%. The dialysis data were not corrected for recovery. Dialysates were collected every 30 minutes and immediately injected into a microbore high-performance liquid chromatography (HPLC) system. The HPLC system consisted of a BAS PM-60 pump connected via a column splitter to a reverse-phase analytical column (Spherisorb 30DS2; 10 cm $\times$ 1 mm inner diameter). The mobile phase, flowing at 70 μ l/min, was composed of 152 mmol/L citric acid, 15 mmol/L sodium acetate, 1 mmol/L octyl sulphate, 0.8 mmol/L ethylenediaminetetraacetic acid and 8% methanol, pH 3.6. Separated compounds were detected using a BAS LC-4B cell, fitted with a glassy carbon electrode held at a potential of +0.7 V relative to Ag/AgCl reference electrode. The output from the detector was analyzed by an integrator. Identification and quantification of the DA peak was achieved by comparison with external standards of known concentration. The identity of the DA peak was further confirmed by spiking with an internal standard. Under the conditions described the maximum sensitivity for DA was 0.25 pg (signal-to-noise ratio of 3).

Dialysis Procedures

All the dialysis experiments followed the same pattern. After a 1-hour postimplantation period, 3 hours of baseline determinations were taken followed by a further 3 hours of collection after the administration of *d*-amphetamine (free base, 2 mg/kg in 0.5 ml saline IP). Under the general conditions described, both basal and stimulus-evoked (75 mmol/L K⁺) release of DA were calcium dependent (60% reduction in basal levels, 75% reduction in stimulus-evoked levels; data not shown). Throughout the experiment, the core temperature of the animal was monitored with a rectal probe and maintained at 37°C.

Histologic Verification of Probe Placements

Immediately after the final dialysis sample, animals were killed by intraperitoneal injection of pentobarbital and perfused transcardially with phosphate buffered saline (PBS), pH 7.4, containing 1% sodium nitrite for 2 minutes, followed by 250 ml of 4% paraformaldehyde in PBS containing 0.4% picric acid. The brains were removed from the skull, postfixed in the same solution for 1 hour, and transferred to 25% sucrose in Tris-

buffered saline. After equilibration in the sucrose, sections were cut on a freezing stage sledge at 4°C prior to staining. One in five series was mounted on gelatin-coated slides and stained for cresyl violet and acetylcholinesterase, respectively. After staining, slides were dehydrated in alcohols, cleared in xylene, and mounted in DPX.

Statistical Analyses

Spontaneous Activity Scores. The activity data (beam breaks) were analyzed using a three-way analysis of variance (ANOVA), with two between-subjects factors, Social Experience (i.e., social or isolate) and Age (i.e., weanling or adult when treatment commenced), and one repeated within-subjects factor, Time Bin.

Startle and Prepulse Inhibition of Startle. Startle responses were analyzed using two- and three-way ANOVAs, with two between-subjects factors, Social Experience and Age, and one within-subjects factor, Block. Prepulse inhibition was defined as the difference between the responses on the 120-dB pulse-alone trials and the responses on the prepulse + 120-dB pulse trials divided by the average response on the corresponding 120-dB pulse-alone trials multiplied by 100. Prepulse inhibition data for each prepulse-type (i.e., 2, 4, 8, or 16 dB above background) were analyzed using two- and three-way ANOVAs with two between-subjects factors, Social Experience and Age, and one within-subjects factor, Block. Where appropriate, post hoc specific comparisons were made using Duncans Multiple Range Test (Winer, 1971).

Latent Inhibition. Because the experiments presented here used a within-subjects design, the analyses were performed on the number of lever presses made during the conditioned stimulus. The standard measure of conditioned suppression, namely a suppression ratio, is designed to reduce between-subject variance. Because the comparison of major interest in the present study is a within-subjects comparison and there were no differences in baseline rates of lever pressing between isolation- and socially reared groups in any of the experiments (see Results), use of suppression ratios would have been inappropriate and unnecessary. The variance of the data was found to increase with the mean. Accordingly, all lever press data presented were subjected to a square-root transformation prior to two- and three-way ANOVAs, with one between-subjects factor, Social Experience, and three within-subjects factors, Preexposure, Trial, and Response Bin.

Dialysis Data. Extracellular levels of DA were expressed in picomoles per 60 µl of dialysate and were not corrected for probe recovery. The data were analyzed using a two-way ANOVA with one between-

subjects factor, Social Experience, and one within-subjects factor, Time Bin. To meet the assumptions of equal variance and normality, basal and drug response data were analyzed separately and log transformed. Where appropriate, post hoc specific comparisons were made using Duncans Multiple Range Test (Winer, 1971).

RESULTS

Spontaneous Activity in Novel Environment

Figure 1 shows the activity scores in 10-minute time bins for isolation-reared and isolation-housed animals, together with their respective social controls on exposure to the activity cages. All four groups showed evidence of habituation over the 2-hour test period, as indicated by a significant main effect of Time Bin ($F[11,660] = 80.41, p < .0001$) in the three-way repeated-measures ANOVA with factors Social Experience, Age, and Time Bin. The ANOVA also revealed a significant main effect of Social Experience ($F[1,60] = 19.89, p < .0001$), reflecting increased activity in isolated animals. This effect was independent of the age of the animal when isolated; there was no significant Age by Social Experience interaction and was not related to any group differences in the rate of change of activity over the test period;

Figure 1. Time course of the locomotor activity of isolation-reared (■) and isolation-housed (●) animals together with their social control groups, socially reared (□) and socially housed (○). The data are expressed as the number of beam breaks per 10-minute bin ($n = 12$). The error bar represents twice the standard error of the difference between means (SED) for the three-way interaction with factors Social Experience, Age, and Time Bin. The SED is the denominator for making pairwise comparisons post hoc. Hence, the error bar provides an appropriate means of visually evaluating significance between pairs of points. The relevant formulae are given in Cochran and Cox (1957).

Figure 2. **A:** Startle responses for the initial three successive, 120-dB startle stimuli of the startle session as exhibited by isolation-reared (IR), isolation-housed (IH), socially reared (SR), and socially housed (SH) animals. The data are expressed in arbitrary startle units and are the mean plus or minus standard error of the mean (SEM) ($n = 12$). * Denotes a significant difference ($p < .05$) between the mean values obtained for isolates (both isolation reared and isolation housed) and social controls (Duncans Multiple Range Test). **B:** Mean startle responses to the 120-dB startle stimuli measured over the five blocks of trials comprising the remainder of the startle session and from which the extent of any PPI was assessed. The key to experimental groups is the same as in Figure 1. The bar represents twice the SED between means for the three-way interaction with factors Social Experience, Age, and Block.

there was no significant interaction involving Time Bin. Therefore, social isolation for a period of 8 weeks, in both the weanling and the adult rats, led to increased activity in the novel surroundings of the activity cages.

Startle and Prepulse Inhibition of Startle

Startle. Figure 2A illustrates acoustic startle responses for isolation-reared and isolation-housed animals, to-

gether with their respective social controls, for the first three presentations of the 120-dB startle stimulus. The two-way ANOVA of these data with factors Social Experience and Age revealed a significant main effect of Social Experience ($F[1,44] = 5.51, p < .05$) and no interaction, reflecting the fact that isolated animals (both isolation reared and isolation housed) were hyperreactive to these initial stimuli. Specific post hoc comparisons confirmed this conclusion. By contrast, there were no significant between-group differences in the subsequent responses to the 120-dB stimuli used to assess PPI (see Fig. 2B). Clearly, however, there was some degree of habituation on repeated presentation of the startle stimulus in all groups. This habituation was confirmed by a main effect of Block ($F[4,176] = 20.37, p < .0001$) in the three-way ANOVA with factors Social Experience, Age, and Block. No-stimulus values, which averaged approximately 8 startle units, did not differ over Block or between groups.

Prepulse Inhibition of Startle. Because the 2-dB and 4-dB prepulse stimuli failed to alter startle responses in all experimental groups, these data are not shown. Figure 3A illustrates the effects of the 8-dB prepulse stimulus on startle response. It is important to note that the data are collapsed across blocks of trials. This maneuver was justified because the three-way ANOVA with factors Social Experience, Age, and Block failed to detect any effects of, or interactions with Block. That is, for all groups, prepulse effects were remaining constant over the session. Hence, any between-group differences were not due to differential rates of habituation of the startle response itself. From Figure 3A, it can be seen that in three of the experimental groups, the 8-dB prepulse reduced startle responses by some 50%, a typical PPI effect. By contrast, in the isolation-reared group, this prepulse essentially failed to modify responding to the startle stimulus. The two-way ANOVA, with factors Social Experience and Age, revealed a significant Social Experience by Age interaction ($F[1,44] = 6.44, p < .002$). Specific post hoc comparisons indicated that the isolation-reared group was significantly different from the other three groups, confirming the above conclusions. The effects of the 16-dB prepulse stimulus on acoustic startle are shown in Figure 3B. This larger prepulse stimulus increased the degree of PPI in all experimental groups, although again, the absolute extent of PPI appeared to be less in the isolation-reared group. Nevertheless, increasing the intensity of the prepulse stimulus from 8 dB to 16 dB abolished the significant difference between groups, possibly due to a ceiling effect. Together, these data are consistent with the following conclusions: PPI of acoustic startle was selectively reduced in animals isolated at weaning age; the effect was not dependent on underlying changes in startle itself; and given the ability of more intense prepulse stimuli to elicit inhibition in

Figure 3. A: The percentage inhibition of startle produced by a prepulse stimulus 8 dB above the 65-dB background presented 100 milliseconds prior to the 120-dB startle stimuli, meaned over five blocks of trials. The key to experimental groups is the same as in Figure 2A. * Denotes a significant difference ($p < .05$) between isolation-reared animals and the other three experimental groups (Duncans Multiple Range Test). B: Percentage inhibition of startle produced by a larger prepulse stimulus, 16 dB above background. In both cases, the data are the mean $\pm$ SEM ($n = 12$).

this group (albeit less than that seen in other experimental groups), it was specifically the threshold for inhibition that seemed to have been altered.

Latent Inhibition in a Conditioned-Suppression Paradigm

Baseline Rates. One animal from the tone preexposed isolation-reared group and one from the tone preexposed socially reared group failed to respond following the first shock presentation; thus, they were dropped from the study and all subsequent analyses. Baseline rates of lever pressing in the isolate and social groups were not significantly different. The mean rates of lever pressing during the 30-second period immediately prior to the conditioned-stimulus onsets (preconditioned-stimulus period) across all conditioning trials

for isolation-reared and socially reared controls were: Isolates = 7.8; Socials = 5.7. A two-way ANOVA with Social Experience and Preexposure did not reveal any significant main effects or any interaction.

Unconditioned Suppression. Figure 4A shows the mean number of lever presses made during presentation of the preexposed and nonpreexposed stimuli from the two unconditioned stimulus exposures and from the first conditioning trial. Clearly, both groups of animals could discriminate between the two stimuli and demonstrated a significant degree of unconditioned suppression to the novel stimulus. This unconditioned suppression habituated at the same rate in both the isolated and social groups, and there was no difference in the suppression of lever pressing caused by either stimulus in either group by the first conditioning trial (see Fig. 4B). These conclusions are supported by the three-way ANOVA with Social Experience, Preexposure, and Trial that showed no significant main effect of Social Experience or interaction involving this factor. There were, however, significant main effects of Trial ($F[2,24] = 12.224, p < .001$) and Preexposure ($F[1,12] = 20.945, p < .001$), together with a significant Trial by Preexposure interaction ($F[2,24] = 8.563, p < .005$). Analysis of Simple Effects from this interaction indicated a significant effect of Trial on the preexposed stimulus ($F[2,24] = 11.814, p < .001$) and a significant effect of preexposure on the first unconditioned presentation ($F[1,12] = 23.697, p < .001$) but not on the second or third presentation. It would seem, therefore, that lever pressing was significantly suppressed on the initial presentation of the nonpreexposed stimulus, but that this suppression had habituated by the first conditioning trial.

Conditioned Suppression. Figure 4B shows the mean number of lever presses made by isolates and socials during presentation of the preexposed and nonpreexposed stimuli for the four conditioning trials. As expected, socials showed a strong LI effect manifest as a slowed conditioning to the preexposed stimulus relative to the nonpreexposed stimulus. Isolates showed qualitatively similar effects but the absolute values for lever pressing were higher during both preexposed and nonpreexposed stimuli. Thus, although there was no evidence of any reduction in LI in isolates, it would appear that lever pressing was influenced by some type of generalized response disinhibition in this group. These conclusions were supported by the three-way ANOVA with Social Experience, Preexposure, and Trial, which revealed no main effect of Social Experience but significant main effects of Preexposure ($F[1,12] = 13.233, p < .005$) and Trial ($F[3,36] = 10.866, p < .001$). None of the interactions were significant.

Shock-Induced Suppression. Figure 4C shows the distribution of lever press responding for isolates and so-

Figure 4. A: Mean number of lever presses made during presentations of preexposed and nonpreexposed stimuli prior to conditioning. Closed symbols refer to isolation-reared animals, open symbols to socially reared animals. Circles show responding to the nonpreexposed stimulus, squares to the preexposed stimulus. $n = 4$ in each cell. The bar represents twice the SED between means for the three-way interaction with factors Social Experience, Preexposure, and Trial. B: Mean number of lever press made during presentations of preexposed and nonpreexposed stimuli during conditioning. Note the appearance of LI (retarded conditioning to the preexposed stimulus) in both groups and the general retardation of conditioning in isolation-reared animals. Again, the bar

represents twice the SED between means for the three-way interaction with factors Social Experience, Preexposure, and Trial. C: Pattern of lever press responding across conditioning sessions presented as the mean number of presses made during nine, equally sized time bins across the 40-min session. Arrows indicate the points of postconditioned stimulus (CS) shock presentation (after 10 minutes and 33 minutes). Note the reduced degree of suppression in the isolation-reared animals (■) compared to socially reared controls (○). The bar represents twice the SED between the means for the three-way interaction with factors Social Experience, Response Bin, and Trial.

Extracellular Levels of Dopamine in Nucleus Accumbens

Figure 5 shows the basal and *d*-amphetamine stimulated levels of extracellular DA in the NACs of those animals assessed for LI, as measured by in vivo dialysis. All dialysis probes were placed accurately within the medial portion of the nucleus (see Fig. 6) with an anteroposterior range of less than 1 mm. In both social and isolation-reared animals, basal levels of DA were within the range of values reported by others (Zetterstrom et al. 1983). A two-way ANOVA involving Social Experience and Time Bin failed to reveal any significant effects of either of these factors on basal levels of DA. By contrast, main effects of Time Bin ($F[8,96] = 12.449, p < .0001$) and Social Experience ($F[1,12] = 23.317, p < .0005$), together with a significant Social Experience by Time Bin interaction ($F[8,96] = 5.418, p < .0001$), were present in the postamphetamine data. That is to say, although there were no differences in basal levels of DA, the administration of *d*-amphetamine had a much greater effect in enhancing DA levels in the isolation-reared animals than in the socially reared con-

represents twice the SED between means for the three-way interaction with factors Social Experience, Preexposure, and Trial. C: Pattern of lever press responding across conditioning sessions presented as the mean number of presses made during nine, equally sized time bins across the 40-min session. Arrows indicate the points of postconditioned stimulus (CS) shock presentation (after 10 minutes and 33 minutes). Note the reduced degree of suppression in the isolation-reared animals (■) compared to socially reared controls (○). The bar represents twice the SED between the means for the three-way interaction with factors Social Experience, Response Bin, and Trial.

Figure 5. Levels of extracellular DA in dialysates obtained from NACs of isolation-reared (■) and socially reared (□) animals under basal and *d*-amphetamine stimulated conditions. The animals were anesthetised with 0.9% halothane in O₂. *d*-Amphetamine (free base, 2 mg/kg) was administered intraperitoneally at the time point shown by the arrow. The data are expressed as picomoles per 60 μ l dialysate and are not corrected for probe recovery. Basal and postamphetamine data were analyzed separately. In each case, the bars represent twice the SED between means for the two-way interaction with factors Social Experience and Time Bin. * ($p < .001$), ** ($p < .02$), and *** ($p < .05$) denote significant differences between isolation-reared and socially reared animals (analysis of Simple Effects from the significant interaction between factors Social Experience and Time Bin).

trols. Furthermore, there were additional group differences manifest over time, in that DA levels returned to basal levels 3 hours postamphetamine in the socials, whereas levels were still elevated in the isolates. Specific post hoc comparisons confirmed these conclusions.

DISCUSSION

The main finding of the experiments was that social isolation from the weaning age in rats disrupted performance in only one of the two attentional paradigms that have been suggested as animal models related to schizophrenia. This treatment had no effect on LI but confirming previous observations (Geyer et al. 1993), reduced PPI of the acoustic startle response. Furthermore, there was evidence that the effects of social isolation on PPI were developmentally specific in that social isolation in adult rats was without effect on PPI.

The disruption of PPI following isolation rearing is consistent with the neurochemical changes in DA function produced by this manipulation, namely, enhanced effects of systemic *d*-amphetamine challenge on DA release in the ventral striatum. Both the indirectly acting agonist *d*-amphetamine and direct agonists, such

Figure 6. Distribution of dialysis probe placements within the ventral striatum. Sites were verified from sections stained with cresyl violet that were examined under the microscope. Isolation reared animals (●), socially reared animals (○). NAcc—nucleus accumbens, CPu—caudate putamen.

as apomorphine or quinpirole, have been shown to disrupt PPI, effects that can be blocked by D₂-receptor antagonists (Geyer et al. 1990; Swerdlow et al. 1991). Given the additional evidence that the disruption of PPI by isolation rearing can be reversed by treatment with nonsedative doses of the selective, D₂ antagonist raclopride (Geyer et al. 1993), it would seem probable that the perturbations in DA transmission seen in animals reared in isolation are involved, to some extent, in the impairments seen in PPI.

By contrast, the data provide no support for the involvement of ventral striatal dopaminergic mechanisms in the expression of LI. Although published results have suggested an effect of dopaminergic drugs on LI (Weiner et al. 1988; Killcross and Robbins 1993; Killcross et al. in press), these findings are not inconsistent with the present lack of effect of isolation rearing. A recent study has shown that behaviorally active doses

of *d*-amphetamine infused directly into the ventral striatum have no effect on LI in the same within-subjects conditioned-suppression LI procedure as used here (Killcross and Robbins 1993). Therefore, the lack of effect of isolation rearing might be expected, even though this treatment evidently leads to increases in DA activity in the ventral striatum. Presumably, the disruptive effects of systemic *d*-amphetamine on LI that we have also observed (Killcross and Robbins, 1993) are occurring at a site or sites other than the ventral striatum.

Although not affecting LI, isolation rearing decreased the suppression of lever pressing observed in the presence of aversive conditioned stimuli, suggesting impaired mechanisms of behavioral inhibition. This finding is worthy of note as the disinhibition produced by isolation rearing was paralleled by intraaccumbens *d*-amphetamine (Killcross and Robbins, 1993) and lesions of the ventral subiculum (Clarke et al. 1992), both of which failed to impair LI. This parallel suggests a role for altered mesolimbic DA in the behavioral disinhibition exhibited by isolates, both here, and in other situations (Morgan et al. 1977).

The effects of isolation rearing on PPI were specific in several ways. First, they were independent of any underlying baseline changes in the startle response itself, as the reduction in PPI was present at all stages of habituation to the startle stimulus. Second, the rightward shift in responsivity to the prepulse stimuli, suggestive of an elevated threshold for PPI in isolation-reared animals, was probably not due to basic sensory loss, because the response to the 120-dB startle stimulus was unaffected, whereas it would be expected to be reduced in rats with hearing impairments. Third, although isolation rearing increases general locomotor activity, this behavior cannot account for the reductions in PPI because activity within the startle apparatus (as indicated by the No-Stimulus values) was not affected by isolation, and rats isolated as adults, which were also hyperactive, did not show deficits in PPI.

The sparing of PPI in animals isolated as adults is of particular interest because it occurred, as noted above, in parallel with increases in both initial startle reactivity and spontaneous locomotor activity that were common to both isolated groups. It seems unlikely, therefore, that the differential effect of the two isolation conditions on PPI was due to the fact that 8 weeks of isolation has quantitatively less impact as a stressor when experienced as an adult. Furthermore, adults isolated for periods of up to 24 weeks continue to show normal PPI (our unpublished data). Rather, the behavioral dissociations suggest that the changes in PPI, startle, and locomotor activity were not redundant measures of the same effect of social isolation and that in young animals, this treatment exerts a qualitatively different effect on brain function than in older animals, perhaps as a result of differences in plasticity between

the developing and mature brain. With regard to the impact of isolation housing on DA function in ventral striatum, it would be of clear interest in future studies to conduct dialysis experiments on animals isolated as adults. Such studies would aid in the interpretation of the behavioral differences between isolation-reared and isolation-housed subjects in terms of neural substrates.

The differential effects of isolation rearing on PPI and LI also provide strong evidence that these two phenomena do in fact represent the expression of two quite different psychological functions. Although they have both been considered to reflect attentional mechanisms (Weiner 1990; Braff and Geyer 1990), it is clear that they are psychologically and neurally distinct. Indeed, recent analyses have considered the possibility that LI is a function of associative rather than attentional processes (Killcross et al. in press), whereas PPI has been attributed to a hypothetical "sensory gating" system (Braff and Geyer, 1990). Within this theoretical framework, common deficits in PPI in schizophrenia and in the rat reared in social isolation, perhaps related to ventral striatal dysfunction, could be seen as impairments in sensory filtering of the prepulse stimulus. Alternatively, DA-dependent ventral striatal outflow may exert descending influences on the expression and modulation of sensorimotor reflexes at the brainstem level. According to this view, neurochemical changes in the ventral striatum produced, for example, by dopaminergic drugs or as in the present case, by isolation rearing, occasion changes in state that indirectly affect the processing of prepulse stimuli, rather than modifying perception of the stimuli per se.

In summary, the present investigation directly compared two putative models of the attentional deficits seen in schizophrenia. The results indicate a developmentally specific effect of social isolation on PPI of the acoustic startle response but no effects of isolation rearing on LI. The use of social isolation in the weanling rat is particularly relevant given that it provides a non-pharmacologic, nonlesion means of perturbing ventral striatal DA function thought to be important in the neuropathology of schizophrenia. Further studies will be required to characterize possible neurochemical changes underlying the other aspects of the isolation syndrome described here. In this regard, isolation-induced alterations in serotonin transmission might be of particular significance (Jones et al. 1992).

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