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Different involvement of type 1, 2, and 3 ryanodine receptors in memory processes

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The administration of the ryanodine receptor (RyR) agonist 4-Cmc (0.003–9 nmol per mouse intracerebroventricularly [i.c.v.]) ameliorated memory functions, whereas the RyR antagonist ryanodine (0.0001–1 nmol per mouse i.c.v.) induced amnesia in the mouse passive avoidance test. The role of the type 1, 2, and 3 RyR isoforms in memory processes was then evaluated by inhibiting the expression of the three RyR proteins in the mouse brain. A selective knockdown of the RyR isoforms was obtained by the i.c.v. administration of antisense oligonucleotides (aODNs) complementary to the sequence of RyR1, RyR2 and RyR3 proteins, as demonstrated by immunoblotting experiments. RyR1 (5–9 nmol per mouse i.c.v.) knockdown mice did not show any memory dysfunction. Conversely, RyR2 (1–7 nmol per mouse i.c.v.) and RyR3 (1–7 nmol per mouse i.c.v.) knockdown animals showed an impairment of memory processes. This detrimental effect was temporary and reversible, disappearing 7 d after the end of the aODN treatment. At the highest effective doses, none of the compounds used impaired motor coordination, as revealed by the rota rod test, nor modified spontaneous mobility and inspection activity, as revealed by the hole-board test. In conclusion, the lack of any involvement of cerebral RyR1 was demonstrated. These findings also showed the involvement of type 2 and type 3 RyR in the modulation of memory functions identifying these cerebral RyR isoforms as critical targets underlying memory processes.

An increase in cytoplasmic Ca²⁺ in hippocampal neurons plays an important role in the regulation of neuronal processes, including synaptic transmission and synaptic plasticity such as long-term potentiation (LTP) and long-term depression (Nishiyama et al. 2000; Balkowiec and Katz 2002). It has also been proposed that altered Ca²⁺ regulation might play a role in brain aging and Alzheimer's disease (Gibson and Peterson 1987; Landfield 1987; Khachaturian 1989; Disterhoft et al. 1994). In brain neurons from aging rodents and rabbits, as compared with neurons from younger animals, Ca²⁺ influx associated with action potentials induces a larger Ca²⁺-dependent afterhyperpolarization (AHP) (Landfield and Pitler 1984; Moyer et al. 1992; Potier et al. 1992; Disterhoft et al. 2004; Stutzmann et al. 2006) and impairs short-term synaptic plasticity (Landfield et al. 1986; Thibault et al. 2001). Furthermore, these findings have been reinforced by studies showing that pharmacologically isolated Ca²⁺ action potentials (Pitler and Landfield 1990; Disterhoft et al. 1996), whole-cell Ca²⁺ currents (Campbell et al. 1996), and Ca²⁺ transients during repetitive spike trains also are larger in hippocampal neurons from aged animals (Thibault et al. 2001; Hemond and Jaffe 2005). An apparent excess Ca²⁺ influx via L-type voltage-gated Ca²⁺ channels (L-VGCC) appears to be involved in this Ca²⁺ dysregulation (Landfield 1996; Thibault et al. 1998).

In addition to the accumulating evidence of increased Ca²⁺ influx through L-VGCCs, there is also recent evidence that altered function of intracellular organelles might play a critical role in Ca²⁺ regulation during aging or Alzheimer's disease (Toescu and Verkhratsky 2003). In particular, changes in intracellular Ca²⁺ release from the endoplasmic reticulum appear likely to contribute to brain Ca²⁺ dyshomeostasis and have been associated with changes in [Ca²⁺]_i.

During neuronal activation, one important feature of Ca²⁺ signaling is the release of intracellular Ca²⁺ stores as a consequence of Ca²⁺ influx through voltage- or ligand-gated Ca²⁺ channels, the Ca²⁺-induced Ca²⁺ release (CICR) (Tully and Treisman 2004). The mechanism through which CICR occurs is by activation of ryanodine receptors (RyR), one of the two Ca²⁺ release channels found on the endoplasmic reticulum.

RyRs are widely expressed in mammalian brain (Giannini et al. 1995), where they play a central role in the regulation of intracellular calcium homeostasis (Simpson et al. 1995). RyRs regulate vital brain functions, including protein synthesis (Paschen et al. 1996), neurotransmitter release (Mothet et al. 1998; He et al. 2000), action potential hyperpolarization (Kawai and Watanabe 1989), and axonal retrograde transport (Breuer et al. 1992).

These receptors localize in the endoplasmic reticulum in the cell soma, dendrites, dendritic spines, and synaptic terminals (Sharp et al. 1993). Three main isoforms of the receptor (RyR1, RyR2, and RyR3) are present in brain, the RyR2 being predominant (Zalk et al. 2007). To date little is known on the role of the three neuronal RyR isoforms from behavioral studies. The aim of the present study was to investigate the involvement of RyR1, RyR2, and RyR3 in the intracellular mechanism involved in memory processes in a mouse passive avoidance task. To this purpose, we selectively inhibited the expression of each RyR subtype by using antisense oligonucleotides.

Results

Effect of RyR pharmacological modulators on memory processes

The RyR agonist 4-chloro-*m*-cresol (4-Cmc; 0.003–9 nmol per mouse intracerebroventricularly [i.c.v.], administered 30 min before the training session) increased the latency to enter the dark

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compartment in the retention session of the mouse passive avoidance test. The ANOVA on the retention session latencies revealed a significant group effect ($F_{(8,162)} = 4.506$; $P < 0.0001$). Bonferroni/Dunn multiple comparison analysis showed that mice receiving either 0.03 ($P < 0.05$) or 0.3 nmol ($P < 0.05$) per mouse i.c.v. of 4-Cmc spent significantly more time to enter the dark compartment than controls. (Fig. 1A). In these experimental conditions, diphenhydramine (20 mg/kg intraperitoneally [i.p.]) and clonidine (0.125 mg/kg i.p.), used as reference drugs, produced a statistically significant amnesic effect ($P < 0.05$ vs. control group) (Fig. 1A). Time course experiments on 4-Cmc (0.03 nmol per mouse) revealed an overall significant difference between groups ($F_{(4,75)} = 4.286$; $P < 0.001$). Bonferroni/Dunn multiple comparison analysis showed that a significant 4-Cmc-induced ameliorating effect on memory functions was obtained when administered 30 min before the training session ($P < 0.05$) (Fig. 1B).

The RyR antagonist ryanodine (0.01–1 nmol per mouse i.c.v.) was devoid of any effect when injected 15 min before the training session, as illustrated in Figure 2A. When administered immediately after the training session, an amnesic effect was obtained. The ANOVA on the retention session latencies revealed a

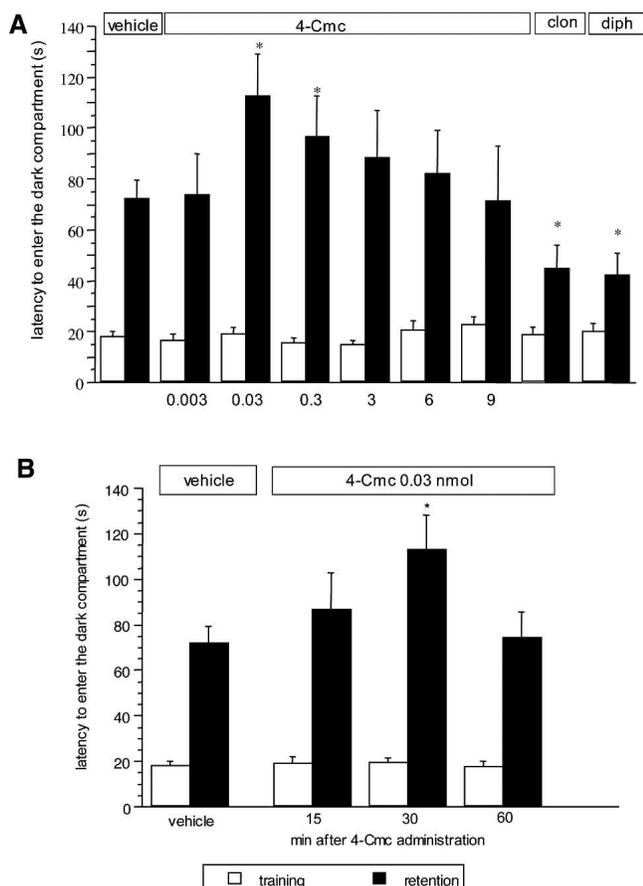


Figure 1. (A) Dose-response curve of 4-Cmc in the mouse passive avoidance test. 4-Cmc (0.003–9 nmol per mouse i.c.v.) was administered 30 min before the training session. Data are expressed as mean \pm SEM; $*P < 0.05$ in comparison with vehicle-treated mice. (Vehicle) Ethanol 0.5%; (clon) clonidine 0.125 mg/kg i.p.; (diph) diphenhydramine 20 mg/kg i.p. (B) Time-course curve of 4-Cmc in the mouse passive avoidance test. 4-Cmc (0.03 nmol per mouse i.c.v.) was administered 15, 30, or 60 min before and immediately after the training session. Data are expressed as mean \pm SEM; $*P < 0.05$ in comparison with vehicle-treated mice. (Vehicle) Ethanol 0.5%

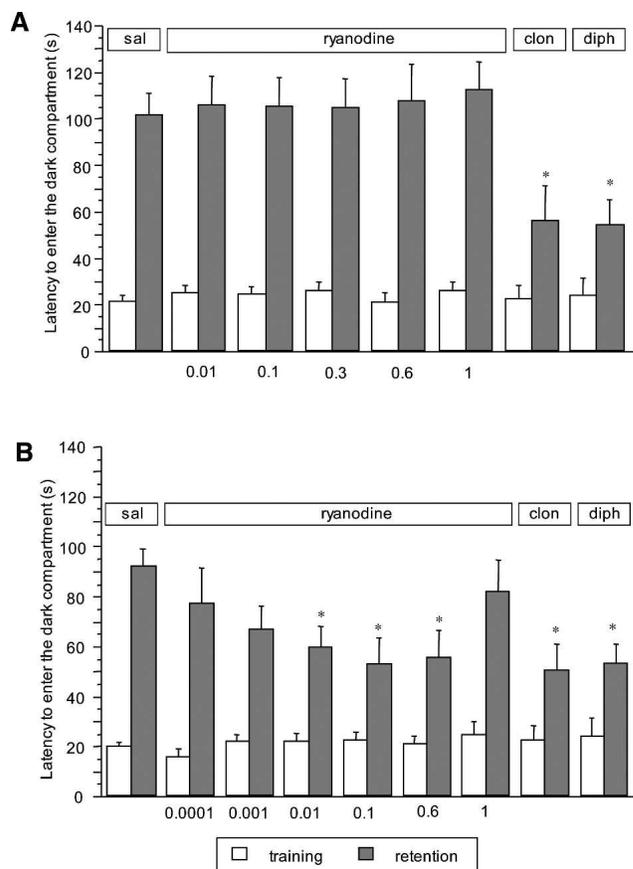


Figure 2. (A) Lack of effect of ryanodine in the mouse passive avoidance test. Ryanodine (0.01–1 nmol per mouse i.c.v.) was administered 15 min before the training session. Data are expressed as mean \pm SEM; $*P < 0.05$ in comparison with saline-treated mice; (clon) clonidine 0.125 mg/kg i.p.; (diph) diphenhydramine 20 mg/kg i.p. (B) Dose-response curve of the amnesic effect of ryanodine in the mouse passive avoidance test. Ryanodine (0.0001–1 nmol per mouse i.c.v.) was administered immediately after the training session. Data are expressed as mean \pm SEM; $*P < 0.05$ in comparison with vehicle-treated mice; (clon) clonidine 0.125 mg/kg i.p.; (diph) diphenhydramine 20 mg/kg i.p.

significant group effect ($F_{(7,135)} = 3.954$; $P < 0.001$). Bonferroni/Dunn multiple comparison analysis showed that mice receiving 0.01 ($P < 0.05$), 0.1 ($P < 0.05$) or 0.6 nmol ($P < 0.05$) per mouse i.c.v. of ryanodine spent significantly less time to enter the dark compartment than controls. The intensity of the amnesia obtained was comparable to that induced by clonidine (0.125 mg/kg i.p.) and diphenhydramine (20 mg/kg i.p.) (Fig. 2B).

The amnesia induced by ryanodine (0.1 nmol per mouse i.c.v. injected immediately after the training session) was prevented by 4-Cmc (injected 30 min before the training session) coadministration ($F_{(5,109)} = 3.941$; $P < 0.01$). Bonferroni/Dunn multiple comparison analysis confirmed the RyR-mediated mechanism of ryanodine-induced amnesia. Indeed, the detrimental effect of ryanodine was prevented by pretreatment with 4-Cmc at the dose of 9 nmol per mouse ($P < 0.05$) (Fig. 3). Both 4-Cmc concentrations used were ineffective when injected alone (Fig. 3).

4-Cmc and ryanodine did not alter the latency to enter the dark compartment in the training session in comparison with control groups as shown in Figures 1, 2, and 3.

Role of type 1, 2, and 3 RyR in memory processes

Pretreatment with an aODN against type 1 RyR (5–9 nmol per mouse i.c.v.) did not modify the entrance latency in the dark

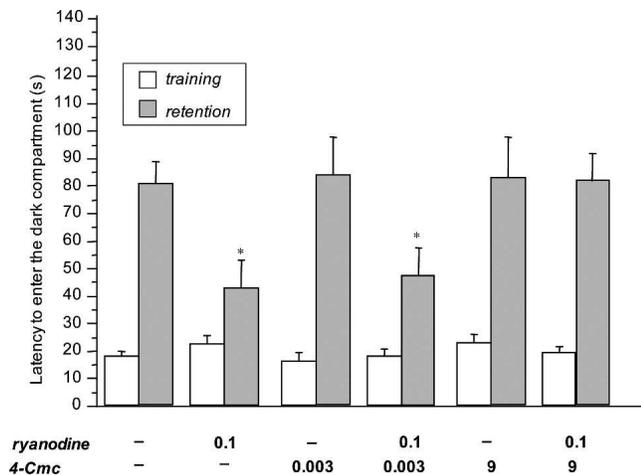


Figure 3. Reversal by 4-Cmc of the amnesia induced by ryanodine in the mouse passive avoidance test. Doses administered (nmol per mouse i.c.v.) are reported below each column. 4-Cmc and ryanodine were injected 30 min before and immediately after the training session, respectively. Data are expressed as mean \pm SEM; * $P < 0.05$ in comparison with saline-treated mice.

compartment in comparison with dODN treated mice. In the same experimental conditions clonidine (0.125 mg/kg i.p.) was able to induce amnesia ($F_{(4,85)} = 3.908$; $P < 0.01$) (Fig. 4A).

Knockdown of the type 2 RyR produced a detrimental effect on memory processes. The ANOVA on the retention session latencies revealed a significant group effect ($F_{(4,80)} = 5.141$; $P < 0.001$). Bonferroni/Dunn multiple comparison analysis showed that mice receiving 7 nmol per mouse i.c.v. showed a significant lower latency to enter the dark compartment in comparison with dODN group ($P < 0.05$). The reduction of the entrance latency in the retention session was of intensity comparable to that produced by clonidine (Fig. 4B). Similarly, the knockdown of the type 3 RyR produced a detrimental effect on memory functions ($F_{(4,83)} = 5.468$; $P < 0.001$) reaching the statistical significance at 7 nmol per mouse i.c.v. (Bonferroni/Dunn multiple comparison analysis: $P < 0.05$), as illustrated in Figure 4C.

Anti-RyR1, anti-RyR2, and anti-RyR3 did not modify the mouse entrance latency in the training session in comparison with the control groups (Fig. 4).

Effect of aODNs on RyR1, RyR2, and RyR3 protein levels

Mice were treated with the aODNs on day 1, 2, and 3. On day 4, 18 h after the last i.c.v. injection, mice were killed, and the cortex and hippocampus were dissected and examined for the protein levels of RyR1, RyR2, and RyR3 in comparison with mice treated with dODN using specific antisera developed against each RyR isoform. Figure 5A shows a representative immunoblot where three distinct prominent protein bands of ~500 kDa were observed, which display a tissue specificity of expression. The anti-RyR1 antiserum selectivity was demonstrated by a specific interaction with RyR1 proteins present in the sarcoplasmic reticulum of skeletal muscle whereas the anti-RyR2 antiserum recognized the RyR2 proteins in cardiac muscle. RyR2 and RyR3 proteins were detected in all brain areas investigated. Conversely, RyR1 isoform was not detected in mouse cortex (Fig. 5A).

Densitometric analysis of all samples revealed a statistically significant reduction of the expression of RyR1 ($P < 0.05$, Student's *t*-test), RyR2 ($F_{(2,18)} = 4.562$; $P < 0.05$) and RyR3 ($F_{(2,18)} = 4.186$; $P < 0.05$) after aODN treatment at the highest concentration used (anti-RyR1 9 nmol i.c.v.; anti-RyR2 7 nmol

i.c.v.; anti-RyR3 7 nmol i.c.v.) in comparison with mice treated with the corresponding dODN (Fig. 6B). Bonferroni/Dunn multiple comparison analysis confirmed that every aODN treatment decreased expression of the corresponding receptors in all cerebral areas investigated ($P < 0.05$) (Fig. 5B).

A dose-dependent influence on RyR expression in aODN treated animals as well as a correlation between degree of RyR subtype knockdown and severity of learning retrieval was observed. The dose of 1 nmol per mouse of each aODN did not significantly reduce the corresponding RyR protein level and was devoid of any effect on learning processes. The dose of 5 nmol per mouse reduced protein levels in anti-RyR1 (82.0 ± 6.8 , hippocampus), anti-RyR2 (76.7 ± 6.9 , cortex; 73.1 ± 7.6 hippocampus), and anti-RyR3 (80.7 ± 6.7 , cortex; 78.7 ± 5.8 hippocampus). This dose produced a reduction of the latency to enter the dark compartment in RyR2 and RyR3 knockdown mice that did not reach the statistical significance (Fig. 4B,C).

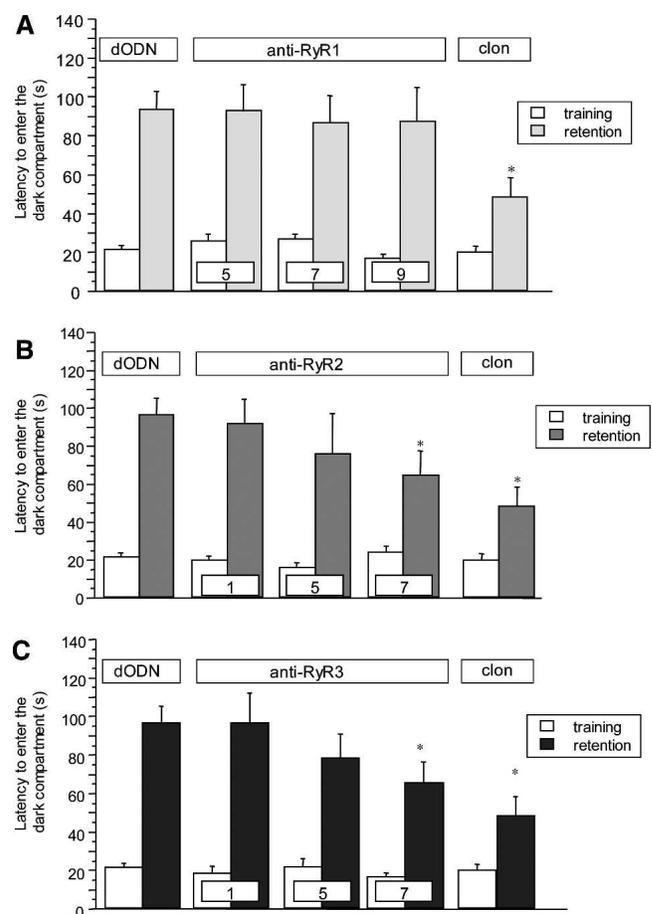


Figure 4. (A) Lack of effect of anti-RyR1 treatment (5–9 nmol per mouse i.c.v.) in the mouse passive avoidance test. Doses of anti-RyR1 administered (nmol per mouse i.c.v.) are reported in each column. Data are expressed as mean \pm SEM; * $P < 0.05$ in comparison with dODN-treated mice; (clon) clonidine 0.125 mg/kg i.p. (B) Dose-response curve of the amnesic effect of anti-RyR2 treatment (1–7 nmol per mouse i.c.v.) in the mouse passive avoidance test. Doses of anti-RyR2 administered (nmol per mouse i.c.v.) are reported in each column. Data are expressed as mean \pm SEM; * $P < 0.05$ in comparison with dODN-treated mice; (clon) clonidine 0.125 mg/kg i.p. (C) Dose-response curve of the amnesic effect of anti-RyR3 treatment (1–7 nmol per mouse i.c.v.) in the mouse passive avoidance test. Doses of anti-RyR3 administered (nmol per mouse i.c.v.) are reported in each column. Data are expressed as mean \pm SEM; * $P < 0.05$ in comparison with dODN-treated mice; (clon) clonidine 0.125 mg/kg i.p.

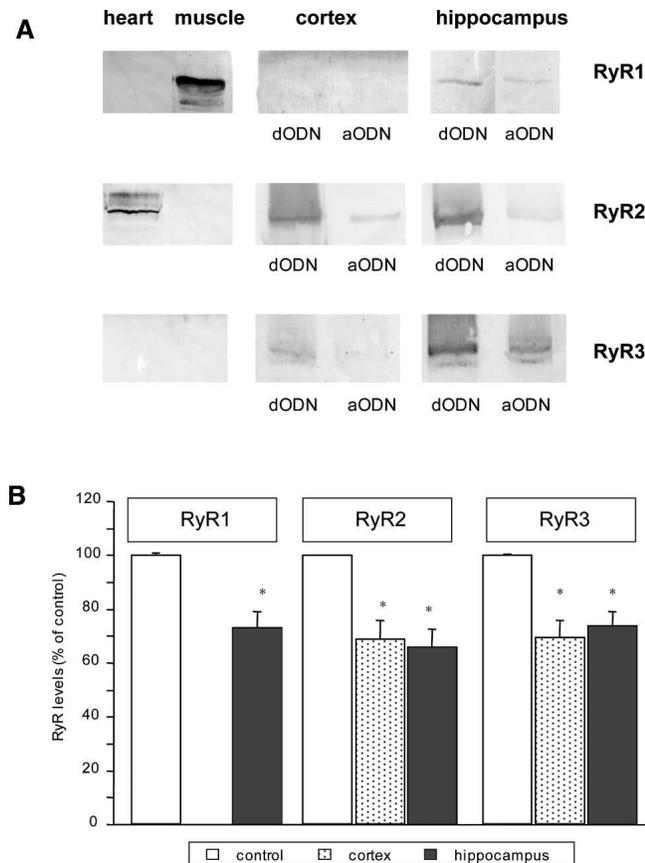


Figure 5. Reduction of RyR1, RyR2, and RyR3 protein expression in mouse muscle, heart, cortex, and hippocampus by aODN treatment in comparison with corresponding dODN-treated mice. (A) Samples (100 μ g protein/lane) of microsomal fractions of mouse cortex and hippocampus, and samples of heart (20 μ g protein/lane) and muscle (50 μ g protein/lane) were resolved on 6% SDS-PAGE, transferred to nitrocellulose, and probed with RyR1, RyR2, or RyR3 antibody. (B) Densitometric quantitation of immunoreactive protein expressed relative to control. Data are expressed as mean \pm SEM of band intensities from each of the three groups ($n = 7$ per group). * $P < 0.05$ vs. control.

The dODN did not significantly change the immunoreactivity when compared with that of naïve animals. Immunoprecipitation experiments also produced similar results (data not shown).

Immunoblots were reprobed for a protein considered to be not regulated as β -actin, and no significant density difference was revealed for this protein between samples from the RyR1, RyR2, and RyR3 downregulated brain region (data not shown).

Cross-reactivity of the primary antibodies used was excluded.

Time-dependent reversibility of RyR protein knockdown and memory impairment

To evaluate the reversibility of the aODN effects, mice were treated with the aODNs on days 1, 2, and 3; 7 d after the last i.c.v. injection, mice were killed, and the cortex and hippocampus were dissected and examined for the protein levels of RyR1, RyR2, and RyR3 in comparison with mice treated with dODN. Figure 6A shows a representative immunoblot where three distinct prominent protein bands of \sim 500 kDa, corresponding to RyR1, RyR2, and RyR3, were observed in the mouse hippocampus and cortex, with the exception of the RyR1 isoform that was not

detected in mouse cortex (Fig. 5A). RyR subtypes protein expression in aODN treated mice was similar to control animals in both brain regions.

A time-course of the amnesia obtained after aODN treatment was performed. The impairment of memory processes produced by anti-RyR2 and anti-RyR3 at the active dose (7 nmol per mouse i.c.v.) disappeared 7 d after the end of the aODN pretreatment (Fig. 6B). The anti-RyR1 treated group showed entrance latency values in the retention session comparable to control, anti-RyR2, and anti-RyR3 treated groups (Fig. 6B).

Effect of RyR pharmacological modulators and aODNs on mouse behavior

The spontaneous mobility and exploratory activity of mice pretreated with anti-RyR1 (9 nmol per mouse i.c.v.), anti-RyR2 (7 nmol per mouse i.c.v.), anti-RyR3 (7 nmol per mouse i.c.v.), ryanodine (0.1 nmol per mouse i.c.v.), and 4-Cmc (9 nmol per mouse i.c.v.) was evaluated by using the hole-board test. Both parameters in all treated groups were not impaired when compared with that of control mice. In the same experimental conditions D-amphetamine (1 mg/kg i.p.), used as the reference drug, increased both spontaneous mobility ($F_{(6,77)} = 4.343$; $P < 0.05$) and exploratory activity ($F_{(6,77)} = 3.998$; $P < 0.05$) (Fig. 7A,B).

The motor coordination of mice was unmodified by administration of the above-mentioned treatments as revealed by the rota rod test. The number of falls from the rotating rod was com-

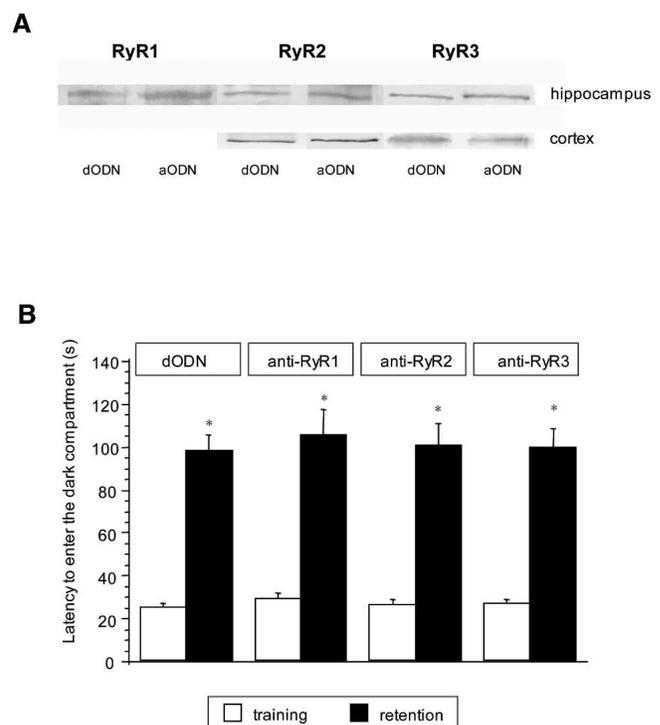


Figure 6. Reversibility of RyR1, RyR2, and RyR3 protein knockdown and memory impairment in aODN-treated mice in comparison with corresponding dODN-treated mice. (A) Samples (100 μ g protein/lane) of microsomal fractions of mouse cortex and hippocampus were resolved on 6% SDS-PAGE, transferred to nitrocellulose, and probed with RyR1, RyR2, or RyR3 antibody. (B) Lack of effect of anti-RyR1 (9 nmol per mouse i.c.v.), anti-RyR2 (7 nmol per mouse i.c.v.), and anti-RyR3 (7 nmol per mouse i.c.v.) in the mouse passive avoidance test performed 7 d after the end of the aODN repeated treatment. Data are expressed as mean \pm SEM; * $P < 0.05$ in comparison with the corresponding training session values (Student's *t*-test).

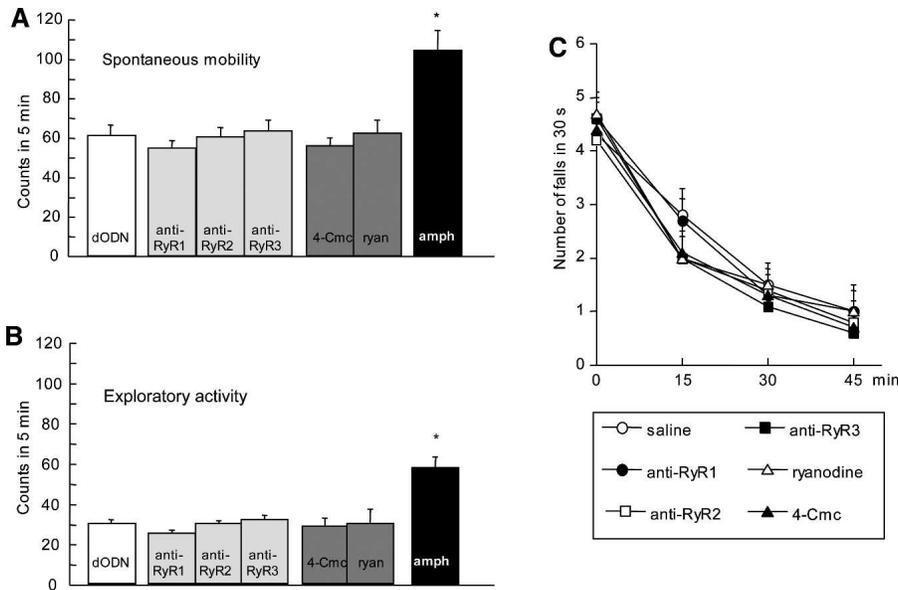


Figure 7. Lack of effect of anti-RyR1 (9 nmol per mouse i.c.v.), anti-RyR2 (7 nmol per mouse i.c.v.), anti-RyR3 (7 nmol per mouse i.c.v.), ryanodine (0.1 nmol per mouse i.c.v.), and 4-Cmc (9 nmol per mouse i.c.v.) on mouse spontaneous mobility (A) and exploratory activity (B) evaluated in the mouse hole board test, and on motor coordination (C) evaluated in the rota rod test. ODNs were administered once daily for 3 d. The test was performed 18 h after the last injection. Amphetamine (amph) was administered at the dose of 1 mg/kg s.c.; data are expressed as mean \pm SEM; * P < 0.05 vs. control group.

parable with those of control mice at every session of the test (Fig. 7C).

Discussion

The role played by the three RyR isoforms in the modulation of memory processes is illustrated by these results. A different involvement of central RyR subtypes was demonstrated. In particular, the integrity and functionality of RyR2 and RyR3 proteins appears essential to avoid memory impairments. By contrast, the RyR1 subtype, in these experimental conditions, appears not to be involved.

The effect produced by ryanodine, a RyR pharmacological antagonist, was investigated in a mouse passive avoidance task. I.c.v. administration of ryanodine induced amnesia in mice. Neurons with pharmacologically reduced slow afterhyperpolarization currents (sAHP) showed facilitated LTP. In particular, in peripheral neurons, a complete block of sAHP was reported with ryanodine (Moore et al. 1998). However, in central neurons the involvement of RyRs in the reduction of sAHP appears to be less evident. The CICR mechanism contributes less to the sAHP (Kumar and Foster 2004; Van de Vrede et al. 2007) or does not participate to generation of sAHP (Zhang et al. 1995). The ryanodine retention loss was observed only when the RyR antagonist was administered immediately after the training session. In accordance with our data, the RyR antagonist dantrolene produced a retention loss in chicks in a single trial passive avoidance task when administered immediately after training (Edwards and Rickard 2006). To confirm the role of RyRs in a mouse passive avoidance paradigm, the effect of 4-Cmc, a RyR agonist (Herrmann-Frank and Varsanyi 1993), was tested. 4-Cmc produced a dose-dependent ameliorating effect on memory processes demonstrated by an increase of the latency to enter the dark compartment in the retention session of the test. Conversely to ryanodine, this effect was obtained when 4-Cmc was administered 30 min before the training test.

This discrepancy can be explained considering that it has been observed that the antinociceptive (Galeotti et al. 2004b) and depressant-like (Galeotti et al. 2006) effects produced by 4-Cmc became significant starting from 90 min after administration. Taking into account this latency to produce its pharmacological effects, the action of 4-Cmc on memory processes might appear approximately in correspondence to the ryanodine effect. Since both RyR pharmacological modulators exerted their effect after the aversive stimulus of the training session was applied, we might hypothesize a possible action on retrieval rather than on the memory acquisition. However, given that 24 h are present between learning the task and recalling the task, any other mechanism cannot be excluded.

Ryanodine is a selective RyR antagonist unable to distinguish among RyR subtypes. To investigate the role played by the RyR isoforms, an antisense strategy was necessary since mice missing the RyR1 and RyR2 gene products die early during embryonic development. Mice carrying a targeted disruption of the RyR1 gene show complete loss of the skeletal muscle excitation-contraction coupling and die perinatally because of respiratory failure (Takeshima et al. 1995). Similar to what observed in RyR1 knockout mice, the generation of mice carrying a targeted disruption of the RyR2 gene indicates a pivotal role for the RyR2 isoform, not only in cardiac excitation-contraction coupling but also during myocardial development, as RyR2 knockout mice die at embryonic day 10 and show morphological abnormalities in the heart tub (Takeshima et al. 1998). These observations ruled out the possibility of using RyR1 and RyR2 knockout mice for behavioral studies. Mice lacking both RyR1 and RyR3 were also generated. These double-mutant mice did not actively move after birth and died, most probably from respiratory failure, as was the case for RyR1-deficient mice (Ikemoto et al. 1997).

RyRs are distributed not only in the central nervous system but also at a peripheral level. RyRs are present in the skeletal and cardiac muscle where their activation represents a crucial step for muscle contraction (Rossi and Sorrentino 2002). To avoid the possible appearance of peripheral effects that could lead to a misinterpretation of the results obtained, the aODNs used were administered directly into the cerebral ventricles.

The capability of the aODN treatment to knock down RyR protein levels was demonstrated by immunoblotting and immunoprecipitation experiments where the levels of RyR1, RyR2, and RyR3 from microsomes prepared from hippocampus and cortex of animals treated with the corresponding aODN, in comparison with the dODN-treated mice, were determined. A selective decrease of RyR1, RyR2, and RyR3 protein levels was demonstrated by using selective antibodies for each RyR isoform.

Knockdown of RyR2 produced memory disruption of intensity comparable to that induced by amnesic drugs such as the α 2-adrenoceptor agonist clonidine (Galeotti et al. 2004a) and the antihistamine diphenhydramine (Galeotti et al. 2003), indicating the importance of the integrity and functionality of the type 2 RyR isoform for the memory processes in a mouse passive avoidance task. In agreement with our results, the importance of

hippocampal microsomal RyR2 isoform in memory functions was postulated, since RyR2 mRNA and protein levels were up-regulated in the hippocampus of rats intensely trained in a water maze (Zhao et al. 2000). Present results confirm and further extend the knowledge on the crucial role played by RyR2 in the mnemonic functions.

To date, discrepant results on RyR3 involvement in memory were obtained by using RyR3 knockout mice. Futatsugi et al. (1999) noted improved spatial ability of RyR3 deficient mice. By contrast, Balshun et al. (1999) observed equivalent learning for both transgenic and wild-type animals. However, the transgenic mice demonstrated impaired ability to relearn a new target. We thought it necessary to investigate the RyR3 role by using a different experimental approach. We generated knockdown mice by using an antisense strategy. RyR3 knockdown animals showed memory impairment in the passive avoidance test demonstrated by a reduced latency to enter the dark compartment in the retention session of the test in comparison with the dODN-treated mice. In the present study, we demonstrated that the functionality of type 3 RyR is required for memory processing. In support to our results, Kouzu et al. (2000) reported that RyR3-deficient mice exhibited impairments of performance in the contextual fear conditioning test, passive avoidance test, and Y-maze learning test further confirming the role of RyR3 in memory functions. The reduction of expression of RyR3 by antisense treatment induced an amnesia of comparable intensity to that observed in anti-RyR2 treated mice and it appeared at the same concentrations. A similar involvement of these two receptor subtypes in memory functions in a mouse passive avoidance task can be postulated.

Conversely to anti-RyR2 and anti-RyR3 treated mice, RyR1 knockdown animals did not show any memory impairment in comparison with dODN-treated mice. The lack of effect of an aODN against RyR1 implies that this receptor subtype is not a major component of transduction mechanisms modulating memory functions. This observation represents the first investigation on the physiological role of cerebral RyR1. The lack of effect by the anti-RyR1 aODN in the mouse memory functions as a consequence to inefficient knockdown of the RyR1 protein by the antisense treatment can be ruled out. Western blotting and immunoprecipitation experiments revealed a reduction of the RyR1 protein levels with the highest dose administered of anti-RyR1. The different role of the three RyR isoforms in memory processes might be due the different cerebral distribution of RyR1, RyR2, and RyR3. Type 2 RyR represents the predominant isoform in the central nervous system found in widespread brain regions (McPherson and Campbell 1993; Giannini et al. 1995). Type 3 RyR has also a wide cerebral distribution including areas involved in memory processes such as hippocampus and cortex. Conversely, RyR1 has a limited distribution being absent in the cortex, as demonstrated by immunoblotting experiments, and in agreement with literature data (Giannini et al. 1995).

aODNs transiently inactivate single genes down-regulating the synthesis of the encoded protein. The reversibility of the aODN effect is fundamental to ensure that the behavioral effects observed were related to the knockdown of the target protein. The memory disruption observed in anti-RyR2 and anti-RyR3 treated animals disappeared 7 d after pretreatment with the aODNs. This return of sensitivity implies both the total reversal of aODN-induced specific inhibition of RyR proteins and a lack of damage or toxicity associated with aODN treatment. Time-course experiments on RyR knockdown mice might also support the argument that acquisition and/or consolidation processes were not altered by the treatments given the reversal of the effect with time.

Pretreatment with aODNs at the highest effective doses did

not modify the entrance latency in the training session, showing the absence of any unspecific effect produced by these pretreatments. This observation ruled out the possibility that the amnesic effect exerted by aODNs may have resulted from a sequence-independent action on cerebral structures.

As RyRs are widely distributed in the neural areas, the function of a variety of cellular receptors might be altered by pharmacological modulators as well as by the impairing effect of the antisense treatment. It has been, therefore, necessary to evaluate the possible induction of side effects by ryanodine, 4-Cmc, and the anti-RyR treatments. All compounds investigated did not modify animals' gross behavior. Additional behavioral tests were, then, performed to unmask any side effect produced by the treatments that cannot be revealed by the researcher through the observation of the animal's spontaneous behavior. In the passive avoidance test, an altered motor activity may shorten the latency to enter the dark compartment giving a false positive effect. The treatments did not impair motor coordination, as revealed by the progressive reduction of the number of falls in the rota rod test. The number of falls in this test progressively decreases since animals learn how to balance on the rotating rod. The lack of variation, or an increase in the number of falls after treatment, indicates an impairment of mice motor coordination that could lead to a misinterpretation of the results obtained in the behavioral test. Moreover, treatments did not modify spontaneous mobility and exploratory activity as indicated by the hole-board test. We can, thus, suppose that the effects observed in the passive avoidance test were not imputable to compromised behavioral paradigms. These results were of particular relevance since it has been observed that deletion of ryanodine receptor type 3 induces an increased speed of locomotion and a mild tendency to circular running (Balschun et al. 1999).

In conclusion, our results demonstrate that stimulation of RyRs ameliorates memory processes whereas a RyR blockade induces amnesia. In particular, knockdown of RyR2 and RyR3 induced a detrimental effect on memory functions indicating their fundamental role in memory processing. Conversely, RyR1 appears not to be endowed with a prominent role in the cellular events involved in learning and memory. Finally, present results indicate that a gene-specific cerebral RyR isoform level reduction by using an antisense strategy is compatible with animal life, is devoid of behavioral side effects, and represents a useful tool to investigate physiological and/or pathological central role of RyR-mediated cellular events.

Materials and Methods

Animals

Male Swiss albino mice (23–30 g) from the Morini (San Polo d'Enza, Italy) breeding farm were used. Fifteen mice were housed per cage. The cages were placed in the experimental room 24 h before the test for acclimatization. The animals were fed a standard laboratory diet and tap water ad libitum and kept at $23 \pm 1^\circ\text{C}$ with a 12 h light/dark cycle, light on at 7 a.m. All experiments were carried out in accordance with the European Communities Council Directive of 24 November 1986 (86/609/EEC) for experimental animal care. All efforts were made to minimize the number of animals used and their suffering.

Intracerebroventricular injection technique

Intracerebroventricular administration was performed under ether anesthesia, as previously described (Galeotti et al. 2006). Briefly, during anesthesia, mice were grasped firmly by the loose skin behind the head. A 0.4-mm external diameter hypodermic needle attached to a 10- μL syringe was inserted perpendicularly through the skull and no more than 2 mm into the brain of the mouse, where 5 μL were then administered. The injection site

was 1 mm to the right or left from the midpoint on a line drawn through to the anterior base of the ears. Injections were performed into the right or left ventricle, randomly. To ascertain that the drugs were administered exactly into the cerebral ventricle, some mice (20%) were injected with 5 μ L of diluted 1:10 India ink, and their brains examined macroscopically after sectioning. The accuracy of the injection technique was evaluated, and the percentage of correct injections was 95.

Passive-avoidance test

The test was performed according to the step-through method described by Jarvik and Kopp (1967). The apparatus consisted of a two-compartment acrylic box with a lighted compartment connected to a darkened one by a guillotine door. As soon as the mouse entered the dark compartment, it received a punishing electrical shock (0.5 mA, 1 sec). The latency times for entering the dark compartment were measured in the training test and after 24 h in the retention test. The maximum entry latency allowed in the training and retention sessions was, respectively, 60 and 180 sec. Between 15 and 22 mice were tested.

Hole-board test

The hole-board test was performed according to Galeotti et al. (2006). The test consisted of a 40-cm square plane with 16 flush mounted cylindrical holes (3 cm diameter) distributed four by four in an equidistant, grid-like manner. Mice were placed on the center of the board one by one and allowed to move about freely for a period of 10 min each. Two electric eyes, crossing the plane from midpoint to midpoint of opposite sides, thus dividing the plane into four equal quadrants, automatically signaled the movement of the animal (counts in 5 min) on the surface of the plane (locomotor activity). Miniature photoelectric cells, in each of the 16 holes, recorded (counts in 5 min) the exploration of the holes (exploratory activity) by the mice. Twelve mice per group were tested.

Rota-rod test

The apparatus consisted of a base platform and a rotating rod with a diameter of 3 cm and a nonslippery surface. The rod was placed at a height of 15 cm from the base. The rod, 30 cm in length, was divided into 5 equal sections by 6 disks. Thus, up to five mice were tested simultaneously on the apparatus, with a rod-rotating speed of 16 revolutions per minute. The integrity of motor coordination was assessed on the basis of the number of falls from the rod in 30 s according to Vaught et al. (1985). Those mice scoring less than three and more than six falls in the pretest were rejected (20%). The performance time was measured before (pretest) and 15, 30, and 45 min after the beginning of the test. Twelve mice per group were tested.

Antisense oligonucleotides

Phosphodiester oligonucleotides (ODNs) protected by terminal phosphorothioate double substitution (capped ODNs) against possible exonuclease-mediated degradation were purchased from Tib-Molbiol (Genoa, Italy). The sequences are the following: anti-RyR1: 5'-T*G*CCCCTGACATGTCC*C*3'; anti-RyR2: 5'-T*T*C*GCCCCATCAGCC*A*T-3'; anti-RyR3: 5'-A*G*ATGCTAATTGCATC*T*C-3' (* indicates the phosphorothioate residues). A 18-mer fully degenerated ODN (dODN), 5'-N*N*NNNNNNNNNNNNNNN*N*N-3' (where N is G, or C, or A, or T), was used as a control ODN. ODNs were vehiculated intracellularly by an artificial cationic lipid (DOTAP, Sigma) to enhance both uptake and stability. aODNs or dODNs were preincubated at 37°C for 30 min with 13 μ M DOTAP and supplied to mice by i.c.v. injection of a 5- μ L solution once daily on days 1, 2, and 3.

Preparation of membranes

Mouse brains were dissected to separate specific areas. Mouse hippocampus and cortex were homogenized in an homogenization buffer containing 25 mM Tris-HCl at pH 7.5, 25 mM NaCl, 5 mM EGTA, 2.5 mM EDTA, 2 mM NaPP, 4 mM PNEF, 1 mM

Na₂VO₄, 1 mM PMSF, 20 μ g/mL leupeptin, 50 μ g/mL aprotinin, 0.1% SDS. The homogenate was centrifuged at 9000g for 15 min at 4°C, and the low-speed pellet was discarded. The microsomal membranes were obtained from the supernatant of the 9000g spin by centrifugation at 100,000g for 1 h at 4°C. Microsomes were resuspended in homogenization buffer and stored at -80°C. Protein concentration of the microsomal fraction was quantified using a protein assay kit (Bio Rad Laboratories). Skeletal muscle microsomes were prepared from the posterior leg of mice. Cardiac microsomes were prepared from the entire heart, cleaned from large vessels.

Western blot analysis

Membrane homogenates (100 μ g) made from cerebellum, hippocampus and cortex regions of control and antisense-treated mice and membrane homogenates from skeletal and cardiac muscle were separated on 6% SDS-PAGE and transferred onto nitrocellulose membranes (180 min at 80 V) using standard procedures. Membranes were blocked in PBST (PBS containing 0.1% Tween) containing 5% nonfat dry milk for 90 min. Following washings, blots were incubated overnight at 4°C with specific antibodies against RyR1, RyR2, or RyR3 (1:3000 dilution), a kind gift of Professor Sorrentino (University of Siena, Italy). After being washed with PBS containing 0.1% Tween, the nitrocellulose membrane was incubated with goat anti-rabbit horseradish peroxidase-conjugated secondary antisera (1:10,000) and left for 1 h at room temperature. Blots were then extensively washed according to the manufacturer's instruction and developed using enhanced chemiluminescence detection system (Pierce). Exposure and developing time used was standardized for all the blots. Densitometric analysis of scanned images was performed on a Macintosh iMac computer using the public domain NIH Image program. Measurements in control samples were assigned a relative value of 100%.

Immunoprecipitation of RyR1, RyR2, RyR3

Immunoprecipitation was carried out on 1 mL of microsomal fraction containing 200 μ g of proteins by incubation for 2 h at 4°C with 10 μ g of specific antibodies against RyR1, RyR2, or RyR3. All tubes then received 20 μ L of 25% (v/v) Protein G-Agarose (Santa Cruz Biotechnology) and incubated for further 2 h at 4°C. Pellets were collected by centrifugation at 1000g for 5 min at 4°C, washed three times with homogenization buffer. Pellets were finally resuspended in 40 μ L of electrophoresis sample buffer and boiled for 5 min, and samples were processed by SDS-PAGE.

Drug administration

The following drugs were used: ryanodine, 4-chloro-*m*-cresol (Calbiochem); diphenhydramine hydrochloride, D-amphetamine hydrochloride (De Angeli, Rome, Italy); clonidine hydrochloride (Sigma). 4-Chloro-*m*-cresol (4-Cmc) was dissolved in 0.5% ethanol. All other drugs were dissolved in isotonic (NaCl 0.9%) saline solution. Drug and ODNs concentrations were prepared in such a way that the necessary dose could be administered in a volume of 5 μ L by i.c.v. injection or 10 mL/kg by subcutaneous (s.c.) or i.p. administration.

To perform time-course experiments, 4-Cmc was administered 15, 30, or 60 min before and immediately after the training session, and ryanodine was administered 15 min before and immediately after the training session. In ryanodine time-course experiments, longer latencies from the training session (30, 60 min) were not used, since, conversely to 4-Cmc, it is endowed with a short lasting effect, as observed in previous behavioral experiments performed in our laboratory (Galeotti et al. 2004b, 2006). From time-course experiments the ideal time of administration of 4-Cmc (30 min before training) and ryanodine (immediately after training) was chosen to perform dose-response curves. The same administration times were used when 4-Cmc and ryanodine were coadministered. Clonidine and diphenhydramine, used as reference drugs, were administered, respectively, 70 min before and immediately after the training session.

ODNs were i.c.v. injected once daily on days 1, 2, and 3. Behavioral tests were performed on day 4, 18 h after the last i.c.v. injection of ODNs, and on day 11. Brain areas to conduct Western blotting experiments were collected on day 4, 18 h after the last i.c.v. injection and on day 11.

Statistical analysis

All experimental results are given as the mean \pm SEM. Student's *t*-test or a two-way ANOVA followed by Bonferroni/Dunn post-hoc test was used for statistical analysis; *P*-values of less than 0.05 were considered significant.

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