**Inhibition of Mitochondrial P53 Abolishes the Detrimental Effects of Social Isolation on Ischemic Brain Injury**

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**Background and Purpose**—Social isolation (SI) increases stroke incidence and delays poststroke recovery. Women may be at greater risk from the negative consequences of SI, but few studies have examined both sexes in experimental models, and none have evaluated the effects of isolation initiated after stroke. The effects of poststroke SI in men and women were examined, and the role of mitochondrial P53 was evaluated.

**Methods**—C57Bl6 mice were pair-housed (PH; male and ovariectomized female) for 2 weeks, subjected to stroke and then assigned to a housing condition (isolated or PH). The effects of housing on infarct volume and recovery were examined. Changes in Bcl-2 and mitochondrial p53 were assessed by Western blot. A mitochondrial p53 inhibitor (pifithrin-μ) was given to mice of both sexes.

**Results**—Compared with pair-housed mice, poststroke SI significantly increased infarct size in both sexes; SI mice also had worse neurological deficits. The detrimental effects of SI paralleled increases in mitochondrial p53 levels. Pharmacological inhibition of mitochondrial p53 using pifithrin-μ abolished the detrimental effects of SI and reduced cell death.

**Conclusions**—Poststroke SI results in increased ischemic injury in both sexes. The effect of housing on infarct was more pronounced in women. Targeting the mitochondrial P53 pathway could minimize the detrimental effects of isolation after stroke. (Stroke. 2014;45:3101-3104.)

**Key Words:** apoptosis ■ ischemia ■ reperfusion injury ■ sex-specific ■ stroke
P53 Inhibitor Treatment
Pifithrin-μ (Calbiochem, CA) was dissolved in dimethylsulfoxide and diluted to 4% in saline (2 mg/kg) and a final volume of 100 μL/10 g body weight of drug or vehicle (4% dimethylsulfoxide in saline) was injected intraperitoneally to randomized mice at 3, 24, and 48 hours after stroke. Pifithrin-μ is a selective and specific mitochondrial inhibitor as determined previously in vivo and in vitro. Binding studies have shown that pifithrin-μ binds to p53. The dose used was determined by pilot experiments and from the literature; a low dose was selected to avoid nonselective infarct reduction in all groups with a higher dose. This dose did not completely abolish p53 levels to sham levels but reduced it to levels comparable with PH stroke mice.

Neurological Deficit Scores, Infarct Analysis, and Survival Rates
After 72 hours of reperfusion, the neurological deficit scores (NDSs) were recorded by a blinded investigator as in Ref. 6 and 7. Infarcts were quantified from 2,3,5-triphenyltetrazolium chloride–stained coronal sections as detailed previously.6,7 Mice that died during reperfusion were included in mortality rates, and if paired, the partner was not used for analysis.

Open Field Analysis
Spontaneous locomotor activity was performed during the light phase of the circadian cycle, between 9:00 AM and 12:00 PM under normal fluorescent room lights. For testing, mice were acclimatized to the room conditions and were individually placed in the open field chamber (15”×15”) equipped with 16 infrared beam emitting LEDs on each side for a duration of 20 minutes. The total number of beam breaks was automatically collected by a computer-operated PAS Open Field system (San Diego Instruments, San Diego, CA).6 The open field chambers were cleaned after each individual test session using 70% ethanol. All animals were assessed by a blinded investigator.

Western Blots
An additional cohort of mice was euthanized at 24 hours after stroke for protein analysis. Brains were homogenized, and a portion of lysate was used for whole cell lysate analysis. The mitochondrial fraction was obtained as described previously.7 Protein levels were assessed for mitochondrial P53 (1:200; abcam) and Bcl-2 (1:500; cell signaling) was used for whole cell lysate analysis. The mitochondrial fraction was obtained as described previously.7 Protein levels were assessed for mitochondrial P53 (1:200; abcam) and Bcl-2 (1:500; cell signaling) using actin and cytochrome c oxidase (COX IV; 1:2000; abcam) as loading controls. Densitometry was performed with ImageJ software.

Results
Infarct Analysis
Poststroke SI significantly exacerbated infarct size compared with PH animals both in males (Figure 1A; P<0.05; t test) and females (Figure 1B; P<0.001; t test). Equivalent blood flow reduction was seen by laser Doppler in both sexes and in both housing conditions. A significant 3-way interaction between SI, infarct size, and sex was also found F(1,44)=4.2, P<0.05, suggesting that the detrimental effect of housing on infarct is more pronounced in ovariectomized females.

NDS and Mortality
The detrimental effect of SI was also reflected in the NDS. SI males had significant worsening of their NDS (Figure 1C; P<0.05) as did females (Figure 1D; P<0.05) compared with PH. A higher mortality was also seen in SI male (29%) and female (36%) mice compared with PH male (7%) and female (0%) mice.

Protein Analysis
Analysis of variance for mitochondrial P53 protein levels revealed a significant effect of stroke in males (Figure 2B; P<0.05) and females (Figure 2D) both by stroke (P<0.05) and by housing (P<0.05). Bcl-2 levels were significantly elevated after stroke in both housing conditions, but this increase was less in SI mice (Figure 2B and 2D; P<0.05) and in SI females (P<0.05) compared with PH mice. Moreover, a significant strokehousing interaction was seen in both sexes (P<0.05), suggesting that stroke-induced increases in p53 and Bcl-2 expression was significantly altered in SI mice versus PH mice.

![Figure 1](http://stroke.ahajournals.org/)

Figure 1. Social isolation (SI) mice had significantly increased infarct size after stroke in males (A) and in females (C). Values are mean±SEM. At 72 h after stroke, neurological deficit scores were higher in SI mice, suggesting poorer recovery compared with pair-housed (PH) mice in males (B) and in females (D), data presented as box-and-whisker plot. *P<0.05, **P<0.01 indicates statistically significant differences.
Pifithrin-μ Treatment Reverses SI Effects

Pifithrin-μ (2 mg/kg) abolished the detrimental effects of SI on infarct (Figure 3A) but had no neuroprotective effect in PH mice. Two-way analysis of variance yielded a significant effect of housing ($F(1,42)=17.1, P<0.05$) and a significant effect of drug ($F(1,42)=18.7, P<0.05$), and a significant interaction between housing and drug ($F(1,42)=10.9, P=0.002$) in males (Figure 3A) and also in females (Figure 3B). The beneficial effect of P53 inhibition was also reflected in the restoration of spontaneous locomotor activity after stroke in SI males. Two-way analysis of variance yielded a significant effect of housing, $F(1,42)=4.49, P<0.05$, and a significant effect of drug, $F(1,42)=4.38, P<0.05$, and a significant interaction between housing and drug, $F(1,42)=5.88, P<0.05$. These findings indicate that drug abolished the detrimental effects of isolation.

Discussion

In this study, we found that poststroke SI significantly worsens stroke damage in both sexes, and this effect can be abolished by inhibition of mitochondrial P53 activation. SI before stroke worsens outcomes in experimental models, as well as in clinical populations, in both sexes.2,6,8,9 However, much less is known about the effects of SI initiated after injury. As most patients who are isolated do not come to medical attention until after the injury occurs, the ability to manipulate poststroke housing environments has broader translational significance to functional recovery. Stroke is a sexually dimorphic disease. Several molecular pathways have been identified as important contributors to cell death after ischemic injury, and these pathways, although not completely distinct, are differentially regulated by sex. Women are more sensitive to caspase-dependent cell death, whereas caspase-independent or poly[ADP-ribose] polymerase-1–mediated cell death predominates in men.4,14 Both these cell death signaling pathways converge on mitochondrial permeability transition pore dysfunction.10,11 This work suggests that mitochondrial stress underlies some of the detrimental effects of SI. Other mechanisms may also contribute such enhancement of inflammation or late effects on cerebral blood flow.

In this work, we found that SI significantly increased cell death after ischemic stroke in both sexes, although women were more susceptible to the detrimental effects of isolation. Although statistically significant, this interaction between isolation, infarct, and sex requires further study to determine its physiological relevance. Both sexes responded to a p53
inhibitor, with a reversal in isolation-induced brain injury. Future studies are needed to specifically examine poststroke depressive phenotypes and cognitive deficits after SI in both sexes, using chronic end points and aged animals. As these deficits often lead to nursing home placement in stroke survivors, especially elderly women, the effect of social factors on stroke recovery deserves investigation.

Conclusions
Our findings suggest that mitochondrial association of p53 is an important underlying mechanism for SI-enhanced ischemic injury in both sexes. These findings demonstrate that poststroke SI has detrimental effects on ischemic injury. Consistent with clinical studies, men seem to have a greater susceptibility to the negative effects of isolation.

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Disclosures
None.

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