Therapeutic Strategy for the Prevention of Pseudorabies Virus Infection in C57BL/6 Mice by 3D8 scFv with Intrinsic Nuclease Activity

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3D8 single chain variable fragment (scFv) is a recombinant monoclonal antibody with nuclease activity that was originally isolated from autoimmune-prone MRL mice. In a previous study, we analyzed the nuclease activity of 3D8 scFv and determined that a HeLa cell line expressing 3D8 scFv conferred resistance to herpes simplex virus type 1 (HSV-1) and pseudorabies virus (PRV). In this study, we demonstrate that 3D8 scFv could be delivered to target tissues and cells where it exerted a therapeutic effect against PRV. PRV was inoculated via intramuscular injection, and 3D8 scFv was injected intraperitoneally. The observed therapeutic effect of 3D8 scFv against PRV was also supported by results from quantitative reverse transcription polymerase chain reaction, southern hybridization, and immunohistochemical assays. Intraperitoneal injection of 5 and 10 μg 3D8 scFv resulted in no detectable toxicity. The survival rate in C57BL6 mice was 9% after intramuscular injection of 10 LD50 PRV. In contrast, the 3D8 scFv-injected C57BL/6 mice showed survival rates of 57% (5 μg) and 47% (10 μg). The results indicate that 3D8 scFv could be utilized as an effective antiviral agent in several animal models.

INTRODUCTION

The pseudorabies virus (PRV) has a broad host range, infecting most mammals and some avian species. PRV infection frequently proves fatal in young piglets and other susceptible species, and infected animals tend to die from central nervous system disorders (Enquist, 1999; Field and Hill, 1974; McCracken et al., 1973; Pensaert and Kluge, 1989). Infection in the natural host is characterized by epidermal lesions, usually on the mucosal surface, with viral spread to the nervous system and the establishment of latent neuronal infections (Ono et al., 2004). As the PRV has inflicted particularly heavy damage on swine livestock farming and the husbandry industry, research into vaccines against PRV has progressed relatively rapidly, but only a few antiviral agents against the virus have been developed thus far (Steigerman et al., 1997). As a novel approach, the DNA vaccine method has been employed, and this method elicits an enhanced general immune response in livestock including swine, in which the levels of interleukin-4 and interferon-gamma increase markedly (Dory et al., 2005; Gerdz et al., 1997; Hong et al., 2002). Several approaches using antiviral agents for DNA replication inhibitors or viral reverse transcriptase inhibitors have also been used in PRV containment efforts. Although generally cost-effective, these antiviral reagents work only in a limited viral spectrum, and their efficacies are generally more prone to viral mutation (Ferraris and Lina, 2008; Morfin and Thouvenot, 2003).

Through collaboration with several research groups, we have developed the 3D8 single chain variable fragment (scFv) catalytic antibody fragment, which exhibits non-specific nuclease activity on both DNA and RNA (Kim et al., 2006). Importantly, 3D8 scFv exhibits antiviral effects against herpes simplex virus (HSV), pseudorabies virus (PRV), classical swine fever virus (CSFV), murine norovirus (MNV), 2 geminiviruses, 5 tobamoviruses and cucumovirus (Hoang et al., 2014; Jun et al., 2010; Lee et al., 2013a; 2013b; 2014). We also observed caveolae-mediated endocytosis, by which the exogenously treated 3D8 scFv protein (10 μM) was internalized and detected in the cell within 2 h and eventually in the nucleus after an extended observation period (Jang et al., 2009).

In a continued effort to develop and refine an effective antiviral agent based on 3D8 scFv, we evaluated the antiviral effects of intraperitoneally (i.p.) injected 3D8 scFv in PRV-infected C57BL/6 mice. The results of our biochemical and cytological analyses demonstrated that 3D8 scFv could be utilized as an effective antiviral agent. Our study also demonstrates that 3D8 scFv can be delivered to target tissues via i.p. injection, and we determined the threshold dosage of the injection that does not
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Materials and Methods

Cells and viruses
Rat C6 glioma and B16F10 mouse melanoma cells were maintained in DMEM media supplemented with 10% fetal bovine serum (Hyclone, USA), 100 U/ml of penicillin-streptomycin (Hyclone), and non-essential amino acids (Sigma, USA), at 37°C in a 5% CO₂ atmosphere. The PRV-YS strain was obtained from the National Veterinary Research and Quarantine Service (NVRQS), Korea (Yoon et al., 2008). The C6 and B16F10 cell cultures were infected with PRV at a multiplicity of infection of 0.1, and then incubated for 1 h in a humidified CO₂ incubator at 37°C.

Mice
Eight-week-old female C57BL/6 mice (Dae Han Bio Link Co., Ltd., Korea), weighing 18-22 g each, were maintained under standard laboratory conditions (tap water, constant room temperature of 22°C). All animal procedures performed in this study (permit number: KU15006) were reviewed, approved, and supervised by the Institutional Animal Care and Use Committee of Konkuk university.

Antibodies
Anti-PRV gD monoclonal antibody was purchased from Jeno Biotech Inc. (Korea). Preparation of the anti-3D8 scFv polyclonal antibody has been described previously (Lee et al., 2013a).

Expression and purification of the 3D8 scFv protein
E. coli strain BL21 DE3 (pLysE) cells containing plg20-3D8 were grown in 1 L of LB broth with 100 μg/ml ampicillin and 20 μg/ml chloramphenicol at 37°C to express the recombinant 3D8 scFv protein (Fig. 1A). When the A₆₀₀ of the culture reached 0.8, IPTG (0.5 mM) was added to induce protein expression. The culture supernatant was obtained by centrifugation at 10,000 × g for 10 min at 4°C and filtered through a 0.45-μm membrane. The 3D8 scFv protein was purified from the filtered supernatant using an IgG-Sepharose column (Amersham Pharmacia, USA). The column was washed with 20 bed volumes of PBS and then with two volumes of 5 mM ammonium acetate (pH 5.0). The 3D8 scFv protein was eluted with 0.1 M acetic acid (pH 3.4) in fractions of 1.5 ml each. The eluate in the fractions was neutralized to pH 7.0 with 0.1 volume of 1 M Tris-base (pH 9.5). The 3D8 scFv protein was analyzed by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions. The 3D8 scFv protein was eluted with 0.1 M acetic acid (pH 3.4) in fractions of 1.5 ml each. The eluate in the fractions was neutralized to pH 7.0 with 0.1 volume of 1 M Tris-base (pH 9.5). The 3D8 scFv protein was analyzed by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions. The 3D8 scFv protein was eluted with 0.1 M acetic acid (pH 3.4) in fractions of 1.5 ml each. The eluate in the fractions was neutralized to pH 7.0 with 0.1 volume of 1 M Tris-base (pH 9.5). The 3D8 scFv protein was analyzed by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis under reducing conditions. The 3D8 scFv protein was eluted with 0.1 M acetic acid (pH 3.4) in fractions of 1.5 ml each. The eluate in the fractions was neutralized to pH 7.0 with 0.1 volume of 1 M Tris-base (pH 9.5).

ssDNA and dsDNA catalytic activity test with the scFv protein
The assay reaction was performed in assay buffer (20 mM Tris-HCl, pH 8.0, 50 mM NaCl, 5 mM MgCl₂). The DNA and RNA binding test was performed dependent on reaction time. DNA and RNA (0.25 μg each) were mixed with 0.2 μg purified scFv protein and BSA, and samples were collected after 0, 1, 2, 3, 4, and 5 h. Agarose gel electrophoresis was performed in 1.0% agarose gel and stained with ethidium bromide.

Immunocytochemistry
Confocal microscopy was conducted as described previously (Jang et al., 2009). Cells on coverslips were washed in PBS and fixed for 10 min in 4% paraformaldehyde in PBS at room temperature. The cells were permeabilized with Perm-buffer (1% BSA, 0.1% saponin, and 0.1% sodium azide in PBS) for 10 min at room temperature (RT). After 1 h of blocking with 3% bovine serum albumin in PBS, 3D8 scFv-treated cells were incubated with rabbit anti-3D8 scFv antibody, followed by incubation with TRITC anti-rabbit Ig. Nuclei were stained with DAPI during the last 10 min of incubation at RT. Cells on coverslips were mounted in Vectashield anti-fade mounting medium (Vector Labs, USA) and observed with a Zeiss LSM 510 laser confocal microscope (Carl Zeiss, Germany) followed by analysis with Carl Zeiss LSM imaging software.

FITC labeling of the 3D8 scFv protein
FITC labeling of 3D8 scFv was conducted using an AnaTag™ 5-FITC microscale protein labeling kit (Anaspec, USA) according to the manufacturer’s instructions. The reaction mixture was labeled with 5-FITC at room temperature for 1 h followed by purification with a spin column (Banks and Paquette, 1995; Brinley, 1992). C57BL/6 mice were injected i.p. at 10 μg/20 g
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(per mouse) with 3D8 scFv-FITC. The organs (liver, muscle, lung, and brain) of each mouse were collected at 0, 20, 40, 60, 120, and 180 min. Total proteins were prepared using Pro-prep solution (Intron, Korea) and measured at 494 nm via fluorometer (Hybrid Multi-Mode Microplate Reader, BioTeck Inst. Inc., USA).

Grouping and viability test
The C57BL/6 mice were divided into seven groups (M, A-F; 21 mice in each group). Group M was a PBS-injected mock group. Groups A-C were infected via intramuscular (i.m.) injection with strain PRV-YS at the LD$_{50}$. After PRV infection, groups A-C were treated i.p. with PBS, 5 μg 3D8 scFv, and 10 μg 3D8 scFv, respectively. Groups D-F were infected i.m. with the PRV strain at 10x the LD$_{50}$. Following PRV infection, groups D-F were treated i.p. with PBS, 5 μg 3D8 scFv, and 10 μg 3D8 scFv, respectively. All groups were treated with PBS and 3D8 scFv protein at 12 h intervals for 3 days. After each group was treated with PRV and 3D8 scFv, they were observed at 12 h intervals to determine the number of dead animals, and muscle and brain were harvested and stored at -70°C for further analyses.

RNA preparation and quantitative real-time reverse transcription polymerase chain reaction (RT-PCR)
RNAs were isolated from the liver, muscle, lung, and brain tissues of C57BL/6 mice using Corezol™ reagent (Corebio System, USA) after viral challenge (Haimov-Kochman et al., 2006). cDNA was synthesized from 5 μg total RNA using random hexamers and Moloney murine leukemia virus reverse transcriptase (SuperBio). All primers were designed with the Primer 3 program (Supplementary Table S1). Expression of the PRV gpD transcript was evaluated via RT-PCR. Expression of C-X-C motif chemokine 10 (CXCL10) and inducible nitric oxide synthase (iNOS) (chemokine-related genes) was assayed via qRT-PCR using a Rotor-Gene 3000 (Corbett Research, Australia). The relative expression of chemokine mRNAs was calculated after normalization to the GAPDH gene using the Delta Ct method (Jung et al., 2013). Each qRT-PCR was representative of data from three replicate experiments. Data represent the average of three individual experiments and error bars indicate standard errors (SEs).

Immunohistochemical staining
Representative 3-μm thick brain tissue sections were mounted on silane-coated slides for immunohistochemical analysis as described previously by Ramos-Vara (2005). Anti-PRV gpD antibody (Jeno Biotech Inc., Korea), which was used as the primary antibody (1:100) in this study, was administered for 60 min at RT. The samples were treated for color development using an iVIEW DAB Detection Kit (Ventana Medical Systems, USA).

Statistical analysis
All analyses were conducted using the GraphPAD Prism program (GraphPAD Software, USA). One-way analysis of variance and Tukey’s post-hoc t-tests were employed for statistical analyses. Data are expressed as mean ± SE. A p < 0.05 was considered significant.

RESULTS
Purification of 3D8 scFv protein and catalytic affinity assay
The purified scFv protein appeared as a single band at approximately 32 kDa on a 10% acrylamide gel (Figs. 1A and 1B). In Limulus Amebocyte Lysate (LAL) assay (PYROGENT™ 25 single tests 0.125 EU/ml sensitivity, Switzerland), the endotoxin concentration was not detected in test tubes which 3D8 scFv protein was added (Values of 3D8 scFv endotoxinity were <0.125 EU ml$^{-1}$). To investigate nuclease activity, DNA and RNA were purified with 3D8 scFv protein. A catalytic affinity assay was performed dependent on reaction time. When BSA as a negative control was treated to different types of nucleic acids (dsDNA, ssDNA, dsRNA, ssRNA), any nucleic acid was not hydrolyzed (Fig. 1C). However, when ssDNA and ssRNA were incubated with endotoxin-free 3D8 scFv, ssDNA and ssRNA were degraded from 1 h after 3D8 scFv treatment. In contrast to ssDNA and ssRNA, dsDNA and dsRNA were gradually degraded by 3D8 scFv as times went on. Taken together, we confirmed the catalytic activity of 3D8 scFv against all nucleotide types even though the hydrolyzing activity was slightly different among the types of nucleotides in time-dependent manner.

Blood chemistry analysis for 3D8 scFv in C57BL/6 mice
Blood samples were collected from mice injected with endotoxin-free 3D8 scFv and toxicity was assayed by measuring several representative parameters, followed by comparison with the samples collected from the control mice. The levels of aspartate aminotransferase and creatine phosphokinase increased significantly upon treatment with 100 μg 3D8 scFv relative to the PBS treatment group used as a control. In contrast, the levels of alkaline phosphatase and glucose decreased upon treatment with 20-100 μg 3D8 scFv relative to that measured in the controls (Supplementary Fig. S1 and Supplementary Table S2). Taken together, the results that i.p. injection of 20-100 μg 3D8 scFv caused toxic effects; therefore, treatments of 3D8 scFv in all subsequent experiments were administered at concentrations well below this threshold (5-10 μg).

Diffusion of FITC-labeled 3D8 scFv in the organs of C57BL/6 mice and verification of 3D8 scFv delivery to target cells
The relative quantities of endotoxin-free 3D8 scFv were measured in muscle, liver, lung, and brain tissues after the initial injection to determine if i.p. injected 3D8 scFv penetrates to target tissues. FITC-labeled 3D8 scFv protein was injected i.p. at 10 μg/20 g mouse. Total protein from each of the tissue samples was subsequently prepared at time intervals of 0, 20, 40, 60, 120, and 180 min, and the levels of FITC-3D8 scFv were measured by flow cytometry at an excitation wavelength of 494 nm (Fig. 2A). The 3D8 scFv protein was detected in all tissues 20 min after injection, and the protein level peaked at 60 min post-injection (20-30 fluorescence density). The protein level maintained at a fluorescence density of approximately 15-20 for up to 120 min after injection. These findings demonstrate that the injected 3D8 scFv protein was transported through the bloodstream to reach target tissues, and that was stably maintained in these cells for at least a short time (Fig. 2A). This experiment was conducted using different mice, and the same results were consistently observed in all cases.

Actual penetration of 3D8 scFv was monitored in vivo using the B16F10 and C6 cell lines derived from mouse epithelia and glioma, respectively. Exogenously treated 3D8 scFv protein was detected in the cytoplasm of the cell lines approximately 48 h later (Fig. 2B). The results of these in vitro cell culture and in vivo mice experiments indicate that 3D8 scFv could transport to systemically each organ of C57BL/6 mice and 3D8 scFv could penetrate into cells (B16F10 and C-6).
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Fig. 2. The 3D8 single chain variable fragment (scFv) protein penetrates tissues (liver, muscle, lung and brain) and cells (B16F10 and C6). (A) FITC-labeled 3D8 scFv protein was injected intraperitoneally at 10 μg/20 g mouse into C57BL/6 mice. The organs (liver, muscle, lung and brain) of all mice were collected at 0, 20, 40, 60, 120, and 180 min, and measured at 494 nm via flow cytometry. Fluorescence density (a.u.: arbitrary unit) of three independent trials is shown. (B) Penetration of 3D8 scFv into cultured cells (B16F10 and C6) was evaluated by immunocytochemistry with confocal microscopy. 3D8 scFv was visualized by TRITC (rhodamine)-labeled anti-3D8 polyclonal scFv antibody. Upper panels: PBS treatment, bottom panels: 3D8 scFv treatment.

3D8 scFv-injected C57BL/6 mice survive exposure to PRV-YS
The effect of 3D8 scFv as an antiviral agent against PRV in a mouse model was evaluated at protein injection concentrations of 5 and 10 μg (Fig. 3A). The virus was inoculated i.m. in parallel with an i.p. injection of 3D8 scFv. The mice generally began to exhibit clinical signs of illness 3-5 days post-challenge. Approximately 57% of the group inoculated with the LD₅₀ of the virus without 3D8 scFv (group A) survived, whereas the groups treated with the 3D8 scFv at a concentration of 5 μg (group B) or 10 μg (group C) evidenced survival rates of 81% and 76%, respectively, at the same lethal viral dose. Survival rate dropped significantly to 9% in the group that did not receive 3D8 scFv (group D) when inoculated with a much higher viral titer (10× LD₅₀). However, the 3D8 scFv-treated groups had survival rates of 57% (5 μg; group E) and 47% (10 μg; group F) (Fig. 3B). A typical symptom of PRV infection in each group is shown in Fig. 3C. Lesions in the femoral muscles of mice in group D were induced as a result of viral infection, leading to death of the mice, which represented typical symptoms of the disease (Fig. 3C).

The 3D8 scFv protein has antiviral therapeutic effects
Viral replication and gene and protein expression levels were measured in each group using RT-PCR and immunohistochemistry to confirm that the observed difference in survival rate after treatment with 3D8 scFv was the result of reduced viral pathogenicity. The RT-PCR analyses conducted on RNA isolated from the groups described in Fig. 4 showed that the viral transcript glycoprotein D (gpD) was readily detected in the RNA samples isolated from dead mice in the groups treated with 3D8 scFv (groups E and F). However, no gpD transcripts were detected in samples isolated from survivors of the same groups (Fig. 4A). Similarly, the gpD protein, as measured by immunohistochemistry, was detectable only in the mice that died from viral infection, whereas the protein was not detected in any of the surviving mice, regardless of their 3D8 scFv treatment status (Fig. 4B).

The antiviral effect is linked to 3D8 scFv catalytic activity
The expression levels of genes involved in the inflammation response were assessed via qRT-PCR with RNA samples obtained from the groups described in Fig. 3A to determine whether the antiviral mechanisms of 3D8 scFv include any indirect effects such as triggering of endogenous antiviral responses. As demonstrated in Fig. 5, the levels of two transcripts associated with the inflammation response, CXCL10 and iNOS, were detected at elevated levels in the PRV-infected samples (groups D-F). However, no significant differences in the expression levels of these transcripts were observed between the survivors and dead among these groups, thereby confirming the absence of any indirect effect of the antiviral mechanism exploited by 3D8 scFv (Figs. 5A and 5B).

DISCUSSION
A number of studies have developed effective antiviral agents, including DNA vaccines, viral DNA replication inhibitors, and viral neuraminidase inhibitors (Dory et al., 2005; Erlich and Mills, 1985; Fox et al., 2006; Koziel and Peters, 2007; Laskin, 1984; Le et al., 2005). 3D8 scFv can be distinguished from these antiviral agents, as it can digest the viral genome itself with no sequence specificity. Thus, this agent may be the vanguard of a new paradigm in antiviral therapeutics, in which the host is protected against a broad spectrum of viruses, including DNA and RNA viruses. Using the findings of prior studies, we determined that 3D8 scFv induces a series of antiviral effects against RNA viral genomes at the cellular level; thus, implicating 3D8scFv as a potentially valuable antiviral therapeutic (Jun et al., 2010). In this study, we extended our ongoing research by
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Fig. 3. 3D8 scFv-treated C57BL/6 mice demonstrate high survival rates against pseudorabies virus (PRV) infection. (A) Schematic diagram for PRV and 3D8 scFv challenge protocol. M: Mock-treated group; A, B, and C: Groups challenged with PRV at LD50; D-F: Groups challenged with 10 LD50. Different amounts of 3D8 scFv administered to each group, indicated at right. (B) Viability of mouse groups showing correlation with the amount of 3D8 scFv treatment. (a) Effect of 3D8 scFv on the survival rate of LD50 PRV infection groups (A: PBS, B: 5 μg 3D8 scFv, C: 10 μg 3D8 scFv) and (b). Effect of 3D8 scFv on the survival rate of 10 LD50 PRV infection groups (D: PBS, E: 5 μg 3D8 scFv, F: 10 μg 3D8 scFv). Results obtained from 21 mice per group (M group: eight mice) for up to 5 days after PRV infection. (C) Varying degrees of symptom development after PRV infection in the mice groups, as described in (A). Dissection picture of group D shows the most severe dermatitis formation, as indicated by red arrows.

Fig. 4. 3D8 scFv protein exerts antiviral therapeutic effects. (A) Absence of viral transcript expression (pseudorabies virus [PRV] gpD) in mice treated with the 3D8 scFv, as shown by reverse transcription-polymerase chain reaction analysis results. RNA samples from various organs (liver, muscle, lung, and brain) of different mouse groups were analyzed for PRV gpD expression. M: Mock-treated group; D challenged with 10 LD50 PRV followed by PBS treatment; E challenged with 10 LD50 PRV followed by 5 μg 3D8 scFv treatment; F challenged with 10LD50 PRV followed by 10 μg 3D8 scFv treatment. Both surviving (O) and dead (X) mice were analyzed for PRV gpD expression in each group. Data from surviving mice in groups E and F are provided in (a) and data from the dead mice in groups E and F are shown in (b). (B) Absence of viral protein expression (PRV gpD) in mice treated with 3D8 scFv as detected via immunohistochemistry. Brain cerebellum tissues of M, D, and E mouse groups were stained with anti-PRV monoclonal antibody visualized with biotinylated anti-mouse secondary antibody followed by DAB staining. Magnification: 100× for upper panel, 400× for bottom panel. Presence of PRV gpD protein (shown as brown staining) is detectable only in the Purkinje layer cells of the group D mice.
fragment, such as 3D8 scFv, to different organs and tissues subsequent to i.p. injection is likely to reflect a more general phenomenon.

The effect of 3D8 scFv on viral titer was evaluated in different mouse organs infected with PRV alone or coupled with 3D8 scFv treatments. In the groups with high survival rates following viral infection due to the effects of 3D8 scFv (e.g., group E and F), no viral protein PRV-gpD was detectable in any of the organs of the surviving mice. In contrast, the viral protein was detected in dead individuals of the same group, but only in muscle and brain and not in liver or lung tissues. These results suggest that the injected 3D8 scFv was transmitted to virtually all parts of the mouse, conferring resistance against the virus in all organs examined herein. This finding also reflects that transmission of PRV principally occurs through the nervous system, and little viral replication actually occurs within the liver and lung. The finding of abundant PRV within the muscle tissues is probably attributable to the initial i.m. introduction of the virus. Our observed path of PRV transmission is consistent with the findings by Breshears et al. (2005), in which Saimiriine herpesvirus 1 infected via i.m. injection translocated to the central nervous system.

The expression levels of the genes involved in the inflammation pathway were monitored to determine whether the viral genome was destroyed directly by the nuclease activity of 3D8 scFv, or indirectly by turning on the endogenous antiviral responses of the host cell. Tight correlations between induction of the genes involved in the inflammation response and increased resistance against viral infection have been reported in a number of studies. iNOS and CXCL10 transcript levels are maintained at high levels in cells infected with several viruses, including herpes simplex virus and the influenza virus (Akaike et al., 1996; Daigle and Carr, 1998; Haas et al., 1999; Hsieh et al., 2006; Su et al., 1996). Therefore, we assayed the levels of iNOS and CXCL10 transcripts in cells inoculated with PRV; thus, allowing us to determine whether treatment with 3D8 scFv altered the expression levels of these transcripts in these cells. As shown in Fig. 5, although PRV-infected cells generally showed much higher levels of these transcripts relative to controls, 3D8 scFv treatment resulted in no additional changes in the transcript levels in virally-infected cells. Also, when we treated the inhibitor of type I interferon receptor as a mediator of intrinsic antiviral activity to Hantaan virus (HTNV)-infected HEK and 3D8 scFv transgenic cells, no differences were observed between presence or absence of inhibitor in 3D8 scFv transgenic cells (data not shown). Thereby, it is demonstrate that the observed antiviral effects of 3D8 scFv were induced directly by the nuclease activity of 3D8 scFv. (A) The model shows the pseudorabies virus (PRV) replication cycle. Entry begins with attachment of the virus particle to the cell surface. The viral DNA enclosed by nucleocapsid delivers into nucleus. Viral DNA synthesis begins after the expression of the beta proteins and the temporal program of viral gene expression ends with the appearance of the gamma or late proteins, which constitute the structural proteins of the virus. Finally, the virus initiates a lytic cycle. (B) Mice were infected intraperitoneally with purified 3D8 scFv protein. The treated 3D8 scFv proteins penetrated the cell via a caveolae-mediated endocytosis mechanism. Consequently, 3D8 scFv proteins are localized in cytoplasm. Viral DNA is degraded by RNase activity and inhibit the expression of the viral proteins. The RNase activity of 3D8 scFv exhibits the therapeutic effects against PRV infection.
the activities of 3D8 scFv itself.

In conclusion, as an extension of our previous studies in which the antiviral effect of 3D8 scFv was demonstrated by in vitro experiments, we further showed that nucleic activity of 3D8 scFv could be effectively employed in vivo as an antiviral therapeutic agent using the mouse model system. Purified 3D8 scFv protein from E. coli was i.p. injected into PRV infected mice and localized to the cytoplasm by caveloic-mediated endocytosis (Jun et al., 2010). Then, the PRV transcriptome was degraded by 3D8 scFv with intrinsic RNase activity in cytoplasm (Fig. 6). The finding that 3D8 scFv nuclease activity acts on both DNA and RNA demonstrates the utility of 3D8 scFv as a multifaceted, broad-spectrum antiviral agent that works against DNA and RNA viruses.

Note: Suplementary information is available on the Molecules and Cells website (www.molcells.org).

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