Improved development of mouse somatic cell nuclear transfer embryos by chlamydocin analogues, class I and IIa histone deacetylase inhibitors†

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Abstract

In mammalian cloning by somatic cell nuclear transfer (SCNT), the treatment of reconstructed embryos with histone deacetylase (HDAC) inhibitors improves efficiency. So far, most of those used for SCNT are hydroxamic acid derivatives—such as trichostatin A—characterized by their broad inhibitory spectrum. Here, we examined whether mouse SCNT efficiency could be improved using chlamydocin analogues, a family of newly designed agents that specifically inhibit class I and IIa HDACs. Development of SCNT-derived embryos in vitro and in vivo revealed that four out of five chlamydocin analogues tested could promote the development of cloned embryos. The highest pup rates (7.1–7.2%) were obtained with Ky-9, similar to those achieved with trichostatin A (7.2–7.3%). Thus, inhibition of class I and/or IIa HDACs in SCNT-derived embryos is enough for significant improvements in full-term development. In mouse SCNT, the exposure of reconstructed oocytes to HDAC inhibitors is limited to 8–10 h because longer inhibition with class I inhibitors causes a two-cell developmental block. Therefore, we used Ky-29, with higher selectivity for class IIa than class I HDACs for longer treatment of SCNT-derived embryos. As expected, 24-h treatment with Ky-29...
up to the two-cell stage did not induce a developmental block, but the pup rate was not improved. This suggests that the one-cell stage is a critical period for improving SCNT cloning using HDAC inhibitors. Thus, chlamydocin analogues appear promising for understanding and improving the epigenetic status of mammalian SCNT-derived embryos through their specific inhibitory effects on HDACs.

**Summary sentence**

Chlamydocin analogues, a novel family of inhibitors specific for class I and IIb HDACs, significantly improved the ability of mouse SCNT-derived embryos to produce offspring.

**Key words:** cloned embryo, histone deacetylase, histone deacetylase inhibitor, mouse, somatic cell nuclear transfer.

**Introduction**

Cloning by somatic cell nuclear transfer (SCNT) is a technique that produces embryos or animals that are genetically identical to the donor cells. Since the birth of Dolly the sheep derived from an adult mammary gland cell, SCNT has been heralded as a promising assisted reproductive technology with a broad range of applications in regenerative medicine, livestock industries, and pharmaceutical manufacturing [1–5]. Recent advancements in gene-editing technologies, especially the CRISPR/Cas9 system, have further enhanced the use of SCNT in livestock animals including pigs, goats, cattle, and sheep for the production of useful phenotypes such as disease resistance and high meat/milk production [6]. Meanwhile, there have been many attempts to increase the efficiency of SCNT, especially in mice. Cloned mouse embryos, fetuses, and placenta have provided invaluable information on epigenetic aberrations associated with SCNT, and their corrections lead to significant improvements in cloning efficiency [3, 7]. However, many of the effective treatments require genetically modified donor cells or further micromanipulation of reconstructed embryos [8–11]. In this respect, one of the most feasible measures that has been shown to improve mouse cloning is the treatment of SCNT-derived embryos with histone deacetylase (HDAC) inhibitors. HDACs are a family of enzymes involved in the regulation of a number of cellular processes including the control of gene expression through histone acetylation. Therefore, HDAC inhibitors are expected to induce an open chromatin structure of the donor cell-derived genome, leading to efficient transcription of developmentally related genes, especially transcription factors [12, 13]. The first HDAC inhibitor used successfully for mouse SCNT was trichostatin A (TSA), a naturally derived hydroxamate, which has a broad spectrum for inhibiting different HDAC families [14]. Mouse SCNT-derived embryos treated with TSA showed a high potency to develop into blastocysts in vitro and to term offspring in vivo, with pup rates reaching more than 5% per embryos transferred [15]. TSA was also reported to be effective for SCNT cloning in bovines [16, 17], pigs [18, 19], and rabbits [20, 21].

HDACs are grouped into four classes based on structural and functional similarities. The class I isoforms (HDAC1, −2, −3, and −8), class IIa (HDAC4, −5, −7, and −9), and class IIb (HDAC6 and −10) are Zn-dependent enzymes, whereas class III isoforms (SIRT1 to 7) are NAD⁺-dependent [22]. HDAC inhibitors can also be characterized as pan-HDAC inhibitors or selective HDAC inhibitors according to their selectivity for specific classes. In mouse cloning experiments, it is desirable to use HDAC inhibitors that have selective inhibitory activities for the most efficient and safe SCNT outcomes. However, only hydroxamates—typical pan-HDAC inhibitors including TSA [15, 23], scriptaid [24], suberoylanilide hydroxamic acid (SAHA) [25], oxamflatin [25], m-carboxycinnamic acid bishydroxamide [26], and PXD101 (belinostat) [27]—have been proven to increase mouse cloning efficiencies.

Besides hydroxamates, there are HDAC inhibitors that belong to other molecular families such as short-chain fatty acids, cyclic peptides, and benzamide [28], but none of them have been used effectively for improving mouse SCNT cloning. HDAC inhibitors generally exhibit strong binding to HDACs. For example, TSA is thought to chelate the zinc ion in the active site in a bidentate fashion and binds strongly to HDAC through an aromatic “Cap” structure [29]. Chlamydocin is also a naturally occurring HDAC inhibitor with a cyclic tetrapeptide structure that can bind strongly to HDACs. Based on the structure of chlamydocin, a series of chlamydocin analogues containing various metal-binding groups have been developed as potent HDAC inhibitors [30–32]. These synthetic compounds inhibited class I and IIa HDACs, but not IIb HDACs, at nanomolar concentrations [31] (Figure 1). FK228 (romidepsin), an anti-T-cell lymphoma agent that also has a cyclic tetrapeptide structure (Figure 1), is known to inhibit class I HDACs (HDAC1 and HDAC2) [33]. Here, we examined whether the development of mouse SCNT-derived embryos could be improved by treatment with these newly developed chlamydocin analogue HDAC inhibitors and FK228, which have more selective inhibitory effects than the hydroxamate HDAC inhibitors such as TSA.

**Materials and methods**

**Animals**

Eight- to 10-week-old B6D2F1 (C3HBL/6 × DBA/2 hybrid) female and male mice were purchased from Japan SLC (Shizuoka, Japan) and used for the collection of oocytes and donor cumulus cells. Eight- to 10-week-old C3HBL/6N male mice purchased from Japan SLC were used for in vitro fertilization (IVF). ICR strain female mice (CLEA Japan, Inc., Tokyo, Japan), 8–12 weeks old, were used as embryo transfer recipients. The animals were housed under a controlled lighting condition (daily light 07:00–21:00 h) and were maintained under specific pathogen-free conditions. All animal experiments were approved by the Animal Experimentation Committee at the RIKEN Tsukuba Institute and were performed in accordance with the committee’s guiding principles.

**Somatic cell nuclear transfer**

SCNT was carried out as described previously [34, 35]. Briefly, B6D2F1 female mice were superovulated by the injection of 7.5 IU of equine chorionic gonadotropin (Sankyo Yell Yukuhin, Co., Tokyo, Japan) and 7.5 IU of human chorionic gonadotropin (hCG; ASKA Pharmaceutical Co., Ltd., Tokyo, Japan) at a 48-h interval.
Figure 1. HDAC inhibitory activity and the structure of HDAC inhibitors used in this study. The HDAC inhibitory activities (IC₅₀ values) of HDAC inhibitors were measured as described in the Materials and methods or are quoted from references [30–33, 57].

At 15 h after hCG injection, the mice were euthanized by cervical dislocation and cumulus-oocyte complexes were collected from the oviducts. Cumulus cells were removed by suspending the complexes in potassium-modified simplex optimized medium (KSOM) [36] containing 0.1% bovine testicular hyaluronidase. The oocytes were enucleated in Hepes-buffered KSOM containing 7.5 μg/ml cytochalasin B (CB) using a piezo-driven micropipette (PMM-150FU; Primetech, Ibaraki, Japan). After culture in KSOM under 5% CO₂ in air at 37°C for at least 1 h, the enucleated oocytes were injected with donor cumulus cells using a piezo-driven micropipette in Hepes-buffered KSOM without bovine serum albumin (BSA). After culture in KSOM for about 1 h, injected oocytes were incubated in Ca²⁺-free KSOM containing 2.5 mM SrCl₂ and 5 μg/ml CB with or without a HDAC inhibitor (TSA, chlamydocin analogues, or FK228) for 1 h. Next, the oocytes were incubated in KSOM containing 5 μg/ml CB and a HDAC inhibitor for 5 h followed by incubation in KSOM containing a HDAC inhibitor alone (where appropriate) for 2 h. After washing, the SCNT-derived embryos were cultured in KSOM under 5% CO₂ in air at 37°C. The embryos reached the two-cell stage after 24 h in culture and were transferred into the oviducts of recipient females, as described above. The pregnant females were euthanized at 19.5 dpc and examined for fetuses and placentas in their uteri. In some experiments, SCNT-derived embryos treated with Ky-9 were additionally treated with 10 μg/ml vitamin C and 0.4% deionized BSA for 8 h [37] (Supplementary Figure S1). The embryos that reached the two-cell stage after 24 h in culture were transferred into the oviducts of recipient females, as described above. The pregnant females were euthanized at 19.5 dpc and examined for fetuses and placentas in their uteri.

In vitro fertilization
IVF-derived embryos were obtained as described previously [38]. Briefly, cumulus-oocyte complexes were collected from the oviducts of B6D2F1 female mice that had been superovulated as described above and were transferred into human tubal fluid (HTF) medium at 37°C under 5% CO₂ in humidified air. Spermatozoa were collected from the cauda epididymis of 12-week-old C57BL/6 male mice and preincubated in HTF medium at concentrations of 200–400 spermatozoa/μl. After 6 h of culture, pronuclear-stage embryos were transferred into the oocyte culture medium at concentrations of 200–400 spermatozoa/μl. After 6 h of culture, pronuclear-stage embryos were transferred into KSOM and used for immunostaining as described below.

Intracytoplasmic sperm injection
Intracytoplasmic sperm injection (ICSI)-derived pups were obtained as reported [39]. Briefly, cumulus-free oocytes prepared as described above were each injected with an epididymal sperm head in Hepes-buffered KSOM without BSA. The injected oocytes were cultured in

| Compound | HDAC1 (Class I) IC₅₀ (μM) | HDAC4 (Class Iia) IC₅₀ (μM) | HDAC6 (Class Iib) IC₅₀ (μM) |
|----------|---------------------------|-----------------------------|-----------------------------|
| Ky-2     | 0.0087 ± 0.0006           | 0.020 ± 0.002               | 0.16 ± 0.01                 |
| Ky-9     | 0.038 ± 0.004             | 0.028 ± 0.002               | >100                        |
| Ky-29    | 1.4 ± 0.2                 | 0.36 ± 0.06                 | 35.9 ± 17.2                 |
| Ky-72    | 1.8 ± 0.33                | 1.6 ± 0.2                   | 19.3 ± 6.6                  |
| Ky-309   | 0.0058                    | 0.0045                      | 0.26                        |
| FK228    | 0.036 ± 0.016             | 0.510 ± 0.340               | 14.000 ± 3.100              |
| TSA      | 0.0060 ± 0.0025           | 0.038 ± 0.004               | 0.0086 ± 0.0014             |
Production of parthenogenetic embryos

Metaphase II (MII) oocytes were collected from B6D2F1 female mice as described above. They were activated by treatment with Ca^{2+}-free CZB medium containing 2.5 mM SrCl_{2} and 5 μg/ml CB for about 15 min and then transferred into CZB medium containing 5 μg/ml CB with or without an HDAC inhibitor for 6 h followed by incubation in CZB medium containing an HDAC inhibitor alone (where appropriate) for 2 h. The embryos were washed and cultured in CZB medium as described above until analysis.

Immunostaining

Histone acetylation levels of SCNT-derived or IVF-derived embryos treated with HDAC inhibitors were analyzed by immunostaining. The embryos were fixed in 100% ethanol (FUJIFILM Wako Pure Chemical Corporation, Osaka, Japan) at −20°C for 20 min and permeabilized with phosphate-buffered saline (PBS) containing 0.2% Triton X-100 (Merck KGaA) at room temperature (RT) for 1 h. After blocking with 3% BSA in PBS containing 0.2% Triton X-100 for 1 h, they were incubated with primary antibodies (anti-acetyl-Histone H3 antibody; Merck Millipore, Darmstadt, Germany) in PBS containing 30 μg/ml BSA at 4°C overnight. After washing with PBS, the embryos were reacted with secondary antibodies (Anti-rabbit IgG-Alexa 555; Thermo Fisher Scientific) at RT for 1 h. The embryos were mounted on glass slides in Vectashield Mounting Medium with DAPI (Vector Laboratories, Burlingame, CA). The slides were imaged using a CellVoyager CV1000 confocal scanner system (Yokogawa Electronic, Tokyo, Japan) and fluorescence intensity was measured using image processing software [Image; https://imagej.net/Welcome].

Transcriptional activity assay

For detecting nascent RNA synthesis during zygotic gene activation (ZGA) at the two-cell stage, diploid parthenogenetic embryos were generated as above and treated with or without an HDAC inhibitor (Ky-9, Ky-309, TSA, or FK228). Transcription was detected by Click-iT RNA Alexa Fluor 594 Imaging Kits (Thermo Fisher Scientific) according to the manufacturer’s instructions. Briefly, parthenogenetic embryos at 5 h after activation were cultured in CZB medium containing 2 mM 5-ethyluridine (EU) until 24 h after activation (19 h exposure). They were fixed with 4% paraformaldehyde (FUJIFILM Wako Pure Chemical Corporation) and permeabilized with 0.2% Triton X-100 in PBS containing 30 μg/ml BSA. The parthenogenetic embryos were then incubated in the Click-iT reaction cocktail at RT for 30 min and washed with the Click-iT reaction rinse buffer. Fluorescence images of EU were detected using a CellVoyager CV1000 confocal microscope scanner system with consistent laser strength and fluorescence intensity was measured using ImageJ.

Statistical analysis

The developmental rates of embryos (Tables 1, 2, and 4; Supplementary Table S1 and Figure S1) were analyzed using Fisher exact probability test. Immunofluorescence (Figure 2) and body and placental weights (Figures 2 and 3; Table 3) were analyzed using Turkey test for multiple comparisons between groups. P values <0.05 were considered statistically significant.
Table 1. Development in vitro and in vivo (morula/blastocyst transfer) of SCNT-derived embryos treated with HDAC inhibitors

| HDAC inhibitor | Concentration of inhibitor (nM) | No. of embryos cultured | No. (%) of two-cell embryos at 24 h | No. of four-cell embryos at 48 h (% per 2 cells) | No. of morula/blastocyst at 72 h (% per 2 cells) | No. of embryos transferred (no. of recipients) | No. of implantations (%) | No. of cloned offspring (%) | No. of placentas only (%) |
|----------------|---------------------------------|-------------------------|-----------------------------------|-----------------------------------------------|-----------------------------------------------|---------------------------------------------|--------------------------|---------------------------|--------------------------|
| Control        | –                               | 530                     | 484 (91.3)                         | 382 (78.9)                                    | 271 (56.0)                                    | 210 (14)                                    | 83 (39.5)                | 0 (0)                     | 3 (1.4)                  |
| TSA            | 5                               | 64                      | 49 (76.5)                         | 45 (91.8)†                                    | 32 (65.3)                                    | 15 (1)                                      | 6 (40.0)                 | 1 (6.7)                   | 0 (0)                    |
| Ky-2           | 100                             | 63                      | 52 (82.5)                         | 45 (86.5)†                                    | 31 (59.6)                                    | 31 (2)                                      | 7 (22.6)†                 | 1 (3.2)                   | 1 (3.2)                   |
| Ky-9           | 1600                            | 69                      | 57 (82.6)                         | 44 (77.2)†                                    | 28 (49.1)                                    | 28 (2)                                      | 19 (67.9)†                 | 2 (7.1)†                   | 3 (10.7)†                 |
| Ky-29          | 2000                            | 61                      | 58 (88.5)                         | 51 (94.4)†                                    | 37 (68.5)                                    | 37 (2)                                      | 21 (56.8)†                 | 1 (2.7)†                   | 4 (10.8)†                 |
| TSA            | 5                               | 66                      | 60 (90.9)                         | 35 (58.3)†                                    | 25 (2)                                       | 10 (40.0)                                   | 0 (0)                     | 1 (4.0)                   | 0 (0)                    |

Table 2. Development in vitro and in vivo (two-cell transfer) of SCNT-derived embryos treated with HDAC inhibitors

| HDAC inhibitor | Concentration of inhibitor (nM) | No. of embryos cultured | No. (%) of two-cell embryos at 24 h | No. of embryos transferred (no. of recipients) | No. of implantations (%) | No. of offspring (%) | No. of placentas only (%) |
|----------------|---------------------------------|-------------------------|-----------------------------------|---------------------------------------------|--------------------------|---------------------|--------------------------|
| Control        | –                               | 487                     | 416 (85.4)                         | 213 (8)                                     | 93 (43.7)                | 6 (2.8)             | 6 (2.8)                  |
| TSA            | 5                               | 153                     | 133 (86.9)                         | 109 (6)                                     | 69 (63.3)†               | 8 (7.3)             | 3 (2.8)                  |
| Ky-2           | 100                             | 153                     | 142 (92.8)                         | 137 (6)                                    | 55 (40.1)                | 3 (2.2)             | 3 (2.2)                  |
| Ky-9           | 3200                            | 232                     | 203 (87.5)                         | 135 (6)                                    | 78 (57.8)†               | 9 (6.7)             | 2 (1.5)                  |
| Ky-29          | 2000                            | 335                     | 292 (87.2)                         | 217 (9)                                    | 85 (39.2)                 | 3 (1.4)             | 3 (1.4)                  |
| TSA            | 5                               | 113                     | 106 (93.8)                         | 89 (84.0)                                   | 61 (57.5)                 | 21 (1)              | 4 (19.0)†                 |
| Ky-29          | 0.5                             | 288                     | 63 (94.0)                          | 2 (3.2)†                                    | 0 (0)                     | N.D.                | N.D.                     |

<ref>P < 0.05 (vs. control, Fisher exact probability test).

NR, not recorded.<ref>

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Developmental ability of SCNT-derived embryos treated with chlamydocin analogues

Next, we examined whether these analogues could improve the in vitro developmental ability of SCNT-derived embryos. In addition to the concentrations above, lower concentrations were also tested for some analogues. Compared with untreated SCNT-derived embryos (controls), only Ky-29 at the higher concentration (2000 nM) significantly increased both four-cell and morula/blastocyst formation rates (*P < 0.05; Table 1). By contrast, Ky-309 at the higher concentration (100 nM) significantly decreased both four-cell and morula/blastocyst formation rates (*P < 0.05; Table 1). FK228 induced a strong developmental arrest of SCNT-derived embryos at the two-cell stage and no morulae were obtained (Table 1). In other experimental groups including controls, about 80 and 50% of embryos (per two-cell stage) developed into four-cell and morulae/blastocyst stages, respectively (Table 1). After transfer of these morulae/blastocysts into recipient female mice, at least one cloned offspring was born at term, except for the control group, and in the lower concentration Ky-9 and Ky-309 groups. Of these, the higher concentration Ky-9, lower concentration Ky-29, and higher concentration Ky-72 and Ky-309 treatment groups resulted in significantly higher pup rates than in the control group. Except for Ky-309, which seemed to have a loosened chromatin structure, reflecting a more reprogrammable genomic state following SCNT.
embryonic toxicity, Ky-9 gave the best pup rate (7.1%), similar to that of TSA (6.7%).

Our results indicate that these novel chlamydocin analogues can have beneficial effects on the in vivo development of SCNT-derived embryos. To further examine this possibility, we undertook embryo transfer experiments using two-cell embryos. As expected, Ky-9 at 1600 nM gave the highest pup rate (7.2%), which was significantly higher than in control (2.8%, \( P < 0.05 \); Table 2). We then tested a higher concentration of Ky-9 (3200 nM). This treatment did not further increase the pup rate of SCNT-derived pups, but it was still
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Figure 3. Transcriptional activity assay at the two-cell stage in parthenogenetic embryos treated with FK228 and other HDAC inhibitors. (A) Scheme of the transcriptional activity assay using parthenogenetic embryos. 5-EU was added to the medium at 5 h after activation and the embryos were cultured until 24 h. Nascent RNA transcription during the major zygote gene activation (ZGA) phase at the two-cell stage was measured by incorporation of EU detected by specific fluorescence. (B) Representative images of fluorescence specific for incorporated EU in control and HDAC inhibitor-treated parthenogenetic embryos. Bar = 20 μm. (C) Relative fluorescent intensity for incorporated EU (see Materials and methods). Each dot represents one nucleus (two nuclei per embryo). The fluorescence level of the FK228-treated embryos was significantly lower that of the embryos treated with other HDAC inhibitors (P < 0.0001 by Tukey test).

Table 3. Body and placental weights of pups derived from HDAC inhibitor-treated SCNT-derived embryos

|                      | Control | Ky-2  | Ky-9  | Ky-29 | Ky-72 | Ky-309 | TSA  |
|----------------------|---------|-------|-------|-------|-------|--------|------|
| No. of offspring and placentas | 4       | 5     | 25    | 13    | 3     | 2      | 7    |
| Body weight at term (g ± SD) | 1.49 ± 0.18 | 1.64 ± 0.19 | 1.39 ± 0.29 | 1.41 ± 0.27 | 1.78 ± 0.16 | 1.61 ± 0.21 | 1.57 ± 0.15 |
| Placental weight at term (g ± SD) | 0.36 ± 0.09 | 0.41 ± 0.09 | 0.30 ± 0.11 | 0.36 ± 0.06 | 0.40 ± 0.04 | 0.31 ± 0.05 | 0.28 ± 0.07 |

There were no significant differences between the HDAC inhibitor groups and the control group for each parameter (Tukey multiple comparisons test).

High (6.7%, P = 0.075 vs. control; Table 2). This result indicates that the embryotoxicity of Ky-9 was low even at a concentration higher than the IC50 (see Figure 1). The pup rates obtained with Ky-9 were similar to that produced with TSA (7.3%, P = 0.059 vs. control), although TSA promoted the development of SCNT-derived embryos at a much lower concentration (5 nM) than Ky-9. We also tested an intermediate concentration of Ky-9 (320 nM) and found that the effect was modest (4.6% pup rate; Table 2).

Cloned pups obtained following HDAC inhibitor treatment commenced respiration soon after Caesarian section and showed no obvious abnormalities in appearance. The HDAC inhibitor treatment did not affect the mean body or placental weights at birth when compared with those of the control group (Table 3). We also observed the growth of pups obtained from SCNT embryos treated with Ky-9 or TSA and found that they were weaned at 3 weeks of age at rates similar to those of ICSI-derived pups (Supplementary Table S1). They showed no discernible abnormalities in behavior and appearance. All the Ky-9-derived clones tested for fertility (n = 4) produced offspring after mating with male mice (Supplementary Figure S2).

Developmental block induced by FK228 was associated with the inhibition of ZGA

As mentioned above, FK228 induced a strong two-cell developmental block following treatment for SCNT-derived embryos at the one-cell stage (Table 1). It is known that inhibition of the major ZGA phase that occurs at the two-cell stage in mouse embryos causes a developmental block [44]. To evaluate this possibility, we performed...
Synergistic effect of vitamin C and deionized BSA on the development of embryos derived from SCNT-derived embryos treated with Ky-9

It has been reported that an additional treatment with vitamin C and deionized BSA to TSA-treated SCNT embryos further improved their developmental efficiency [37]. To test whether this was also true for Ky-9-treated SCNT-derived embryos, we cultured them in the presence of vitamin C and deionized BSA for 8 h following treatment with Ky-9. As expected, the cloning efficiency was significantly increased by additional treatment with vitamin C and deionized BSA (Supplementary Figure S1).

**Table 4. In vitro development of SCNT-derived embryos following extended treatment with HDAC inhibitors for 24 h**

| Group       | Time (h) of inhibitor treatment | No. of embryos cultured | No. (%) of two-cell embryos at 24 h (%) | No. (%) of four-cell embryos at 48 h (per two cells) | No. of embryos at 96 h (% per two cells) |
|-------------|---------------------------------|--------------------------|----------------------------------------|------------------------------------------------------|----------------------------------------|
| Control     | None                            | 20                       | 20 (100)                               | 16 (80.0)                                            | 0 (0)                                  |
| TSA         | 8                               | 22                       | 22 (100)                               | 19 (86.4)                                            | 0 (0)                                  |
| Ky-29       | 8                               | 23                       | 23 (100)                               | 20 (87.0)                                            | 0 (0)                                  |
| TSA + TSA   | 24                              | 20                       | 20 (100)                               | 3 (15.0)*                                            | 3 (15.0)                               |
| TSA + Ky-29 | 24                              | 21                       | 19 (90.5)                              | 9 (47.4)*                                            | 0 (0)                                  |
| Ky-29 + Ky-29 | 24                          | 20                       | 20 (100)                               | 17 (85.0)                                            | 0 (0)                                  |

*P < 0.05 (vs. control, Fisher exact probability test).

Discussion

Since the first reports on the beneficial effect of TSA treatment on the development of mouse SCNT-derived embryos [15, 23], this method has become one of the standard cloning approaches. The same idea prompted cloning researchers to search for other HDAC inhibitors that might promote the development of SCNT-derived embryos. Those include scriptaid [24], SAHA [25], oxamflatin [25], and CBHA [26], which were found to significantly improve cloning efficiency in the mouse. All these HDAC inhibitors are hydroxamic acid derivatives and have a broad inhibitory spectrum against class I, IIa, and Ib HDACs. Here, we tested the activity of chlamydocin analogues, which include nonhydroxamic acid-type HDAC inhibitors and preferentially inhibit class I and IIa HDACs, and demonstrated that at least some of them—such as Ky-9, which possesses a methoxymethyl ketone moiety as the zinc-binding group—strongly promoted the development of SCNT-derived embryos to a level similar to that of TSA. Therefore, our results appear to have narrowed down the target HDACs for successful mouse SCNT to these two classes. By contrast, Ono et al. [25] considered class Ib HDAC to be the key for improving mouse SCNT. This was because valproic acid (VPA), an inhibitor for class I and IIa HDACs, had no effect on mouse SCNT. As our results using FK228 infer, HDAC inhibitors can have unexpected adverse effects on embryonic development. Indeed, the effective dose of VPA is high (in the mM range) and might not have exerted its beneficial effect on SCNT-derived embryos within its safe range. Indeed, VPA even at the minimal concentration that enhanced nuclear acetylation levels showed significant cell toxicity in vitro [45].

We expected that the use of FK228, an inhibitor of class I HDACs, would provide important clues to understanding how such inhibitors might affect mouse SCNT. Unexpectedly, FK228 treatment of SCNT-derived embryos resulted in a strong developmental block at the two-cell stage. Our EU incorporation assay revealed that this was a typical two-cell block caused by failure of major ZGA. Although it is not known how the inhibition of HDAC by FK228 explains this adverse effect at a molecular level, it is known that the sensitivity of cells to FK228 is increased with the intracellular level of glutathione [33]. As oocytes and early preimplantation embryos contain high amounts of glutathione [46], they might be particularly sensitive to FK228. Although this HDAC inhibitor is approved for the treatment of cutaneous and peripheral T-cell lymphomas [47], it is reported to be associated with a risk of cardiac toxicities [48]. As FK228 is a natural product, its analogues have been developed by refining its HDAC inhibitor activity and isoform selectivity [49]. It would be interesting to know whether the use of these FK228 analogues might help...
Figure 4. Analysis of the developmental ability of SCNT-derived embryos treated with HDAC inhibitor (s) for 24 h. (A) Scheme of the experiment. SCNT-derived embryos treated with an inhibitor for 8 h were further treated with the same or different inhibitor until 24 h. (B) The experimental groups including three groups of 24-h HDAC inhibitor treatment. (C) Bright-field images of SCNT-derived embryos treated with HDAC inhibitors for 24 h. The images were taken at 72 h after oocyte activation. For the details of the results, see Table 4.

Table 5. Full-term development of SCNT-derived embryos following sequential treatment with TSA and Ky-29 for 24 h

| Group       | No. of embryos cultured | No. of embryos at 24 h (%) | No. of embryos transferred (no. of recipients) | No. of implantations (%) | No. of offspring (%) | No. of placentas only (%) |
|-------------|-------------------------|---------------------------|-----------------------------------------------|-------------------------|---------------------|-------------------------|
| TSA +Ky-29  | 184                     | 179 (97.3)                | 179 (6)                                       | 62 (34.6)               | 2 (1.1)             | 6 (3.3)                 |

Although we do not know the exact cause of the developmental block induced by Ky-309, it might be related to the unique structure of Ky-309. This molecule is characterized by a bicyclic tetrapeptide structure with a CH$_2$ ($n = 10$) loop (Figure 1). This long CH$_2$ loop brings a bulky, hydrophobic region situated out of the active
These, HDAC4 is known to be expressed in mouse oocytes [56]. Low toxicity.

Primary function is histone deacetylation in oocytes and embryos, [56]. Thus, it is important to accurately identify those HDACs whose technical help with HDAC enzyme assays. Drug Discovery Platform, RIKEN Center for Sustainable Resource Science, for Acknowledgments.

Supplementary data are available at Supplementary material.

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Conflict of interest

The authors have declared that no conflict of interest exists.

Data availability

All data are incorporated into the article and its online supplementary material.

Author contributions

KI, NN, TW, MY, and AO conceived the project and designed the experiments. SK, AI, NN, MY, and AO wrote the manuscript. SK, EM, JM-K, KI, HI, NO, KM, SI, NI, AI, NN, and AO performed the experiments.

Supplementary material

Supplementary data are available at BIOLRE online.

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