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Interaction of reactive astrocytes with collagen type I induces astrocytic scar formation through the integrin/N-cadherin pathway after spinal cord injury

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1 **Abstract**

2 Central nervous system (CNS) injury activates naïve astrocytes into reactive astrocytes, which
3 eventually transform into scar-forming astrocytes that can impair axonal regeneration and
4 functional recovery. This sequential phenotypic change, known as reactive astrogliosis, has
5 long been considered unidirectional and irreversible. However, we report here that reactive
6 astrocytes isolated from injured spinal cords retrogradely reverted to naïve astrocytes when
7 transplanted into a naïve spinal cord, whereas they formed astrocytic scars when transplanted
8 into an injured spinal cord, indicating the environment-dependent reversibility of reactive
9 astrogliosis. We also found that collagen type I was highly expressed during the scar-forming
10 phase and induced astrocytic scar formation via the integrin/N-cadherin pathway.
11 Pharmacological blockade of reactive astrocyte-collagen type I interaction prevented astrocytic
12 scar formation, thereby leading to improved axonal regrowth and better functional outcomes
13 in a mouse model of spinal cord injury. Our findings reveal environmental cues regulating
14 astrocytic fate decisions, thereby providing a potential therapeutic target for CNS injury.

1 **Introduction**

2 Spinal cord injury (SCI) is a devastating trauma that causes persistent severe motor/sensory
3 dysfunction^{1,2}. After SCI, astrocytes, the most abundant resident cells in the central nervous
4 system (CNS), play a crucial role in the SCI pathology through a phenotypic change known as
5 reactive astrogliosis. In this process, naïve astrocytes (NAs) sequentially exhibit opposite
6 phenotypes: first as reactive astrocytes (RAs) and then as scar-forming astrocytes (SAs). In the
7 subacute phase of SCI (4-14 days post-injury (dpi) in the mouse), RAs migrate to the lesion
8 epicenter and seclude inflammatory cells, leading to tissue repair and functional improvement
9 after SCI². However, RAs gradually transform into SAs that form astrocytic scars, the main
10 impediment for CNS axonal regeneration, resulting in a limited functional recovery in the
11 chronic phase of SCI^{3,4} (more than 14 dpi in the mouse). Astrocytic scars have been shown to
12 be irreversible and permanently inhibit axonal regrowth in both rodents and humans with SCI⁴
13 ⁶, although there is a different opinion that attenuating astrocytic scar formation failed to
14 promote axonal regeneration after SCI^{7,8}. As such, clarifying the mechanism of astrocytic scar
15 formation and regulating this scar formation may be a potential therapeutic strategy for SCI.

16 Astrocytic scars have been studied for more than half a century^{9,10}, and their formation has
17 been suggested to be regulated by complex and combinatorial inter- and intra-cellular signaling
18 mechanisms^{3,5,11}. However, despite the large number of studies examining these astrocytic
19 changes^{5,11,12}, the mechanism underlying astrocytic scar formation remains unclear. One factor
20 limiting basic research in this area is the lack of clear definitions of NAs, RAs, and SAs. A
21 conventional method for distinguishing between these cells is a histological analysis, but this
22 method is neither objective nor quantitative.

23 In this study, we established a clear distinction between NAs, RAs, and SAs based on marker
24 gene expression and investigated the regulatory mechanism underlying astrocytic scar

1 formation after SCI.

1 **Results**

2 **In situ isolation of astrocytes after SCI enables the discovery of RA- and SA-specific** 3 **marker genes**

4 We employed the contusion SCI model in female C57BL/6J mice (8-10 weeks old). This
5 approach is reported to resemble human SCI more closely than transection models, because
6 most human SCI are due to blunt trauma¹³. After SCI, NAs exhibited drastic changes around
7 the lesion area (**Supplementary Fig. 1a**). They underwent a typical change of hypertrophy,
8 process extension and increased GFAP expression within several dpi, characteristic of RAs
9 (**Supplementary Fig. 1b**)². Thereafter, at 14 dpi, the RAs overlapped and transformed into
10 SAs, resulting in astrocytic scar formation (**Supplementary Fig. 1b**)^{3,12}. Although this
11 sequential astrocytic transformation is roughly referred to as “reactive astrogliosis”, identifying
12 the point at which the astrocytes are situated in this process is difficult: RAs and SAs both
13 highly express their hallmark protein (GFAP, Nestin, β -catenin, N-cadherin and Sox9), but
14 clearly distinguishing between each form of astrocytes (NAs, RAs, and SAs) based solely on
15 their hallmark expression is difficult with a histological analysis (**Supplementary Fig. 2a,b,**
16 **Supplementary Fig. 3a,b**)^{12,14-16}.

17 Therefore, we selectively isolated each astrocyte type by laser microdissection (LMD)
18 combined with immunohistochemistry: NAs, morphologically identical to resident astrocytes
19 in the naïve spinal cord; RAs, with the morphological characteristics described above (observed
20 in the injured spinal cord at 7 dpi); and SAs, scar-forming GFAP-positive cells observed in the
21 injured spinal cord at 14 dpi in a mouse model of contusion SCI (**Fig. 1a-g**). Excellent RNA
22 qualities as well as no contamination of other cell type were confirmed with a bioanalyzer and
23 a PCR analysis in the collected samples of astrocytes (**Fig. 1h, i**). We investigated the
24 expression levels of RA- and SA-characteristic genes that were reported previously, and

1 defined the specific genes both significantly and selectively upregulated in LMD-isolated NAs,
2 RAs and SAs as their respective marker genes.

3 Regarding RAs, we confirmed elevated gene expressions of *Gfap*, *Nes*, *Vim*, and *Ctnnb1*
4 (encoding β -catenin) compared with NAs. However, increased expressions of *Gfap* and *Vim*
5 were also observed in SAs, indicating that only *Nes* and *Ctnnb1* were suitable as RA marker
6 genes (**Fig. 1j**). In addition, we found the β -catenin-related genes suitable as RA markers from
7 a functional aspect. In SCI, RA migration is crucial for both functional and tissue repair², and
8 recent reports have suggested a possible molecular link between β -catenin and cell movement
9 via MMPs^{17,18}. After β -catenin nuclear translocation, the β -catenin-LEF/TCF complex binds
10 the enhancer of urokinase-type plasminogen activator receptor (uPAR) gene, resulting in the
11 activation of MMPs and cell migration. We demonstrated selectively elevated expression of
12 *Plaur* (encoding uPAR), *Mmp2*, and *Mmp13* in RAs compared with NAs and SAs (**Fig. 1j**).
13 Furthermore, the expression of *Axin2*, a negative feedback regulator of β -catenin, was
14 significantly higher in RAs than in NAs or SAs (**Fig. 1j**). These results show that *Plaur*, *Mmp2*,
15 *Mmp13*, and *Axin2* as well as *Nes* and *Ctnnb1* are suitable as RA marker genes, providing
16 further insight into the mechanism of RA migration through β -catenin/MMPs signaling.

17 Regarding SAs, we confirmed significantly elevated expression of *Cdh2* (encoding N-
18 cadherin) and *Sox9* compared with NAs and RAs (**Fig. 1k**). We also detected significantly
19 higher expression of chondroitin sulfate proteoglycan (CSPG)-related genes, such as *Xylt1*,
20 *Chst11*, *Csgalnact1*, *Acan*, and *Pcan*, in SAs than in NAs and RAs (**Fig. 1k**). These results
21 were biologically plausible, because CSPGs are the most abundant extracellular inhibitor of
22 axonal regeneration, and their expression has been selectively observed in the astrocytic scar
23 area after SCI^{3,12,16,19}. In addition, the strong repulsive axon guidance gene *slit2*^{3,20} was also
24 found to be suitable as a SA marker gene (**Fig. 1k**). These marker gene profiles combined with

1 the morphological definitions are useful for identifying the astrocyte phenotypes present, as
2 reactive astrogliosis is a graded continuum of changes in gene expression and forms.

3

4 **Dependence of reactive astrogliosis on the surrounding environment**

5 The phenotypic changes in mammalian cells are roughly driven in a cell-autonomous or a non-
6 cell-autonomous manner^{21,22}; however, the mechanism of reactive astrogliosis remains unclear.
7 To clarify the mechanism, we prepared GFP-positive NAs from primary astrocyte cultures of
8 *CAG-EGFP* mice and transplanted them into the spinal cord of either naïve mice or
9 immediately after spinal contusion in SCI mice (**Fig. 2a, Supplementary Fig. 4a-c**). At 7 and
10 14 days after transplantation (dat), we examined morphological transformations of the grafted
11 NAs in response to the environment (**Fig. 2b**). The morphology of transplanted NAs, such as
12 the cell diameter and ratio of cells with overlapping of neighboring astrocytic processes, was
13 unchanged in the naïve spinal cord, whereas they had transformed along with the host astrocytic
14 changes and were integrated into reactive astrogliosis when transplanted into the injured spinal
15 cord (**Fig. 2c-j**).

16 In addition to these morphological changes, we examined the sequential changes in the gene
17 expression by a cell-selective analysis of grafted NAs using LMD (**Fig. 3a,b**). Consistent with
18 the morphological transition, gene expression of the grafted NAs remained unchanged when
19 transplanted into the naïve spinal cord, whereas the expression of the RA marker genes at 7 dat
20 and SA marker genes at 14 dat were significantly increased compared to those at other time
21 points when transplanted into the injured spinal cord (**Fig. 3c,d**). These findings indicate that
22 astrocytes change their phenotypes in an environment-dependent manner (**Supplementary Fig.**
23 **5a**).

24

1 **Reversibility of reactive astrogliosis**

2 After confirming the environment-dependent properties of reactive astrogliosis after SCI, we
3 examined the reversibility of the sequential astrocytic changes and whether or not RAs could
4 retrogradely revert to NAs after transplantation into naïve spinal cord. To selectively isolate
5 RAs, we developed *Nes-Cre; CAG-CAT^{loxP/loxP}-EGFP* (*Nes-EGFP*) mice, which show selective
6 EGFP expression in RAs upon Cre-mediated recombination². We confirmed that these mice
7 showed no GFP-positive cells in the naïve spinal cord, but had high Cre-mediated EGFP
8 expression in RAs around the lesion area at 7 dpi in SCI mice (**Supplementary Fig. 6a-d, Fig.**
9 **4a,b**). After dissociating the injured spinal cords of *Nes-EGFP* mice at 7 dpi, we selectively
10 isolated the GFP-positive RAs by fluorescence-activated cell sorting (FACS) and confirmed
11 significantly increased expressions of the RA marker genes in the isolated RAs
12 (**Supplementary Fig. 7a,b**). We then transplanted these GFP-positive RAs into either naïve or
13 injured spinal cord models and performed immunohistological and cell-selective gene
14 expression analyses with LMD at 7 dat (**Fig. 4c,d**). The grafted RAs formed astrocytic scars
15 when transplanted into the injured spinal cord but histologically reverted to NAs when
16 transplanted into the naïve spinal cord at 7 dat. (**Fig. 4e-h**). Consistent with this morphological
17 conversion, a selective gene expression analysis of the grafted cells demonstrated that the SA
18 marker gene expressions were upregulated when transplanted into the injured spinal cord, while
19 the original expression patterns of SA marker genes were significantly suppressed when
20 transplanted into the naïve spinal cord (**Fig. 4i**). Moreover, the whole transcriptome analysis of
21 isolated RAs, transplanted RAs in the naïve spinal cord at 7 dat, and host astrocytes in the naïve
22 spinal cord, revealed that the global gene expression profile of transplanted and host astrocytes
23 was comparable, while it was markedly different from isolated RAs (**Fig. 4j, Supplementary**
24 **Fig. 8a**), suggesting that the *in situ* phenotype reprogramming of astrocytes depended on the

1 environment. These results clearly indicate that the process of reactive astrogliosis is reversible
2 under certain conditions (**Supplementary Fig. 9a**).

3

4 **Upregulation of collagen type I genes in the lesion area after SCI**

5 RAs are beneficial, while SAs are harmful for the repair process after SCI; as such, preventing
6 the transformation of RAs to SAs through environmental regulation may be a new therapeutic
7 strategy for SCI. Given that grafted RAs reverted to NAs in the naïve spinal cord but
8 transformed into SAs in the injured spinal cord, the fate of RAs seems to be determined by
9 environmental differences between the naïve and injured spinal cord.

10 We therefore performed a time-course genome-wide expression analysis in naïve and injured
11 spinal cord models after SCI, as described in our previous studies^{23,24}. The scatter plots showed
12 both the upregulation and downregulation of all genes between the naïve and injured groups
13 (**Fig. 5a**). The expressions of nearly 5% of the genes were considerably increased in the injured
14 spinal cord at 14 dpi compared to those in the naïve spinal cord (>5-fold change; **Fig. 5a**). We
15 originally expected that the genes involved in intracellular signaling or cytokines guiding cell
16 transformation would be enriched in the scar-forming phase of SCI. However, interestingly, a
17 GO term analysis showed that the genes associated with the extracellular matrix (ECM) were
18 highly enriched at 14 dpi, indicating the involvement of the ECM in scar formation after SCI
19 (**Fig. 5b**). Among the ECM genes, the collagen type I (Col I) genes (*Colla1* and *Colla2*) were
20 the most highly expressed in the injured spinal cord at 14 dpi (**Fig. 5c**). In addition, the mRNA
21 expressions of Col I genes were significantly greater in the injured spinal cord than in the naïve
22 spinal cord (**Fig. 5d**). Furthermore, in immunohistochemistry, the Col I expression was
23 abundantly observed at the lesion area at 14 dpi in the injured spinal cord but not in the naïve
24 spinal cord (**Fig. 5e**). Interestingly, astrocytic scars were formed by SAs colocalizing with Col

1 I; however, the RAs retained their phenotypes in the Col I-absent area at 14 dpi (**Fig. 5e-g**).
2 These findings suggest that Col I is directly involved in the transformation of RAs into SAs as
3 well as astrocytic scar formation after SCI.

4 5 **Induction of N-cadherin-mediated adhesion by collagen type I in astrocytes**

6 To clarify the influence of Col I on RA fate determination, we cultured RAs *in vitro* on Col I-
7 coated dishes (Col I-coated group) or non-coated dishes (non-coated group) and performed an
8 immunocytological analysis. While the RAs of the non-coated group showed process retraction
9 and decreased GFAP expression, the RAs of the Col I-coated group adhered closely to the
10 adjacent cells and showed increased GFAP expression, exhibiting a phenotype similar to that
11 of SAs (**Fig. 5h**). In addition, we found significantly increased expression of N-cadherin,
12 characteristic of SA, in only the Col I-coated group (**Fig. 5h,i**). Given that astrocytes form N-
13 cadherin-mediated adhesion^{25,26} and that N-cadherin upregulation had been observed at
14 astrocytic scars (**Supplementary Fig. 3a**)¹⁵, these results suggest that Col I enhances N-
15 cadherin-mediated contacts in RAs and induces their transformation into SAs.

16 17 **Induction of astrocytic transformation by the collagen type I-integrin axis**

18 For ECM proteins, including Col I, their counterpart receptors are necessary in order to interact
19 with cells²⁷. Among ECM receptors, $\alpha 1\beta 1$, $\alpha 2\beta 1$, $\alpha 10\beta 1$, and $\alpha 11\beta 1$ integrin are reported to be
20 a major class of collagen-binding integrins that play an important role in cell-Col I interaction²⁷.
21 Since these collagen-binding integrins are expressed on astrocytes^{28,29}, we speculated that they
22 directly mediated the RA-Col I interaction and induced the transformation into SAs. We
23 therefore analyzed the morphological and gene expression changes in RAs of the Col I-coated
24 group treated with or without anti-collagen-binding integrin antibody (anti- $\beta 1$ Ab) (**Fig. 6a**).

1 Notably, the RAs treated with control Ab showed cell adhesion and the increased expression
2 of N-cadherin and GFAP, while those treated with anti- β 1 Ab remained scattered with no
3 change in the expression of N-cadherin or GFAP (**Fig. 6b,c**). Similarly, N-cadherin
4 neutralizing antibody (N-cadherin nAb) inhibited the transformation of RAs into SAs under
5 the same conditions (**Supplementary Fig. 10a**). These results suggest that Col I induces the
6 transformation of RAs into SAs via the integrin/N-cadherin pathway, indicating a potential
7 therapeutic target.

8

9 **Prevention of astrocytic scar formation and amelioration of functional recovery after SCI** 10 **via the blockade of RA-collagen type I interaction**

11 Therapeutically, whether or not preventing the transformation of RAs into SAs can indeed
12 attenuate astrocytic scar formation after SCI is particularly important, because axons cannot
13 regenerate beyond astrocytic scars, thereby limiting the functional recovery^{3,4,6,12}. We
14 investigated the effects of inhibition of integrin-mediated RA-Col I interaction on *in vivo*
15 astrocytic scar formation and functional recovery after SCI (**Fig. 6a**).

16 Anti- β 1 Ab was administered from 9 to 13 dpi after SCI in order to selectively attenuate the
17 transformation of RAs to SAs (**Fig. 6d**). At 14 dpi in the anti- β 1 Ab injection group, we
18 observed significantly lower expression of both N-cadherin and GFAP at the lesion area,
19 resulting in reduced cell adhesion and astrocytic scar formation compared to the control group
20 (**Fig. 6e,f**). We also confirmed that the injection of N-cadherin nAb attenuated astrocytic scar
21 formation similarly to the administration of anti- β 1 Ab (**Supplementary Fig. 10b-d**).
22 Regarding the axonal regrowth, immunostaining using an antibody against GAP43, a marker
23 for regenerating axons, showed that the number of GAP43-positive axons was significantly
24 increased at both the lesion epicenter and the caudal area of the lesion in mice treated with anti-

1 β 1 Ab compared with those treated with control Ab (**Fig. 6g,h**). The numbers of serotonin (5-
2 HT)-positive and tyrosine hydroxylase (TH)-positive fibers in mice treated with anti- β 1 Ab
3 were also larger in comparison to mice treated with control Ab, and most of these fibers in mice
4 with anti- β 1 Ab expressed GAP43 (**Supplementary Fig. 11a-d**). These results indicate that
5 the number of regenerative fibers passing through the lesion epicenter was significantly
6 increased by the attenuation of astrocytic scar formation. To investigate whether or not these
7 regenerating axons contribute to the behavioral improvements, we performed an open-field
8 locomotor test and a footprint analysis, and demonstrated that the motor functional recovery
9 with anti- β 1 Ab injection was significantly greater than that achieved with control Ab injection
10 (**Fig. 6i-l**). These results indicate that the inhibition of integrin-mediated RA-Col I interaction
11 prevents astrocytic scar formation via N-cadherin downregulation and promotes the axonal
12 regeneration and functional recovery after SCI.

1 **Discussion**

2 Astrocytic scars, the final form of reactive astrogliosis, are widely regarded as a principal cause
3 for axonal regrowth failure and a poor functional outcome^{3,12}. Reactive astrogliosis has long
4 been considered unidirectional and irreversible in the SCI pathology^{2,4,5}. However, our findings
5 conversely showed the environment-dependent reversibility of reactive astrogliosis,
6 highlighting the therapeutic potential of regulating this astrocytic change through
7 environmental intervention. In particular, a time-course RNA-seq analysis identified Col I as a
8 prominent factor for the scar-forming phase of SCI, and Col I was crucial for astrocytic scar
9 formation via the integrin/N-cadherin pathway. Furthermore, the blockade of RA-Col I
10 interaction attenuated astrocytic scar formation and promoted functional improvement after
11 SCI, providing a novel therapeutic target for the treatment of SCI.

12 Using LMD, we demonstrated for the first time the *in vivo* transcriptional profiles of NAs,
13 RAs, and SAs and defined their marker genes. FACS or translating ribosomal affinity
14 purification (TRAP) have thus far been mainly used for cell-specific analyses^{23,30}; however,
15 LMD had an advantage over these methods in our study. While FACS and TRAP require the
16 separation of solid organ tissues into single cells³¹, LMD makes it possible to capture cells
17 while maintaining and observing cell-specific structures³². Therefore, LMD combined with
18 immunohistochemistry was indispensable for selectively isolating each form of astrocyte based
19 on the recognition of their antigens and morphologies, because the astrocytes exhibited
20 phenotypic heterogeneity around the SCI lesion (**Supplementary Fig. 1a, Fig. 5f-h**).

21 Although the completed astrocytic scars strongly inhibit the regenerative response after CNS
22 injury, RAs, transitional form of reactive astrogliosis, play beneficial roles in the subacute
23 phase of SCI, in processes such as local immune regulation, neuroprotection, and tissue
24 repair³³⁻³⁶. In addition to the definition of astrocytic marker genes, a cell-selective analysis with

1 LMD demonstrated that the signaling pathways related to these beneficial functions were
2 activated in RAs (**Fig. 1j, Supplementary Fig. 12a**). For example, the β -catenin/MMPs
3 pathway was reported to be involved in astrocytic migratory activity as well as tissue repair
4 after SCI. TGF- β signaling contributes to both the neuronal survival and axonal
5 regeneration^{37,38}. PDGF reduces the inflammatory response by enhancing the phagocytic
6 activity of microglia/macrophages after SCI³⁹. We found that these pathways were significantly
7 and selectively activated in RAs, providing deeper insight into the beneficial influence of RAs
8 on tissue repair and functional recovery after SCI.

9 As shown in **Figs. 5 and 6**, we demonstrated that Col I was involved in the N-cadherin-
10 mediated cell-cell adhesion in astrocytes through integrin signaling and astrocytic scar
11 formation. To date, pericytes and fibroblasts have been reported to be the cell types that
12 produce Col I after SCI^{12,40,41}, and we actually confirmed that these cells were clearly merged
13 with Col I in the lesion epicenter (**Supplementary Fig. 13a,b**). Regarding a possible molecular
14 link between the Col I-integrin axis and increased N-cadherin-dependent contact in astrocytes,
15 c-Jun-N-terminal kinase (JNK) may play a role. In previous studies of epithelial and cancer
16 cells, the Col I-integrin axis was reported to activate the downstream JNK pathway, which
17 influenced the N-cadherin-dependent intercellular adhesion^{42,43}. Similarly, in astrocytes, Col I-
18 integrin axis stimulation was recently reported to induce JNK activation⁴⁴, and upregulation of
19 both JNK and N-cadherin has been observed in astrocytes after CNS injury (**Supplementary**
20 **Fig. 3a**)^{15,45,46}. Given that JNK inhibitor attenuates astrocytic scar formation in the injured
21 spinal cord⁴⁷, the integrin/JNK/N-cadherin pathway is crucial for astrocytic scar formation after
22 SCI. In this study, we also confirmed both attenuated astrocytic scar formation via Col I-
23 integrin axis inhibition by anti- β 1 Ab as well as an improved functional outcome (**Fig. 6d-l**),
24 providing a novel potential therapeutic target for SCI.

1 N-cadherin regulated reactive astrogliosis by increasing the astrocytic cell-cell adhesion. N-
2 cadherin belongs to the classic cadherins that are calcium-dependent intercellular adhesion
3 molecules^{25,26} and has many functions in the CNS, such as differentiation, migration, and cell
4 polarization^{25,26,48}. In addition to cell adhesion, N-cadherin in astrocytes has also been reported
5 to be involved in the migratory activity and cellular morphology. For example, both a lack of
6 and the overexpression of N-cadherin result in the impairment of astrocytic migration^{48,49}.
7 Kanemura et al. also reported that astrocyte-specific knockout of N-cadherin results in
8 insufficient morphological changes in both RAs and SAs after CNS injury⁴⁸. These results
9 indicate that the N-cadherin expression in astrocytes is finely tuned through the process of
10 reactive astrogliosis. Indeed, we confirmed the time-phase-dependent expression of N-cadherin
11 concomitant with cellular hypertrophy and process extension in RAs and SAs (**Fig. 1k, Fig.**
12 **5h, Fig. 6b,e, Supplementary Fig. 3a**). Although both anti- β 1 Ab and N-cadherin nAb
13 prevented the transformation of RAs into SAs and attenuated astrocytic scar formation in this
14 study, there is a possibility that N-cadherin nAb impairs the beneficial function of RA more
15 severely than anti- β 1 Ab, since N-cadherin has significant influence on the cellular morphology
16 and migration of RA as described above. We therefore used anti- β 1 Ab. In the present study,
17 anti- β 1 was administered during only the scar-forming phase to attenuate scar formation
18 without influencing the morphological changes or migratory activity of RAs. Our results have
19 important implications for the optimization of therapeutic intervention from the time-phase-
20 dependent view of reactive astrogliosis.

21 Regarding the role of astrocytic scars after SCI, Anderson et al. recently reported that the
22 attenuation of astrocytic scar formation failed to result in spontaneous axonal regrowth after
23 SCI^{7,8}, suggesting that astrocytic scar formation aids axonal regeneration. However, the
24 impairment of “the whole process of reactive astrogliosis” has already been reported to cause

1 the deterioration of axonal regrowth and functional recovery after SCI in several previous
2 studies^{3,50,51}, and there has been no study demonstrating that astrocytic “scars” directly promote
3 axonal regeneration after SCI. In contrast, Silver and many other investigators demonstrated
4 that astrocytic “scars” directly impede axonal regeneration^{3-6,12,51}, and our results supported
5 this finding. In this study, we therefore prevented only the transformation of RAs into SAs
6 without affecting RAs, leading to axonal elongation and functional improvement after SCI.
7 These findings suggest the significance of the time-phase-dependent regulation of reactive
8 astrogliosis for the SCI treatment.

9 In conclusion, we demonstrated the reversibility of reactive astrogliosis by a selective gene
10 expression analysis with LMD and transplantation experiments. We also clarified a mechanism
11 of astrocytic scar formation through RA-Col I interaction, supporting a novel therapeutic
12 strategy for attenuating astrocytic scar formation and improving the functional outcomes after
13 SCI.

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8

9 **Author contributions**

10 M.H. designed and performed most of the experiments with technical help from H.K., T.S., and
11 S.Y. Y.O. performed the RNA-Seq analysis. K.Y. and K. Kijima analyzed the data of LMD. K.
12 Kobayakawa, K.H. and Y.N. designed the studies and supervised the overall project. S.O.
13 designed the studies, supervised the overall project, and performed the final manuscript
14 preparation.

15

16 **Competing financial interests**

17 The authors declare no competing financial interests.

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7

1 **Figure legends**

2 **Figure 1. Selective *in situ* isolation and gene expression analysis of astrocytes from the**
3 **naïve and injured spinal cords. (a, c and e)** Example images depicting GFAP (green) staining
4 of the naïve (a) and injured spinal cords at 7 (c) and 14 (e) dpi. The asterisk indicates the lesion
5 epicenter in SCI mice (c and e). Scale bars, 500 μ m. (b, d and f) Example high magnification
6 images depicting immunohistochemical analysis of GFAP-positive astrocytes (green) stained
7 with Hoechst (blue) in the naïve (b) and injured spinal cords at 7 (d) and 14 (f) dpi. The
8 arrowheads and dashed-outlines indicate the astrocytes before (left) and after (right) LMD. The
9 images are magnifications of the boxed areas in (a), (c) and (e), respectively. Scale bars, 50 μ m.
10 (g) A schematic representation of the selective isolation of GFAP-positive astrocytes using
11 LMD. (h) Bioanalyzer gel image (left) and electropherogram with the associated RNA integrity
12 number (RIN) values (right) of RNA from LMD-isolated astrocytes. (i) Example PCR analysis
13 of selected transcripts in whole spinal cord (top) and LMD-isolated astrocytes (bottom). (j)
14 Quantification of mRNA expression levels of RA-related genes in NAs, RAs, and SAs ($n = 4$
15 mice per group). (k) Quantification of mRNA expression levels of SA-related genes in NAs,
16 RAs, and SAs ($n = 4$ mice per group). * $P < 0.05$, analysis of variance (ANOVA) with the
17 Tukey-Kramer post hoc test (j and k); n.s., not significant. Data represent mean \pm SEM.

18

19 **Figure 2. Environment-dependent morphological changes of NAs transplanted into naïve**
20 **or injured spinal cords. (a and b)** The NA transplantation protocol. NAs from primary
21 astrocyte cultures of *CAG-EGFP* mice were transplanted into the naïve and injured spinal cord
22 immediately after SCI. An immunohistochemical (IHC) analysis of the grafted NAs was
23 performed at 0, 7, and 14 dat. (c, e and g) Example images depicting IHC analysis of GFP-
24 positive NAs (green) transplanted into the naïve spinal cord at 7 dat (c) and the injured spinal

1 cord at 7 (e) and 14 (g) dat, stained with GFAP (red). The asterisk indicates the lesion epicenter
2 in SCI mice (e and g). Scale bars, 500 μ m. (d, f and h) Example high magnification images
3 depicting IHC analysis of GFP-positive grafted NAs (green) stained with GFAP (red) and
4 Hoechst (blue) in the naïve spinal cords at 7 dat (d) and the injured spinal cords at 7 (f) and 14
5 (h) dat. The images are magnifications of the boxed areas in (c), (e) and (g), respectively. Scale
6 bars, 50 μ m. (i) A morphometric analysis of the cell diameter in host or GFP-positive
7 transplanted astrocytes in naïve or injured spinal cords ($n = 8$ mice per group). (j) The
8 percentage of host or GFP-positive transplanted astrocytes with overlapping of neighboring
9 astrocytic processes in naïve or injured spinal cords ($n = 8$ mice per group). Wilcoxon's rank
10 sum test (i and j); n.s., not significant. Data represent mean \pm SEM.

11

12 **Figure 3. Environment-dependent transcriptional profiles of NAs transplanted into naïve**
13 **or injured spinal cords.** (a and b) The NA transplantation and selective isolation protocol.

14 NAs from primary astrocyte cultures of *CAG-EGFP* mice were transplanted into naïve or
15 injured spinal cords immediately after SCI. GFP-positive grafted NAs were selectively isolated
16 by LMD for a gene expression analysis at 0, 7, and 14 dat. The arrowheads and dashed-outlines
17 indicate the astrocytes before (left) and after (right) LMD. Scale bar, 50 μ m (a). (c and d) Time-
18 course quantification of mRNA expression levels of RA (c) and SA marker genes (d) in NAs
19 transplanted into naïve or injured spinal cords ($n = 4$ mice per group). Gene expression levels
20 were normalized to that of the NAs transplanted into the naïve spinal cord at 0 dat. $\star P < 0.05$,
21 ANOVA with the Tukey-Kramer post hoc test; n.s., not significant. Data represent mean \pm SEM.

22

23 **Figure 4. Reversibility of reactive astrogliosis depends on the surrounding spinal cord**
24 **environment.** (a) Example images depicting GFP (green) staining of the sagittal section of

1 *Nes-EGFP* mice at 7 dpi after SCI. The asterisk indicates the lesion epicenter in SCI mice.
2 Scale bar, 500 μ m. **(b)** Example high magnification images depicting immunostaining of GFP
3 (green), GFAP (red), and Nestin (blue) triple-positive RAs in the perilesional area of *Nes-EGFP*
4 mice at 7 dpi. The asterisk indicates the lesion epicenter in SCI mice. Scale bar, 50 μ m. **(c and**
5 **d)** The protocol showing selective isolation and transplantation of RAs. **(e and f)** Example high
6 magnification images depicting immunostaining of the GFP-positive RAs (green) transplanted
7 into naïve or injured spinal cords at 7 dpi, stained with GFAP (red) and Hoechst (blue). The
8 asterisk indicates the lesion epicenter in SCI mice **(e)**. Scale bars, 50 μ m. **(g)** A morphometric
9 analysis of the cell diameter in host or GFP-positive transplanted astrocytes in naïve or injured
10 spinal cords ($n = 8$ mice per group). **(h)** The ratio of host or GFP-positive transplanted
11 astrocytes with overlapping of neighboring astrocytic processes in naïve or injured spinal cords
12 ($n = 8$ mice per group). **(i)** Time-course quantification of mRNA expression levels of SA
13 marker genes in RAs transplanted into naïve or injured spinal cords ($n = 4$ mice per group).
14 Gene expression levels were normalized to that of NAs in the naïve spinal cord of *Nes-EGFP*
15 mice. **(j)** The hierarchical clustering of the whole transcriptome analysis data from RAs,
16 transplanted astrocytes and host astrocytes ($n = 3$ mice per group). The values indicate
17 Pearson's correlation coefficients. $\star P < 0.05$, ANOVA with the Tukey-Kramer post hoc test (**g,**
18 **h and i**); n.s., not significant. Data represent mean \pm SEM.

19
20 **Figure 5. RNA-seq analysis reveals the role of Col I in the induction of N-cadherin-**
21 **dependent adhesion in RAs. (a)** Scatter plots between the naïve and injured spinal cord ($n =$
22 1 mice per group). These represent many overexpressed (>5 -fold change) genes (red). **(b)** A
23 GO term analysis of the overexpressed (> 5 -fold change) genes in the RNA-seq analysis of the
24 injured spinal cord (14 dpi) compared to those of the naïve spinal cord ($n = 1$ mice per group).

1 The lists show the top seven obtained GO terms ranked by *p*-value (Fisher's exact test with the
2 Benjamini-Hochberg correction). (c) The FPKM values of the top 20 extracellular matrix genes
3 at 14 dpi (*n* = 1 mice per group). (d) Quantification of mRNA expression levels of the Col I
4 genes (*Colla1* and *Colla2*) of spinal cords before injury and at 14 dpi (*n* = 4 mice per group).
5 (e, f, and g) Example images depicting immunostaining of the naïve and injured spinal cord at
6 14 dpi, stained with GFAP (green), Col I (red), and Hoechst (blue). The asterisk indicates the
7 lesion epicenter in SCI mice (e). Scale bars, 100 μ m (e), 50 μ m (f and g). (h) Example images
8 depicting immunocytochemical analysis of the RAs on the non-coated or Col I-coated dishes,
9 stained with GFAP (green), N-cadherin (red), and Hoechst (blue). Scale bar, 50 μ m. (i)
10 Quantification of *Cdh2* mRNA expression levels in the RAs on the non-coated or Col I-coated
11 dishes (*n* = 4 wells per group). **P* < 0.05, Wilcoxon's rank sum test (d and i). Data represent
12 mean \pm SEM.

13

14 **Figure 6. Inhibition of integrin-mediated RA-Col I interaction prevents astrocytic scar**
15 **formation and promotes functional recovery after SCI.** (a) A schematic representation of
16 the SA transformation induced by the Col I/integrin/N-cadherin pathway. Anti- β 1 Ab blocks
17 the Col I-integrin axis in RAs. (b) Example images depicting immunocytochemical analysis of
18 RAs on Col I-coated dishes after anti- β 1 Ab or control Ab administration, stained with GFAP
19 (green), N-cadherin (red), and Hoechst (blue). Scale bar, 50 μ m. (c) Quantification of *Cdh2*
20 mRNA expression levels in RAs after anti- β 1 Ab or control Ab administration (*n* = 4 wells per
21 group). Gene expression levels were normalized to that of the RAs on non-coated dishes
22 without any Ab administration. (d) The schedule of *in vivo* anti- β 1 Ab injection to wild-type
23 mice after SCI. (e) Example images depicting IHC analysis of the injured cords receiving anti-
24 β 1 Ab or control Ab injection, stained with GFAP (green), N-cadherin (red), and Hoechst (blue).

1 The asterisk indicates the lesion epicenter in SCI mice. Scale bar, 50 μm . **(f)** The ratio of GFAP-
2 positive astrocytes with overlapping of neighboring astrocytic processes in the lesion area of
3 SCI mice after anti- β 1 Ab or control Ab administration ($n = 8$ mice per group). **(g)** Example
4 images depicting GAP43 (white) and Col I (red) staining of the injured cords receiving anti- β 1
5 Ab or control Ab injection at 42 dpi. Scale bars below the images, 200 μm , insets 40 μm . **(h)**
6 Quantification of GAP43-positive area per $4.0 \times 10^5 \mu\text{m}^2$ ($n = 8$ mice per group). **(i)** Time
7 course of the BMS score after SCI ($n = 12$ mice per group). **(j)** Example images depicting
8 footprint analysis at 42 dpi. **(k)** Quantification of the footprint analysis findings in each mouse
9 ($n = 8$ mice per group). **(l)** Scores for the grip walk test at 42 dpi ($n = 8$ mice per group). $\star P <$
10 0.05 , $\star\star P < 0.005$, ANOVA with the Tukey-Kramer post hoc test **(c)**, Wilcoxon's rank sum test
11 **(f, h, k and l)**, and two-way repeated measures ANOVA with the Tukey-Kramer post hoc test
12 **(i)**. Data represent mean \pm SEM.

13

14

1 **Online Methods**

2 **Animals**

3 All animal studies were approved by the Committee of Ethics on Animal Experimentation of
4 the Faculty of Medicine, Kyushu University. The experiments were conducted in accordance
5 with the National Institutes of Health guidelines for the care and use of animals. Female
6 C57BL/6J mice (8–10 weeks old) were used in this study. *CAG-EGFP* mice, *Nes-Cre* mice,
7 and *CAG-CAT^{loxP/loxP}-EGFP* mice were as described previously^{2,52}. To generate *Nes-Cre; CAG-*
8 *CAT^{loxP/loxP}-EGFP* (*Nes-EGFP*) mice, we crossed *Nes-Cre* mice and *CAG-CAT^{loxP/loxP}-EGFP*
9 mice.

10

11 **Primary astrocyte cultures**

12 Purified primary astrocyte cultures were prepared from *CAG-EGFP* mice or C57BL/6J mice,
13 as described previously^{2,53}. In brief, after the removal of the meninges, postnatal day 2 mouse
14 brain tissues were minced and incubated in a rocking water bath at 37 °C for 30 min in
15 Dulbecco's modified Eagle's medium (DMEM; Invitrogen, Carlsbad, California) in the
16 presence of 0.25% trypsin (Sigma-Aldrich) and 300 g ml⁻¹ DNase I (Sigma-Aldrich). The
17 dissociated cells were triturated with 0.25% fetal bovine serum (FBS) and centrifuged at 300
18 × g for 5 min. Following dilution with astrocyte-specific medium (DMEM containing 10%
19 FBS, 0.2 mL-glutamine, and 1% penicillin-streptomycin), the cells were plated on non-
20 coated culture dishes. After 7-10 days in a humidified CO₂ incubator at 37 °C, the astrocytes
21 were used for experiments. For the activation to RAs, the astrocytes were stimulated with 50
22 ng ml⁻¹ IL-6 (R&D Systems Inc.) and 200 ng ml⁻¹ soluble IL-6 receptor (R&D Systems Inc.).
23 For the preparation of Col I-coated dishes, the dishes were incubated at 37 °C for 2 h with
24 mouse collagen type I (50 µg ml⁻¹; AbD Serotec), as described previously⁵⁴.

1

2 **Spinal cord injury**

3 The animals were anesthetized via an intraperitoneal injection of pentobarbital (75 mg kg⁻¹)
4 and subjected to a contusion injury (70 kilodyne) at the 10th thoracic level using an Infinite
5 Horizons Impactor (Precision Systems Instrumentation)³². During the period of recovery from
6 anesthesia, the mice were placed in a temperature-controlled chamber until thermoregulation
7 was reestablished.

8

9 **Flow cytometry and cell sorting**

10 The spinal cord samples (4 mm in length, centered around the lesion) were dissociated in
11 collagenase type I (Invitrogen), followed by an analysis using a cell sorter, SH800 (Sony), as
12 described previously^{23,55}. GFP-positive RAs isolated from the injured spinal cord of *Nes-EGFP*
13 mice were used for transplantation.

14

15 **Cell transplantation**

16 Using a stereotaxic injector (KDS 310; Muromachi-kikai), a 25- μ l Hamilton syringe
17 (Hamilton) with a glass tip was inserted into the injured spinal cord (1 mm rostral to the lesion
18 epicenter) or the naïve spinal cord. A total volume of 2 μ l of cell suspension (2.0×10^4 viable
19 cells per microliter) was injected at a rate of 0.5 μ l min⁻¹.

20

21 **Behavioral analyses**

22 The motor function was evaluated with a locomotor open-field rating scale on the Basso Mouse
23 Scale (BMS)³². A footprint analysis was also performed, as described previously³². The
24 forelimbs and hindlimbs of the mice were dipped in red and green dyes, respectively. For the

1 grip walk test, a grid walk was reconstructed for mice using two parallel pieces of wood. After
2 3 days of training, each mouse was evaluated using 50 cm of the grid with three patterns: easy
3 (50 steps, 1 cm apart), medium (every third step removed), and hard (every other step
4 removed)²⁴. The sum of the number of grips for all three patterns was calculated in the analysis.
5 All tests were performed in a double-blinded fashion.

6

7 **Histopathological examination**

8 After the animals were reanesthetized and transcardially fixed with 4% paraformaldehyde, the
9 spinal cord was removed, dehydrated, and embedded in an OCT compound. The frozen
10 sections were cut in the sagittal plane at 16 μm or the axial plane at 20 μm . The sections were
11 subsequently stained with primary antibodies in the blocking solution at 4 °C overnight and
12 then incubated with Alexa Fluor secondary antibodies 1 (1:200; Invitrogen) and Hoechst 33258.
13 The following primary antibodies were used: GFAP (1:200, rabbit; Dako, Z0334), GFAP
14 (1:200, rat; Life Technologies, 130300), Nestin (1:200, rat; a gift from Dr. Taro Tachibana,
15 Osaka City University, Osaka, Japan), β -catenin (1:200, mouse; Millipore, 05-665), N-cadherin
16 (1:200, mouse; BD Transduction Laboratories, 610921), Sox9 (1:200, rabbit; Abcam, ab26414),
17 COL1A1 (1:200, mouse; Sigma-Aldrich, SAB1402151), GAP-43 (1:200, rabbit; Novus
18 Biologicals, NB300-143), serotonin (5-HT; 1:200, goat; ImmunoStar, 20079), tyrosine
19 hydroxylase (TH; 1:200, mouse; Millipore, MAB318), platelet-derived growth factor receptor
20 β (PDGFR β ; pericyte marker, 1:200, rabbit; Abcam, ab32570), ER-TR7 (fibroblast marker,
21 1:200, rat, Santa Cruz Biotechnology, sc-73355). The GAP43-positive, 5-HT-positive or TH-
22 positive area per $4.0 \times 10^5 \mu\text{m}^2$ at 42 dpi was measured using the BZ-II analyzer software
23 program (Keyence). We randomly selected 20 cells in each sample and measured the astrocyte
24 cell diameter or ratio of cells with overlapping of neighboring astrocytic processes using the

1 Image J software program (U.S. National Institutes of Health). All images were captured using
2 a BZ-9000 digital microscope system (Keyence) or epifluorescence microscope equipped with
3 a digital camera (BX51; 12 Olympus).

5 **Laser-capture microdissection**

6 Fresh spinal cords were immediately frozen in dry ice/hexane and stored in a deep freezer at -
7 80 °C. The tissues were cut at 16 µm using a cryostat at -20 °C and mounted on PEN membrane
8 slides. The sections were then fixed in ice-cold acetone for 2 min and stained with the antibody
9 against GFAP (1:50, rabbit; Dako) for 5 min. GFP-positive RAs and GFAP-positive astrocytes
10 were dissected with an LMD 6500 system (Leica Microsystems) and transferred by gravity into
11 separate microcentrifuge tube caps placed directly beneath the section. The tube caps were then
12 filled with 75 µl of RLT buffer (Qiagen). For each sample, 1000 cells were dissected from 1
13 series of sagittal sections and subjected to RNA extraction.

15 **Reverse transcription (RT)-PCR**

16 Total RNA was isolated from the astrocytes obtained from the spinal cord tissue using the
17 RNeasy Micro Kit (Qiagen) or from the injured spinal cord (4 mm in length) using the RNeasy
18 Mini Kit (Qiagen). For the complementary DNA (cDNA) synthesis, a reverse transcription
19 reaction was performed using a PrimeScript first-strand cDNA Synthesis Kit (Takara Bio).
20 Quantitative RT-PCR was performed using primers specific to the genes of interest (Table S1)
21 and SYBR Premix Dimmer-Eraser (Takara Bio). The data were normalized to the expression
22 of glyceraldehyde-3-phosphate dehydrogenase. RT-PCR was conducted using a Thermocycler
23 (Biometra), and the products were detected by electrophoresis and ethidium bromide staining.

24

1 **RNA sequencing**

2 The sample preparation and data analyses were performed as described previously⁵⁶. In brief,
3 the mRNA-seq library was prepared using the NEBNext Ultra Directional Library Prep Kit for
4 Illumina (NEB) and sequenced on an Illumina HiSeq-1500 system. The sequence reads were
5 aligned to the mouse reference genome (mm9, build 37), and the gene expression (fragments
6 per kilobase of exon per million mapped sequence reads; FPKM) was calculated using the
7 TopHat (version 2.0.11) and Cufflinks (version 2.1.1) software programs⁵⁷. The selected genes
8 of interest were converted into lists of GO terms on GO database annotations with DAVID⁵⁸.
9 The calculated $-\log_{10}(P\text{-values})$ are displayed on the x-axis of the graph.

10

11 **The sequencing and data analysis (CEL-Seq2)**

12 We performed CEL-Seq2 for the whole transcriptome analysis of RAs (isolated by FACS at 7
13 dpi), transplanted and host astrocytes (both isolated by LMD at 7 dpi), in accordance with a
14 previously described protocol⁵⁹. We sequenced the libraries using an Illumina MiSeq system.
15 The reads were mapped to the mouse reference genome (mm10) using the Bowtie 2 software
16 program (version 2.2.6)⁶⁰. Pearson's correlation coefficients were calculated using the common
17 logarithm of the read counts for genes with non-zero counts in at least one sample.
18 Differentially expressed genes (DEGs) were extracted using the DESeq2 library (version
19 1.10.1) for R (with adjusted p-values of < 0.1)⁶¹.

20

21 **Anti- β 1 Ab injection**

22 SCI mice received an injection of 1 μ g function-blocking monoclonal antibody against β 1-
23 integrin (anti- β 1 Ab; BD Pharmingen, 555003) at the epicenter using a stereotaxic injector
24 (KDS 310; Muromachi-kikai) every 2 days from 9 to 13 dpi. Control mice were given an

1 equivalent amount of isotype control (control Ab; BD Pharmingen, 553958). For *in vitro*
2 blocking of collagen-integrin signaling, anti- β 1 Ab or control Ab was added at $1 \mu\text{g ml}^{-1}$, and
3 an immunocytological analysis and mRNA extraction were performed at 12 h after
4 administration.

5

6 **The administration of N-cadherin nAb**

7 We injected $10 \mu\text{g}$ of neutralizing monoclonal antibody against N-cadherin (N-cadherin nAb;
8 Sigma-Aldrich, C3865) at the lesion epicenter in SCI mice using a stereotaxic injector (KDS
9 310; Muromachi-kikai) every 2 days from 9 to 13 dpi. Control mice received an equivalent
10 amount of isotype control IgG (control IgG; BioXCell, BE0083). For the *in vitro* N-cadherin
11 blockade experiments, N-cadherin nAb or control IgG was added at $10 \mu\text{g ml}^{-1}$, and an
12 immunocytological analysis was performed at 12 h after administration.

13

14 **Statistical analysis**

15 Wilcoxon's rank-sum test was used to compare the medians of the data of qPCR, the cell
16 diameter, the ratio of overlapping-positive cells, the GAP43-positive, 5-HT-positive or TH-
17 positive area, and the functional outcome scores. For multiple comparisons in qPCR, an
18 analysis of variance (ANOVA) with the Tukey-Kramer post hoc test was applied. For the
19 analysis of the differences in the BMS scores between the groups over time, a two-way repeated
20 measures ANOVA with the Tukey-Kramer post hoc test was performed. In all statistical
21 analyses, the level of significance was set at 0.05. The values for groups are presented as the
22 average \pm SEM. All of the statistical analyses were carried out using the JMP software program
23 (version 11; SAS Institute).

24

1 **Data availability**

2 RNA-seq (whole spinal cords) and CEL-Seq2 (isolated astrocytes) data are deposited with

3 accession codes DRA004535 (DDBJ) and DRA005561 (DDBJ), respectively.

4

5

6

7

1 **Methods-only References**

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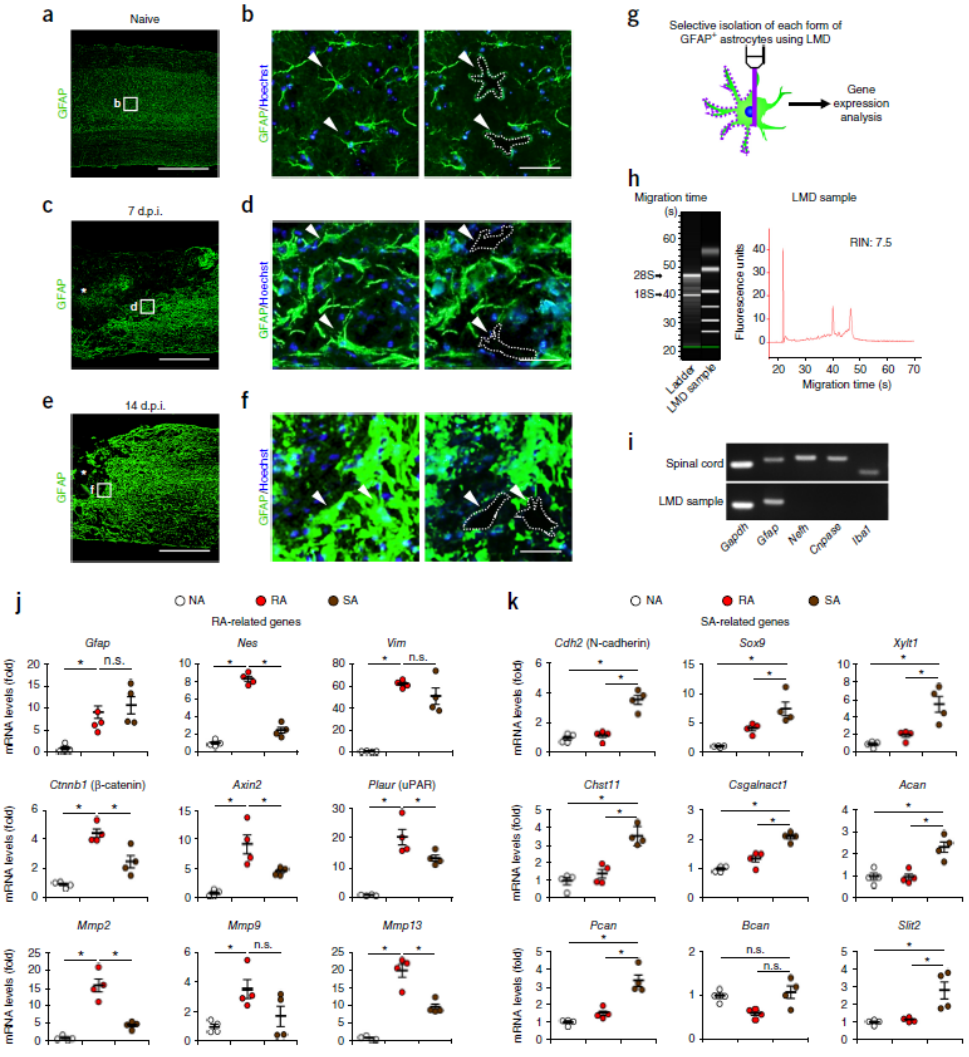


Figure 1.

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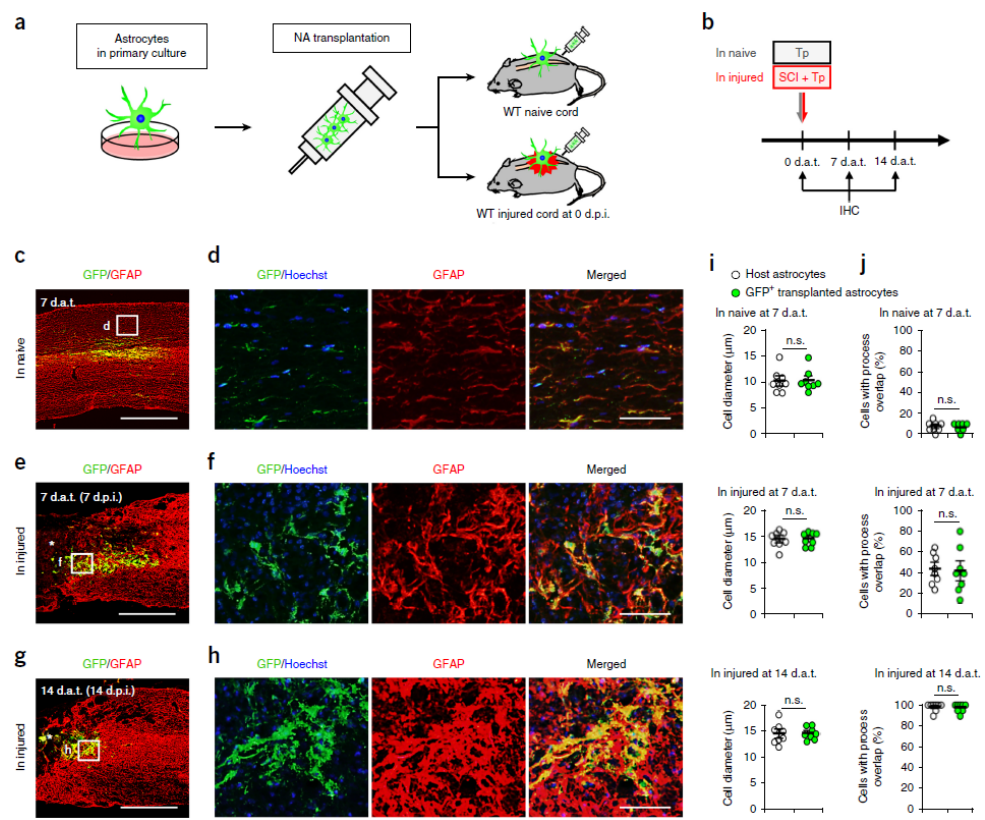


Figure 2.

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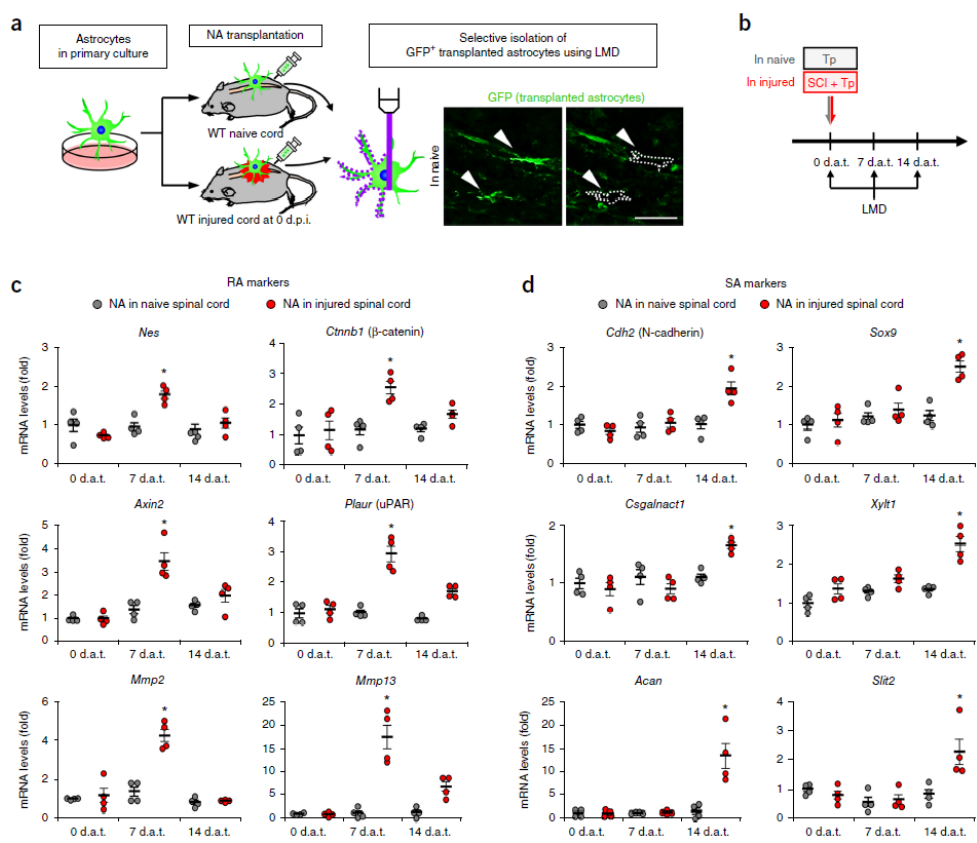


Figure 3.

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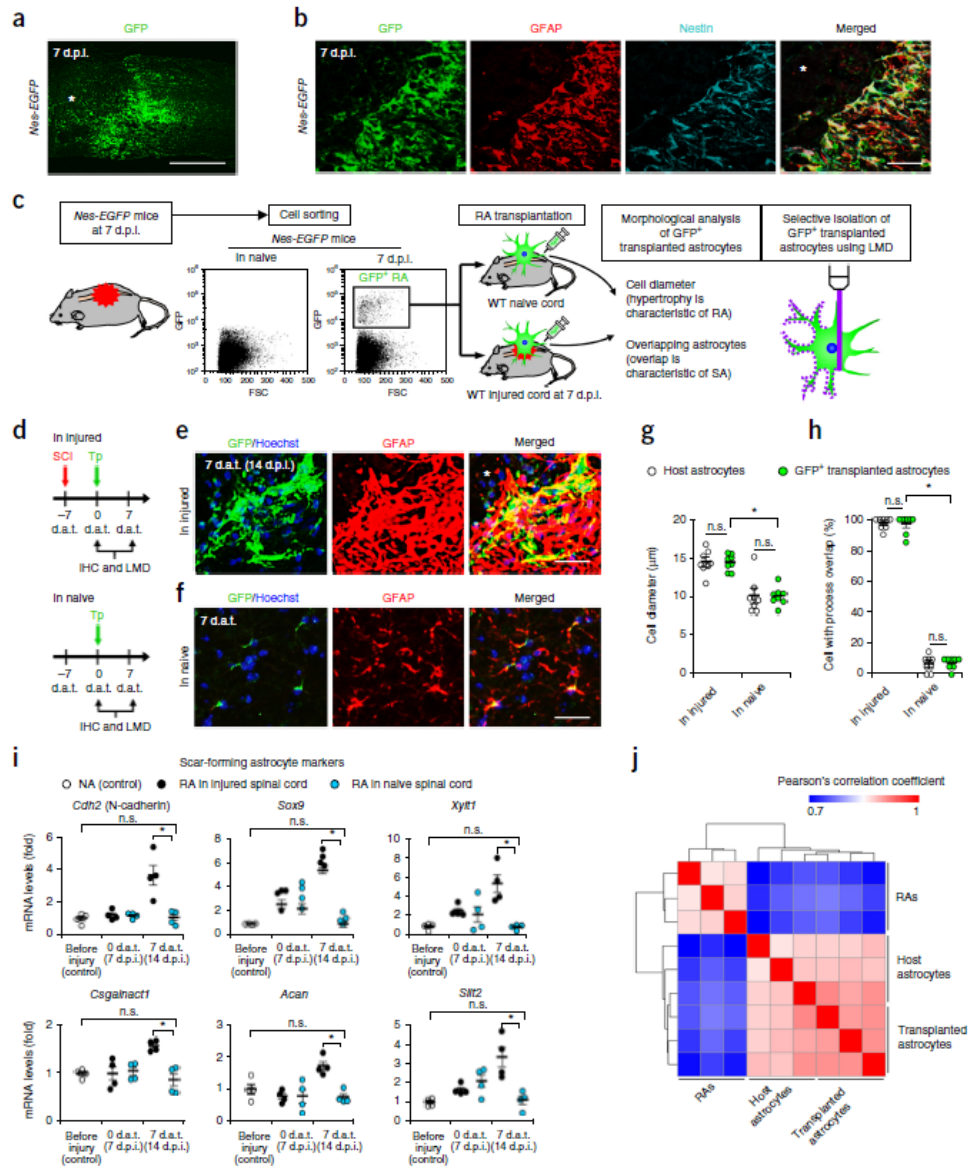


Figure 4.

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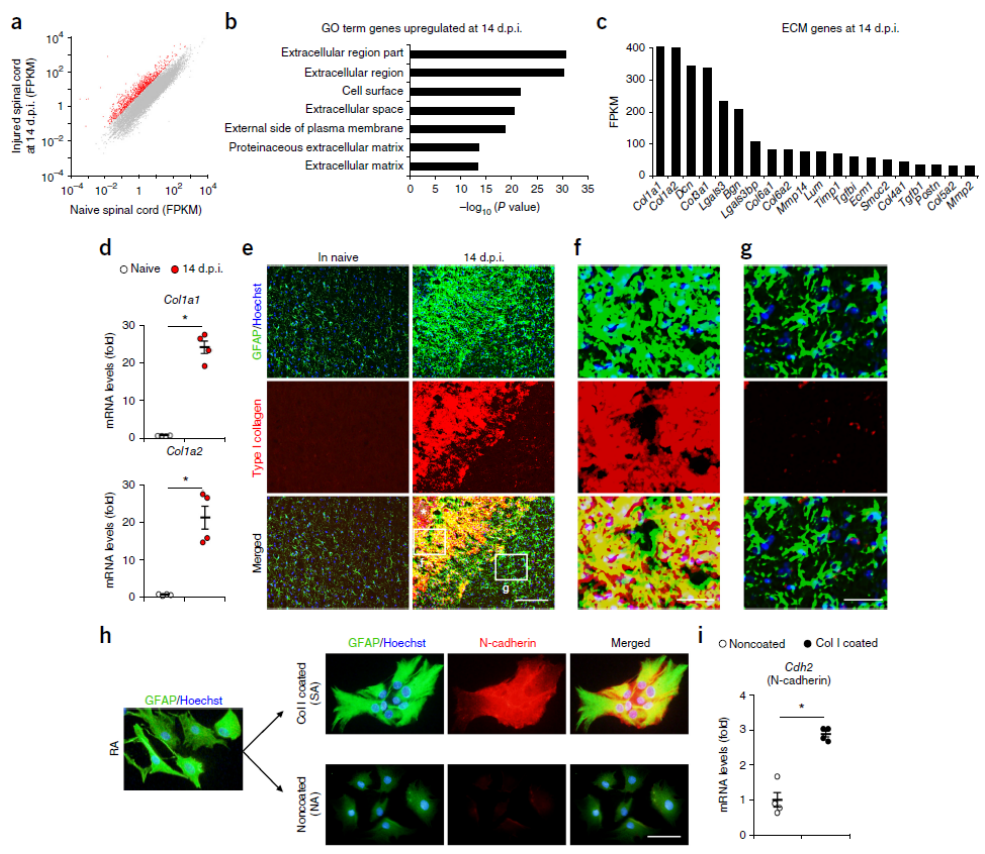


Figure 5.

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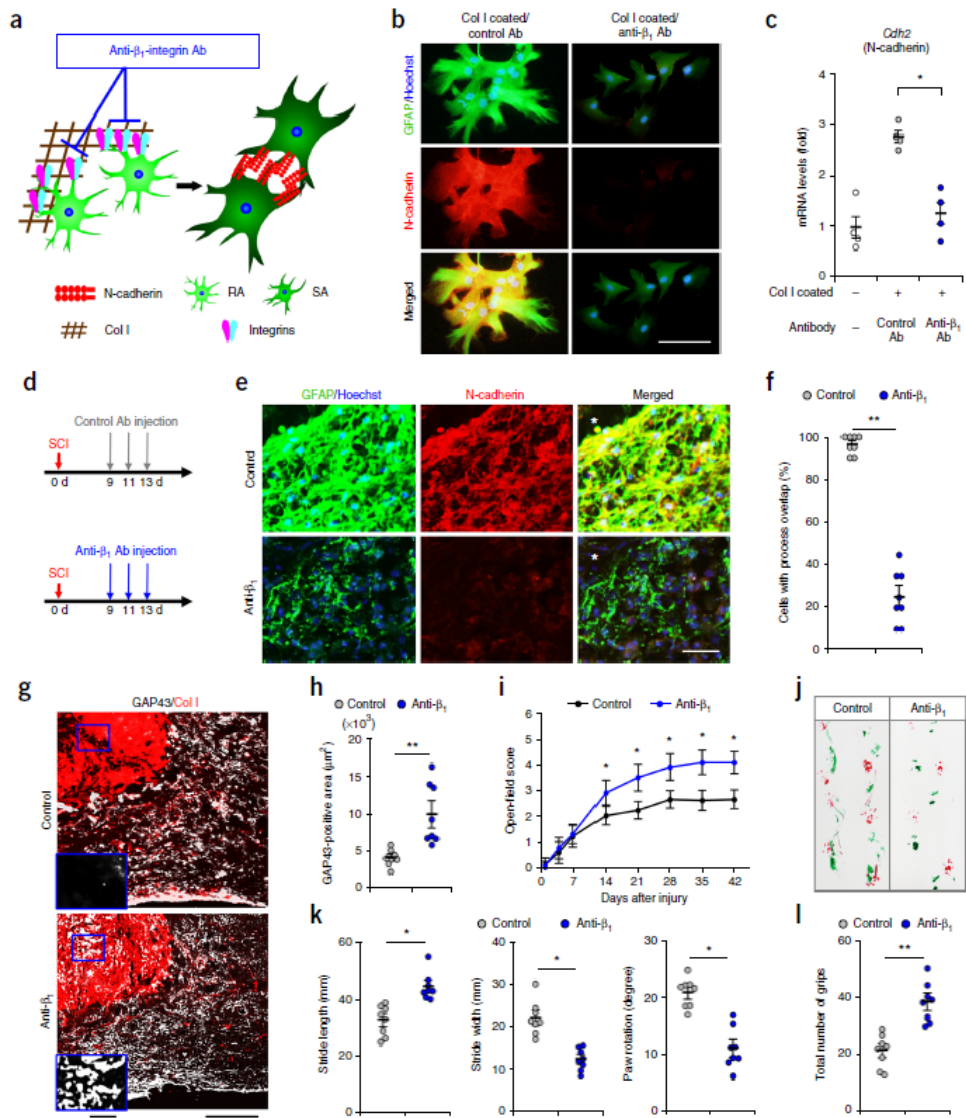
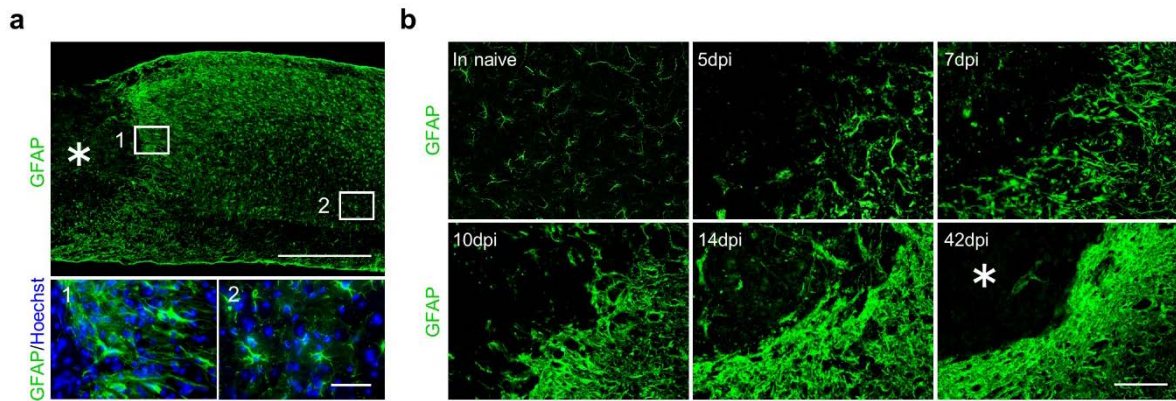


Figure 6.

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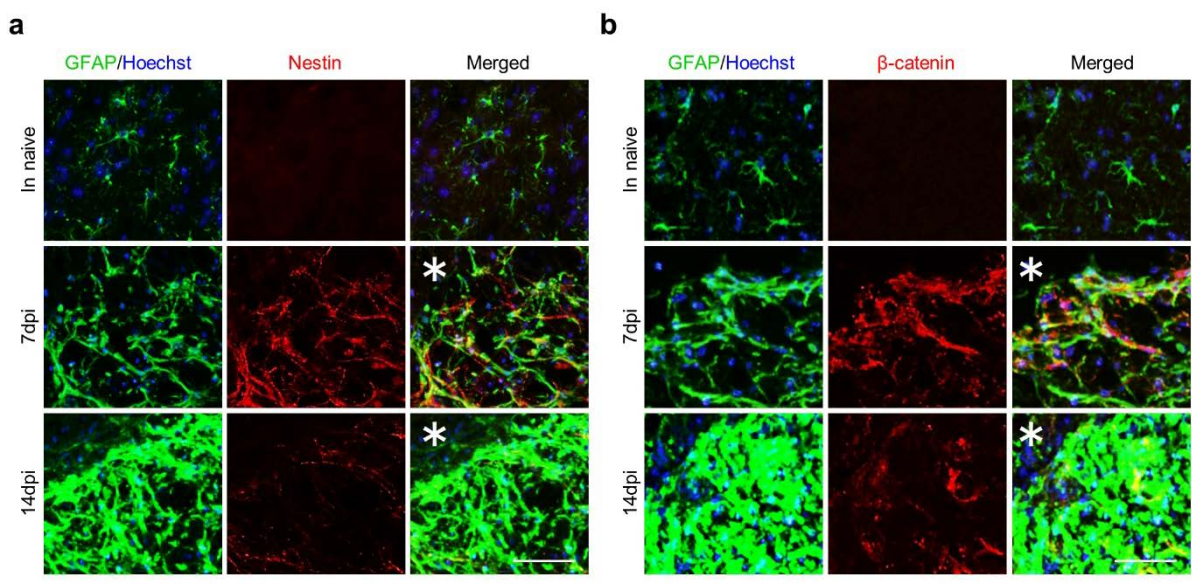
Supplementary Figure 1



Supplementary Figure 1. Dramatic changes in the astrocytic morphology around the lesion area. (a) Example images depicting GFAP (green) staining of the injured spinal cords at 7 dpi after SCI. The asterisk indicates the lesion epicenter in SCI mice. The inset (1) and (2) are magnifications of the boxed areas in (a). Scale bars, 500 μm, insets 50μm. **(b)** Example images depicting time course of GFAP-positive astrocytes (green) before and after SCI. The asterisk indicates the lesion epicenter in SCI mice. Scale bar, 100 μm.

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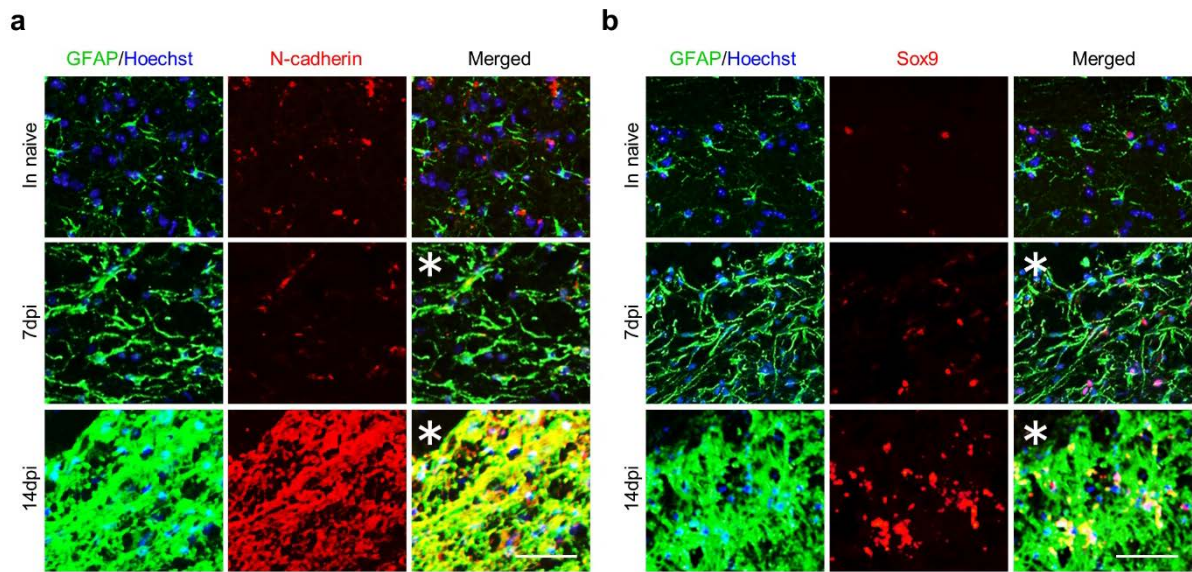
Supplementary Figure 2



Supplementary Figure 2. RAs expressing their hallmark proteins. (a and b) Example images depicting immunohistological analysis of the GFAP-positive astrocytes (green) stained with the RA markers Nestin or β -catenin (red) before injury and at 7 and 14 dpi. The nuclei are counterstained with Hoechst (blue). The asterisk indicates the lesion epicenter in SCI mice. Scale bars, 50 μ m.

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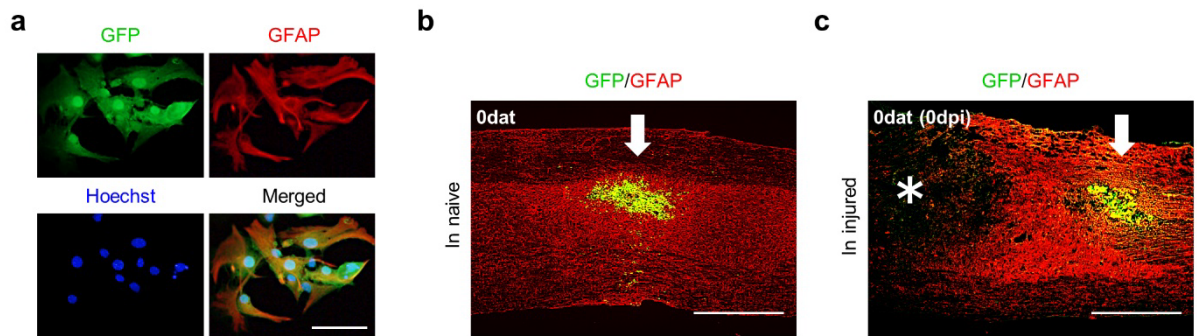
Supplementary Figure 3



Supplementary Figure 3. SAs expressing their hallmark proteins. (a and b) Example images depicting immunohistological analysis of the GFAP-positive astrocytes (green) stained with the SA markers N-cadherin or Sox9 (red) before injury and at 7 and 14 dpi. The nuclei are counterstained with Hoechst (blue). The asterisk indicates the lesion epicenter in SCI mice. Scale bars, 50 μ m.

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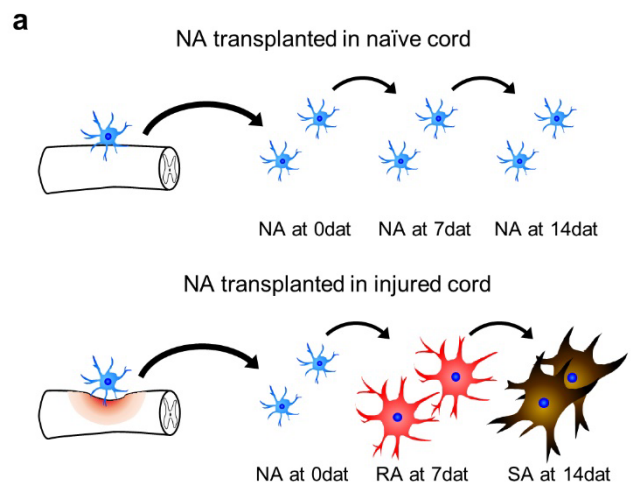
Supplementary Figure 4



Supplementary Figure 4. Transplantation of NAs into naïve and injured spinal cords. (a) Example images depicting immunocytochemical analysis of primary astrocyte cultures prepared from *CAG-EGFP* mice, stained with GFP (green), GFAP (red), and Hoechst (blue). Scale bar, 50 μ m. (b and c) Example images depicting immunostaining of GFP-positive NAs (green) transplanted into the naïve (b) and injured spinal cord (c) at 0 dat, stained with GFAP (red). The arrows indicate the transplantation points of GFP-positive NAs. The asterisk indicates the lesion epicenter in SCI mice (c). Scale bars, 500 μ m.

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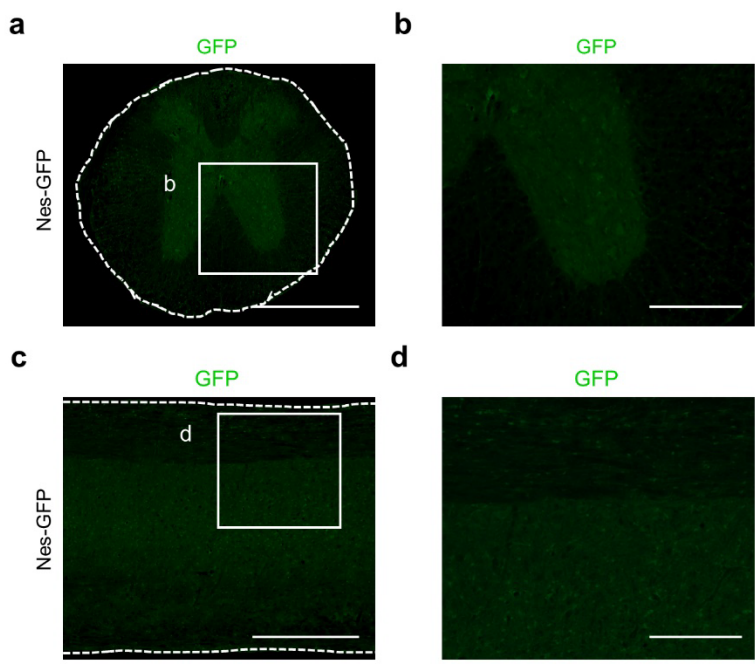
Supplementary Figure 5



Supplementary Figure 5. Environment-dependent transformation of NAs into naïve and injured spinal cords. (a) A schematic representation of the environment-dependent transformation of the grafted NAs.

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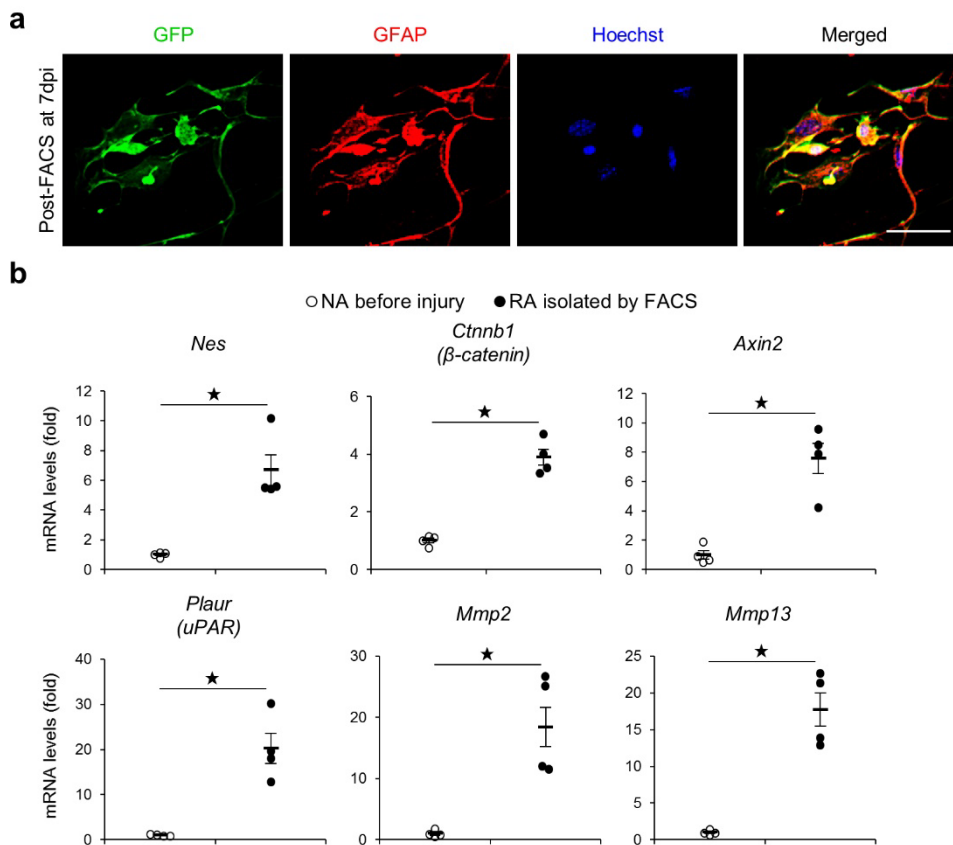
Supplementary Figure 6



Supplementary Figure 6. No GFP-positive cells in the naïve spinal cord of *Nes-EGFP* mice. (a-d) Example images depicting GFP (green) staining of the sagittal (a and b) and axial (c and d) sections of the naïve spinal cord of *Nes-EGFP* mice. No GFP-positive cells are present. The images (b) and (d) are magnifications of the boxed areas in (a) and (c), respectively. Scale bars, 500 μ m (a and c), 200 μ m (b and d).

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Supplementary Figure 7



Supplementary Figure 7. The direct isolation and gene expression analysis of GFP-positive RA from *Nes-EGFP* mice. (a) Example images depicting immunocytochemical analysis of GFP-positive RAs (green) isolated from the injured spinal cord of *Nes-EGFP* mice at 7 dpi using FACS, stained with GFAP (red) and Hoechst (blue). Scale bar, 50 μ m. (b) Quantification of mRNA expression levels of RA marker genes in NAs before injury and RAs isolated by FACS ($n = 4$ mice per group). $\star P < 0.05$, Wilcoxon's rank sum test. Data represent mean \pm SEM.

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Supplementary Figure 8

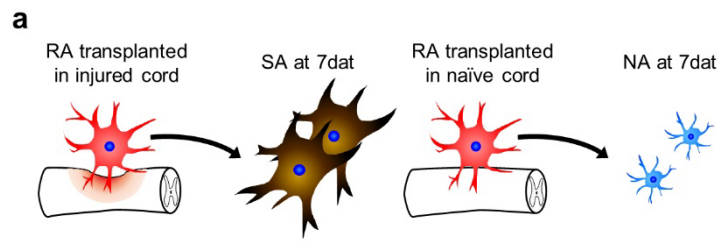
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	# of DEGs	
Host astrocytes versus Transplanted astrocytes	0	
RAs versus Host astrocytes	2386	} 2060 (# of common DEGs)
RAs versus Transplanted astrocytes	2670	

Supplementary Figure 8. The whole transcriptome analysis of RAs, transplanted astrocytes and host astrocytes. (a) The number of significant differentially expressed genes (DEGs) between RAs, transplanted astrocytes and host astrocytes. There were no significant DEGs between transplanted and host astrocytes (with adjusted p-values of < 0.1).

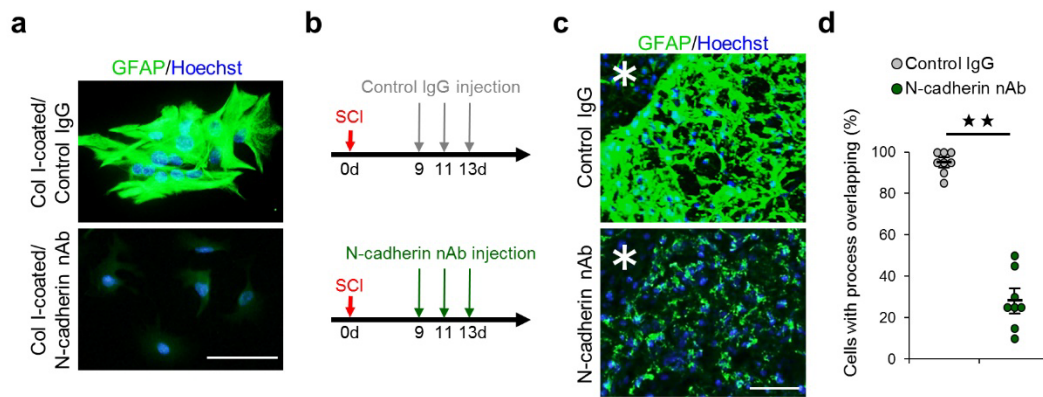
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Supplementary Figure 9



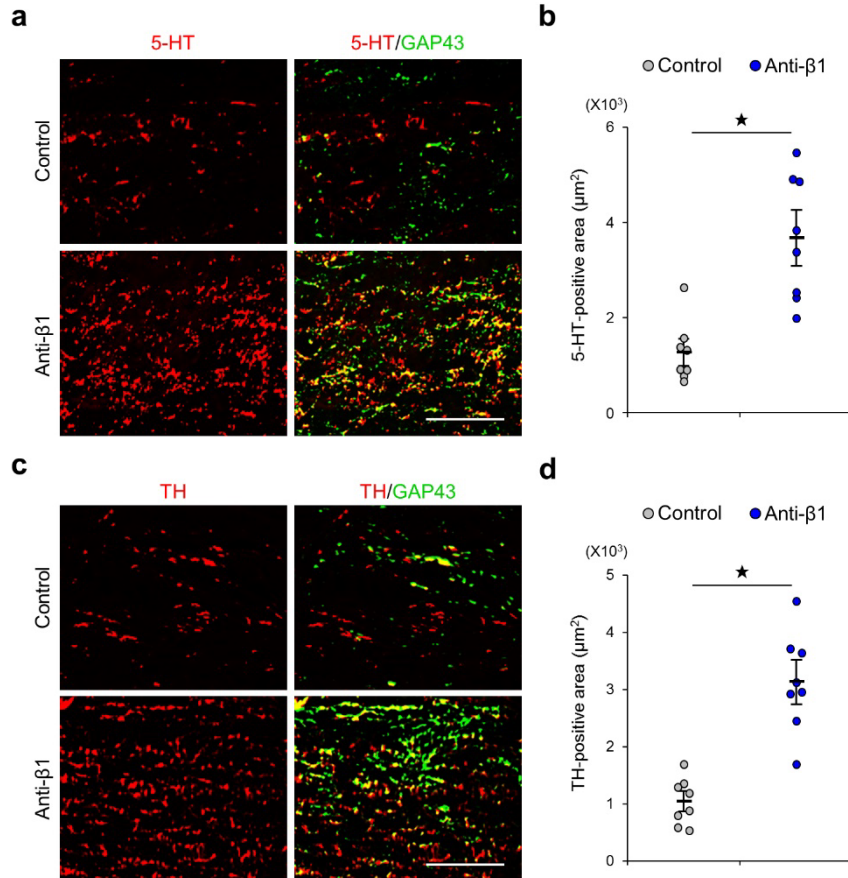
Supplementary Figure 9. Environment-dependent reversibility of reactive astrogliosis. (a) A schematic representation of the environment-dependent reversibility of reactive astrogliosis.

Supplementary Figure 10



Supplementary Figure 10. The functional blockade of N-cadherin inhibited the transformation of RAs into SAs and attenuated astrocytic scar formation after SCI. (a) Example images depicting immunocytochemical analysis of RAs on Col I-coated dishes after the administration of N-cadherin nAb or control IgG, stained with GFAP (green) and Hoechst (blue). Scale bar, 50 μm. (b) The schedule of the injections of N-cadherin nAb or control IgG into WT mice after SCI in the *in vivo* experiments. (c) Example images depicting immunostaining of the lesion area of the injured spinal cord receiving N-cadherin nAb or control IgG injection at 14 dpi, stained with GFAP (green) and Hoechst (blue). The asterisk indicates the lesion epicenter in SCI mice. Scale bar, 50 μm. (d) The ratio of GFAP-positive astrocytes with overlapping of neighboring astrocytic processes ($n = 8$ mice per group). ** $P < 0.005$, Wilcoxon's rank sum test. Data represent mean \pm SEM.

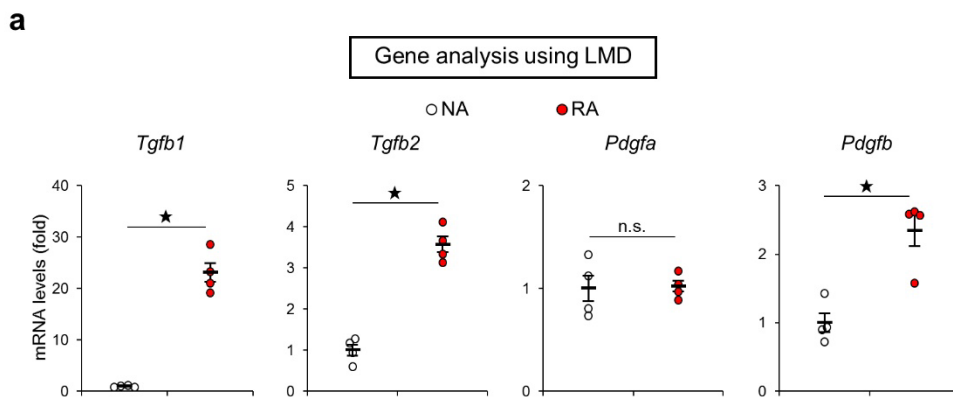
Supplementary Figure 11



Supplementary Figure 11. Increased numbers of 5-HT-positive and TH-positive axons were observed in the area caudal to the lesion epicenter. (a and c) Example images depicting immunostaining of the area caudal to the lesion epicenter of the injured spinal cord receiving anti-β1 Ab or control Ab injection at 42 dpi, stained with GAP43 (green) and 5-HT or TH (red). Scale bars, 100 μm. (b and d) Quantification of the 5-HT-positive or TH-positive area per $4.0 \times 10^5 \mu\text{m}^2$ (n = 8 mice per group). *P < 0.05, Wilcoxon's rank sum test. Data represent mean ± SEM.

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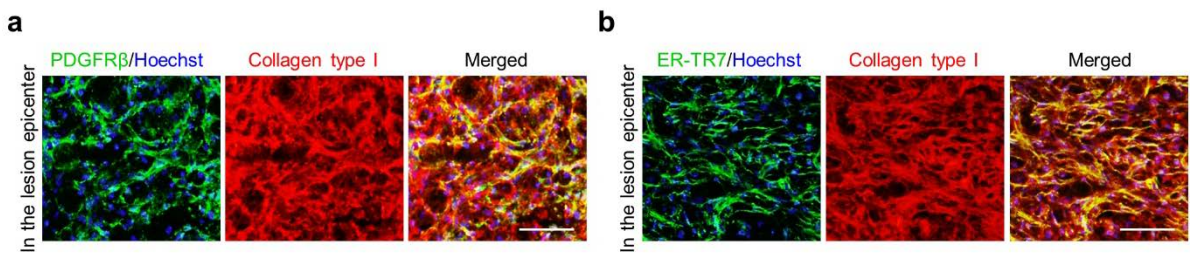
Supplementary Figure 12



Supplementary Figure 12. Upregulation of neuroprotective factors in RAs. (a) Quantification of mRNA expression levels of *Tgfb1*, *Tgfb2*, *Pdgfa*, and *Pdgfb* in NAs and RAs ($n = 4$ mice per group). ★ $P < 0.05$, Wilcoxon's rank sum test. Data represent mean \pm SEM.

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Supplementary Figure 13



Supplementary Figure 13. Pericytes and fibroblasts produced Col I in the lesion epicenter after SCI. (a and b) Example images depicting immunohistochemical analysis of the injured spinal cord of WT mice at 14 dpi, stained with collagen type I (red), Hoechst (blue), and PDGFR β or ER-TR7 (green). Scale bars, 50 μ m.

Supplementary Table 1

Primers used for qPCR

Gene symbol	Accession number	5'- Forward primer -3'	5'- Reverse primer -3'
Acan	NM_007424.2	tftgattcccaccgtgcctttcc	ttcctggctcctgtcttcttcagc
Axin2	NM_015732.4	gagaagttggattgtgtccacg	accatttctgcatgtgtcatgg
Bcan	NM_001109758.1	aattctgctgaaggctcaatgcc	cggaagtgcagaatggaagatcc
Cdh2	NM_007664.4	tacgcagctggtgcagataaagg	tctgactcctccatagtctatgc
Chst11	NM_021439.2	gttctggtgaagtccacaaactgc	tgtcaccatgggattctacacacg
Cnpase	NM_001146318.1	aatggcagaccagtatcagtacc	gtctcagaactcttttggcagg
Col1a1	NM_007742.3	catggagacaggtcagacctgtgt	ggacattaggcgcaggaaggctcag
Col1a2	NM_007743.2	atccaac taagtctcct ccctgg	ggcttctgacta tctccacag ag
Csgalnact1	NM_001252623.1	cttgagacagtctgtcacagagc	cagtccttagatcagatctccagg
Ctnnb1	NM_001165902.1	gggtgaatactttactctgctgc	gtataacgctgcaaaagctgtggc
Gapdh	NM_008084.3	gactcaacagcaactcccactct	ggtttctactcctggaggccat
Gfap	NM_001131020.1	tgtactaacagagcgagcctatgc	gggacttgctgccttaacattgg
Iba1	NM_019467.2	caaagaacacaagaggccaactgg	ttcatgctgctgtcatcagaagc
Mmp13	NM_008607.2	gagagcttagttctgtgaacgagc	aaagcagatggaccccatgtttgc
Mmp2	NM_008610.2	ctatcatctcatcgctgcacacc	gtacagtcagcaccttctttggg
Mmp9	NM_013599.3	aaggattcagttgccctactgg	acacggagaatctctgagcaatcc
Nefh	NM_010904.3	tagcaagagaagataaccctgagc	tcatctgtcagttggacatacagg
Nes	NM_016701.3	gtcagctgagcctatagtccaacg	agagtcactcatcattgtgtctcc
Pcan	NM_001081306.1	taatggtgcagcttgctgatgg	cctgacagtaactcattctgtctgc
Pdgfa	NM_008808.3	agacagatgtgaggtgagatgagc	acggaggagaacaaagaccgcacg
Pdgfb	NM_011057.3	tacctcactctgtgtcttctcc	catcccattacaacctgtctcacc
Plaur	NM_011113.3	tctggatcttcagagcttccacc	cttacggataactccgggttccc
Slit2	NM_001291227.1	cgtctctagaagcttctagcttcg	tgtagggggagctttagtacaagc
Sox9	NM_011448.4	gaaggtaacgattgtctggattcc	cgctcctcatgttaactctgaagg
Tgfb1	NM_011577.1	gtgacagcaagataacaaactcc	gagctgaagcaatagttggtatcc
Tgfb2	NM_009367.3	tctgagattacagcaacaacaacc	caatacgtacaactcactgaacg
Vim	NM_011701.4	tgtaactaccaggacactattgg	aggtagttctctcaggttcagg
Xylt1	NM_175645.3	cagtgaagattctccatcactggg	tctgaaactctgtccatgtagg