Angiotensin II receptor blockade promotes repair of skeletal muscle through down-regulation of aging-promoting C1q expression

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Disruption of angiotensin II type 1 (AT1) receptor prolonged life span in mice. Since aging-related decline in skeletal muscle function was retarded in Atgr1a−/− mice, we examined the role of AT1 receptor in muscle regeneration after injury. Administration of AT1 receptor blocker irbesartan increased the size of regenerating myofibers, decreased fibrosis, and enhanced functional muscle recovery after cryoinjury. We recently reported that complement C1q, secreted by macrophages, activated Wnt/β-catenin signaling and promoted aging-related decline in regenerative capacity of skeletal muscle. Notably, irbesartan induced M2 polarization of macrophages, but reduced C1q expression in cryoinjured muscles and in cultured macrophage cells. Irbesartan inhibited up-regulation of Axin2, a downstream gene of Wnt/β-catenin pathway, in cryoinjured muscles. In addition, topical administration of C1q reversed beneficial effects of irbesartan on skeletal muscle regeneration after injury. These results suggest that AT1 receptor blockade improves muscle repair and regeneration through down-regulation of the aging-promoting C1q-Wnt/β-catenin signaling pathway.

Angiotensin (Ang) II is the crucial bioactive molecule of the renin-angiotensin system, and exerts most of its pathophysiological actions through binding to Ang II type 1 (AT1) receptor1,2. While humans have a single AGTR1 gene that encodes AT1 receptor, mice have Agtr1a and Agtr1b genes encoding two isoforms (AT1a and AT1b) of AT1 receptor, and the major isoform of mouse AT1 receptor is AT1a receptor1,2. AT1 receptor is a member of the G protein-coupled receptor family, which shares typical conformation of seven transmembrane-spanning α-helices3. Upon stimulation by binding to Ang II or mechanical stretch4,5, AT1 receptor activates multiple G protein-dependent and -independent signaling pathways and promote the formation of reactive oxygen species (ROS), leading to a variety of responses such as...
cellular hypertrophy and proliferation, vascular contraction, inflammatory responses, and salt and water retention.

Besides homeostatic regulation of blood pressure and electrolyte and water balance, AT1 receptor plays physiological roles in normal organ development. Targeted disruption of both Agtr1a and Agtr1b genes in mice resulted in abnormal kidney development characterized by tubular atrophy and interstitial expansion, papillary atrophy, and severe impairment of concentrating urine. In this sense, AT1 receptor is essential during embryogenesis and beneficial for organismal survival. However, sustained and excessive activation of AT1 receptor is detrimental, promoting various aging-related diseases such as cardiovascular diseases, diabetes, chronic kidney disease, dementia, osteoporosis, and cancer. Furthermore, recent studies unraveled the involvement of AT1 receptor in aging process per se. Targeted disruption of Agtr1a gene prolonged life span in mice, which was associated with prevention of aging-related progression of cardiac hypertrophy and fibrosis. Accordingly, AT1 receptor plays antagonistic and pleiotropic roles according to the ages and pathophysiological conditions.

Benigni A. et al. reported that Agtr1a−/− mice showed a decrease in oxidative stress, an increase in mitochondrial number, and an increase in expression levels of prosurvival genes such as Nampt and Sirt3, which in combination might contribute to a prolongation of life span. However, it remains precisely unknown how AT1 receptor regulates life span. We observed that aging-related decline in skeletal muscle function was prominently milder in Agtr1a−/− mice, as compared with Agtr1a+/+ littermate mice, and that systemic administration of an AT1 receptor blocker (ARB) irbesartan significantly enhanced regression of cardiac hypertrophy and fibrosis. Accordingly, AT1 receptor plays antagonistic and pleiotropic roles according to the ages and pathophysiological conditions.

Results

Aging-related decline in skeletal muscle function was milder in Agtr1a−/− mice. First, we performed a prospective observational study in 66 Agtr1a−/− mice and 55 Agtr1a+/+ littermate mice, and found that Agtr1a−/− mice significantly lived longer than Agtr1a+/+ mice (Supplementary Fig. 1) as previously reported. The average life span of Agtr1a−/− and Agtr1a+/+ mice was 760.0 ± 20.9 and 651.8 ± 21.7 days, respectively (P < 0.05). In association with the elongation of life span, aging-related changes were significantly milder in multiple tissues of Agtr1a−/− mice. For example, aging-related thinning of epidermal and fat layers was milder in Agtr1a−/− mice (Supplementary Fig. 2a). After shaving of dorsal hair in the same area, hair growth was significantly more rapid in Agtr1a−/− mice than in Agtr1a+/+ mice at the age of 18 months, although it was comparable at the age of 3 months (Supplementary Fig. 2b). Especially, aging-related changes in locomotive activity and coordination skill were milder in Agtr1a−/− mice. In both vertical pole test and hanging wire test, the time to fall off was significantly longer in Agtr1a+/+ mice at the age of 12 and 18 months, although it was comparable at the age of 3 months (Supplementary Fig. 2a,b). In addition, aging-associated fibrosis in skeletal muscle at the age of 24 months was significantly less severe in Agtr1a−/− mice, as revealed by Masson’s trichrome staining (Supplementary Fig. 3c). These results suggest that aging-related decline in skeletal muscle function is milder in Agtr1a−/− mice.  

Irbesartan enhanced repair and functional recovery of skeletal muscle after cryoinjury. Since impairment of regeneration potential is one of the most important features observed during aging of skeletal muscles, 19, we investigated the role of AT1 receptor signaling in skeletal muscle regeneration in response to injury. We previously reported that AT1 receptor signaling regulates the hypothalamic neurocircuit that is involved in the control of food intake, and that Agtr1a−/− mice were hyperphagic and obese with increased adiposity on an ad libitum diet, as compared with Agtr1a+/+ mice. To avoid the metabolic effects on skeletal muscle regeneration in Agtr1a−/− mice, we cryoinjured tibialis anterior (TA) muscle of wild-type mice and examined the effects of treatment with an AT1 receptor blocker irbesartan on muscle regeneration after injury. We first confirmed that orally administered irbesartan (20 mg/kg/d) had no effect on daily food intake and body weight (Supplementary Fig. 4), as well as heart rate and blood pressure (Supplementary Table 1).

We next compared repair and functional recovery of skeletal muscle between irbesartan- and vehicle-treated mice. Irbesartan-treated mice showed a significantly higher weight of TA muscle than vehicle-treated mice at 4 d after cryoinjury, although the TA weight-to-tibia length ratios were comparable at 14 d (Fig. 1a). Total running distance during treadmill testing at 14 d was significantly longer in irbesartan-treated mice than in vehicle-treated mice (Fig. 1b). Histological analysis revealed that the size of centronuclear regenerating myofibers at 14 d was significantly larger in irbesartan-treated mice than in vehicle-treated mice (Fig. 1c,d). In addition, the number and size of embryonic myosin heavy chain (eMHC)-positive myofibers at 10 d were significantly increased in irbesartan-treated mice, as compared with vehicle-treated mice (Fig. 1e,f). Fibrotic area in skeletal muscle at 14 d after injury was significantly smaller in irbesartan-treated mice than in vehicle-treated mice, as revealed by Masson’s trichrome staining (Fig. 2a,b). qRT-PCR analysis also showed a significant decrease in mRNA expression.
of fibrosis-related genes such as \(\text{Tgfb1, Postn, Colla1, and Col3a1}\) in TA muscles of irbesartan-treated mice at 10 d after injury, as compared with vehicle-treated mice (Fig. 2c). These results suggest that AT\(_1\) receptor blockade promotes myogenic growth after injury.

**Irbesartan increased Pax7+ satellite cells in regenerating skeletal muscle after cryoinjury.** Satellite cells, characterized by the expression of the paired box protein Pax7, are crucial for skeletal muscle growth and regeneration\(^{19}\). In resting muscles, satellite cells are located juxtaposed to
muscle fibers in a quiescent state, but re-enter the cell cycle and proliferate to form sufficient number of myofibers in response to injury19. Immunohistochemical analysis revealed that the number of Pax7+ satellite cells was significantly increased in TA muscles at 4 d after cryoinjury of irbesartan-treated mice, as compared with vehicle-treated mice (Fig. 3). These results suggest that AT1 receptor blockade promotes activation and cell cycle re-entry of Pax7+ satellite cells following injury.

Irbesartan induced M2 polarization of macrophages in regenerating skeletal muscle after cryoinjury. Macrophages in inflammatory microenvironments have a regulatory role in the muscle response to injury, not only by removing necrotic tissue but also by promoting muscle growth and regeneration21. Histological analysis of TA muscles revealed no significant difference at 6 d, but the number of remaining inflammatory cells at 14 d was significantly lower in irbesartan-treated mice than in vehicle-treated mice (Fig. 4a). Consistently, mRNA levels of Cd68 encoding the macrophage marker CD68 were comparable in TA muscles at 6 d, but Cd68 mRNA levels were significantly lower in irbesartan-treated mice than vehicle-treated mice at 14 d (Fig. 4b), suggesting that inflammatory responses were attenuated in irbesartan-treated mice. Since the M1 to M2 transition of macrophages is important not only for dampening of inflammation, scavenging of debris, and tissue healing, but also for the transition from proliferative to differentiation stages of myogenesis21, we next examined the polarity of infiltrating macrophages at 6 d. Although the total number of CD11b+ and F4/80+ macrophages was unchanged (Fig. 4c,d), flow cytometric analysis revealed that the number of RELMα+ M2 macrophages in CD11b+ and F4/80+ macrophages was significantly higher in TA muscles at 6 d after cryoinjury of irbesartan-treated mice than in vehicle-treated mice (Fig. 4e,f). These results suggest that AT1 receptor blockade induces M2 polarization of macrophages in injured skeletal muscle, and thereby attenuates inflammatory responses and promotes tissue repair.

Irbesartan decreased C1q mRNA expression in macrophages both in vivo and in vitro. Recently, we reported that complement C1q activates canonical Wnt/β-catenin signaling and promotes aging-associated decline in skeletal muscle regeneration18. Since macrophages are one of the major sources of C1q biosynthesis22, we next explored whether C1q expression was altered by treatment with irbesartan. The mRNA levels of C1qa in CD11b+ and F4/80+ macrophages sorted from TA muscles at 6 d after cryoinjury were significantly lower in irbesartan-treated mice than in vehicle-treated
Figure 3. Pax7 + satellite cells in cryoinjured skeletal muscle of irbesartan-treated mice. (a) Immunostaining of TA muscles of mice treated with irbesartan or vehicle at 4 d after cryoinjury or sham operation. Laminin, Pax7 are represented in green and red, respectively, with TO-PRO-3 staining of the nucleus (blue). Scale bar, 20 µm. (b) The number of Pax7 + satellite cells in TA muscles of mice treated with irbesartan or vehicle at 4 d after cryoinjury or sham operation (n = 11, in each group). Data are presented as mean ± SEM. *P < 0.05, **P < 0.01.
Figure 4. M2 polarization of macrophages in cryoinjured skeletal muscle of irbesartan-treated mice. (a) Histological sections with HE staining of TA muscles in mice treated with irbesartan or vehicle at 6 d and 14 d after cryoinjury. Scale bars, 100 μm. (b) The mRNA levels of Cd68 in TA muscles of mice treated with irbesartan or vehicle at 6 d (n = 8, in each group) and 14 d (n = 16, in each group) after cryoinjury. Data are shown as fold induction over sham + vehicle at 6 d (mean ± SEM). **P < 0.01. NS, not significant. (c) Representative flow cytometric analysis demonstrating CD11b+ and F4/80+ macrophages in TA muscles of mice treated with irbesartan or vehicle at 6 d after cryoinjury. (d) The percentage of CD11b+ and F4/80+ macrophages in flow cytometric analysis of TA muscles of mice treated with irbesartan or vehicle at 6 d after cryoinjury. (n = 4, in each group). Data are presented as mean ± SEM. NS, not significant. (e) Representative flow cytometric analysis demonstrating F4/80+ and RELMα+ M2 macrophages in CD11b+ and F4/80+ macrophages sorted from TA muscles of mice treated with irbesartan or vehicle at 6 d after cryoinjury. (f) The percentage of CD11b+, F4/80+, and RELMα+ M2 macrophages in flow cytometric analysis of TA muscles of mice treated with irbesartan or vehicle at 6 d after cryoinjury (n = 4, in each group). Data are presented as mean ± SEM. **P < 0.01.
mice (Fig. 5a). In vitro, stimulation with LPS induced M1 polarization in macrophage Raw264.7 cells, as revealed by an increase in mRNA expressions of Tnf and Nos2 (Fig. 5b). In contrast, stimulation with IL-4 induced M2 polarization in Raw264.7 cells, as revealed by an increase in mRNA expressions of Mrc1 and Retnla (Fig. 5c). We found that irbesartan treatment significantly decreased C1qa mRNA expression in Raw264.7 cells, when they were either M1-polarized by LPS or M2-polarized by IL-4 (Fig. 5d,e). These results suggest the hierarchical relationship between AT1 receptor signaling and C1q/Wnt-β-catenin signaling.

Irbesartan suppressed the activation of C1q-Wnt/β-catenin signaling pathway after cryoinjury. Serum C1q concentration was elevated at 4 d after cryoinjury, but irbesartan treatment attenuated the increase in the serum C1q concentration in cryoinjured mice (Fig. 6a). Irbesartan treatment also inhibited the increase in mRNA expressions of C1qa and Axin2, which is a downstream molecule of the Wnt/β-catenin signaling pathway, in cryoinjured TA muscles at 4 d after cryoinjury, as revealed by real-time RT-PCR analysis (Fig. 6b). X-gal staining also revealed that the number of LacZ+ cells in histological sections of TA muscles in Axin2−/− mice at 4 d was significantly decreased by irbesartan treatment (Fig. 6c,d). These results suggest that AT1 receptor blockade suppresses activation of C1q-Wnt/β-catenin signaling pathway in injured skeletal muscle.

Topical administration of C1q reversed the beneficial effects of irbesartan on skeletal muscle repair after cryoinjury. Finally, we examined the effects of administration of C1q on irbesartan-induced enhancement of repair and regeneration in TA muscles after cryoinjury. Irbesartan treatment significantly decreased Axin2 mRNA expression in TA muscles at 2 d after cryoinjury, but topical administration of C1q in PuraMatrix hydrogel reversed the suppressive effect of irbesartan on Axin2 mRNA expression (Fig. 7a). Similarly, immunostaining revealed that the increase in the size of eMHC-positive myofibers and the decrease in Collagen 1+ fibrotic area in TA muscles of irbesartan-treated mice at 10 d after cryoinjury were reversed by topical administration of C1q (Fig. 7b–d). These results suggest that AT1 receptor blockade improves muscle repair and regeneration through down-regulation of the aging-promoting C1q.

Discussion
In the present study, we demonstrated that genetic blockade of AT1 receptor in mice led to a prolongation of chronological life span (Supplementary Fig. 1), as reported by Benigni A. et al.17. They described morphological differences in several organs between aged Agtr1a−/− mice and wild-type controls, such as lower cardiomyocyte size and collagen deposition in the heart, fewer atherosclerotic lesions in the aorta, and fewer lymphoid aggregates in the pancreas of Agtr1a−/− mice17. Our analysis revealed that aging-related decline in skeletal muscle function was remarkably milder in Agtr1a−/− mice (Supplementary Fig. 3). One of the most important aging-related features in our body is the locomotive decline, which has a great impact on individual health span and quality of life. Locomotive decline is primarily caused by skeletal muscle aging, which is characterized by the loss of muscle mass and strength23. The life-long maintenance of skeletal muscle is ensured by the continuous and balanced renewal of myofibers19. Therefore, we investigated the role of AT1 receptor signaling in skeletal muscle regeneration by applying cryoinjury on TA muscles. Our analysis revealed that irbesartan significantly enhanced regenerative capacity of skeletal muscle with better muscular differentiation and growth, and that it induced M2 polarization of macrophages and down-regulated aging-promoting C1q expression. According to an observational study, elderly women with hypertension continuously using Ang converting enzyme (ACE) inhibitors showed a lower decline in muscle strength than intermittent or never users of ACE inhibitors24. Cross-sectional analysis also demonstrated that the use of ACE inhibitors was associated with larger muscle mass of lower extremities in the elderly25. Our study may provide mechanistic insights into the beneficial effects of pharmacological inhibition of renin-angiotensin system in the prevention of age-related sarcopenia.

To explore the impact of AT1 receptor blockade on skeletal muscle regeneration, we examined AT1 receptor activation in cryoinjured mice by treatment with irbesartan, not by genetic disruption of Agtr1a gene, to eliminate possible metabolic effects of hyperphagia and obesity in Agtr1a−/− mice39. Irbesartan is one of the ARBs that are commercially available as highly effective and well-tolerated drugs for the management of hypertension26,27. Although peripheral administration of irbesartan was reported to have access to central nervous system28, we observed that irbesartan, orally administered at a subpressor dose, showed no significant effect on daily food intake and body weight (Supplementary Fig. 4). There has been conflicting evidence for AT1 receptor blockade on skeletal muscle regeneration29. Bedair HS, et al. and Burks TN. et al. reported that treatment with the ARB losartan in drinking water led to histological improvement in muscle regeneration after laceration-induced injury in mice30,31, but in contrast, Johnston APW. et al. reported that treatment with the angiotensin-converting enzyme inhibitor captopril or genetic disruption of Agtr1a gene led to significant impairment in muscle growth after cardioxotin-induced injury in mice32. Meanwhile, Murphy KT. et al. reported that Agtr1a−/− mice exhibited impaired muscle regeneration after notoxin-induced injury, while they had enhanced muscle strength, mobility, and locomotor activity at baseline33. Although the reasons for these discrepant results are currently unclear, irbesartan, in our hands, promoted repair and regeneration of skeletal muscle after cryoinjury (Figs 1 and 2). Besides selectively binding to the AT1 receptor and inhibiting Ang II-induced receptor activation, irbesartan
Figure 5. Decreased expression of C1q mRNA expression in macrophages by treatment with irbesartan both in vivo and in vitro. (a) The mRNA levels of C1qa in CD11b + and F4/80 + macrophages sorted from TA muscles of mice treated with irbesartan or vehicle at 6 d after cryoinjury (n = 6, in each group). Data are shown as fold induction over vehicle (mean ± SEM). *P < 0.05. (b) The mRNA levels of Tnf and Nos2 in Raw264.7 cells after stimulation with LPS or vehicle (n = 9, in each group). Data are shown as fold induction over vehicle (mean ± SEM). *P < 0.05. (c) The mRNA levels of Mrc1 and Retnla in Raw264.7 cells after stimulation with IL-4 or vehicle (n = 9, in each group). Data are shown as fold induction over vehicle (mean ± SEM). *P < 0.05. (d) The mRNA levels of C1qa in LPS-stimulated Raw264.7 cells after treatment with irbesartan or vehicle (n = 9, in each group). Data are shown as fold induction over vehicle (mean ± SEM). **P < 0.01. (e) The mRNA levels of C1qa in IL-4-stimulated Raw264.7 cells after treatment with irbesartan or vehicle (n = 9, in each group). Data are shown as fold induction over vehicle (mean ± SEM). **P < 0.01.
has been reported to exert pleiotrophic actions such as activation of peroxisome proliferator-activated receptor-γ and inhibition of nuclear factor-κB activity. Further studies will be required to elucidate whether the actions independent of the AT1 receptor inhibition make significant contributions to the beneficial effects of irbesartan on skeletal muscle repair and regeneration.

Regenerative capacity of skeletal muscle could be caused by perturbations to the processes that regulate activation, proliferation, and differentiation of satellite cells. In response to injury, neutrophils and M1 macrophages infiltrate the damaged muscles and release proinflammatory cytokines, leading to further damage of myofibers and activation and proliferation of satellite cells. Subsequently, M1 macrophages are replaced by a population of M2 macrophages that attenuate the inflammatory response and promote differentiation and growth of satellite cells. Therefore, myogenic regeneration is tightly linked to inflammatory microenvironments. We observed that treatment with irbesartan promoted repair and growth of skeletal muscle after cryoinjury (Fig. 1). Irbesartan also induced M2 polarization of macrophages in injured muscles (Fig. 4), which was consistent with the previous findings that treatment with ARB skewed intrarenal macrophages from M1 to M2 phenotype in a mouse model of obesity-related...
Figure 7. Effects of topical administration of C1q on irbesartan-induced enhancement of skeletal muscle repair after cryoinjury. (a) The mRNA levels of Axin2 in TA muscles of irbesartan- or vehicle-treated mice with topical administration of C1q or PBS in PuraMatrix hydrogel at 2 d after cryoinjury (vehicle + PuraMatrix-PBS, n = 12; vehicle + PuraMatrix-C1q, n = 7; irbesartan + PuraMatrix-PBS, n = 9; irbesartan + PuraMatrix-C1q, n = 7). Data are shown as fold induction over vehicle + PuraMatrix-PBS (mean ± SEM). *P < 0.05, **P < 0.01. NS, not significant. PuraM, PuraMatrix. (b) Immunostaining of TA muscles of irbesartan- or vehicle-treated mice with topical administration of C1q or PBS in PuraMatrix hydrogel at 10 d after cryoinjury. Embryonic myosin heavy chain (eMHC) and Collagen 1 are represented in red and green, respectively. Scale bar, 100 μm. (c) The sizes of eMHC+ fibers in TA muscles of irbesartan- or vehicle-treated mice with topical administration of C1q or PBS in PuraMatrix hydrogel at 10 d after cryoinjury (vehicle + PuraMatrix-PBS, n = 284; vehicle + PuraMatrix-C1q, n = 214; irbesartan + PuraMatrix-PBS, n = 338; irbesartan + PuraMatrix-C1q, n = 286). Data are presented as mean ± SEM. *P < 0.05, **P < 0.01. NS, not significant. (d) The percent area stained for Collagen 1 in TA muscles of irbesartan- or vehicle-treated mice with topical administration of C1q or PBS in PuraMatrix hydrogel at 10 d after cryoinjury (n = 7, in each group). *P < 0.05, **P < 0.01. NS, not significant.
kidney injury and a rat model of anti-glomerular basement membrane glomerulonephritis. Although little is known about the mechanisms how AT₁ receptor blockade induces M2 polarization of macrophages, the shift in macrophages from M1 to M2 phenotype might contribute to the beneficial effects of irbesartan on muscle growth and repair after cryoinjury.

Our analysis indicated that irbesartan promoted activation and proliferation of Pax7+ satellite cells following injury (Fig. 3). It has been reported that the proliferative potential declined with aging because of an increase in transforming growth factor-β (TGF-β) signaling and a decrease in Notch signaling pathway, and that AT₁ receptor blockade suppressed TGF-β signaling, and thereby attenuated TGF-β-mediated impairment of muscle regeneration in myopathic mice with Fibrillin 1 mutation or dystrophin deficiency. Since AT₁ receptor is expressed in satellite cells, AT₁ receptor blockade may also inhibit the direct anti-proliferative actions of Ang II on satellite cells. In the present study, we demonstrated that down-regulation of the aging-promoting C1q mediated beneficial effects of AT₁ receptor blockade on muscle repair and regeneration (Figs 5–7). We recently identified C1q as a diffusible factor in the serum that activates canonical Wnt/β-catenin signaling by binding to Wnt receptor Frizzled and by inducing cleavage of Wnt coreceptor LRP6. Serum and tissue concentrations of C1q showed a significant increase with aging, and C1q-mediated activation of Wnt/β-catenin signaling led to aging-related decline in regeneration capacity of skeletal muscles. C1q treatment attenuated satellite cell proliferation and stimulated fibroblast proliferation both in vitro and in vivo. We also reported that M2 macrophages were recruited to the aorta and secreted C1q, which promoted arterial remodeling through activation of Wnt/β-catenin signaling, in Ang II-infused mice. We propose that AT₁ receptor blockade reduces systemic and local levels of C1q through inhibiting C1q production by infiltrated macrophages, and thereby enhances activation and proliferation of satellite cells. Inasmuch as AT₁ receptor is expressed in both myocytes and macrophages, further studies will be required to elucidate the cell type-specific role of AT₁ receptor in inducing C1q expression in macrophages in injured muscles.

In conclusion, AT₁ receptor blockade prolongs life span, and improves muscle repair and regeneration through modulating the inflammatory microenvironment, including M2 polarization of macrophages and down-regulation of the aging-promoting C1q-Wnt/β-catenin signaling pathway. Elucidation of detailed relationship between AT₁ receptor signaling and C1q/Wnt/β-catenin signaling will be of interest, and will contribute to understanding the mechanisms of functional decline of multiple organs with aging and developing preventive and therapeutic measures against aging-related diseases.

Methods
Mice, and cryoinjury. All of the experiments were approved by the Institutional Animal Care and Use Committees of Osaka University and Chiba University, and carried out in accordance with the guidelines of Osaka University and Chiba University. C57BL/6J mice were purchased from CLEA Japan, Inc., and generation of Agtr1a-/- mice and Axin2+/2 mice has been described previously. The mice were fed a standard chow (CRF-1; Charles River Laboratories Japan, Inc.) or a chow containing irbesartan (Shionogi & Co., Ltd.). Treatment with irbesartan (20 mg/kg/day) was started one day prior to cryoinjury operation, and continued until cessation of the experiment. To produce cryoinjury, we anesthetized 7 to 10-week-old male mice by intraperitoneal injection of medetomidine hydrochloride (0.3 mg/kg), midazolam (4 mg/kg), and butorphanol (5 mg/kg), and anesthesia was monitored by pinching the toe. We cooled a metal probe (2 mm in diameter and 15 cm in length), and the wire was placed 30 cm above the floor. The mouse was used in conscious mice noninvasively by a programmable sphygmomanometer (BP-98A; Softron) using the tail-cuff method.

Blood pressure measurement. The systolic and diastolic blood pressures and pulse rates were measured in conscious mice noninvasively by a programmable sphygmomanometer (BP-98A; Softron) using the tail-cuff method.

Analysis of hair growth. Age-matched young (12 to 16-week-old) and old (72-week-old) mice were shaved on their dorsal surface (2.0 cm²) using an electric razor for hair growth assay, as described previously. We measured an area with hair growth 28 d after shaving.

Analysis of skeletal muscle function. A vertical pole test was used to test the motor balance of mice, as described previously. A mouse was placed onto a horizontally positioned wood pole (2 cm in diameter and 50 cm long), and the pole was raised gently and slowly to a 90° position. The length of time for the mouse to fall off the pole was recorded. A hanging wire test was used to test the muscle strength, tone, and equilibrium, as described previously. A mouse was placed onto a horizontally positioned wire (1.5 mm in diameter and 15 cm in length), and the wire was placed 30 cm above the floor. The mouse was required to grasp the middle part of the wire with their forepaws, and the length of time for the mouse to fall off the wire was recorded. A cutoff time of 1 min was used for the vertical pole and hanging wire tests. A treadmill test was used to assess running performance. Mice were subjected to a horizontal treadmill at 13 m/min after acclimatization to a rodent treadmill device (KN-73; Natsume Seisakusho, Co. Ltd.) by 5 min rest on the conveyor belt and subsequent 5 min running. Running was terminated...
when mice touched the electric shock grid at the back of the treadmill more than 10 times/min, and the total running distance was recorded.

**Cell culture.** Raw264.7 cells (American Type Culture Collection) were plated at a density of 1 × 10^5 cells/3.5 cm dish in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum. After 12 h of culture, Raw264.7 cells were stimulated with 0.5 ng/ml of LPS (Sigma-Aldrich) or 0.4 ng/ml of IL-4 (PeproTech, Inc.) for 24 h and then treated with 10^{-7} M of irbesartan or vehicle for 3 h.

**Histological analysis.** For histological analysis, TA muscles were excised, fixed in 10% neutralized formalin, and embedded in paraffin. Serial sections at 5 μm were deparaffinized and stained with hematoxylin and eosin for morphological analysis and Masson's trichrome for evaluation of fibrosis. Images were acquired with a microscope (FSX100; Olympus), and for measurement of the cross-sectional area of centronuclear myofibers, images of random fields of injured area were analyzed using WinROOF software (Mitani Corporation). We evaluated a ratio of fibrotic area to injured area for quantification of the fibrotic area (%) in images acquired with a microscope (BZ-X700; Keyence Corporation) using BZ-X analysis application (Keyence Corporation).

**Immunohistochemical analysis.** In vivo cryotechnique was used to freeze TA muscles promptly for immunohistochemical analysis. TA muscles of anesthetized mice were excised, and treated with DNase to remove contaminating genomic DNA using TURBO DNA-free Kit (Ambion, Inc.), and washed in acetone, immersed in 30% sucrose overnight, and frozen with isopentane-propane cryogen (−193 °C) precooled in liquid nitrogen. The frozen muscles were removed with an electric dental drill in liquid nitrogen, and processed for freeze-substitution fixation in acetone containing 0.2% glutaraldehyde at −80 °C for 24 h and then −30 °C and 4 °C for 2 h each. After being left at room temperature for 1 h, they were washed in acetone, immersed in 30% sucrose overnight, and frozen with isopentane precooled in dry ice for 1 h. The frozen sections (7 μm) were fixed in acetone, incubated with the following primary antibodies overnight at 4 °C: mouse monoclonal anti-eHIC antibody (clone F1.652, Developmental Studies Hybridoma Bank), rabbit polyclonal anti-Collagen I antibody (Abcam), mouse monoclonal anti-Pax7 antibody (clone PAX7, R&D Systems, Inc.), and rat monoclonal anti-Laminin 2 alpha antibody (clone 4H8-2, Abcam). Secondary antibodies (Alexa Fluoro 488-conjugated anti-rabbit IgG antibody, Alexa Fluoro 488-conjugated anti-rat IgG antibody, or Alexa Fluoro 594-conjugated anti-mouse IgG antibody (Life Technologies, Inc.)) and TO-PRO-3 (Life Technologies, Inc.) were applied to visualize expression of specific proteins and nuclei, respectively. Sections were mounted with ProLong Gold Antibody (Life Technologies, Inc.), and images were acquired with either a fluorescence microscope (BZ-X700; Keyence Corporation) or an LSM 700 confocal microscope (Carl Zeiss), and analyzed using WinROOF software (Mitani Corporation).

**X-gal staining.** X-gal stainings were performed, as described previously. TA muscles were excised from 10-week-old Axin2lacZ/+ mice 4 d after cryoinjury. Cryostat sections (10 μm) were stained with X-gal, and counterstained with nuclear fast red.

**Real time RT-PCR analysis.** Total RNA was extracted by using the TRIzol Reagent (Life Technologies, Inc.), and treated with DNase to remove contaminating genomic DNA using TURBO DNA-free Kit (Life Technologies, Inc.). Single-stranded cDNA was transcribed by using the SuperScript VILO cDNA Synthesis Kit (Life Technologies, Inc.) according to the manufacturer's protocol. We conducted quantitative real-time PCR analysis using Light Cycler TaqMan Master Kit (Roche Applied Science, Inc.) with the target-specific primers and the matching probes designed by the Universal ProbeLibrary System (Roche Applied Science, Inc.) applied to visualize expression of specific proteins and nuclei, respectively. Images were acquired with either a fluorescence microscope (BZ-X700; Keyence Corporation) or an LSM 700 confocal microscope (Carl Zeiss), and analyzed using WinROOF software (Mitani Corporation).

**Flow cytometric analysis.** TA muscles were minced and digested in high glucose Dulbecco’s modified Eagle’s medium containing 0.2% collagenase type 1 (Wako Pure Chemical Industries, Ltd.). After digested tissues were further dissociated with 18 G needle and remaining debris was sedimented, the supernatant was collected after filtering through 100- and 40-μm cell strainer (Corning, Inc.), and cells were suspended in PBS containing 3% fetal bovine serum. After Fc receptor blocking using rat anti-mouse CD16/CD32 monoclonal antibody (clone 2.4G2, BD Biosciences), cells were incubated with...
PE-conjugated anti-F4/80 antibody (Biolegend, Inc.) and PerCP-Cy5.5-conjugated anti-CD11b antibody (BD Biosciences) for 30 min on ice, and washed with PBS containing 3% fetal bovine serum. Dead cells were excluded using LIVE/DEAD Fixable Dead Cell Stain Kit (Life Technologies, Inc.). The percentages of CD11b+ and F4/80+ cells were analyzed by the BD FACSAria II (BD Biosciences) using BD FACSDiva software (BD Biosciences). CD11b+ and F4/80+ cells were sorted and collected using the BD FACSAria II (BD Biosciences) for real time RT-PCR analysis and further staining with RELMα. CD11b+ and F4/80+ cells were fixed and permeabilized in BD Cytofix/Cytoperm buffer (BD Biosciences) for 20 min on ice, and washed in BD Perm/Wash solution (BD Biosciences). Cells were incubated with biotinylated rabbit polyclonal anti-RELMα antibody (Abcam) for 30 min on ice, followed by APC-conjugated streptavidin (BD Biosciences) for 30 min on ice, and washed with PBS containing 3% fetal bovine serum. The percentages of CD11b+ and RELMα+ cells were analyzed by the BD FACSAria II (BD Biosciences) using BD FACSDiva software (BD Biosciences).

**ELISA for C1q.** Serum C1q concentration was determined by using C1q, Mouse, ELISA kit (Hycult Biotech), according to the manufacturer’s protocol.

**Statistical analysis.** All of the data are presented as mean ± SEM. Two-group comparison was analyzed by unpaired 2-tailed Student’s t test and Welch’s t test, and multiple-group comparison was performed by 1-way ANOVA followed by the Tukey-Kramer HSD test and Steel-Dwass test for comparison of means. We estimated survival curves by the Kaplan-Meier method, and compared the groups using the log-rank test. Values of P < 0.05 were considered statistically significant.

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How to cite this article

doi: 10.1038/srep14453 (2015).

Supplementary information

accompanies this paper at http://www.nature.com/srep

Acknowledgements

We thank S. Saitoh (University of Yamanashi) for technical advice, and M. Shimizu, H. Taniwaki, K. Kawaguchi, N. Miyagawa, and Y. Ueda for their excellent technical assistance. This work was supported in part by grants from Japan Society for the Promotion of Science (KAKENHI 23390213, 24659390, 26670395 to H.A., KAKENHI 21229010 and AMED-CREST, Japan Agency for Medical Research and Development to I.K.); Health and Labor Sciences Research Grants (to I.K.).

Author Contributions

H.A. and I.K. planned and designed the experiments. I.K. supervised the project. C.Y., R.Y., M.Y., Y.K.-S. and T.S. performed the experiments. T.K., H.Y., Y.Sh. and A.S.-K. analyzed the data. A.T.N., T.O., J.-K.L., J.S., Y.Sa. and E.U. advised on the experiments. C.Y., H.A. and I.K. wrote the manuscript.

Additional Information

Supplementary information accompanies this paper at http://www.nature.com/srep

Competing financial interests: H.A. has received trust research/joint research funding from Shionogi & Co., Ltd., and research funding from Takeda Pharmaceutical Co., Ltd., Daiichi Sankyo Co., Ltd., and Nippon Boehringer Ingelheim Co., Ltd. I.K. has received research funding from Astellas Pharma Inc., Daiichi Sankyo Co., Ltd., Nippon Boehringer Ingelheim Co., Ltd., and Takeda Pharmaceutical Co., Ltd., and has affiliations with endowed department sponsored by Shionogi & Co., Ltd.

How to cite this article: Y abumoto, C. et al. Angiotensin II receptor blockade promotes repair of skeletal muscle through down-regulation of aging-promoting C1q expression. Sci. Rep. 5, 14453; doi: 10.1038/srep14453 (2015).
