Glucagon-Like Peptide 1 Inhibits the Sirtuin Deacetylase SirT1 to Stimulate Pancreatic β-Cell Mass Expansion

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OBJECTIVE—The glucocincretin hormone glucagon-like peptide 1 (GLP-1) enhances glucose-stimulated insulin secretion and stimulates pancreatic β-cell mass expansion. We have previously shown that the forkhead transcription factor FoxO1 is a prominent transcriptional effector of GLP-1 signaling in the β-cell. FoxO1 activity is subject to a complex regulation by Akt-dependent phosphorylation and SirT1-mediated deacetylation. In this study, we aimed at investigating the potential role of SirT1 in GLP-1 action.

RESEARCH DESIGN AND METHODS—FoxO1 acetylation levels and binding to SirT1 were studied by Western immunoblot analysis in INS832/13 cells. SirT1 activity was evaluated using an in vitro deacetylation assay and correlated with the NAD+-to-NADH ratio. The implication of SirT1 in GLP-1–induced proliferation was investigated by BrdU incorporation assay. Furthermore, we determined β-cell replication and mass in wild-type and transgenic mice with SirT1 gain of function assay. Furthermore, we determined β-cell replication and mass in wild-type and transgenic mice with SirT1 gain of function assay.

RESULTS—Our data show that GLP-1 increases FoxO1 acetylation, decreases the binding of SirT1 to FoxO1, and stunts SirT1 activity in β-INS832/13 cells. GLP-1 decreases both the NAD+-to-NADH ratio and SirT1 expression in INS cells and isolated islets, thereby providing possible mechanisms by which GLP-1 could modulate SirT1 activity. Finally, the action of GLP-1 on β-cell mass expansion is abolished in both transgenic mice and cultured β-cells with increased dosage of SirT1.

CONCLUSION—Our study shows for the first time that the glucocincretin hormone GLP-1 modulates SirT1 activity and FoxO1 acetylation in β-cells. We also identify SirT1 as a negative regulator of β-cell proliferation.

The glucocincretin hormone glucagon-like peptide 1-[7–36]amide (GLP-1) (1–3) is a potent therapeautic agent in the treatment of diabetes (4). GLP-1 improves insulin secretion in subjects with impaired glucose tolerance and type 2 diabetes (5). It also stimulates insulin gene expression and insulin biosynthesis (6), in part via increased expression and activity of the β-cell–specific transcription factor Pdx1 (7,8). Moreover, GLP-1 has been shown to promote β-cell mass expansion in both experimental animal models (8,9) and cultured β-cells (7,10–14). However, the molecular mechanism by which GLP-1 exerts its action is not fully elucidated.

We have previously shown that GLP-1 transactivates the epidermal growth factor receptor (12) to subsequently activate phosphatidylinositol-3 kinase and Akt signaling (7,11). Activation of epidermal growth factor receptor/phosphatidylinositol-3 kinase/Akt signaling by GLP-1 stimulates β-cell proliferation (7,11) and survival (13,14). Of interest, this signaling pathway has been suggested to play a role in the glucocincretin effect of GLP-1 as well (15). We have also demonstrated that the forkhead transcription factor FoxO1, an important regulator of β-cell mass (16–18), is a prominent transcriptional effector of GLP-1 action in β-cells (10). Thus, GLP-1 inhibits FoxO1 via Akt-mediated phosphorylation and nuclear exclusion. Inhibition of FoxO1 by GLP-1 increases both Pdx1 and Foxa2 expression and triggers β-cell mass expansion (10).

FoxO1 activity is regulated in a complex fashion by various posttranslational modifications, including reversible Ser-Thr phosphorylation and Lys acetylation (19). Acetylation at Lys-242, -245, and -262 of FoxO1 attenuates its ability to bind cognate DNA sequence and increases its susceptibility to phosphorylation by Akt (20). Conversely, deacetylation of FoxO1 by the NAD+-dependent protein deacetylase SirT1 increases its transcriptional activity (21–23).

We therefore sought to test the possible implication of SirT1 in GLP-1 action. The current study shows that GLP-1 stunts SirT1-mediated FoxO1 deacetylation, thereby relieving a molecular brake on β-cell mass expansion. Our work describes a novel mechanism for GLP-1 action. It also identifies SirT1 as a negative regulator of β-cell proliferation.

RESEARCH DESIGN AND METHODS

Reagents. Human GLP-1 fragment 7–36 amide, exendin-4, nicotinamide, and resveratrol were obtained from Sigma (St. Louis, MO). RPMI-1640 medium, FCS, and other culture media were purchased from Invitrogen (Burlington, ON, Canada). Anti-FK228 antibody was purchased from Cell Signaling (Beverly, MA). Anti-actin and anti-SirT1 antibodies were obtained from Millipore (Bedford, MA). Anti-guinea-pig insulin was purchased from Sigma.

Cell culture. INS832/13 cells (24) were grown in RPMI-1640 medium supplemented with 10 mmol/L HEPES, 10% heat-inactivated FCS, 2 mmol/L L-glutamine, 1 mmol/L sodium pyruvate, 50 μmol/L β-mercaptoethanol, 100 μg/mL penicillin, and 100 μg/mL streptomycin at 37°C in a humidified 5% CO2 atmosphere. Cells at 70% confluence were washed with phosphate-buffered saline and preincubated in serum-free RPMI-1640 medium supplemented with 3 mmol/L glucose and 0.1% BSA (Sigma) for at least 4 h before treatment. This condition mimics calorie restriction and was shown to activate SirT1.

Islet isolation. Rat islets were isolated from male Wistar rats (250 g) by collagenase digestion. Islets were subsequently purified over a Histopaque gradient and handpicked under a microscope.

Human islets were isolated from organ donors at the Department of Surgery, Montreal General Hospital, McGill University Health Center, Montreal, Quebec, Canada (three separate donors were received). Human ethics approval was obtained through the McGill University Health Center ethics committee. Donors were between ages 42 and 65, and none had a history of diabetes or metabolic disorder. Islets were isolated by digestion with Liberase CI (Boehringer Mannheim, Indianapolis, IN) followed by purification over a Ficoll gradient using a CORE-2901 Cell Processor (COBE BCT, Denver, CO). After isolation, islets were rested overnight in complete CMRL medium.
indicated periods of time.

**Western blot.** Proteins were extracted and quantified by BCA assay (Roche, Indianapolis, IN). FoxO1 protein complexes were immunoprecipitated from 1 mg of total protein extracts and subjected to electrophoresis on 8 or 10% polyacrylamide gels.

**Chromatin immunoprecipitation.** Cells were fixed to isolate intact chromatin and sheared in 200-base pair fragments, and SirT1 DNA/protein complexes were immunoprecipitated using an anti-SirT1 antibody (Millipore). Bound DNA was amplified by PCR using sequence-specific primers designed for amplification of the forkhead binding site region of the Foxa2 promoter (16).

**SirT1 activity.** Cells were treated for 15 min, and SirT1 activation was measured using a commercially available fluorometric assay (Biomol, Plymouth Meeting, PA). In brief, the kit uses a SirT1 substrate consisting of amino acids 379–382 of human p53 (Arg-His-Lys-Lys[Ac]). The assay’s fluorescence signal is generated in proportion to the amount of deacetylation of Lys-382, a known in vivo target of SirT1. Cell extracts were incubated for 10 min at 37°C with a 25 μmol/L concentration of the fluorogenic SirT1 substrate. Reactions were stopped by the addition of 1 mmol/L nicotinamide, and the fluorescence resonance energy transfer signal was determined.

**NAD^+/-NADH ratio and cellular ATP concentrations.** NAD^+, NADH, and ATP levels were measured in cell extracts using commercial kits (NAD/NADH quantification kit and ATP fluorometric kit from Biovision, Mountain View, CA).

**Nicotinamide phosphoribosyltransferase activity.** Nicotinamide phosphoribosyltransferase (Nampt) activity was measured by the conversion of ^14C-labeled nicotinamide to ^14C]nicotinamide mononucleotide as described previously (25). In brief, cell lysates (10 μg total protein) were incubated with ATP, 5-phosphoribosyl-1-pyrophosphate, and ^14C]nicotinamide for 30 min; filtered through glass fiber filters; and washed with acetone. Filters were air-dried and transferred into scintillation vials for quantification.

**Transfection and cell proliferation.** Cells were transfected with either control green fluorescent protein (GFP), wild-type SirT1, or a catalytically inactive SirT1 mutant (26) using Lipofectamine 2000 (Invitrogen) as described previously (16). Transfection efficiency was 70%. The following day, proliferation of transfected cells was evaluated using BrdU incorporation assay (Roche). In brief, cells were treated for 24 h in the absence or presence of 1 nmol/L GLP-1, and BrdU was added to the culture medium for the last 60 min of the incubation period.

**Animal studies.** Mice were maintained on a mixed background (129sv and C57BL/6). Wild-type and SirBACO transgenic mice (27) aged 6–12 months were treated with daily intraperitoneal injections of exendin-4 (10 nmol/kg) or saline for 7 days (n = 5 each). Mice were killed, and pancreas sections were processed for insulin and Ki67 immunohistochemistry to assess β-cell proliferation. Cross-sectional islet area was measured using Image Pro Plus software (Media Cybernetics, Silver Spring, MD). Results are presented as relative β-cell areas (i.e., percent of insulin + areas relative to total pancreas areas). For each animal, at least four sections spaced 50 μm apart were studied.

**Calculations and statistics.** Data are presented as means ± SEM. Statistical analyses were performed with SPSS (SPSS Inc., Chicago, IL) using ANOVA.

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**RESULTS**

GLP-1 increases FoxO1 acetylation via inhibition of the sirtuin deacetylase SirT1. We have previously shown that inhibition of the forkhead transcription factor FoxO1 mediates the pleiotropic action of the glucoincretin hormone GLP-1 (10). FoxO1 is subject to various posttranslational modifications, including Akt-dependent phosphorylation and SirT1-mediated deacetylation. Here, we sought to test the hypothesis that GLP-1 inhibits SirT1 to enhance FoxO1 acetylation and to stimulate β-cell mass expansion.

We first evaluated FoxO1 acetylation and binding to SirT1 by Western blot in INS832/13 cells. We resorted to the use of a pancreatic β-cell line because of the high amount of biological material required to perform the assays. INS cells were incubated in serum-free medium, a condition known to activate SirT1, in the presence or absence of GLP-1 and the SirT1 activators H_2O_2 or resveratrol. Figure 1A and B shows that GLP-1 induced FoxO1 acetylation was PCR-amplified using oligonucleotides flanking the forkhead binding site in the Foxa2 promoter. Western blots showing the subcellular localization of SirT1 and FoxO1 after 45-min incubation with either 10 nmol/L GLP-1 (GLP) or 10 μmol/L resveratrol. Actin is shown as control. Representative images of at least three experiments are shown. Res, resveratrol; Ack, acetyl-lysine, Ctl, control.
acetylation, an action prevented by resveratrol but not by 
H₂O₂. Coimmunoprecipitation experiments revealed that 
GLP-1 inhibited the physical association between FoxO1 
and SirT1 (Fig. 1A). However, this action was inhibited 
by both H₂O₂ and resveratrol. The action of GLP-1 on FoxO1 
acetylation was transient, with a maximal effect observed 
between 15 and 45 min. FoxO1 acetylation returned to 
basal value after ~180 min (Fig. 1C). Taken together, these 
results suggest that GLP-1 inhibits SirT1-mediated FoxO1 
deacetylation.

Chromatin immunoprecipitation assay was performed to 
assess whether SirT1 is recruited to the promoter region of 
FoxO1 target genes. Figure 1D shows that SirT1 was 
physically associated to a region of the Foxa2 promoter 
comprising the forkhead binding site (10). Moreover, 
SirT1 binding to the Foxa2 promoter was reduced upon 
GLP-1 treatment. This result indicates that SirT1 binding to 
the Foxa2 promoter mirrors that of FoxO1, as we have previously demonstrated (10). Altogether, these findings 
demonstrate that FoxO1 and SirT1 are recruited to the 
promoter of FoxO1 target genes in a GLP-1–inhibitable 
manner. These data suggest a role for SirT1 in the tran-
scriptional response to the glucoincretin hormone in 
β-cells. We next performed subcellular fractionation to 
address whether GLP-1 provokes SirT1 nuclear exclusion, 
as it does for FoxO1 (Fig. 1E). Our results show that SirT1 
is predominantly nuclear in INS cells and that its 
subcellular localization was not altered by GLP-1 treatment. 
Conversely, GLP-1 induced cytoplasmic translocation of 
FoxO1, as we have published previously (10).

We next sought to directly test whether GLP-1 affects 
SirT1 activity using an in vitro deacetylase assay. Figure 2A 
shows that GLP-1 blunted SirT1 activity to the same extent 
as the SirT1 inhibitor nicotinamide. Conversely, SirT1 ac-
activity was increased in resveratrol-treated cells. Because 
SirT1 activity is dependent on NAD⁺, we measured changes 
in the NAD⁺-to-NADH ratio in response to GLP-1 treatment 
in INS cells and isolated human islets. Figure 2B shows 
that GLP-1 decreased the NAD⁺-to-NADH ratio by 30% in INS 
cells, an action mimicked by the addition of 10% serum. 
Conversely, resveratrol increased the NAD⁺-to-NADH ratio 
by 60%. Both GLP-1 and serum also elicited a reduction in 
the NAD⁺-to-NADH ratio in human islets (Fig. 2C), whereas 
the effect of resveratrol (~25%) did not reach statistical 
significance. The reduction in NAD⁺ availability provides 
a possible mechanism by which GLP-1 could acutely 
regulate SirT1 activity. To directly test this possibility, we 
supplemented the assay buffer with 450 μmol/L NAD⁺ 
(Fig. 2D). Supplementation of NAD⁺ suppressed GLP-1 as 
well as serum inhibition of SirT1 activity. In an effort to 
unravel the mechanism by which GLP-1 could alter the 
NAD⁺-to-NADH ratio, we investigated its action on the 
activity of Nampt, the rate-limiting enzyme for the bio-
synthesis of NAD, and on the cellular ATP content. GLP-1 
decreases the activity of Nampt (Fig. 2E), thereby revealing 
a novel mechanism by which GLP-1 could decrease the 
NAD⁺-to-NADH ratio. FK-866 (10 nmol/L), a specific Nampt 
pharmacological inhibitor, was used as a positive control. 
Conversely, GLP-1 did not significantly change the cellular 
ATP content (Fig. 2F), consistent with a recent publication 
by Peyot et al. (28). It is interesting that the addition of se-
rum did not alter Nampt activity but significantly increased 
the cellular ATP concentration. This suggests that serum and 
GLP-1 could inhibit SirT1 via distinct mechanisms.

We next measured SirT1 expression by quantitative PCR 
after overnight incubation in serum-free medium in the 
absence or presence of GLP-1 (Fig. 3A). Our data show 
that glucose and serum deprivation, a condition mimicking 
calorie restriction, increased SirT1 expression by fivefold 
compared with complete growth medium in INS cells. 
GLP-1 treatment completely abolished the rise in SirT1 
expression. We next sought to confirm these observations 
in isolated rat islets (Fig. 3B). Both GLP-1 and serum de-
creased SirT1 expression in rat islets as well, although the 
detected changes were more modest. Thus, SirT1 expres-
sion was repressed by 30% in the presence of 10 mmol/L GLP-1 
and by 40% in response to serum. We assessed SirT1 protein levels 
by Western blot to test whether alterations in SirT1 expres-
sion were translated into changes in protein levels. 
Figure 3C and D shows that GLP-1 decreased SirT1 protein 
levels by 70% in INS cells, an effect mimicked by 
serum.

**FIG. 2.** GLP-1 inhibits SirT1 deacetylase activity. A: Cells were 
incubated in the presence or absence of 10 mmol/L GLP-1, 10 mmol/L 
nicotinamide (NAM), or 10 μmol/L resveratrol (Res). Proteins were 
extracted after 15 min to perform the in vitro SirT1 deacetylase assay 
using a fluorogenic SirT1 substrate. B and C: NAD⁺ and NADH levels 
were determined in extracts from cells (B) or human islets (C) 
in the presence or absence of 10 mmol/L GLP-1, 10% serum, or 
10 μmol/L resveratrol for 15 min. D: SirT1 activity was measured as 
described in A with the modification that the assay buffer was supple-
mented with 25 μmol/L NAD⁺. E: Nampt activity was measured by the 
conversion of 14C-labeled nicotinamide to [14C]nicotinamide mono-
nucleotide. The specific Nampt pharmacological inhibitor FK-866 (10 
μmol/L) was used as control. F: Total cellular ATP was measured in 
cells incubated as described above. Means ± SE of three experiments, 
each comprising duplicates. *P < 0.05.

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Overexpression of SirT1 abolishes GLP-1 action on β-cell mass expansion. To test whether SirT1 inactivation mediates GLP-1 action on β-cell proliferation, we sought to perform SirT1 gain of function both in vitro and in vivo. INS832/13 cells were either transfected with wild-type SirT1, a catalytically inactive SirT1 mutant, or with control GFP, and β-cell replication was evaluated by BrdU incorporation. Transfection of the cells with SirT1 led to a fivefold increase in protein levels as compared with GFP (Fig. 4A). Overexpression of wild-type SirT1 inhibited GLP-1–induced β-cell proliferation but failed to significantly affect the effect of serum (Fig. 4B). Overexpression of a catalytically inactive SirT1 mutant did not significantly alter β-cell proliferation, indicating that the effects of SirT1 on β-cell proliferation are dependent on its deacetylase activity.

To demonstrate the implication of SirT1 in GLP-1 action in vivo, we determined whether the effects of GLP-1 on β-cell mass were curtailed in transgenic mice with increased dosage of SirT1 (SirBACO mice) as previously described (27). Thus, mice were administered with either saline or the long-acting GLP-1 analog exendin-4 (10 nmol/kg) daily for 7 days. Cross-sectional β-cell areas were estimated by morphometry of pancreatic sections stained for insulin (Fig. 5A), and β-cell proliferation was evaluated by Ki67 staining (Fig. 5B). Exendin-4 administration increased the relative β-cell area by ~2.5-fold and stimulated β-cell replication by ~50% in wild-type animals. However, GLP-1 effects on both β-cell area and β-cell replication were abolished in SirBACO mice. Taken together, our results indicate that SirT1 gain of function prevents the mitogenic action of GLP-1 and identify SirT1 as a negative regulator of β-cell mass expansion.

DISCUSSION
Restoration and maintenance of β-cell mass is a major goal of diabetes treatment (29–31). GLP-1, a potent antidiabetes medication (4), enhances glucose-stimulated insulin secretion and promotes β-cell mass expansion, at least in rodents. However, the molecular mechanism by which GLP-1 exerts its action on β-cell mass is not fully elucidated. We have previously demonstrated that the transcription factor FoxO1, an important regulator of β-cell mass (16–18), is a prominent transcriptional effector of GLP-1 action (10). Thus, GLP-1 provoked nuclear exclusion of FoxO1 to relieve a molecular brake on Pdx1 and Foxa2 expression. Also, the effect of exendin-4 on β-cell mass expansion was abolished in transgenic mice with β-cell–specific FoxO1 gain of function (10). Acetylation/deacetylation of FoxO1 regulates its transcriptional activity (19). On one hand, acetylation of FoxO1 by CBP/p300 reduces its DNA binding activity (19), and deacetylation by SirT1 and Sirtuin 2 increases its DNA binding activity (28). On the other hand, GLP-1 increased deacetylation of FoxO1 by Sirtuin 1 (28). Therefore, GLP-1–induced deacetylation of FoxO1 may be important for the action of GLP-1 on β-cell mass.
activity and increases its susceptibility to phosphorylation by Akt (20). On the other hand, acetylated FoxO1 binds to the promyelocytic leukemia–associated protein Pml and is targeted to specific nuclear subdomains (Pml nuclear bodies) (32). Upon calorie restriction or oxidative stress, FoxO1 is deacetylated by the NAD+-dependent sirtuin deacetylase SirT1, which renders FoxO1 immobile within the nuclear compartment and results in FoxO1 transcriptional activity (20). This prompted us to investigate the role of SirT1 in GLP-1 action.

Our findings show that GLP-1 increases FoxO1 acetylation via inhibition of the NAD+-dependent SirT1 deacetylase. Our study provides two possible mechanisms by which GLP-1 could regulate SirT1. First, we show that GLP-1 acutely decreases the cellular NAD+-to-NADH ratio, possibly via its hitherto uncharacterized action on Nampt. Second, GLP-1 downregulates SirT1 expression, thereby providing a long-term mechanism by which chronic or prolonged exposure to GLP-1 receptor agonists could also inhibit SirT1. The fact that GLP-1 failed to increase the intracellular ATP content seemingly refutes the hypothesis that GLP-1 could inhibit SirT1 via the stimulation of β-cell glucose metabolism. However, because our assay did not take into account compartmentalization of ATP production or breakdown, this possibility cannot be completely ruled out. We also demonstrate that the effects of GLP-1 and its long-acting analog exendin-4 on β-cell mass were blunted in a pancreatic β-cell line with increased dosage or SirT1 as well as in transgenic mice with SirT1 gain of function. Our results are consistent with a model in which GLP-1 inhibits SirT1-mediated FoxO1 deacetylation to relieve the constraint on β-cell proliferation and β-cell mass expansion (Fig. 6). Thus far, the action of GLP-1 agonists on β-cell mass has not been demonstrated in humans in vivo. Moreover, the reported increase in β-cell mass in mice has been shown to be restricted after the initial burst of proliferation in the postnatal period (33,34).

However, a recent publication suggests that exendin-4 can stimulate β-cell replication in human islet grafts (35).

Previous studies using genetically engineered mice with SirT1 gain (36) or loss of function (37) indicate that SirT1 could enhance insulin secretion, an effect mediated via reduction of Ucp2 expression. These findings garnered much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already much attention by identifying SirT1, a molecule already.
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