Signal Transduction: 
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Characterization of the Raptor/4E-BP1 Interaction by Chemical Cross-linking Coupled with Mass Spectrometry Analysis*

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Background: mTORC1 recruits its substrate 4E-BP1 via Raptor/4E-BP1 interaction. Chemical cross-linking/mass spectrometry permits characterization of protein-protein interactions.

Results: Cross-linked peptides between Raptor and 4E-BP1 were identified. Raptor intramolecular cross-links were also identified.

Conclusion: Raptor N-terminal region containing RNC1 is implicated in the interaction with the central region of 4E-BP1.

Significance: Our study provides novel insight into how mTORC1 recognizes 4E-BP1.

mTORC1 plays critical roles in the regulation of protein synthesis, growth, and proliferation in response to nutrients, growth factors, and energy conditions. One of the substrates of mTORC1 is 4E-BP1, whose phosphorylation by mTORC1 reverses its inhibitory action on eIF4E, resulting in the promotion of protein synthesis. Raptor in mTOR complex 1 is believed to recruit 4E-BP1, facilitating phosphorylation of 4E-BP1 by the kinase mTOR. We applied chemical cross-linking coupled with mass spectrometry analysis to gain insight into interactions between mTORC1 and 4E-BP1. Using the cross-linking reagent bis[sulfosuccinimidyl] suberate, we showed that Raptor can be cross-linked with 4E-BP1. Mass spectrometric analysis of cross-linked Raptor-4E-BP1 led to the identification of several cross-linked peptide pairs. Compilation of these peptides revealed that the most N-terminal Raptor N-terminal conserved domain (in particular residues from 89 to 180) of Raptor is the major site of interaction with 4E-BP1. On 4E-BP1, we found that cross-links with Raptor were clustered in the central region (amino acid residues 56–72) we call RCR (Raptor cross-linking region). Intramolecular cross-links of Raptor suggest the presence of two structured regions of Raptor: one in the N-terminal region and the other in the C-terminal region. In support of this idea that the Raptor N-terminal conserved domain and the 4E-BP1 central region are closely located, we found that peptides that encompass the RCR of 4E-BP1 inhibit cross-linking and interaction of 4E-BP1 with Raptor. Furthermore, mutations of residues in the RCR decrease the ability of 4E-BP1 to serve as a substrate for mTORC1 in vitro and in vivo.

The mammalian target of rapamycin (mTOR) signaling pathway has attracted attention because of its involvement in a variety of cellular processes in response to stimuli such as growth factors, nutrients, and energy conditions (1–4). Up-regulation of this signaling pathway is frequently found in a variety of human cancers (5–7). Single amino acid changes can confer constitutive activation of mTOR, and we have identified activating mutations R2505P and S2215Y by mining the human cancer genome database (8). mTOR is a protein kinase belonging to the PI3K-related kinase family that includes PI3K-, ATM-, ATR-(ATM- and Rad3-related), and DNA-dependent protein kinase (9). Protein kinases in this group share similar structural features including the presence of the HEAT repeat, FAT, FATC, and the kinase domains. mTOR forms two different multiprotein complexes, mTORC1 and mTORC2 (10, 11); mTORC1 consists mainly of mTOR, Raptor, and mLST8 (G3P), whereas mTORC2 consists of mTOR, Rictor, mLST8, and Sin1. Substrates of mTORC1 include 4E-BP1 and S6K, whose phosphorylation causes promotion of protein synthesis. In the case of 4E-BP1, its phosphorylation results in relieving inhibitory action of this protein on eIF4E, a cap-binding protein (1–4, 12). S6K stimulates protein synthesis by phosphorylating ribosomal protein S6. In cancer cells, it is believed that 4E-BP1 has an important role in growth and proliferation (13, 14).

Phosphorylation of 4E-BP1 by mTOR is assisted by Raptor, which functions as a scaffold protein. Raptor recruits substrates

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proteins to mTORC1 so that they can be phosphorylated by mTOR (15, 16). Raptor is a protein of 150 kDa that contains the Raptor N-terminal conserved (RNC) motif, the HEAT repeat, and the WD40 motif (15, 16). Mutations in the RNC motif decrease interaction with 4E-BP1 (15, 16). We wanted to further investigate how mTORC1 interacts with 4E-BP1. Specifically, we sought to gain insight into the region of proximity between Raptor and 4E-BP1.

The recognition of substrate proteins by Raptor is believed to involve a five-amino acid sequence called the TOR signaling (TOS) motif, because mutations of this motif result in decreased interaction with Raptor (17–19). Another motif called “RAIP” is also proposed to be involved (17–19). Although these motifs are important, other regions of 4E-BP1 are also implicated in the interaction with Raptor (19).

Chemical cross-linking coupled with mass spectrometry analysis (CXMS) has emerged as a powerful method to characterize protein–protein interactions (20–26). A basic principle of the method is to use chemical cross-linkers to link reactive amino acid residues that are closely located. After protease digestion, cross-linked peptide pairs are separated by liquid chromatography and can be identified using mass spectrometry. Because only the residues located in close proximity can be cross-linked, intramolecular cross-links can provide insights into the folding of a protein, and intermolecular cross-links can reveal the site(s) of interaction between two proteins. As such, CXMS has been applied to gain information about subunit interactions within multiprotein complexes, which is particularly advantageous for protein complexes that are difficult to purify and crystallize. One of the challenges concerning CXMS is to develop a software tool for identification of cross-linked peptides from their fragmentation spectra (21, 25). The pLink software (21) has been optimized using a large data set and shown to be effective on a variety of samples including protein complexes.

In this paper, we report analysis of Raptor-4E-BP1 interaction using CXMS. We show that Raptor cross-links with 4E-BP1 and that several cross-linked peptides can be identified. Our results show that the cross-links mainly involve the RNC1 domain of Raptor and the central region of 4E-BP1, suggesting their close proximity. Identification of intramolecular cross-links raises the possibility that Raptor contains two structured domains. Finally, we have been able to block the formation of Raptor-4E-BP1 cross-links by synthetic peptides corresponding to the sequence within the RCR of 4E-BP1. We also show that mutations of 4E-BP1 residues within the RCR decrease the ability of 4E-BP1 to serve as a substrate of mTORC1 in vitro and in vivo.

**EXPERIMENTAL PROCEDURES**

**Reagents**—Cross-linkers, bis[sulfo succinimidyl] suberate (BS3) and diisuccinimidyl suberate were obtained from Pierce. Anti-FLAG M2 affinity gel and 3× FLAG peptide were obtained from Sigma. Full-length recombinant 4E-BP1 protein was obtained from Santa Cruz Biotechnology (Dallas, TX). Phosphorylated 4E-BP1 purified from Sf9 cells was obtained from CalBiochem (EMD Millipore, Billerica, MA). Anti-mTOR, anti-mLST8, and anti-4E-BP1 antibodies were obtained from Cell Signaling (Danvers, MA). Anti-Raptor antibody was obtained from BETHYL Lab (Montgomery, TX). Peptides were synthesized by Peptibody Inc. (Charlotte, NC).

**Purification of mTORC1 and Raptor**—Cells stably expressing FLAG-Raptor protein were established using the pLJM1 FLAG-Raptor plasmid (Addgene no. 26633). The lentivirus containing this plasmid was generated by the UCLA Vector Core Facility. HEK293T cells were infected with the lentivirus, and cells stably expressing FLAG-Raptor were obtained by puromycin selection. The resulting cell line, FLAG-Raptor 2, was cultured using Dulbecco’s modified Eagle’s medium enriched with 10% fetal bovine serum and 1× penicillin/streptomycin at 37 °C and 5% CO2. Cells expressing FLAG-Raptor were grown, washed with PBS, and stored at −80 °C for subsequent experiments. Cells (3.3×107) were lysed in 1 ml of 50 mM HEPES, pH 7.4, 150 mM NaCl, 0.4% CHAPS, 1× Complete, EDTA-free protease inhibitor mixture (Roche Applied Science), and 1 mM Na3VO4. In experiments where Raptor was isolated without the other components of mTORC1, 1% Triton X-100 was substituted for 0.4% CHAPS in the lysis buffer. The cell suspension was cleared of insoluble debris by centrifugation at 16,000×g for 10 min before being mixed with anti-FLAG M2 affinity gel (Sigma) for affinity purification. The affinity gel was collected, washed twice with wash buffer including ATP (50 mM HEPES, pH 7.4, 150 mM NaCl, 2 mM DTT, 2 mM ATP, 0.1% CHAPS), and washed three times with wash buffer without ATP (50 mM HEPES, pH 7.4, 150 mM NaCl, 2 mM DTT, 0.1% CHAPS). Bound proteins were eluted using 62 µg/ml 3× FLAG peptide (Sigma) in 50 mM HEPES, pH 7.4, 150 mM NaCl, 0.4% CHAPS. Eluted proteins were concentrated using Amicon Ultra 0.5-ml centrifugal filters NMWL 100K (EMD Millipore, Billerica, MA). If necessary, we subjected the preparation to another round of affinity purification using anti AU1 beads. In this case, we transfected HEK293T cells stably expressing FLAG-Raptor with AU1-mTOR DNA (8) before collecting cells. The presence of mTOR, Raptor, and mLST8 in the final preparation was evaluated by staining and Western blotting.

**mTORC1 Kinase Assay**—Protein kinase activity of mTORC1 was examined by carrying out kinase reaction using unphosphorylated 4E-BP1 (Santa Cruz Biotechnology, Dallas, TX) as described previously (27). Briefly, mTORC1 purified above was incubated with 0.25 µg of 4E-BP1 protein in kinase buffer (20 mM Tris-HCl, pH 7.5, 10 mM MgCl2, 0.2 mM ATP) for 20 min at 37 °C. The product was analyzed by Western blot using antibodies against phospho-4E-BP1 (Thr37/46), phospho-4E-BP1 (Ser46), or total 4E-BP1.

**4E-BP1 Binding Assay**—Binding of 4E-BP1 to mTORC1 was examined by the pulldown of 4E-BP1 with mTORC1 bound to beads. mTORC1 was purified from HEK293T cells expressing FLAG-Raptor using anti-FLAG beads. The beads were mixed with 4E-BP1 in a buffer containing 20 mM Tris-HCl, pH 7.5, 10 mM MgCl2, 100 mM NaCl. After the incubation, the beads were collected, washed, and then suspended in Laemml buffer. The sample was run on a SDS-polyacrylamide gel and probed with an antibody against 4E-BP1 as described (27).

**Cross-linking Procedure**—Purified proteins were combined with 4E-BP1 (Santa Cruz Biotechnology, Dallas, TX) and cross-
linked using BS³ or disuccinimidyl suberate. The reaction condition was the same as that used for the kinase reaction (27), except that ATP and MgCl₂ were omitted and Tris was replaced with HEPES. The water-soluble characteristics of BS³ made this cross-linker more compatible with the experimental work flow than its water-insoluble counterpart disuccinimidyl suberate. The cross-linking reagent was reacted with the proteins at room temperature for 20 min. Then the solution was adjusted to 20 mM NH₄HCO₃ and kept at room temperature for 20 min to inactivate the remaining free cross-linker reagent. The cross-linked proteins were stored at −80 °C in Laemmli buffer. Cross-linked proteins were separated by SDS-PAGE and stained with GelCode Blue (Pierce) according to the manufacturer’s instructions. Protein bands corresponding to areas of interest identified by Western blot were excised and subjected to mass spectrometry analyses. For peptide inhibition experiments, peptides with the amount indicated in the legend to Fig. 6 were added to cross-linking reactions.

**Mass Spectrometry Analysis and Database Search** —In-gel digestions were carried out as described before (28) with slight modifications. Briefly, gel slices of cross-linked protein bands were destained and replaced with 200 μl of 10 mM DTT at 56 °C for 40 min. After the removal of the DTT solution, the gel slices were incubated with 200 μl of 55 mM iodoacetamide in the dark for 1 h. Washed and dehydrated gel pieces were rehydrated with trypsin alone or with trypsin and ASP-N solution (10 ng/μl for each protease).

Extracted peptides were loaded onto a precolumn (75-μm inner diameter, 6 cm long, packed with 10 μm of C18 resin) before it was connected with an analytic column (75-μm inner diameter, 10 cm long, packed with 1.8 μm of C18 resin). The HPLC gradient was generated by Easy-nLC 1000 (Thermo Fisher Scientific): 300 nl/min flow rate, a 45-min linear gradient from 100% buffer A (0.1% formic acid in water) to 30% buffer B (100% acetonitrile, 0.1% formic acid), followed by a 5-min gradient from 30% to 100% buffer B, then from 100% buffer B to 0% buffer B over 5 min, and lastly a 10-min wash with buffer A. Mass spectrometry data were collected on Q-Exactive (Thermo Scientific): 300 nl/min flow rate, a 45-min linear gradient from 30% to 100% buffer B over 5 min, and lastly a 10-min wash with buffer A. The search results were filtered with a false discovery rate of less than 5% and E values of less than 0.01.

**Structure Modeling** —The tertiary structures for Raptor RNC1 domain and 4E-BP1 fragment were modeled using the I-TASSER server (29). The online program implements a homolog threading protocol followed by fragment assembly and refinement. Five three-dimensional models for each protein were generated based on C-score, a quality measurement for I-TASSER prediction, and the one with the highest C-score was used for subsequent modeling. The structure of the RNC1-4E-BP1 complex was modeled with XPLOR-NIH (30), by refining against knowledge-based torsional angle potential (31) and distance restraints derived from both intra- and intermolecular cross-linking. For each cross-linking, the distances between lysine Ca atoms were constrained <22 Å, and the distances between NZ atoms were constrained <10 Å. The secondary structures, mainly helices, from I-TASSER prediction were kept fixed, and the complex was heated up to 3000 K before gradual cooling to 25 K in a simulated annealing protocol (32).

A set of structure models was generated, all of which satisfy experimental data. The most compact structure that appears most energetically reasonable was selected using program ZRANK (33), which utilizes a scoring function consisting of van der Waals, electrostatics, and desolvation. The structure model was illustrated with the program PyMOL (34).

**Construction of Mutant 4E-BP1 Plasmid** —pET14b PHAS-1 plasmid (Addgene 15679) was used as template for construction of mutant 4E-BP1. A QuikChange Lightning multisite-directed mutagenesis kit (Agilent Technologies, Santa Clara, CA) was used to construct double-site mutations in the region of 4E-BP1 that cross-links with Raptor. PCR primers were designed as 5′-gtcctttgggggtgttaaggccacaggcgagttc-3′ and 5′-ccg-acacctcatcagaaatccgcatgttctg-3′ (obtained from Invitrogen) to double mutate lysine 56 and lysine 68 to leucine. Original plasmids were digested with DpnI restriction enzymes. Mutations were confirmed by sequencing analysis (GENEWIZ, Inc.) (supplemental Fig. S2). Mutant plasmids were transformed into BL21 Star DE3 One Shot® chemically competent cells (Invitrogen) for protein expression.

**Expression and Purification of Mutant 4E-BP1** — Cultures were grown to an A₆₀₀ of 0.5, induced with 1 mM isopropyl β-D-thiogalactopyranoside, harvested by centrifugation, and frozen at −20 °C overnight. The pellet was then thawed on ice and resuspended in lysis buffer (4 ml/g). 1 mg/ml lysozyme was added, and the suspension was incubated on ice for 30 min followed by three periods of sonication. Soluble and insoluble fractions were separated by centrifugation at 100,000 × g for 30 min. The supernatant was retained for purification of soluble proteins. Soluble proteins were purified with nickel-nitrilotriacetic acid-agarose slurry according to the manufacturer’s protocol (Qiagen). Purity was confirmed by SDS-PAGE followed by GelCode Blue staining.

**Raptor/4E-BP1 Far Western Blot** — Far Western blot (overlay) was carried out following published protocol (19). Briefly, 0.2–2 μg of bacterially expressed His-tagged 4E-BP1 was separated on a 15% SDS-PAGE gel and transferred onto a PVDF membrane by electroblotting. The membrane was blocked for 1 h with 5% (w/v) fat-free milk in PBS and 0.02% (v/v) Tween 20.

Using the pLink search engine (21), we searched the mass spectrometry data against a protein sequence database containing human Raptor, human mTOR, and rat 4E-BP1. The following pLink parameters were used: precursor mass tolerance, 50 ppm; fragment mass tolerance, 20 ppm; cross-linker BS³ (cross-linking sites K and protein N terminus; xlink mass shift, 183,0680796; monolink mass shift, 156,0786442); fixed modification C, 57,02146; peptide length minimum, 4 amino acids per chain; peptide length maximum, 100 amino acids per chain; peptide mass minimum, 400 Da per chain; and peptide mass maximum, 10,000 Da per chain). No restriction on peptide terminal residues was applied in pLink search (i.e., treat the peptides as protein digestion products using a nonspecific protease).
The membrane was then incubated overnight at 4 °C with FLAG-Raptor lysate diluted in blocking buffer (1:10). The FLAG-Raptor lysate was obtained by lysing FLAG-Raptor expressing HEK293T cells in a 15-cm dish with 1 ml of lysis buffer (50 mM HEPES, pH 7.4, 150 mM NaCl, 0.1% CHAPS) and centrifuged for 30 min to remove insoluble fraction. The PVDF membrane was washed in blocking buffer and probed with anti-FLAG antibody (1:500) for 1 h, washed again, probed with secondary HRP-conjugated anti-rabbit antibody (1:10,000) (GE Healthcare) for 1 h, incubated with enhanced chemiluminescence solution (ECL) (Pierce) for 60 s, and exposed to autoradiography film.

**RESULTS**

**Cross-linking of Raptor with 4E-BP1 and Identification of Cross-linked Peptides by Mass Spectrometry—**mTORC1 consists of three major proteins: mTOR, Raptor, and mLST8 (G_{mL}). This multiprotein complex functions to phosphorylate substrate proteins including 4E-BP1. This occurs when Raptor recognizes 4E-BP1, which results in the recruitment of 4E-BP1 to the mTOR kinase. To investigate the interaction between mTORC1 and 4E-BP1, we first purified mTORC1 from HEK293T cells stably expressing FLAG-tagged Raptor (established by infecting cells with lentivirus carrying FLAG-Raptor). Cells were lysed in a buffer containing 0.4% CHAPS, and mTORC1 was affinity-purified by using anti-FLAG beads and eluted by using FLAG peptides as described under “Experimental Procedures.” The preparation contained mTOR, Raptor, and mLST8, as detected by staining and Western analysis (Fig. 1, A and B). The purified mTORC1 was active in phosphorylating 4E-BP1, as evidenced by the appearance of phospho-4E-BP1 after incubating mTORC1 with unphosphorylated 4E-BP1 (Fig. 1C). The kinase activity was robust in the presence of Mg^{2+} as well as in the presence of Mn^{2+}.

To examine the interaction of 4E-BP1 with mTORC1, we mixed mTORC1 with 4E-BP1 in the presence of increasing concentration of a cross-linker BS³ that can covalently connect two closely spaced lysine residues (Ca-Ca distance, <24 Å). Western blotting was carried out to identify 4E-BP1 bands on a SDS-polyacrylamide gel. As can be seen in Fig. 2A, incubation with BS³ led to the appearance of new bands that had molecular weights much higher than that of 4E-BP1. A prominent band was detected just above the molecular weight of Raptor. Western analyses using anti-Raptor antibody confirmed the appearance of BS³-dependent Raptor bands that coincide with the bands detected by the anti-4E-BP1 Western (data not shown).

To evaluate whether free Raptor can be cross-linked to 4E-BP1, we prepared Raptor protein separated from the complex. Because Triton X-100 is known to disrupt mTORC1 complex (36), we lysed HEK293T cells stably expressing FLAG-Raptor in a buffer containing 1% Triton X-100 and purified Raptor by the use of anti-FLAG beads and then eluted Raptor using FLAG peptides. Analysis of this preparation by SDS-polyacryl-
amide gel electrophoresis showed only a single band corresponding to the size of Raptor (data not shown). Raptor was then incubated with 4E-BP1 in the presence of BS⁵. As shown in Fig. 2B, a pattern of cross-linked bands obtained was similar to that seen when mTORC1 was used, indicating that it is Raptor that cross-links with 4E-BP1.

The major cross-linked band was cut out, digested with proteases, and subjected to mass spectrometry. Database search was carried out using the pLink program. Fig. 3 shows four representative, high resolution fragmentation spectra of cross-linked peptides between Raptor and 4E-BP1. For example, the cross-linked peptide spectrum shown in Fig. 3A depicts a cross-link between Raptor (KLCLTSLR) and 4E-BP1 (NSPVAKTPPK). Other spectra are shown in supplemental Fig. S1.

The Raptor/4E-BP1 Cross-links Involve the N-terminal Conserved Region 1 of Raptor and a Central Region of 4E-BP1—Table 1 summarizes all the cross-linked peptides between Raptor and 4E-BP1 that we have identified. A total of seven intramolecular cross-links were identified, and six of the spectra show excellent spectral quality. The table also shows how many times these peptide pairs were identified (spectral counts) and the best E values associated with their detection. Fig. 4A shows these cross-links. It is striking that six of the seven cross-links are clustered in a small region on each protein, namely, the RNC1 and the central region of 4E-BP1.

Raptor, a protein of 1335 amino acids, has 52 lysine residues that are located throughout the protein. Only six of these were cross-linked to 4E-BP1. The protein has multiple structural features, the RNC regions 1–3, the HEAT repeats in the middle, and the WD40 repeats that are located in the C terminus. The sites of cross-linking with 4E-BP1 are all located in the RNC1 region except for one that is located in the RNC2 region. For 4E-BP1, we find that all the cross-links with Raptor took place at the central region of this protein. 4E-BP1 is a small protein of 117 amino acid residues, including four lysine residues that are located at the N-terminal and the C-terminal regions as well as two that are located in the internal regions. One of them, Lys⁶⁸, was the major cross-linking residue, whereas another, Lys⁹⁶, also cross-linked to Raptor. These results suggest that the central region of 4E-BP1 defined as the RCR is located closely to Raptor.

In Fig. 4B, we present a model of Raptor/4E-BP1 interaction based on the above cross-linking study as well as by using structure prediction described under “Experimental Procedures.” RNC1 was predicted to have two helices, encompassing residues 95–112 and 127–140, arranged in a parallel fashion. For the 4E-BP1 fragment, residues 57–62 likely form a short helix, followed by a loop structure. When interacting with RNC1, the short helix of 4E-BP1 mainly interacts with the second helix of RNC1 with a large angle between the two helices, whereas the helix C terminus and the following loop swing toward the first helix of RNC1. The structure of a slightly shorter fragment of 4E-BP1 in complex with eukaryotic translation initiation factor 4E has been previously determined using x-ray crystallography in several independent studies. The structural model of Raptor-4E-BP1 complex bears high similarity to these high resolution structures, which further corroborates our findings.

Raptor Intramolecular Cross-links Define Two Regions of Structured Domains in Raptor—In addition to the cross-linked peptides between Raptor and 4E-BP1, our analyses revealed the presence of Raptor-Raptor cross-links. A total of 17 different cross-links were identified, and they are listed in Table 2. Because we cut out the bands that were located just above the Raptor band of 150 kDa but well below 250 kDa, these cross-links most likely represent those that are formed within the same polypeptide, not from a Raptor dimer. Some of the cross-links were repeatedly identified. For example, the cross-link between Lys²⁰⁷ and Lys³³⁵ was observed 132 times. Most cross-links were formed between lysine residues that are both in the N-terminal half of the protein or the C-terminal half except for the cross-link between Lys¹²⁰ and Lys⁸⁸⁴. Fig. 5 depicts these intramolecular cross-links within a Raptor molecule, based on the idea that two lysine residues will cross-link only if they are located nearby. Fig. 5A depicts an extended Raptor structure to indicate the N-terminal and C-terminal regions. Two structured domains, one located in the N-terminal region and the other located in the C-terminal region, are seen. The central region of Raptor (residues 336–689) that includes the HEAT domain had no cross-linked residues, although this area contains six lysines. Interestingly, the sites where the interaction with 4E-BP1 takes place (Lys¹³⁰, Lys¹³¹, and Lys¹⁶⁷) appear to be clustered in one area within the N-terminal portion. Because we have identified a cross-link between residues 120 and 894, it is possible that the N-terminal and C-terminal regions are located near each other. This is depicted in Fig. 5B, and we mention its possible significance under “Discussion.”

Peptides Harboring the RCR of 4E-BP1 Inhibit Raptor-4EBP1 Cross-link—To gain further insight into the site of interaction between Raptor and 4E-BP1, we decided to use synthetic peptides and ask whether they inhibit cross-linking between the two proteins. To do this, we first synthesized a peptide corresponding to 4E-BP1 (amino acids 56–72), which has the sequence KFLMCRNNSPVAKTPPK (peptide 1), and a scrambled version of this sequence as a control (control 1). Two different concentrations of these peptides were added separately to the cross-linking reactions, and the amount of cross-linked Raptor-4E-BP1 was examined on a SDS-polyacrylamide gel. As can be seen in Fig. 6A, 48 μM of peptide 1 indeed inhibited the cross-link between Raptor and 4E-BP1. On the other hand, the
FIGURE 3. Representative high resolution fragmentation spectra of cross-linked peptides between Raptor and 4E-BP1. A, 4E-BP1[NSPVAKTPPK(6)]-Raptor[KLCTSLR(1)]. B, 4E-BP1[KFLMECR(1)]-Raptor[NAKEER(3)]. C, 4E-BP1[NSPVAKTPPK(6)]-Raptor[NAKEER(3)]. D, 4E-BP1[NSPVAKTPPK(6)]-Raptor[YKQSL(3)]. The numbers in parentheses indicate the positions of the cross-linked lysine residues in the peptide sequences. Observed C-terminal fragments (y ions, typically the dominant species) resulting from a peptide bond cleavage of the \( /H9251 \) peptide (the one with a higher mass) are colored purple, and those of the \( /H9252 \) peptide are dark yellow. The corresponding N-terminal fragments are called b ions, from which a further loss of carbonyl results in a ions. The b ions are colored green if the cleavage occurs on the \( /H9251 \) peptide or light blue if the cleavage occurs on the \( /H9252 \) peptide. The a ions are highlighted dark purple if generated from the breakage of the \( /H9251 \) peptide or dark blue if from the \( /H9252 \) peptide.

TABLE 1
Cross-linked peptide pairs between Raptor and 4E-BP1

<table>
<thead>
<tr>
<th>Raptor Sequence</th>
<th>4E-BP1 Sequence</th>
<th>Spectral counts</th>
<th>Best ( E ) value</th>
<th>Spectral quality (manual evaluation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>KLCTSLR</td>
<td>NSPVAKTPPK</td>
<td>68</td>
<td>7.09 ( \times 10^{-11} )</td>
<td>High (Fig. 3A)</td>
</tr>
<tr>
<td>NAKEER</td>
<td>KFLMECR</td>
<td>56</td>
<td>1 ( \times 10^{-6} )</td>
<td>High (Fig. 3B)</td>
</tr>
<tr>
<td>DPLIGMPQK</td>
<td>NSPVAKTPPK</td>
<td>68</td>
<td>2.36 ( \times 10^{-8} )</td>
<td>High (Fig. 3A)</td>
</tr>
<tr>
<td>NAKEER</td>
<td>NSPVAKTPPK</td>
<td>68</td>
<td>5.56 ( \times 10^{-6} )</td>
<td>High (Fig. 3B)</td>
</tr>
<tr>
<td>DLEKKIPQR</td>
<td>NSPVAKTPPK</td>
<td>68</td>
<td>3.23 ( \times 10^{-5} )</td>
<td>High (Fig. 3B)</td>
</tr>
<tr>
<td>YKQSL</td>
<td>NSPVAKTPPK</td>
<td>68</td>
<td>3.75 ( \times 10^{-4} )</td>
<td>High (Fig. 3B)</td>
</tr>
<tr>
<td>PTNGEVWFPKNTQYIPLSIY</td>
<td>NSPVAKTPPK</td>
<td>169</td>
<td>2.37 ( \times 10^{-4} )</td>
<td>Low (Fig. 3C)</td>
</tr>
</tbody>
</table>
same concentration of the control peptide (control 1) did not inhibit the cross-link. We then synthesized another peptide that is located slightly closer to the N terminus. This peptide “GTRIILDRKFLMECRNS” (peptide 2) corresponds to the 4E-BP1 sequence between residues 48 and 64. A scrambled version of this peptide (control 2) was also synthesized. As shown in Fig. 6B, this peptide exhibited strong inhibition of the Rap- tor-4E-BP1 cross-link. Almost complete inhibition was seen in these experiments. In contrast, little inhibition was observed with the scrambled control peptide (control 2). To obtain IC_{50} values of the inhibition, we varied the concentration of peptides. As shown in Fig. 6C, peptide 2 inhibited the formation of Raptor-4E-BP1 cross-link at an IC_{50} value of 15 μM. To examine whether peptide 2 inhibits binding of 4E-BP1 to mTORC1, we purified mTORC1 from HEK293T cells expressing FLAG-Raptor by using anti-FLAG beads. After mixing 4E-BP1 with the beads in the presence of peptide 2 or control peptide, the amount of 4E-BP1 pulled down was examined by Western analysis. As shown in Fig. 6D, the amount of 4E-BP1 bound to mTORC1 was significantly decreased by peptide 2, whereas this effect was less with the control peptide. These results support the results obtained from the CXMS analyses.

Mutations of Residues within the RCR of 4E-BP1 Decrease Its Ability to Serve as a Substrate for mTORC1 in Vitro—We mutated the residues within the RCR of 4E-BP1 to assess the significance of this region for its interaction with Raptor. 4E-BP1 is a small protein that functions without folded structure (43). By PCR-mediated in vitro mutagenesis, we changed

DPLSMGPKQALETIAGNLKQYENWQPRNRYQGSLDPTVEYVKLCNLSRNKAEEVFHYNFGHVRRPTRNGEVEWWFNKNYITQYIPLSY

Mutations at positions 1335RNC1 RNC2 RNC3 HEAT WD40 89 180 117 56 73 117 RAIP TOS
two lysine residues within the RCR of 4E-BP1. The mutant protein was purified with His$_6$ nickel-nitrilotriacetic acid matrix (supplemental Fig. S2). Both the wild type and mutant 4E-BP1 were used for kinase assay to detect phosphorylation of 4E-BP1. As shown in Fig. 7A, although the wild type 4E-BP1 showed strong phosphorylation of 4E-BP1 as detected by the use of anti-phospho (Thr$^{37/46}$) antibody as well as by anti-phospho (Ser$^{65}$) antibody, the phosphorylation was dramatically reduced with the mutant 4E-BP1 protein. We also examined the binding of the 4E-BP1 mutant to mTORC1. The results shown in Fig. 7B demonstrate that the mutant protein binds much less efficiently than the wild type protein. Furthermore, Far Western blot study shows that the mutant 4E-BP1 interacts with FLAG-Raptor significantly less than the wild type (Fig. 7C). The Western blot in the upper panel shows the amount of total 4E-BP1 protein loaded onto the gel, whereas the lower panel shows the amount of FLAG-Raptor binding to the wild type or the mutant 4E-BP1.

**Mutations of Residues within the RCR of 4E-BP1 Decrease Its Ability to Serve as a Substrate for mTORC1 in Vivo**—Next, we expressed the mutant 4E-BP1 in HEK293T cells. For this purpose, we inserted FLAG epitope-tagged 4E-BP1 (wild type and mutant) into pcDNA3 plasmid by PCR as described under “Experimental Procedures.” FLAG-4E-BP1 plasmids (wild type and mutant) were then transfected with Lipofectamine 2000 into HEK293T cells. 24 h after transfection, cells were collected and lysed, and FLAG-4E-BP1 was immunoprecipitated with anti-FLAG M2 affinity gel. The eluted proteins were analyzed.

### TABLE 2

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**FIGURE 5.** Two views of Raptor based on the identification of Raptor-Raptor cross-linked peptides. Because the residues that are cross-linked suggest that they are in close proximity, it is possible to place cross-linked residues closeby. A, a scheme showing a stretched out structure to demonstrate two structured domains. B, a scheme showing a folded structure based on the proximity of residues 120 and 894. Gray circles show the lysine residues that are identified to be cross-linked with 4E-BP1. Black squares show the lysine residues in Raptor that cross-linked with mTOR.
by Western blot for phosphorylation of 4E-BP1 as well as for interaction with Raptor. As shown in Fig. 8A, phosphorylation of 4E-BP1 was dramatically reduced with the mutant 4E-BP1 protein compared with the wild type. We also examined their co-immunoprecipitation with Raptor. Significant reduction of Raptor co-immunoprecipitation with the mutant 4E-BP1 compared with the wild type was observed (Fig. 8B).

DISCUSSION

In this paper, we characterized the interaction of mTORC1 and its substrate 4E-BP1 using the CXMS technology. First, we found that 4E-BP1 could be cross-linked with Raptor using purified mTORC1 as well as free Raptor. We then characterized the cross-linked band by mass spectrometry and identified cross-linked peptides that contain both Raptor and 4E-BP1 sequences. Multiple cross-linked peptides were identified that were clustered primarily to a single region for each protein. These results represent the first observation of Raptor and 4E-BP1 cross-links, and the study revealed that the RNC of Raptor and the RCR of 4E-BP1 are in close proximity. Computational structural prediction (Fig. 4B) can be made regarding the residues in the two proteins.

Our cross-link results were further substantiated by first carrying out peptide inhibition studies. Two peptides that encompass the Raptor interaction site of 4E-BP1 inhibited Raptor–4E-BP1 cross-links, whereas the control peptides each having a scrambled sequence did not. Peptide 2 gave strong inhibition at 15 \( \mu \text{M} \) concentration. Second, we have mutated residues within the 4E-BP1 region we defined by our cross-linking study and showed that the mutations dramatically decrease binding of 4E-BP1 to mTORC1 and reduce phosphorylation by mTORC1. In cells, phosphorylation of the mutant 4E-BP1 was significantly decreased compared with that of the wild type protein. In addition, Raptor binding was significantly decreased with the mutant 4E-BP1.

Our results on 4E-BP1 suggest that a region encompassing residues 56 and 72 is in close proximity with Raptor. This region is highly conserved among 4E-BP family members (44). The sequence KFLMECRNSPVAKTPP in 4E-BP1 corresponds to the sequence KFLDLRRNSPMATTPP in 4E-BP2 and KFLLRECKSPARPTTPP in 4E-BP3. Two mTORC1-mediated phosphorylation sites (45) are present in this region. A sequence that is known to interact with eIF4E, as defined by the co-crystallization of eIF4E and a 4E-BP1 peptide (46), is located on the N-terminal side of the region. The TOS and RAIP motifs have been proposed to play important roles for the interaction between Raptor and mTORC1 substrates. However, the loca-


The same amount of the mutant and the wild type protein was examined by using mTORC1 attached on beads as described under “Experimental Procedures.” The same antibody was raised against the human protein, Rattus 4E-BP1 residue numbering should be shifted by one residue (−1 residue). The amount of mTORC1 used was 0 μl (lanes 1 and 4), 2 μl (lanes 2 and 5), and 4 μl (lanes 3 and 6) of purified protein. Anti-4E-BP1 antibody was used to estimate the amount of total 4E-BP1, B, binding of the mutant and the wild type 4E-BP1 with mTORC1. The ability of 4E-BP1 to bind mTORC1 was examined by using mTORC1 attached on beads as described under “Experimental Procedures.” The same amount of the mutant and the wild type protein was used for this assay. C, binding of the mutant and wild type 4E-BP1 protein with Raptor. The wild type and mutant 4E-BP1 were run on a 15% SDS-PAGE gel and probed for total 4E-BP1 with anti-4E-BP1 antibody (upper panel). Protein was loaded in incrementally increasing concentrations. The lower panel shows the result of Far Western blot (overlay) in which the mutant and the wild type 4E-BP1 were run on a 15% SDS-PAGE gel, incubated with the lysate of FLAG-Raptor expressing HEK293T cells, and probed with anti-Raptor antibody to detect binding interactions taking place between Raptor and 4E-BP1.

Our study also revealed intramolecular cross-links that occur within the Raptor protein. The locations of these intramolecular cross-links suggest that the Raptor protein has two separate structured domains: one in the N-terminal half and the other in the C-terminal half of the protein. The N-terminal structured domain encompasses RNC1 and RNC2 domains. Interestingly, the lysine residues found to be cross-linked with 4E-BP1 are all located in a small region within this structured domain (Fig. 5). On the other hand, our recent preliminary results raise the possibility that the interaction with mTOR takes place mainly at the C-terminal region. We have recently identified Raptor-mTOR cross-linked peptides. Three cross-linked peptides were identified, and all cross-linked lysines in Raptor were located within the C-terminal structured domain (Lys1332 and Lys1008). Two cross-linked peptides had Raptor cross-linked to the mTOR residue 2507, a residue located within the kinase domain of mTOR. These results suggest that 4E-BP1 and mTOR interact with different regions of Raptor: 4E-BP1 at the N-terminal structured region and mTOR at the C-terminal structured region. Based on our identification of Raptor-Rap2507 cross-links, Raptor is likely to be folded as depicted in Fig. 5B. If this were the case, the N-terminal region and the C-terminal region of Raptor are located in close proximity, and this will result in bringing the mTOR kinase domain and 4E-BP1 together.

4E-BP1 in its nonphosphorylated form functions to inhibit protein synthesis by binding to eIF4E, a cap-binding protein that facilitates initiation of protein synthesis. eIF4E is also an oncogene as demonstrated by its ability to confer transformed phenotypes and tumor formation in mice (47), and overexpres-
sion of eIF4E is observed in a variety of cancer cells, most notably in acute myelogenous leukemia (48). Ectopic expression of nonphosphorylated form of 4E-BP1 partially reverses transformed phenotypes (49). Thus, blocking the interaction between Raptor and 4E-BP1 is expected to increase the concentration of nonphosphorylated 4E-BP1, which in turn would inhibit eIF4E and tumor growth when the tumor is driven by the overexpression of eIF4E. Small molecule inhibitors have been identified to inhibit the action of eIF4E (35, 50). Further understanding of how Raptor interacts with 4E-BP1 may open up a novel approach to inhibit eIF4E-mediated tumor formation.

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REFERENCES

translation initiation factors eIF4E and eIF4G. 


