Structural basis for shieldin complex subunit 3–mediated recruitment of the checkpoint protein REV7 during DNA double-strand break repair

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Shieldin complex subunit 3 (SHLD3) is the apical subunit of a recently-identified shieldin complex and plays a critical role in DNA double-strand break repair. To fulfill its function in DNA repair, SHLD3 interacts with the mitotic spindle assembly checkpoint protein REV7 homolog (REV7), but the details of this interaction remain obscure. Here, we present the crystal structures of REV7 in complex with SHLD3’s REV7-binding domain (RBD) at 2.2–2.3 Å resolutions. The structures revealed that the ladle-shaped RBD in SHLD3 uses its N-terminal loop and C-terminal α-helix (αC-helix) in its interaction with REV7. The N-terminal loop exhibited a structure similar to those previously identified in other REV7-binding proteins, and the less-conserved αC-helix region adopted a distinct mode for binding REV7. In vitro and in vivo binding analyses revealed that the N-terminal loop and the αC-helix are both indispensable for high-affinity REV7 binding (with low-nanomolar affinity), underscoring the crucial role of SHLD3 αC-helix in protein binding. Moreover, binding kinetics analyses revealed that the REV7 “safety belt” region, which plays a role in binding other proteins, is essential for SHLD3–REV7 binding, as this region retards the dissociation of the RBD from the bound REV7. Together, the findings of our study reveal the molecular basis of the SHLD3–REV7 interaction and provide critical insights into how SHLD3 recognizes REV7.

DNA double-strand breaks (DSBs)3 are extremely vicious DNA lesions that may cause carcinogenesis or cell death if not properly repaired. In vertebrate cells, two main repair pathways, nonhomologous end-joining (NHEJ) and homologous recombination (HR), are employed in eliminating the cytotoxic DSBs and thereby ensuring genomic integrity (1, 2). The decision-making process of repair pathways is a critical step during DSB response, which involves several elements such as the cell cycle state, the epigenetic regulation, and the DNA end resection (3, 4). 53BP1, in concert with its upstream regulators and downstream effectors, spearheads the restriction of end resection to promote NHEJ and antagonizes BRCA1-mediated HR repair, whose initiation step relies on resected DNA termini (5–14). Recently, several elegant studies have identified a previously-uncharacterized protein complex, named Shieldin, working as a downstream effector of 53BP1 to protect DNA end from resection and to facilitate NHEJ repair in a 53BP1-dependent manner (15–21). Similar to loss of 53BP1, deficient Shieldin leads to reactivation of end resection, impairs NHEJ activities, and affects the immunoglobulin class switching process (22).

Shieldin consists of four subunits: SHLD3 (also known as CTC-534A2.2 or RINN1); SHLD2 (also known as FAM35A or RINN2); SHLD1 (also known as C20ORF196 or RINN3); and REV7 (also known as MAD2L2). Apart from REV7, the other three Shieldin components had not been characterized before. Among these subunits, SHLD3 is the most apical element (16, 19, 22) that bridges REV7 to RIF1 and fills the gap in the 53BP1–RIF1–REV7 axis. Therefore, SHLD3 and REV7 form a sub-complex, which is considered as the DSB-recruitment module, while SHLD2, along with SHLD1, functions as the ssDNA-binding module within the complex (16). Knockdown of SHLD3 largely reduced the localization intensities of REV7/SHLD2/SHLD1 to the ionizing radiation-induced foci region, highlighting its RIF1-proximal role (16, 19). Importantly, depletion of SHLD3 dampens NHEJ-related events, activates DNA end resection, and restores HR in a BRCA1-mutated context, thus rendering these cells poly(ADP-ribose) polymerase inhibitors–resistant (15, 16, 19). Although SHLD3 is essential for recruitment and function of Shieldin by directly interacting with REV7, the underlying molecular mechanism remains elusive.

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This article contains Figs. S1–S4.

1 The abbreviations used are: DSB, double-strand break; RBD, REV7-binding domain; PDB, Protein Data Bank; NHEJ, nonhomologous end-joining; HR, homologous recombination; ITC, isothermal titration calorimetry; MBP, maltose-binding protein; SPR, surface plasmon resonance; RMSD, root mean square deviation; αC-helix, C-terminal α-helix; ssDNA, single-strand DNA; RBM, REV7-binding motif; IMAC, immobilized metal-affinity chromatography; Ni-NTA, nickel-nitrilotriacetic acid; IPTG, isopropyl β-D-thiogalactoside; TLS, translesion DNA synthesis; CAMP, chromosome alignment-maintaining phosphoprotein; SFB, S protein, FLAG, and streptavidin-binding peptide tag.
In this study, we report two crystal structures of REV7 in complex with the SHLD3–RBD. The complex structure shows that SHLD3–RBD forms a “ladel” configuration when it binds to REV7, comprising the N-terminal loop and the C-terminal α-helix (αC-helix) of SHLD3–RBD, which constitute the corresponding “stem” and “base” part of the ladle-shaped structure. The N-terminal loop is structurally similar to other REV7-binding proteins due to the presence of a well-established REV7-binding motif, but the αC-helix displays a binding mode distinct from known REV7-binding partners. Biochemical analyses, coupled with thermodynamic measurements, have verified their contribution to REV7 binding. Importantly, we demonstrated that the αC-helix region of SHLD3–RBD is a crucial contributing factor of robust SHLD3–REV7 interaction. Additionally, we present the first kinetic view on the “safety-belt” binding mechanism underlying the SHLD3–REV7 complex via surface plasmon resonance (SPR) assays. In sum, this work is the first structural study toward the role of REV7 in DSB repair and provides structural insights into how the DSB recruitment module assembles within the Shieldin complex.

Results

SHLD3 employs REV7-binding domain to interact with REV7

SHLD3 is a 250-amino acid polypeptide with a translational elongation initiation factor eIF4E-like domain in its C terminus (15) and contains two presumptive REV7-binding motifs (RBM, defined by PXxP sequence, where X represents any amino acid residue and the lowercase p is a less important proline residue and can be replaced by an alanine in some cases) in the N-terminal region (Fig. 1a). Previous work has mapped the region (residues 29–83), which harbors the second RBM in the N-terminal region of SHLD3 that interacts with REV7 (19). In agreement, we also observed this fragment (residues 29–83) displayed a tight binding to REV7 with a dissociation constant ($K_D$) of 13.8 ± 0.4.nM and a 1:1 binding stoichiometry (Fig. 1b), as measured by ITC assays. However, no binding was detected between the SHLD3 RBM1-containing peptide (residues 1–28) and REV7 (Fig. 1b), suggesting that the second RBM-containing region (residues 29–83) of SHLD3 plays a dominant role in dictating REV7 binding, although the first RBM may have other functions and is dispensable for this interaction. Multiple sequence alignment results also support this finding because the second RBM is invariably conserved in vertebrates, in stark contrast to the first RBM of poor conservation (Fig. S1a). We therefore generated the REV7–SHLD3(29–83) complex and performed crystallization trials. Analogous to previous reports, one (R124A) or two (R124A/A135D) dimer-breaking mutations were introduced into REV7 to improve protein homogeneity and promote crystal packing (23–25). But unfortunately, no crystals were obtained. Secondary structure prediction results suggested that the 29–60- and 74–83-amino acid regions of SHLD3 are mainly disordered (Fig. S1a), and the flexibility of these regions, we propose, may prejudice crystal packing. Hence, we next co-purified REV7 complexed with various truncated constructs of SHLD3 and found that the SHLD3(46–74) is sufficient for REV7 binding (Fig. S1b). Consistently, the SHLD3(46–83) peptide retained full binding affinity as SHLD3(29–83) with a dissociation constant of 15.7 ± 0.8 nM in ITC measurement (Fig. 1b), indicating that the amino acid residues ranging from 29 to 45 of SHLD3, although conserved, are dispensable for REV7 interaction. Therefore, we named residues 46–74 of SHLD3 as the RBD and purified the REV7–SHLD3 (RBD) complex for the next round of crystal screening. Encouragingly, the REV7–SHLD3 (RBD) complex yielded diffraction-quality crystals.

Overall structure of the SHLD3–REV7 complex

The structures of the REV7R124A–SHLD3 (RBD) and REV7R124A, A135D–SHLD3 (RBD) complex were determined by the molecular replacement method using 2.2 and 2.3 Å resolution data, respectively. Both crystals contain one single copy of the SHLD3–REV7 complex per asymmetric unit, and residues 9–207 of REV7 and 49–73 of SHLD3 were successfully modeled, whereas other residues were not included in the final models due to indiscernible electron density. The crystallographic and geometric statistics of the refined models are exhibited in Table 1. Because these two structures are nearly identical, the REV7R124A–SHLD3 (RBD) structure with higher resolution was chosen as a representative for further description.

Within the structure, REV7 displays a canonical HORMA (Hop1, Rev7, and Mad2) fold and adopts the closed conformation with SHLD3 peptide tightly held by the safety-belt loop of REV7 (Fig. 1c). Superimposition of REV7 in our complex with the previously reported structures of REV7/REV3–RBM1 (PDB code 3ABD) (24), REV7/CAMP (PDB code 5XPT) (26), and REV7/REV3–RBM2 (PDB code 6BC8) (25) revealed overall root mean square deviation (RMSD) values of 0.830, 0.953, and 0.915 Å, respectively, suggesting the global similarity of REV7 molecules in these protein complexes, except for some zones showing a high degree of flexibility (Fig. S2). The most striking difference occurs at the REV7 adaptor region (residues 156–174), which forms a short α-helix induced by SHLD3 binding (Fig. S2a), similar to the REV7 structure within the context of the TLS complex (Fig. S2e) (27–29) but distinct from the other reported REV7–RBM heterodimer complex where this region was either missing (24, 26) or subject to conformational changes (Fig. S2, b–d) (25). The structure diversity in this area reveals that the adaptor region of REV7 is capable of reorganizing its shape to accommodate various binding partners. Furthermore, the relative higher B factors of this region also reflect the structural flexibility and imply elasticity suitable for different binding sequences (Fig. S3).

Interestingly, SHLD3–RBD forms a ladle-shaped structure upon binding to REV7, with residues 49–58 (N-terminal loop region) and 59–73 (αC-helix region), respectively, constituting the “stem” and “base” of the ladle (Fig. 1c). The “stem” part contains a well-established RBM, and thus overlays well with other reported RBM, whereas the “base” region protrudes toward the back of REV7 and displays conformational variance compared with other REV7-binding peptides (see below).

Structural basis for REV7–SHLD3 interaction

The observed interaction between the N-terminal loop of SHLD3–RBD and REV7 is similar to other reported complexes due to the presence of RBM. In detail, residue Pro-50 of SHLD3
**Figure 1. SHLD3 employs REV7-binding domain to interact with REV7.**

**a.** schematic domain architecture of SHLD3. The RBD and EIF4E-like domain of SHLD3 are depicted in magenta and green, respectively. The protein sequences of the presumptive REV7-binding motif 1 (RBM1) and RBD used in crystallization are displayed with the defined prolines in two RBMs colored in red. The secondary structure illustration of the RBD region is also shown above the protein sequence.

**b.** isothermal titration calorimetric analysis of three different SHLD3 fragments binding to REV7. Raw data and fitting curves of the integrated data for various SHLD3 proteins are shown together with corresponding KD values.

**c.** overall structure of REV7 in complex of SHLD3 presented in cartoon mode. The SHLD3 fragment is shown in magenta, and the REV7 molecule is colored light blue, except for the C-terminal region of REV7, which is termed the safety-belt region (residues 153–207), highlighted in pale green. The N-terminal loop and αC-helix region of SHLD3–RBD are indicated.
Table 1  

|                      | SHLD3–REV7 (R124A) | SHLD3–REV7 (R124A/A135D) |
|----------------------|---------------------|---------------------------|
| **Data collection**  |                     |                           |
| Space group          | P3_21               | P3_21                     |
| Cell dimensions      |                     |                           |
| a, b, c (Å)          | 61.08, 61.08, 133.24 | 60.64, 60.64, 133.48      |
| Resolution (Å)       | 49.16–2.24 (2.31–2.24)* | 44.99–2.31 (2.39–2.31)   |
| Rmerge/Rfree (%)     | 0.042 (1.019)       | 0.049 (0.935)             |
| I/σ(I)               | 19.6 (2.4)          | 16.7 (2.4)                |
| CC1/2 (%)            | 0.999 (0.959)       | 0.999 (0.951)             |
| Completeness (%)     | 99.8 (99.6)         | 99.9 (99.9)               |
| Multiplicity (%)     | 9.5 (10.1)          | 9.3 (9.8)                 |
| **Refinement**       |                     |                           |
| Resolution (Å)*      | 49.16–2.24          | 44.49–2.31                |
| Anisotropy (Å)*      | 2.5, 2.5, 2.2       | 2.6, 2.6, 2.3             |
| No. of reflections   | 11,810              | 10,644                    |
| Rwork/Rfree (%)      | 20.4/24.6           | 21.0/26.2                 |
| No. of atoms         |                      |                           |
| Protein              | 1721                | 1728                      |
| Water                | 11                  | 14                        |
| Ion/ligand           | 5                   | 5                         |
| B-factors            |                      |                           |
| Protein              | 67.1                | 69.9                      |
| Water                | 59.6                | 57.8                      |
| Ion/ligand           | 126.2               | 153.0                     |
| RMSD                 |                      |                           |
| Bond lengths (Å)     | 0.010               | 0.003                     |
| Bond angles (°)      | 1.059               | 0.626                     |
| Ramachandran plot    |                      |                           |
| Favored (%)          | 94.55               | 96.82                     |
| Allowed (%)          | 5.45                | 3.18                      |
| Outliers (%)         | 0                   | 0                         |
| MolProbity score     | 1.74                | 1.43                      |

* Values in parentheses are for the highest-resolution shell.

To appraise the contribution of residues in the N-terminal loop to the SHLD3–REV7 interaction, we employed an in vitro co-purification-based Ni-pulldown assay that was successfully used in previous studies (24, 26). Interfacial residues in SHLD3 or REV7 were mutated into alanine, and the amount of SHLD3 peptide co-purifying with immobilized His-tagged REV7 was evaluated and quantified. As expected, single alanine substitutions of REV7 Tyr-63 and Trp-171 showed dramatically reduced binding between REV7 and SHLD3, with relative band intensities being ~67.8 and 65.5% of the control level, respectively (Fig. 2b), whereas alanine substitution of Phe-169 had no significant effect on binding (Fig. 2b), suggesting that it makes a minor contribution to the SHLD3–REV7 interaction, in contrast to the significant contribution of this residue in formation of the CAMP–REV7 complex (26). Similar results were also observed in single Y37A, H57A, and F146A REV7 mutants that displayed only small changes in band intensities (Fig. 2b). But double mutation of Tyr-37 and Phe-146 to alanine decreased the binding by 21.4% (Fig. 2b), underlining their synergistic effect on mediating hydrophobic interaction with Pro-57 and Pro-58 of SHLD3. Notably, simultaneous alanine substitutions of Tyr-63 and Trp-171 in REV7 brought the most remarkable effect, with SHLD3–REV7 binding sharply reduced to about 11.2% of the control level (Fig. 2b), further suggesting van der Waals interactions formed by the two aromatic residues are important for maintaining SHLD3–REV7 interaction.

As for SHLD3, we found that single alanine substitutions of Pro-53, Pro-57, or Pro-58 in RBM almost retain full-binding ability to REV7 as the WT protein (Fig. 2c), a result distinct from CAMP–REV7 interaction where a single mutation of Pro-341 (also in RBM) of CAMP strikingly hinders its binding to REV7 (26). P53A/P57A double mutant also had no significant impact on the interaction between REV7 and SHLD3 (Fig. 2c). However, severely diminished interaction was observed when both Pro-53 and Pro-58 were replaced by alanine, and the Pro-57/Pro-58 double mutant also impaired the binding of SHLD3 to REV7 to 66.1% of that WT level (Fig. 2c). These results revealed Pro-58 is the most important one among the three prolines of RBM, whereas Pro-53 and Pro-57 exhibit relatively less significance during SHLD3–REV7 interaction. Additionally, the P50A mutant showed no decrease in band intensities, indicative of a minor role it also plays in interaction. To cross-validate the effects of the N-terminal loop mutants on REV7 interaction, we also conducted ITC measurements to observe the change of their binding affinity to REV7. In agreement with those observations in pulldown assays, the P53A/P57A double point mutant only had a subtle effect on binding with a $K_D$ of 80.0 ± 2.5 nm, and the P57A/P58A mutant moderately diminished the binding by about 20.3-fold, with a $K_D$ of 280.1 ± 4.6 nm (Fig. 2d). Once more, the largest defect was generated by the integration of two alanines into Pro-53 and Pro-58, which displayed a substantially reduced binding affinity of 2.1 ± 0.1 μM, nearly a 152.2-fold decrease that of the WT protein (Fig. 2d).

To determine the effects of these mutations in human cells, the Myc-tagged full-length SHLD3 harboring these mutations was co-transfected with SFB-tagged REV7. Consistent with previous reports (15, 19) and our in vitro results, the SHLD3 P53A/P58A double mutant largely decreased its binding to REV7 (Fig. 2e). Moreover, we observed that SHLD3 P57A/P58A double mutant also dramatically impaired the SHLD3–REV7 interaction, whereas SHLD3 P53A/P57A double mutant did not notably affect its binding to REV7 (Fig. 2e). However, as shown in Fig. 2f, SHLD3 retained binding ability to SFB-tagged REV7 F169A/W171A but failed to pull down SFB-tagged REV7.
Structural insights into REV7 recruitment by SHLD3

**Figure a**

Structural model of REV7 and SHLD3 interactions.

**Figure b**

Western blot analysis of REV7 mutants with SHLD3.

**Figure c**

Western blot analysis of SHLD3 mutants with REV7.

**Figure d**

Heat of injection for SHLD3 mutants.

**Figure e**

Pull-down assay of Myc-SHLD3 with SFP-REV7 and SFB-REV7.

**Figure f**

Pull-down assay of MBP-SHLD3 with anti-FLAG antibodies.
Y63A/W171A, indicating that the interaction between Tyr-63 and Pro-57/Pro-58 of SHLD3 plays a more vital role in SHLD3 binding in vivo than the interaction mediated by Trp-171 and Pro-53 of SHLD3. Collectively, these results corroborated that RBM in the N-terminal loop region of SHLD3–RBD is critical for REV7 binding.

αC-helix enables SHLD3’s high-affinity recognition of REV7

Intriguingly, we observed the αC-helix of SHLD3–RBD interacts with REV7 through fitting into a shallow groove in the REV7 surface, acting as the base of the ladle configuration (Fig. 3a). This special conformation has not been observed in other known REV7-binding partners, as the “base” region of SHLD3–RBD points toward the backside of REV7 and forms a torsion angle of about 96.5°, whereas previous structures were either lacking the corresponding α-helix (CAMP) or adopting a relatively linear configuration (REV3–RBM1 and REV3–RBM2) (Fig. 3, a and b). Superimposition of the N-terminal loop region of SHLD3 RBD revealed the RMSD of 1.107 and 1.213 Å, although structural differences of the full-length RBD from SHLD3 and from REV3–RBM1 or REV3–RBM2 leads to substantially increased RMSD for REV3–RBM1 (2.610 Å) and for REV3–RBM2 (2.649 Å), further suggesting conformational variations in the αC-helix region of SHLD3 RBD. Moreover, this unique conformation ensures a larger contact interface between SHLD3 and REV7, with an interacting area of 359.2 Å², as compared with the binding interface for REV3–RBM1 (166.5 Å²), REV3–RBM2 (241.7 Å²), and CAMP (78.3 Å²) (Fig. S4a).

It is conceivable that the larger interaction interface in this region contributes to stabilize and promote the binding between SHLD3 and REV7. Indeed, we observed that a SHLD3 fragment lacking the αC-helix region shows a remarkably reduced binding affinity in ITC assays, with a $K_D$ of 505.1 ± 9.7 nM (Fig. 3c). Furthermore, the WT SHLD3 fragment interacts with REV7 in an enthalpically and entropically favorable manner, and the binding is mostly entropy-driven, but deletion of the αC-helix region largely compromised the favorable entropy (Fig. S4b), suggesting that the hydrophobic contacts conducted by this interface is crucial for the high-affinity SHLD3–REV7 interaction. A recent study reported that Ran also binds to REV7 via the classical RBM, but it is devoid of the corresponding SHLD3 αC-helix (30). The binding affinity between Ran and REV7 is about 900 nM, a level parallel to the SHLD3 (Δ59–74) fragment (505.1 ± 9.7 nM) but considerably lower than WT peptide (13.8 ± 0.4 nM), which further accentuates the importance of SHLD3 αC-helix region in dictating high-affinity REV7 binding.

In this region of SHLD3, Ile-60 and Val-68 are two anchor residues as their hydrophobic side chains resemble a clamp that tightly holds the corresponding cavity in the REV7 surface (Fig. 2a and Fig. S4a). To dissect the roles of Ile-60 and Val-68 in mediating REV7 association, we made mutations to see whether the mutants displayed impaired binding. As shown in Fig. 3c, I60A mutation had an effect on SHLD3–REV7 interaction, reducing the binding to ~74.3% of the WT level. Furthermore, substitution of Ile-60 with an electrostatic residue markedly impaired the binding to REV7, with I60K and I60D mutants displaying band intensities of 40.6 and 51.1% that of WT SHLD3 (Fig. 3d). Even though the I60A mutant did not display an obvious binding defect in vivo, SFB-tagged SHLD3 I60K indeed exhibited an impaired binding to REV7, and the SHLD3 P53A/I60K dual mutant showed a severely reduced capability to pull down REV7 (Fig. 3e). In addition, the I60K mutant also exhibited impaired binding affinity ($K_D$ of 236.9 ± 10.5 nM) and decreased favorable entropy in ITC assays (Fig. 3c, Fig. S4b). These results validated the critical role of Ile-60 in maintaining SHLD3–REV7 cross-talk both in vitro and in vivo. It should be reminded that the SHLD3 Ile-60 is highly conserved in vertebrates (Fig. S1), implying this hydrophobic interaction is conserved and important in those species, too.

Moreover, the V68D mutant displayed an even weaker binding affinity (781.3 ± 9.2 nM) when compared with the I60K mutant (Fig. 3c), and the favorable entropy was also largely compromised (Fig. 4b).

The contribution of these two anchor residues is further affirmed by the Ni-pulldown assay using corresponding REV7 mutants. As shown in Fig. 3f, the dual mutation Y37S/H57A had no prominent effect on SHLD3–REV7 binding, as did the aforementioned H57A mutant. However, integration of Y37S/H57A with an extra I41D mutation markedly diminished the binding of SHLD3 to REV7 to ~28.9% that of the WT level, again demonstrating the hydrophobic interactions mediated by Ile-60 and Val-68 of SHLD3 are central to the high-affinity association between SHLD3 and REV7. Together, these results suggested that the αC-helix region of SHLD3–RBD underlies robust REV7 binding.

Safety-belt region promotes SHLD3–REV7 binding by retarding dissociation process

Like other RBM-containing protein, SHLD3 is also entangled with REV7 through a well-established “safety-belt” mode, but little is known about the mechanism of this intriguing process. To get better insights into the binding mechanism underlying the SHLD3–REV7 complex, we used SPR assays to study the

Figure 2. Structural basis of SHLD3(RBD)–REV7 complex. a, stereo view of the interaction details between SHLD3 and REV7. Colors of molecules correspond to those in Fig. 1c, and residues involved in the interaction are labeled. The hydrogen bonds are shown as yellow dotted line. b and c, Ni pulldown assays using His–REV7 co-expressed with SHLD3(29–83) fragments. After rounds of extensive washing, the bound proteins were resolved by SDS-PAGE stained with Coomassie Brilliant Blue (upper panel). His–REV7(B12A4) was used as control sample. The bottom plot panel shows the relative abundance of the pulled down SHLD3 proteins. The relative band intensities of SHLD3(29–83) divided by band intensities of REV7 were normalized to those of the control. Values are represented as means ± S.D. from three independent experiments. One-tailed Student’s tests are indicated: *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; and ****, $P < 0.0001$. d, representative ITC results of the interactions between SHLD3 RBM mutants and REV7. Representative binding curves for WT, P53A/P57A, P53A/P58A, and P57A/P58A SHLD3 are shown as black circle, yellow circle, green circle, and blue circle, respectively. The binding affinity of each mutant is shown accordingly. e, 293T cells were transfected with plasmids encoding Myc-tagged SHLD3 WT and its mutants together with the plasmid encoding SFB-tagged REV7. Immunoprecipitation reactions were conducted using S protein beads and then subjected to Western blotting using the indicated antibodies. f, beads coated with bacterially expressed MBP-fused SHLD3 was incubated with cell lysates containing exogenously expressed SFB-tagged REV7 WT and its mutants. Immunoblotting experiments were carried out using the indicated antibodies.
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b

SHLD3
REV3-RBM1
CAMP
REV3-RBM2

Heat (Kcal/mol of injected)
Molar ratio
0
1
2
3

SHLD3
Kd
WT 13.8±0.4 nM
I60K 236.9 ± 10.5 nM
V68D 781.3 ± 9.2 nM
Δ59-74 505.1 ± 9.7 nM

d

SHLD3 mutants

Relative band intensity (%)
0
20
40
60
80
100

Control
I60A
I60K
I60D

His-REV7
SHLD3

f

REV7 mutants

Relative band intensity (%)
0
20
40
60
80
100

Control
Y75S
Y75M
Y75M + D57A

His-REV7
SHLD3
binding kinetics of this interaction. Purified SHLD3(29–83) fragment was immobilized onto a CM5 sensor, and 2-fold increasing concentrations of purified MBP–REV7 or mutants were injected and analyzed. As presented in Fig. 4a, the association rate constant ($k_{\text{on}}$) and the dissociation rate constant ($k_{\text{off}}$) of REV7 WT are $8.11 \times 10^3$ 1/ms and $1.24 \times 10^{-4}$ 1/s, respectively. These rate constants generate a calculated equilibrium dissociation constant ($K_D$) of 15.3 nM (Fig. 4e), a number highly consistent with the ITC result (13.8 ± 0.4 nM) described above, confirming that the interaction between SHLD3 and REV7 has a $K_D$ value in the low nanomolar, tight-binding range. Importantly, the safety-belt mutant W171A displayed an 100-fold
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higher dissociation rate constant of $1.20 \times 10^{-2}$ 1/s and a 1.65-fold lower association rate constant of $4.89 \times 10^3$ 1/ms in contrast to WT protein (Fig. 4, b and e). As a result, the calculated $K_d$ of the W171A mutant is 2.46 $\mu$m, and this binding affinity is in accordance with the binding defect observed in in vitro Ni-pulldown assays (Fig. 2b). The prominent difference in dissociation rate manifested that the Trp-171 in the adaptor region of REV7 serves as a contributing factor of the tight-binding nature of the SHLD3–REV7 interaction by slowing down the dissociation process between REV7 and its cargoes. Moreover, we did not observe the interaction between an adaptor region deletion mutant ($\Delta$156–174) and SHLD3 (Fig. 4, c and e), further suggesting this safety-belt loop contributes to REV7’s interaction with its partners. Notably, REV7 Y37S/I41D/H57A mutant also showed an undetected binding to the SHLD3 peptide (Fig. 4, d and e), underscoring the contribution of interactions mediated by these residues and the $\alpha$C-helix region of SHLD3 to complex formation.

**Discussion**

Recently, several contemporary reports exhaustively described the discovery of a four-subunit complex, termed Shieldin, working as the downstream effector of 53BP1–RIF1 to directly bind ssDNA and recruit CTC1-STMN1-TEP1–polymerase $\alpha$ complex at damage sites that directs 3’ to 5’ fill-in DNA synthesis in opposition of the action of those end-processing nucleases (15–21, 31, 32). Among the four Shieldin subunits, SHLD3 is the most apical component that bridges REV7 to RIF1 and is crucial for correct localization of Shieldin at damage sites (16, 19, 22). Therefore, SHLD3–REV7 is considered as the DSB recruitment module within the Shieldin complex.

In this report, we determined the complex structure of SHLD3–RBD and REV7, where SHLD3–RBD adopts an unexpected ladle configuration to recognize REV7. Elegant studies by Gupta et al. (19) and Ghezraoui et al. (15) have shown that SHLD3–REV7 interaction relies predominantly on Pro-53 and Pro-58 in RBM, using either the immunofluorescence assay or the yeast two-hybrid system. Here, our findings provide the first structural evidence to explain their in vivo results, and they substantiate the RBM in the N-terminal loop of SHLD3 is required for REV7 binding. Meanwhile, the $\alpha$C-helix region of SHLD3–RBD underpins the high-affinity interaction, as a SHLD3 fragment devoid of this part displayed largely compromised binding affinity (Fig. 3c). We then found Ile-60 and Val-68 act as two anchors in the $\alpha$C-helix region of SHLD3, and this binding pattern is important for high-affinity REV7 binding.

Interestingly, we noticed that the double-anchor binding pattern similar to SHLD3 is also presented in two REV3 fragments, whereas CAMP adopts the single-anchor binding model in which an equivalent valine fills the same hydrophobic pocket in REV7 as does Ile-60 of SHLD3 (Fig. 5a). As the contribution of the second hydrophobic anchor in SHLD3–RBD to REV7 binding is significant, it is reasonable to infer that the binding affinity between REV3 and REV7 is comparable with SHLD3, and the interaction between CAMP and REV7 is much weaker due to the lack of corresponding structural elements. Of note, Ran interacts with REV7 via its RBM and exhibits a moderate binding affinity of about 900 nm (30), which parallels the binding affinity between REV7 and SHLD3 N-terminal loop-containing RBM but is much lower than intact SHLD3–RBD. Therefore, the presence of an extra binding part in those proteins strikingly enhances their interaction with REV7. The variance in binding affinity, we surmise, is correlated with different biological functions. For example, during cell-cycle control and mitosis, REV7-binding partners are supposed to interact with REV7 at a given moment, and moderate binding affinity enables timely assembly/disassembly process between them and REV7. In addition, an ample amount of these proteins, such as Ran, makes it less necessary to possess tight binding. However, the abundance of SHLD3 in cells is ultra-low (undetectable in mass spectrometry–based global proteome analyses to a depth of ~10,000 proteins) (19). When DSB arises, the robust binding level ensures the recruitment of SHLD3 to REV7 and thus the proper repair of DNA damages.

Besides its role in DSB repair, the versatile REV7 protein also participates in other cellular pathways, such as TLS and cell-cycle control. The safety-belt region of REV7 plays a central role in various binding partners through an intriguing “open/close” switch, but lack of kinetic information impedes further understanding toward this process. Here, using SPR assays, we found the “safety belt” serves as a contributing factor of the tight-binding nature underlying the SHLD3–REV7 interaction by retarding the dissociation rate (Fig. 4a) because the mutation in safety-belt region impairs interaction mainly via drastically increasing the dissociation process (Fig. 4b). Additionally, we indeed observed the W171A mutant showed SHLD3-binding defects both in SPR assays using purified proteins and Ni-pulldown assays (Figs. 2b and 4b), but the W171A mutant did not have a detectable effect on SHLD3 binding in vivo (Fig. 2f) (15). This scientific disagreement could be explained by the possibility that other regulating factors in the complicated cell environment may modulate the safety-belt region of REV7 and affect its function during DSB repair. The potential regulators and mechanisms will be defined in a future study.

The safety belts are typical structural features among HORMA family proteins, which have structural plasticity and could be tuned between the open and closed state when interacting with different partners. In our structure, SHLD3 RBD is tightly locked by the safety belt of REV7, and this interaction, in turn, stabilizes the closed conformation of this region. But how REV7 dissociates from SHLD3 remains an open question, and other unknown factors or post-translational modifications, we speculate, may be involved in this process, as the safety belt of Mad2, the REV7 paralog, can be remodelled by ATPase TRIP13 to change from a closed conformation to the open conformation, thereby regulating the binding of Mad2 to its partners (33–35). Thus, future efforts are required to identify the regulator of SHLD3–REV7 dissociation and the corresponding molecular mechanism.

Recently, a structure-based screening has identified a potent small molecule that specifically targets the REV1 subunit within the mutagenic TLS complex, thus enhancing cisplatin-induced toxicity in human cell lines and restricting the growth of xenograft human melanomas in mice (36). REV7 is not only a component of the TLS complex but also a pivot in Shieldin complex,
and the SHLD3–REV7 structure exhibited some shallow grooves or cavities on REV7 surface suitable for various binding partners. Those are highly attractive sites for designing therapeutically inhibitors specific for blocking SHLD3–REV7 interaction. Importantly, it will provide feasible treatment strategies for those BRCA1-deficient patients that have developed resistance to the poly(ADP-ribose) polymerase inhibition remedy, as cells harboring defected Shieldin and BRCA1 mutants are more sensitive to ionizing radiation and cisplatin treatment (16, 17). Together, our work elucidated the molecular basis of the SHLD3–REV7 interaction and paved the way for the development of medicines toward cancer treatments.

**Experimental procedures**

**Protein expression and purification**

DNA sequences encoding human REV7(6–211) and SHLD3 fragments varying in length were amplified from human 293T cDNA and cloned into a pET-Duet-1 vector using BamHI/NotI and NdeI/XhoI sites, respectively, which generate the His-tag REV7 and nontag SHLD3 peptide. Mutations of R124A and/or A135D were introduced into REV7 to stabilize the protein (23–25), allowing subsequent crystal screen trials. For co-expression of various His–REV7(6–211)–SHLD3, the plasmids were transformed into *Escherichia coli* BL21-Codon PLUS(DE3)-RIPL cells. When the cells grown at 37 °C reached an OD600 of 0.8, 0.5 mM isopropyl β-D-thiogalactoside (IPTG) was added to induce protein expression at 16 °C for 18 h. Then, the cells were collected and lysed in buffer A (50 mM Hepes, pH 7.4, 500 mM NaCl, 20 mM imidazole). Supernatants separated by centrifugation were applied onto a self-packed Ni-NTA column (Qiagen) equilibrated with buffer A. After washing steps using buffer A and buffer B (20 mM Hepes, pH 7.4, 1.5 M NaCl), the targeted proteins were eluted with buffer C (20 mM Tris-HCl, pH 8.0, 100 mM NaCl, 300 mM imidazole), immediately diluted with buffer D (20 mM Tris-HCl, pH 8.0), and then loaded onto a 5-ml HiTrap Q HP column (GE Healthcare) for further purification. The bound proteins were eluted against a linear gradient from 50 to 800 mM NaCl. The eluate containing REV7–SHLD3 complexes was concentrated using 10,000 MWCO centrifugal filter units (Millipore) and subjected to size-exclusion chromatography (GE Healthcare) equilibrated with buffer E (20 mM Hepes, pH 7.4, 100 mM NaCl) for final purification. The purified protein complexes were fast-frozen in liquid nitrogen and stored at −80 °C until use.

For protein used in ITC assays, full-length REV7 was constructed into a homemade pET28a vector to have a His–MBP fusion tag at the N terminus, which greatly improved the solubility. To effectively avoid degradation and promote purification, SHLD3(29–83) fragment was cloned into another designed pET28a vector to make a recombinant SHLD3 fusion protein with a His-tag at the N terminus and a yeast Smt3 tag at the C terminus. Constructs were expressed in BL21–Codon PLUS(DE3)–RIPL cells, in LB medium induced with 0.4 mM IPTG for 18 h at 16 °C. The proteins were purified similarly as described above. The purified protein samples were concentrated and stored at −80 °C after flash-freezing in liquid nitrogen until use. Mutations were generated using the QuikChange site-directed mutagenesis kit (Stratagene) and were verified by sequencing. Resulting protein mutants were purified in the same way as WT protein.

**Crystallization, data collection, and structure determination**

For crystallization attempts, REV7 complexed with SHLD3 segments of various lengths were used. The crystallization was performed by mixing 1.2-μl protein complexes (20 mg/ml) with a 1.2-μl reservoir solution using the hanging-drop vapor-diffusion method. Nearly 3 days later, crystals appeared at several conditions. After optimization of the reservoir conditions, crystals of good quality come from the REV7(R124A)–SHLD3(46–74) complex in a reservoir solution containing 0.1 M sodium acetate, pH 5.2, 2.1 M ammonium sulfate, and REV7(R124A/A135D)–SHLD3(46–74) complex in a reservoir solution consisted of 0.1 M sodium citrate tribasic, pH 5.0, 1.9 M ammonium sulfate. Then, each crystal was collected within a nylon loop and transferred to liquid nitrogen with cryoprotectant made from the corresponding reservoir solution plus 25% glycerol until use. Diffraction data were collected at the Shanghai Synchrotron Radiation Facility (SSRF) BL17U1 beamline using the Eiger X 16M detector or BL19U1 beamline using the PILATUS 6M detector. All collected data were processed and scaled using the XDS package (37) and AIMLESS (38) within the CCP4 program suite (39). As both crystals displayed anisotropy, the data were further corrected with UCLA diffraction anisotropy server (40), which improved the resulting electron density maps and enabled unambiguous model building.

For REV7(R124A)–SHLD3(46–74) complex, the structure was determined by the molecular replacement method using REVE7–REV3 (PDB code 3ABD) as a template with the program Phaser (41). Manual model building and subsequent refinement were performed in program COOT (42) and program PHENIX (43), respectively. The final structure, refined to $R_{work}/R_{free} = 20.4/24.6\%$, contains REV7 residues 9–207 and SHLD3 residues 49–73. Residues without discernable side-chain density were truncated at the β-carbon. As validated by MolProbity (44), the final model showed good stereochemistry with 94.34% residues in the Ramachandran-favored region, 5.66% residues in Ramachandran-allowed zones, and no outliers. Other statistics during data collection and processing were summarized in Table 1.

For the REV7(R124A/A135D)–SHLD3(46–74) complex, the structure was determined in the same procedure as described for the REV7(R124A)–SHLD3(46–74) complex. The final structure, validated with MolProbity program, showed 96.82% favored, 3.18% allowed, and no outliers in the Ramachandran plot. Details of the structural statistics are shown in Table 1. All structural figures were prepared using the PyMOL program (The PyMOL Molecular Graphics System, Version 2.2.0 Schrödinger, LLC).

**IMAC-based interaction assays**

Mutations in REV7 or SHLD3 were introduced using the QuikChange site-directed mutagenesis kit (Stratagene) into a pET-Duet-1 vector, enabling co-expression of His–REV7 and SHLD3(29–83) peptides. Interaction analysis by IMAC was performed, as described previously (24, 26), with minor differ-
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ences. Briefly, after cell lysis and centrifugation, the soluble fractions were incubated with Ni-NTA resins (Qiagen) at 4 °C for 2 h. The beads were first washed with buffer F (20 mM Hepes, pH 7.4, 1.5 mM NaCl, 20 mM imidazole) three times and then washed using buffer B another three times. Next, the bound proteins were eluted with buffer C and analyzed by SDS-PAGE stained with Coomassie Brilliant Blue. Band intensities were quantified in program Quantity One (Bio-Rad). The quantification results came from three independent experiments, and the statistical significance of differences between samples was determined by an unpaired Student’s t test, using GraphPad Prism 6 (GraphPad Software).

Isothermal titration calorimetry experiments

Purified proteins used here were dialyzed overnight against ITC buffer (20 mM Tris-HCl, pH 7.5, 150 mM NaCl, 2 mM β-mercaptoethanol), and all measurements were done using an ITC200 titration calorimeter (MicroCal) at 20 °C. A pre-drop of 0.4 μL and a subsequent 19 2.0-μL drops of His–SHLD3–Smt3 (300–500 μM) were stepwise-injected into the calorimetric cell containing His–MBP–REV7 (20–50 μM), at intervals of 120 s. The stirring speed remained constant at 750 rpm to thoroughly mix the samples in the cell. In all cases, two independent experiments were performed. As described previously (45), the raw data were analyzed and fitted into a one-site binding model using Origin 7.0 software (Microcal). The relationships between change in enthalpy (ΔH), entropy (ΔS), and binding energy (ΔG) were calculated as ΔG = ΔH − TΔS.

Surface plasmon resonance measurements

Binding kinetics assays were performed on a BIAcore 8K instrument (BIAcore AB, Uppsala, Sweden). Purified SHLD3(29–83) fragment was immobilized on a research-grade CMS chip using EDC–NHS coupling chemistry according to the BIAcore manual, and 2-fold increasing concentrations of purified MBP–REV7 or mutants were injected and analyzed. The SPR running buffer is 20 mM Hepes-Na, pH 7.5, 150 mM NaCl, 0.05% v/v Tween 20, and analyses were recorded at 25 °C, whereas a 20 mM NaOH buffer was used as regeneration buffer. All of the data collected were analyzed in BLAevaluation software (BIAcore). The association (k_on) and dissociation (k_off) rates and the equilibrium dissociation constant (K_D) were obtained using a 1:1 Langmuir binding model.

Co-immunoprecipitation and Western blotting

Cells were lysed with NTEN buffer (20 mM Tris-HCl, pH 8.0, 100 mM NaCl, 1 mM EDTA, 0.5% Nonidet P-40) containing pepstatin A and aprotinin on ice for 30 min. Clear cell lysates were incubated with S protein beads for 2 h at 4 °C. Beads were then washed and boiled in 2× Laemmli buffer, and proteins were separated by SDS-PAGE. Polyvinylidene difluoride membranes were blocked in 5% milk in TBST buffer and then probed with antibodies as indicated.

Data availability

Atomic coordinates and structure factors for the REV7 (R124A)–SHLD3 and REV7(R124A/A135D)–SHLD3 complexes have been deposited to the Protein Data Bank under accession numbers 6K07 and 6K08, respectively.

Author contributions—Y. D. and Z. Z. conceptualization; Y. D., F. Z., and L. W. investigation; Y. D. and Z. Z. writing original draft; S. S., Z. G., and Z. Z. supervision; S. S., Z. G., and Z. Z. funding acquisition; Z. G. and Z. Z. project administration; Z. G. and Z. Z. writing review and editing; Y. D. performed complex assembly, crystallization, structure determination, ITC assays, SPR, and Ni-pulldown assays; F. Z. performed in vivo pulldown assays; L. W. purified protein mutants for ITC assays; S. S. participated in the discussions; Y. D., Z. G., and Z. Z. data analysis.

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