



COMMUNICATION

Molecular Basis for Complement Recognition and Inhibition Determined by Crystallographic Studies of the Staphylococcal Complement Inhibitor (SCIN) Bound to C3c and C3b

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The human complement system plays an essential role in innate and adaptive immunity by marking and eliminating microbial intruders. Activation of complement on foreign surfaces results in proteolytic cleavage of complement component 3 (C3) into the potent opsonin C3b, which triggers a variety of immune responses and participates in a self-amplification loop mediated by a multi-protein assembly known as the C3 convertase. The human pathogen *Staphylococcus aureus* has evolved a sophisticated and potent complement evasion strategy, which is predicated upon an arsenal of potent inhibitory proteins. One of these, the staphylococcal complement inhibitor (SCIN), acts at the level of the C3 convertase (C3bBb) and impairs downstream complement function by trapping the convertase in a stable but inactive state. Previously, we have shown that SCIN binds C3b directly and competitively inhibits binding of human factor H and, to a lesser degree, that of factor B to C3b. Here, we report the co-crystal structures of SCIN bound to C3b and C3c at 7.5 and 3.5 Å limiting resolution, respectively, and show that SCIN binds a critical functional area on C3b. Most significantly, the SCIN binding site sterically occludes the binding sites of both factor H and factor B. Our results give insight into SCIN binding to activated derivatives of C3, explain how SCIN can recognize C3b in the absence of other complement components, and provide a structural basis for the competitive C3b-binding properties of SCIN. In the future, this may suggest templates for the design of novel complement inhibitors based upon the SCIN structure.

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Abbreviations used: C3, complement component 3; SCIN, staphylococcal complement inhibitor; fH, factor H; fI, factor I; RCSB, Research Collaboratory for Structural Bioinformatics; SPR, surface plasmon resonance; Cvf, cobra venom factor.

The complement cascade is a centerpiece of the immune system and functions not only as the first line of response to foreign materials in the body but also in the initiation of inflammation and stimulation of adaptive immunity. This versatility of complement is based on a highly regulated network of soluble or cell-surface-bound proteins that serve as substrates, enzymes, or regulators of a hierarchical

series of proteolytic and protein–protein recognition events. Initiation of the cascade may be triggered by three distinct pattern recognition mechanisms, which have been denoted the classical, lectin, and alternative pathways. Despite the precise nature of the initiating event, all three of these pathways lead to the formation of homologous multi-protein complexes [named complement component 3 (C3) convertases] that enzymatically cleave C3 (184 kDa) into its active fragments C3a (10 kDa) and C3b (175 kDa) (reviewed in Ref. 1). Release of the C3a anaphylatoxin fragment exposes an occluded but reactive thioester moiety in C3b. It is the nucleophilic attack of this group that results in covalent deposition of the C3b on the surface of nearby biomaterial and ultimately triggers its phagocytosis via specific complement receptor-expressing cells.^{2,3}

Structural studies have shown that the conversion from C3 to C3b is accompanied by substantial conformational changes, which themselves expose a number of critical ligand binding sites.^{4–6} A particularly prominent example of this is typified by the alternative pathway, where interaction of the pro-enzyme factor B (fB) with one of these sites enables its cleavage by factor D to generate the active alternative pathway C3 convertase complex (C3bBb). Subsequent binding and cleavage of C3 by the convertase produces additional C3b and thereby propagates a self-amplification loop of complement activation on foreign surfaces.⁷ Unhindered complement activation in this manner leads to the formation of C5 convertases; generation of additional anaphylatoxins (i.e., C3a and C5a); induction of signaling, chemotaxis, and phagocytosis; and assembly of a lytic membrane attack complex. However, the convertase assemblies by themselves are only transiently stable and undergo irreversible dissociation with a half-life of approximately 1 min.⁸ Moreover, on host cells, this dissociation is further catalyzed by a series of “regulators of complement activation” that either destabilize the convertases or mediate the stepwise degradation of the essential C3b substituent (reviewed in Refs. 1 and 9). One pivotal regulator of this activation cycle is factor H (fH), which both accelerates the decay of the convertase complex and acts as a cofactor for the factor I (fI)-mediated proteolysis of C3b that produces the inactive C3c fragment.² Given this tenuous balance that leads to the “double-edged sword” nature of the immune system, many pathological conditions are now known to arise from insufficient regulation or aberrant activation of complement.¹⁰ Therefore, inhibition of complement initiation or amplification has been suggested as an attractive means of protection against these diseases.¹¹

Staphylococcus aureus is a widespread and versatile human pathogen capable of colonizing a remarkable number of different tissues.¹² Diseases caused by *S. aureus* span a broad landscape of relatively minor,

localized skin infections to more life-threatening conditions such as disseminated bacteremia and endocarditis.¹² *S. aureus* cells activate all three complement pathways and stimulate chemotaxis of leukocytes due to the presence of pathogen-associated molecular patterns. However, it has become increasingly clear over the last decade that this pathogen has evolved an array of successful strategies to evade the surveillance and elimination mechanisms that are fundamental to innate immune system function.^{13,14} In this regard, *S. aureus* secretes nearly one dozen factors that bind to and interfere with the various steps needed to activate C3 and to propagate various inflammatory responses within its host.^{14,15} Subsequent structure/function analyses on several of these proteins have demonstrated that they are diverse in their mechanisms of action and, in some cases, has identified entirely novel modes of complement modulation.^{14,15}

The staphylococcal complement inhibitors (SCINs) are a family of small (~9.8 kDa), secreted proteins that adopt an antiparallel three- α -helical bundle fold,¹⁶ variants of which appear to be common among a larger group of staphylococcal immune modulators.^{14–16} The prototypic member of this family, SCIN (also called SCIN-A), is a human-specific complement inhibitor that disrupts bacterial opsonization as well as associated downstream effects of complement activation by blocking function of the solid-phase C3 convertases derived from all three initiation pathways.^{17,18} Previous studies originally suggested that SCIN binds only to fully assembled C3 convertases (e.g., C3bBb);¹⁷ however, this raised significant questions as to the nature of this interaction because all other characterized C3 convertase regulators or inhibitors bind to one or more of the complement components directly. Indeed, while SCIN does not bind native C3, we have recently shown that SCIN binds directly to activated C3b with an apparent nanomolar affinity in near-physiological buffers ($K_d \approx 180$ nM).¹⁹ Functional dissection of the direct interaction between SCIN and C3b has revealed a multi-faceted nature to the SCIN inhibitory mechanism. First, while SCIN impairs the initial rate of C3 convertase assembly, it nevertheless stabilizes the assembled convertase against decay acceleration by fH. Secondly, SCIN reduces the efficiency of fI- and fH-mediated degradation of C3b to iC3b. Finally, SCIN also prevents the proteolytic cleavage, but not the initial binding of the C3 substrate by the convertase.¹⁹

In this manner, SCIN contributes to staphylococcal complement inhibition by trapping the C3 convertase in a stabilized yet inactive state.^{19,20} Taking advantage of these properties, Rooijakkers *et al.* recently described the structure of the SCIN-inhibited alternative pathway C3 convertase.²⁰ While this work gave important insights into the molecular architecture and function of the

Table 1. Diffraction data collection and refinement statistics for the C3c–SCIN and C3b–SCIN complexes

	C3c–SCIN			C3b–SCIN ^a
<i>Data collection</i>				
Beamline	22-BM	22-ID	22-BM	22-ID
Wavelength (Å)	1.00	1.00	1.00	1.00
Space group	<i>P</i> 2 ₁ 2 ₁ 2	<i>P</i> 2 ₁ 2 ₁ 2	<i>P</i> 2 ₁	<i>P</i> 4 ₁ 2 ₁ 2
Cell dimensions				
<i>a</i> , <i>b</i> , <i>c</i> (Å)	231.16, 231.50, 68.89	214.07, 68.43, 114.93	69.11, 217.03, 115.65	128.31, 128.31, 469.76
β (°)			89.99	
Resolution (Å)	50–3.5	50–4.1	50–3.4	50–7.5
Unique reflections	44,393	13,684	45,397	5337
Completeness (%)	93.4 (85.1)	98.4 (99.0)	97.5 (93.1)	97.2 (81.4)
<i>R</i> _{merge} (%) ^b	12.1 (46.6)	18.1 (49.6)	16.0 (50.0)	11.6 (45.3)
Redundancy	7.4 (3.7)	5.4 (5.1)	3.4 (2.8)	10.4 (7.8)
<i>I</i> / σ <i>I</i>	10.5 (2.4)	8.7 (4.1)	5.0 (2.4)	13.2 (3.0)
<i>Refinement statistics</i>				
RCSB accession code	3NSA	3NMS	3L3O	3L5N
Complexes per asymmetric unit	2	1	2	1
<i>R</i> _{work} / <i>R</i> _{free} (%) ^c	26.6/28.8	27.1/28.9	22.3/27.1	26.5/26.8
Mean <i>B</i> -factors (Å ²)	68.6	121.2	108.1	172.3
RMSD				
Bond lengths (Å)	0.012	0.012	0.009	0.008
Bond angles (°)	1.31	1.37	1.20	1.14
Ramachandran plot (%)				
Favored	85.9	86.7	88.9	85.7
Allowed	11.5	11.1	10.4	12.7
Outlier	2.6	2.2	0.8	1.6

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

Methods. Recombinant SCIN was expressed in *Escherichia coli* and purified as previously described.²¹ Purified C3b was prepared by activated-thiol Sepharose (GE Biosciences) separation of a limited trypsin digestion of isolated human C3 according to previously established protocols.²² Purified C3c was obtained from Complement Technology (Tyler, TX, USA). Solutions of each purified monomer were mixed to yield an equimolar complex of C3b–SCIN and C3c–SCIN prior to buffer exchange by ultrafiltration into 10 mM Hepes–NaOH (pH 7.4) to a final concentration of 5 mg mL^{−1} complex. Crystallization was achieved using hanging drop vapor diffusion. For each complex, the precipitant was diluted 1:1 with ddH₂O, and 1 μ L of this solution was mixed with 1 μ L of protein solution. C3b–SCIN co-crystals grew over the course of 21 days at 293 K using 0.1 M Hepes–NaOH (pH 7.0), 30% (v/v) Jeffamine ED-2001–HCl as a precipitant; this precipitant solution was used directly for cryopreservation of crystal samples. C3c–SCIN co-crystals were obtained in 2–5 days at 277 K using 0.1 M Hepes–NaOH (pH 7.5), 10% (w/v) polyethylene glycol 6000, and 5% (v/v) 2-methyl-1,3-propanediol as a precipitant; these crystals were cryoprotected by a brief incubation in an analogous buffer that contained 20% (v/v) glycerol. Following flash-cooling to 93 K, diffraction data were collected using either beamline 22-ID or beamline 22-BM of the Advanced Photon Source (Argonne National Laboratory) and processed using HKL2000.²³ Three different lattice types of C3c–SCIN co-crystals were identified from a single crystallization condition (one monoclinic and two orthorhombic; Table 1). Structures were solved by molecular replacement with Phaser²⁴ using the previously published structures of SCIN (RCSB code 2QFF),¹⁶ C3b (RCSB code 2IO7),⁴ and C3c (RCSB code 2A74)⁵ as search models. The final models were obtained after manual building in Coot²⁵ and refinement in PHENIX,^{26,27} non-crystallographic symmetry restraints were employed where applicable during all steps of solving the C3c–SCIN structures. Models were analyzed and validated using MOLPROBITY.²⁸ All structural representations were prepared using PyMOL (<http://www.pymol.org/>), while quantitative comparisons of superimpositions were determined using STAMP.²⁹

^a Crystallization and data collection on the C3b–SCIN crystals have been published separately.²¹

^b $R_{\text{merge}} = \sum_i \sum_j |I_i(h) - \langle I(h) \rangle| / \sum_i \sum_j I_i(h)$, where $I_i(h)$ is the *i*th measurement of reflection *h* and $\langle I(h) \rangle$ is a weighted mean of all measurements of *h*.

^c $R = \sum_h |F_o(h) - F_c(h)| / \sum_h |F_o(h)|$. *R*_{cryst} and *R*_{free} were calculated from the working and test reflection sets, respectively. The test set constituted 5% of the total reflections not used in refinement.

alternative pathway C3 convertase, it provided no conclusive structural information regarding the direct interaction between C3b and SCIN. Because of this, the precise nature of the SCIN binding site on C3b and C3c, its relationship to native C3, and the structural basis for several important effects of SCIN on C3 convertase assembly and dynamics remained incompletely described. Here, we report the crystal structures of SCIN bound to the C3 derivatives C3b and C3c. Analysis of these structures provides a molecular framework to understand SCIN inhibi-

tion of C3b binding to fH and fB and demonstrates the structural basis for specific recognition of activated C3 fragments by SCIN.

Structures of SCIN bound to human complement fragments C3b and C3c

To gain further insight into molecular mechanisms of SCIN activity, we pursued crystal structures of SCIN bound to complement fragments C3b and C3c.

Binary complexes of SCIN bound to C3b and C3c were reconstituted by incubation of purified components followed by concentration by ultrafiltration.²¹ The presence of each protein component in the resulting samples was verified by both native and SDS-polyacrylamide gel electrophoresis²¹ prior to

crystallization of both complexes by vapor diffusion of hanging drops. Three distinct C3c–SCIN crystal forms that diffracted X-rays to between 3.4 and 4.1 Å limiting resolution were identified, while crystals of C3b–SCIN provided reasonably complete data to 7.5 Å resolution (Table 1). Initial phases were

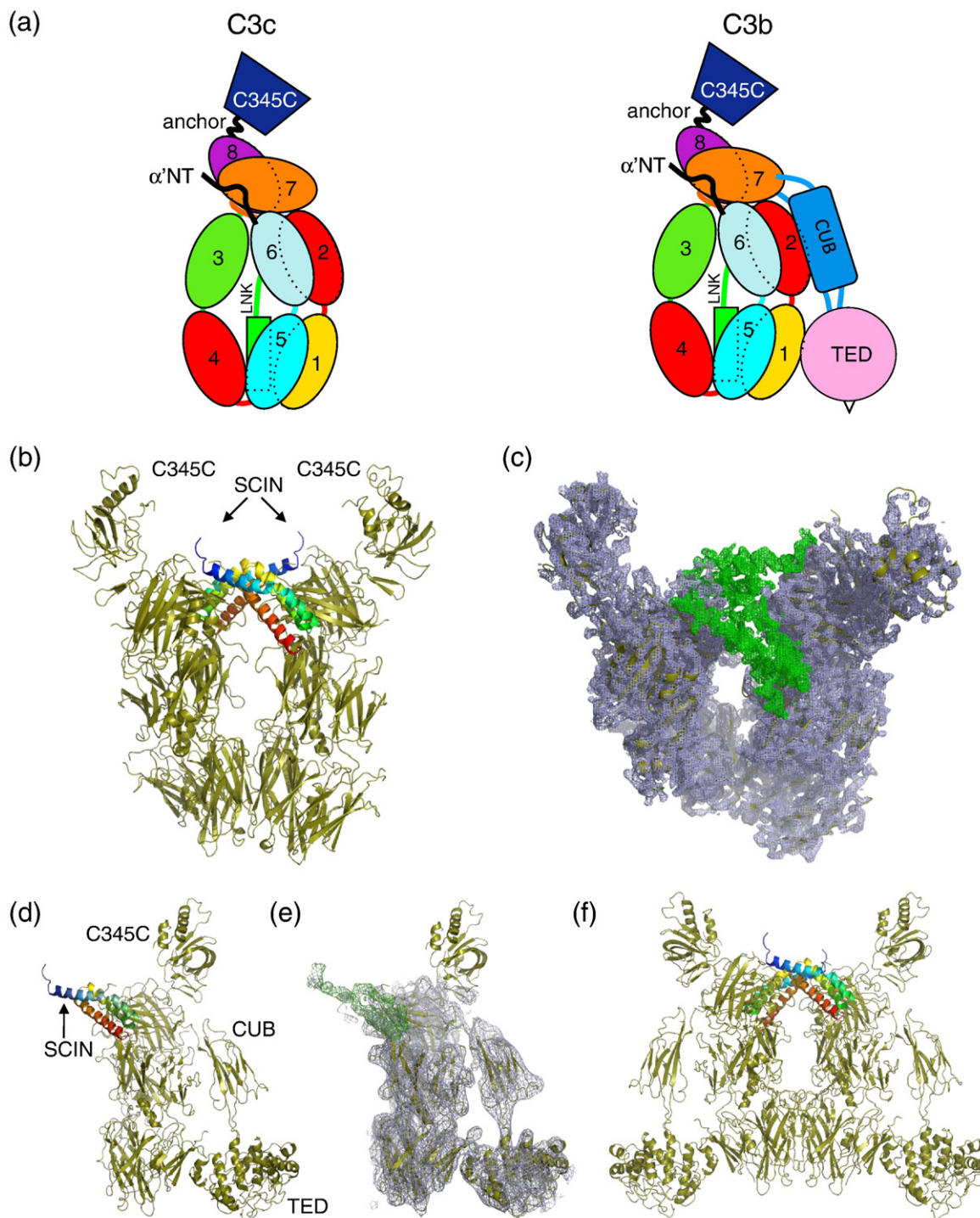


Fig. 1 (legend on next page)

obtained by molecular replacement, and models for each crystal form were iteratively improved and refined to the values reported in Table 1.

The C3c–SCIN structure is composed of a heterotetramer, where two copies of the C3c–SCIN complex are aligned with, but rotated 180° about the long axis of the complement component (Fig. 1b and c). This arrangement is observed in all three C3c–SCIN structures, even though the heterotetramer is divided between neighboring asymmetric units in the second orthorhombic crystal form (Supplemental Fig. 1, Table 1, and Supplemental Table 1). Similarly, though the asymmetric unit of the C3b–SCIN structure is defined by a single copy of each protein arranged as a heterodimer (Fig. 1d and e), inspection of nearby symmetry-related complexes likewise indicates that a similar mode of tetramerization to that of C3c–SCIN is present within the C3b–SCIN crystal lattice (Fig. 1f). In fact, superposition of the symmetry-related C3b–SCIN tetramer with that of C3c–SCIN indicates that these two structures share a high level of identity to one another.

With the exception of the C345C domains, readily interpretable density is observed in $2F_o - F_c$ electron density maps (contoured at 1.5 σ) for all functional domains of C3c and C3b present in the final models of all structures (Fig. 1c and e). In all C3c–SCIN structures, the model *versus* map correlation is somewhat weaker for both copies of the C345C domain; however, these areas are clearest for orthorhombic form 1, and thus, this crystal is discussed in the remainder of the text unless otherwise indicated. In the case of C3b–SCIN, little contiguous density is observed in this region; therefore, in the final model, this domain has zero occupancy (residues 1497–1641). Conformational flexibility of C345C has been reported previously, as this domain adopts alternative conformations relative to the remainder of the molecule in various crystal structures of C3 and its activation and degradation fragments²⁰ (Supplemental Fig. 1f). This is perhaps not surprising since this domain projects away from the “key-ring”-like core of C3c, and it does not make stable contacts with SCIN in either of the structures reported here or in that of the SCIN-inhibited C3 convertase [Research Collaboratory for Structural Bioinformatics (RCSB) code:

2WIN].²⁰ Most significantly, however, SCIN occupies the same binding site and orientation relative to its complement target in both structures, and calculation of $F_o - F_c$ difference maps where the SCIN chains were omitted from either model reveals contiguous regions of positive density (contoured at 2.0 σ) that correspond to the location of the SCIN polypeptide in each complex (Fig. 1c and e). Together, this strongly suggests that the C3c–SCIN and C3b–SCIN structures are directly comparable to one another.

A series of crystallographic studies have demonstrated that significant structural changes accompany activation of native C3 into C3b,^{5,6,30} and interactions of C3b with certain ligands, in particular the distinct staphylococcal complement inhibitor proteins Efb³¹ and Ehp,³² have been shown to induce additional conformational changes in the complement component. We therefore used superimposition analysis to examine whether additional conformational changes accompany SCIN binding in the structures described above. Quantitative comparisons of each SCIN-bound C3 fragment with the structure of its unbound counterpart^{4,5} reveal no obvious changes to the complement components upon SCIN binding, however; without including the poorly ordered C345C domains in the calculations, the main-chain atoms’ root-mean-square deviations (RMSDs) in the SCIN-bound to unbound structures are 1.80 and 2.60 Å for the C3c⁵ and C3b⁴ complexes, respectively. Separately, a recent crystal structure of a SCIN-inhibited form of the alternative pathway convertase (C3bBb–SCIN) has also been reported.²⁰ Although SCIN is indeed bound to C3b in this structure, the additional presence of the large Bb fragment (~57 kDa) raised the possibility that structural rearrangements of the C3b–SCIN interaction may be necessary to form the resulting ternary complex. Because of this possibility, no direct conclusions regarding the nature of C3b–SCIN binding from this previous work could be drawn. We again used superimposition analysis to address these questions (Supplemental Fig. 2). By and large, all three SCIN-bound structures superimpose well with one another; the main-chain atoms’ RMSD in the SCIN–C3c and SCIN–C3b binary complexes when compared to those of C3bBb–SCIN are 0.65 and 0.80 Å, respectively.

Fig. 1. Crystal structures of complement fragments C3c and C3b bound to SCIN. (a) Schematic representation of the domains that comprise C3c and C3b. (b) Crystal structure of C3c–SCIN (drawn from RCSB code 3NSA), where two copies of SCIN (blue to red rainbow from amino to carboxy terminus) and two copies of C3c (sand) are found in the asymmetric unit. (c) Electron density map of the same structure generated by refinement of models not containing SCIN molecules. $2F_o - F_c$ density (blue cage; 1.5 σ contour) fits two copies of C3c, while the $F_o - F_c$ map (green cage; 2.0 σ contour) shows clear positive density corresponding to the location of the SCIN polypeptide. Note that the viewing plane of this panel is inclined slightly relative to (b) for clarity. (d) Crystal structure of C3b–SCIN, where a single copy of both SCIN (blue to red rainbow from amino to carboxy terminus) and C3b (sand) are present in the asymmetric unit. (e) Electron density map of the same structure generated by refinement of the model without the SCIN polypeptide. Map parameters are identical with those in (c). (f) Generation of the symmetry mate ($x, y, -z$) from the C3b–SCIN crystal indicates a mode of tetramerization similar to that seen in all C3c–SCIN structures (please see Supplemental Fig. 1).

Differences are visible between the structures in terms of the relative orientations of the C345C domain (which was not modeled in the C3b–SCIN structure), although this could be due to packing effects in the C3c–SCIN crystal lattices. Still, it is worth noting that this domain appears to serve as a swinging platform, which binds and orients Bb in a manner so that its proteolytic active site extends away from the remainder of the C3b molecule.²⁰ Because the location of the SCIN protein is similar in both the binary and ternary complexes, it seems that SCIN binding is accommodated, even in an assembled and active convertase, through minor reorientation of this C345C domain; indeed, real-time measurements using surface plasmon resonance (SPR) have shown that SCIN binds readily to C3bBb.¹⁹ When these data are considered along with recent functional analyses,¹⁹ the structures presented here provide firm evidence that SCIN is capable of binding C3c and C3b in the absence of any additional complement proteins or cofactors.

Nature of SCIN binding to activated forms of C3

The structures reported here and elsewhere reveal that each molecule of SCIN contacts both copies of the complement protein found in the rotationally symmetric tetramers presented in Fig. 1. An examination of available data strongly suggests that each of these SCIN binding sites on C3b is relevant: both isothermal titration calorimetry and SPR analyses have shown that the C3b–SCIN interaction is most accurately described by a two-site binding model,¹⁹ and small-angle X-ray scattering studies on the C3b–SCIN complex similarly indicate a mixture between 1:1 and 2:2 species in solution.¹⁹ To better understand the physical basis for each SCIN binding site, we used the Protein Interfaces, Surfaces, and Assemblies (PISA)³³ server to analyze the intermolecular contacts found in the C3c–SCIN and C3b–SCIN structures.

The first SCIN “footprint” on the C3b surface masks approximately 900 Å², where 700 and 200 Å² are derived from the α and β chains of the complement component, respectively (Fig. 2a and Supplemental Fig. 3a). Closer inspection of this interface (as judged by the C3c–SCIN structures that provide higher resolution) indicates that it arises from contacts made almost entirely by the second α -helix of SCIN. While 22 SCIN residues participate in complex formation in this manner, 11 of these form interactions that are polar in nature (e.g., hydrogen bonds and/or salt bridges). Conversely, the second SCIN “footprint” arises through interaction of the first and third α -helices of SCIN with the symmetry-related complement protein (Fig. 2a and Supplemental Fig. 3b). Interface analysis reveals that this additional site buries approximately 750 Å² and that all 20 residues that comprise it are located on C3b α -

chain. While the 25 total SCIN residues that form this interaction are derived from both the first and third helices of the bacterial protein, the physical nature of the contacts formed is very much similar to that of the first site described above (Supplemental Fig. 3b). In particular, the eight polar interactions found here are similarly distributed between both side chain-to-side chain and side chain-to-main chain contacts. In general, the proportion of polar contacts found in the SCIN interfaces are somewhat higher than that of the related staphylococcal complement inhibitor protein, Efb, which binds to the C3d/TED domain of C3/C3b via 23 total residues that form a total of eight polar interactions.³¹ Such a difference may partly explain the lower apparent affinity of SCIN for its C3b target relative to Efb ($K_d \approx 180$ nM¹⁹ versus 10 nM³¹ in near-physiological buffers, as judged by SPR).

Considerations of SCIN binding aside, activation of native C3 to C3b is accompanied by significant conformational changes that alter the large set of interactions that give rise to the nearly 5000-Å² interface between the α and β chains of C3.⁶ One consequence of such changes is that many residues previously buried in the C3 α/β interface become solvent accessible in C3b. This raised questions as to the nature of the SCIN binding sites in native C3 relative to its activated counterpart, C3b. Significantly, whereas 15 of the 23 C3b-derived residues that constitute the first C3b–SCIN interaction described above are buried in the large α/β interface of native C3, all of the C3b residues that comprise the second SCIN binding site are solvent exposed in the structure of the native complement protein (Fig. 2).

At first glance, this appears to suggest that the second site is actually the primary site of C3b–SCIN interaction. However, three independent lines of evidence reveal that this is not the case. First, unlike the related staphylococcal complement inhibitors Efb, Ehp, and Sbi,^{14,15} SCIN itself cannot bind to native C3.^{17,19} The fact that the second binding pocket is exposed in native C3 but that the first is assembled only upon activation to C3b strongly suggests that this neoantigen is required for initial C3b–SCIN binding. Second, a previous study using chimeric proteins of SCIN and its non-functional homolog, ORF-D, has shown that residues 31–48 of SCIN are critical for complement inhibition.¹⁶ The structures presented here demonstrate that this region of SCIN, in particular the side chain of R42, forms several contacts with the first binding pocket on C3b (Supplemental Fig. 3). Finally, recent data strongly suggest that SCIN blocks C3 cleavage (but not substrate binding) by surface-bound C3 convertases in their monomeric form.¹⁹ Because complement inhibition requires the SCIN region that binds to the first site on C3b, it follows that this interaction is responsible for the convertase inhibitory activity of SCIN.

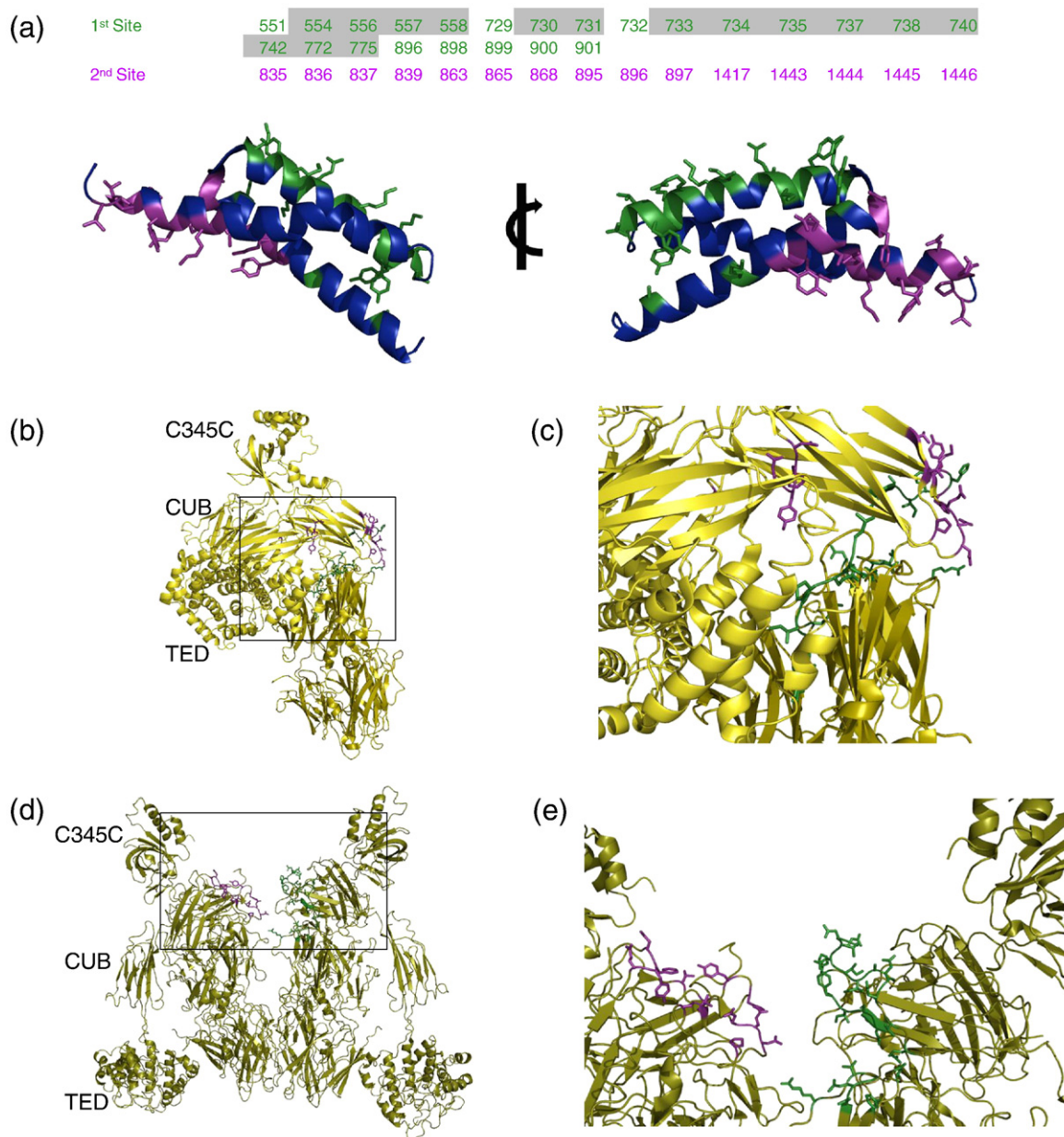


Fig. 2. Molecular nature of SCIN binding sites to C3c and C3b. The EBI PISA server was used to identify and analyze intermolecular contacts within the C3b–SCIN and C3c–SCIN structures. (a) In the top panel, residues from the complement components are classified as belonging to either the first (green) or the second (purple) SCIN binding site described in the text; the residues shown here are greater than 5% buried in the structure of C3b–SCIN relative to unbound C3b. Numbering is based upon the sequence of human pre-pro-C3, and those that are buried in the structure of native C3 are shaded gray. Two rotated views of the SCIN protein are provided in the bottom panel. The side chains from residues comprising the first site (green) and the second site (purple) are represented in ball-and-stick convention. (b) Location of residues comprising the first (green) and second (purple) SCIN binding sites in native C3. (c) Magnified view of the boxed region from (b). (d) Location of residues comprising the first (green) and second (purple) SCIN binding sites within a C3b dimer generated by crystal symmetry. (e) Magnified view of the boxed region from (d).

The observations above suggest that the secondary SCIN binding site on C3b is involved in tetramerization of this complex rather than convertase inhibition *per se*. However, aside from its potent anti-complement properties, SCIN has also

been shown to function as an effective inhibitor of phagocytosis.^{16–18} A mechanism for this activity has recently been proposed, which requires the formation of inhibited convertase pseudo-dimers (C3bBb–SCIN)₂ to block C3b binding by the complement

receptors, CR1 and CR1g.³⁴ Although no crystal structure is currently available for C3b–CR1, examination of the C3b–CR1g complex³⁵ reveals that the quaternary structure of C3b dimers apparently masks the CR1g binding site.³⁴ In this manner, C3b binding by complement receptors is disrupted, and phagocytosis of C3b-opsonized bacteria becomes less efficient. While this clearly suggests that the second C3b binding site of SCIN plays an important role in blocking phagocytosis of C3b-opsonized bacteria, it is worth noting that these (C3b–SCIN)₂ tetramers form in the absence of contributions by fB

(Fig. 1 and Supplemental Fig. 1).¹⁹ Although this hypothesis may form the basis for future work, whether or not these (C3b–SCIN)₂ structures alone are sufficient to inhibit phagocytosis has not been established at this time.

Structural basis for SCIN effects on C3b binding by fH and fB

Precise regulation of complement activation is critical to maintaining the delicate balance between the ability to opsonize foreign cells without directing

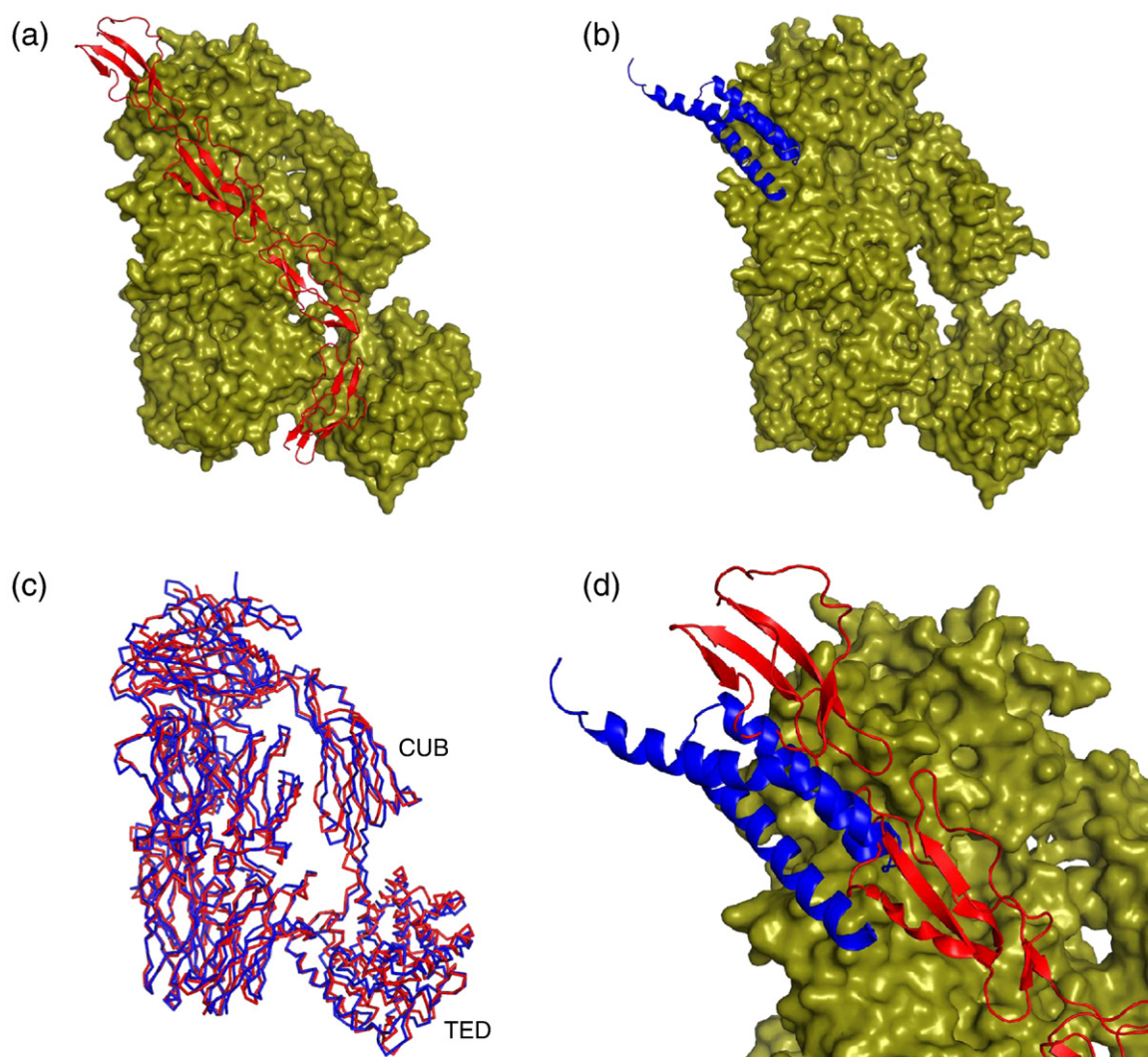


Fig. 3. SCIN masks the C3b binding site for fH. Refined coordinates for the C3b–fH(1–4) (RCSB code 2WII)³⁶ and the C3b–SCIN structure were superimposed by least-squares methods. (a) Representation of the C3b–fH(1–4) structure, with C3b drawn as a molecular surface and fH(1–4) drawn as a red ribbon. The N terminus of fH(1–4) lies toward the top of this panel as drawn. (b) Representation of the C3b–SCIN structure, with C3b drawn as a molecular surface and SCIN drawn as a blue ribbon. (c) Superposition of the C^α positions from the C3b component of (a) (red) and (b) (blue) demonstrates a high degree of structural identity. Note that the C345C domain has been omitted as this region is not readily interpretable in the C3b–SCIN structure. (d) Magnified view of a merged superposition from (a) and (b) illustrates the steric clash between SCIN and the first two domains of fH. The molecular surface shown here is drawn from the C3b–fH(1–4) structure.

complement attack against healthy host cells and tissues. fH acts as an important host regulator of complement activity by both accelerating the decay of convertases and promoting the fI-mediated degradation of C3b to iC3b. The fact that SCIN binds to the C3b substituent of the C3 convertase and renders this otherwise labile multi-component assembly insensitive to both the decay acceleration by fH and impairs its cofactor activity in the degradation of C3b¹⁹ suggested that the SCIN binding site might overlap, at least partly, with a region that is crucial for fH function. Indeed, SPR studies revealed that SCIN can efficiently compete with fH for binding to C3b.¹⁹

To examine the physical basis for the competition between fH and SCIN for C3b binding, we superimposed the C3b–SCIN structure with that of C3b bound to an active fragment of fH [fH(1–4)] (Fig. 3). Aside from the C345C domains (which were omitted from superposition due to poor map/model quality in this area), the remainder of the molecule (including the flexible CUB and TED domains) adopts a nearly identical conformation in each structure (Fig. 3a–c). Interestingly, inspection of the superposed C3b-bound structures of SCIN and fH(1–4) reveals a clear steric clash between these two proteins (Fig. 3d). In particular, binding of the second α -helix of SCIN prevents the extensive contacts made by the first domain of fH(1–4) with the MG7 domain of C3b and, to a lesser extent, those made between the second domain of fH(1–4) and the MG6 β domain of C3b. Even though SCIN is not in position to impede the interactions between the third and fourth domains of fH(1–4) and C3b, the apparently large discrepancy in K_d in near-physiological buffers [180 nM for C3b–SCIN¹⁹ versus 11 μ M for C3b–fH(1–4)³⁶] between these two ligands strongly suggests that SCIN would bind preferentially over fH *in situ*. Thus, it appears that inhibition of fH activity could block both generation of additional C3b by preventing reassembly of functional convertases on that particular C3b molecule and prevent any entrapped C3b from participating in downstream inflammatory events.¹⁹

Though the major effect of SCIN may be attributed to the non-functional trapping of assembled convertases, recent data have also shown that this protein also impedes the rate of C3 convertase formation on surfaces.¹⁹ This feature most likely arises from the direct competition for C3b binding between SCIN and fB, which binds to C3b and is converted to C3bBb through the action of factor D.¹⁹ In the absence of the stabilizing properties of SCIN,^{19,20,37} the C3bB pro-convertase is a kinetically labile complex that dissociates rather rapidly, which has long precluded high-resolution structural studies of this interaction. Cobra venom factor (CVF) is a strong complement activator found in certain cobra species and a structural homolog of C3c that forms

highly stable complexes with fB, which are more amenable to structural analysis.³⁸ To examine the physical basis for competition between SCIN and fB, we superimposed the CVF–fB structure³⁸ with that of C3c–SCIN (Fig. 4a–c). This shows that while much of the fB interface is not perturbed, the loop that connects the first and second α -helices of SCIN would again create a steric clash with fB (Fig. 4d). In particular, the Ba segment of fB appears to be affected to the greatest extent by SCIN binding. This is significant because it is the Ba segment that is responsible for the initial fast and metal-independent loading interaction of fB onto C3b.²⁰ Thus, it seems that by steric impedance of the Ba segment, SCIN can slow the rate of fB loading onto the C3b, which, in turn, lowers the rate of C3 convertase formation.

Conclusions

Throughout their ages-long struggle against formidable host defenses, many pathogens have evolved elegant and powerful means to inhibit, disrupt, or evade the numerous activities required for proper immune function. As a central component of immunity, the complement system has been repeatedly targeted by diverse microbial evasion mechanisms.¹⁴ Detailed analyses on a growing number of these proteins have provided a unique perspective through which the structure/function relationships of central complement proteins can be better understood. In several cases, this work has shown that quite unexpected features can play a critical role in the evasion proteins' mechanism of action. For example, the secreted *S. aureus* proteins Efb and Ehp both bind to C3b and surprisingly induce a conformational change that impairs the activity of the alternative pathway C3 and C5 convertases.^{18,31,32} While SCIN also binds directly to C3b and blocks the activity of surface-assembled convertases,¹⁹ its mechanism of action seems to present a fascinating paradox. Indeed, kinetic stabilization of the ordinarily labile C3 convertase assembly^{17,19} would not at first appear to be advantageous to the pathogen. This strategy actually makes a great deal of sense, however: an assembled but non-functional convertase cannot effectively activate C3 into C3b, which both abrogates the self-amplifying nature of the complement response^{19,39} and prevents efficient generation of downstream pro-inflammatory mediators (i.e., C3a and C5a).¹⁹ Furthermore, by competing with fH and inhibiting its decay acceleration activity,¹⁹ SCIN ensures that the assembled convertases stay locked in their non-functional state. Though many potent regulators of complement activity are known,² none of them appear to involve a mechanism that is conceptually similar to SCIN. This further emphasizes the value of elucidating its functional

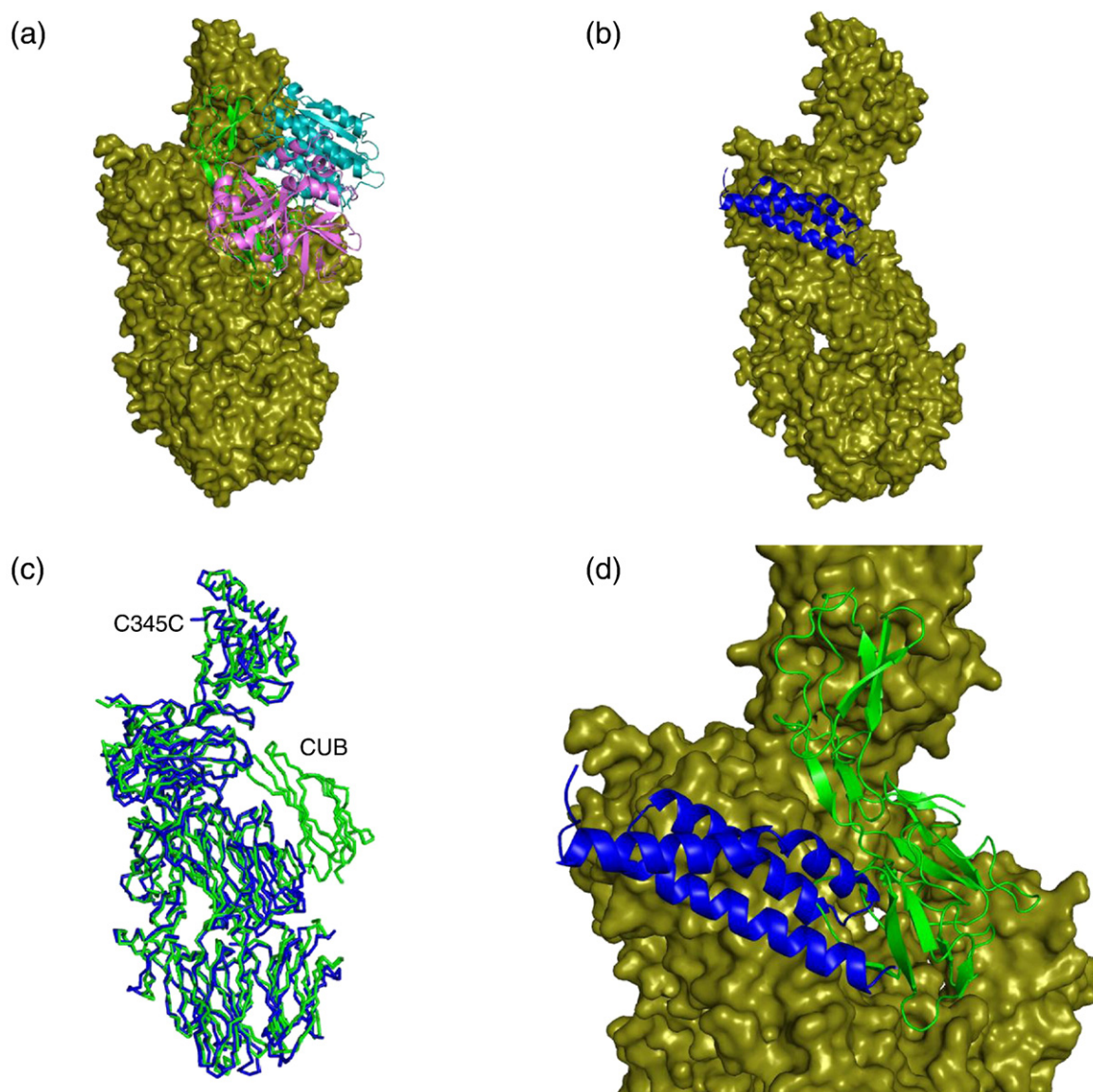


Fig. 4. SCIN partially occludes the C3b binding site for fB. Refined coordinates for the CVF-fB (RCSB code 3HRZ)³⁸ and the C3c-SCIN structure (drawn from RCSB code 3NMS) were superimposed by least-squares methods. (a) Representation of the CVF-fB structure, with CVF drawn as a molecular surface and fB drawn as ribbon. Domains that comprise the Ba (green) and Bb segments (cyan and purple) are colored separately. (b) Representation of the C3c-SCIN structure, with C3c drawn as a molecular surface and SCIN drawn as a blue ribbon. (c) Superposition of the C α positions from the C3c-like component of (a) (green) and (b) (blue) demonstrates a high degree of structural identity. Note that C3c lacks the CUB domain found in CVF. (d) Magnified view of a merged superposition from (a) and (b) illustrates the steric clash between SCIN and the Ba region of fB. The Bb domains have been removed from this panel for clarity. The molecular surface shown here is drawn from the CVF-fB structure.

properties, as well as understanding the precise nature of how SCIN recognizes various complement components at a structural level.

One peculiar feature of microbial immune modulator proteins is that they are often capable of targeting multiple aspects of host defense and repair simultaneously.⁴⁰ On one hand, unique activities can be localized within discrete functional regions of

the same protein. For example, *S. aureus* Efb binds to fibrinogen via its N-terminal region,⁴¹ whereas its C-terminal domain provides its anti-complement activity.^{31,42} On the other hand, a single functional region of a protein can also disrupt independent processes within the host. In this manner, the helical bundle domains of the *S. aureus* complement inhibitors Efb and Ehp not only block the activity

of C3 convertases^{18,31,32} but also block C3d recognition by CR2, which interferes with downstream activation of the adaptive immune response.⁴³ By targeting the activity of complement convertases and phagocytosis simultaneously, SCIN blocks not only complement amplification and inflammation but also clearance of opsonized bacterial cells. The structures presented here reveal that these distinct activities localize to separate regions of SCIN, yet together occupy nearly 1/3 of available surface area of this bacterial protein. The fact that such potent but uniquely valuable activities coexist in a protein of this size further highlights the exquisitely tuned nature of the *S. aureus* immune evasion arsenal.

Insufficiently controlled activation of complement is a major contributor in the pathophysiology of many inflammatory, immune, ischemia/reperfusion, and age-related disorders.¹⁰ In all cases, these events culminate in directing the destructive effects of complement-mediated immune responses against host cells and structures and thereby result in severe tissue damage. Complement inhibitors have long been recognized as a promising therapeutic approach to the prevention of these effects.^{11,44} Surprisingly, bacteria have provided a considerable source of insight into the nature of complement regulation and inhibition. Through its secretion of a collection of small proteins, *S. aureus* deploys ingenious mechanisms to interfere with the central components of the complement cascade, in particular the pivotal conversion of C3 into C3b. Development of complement-directed therapeutics may therefore be greatly aided by the study of the structure, function, and mechanisms of pathogen-derived proteins. In this regard, analysis of the SCIN protein and the physical basis for its interactions with C3b and C3c presented here has provided clues into the recognition of the activated C3b relative its native C3 precursor. In the future, this may suggest templates for the design of novel complement inhibitors that mimic or expand upon SCIN activities toward therapeutic applications.

Accession codes

Refined coordinates and structure factors have been deposited in the RCSB under the accession codes 3NSA, 3NMS, and 3L3O for C3c–SCIN and 3L5N for C3b–SCIN.

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Supplementary Data

Supplementary data associated with this article can be found, in the online version, at [doi:10.1016/j.jmb.2010.07.029](https://doi.org/10.1016/j.jmb.2010.07.029)

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