



## The solution structure of the kallikrein-related peptidases inhibitor SPINK6



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### ABSTRACT

Kallikrein-related peptidases (KLKs) are crucial for epidermal barrier function and are involved in the proteolytic regulation of the desquamation process. Elevated KLK levels were reported in atopic dermatitis. In skin, the proteolytic activity of KLKs is regulated by specific inhibitors of the serine protease inhibitor of Kazal-type (SPINK) family. SPINK6 was shown to be expressed in human *stratum corneum* and is able to inhibit several KLKs such as KLK4, -5, -12, -13 and -14. In order to understand the structural traits of the specific inhibition we solved the structure of SPINK6 in solution by NMR-spectroscopy and studied its interaction with KLKs. Thereby, beside the conserved binding mode, we identified an alternate binding mode which has so far not been observed for SPINK inhibitors.

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### 1. Introduction

Proteases are responsible for the maintenance of the epidermal barrier by proteolytic cleavage of filaggrin and other desmosomal proteins. Among these proteases, the kallikrein-related peptidases (KLKs) are important in the proteolytic regulation of the desquamation process. Members of this family are involved in many of these processes and their insufficient regulation can lead to severe diseases [1].

Kallikrein-related proteases (KLKs) belong to the large class of tryptic and chymotryptic serine proteases. In humans, the KLK family encompasses 15 genes encoded within the single chromosomal locus 19q13.3–13.4, whereas in mice and rat the KLK family consists of 24 and 20 genes, respectively [2]. At least eight KLKs are expressed in human skin [3]. KLK4 has been detected at the mRNA level in human skin, and at the protein level in healthy adult and

fetal skin samples [4–6]. In skin, the enzymatic activity of KLKs is mainly regulated by specific inhibitors of the serine protease inhibitor of Kazal-type (SPINK) family [7]. Spink6, with a single Kazal-type domain, is well conserved among homologs from frogs to humans, suggesting an important role in cellular function [8]. KLK4 was described in human skin, but its functional role in skin biology is poorly understood [5,9]. In the *stratum corneum* of the human skin SPINK6 is cross-linked inter alia to fibronectin by transglutaminases [10,11]. SPINK6 inhibits members of the KLK family with different efficacies. These can be grouped in high (KLK5, 12, 13 and 14), medium (KLK4) and low affinity (KLK6 and 7) inhibition [12]. The medium inhibition of KLK4 activity by SPINK6 is reflected in a  $K_i$  value of 27 nM [12]. In order to elucidate the molecular basis of the KLK4 inhibition by SPINK6 we solved the structure of SPINK6 by NMR-spectroscopy and generated a model of its complex with KLK4 in order to identify the interaction epitope.

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## 2. Materials and methods

### 2.1. Protein expression and purification

SPINK6 was expressed and purified as described before [10].

### 2.2. NMR spectroscopy

For structure determination by NMR-spectroscopy SPINK6 was labeled with  $^{15}\text{N}$ - and  $^{13}\text{C}$ -isotopes by expression in minimal media supplemented with 2 g/l  $^{15}\text{C}$ -glucose and 1 g/l  $^{15}\text{N}$ -ammonium sulfate, and purified as described for the non-labeled material. All spectra were processed with NMRPipe [13] and analyzed with NMRViewJ [14]. NMR measurements were performed on a Bruker Avance 600-MHz spectrometer equipped with a z-gradient triple-resonance cryoprobe. Sequence-specific backbone resonance assignments of SPINK6 were established using the following spectra:  $^1\text{H}$ - $^{15}\text{N}$ -heteronuclear-HSQC, three-dimensional, (H)C(CO)NH, H(CO)NH, HN(CO)CACB, HNCACB,  $^{15}\text{N}$ -edited TOCSY, and  $^{13}\text{C}$ -edited HCCH-TOCSY. For verification of flexible regions ( $^1\text{H}$ - $^{15}\text{N}$  heteronuclear nuclear Overhauser effect (NOE) spectra were measured additionally. Two spectra with on and off resonance  $^1\text{H}$  saturation were recorded in an interleaved manner, and separated before processing with the Bruker AU-program split. The combined recycle and saturation delay was set to 3.2 s. Only well resolved peaks were included in the intensity (peak height) measurements and NOE ratio determination. Distance restraints were obtained from three-dimensional  $^{15}\text{N}$ -edited and  $^{13}\text{C}$ -edited NOE spectra (mixing times, 150 ms).

The spectra were acquired at 303 K and referenced to the water resonance at 4.79 ppm. Chemical shifts and constraints were deposited in the BMRB, accession number: 25691.

### 2.3. Structure calculation

Structure calculations were performed using the program CYANA [15]. The structure calculation was based on 450 interproton distances derived from  $^{15}\text{N}$  and  $^{13}\text{C}$ -edited three-dimensional NOESY experiments. Distances restraints were calibrated using an  $r^6$  function. Three disulfide bonds were defined as 9 distance restraints ranges as follows:  $2.0 \leq d(S_i^\gamma, S_j^\gamma) \leq 2.1 \text{ \AA}$ ;  $3.0 \leq d(C_i^\beta, S_j^\gamma) \leq 3.1 \text{ \AA}$ ;  $3.0 \leq d(S_i^\gamma, C_j^\beta) \leq 3.1 \text{ \AA}$ . TALOS+ [16,17] was used for deriving backbone torsion angles from the NMR chemical shifts of SPINK6 and included in the structure calculations. 100 structures were calculated and subsequently refined in explicit solvent with the CNS program using the RECOORD protocol and parameters [18]. The 20 energetically best structures with an NOE violation energy below a cutoff were selected as the final refined structural ensemble and were deposited (protein data bank accession code: 2n52). All molecular graphical representations were generated using the program RIBBONS [19].

### 2.4. Model building

The region of SPINK6 that is in contact with the protease, KLK4, is very flexible in solution, thereby imposing challenges upon the modeling of the SPINK6-KLK4 complex. To tackle this challenge, we exploited a template-restrained docking protocol, which has been shown to systematically improve superimposed models (Xue, Rodrigues, Dobbs, Honavar, & Bonvin, submitted). This protocol is based on the flexible protein-protein docking method, HADDOCK [20,21] which can incorporate user-provided interface information, either experimental or predicted. Using interface restraints, HADDOCK is able to perform a more detailed flexible refinement of the interface and generate near-native conformations even for

cases with medium to large conformational changes [22]. To identify available templates for SPINK6-KLK4 in the PDB (Protein Data Bank, [www.wwpdb.org](http://www.wwpdb.org)) [23], we used PS-HomPPI [24]. Given the sequences of two interacting query proteins, the newest version of PS-HomPPI – v1.3 searches homologous interacting proteins in the PDB, classifies the templates into Safe Zone (high level of interface conservation), Twilight Zone (medium level) or Dark Zone (low level), clusters the templates based on the structural similarity and returns the inferred clustered interfaces from the templates in the best available zone. For SPINK6-KLK4, PS-HomPPI v1.3 identified 8 templates (Table 2), all of them in the Twilight Zone.

Reliable templates for SPINK6-KLK4 could be identified (with reasonably high sequence identity ranging from 32% to 56%). Superimposed models for SPINK6-KLK4 using the top template identified by PS-HomPPI v1.3: an x-ray crystallography solved complex between a human leukocyte elastase and OMTKY3 (the third domain of turkey ovomucoid inhibitor), PDB ID: 1ppf. These were then refined skipping the initial rigid-body docking stage (it0) and performing only the semi-flexible refinement (it1 and water) of HADDOCK v2.2 with default parameters. The following unambiguous restraints: were used in the refinement: a) CA–CA (C alpha – C alpha) distances calculated by PS-HomPPI v1.3 from the 8 templates in Table 2, b) H-bonds from the template 1ppf, c) dihedral angles extracted from 1ppf to restrain the conformation of the inhibitory loop to the bound form, d) the original NMR NOE restraints, and e) distance restraints extracted from the KLK4 structure to keep the catalytic triad (Asp-His-Ser) in KLK4 in place, whose position is known to be highly conserved. The N-terminus tail (residue 1–22) of SPINK6 was treated as fully flexible during the it1 and water refinement stages. From the 200 calculated SPINK6-KLK4 models we chose top 10 based on HADDOCK scores. These models were further subjected to a refinement round in explicit solvent, without any restraints (using the refinement interface of the HADDOCK server) to remove any strain introduced by the restraints.

## 3. Results

### 3.1. Structure of SPINK6

The  $^1\text{H}$ - $^{15}\text{N}$ -HSQC spectrum and the sequential assignment of SPINK6 is depicted in Fig. 1A showing good dispersion of the resonances. All residues could be unambiguously assigned. 450 distance constraints were derived from the  $^{15}\text{N}$ - and  $^{13}\text{C}$ -edited NOESY

**Table 1**  
Structural statistics for the 20 best structures of SPINK6. No distance constraint was violated by more than 0.5 Å in any structure. Rmsd = root mean square deviation. \*, rmsd was calculated for regions containing helix and strands.

|   | Number      |
|---|-------------|
| <b>Distance restraints</b>  |             |
| Intraresidual ( $ i-j  = 0$ )   | 173         |
| Sequential ( $ i-j  = 1$ )  | 161         |
| Medium range ( $2 \leq  i-j  \leq 4$ )                                | 57          |
| Long range ( $ i-j  \geq 5$ )   | 59          |
| Three disulphide bonds included                                       | 18          |
| Total   | 450         |
| <b>Pairwise rmsd*</b>   |             |
| Average rmsd backbone all   | 0.59 ± 0.12 |
| Heavy atoms rmsd:   | 1.25 ± 0.19 |
| <b>Ramachandran plot (for average structure out of 20 structures)</b> |             |
| Most favored regions (%)  | 77          |
| Additional allowed regions (%)  | 23          |
| Generously allowed regions (%)  | 0.5         |
| Disallowed regions (%)  | 0.1         |

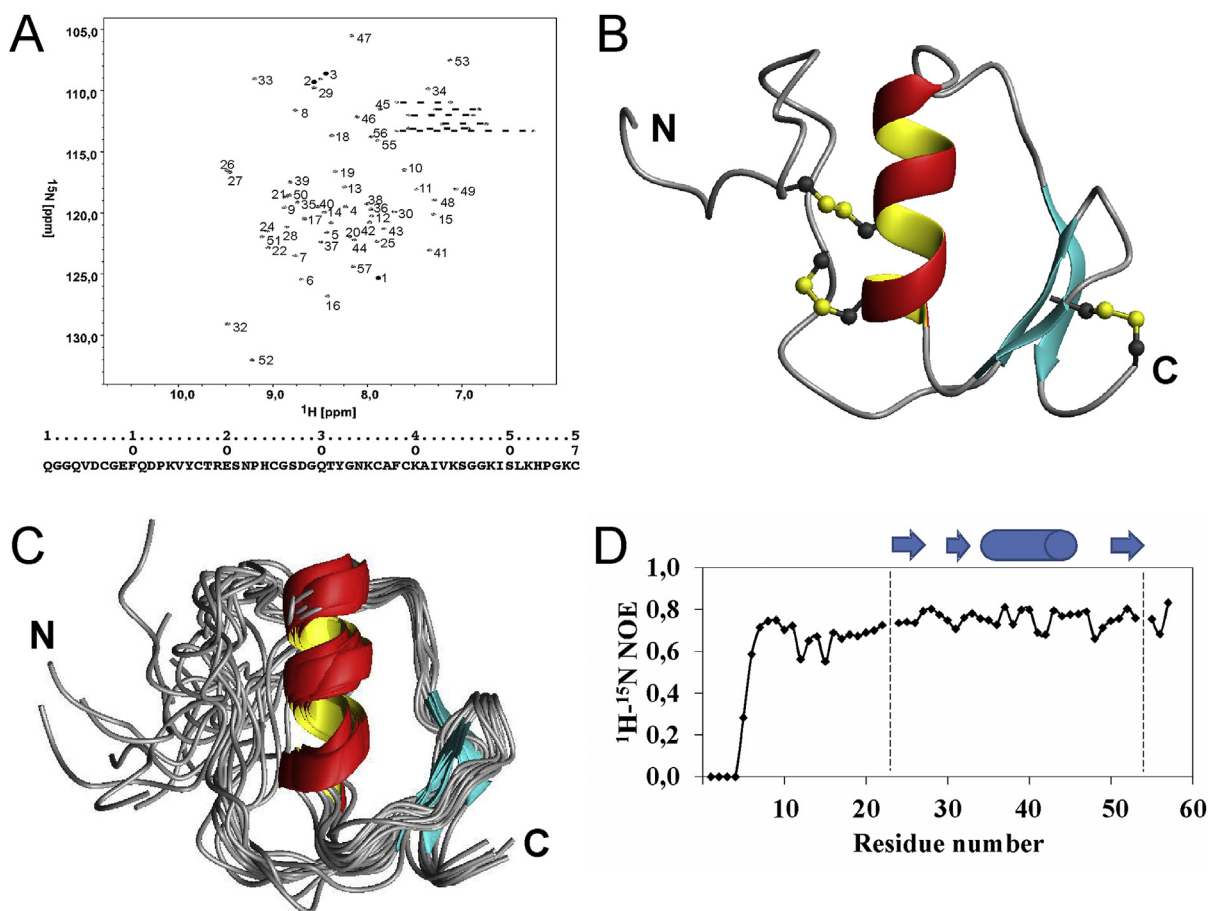
**Table 2**  
Eight templates identified by PS-HomPPI v1.3.

| PDB ID | Chain ID | Identity with SPINK6 | Identity with KLK4 |
|--------|----------|----------------------|--------------------|
| 1ppf   | I:E      | 55.56                | 31.58              |
| 1hja   | I:B      | 56.60                | 34.85              |
| 1z7k   | B:A      | 33.33                | 39.11              |
| 1cho   | I:F      | 55.56                | 34.85              |
| 1cho   | I:G      | 55.56                | 36.46              |
| 1ldt   | L:T      | 41.46                | 39.11              |
| 1an1   | I:E      | 41.46                | 39.11              |
| 1tgs   | I:Z      | 36.59                | 40.44              |

part (residue 1 to 6) confirm the high flexibility and lack of structure in accordance with the results of the structure calculations.

### 3.2. Structural comparison of SPINK6 with SPINK2 and SPINK7

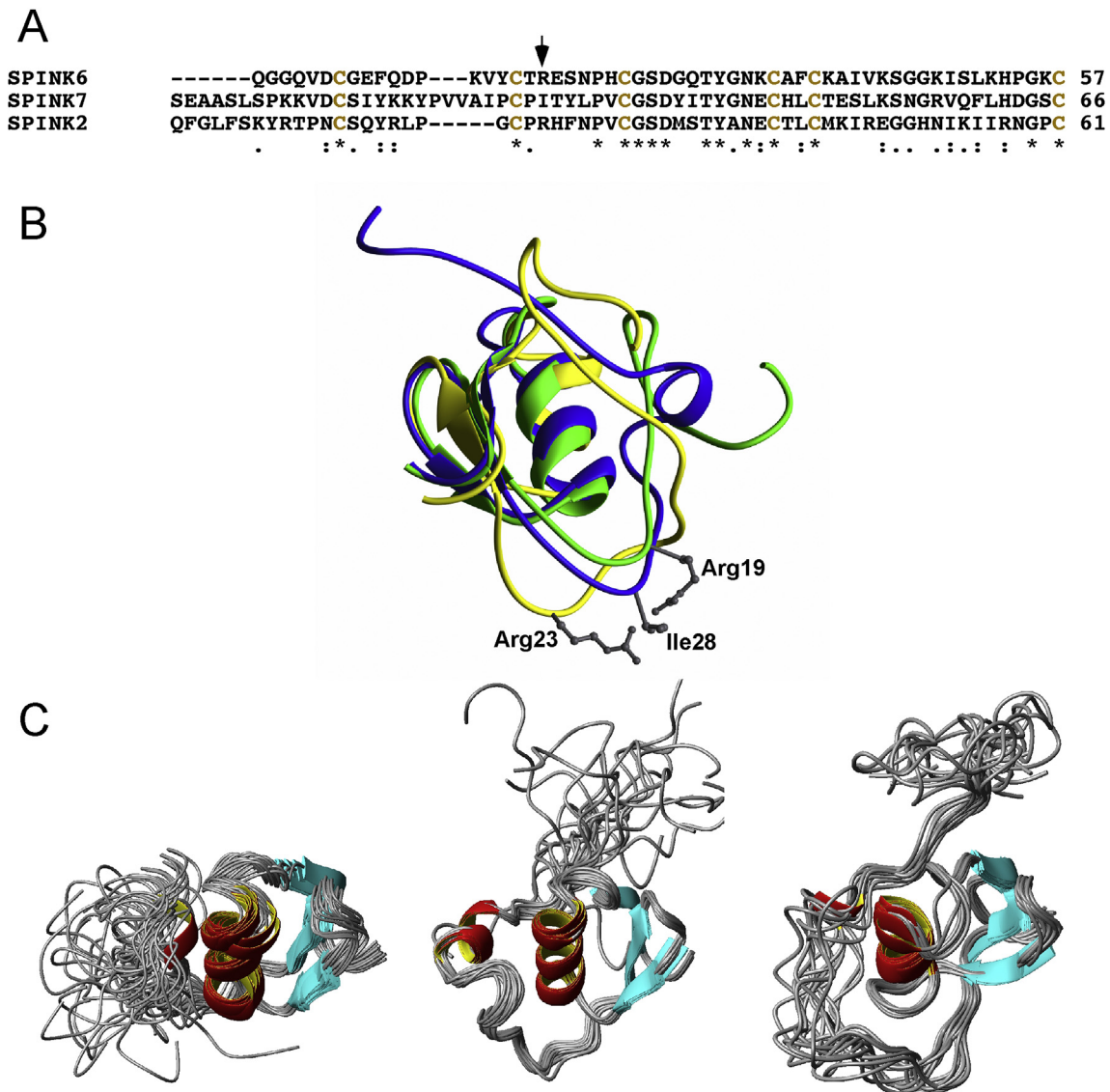
The structures of two other members of the SPINK family have been previously solved – i.e. SPINK2 [25] and SPINK7 [26]. Fig. 2A shows the sequence alignment of the three molecules. The cysteine pattern is conserved in all three molecules. The central residue suggested to be involved in binding to the so-called S1-pocket of



**Fig. 1.** (A)  $^1\text{H}$ - $^{15}\text{N}$  HSQC-spectrum of SPINK6. Resonances are numbered according to the sequence numbers below. Side chain H-N resonances are connected by dotted lines. (B) Ribbon representation of the SPINK6 structure. Side chains of the disulfide-bridges are depicted in yellow and gray and the N-terminus (N) and C-terminus (C) are labeled. (C) Overlay of the final refined 20 structures ensemble. For superposition the secondary structural elements were used and the N-terminus (N) and C-terminus (C) are labeled. (D) Flexibility of the protein backbone measured by heteronuclear NOEs. Positions of helix (cylinder) and  $\beta$ -strands (arrows), respectively, are depicted on top according to the involved residue numbers.

spectra and used to calculate the three-dimensional structure. Fig. 1B shows the representative overall structure of SPINK6. From 100 calculated structures, the 20 energetically best structures were selected for the ensemble which is shown in Fig. 1C. The structural statistics of this ensemble are shown in Table 1. The overall structure of SPINK6 comprises two adjacent  $\beta$ -strands, followed by an  $\alpha$ -helix and a C-terminal  $\beta$ -strand. All secondary elements are well defined as it can be deduced from the ensemble. Since the N-terminus is less defined and exhibited a wide range of different conformations we assessed the backbone flexibility by  $^1\text{H}$ - $^{15}\text{N}$  heteronuclear NOE experiments. Fig. 1D shows the analysis of these data. Whereas for residues 7 to 57 the values around 0.8 are indicative of a rigid structure, lower NOE ratios for the N-terminal

the active site of their target proteases is an arginine in case of SPINK6 and 2, and an isoleucine in case of SPINK7. Arginine residues at this position are typical for trypsin-like proteases, whereas bulky hydrophobic residues, like isoleucine, are typical for chymotrypsin-like proteases. Fig. 2B shows the superposition of the three representative SPINK structures. Whereas the central  $\alpha$ -helix and the  $\beta$ -sheet are structurally conserved, the N-termini differ remarkably. As a consequence, the residues of the SPINK molecules with the S1-pocket of the active site of the proteases differ significantly in their orientation. Fig. 2C shows the side-by-side comparison of the three ensembles of the SPINK structures. Again, the central  $\alpha$ -helix and the  $\beta$ -sheet are very well defined, whereas the N-termini of all three molecules are highly flexible and



**Fig. 2.** Comparison of the structures of SPINK2, 6, and 7. (A) Sequence alignment of SPINK2, 6, and 7. The position of residues known to bind to the active site of the proteases is marked by an arrow. (B) Superposition of the averaged structures of SPINK6 (green), SPINK2 (yellow, PDB ID: 2jxd) and SPINK7 (blue, PDB ID: 2leo). For superposition the secondary structure elements were used. Side chains of the residues known to bind to the active site of the S1-pocket of the proteases are depicted and labeled. (C) Structural ensembles of SPINK6, SPINK7 and SPINK2 (from left to right). In all three molecules, the N-terminal region, which contains the amino-acid residue side chain involved in the inhibition of KLKs, exhibits a high flexibility. Molecules are shown in the orientation with respect to the central helix.

unstructured.

### 3.3. The SPINK6 – KLK4 model

Using the information derived from the identified templates (see Material and Methods) as restraints to define the backbone conformation of the interaction loop and to maintain the conserved arrangement of the catalytic triad in SPINK6, we were able to generate models of the complex of Spink6 and KLK4, which show an improved conformation of the interaction loop of SPINK6 (Fig. 3). The interaction loops of refined SPINK6 models (residues 17–22) are now forming conformations that are similar to the templates; furthermore, the previously disordered N-termini (residues 1–16) of unbound SPINK6 NMR models, which originally pointed in various directions and initially clashed with KLK4 after superimposition, do not clash any more with KLK4. The resulting

models still satisfy the original NOE restraints.

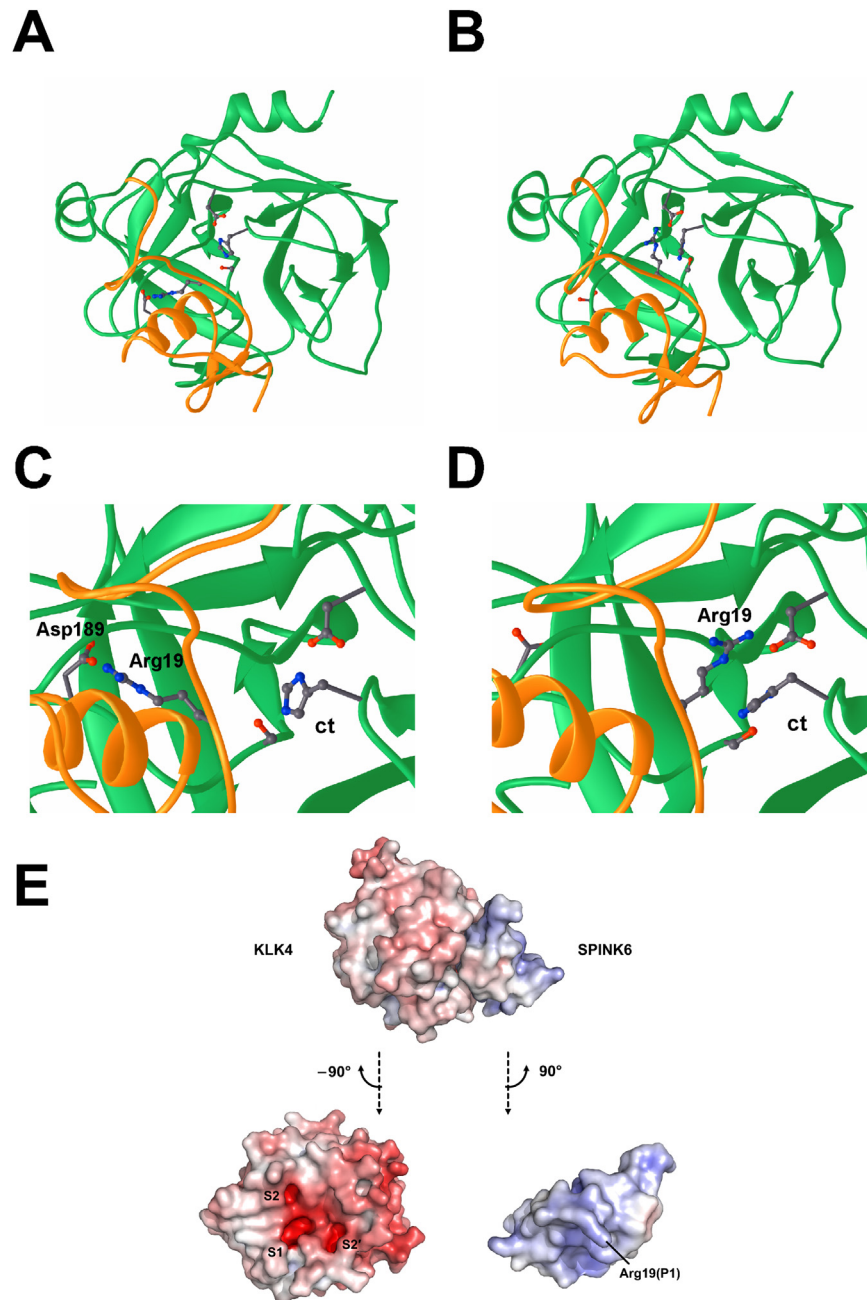
While the backbone interaction conformations of the top 10 models are almost identical, the side chain of the arginine (Arg19) known to bind the active site of KLK4 presents two reasonable but distinct binding conformations<sup>1</sup> (Fig. 3): 1) conserved binding mode: binding the primary pocket of KLK4 as in the templates (Fig. 3C); 2) an alternate binding mode: binding the secondary S2 pocket (Fig. 3D). Models with either binding conformation have reasonable HADDOCK energy score (Table 3).

## 4. Discussion

Proteolytic processes play an important role in many

<sup>1</sup> Two models were discarded due to an unrealistic conformation of Arg19, which pierced deep into the surface of KLK4.





**Fig. 3.** The complex of SPINK6 (yellow) and KLK4 (green). (A and B) Overview of the two conformations of the KLK4-SPINK6 complex. The amino-acid residues constituting the active site of the protease (catalytic triade, ct, Asp102/His57/Ser195), the canonical binding residue (Asp189) and Arg19 of SPINK6 are depicted. (C and D) Close up of the two alternative binding modes of Arg19 of SPINK6 to the active site of KLK4. (E) The KLK4-SPINK6 complex in the canonical binding mode, shown sideways with respect to the standard orientation of the protease. The electrostatic potential is contoured from  $-10$  kBT/e (red) to  $+10$  kBT/e (blue). In the lower panel, both KLK4 and SPINK6 have been turned by  $90^\circ$ , but in opposite directions. The KLK4 active site exhibits highly negative potentials in the S1, S2, and S2' specificity pockets. Additionally, an extended negatively charged patch on the right might serve as exosite for substrate binding. The SPINK6 reactive centre loop and surrounding region is both complementary to KLK4 with respect to shape and its positively charged residues, such as Arg19, which is flexible enough to switch between binding of the S1 and S2 pockets.

physiological and pathophysiological situations. The members of the large superfamily of serine proteases are involved in blood coagulation, activation of the complement system, but also cancer progression. Several kallikrein-related peptidases has been described to be involved in the proteolytic regulation of the desquamation process of the skin. In addition to their physiological roles, the dysregulation of KLK expression, activity or localization, has been described to be associated with pathophysiological disorders, such as skin disorders, respiratory diseases, perturbation of

tooth enamel development, neurological disorders and carcinogenesis [27].

The proteolytic activities of these proteases are regulated by natural inhibitors such as metal ions, serpins, macroglobulins, and Kazal-type inhibitors [4]. Although expression levels of KLK4 in skin are moderate and transient [5], a regulation by SPINK6 seems plausible. However, the major targets of SPINK6 in skin are most likely the two tryptic proteases KLK5 and KLK14, exhibiting high expression levels [10].

**Table 3**  
**HADDOCK statistics for SPINK6-KLK4 models grouped with two distinct conformations of the Arg19 side chain.**  $E_{vdw}$  is Van der Waals energy,  $E_{elec}$  is electrostatic energy, and BSA is Buried Surface Area.

|                                    | ARG_19 (conserved mode) | ARG_19 (alternate mode) |
|------------------------------------|-------------------------|-------------------------|
| $E_{vdw}$ [kJ mol <sup>-1</sup> ]  | -325.1 ± 13.4           | -282.0 ± 14.2           |
| $E_{elec}$ [kJ mol <sup>-1</sup> ] | -725.9 ± 77.0           | -979.1 ± 231.4          |
| BSA [Å <sup>2</sup> ]              | 1948 ± 53               | 1861 ± 57               |
| HADDOCK score [a.u.]               | -113.2 ± 3.7            | -114.9 ± 6.2            |

Kazal-type inhibitors of trypsin-like serine-proteases share a central basic residue, either a lysine or an arginine residue (corresponding to Arg19 of SPINK6, see Fig. 2A) which targets the aspartic acid (Asp189) at the bottom of the S1 specificity pocket in a substrate-like manner. Usually, the S1 pockets accommodate the basic P1 amino acid of a substrate, which is cleaved by trypsin-like proteases. Interestingly, our results suggest a second alternative mode of interaction. Here, Arg19 of SPINK6 binds to the S2 specificity pocket and interacts with the aspartic acid, Asp102. Although binding to the protease may favor the canonical mode, it is remarkable that the alternative conformation is suitably preformed and complementary to the S2 pocket, which exhibits a high negative electrostatic potential, comparable to the one of the S1 pocket (Fig. 3E). Such an interaction has so far not been observed for SPINK inhibitors. Further structural studies of other SPINK/KLK complexes have to be performed, in order to prove whether such an alternative binding mode really exists and how it might contribute to the different inhibitory effects observed for the different KLKs.

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