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Microbiology

Research Article

A large soluble domain of the *Staphylococcus aureus* ESAT-6 Virulence Factor EsaA is stable in the absence of its cognate transmembrane domains

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ABSTRACT

Staphylococcus (s.) aureus is a commensal microorganism with a significant threat to human health. *S. aureus* harbors secretion systems that are utilized to exploit host cells through the secretion of a subset of virulence proteins that cause serious illness. The ESAT6-like or Type VII Secretion System (T7SS) contributes to *S. aureus* virulence. The T7SS secretion system encodes at least twelve proteins categorized as cytosolic, membrane-associated or secreted. Little is known about the exact components of the translocation machinery or translocation mechanism of T7SS substrates across the staphylococcal envelope, but translocation of T7SS substrates across bacterial membranes requires four membrane proteins: EsaA and EssA, B, and C. Topology predictions of EsaA suggest six transmembrane domains with large soluble stretch, likely exposed into *trans* side of the membrane. Whether the large soluble stretch of EsaA is stable without the need for other transmembrane domains, a recombinant 6x-His-tagged soluble domain of EsaA was overproduced in *Escherichia (E.) coli* BL21(DE3) cells. This was followed by affinity purification of the tagged EsaA soluble over a Fast Protein Liquid Chromatography-operated nickel column to apparent homogeneity. SDS-PAGE of the affinity-purified soluble stretch without its cognate transmembrane domains revealed a strong signal, suggesting an independent role for that domain in mediating protein-protein interactions within the ESAT-6 secretion system.

Keywords: *S. aureus*; ESAT-6; Type VII secretion; EsaA-SD

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Citation | Manar M. Ahmed, and Yasser M. Ragab, and Khaled M. Aboshanab. 2017. A large soluble domain of the *Staphylococcus aureus* ESAT-6 Virulence Factor EsaA is stable in the absence of its cognate transmembrane domains. Arch Pharm Sci ASU 1(2): 39-45

DOI: [10.21608/aps.2017.11025](https://doi.org/10.21608/aps.2017.11025)

Online ISSN: 2356-8380

Print ISSN: 2356-8399

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Published by: Ain Shams University, Faculty of Pharmacy.

1. INTRODUCTION

S. aureus is a commensal bacterium with a significant threat to public health [1, 2]. Under some cases of skin injury, innate immune barriers are breached and *S. aureus* gains access into deeper tissues, leading to persistent staphylococcal abscesses in host tissues without promoting immune response [3]. *S. aureus* develops secretion systems

to export their virulence factors extracellularly and into the vicinity of their host environment [4].

ESAT-6 (early secreted antigenic targets of 6 kDa) or Type VII Secretion System (T7SS) is a specialized secretion system that contributes to *S. aureus* virulence [5]. It is evolutionarily related to the ESX-1 secretion system, initially identified

in *Mycobacterium tuberculosis* during the isolation of the BCG vaccine strain *Mycobacterium bovis* [6]. Genomic data analysis show, that ESAT-6 is conserved in other Gram-positive Firmicutes such as most of *S. aureus* strains [7], *Bacillus subtilis* and *S. agalactiae* [8]. The *S. aureus* T7SS encodes at least twelve proteins categorized as cytosolic (EsaB), four membrane-associated proteins (EsaA, EssA, EssB, and EssC), and several secreted proteins (EsxA, EsxB, EsxC, EsxD, and EssD). The two remaining genes encode, EsaE; a membrane platform protein and EssaG, an EssD nuclease [4, 9, 10]. The EsxA and EsxB WXG100 secreted proteins are essential for persistence of staphylococcal abscesses as indicated by murine abscess model experiments [4]. Furthermore, EsxC and EsxD are also secreted and essential for staphylococcal virulence. The hallmark of *S. aureus* T7SS is the secretion of EsxABCD that strongly manipulate host defense mechanisms [4, 8, 10].

Little is known about the translocation machinery or translocation mechanism of T7SS substrates across the staphylococcal envelope, but, translocation of T7SS substrates across biological membranes requires four membrane proteins: EsaA, EssA, EssB and EssC [4, 5, 7]. The molecular mechanism of EssC has been assessed as FtsK/SpoIIIE homolog with ATPase activity; probably acting to energize substrate translocation across staphylococcal envelope [11, 12]. The crystal structure of EssB shows bitopic transmembrane domain residing within the membrane. The C-terminus is likely located into *trans* side of the membrane and is essential for EssB homodimerization, whereas the N-terminus is located in the cytoplasm and its folding pattern resembles that of protein kinases likely recognizing several T7SS substrates [13].

The molecular mechanisms of EssA and EsaA are under ongoing investigation. Topology predictions of EsaA using TMHMM reveal six transmembrane domains embedded within the membrane and spread across protein length. In addition, EsaA contains one large soluble stretch located between TM1 and TM2 and is likely exposed into the *trans* side of the membrane. We

wondered whether the sole large soluble stretch of EsaA membrane protein is stable without the presence of other TM domains, which suggests an independent role in mediating protein-protein interactions. In order to test this hypothesis, a systematic experimental approach has been carried out in this work to confirm this hypothesis.

2. MATERIALS AND METHODS

2.1. Bacterial cultures

E. coli cultures of DH5 α and BL21 (DE3) were grown in Luria Bertani (LB) medium (10 g NaCl, 10 g peptone, 5 g yeast extract) at 37 °C. Ampicillin antibiotic was used at final concentration 100 μ g/mL for plasmid selection in *E. coli*. Isopropyl β -D-1-thiogalactopyranoside (IPTG) with final concentration 0.5 mM was added for gene induction, if necessary. Phenylmethylsulfonylfluoride (PMSF) as protease inhibitor at final concentration 0.5 mM and lysozyme at 1 mg/ml were also added prior to protein purification.

2.2. Bacterial strains and plasmid

E. coli DH5 α cells were used for cloning experiments. *E. coli* BL21 (DE3) cells were also used for protein overproduction experiments. 6x His-tagged *esaA* soluble part (EsaA-SD) was cloned into an expression vector pET15 using the following oligonucleotide sequences (F: 5' CGCAGCATATGACTTTAATTGAAAAACA AAATTCATTAT 3') and (R: 5' CGCAGGGATCCTTATAAAATCACCATTAA GATGAATTTC 3').

2.3. Cloning of *esaA* soluble stretch

esaA soluble stretch was amplified by polymerase chain reaction (PCR) using genomic DNA extracted from *S. aureus* strain USA300 using Qiaminiprep as a PCR template and the aforementioned set of oligonucleotides. Next, PCR products were purified from agarose gels using standard methods and subjected to restriction endonucleases digestion using *NdeI* and *BamHI*. The cleaved products were then ligated into the pET15b expression vector digested with the same restriction endonucleases. Recombinant plasmid (pET15b6x *esaA-SD*) was

then introduced into *E. coli* BL21 (DE3) competent cells for protein overproduction and subsequent affinity purification

2.4. Recombinant protein affinity purification

Plasmid pET15b6x *esaA-SD* was introduced into *E. coli* BL21 (DE3) cells. *E. coli* BL21 (DE3) cultures harboring pET15b6x *esaA-SD* were grown overnight at 37 °C in liquid LB, supplemented with ampicillin for vector propagation. Next day, cultures were subcultured into fresh LB medium containing ampicillin and further incubated at 30 °C until absorbance values A_{600 nm} reached 0.6. Thereafter, IPTG with final concentration 0.5 mM was added, and cells were further incubated for three hours at 30 °C. Cells were harvested by centrifugation (12,000 rpm) while discarding the supernatant. The obtained cell pellet suspended in PBS (phosphate buffer saline) containing 20 mM imidazole, 0.5 mM PMSF and 1mg/ml lysozyme and further subjected to sonication 30 amp (5 min), 10 sec pulse rate and 10 sec break in-between, followed by 30 amp (6 min), 10 sec pulse rate and 10 sec break in-between. Crude lysates were centrifuged in a cooling centrifuge at 4000 rpm for 20 min with the recovery of the filtered supernatant of extracts, producing his-tagged *esaA-SD* along with other native *E. coli* proteins (total cell lysates). For affinity purification, the supernatant was injected using a needle syringe into immobilized metal affinity chromatography (IMAC) over an FPLC-attached His-Trap nickel column (GE Healthcare). The column was washed with phosphate buffer saline, pH 7.5 containing 20 mM imidazole as wash buffer, followed by gradual elution in the presence of phosphate buffer saline (PBS) containing 500 mM imidazole. The eluted fractions of 6x-His-tagged *EsaA-SD* were concentrated with acetone (3:1 and incubated on ice for 30 min, then centrifuged for 45 min at 4000 rpm, room temperature. Acetone treated elution fractions were air dried. Proteins in all fractions (TCL: total cell lysates, FT: flow through, W: wash and EL: elution) were identified using sodium dodecyl sulfate poly Acrylamide gel electrophoresis (SDS-PAGE).

2.5. SDS-PAGE protocol

Following protein affinity purification, standard sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) was performed to validate *EsaA-SD* production and stability. Protein fractions (BL21EV, TCL, FT, W and EL) were mixed with sample buffer and incubated for 5 min at 95 °C. Samples were loaded onto an SDS-PAGE gel [14], followed by coomassie blue staining (300 mL methanol, 100 mL glacial acetic acid, 600 mL distilled water and 1 g Coomassie Blue R250) to detect the protein signal at ~ 70 kDa. De-staining solution (200 mL methanol, 80 mL of glacial acetic acid, and 720 mL distilled water) was used to remove excess staining and manifest band detection.

2.6. Membrane topology predictions

Tied Mixture Hidden Markov Model (TMHMM) server (<http://www.cbs.dtu.dk/services/TMHMM>) was used as software for topology predictions of the *EsaA* membrane protein. *EsaA* amino acid sequence was introduced into the server in a FASTA format, followed by detection of the protein topology as outlined by the web-based tool.

3. RESULTS AND DISCUSSION

3.1. The *EsaA* soluble stretch is stable without its cognate transmembranes

As shown in **Fig. 1**, topology prediction of *EsaA* showed that it is an integral membrane protein with 1009 amino acids comprising six transmembrane domains spread throughout the protein length, with large soluble stretch likely exposed into the *trans* side of the membrane. *EsaA* protein sequence: N- terminus (1-4 amino acids), TM1 (5-27 aa), soluble stretch (28-820 aa), TM2 (821-843 aa), TM3 (864-886 aa), TM4 (901-923 aa), TM5 (928-945 aa), TM6 (978-1000 aa). To confirm whether the sole large soluble stretch of *EsaA* is stable in the absence of cognate TM, an affinity purified 6x-His-tagged *EsaA* soluble stretch (*EsaA His-SD*) was performed upon overproduction in *E. coli* BL21 (DE3) cells using IMAC, over an FPLC-operated His-Trap nickel column (GE Healthcare). Different samples (TCL for total cell lysates, FT

for flow through, WS for wash and EL for elution) were collected based on the chromatogram chart (**Fig. 2**).

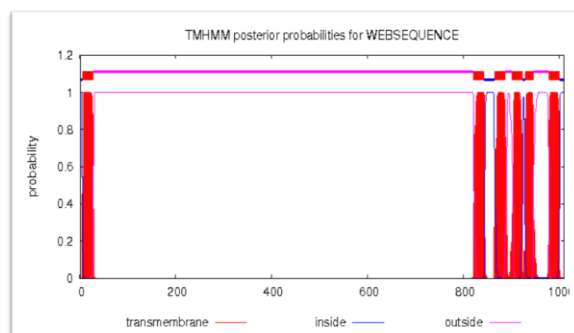


Fig. 1 TMHMM Topology prediction of EsaA integral membrane protein

EsaA is integral membrane protein with 1009 amino acids comprising six transmembrane domains spread throughout the protein length, with large soluble stretch likely exposed into the *trans* side of the membrane. EsaA protein sequence: N- terminus (1-4 amino acids), TM1 (5-27 aa), soluble stretch (28-820 aa), TM2 (821-843 aa), TM3 (864-886 aa), TM4 (901-923 aa), TM5 (928-945 aa), TM6 (978-1000 aa).

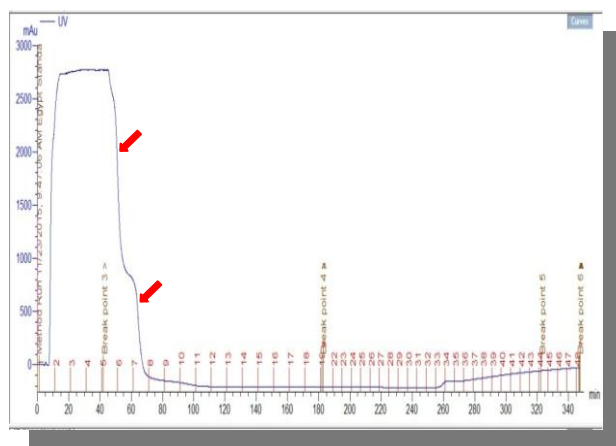


Fig. 2 FPLC Chromatogram chart of the EsaA-SD purification, Arrows are flow through and wash.

Chromatogram represents flow through, wash and elution fractions after injecting the supernatant sample (containing 6x-His-tagged EsaA-SD) into IMAC over an FPLC-attached His-Trap nickel column (GE Healthcare). Flow

through the sample (fraction 5-6), Wash sample (fractions 7-15) and elution samples (36-40).

3.2. SDS-PAGE analysis of EsaA

An additional band of SD-EsaA of approximately ~70 kDa appeared in the total cell lysate (TCL) (Fig. 3, lane 3 left) that refers to successful expression of the SD-EsaA, while it does not show in flow-through sample (FT), as the flow-through sample includes all proteins except His-tagged protein (His-tagged EsaA-SD), as shown in (**Fig. 3 lane 4 left**). The washed sample indicates the successful removal of non 6x-His-tagged proteins traces. EsaA-SD elution fractions (36-40) in the chromatogram (**Fig. 2**) was concentrated using acetone followed by SDS-PAGE, in elution fraction, and a protein band appeared in approximately ~70 kDa, where EsaA-SD is supposed to appear near this molecular size (**Fig. 4, lane 5**).

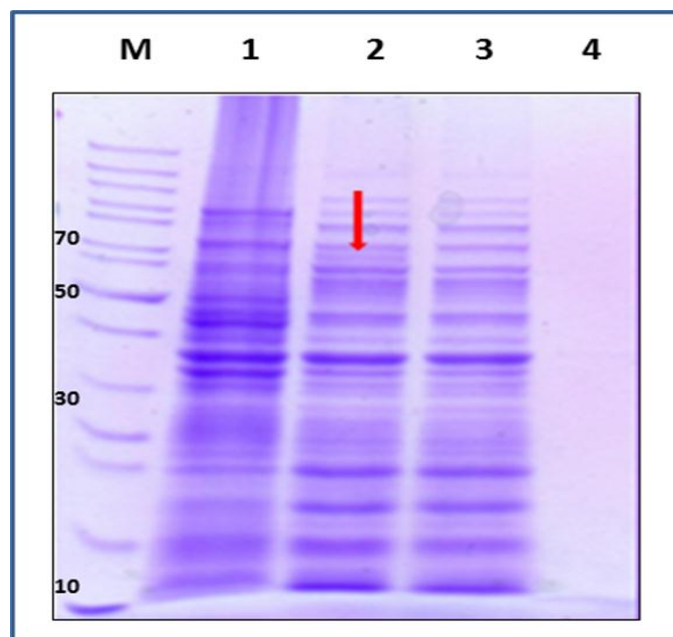


Fig. 3 Coomassie-stained SDS-PAGE of FPLC fractions of affinity purified EsaA-SD.

SDS-PAGE was performed for analysis of total cell lysates containing *E. coli* proteins, including 6x-His-tagged EsaA-SD, Flow through sample and wash sample using *E. coli* BL21 EV as a control. Lanes from left to right: M; Protein

molecular weight marker (10-200 kDa), lane 1; BL21-EV cells following induction with IPTG as a negative control, lane 3; (TCL) crude lysates of BL21-pET15 harboring EsaA (SD), lane 4; (FT) Flow-through,5; Wash.

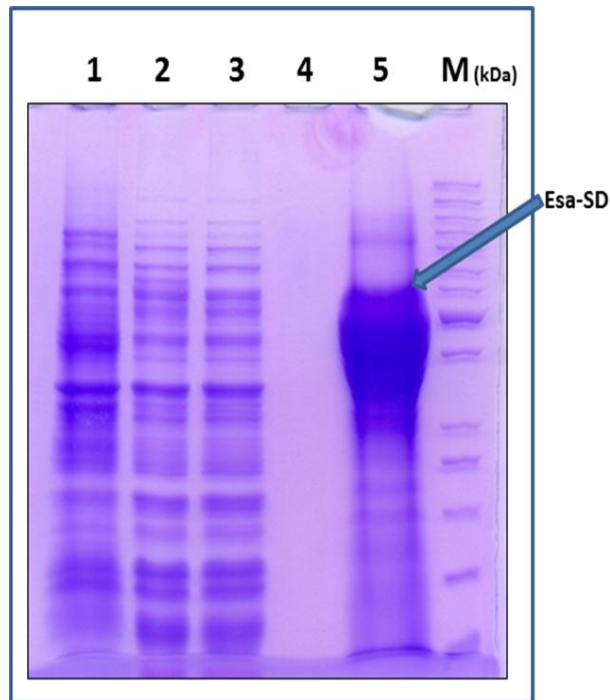


Fig. 4 SDS-PAGE of FPLC concentrated EsaA-SD elution fractions

EsaA-SD elution fractions (36-40) are subjected to concentration with three times absolute acetone solution. SDS-PAGE represents concentrated elution sample with total cell lysates, flow through and wash using BL21-EV as a negative control. Lanes from left to right: Lane 1; BL21-EV cells following induction with IPTG as negative control, Lane 2; crude lysates of BL21-pET15b harboring EsaA (SD), Lane 3; flow-through sample representing all proteins except 6x-His-tagged EsaA-SD, Lane 4; wash, lane 5; concentrated EsaA-SD elution fractions ~70 kDa, lane M; Protein molecular weight marker (10-200 kDa).

4. DISCUSSION

The ESAT-6 secretion pathway is crucial for pathogenicity of many human pathogens, such as *S. aureus* and *M. tuberculosis*. Membrane

proteins often show higher importance in many physiological mechanisms including structural and functional integrity and it is not a coincidence that they are targeted by over 50% of marketed drugs [15]. The *S. aureus* ESAT-6 system is composed of four integral membrane proteins: EsaA, EssBA and EssC, an associated ATPase, which are all required for protein secretion across the staphylococcal envelope [4, 5, 16]. EsaA exhibits six transmembrane domains with a large extracellular loop.

In this study, we have investigated whether the sole large soluble stretch of EsaA is stable without the need for other transmembrane domains using affinity purification. Our results using affinity purification of a 6x-His-tagged EsaA soluble stretch performed over a Fast Protein Liquid Chromatography-operated nickel column reveal the stability of EsaA soluble stretch without its cognate transmembrane domains, suggesting an independent role in protein-protein interactions within the ESAT-6 secretion system.

Previous studies on membrane proteins harboring long extracellular loops develop many roles. For example, the *S. aureus* BraSR/BraDE/VraDE (for Bacitracin resistance associated) system, and the extracellular loop of VraE has been shown to associate bacitracin resistance [17]. Further, the BceB extracellular loop in *B. subtilis* is thought to be necessary for bacitracin sensing and resistance [18, 19].

Similar to the EsaA extracytoplasmic loop, studies have shown that conserved extracytoplasmic loops in other bacteria including *B. subtilis* and *E. coli* may exhibit important functional mechanisms. Interestingly, short deletions in the *E. coli* Sec Dextra cytoplasmic loop result in malfunctioning of the protein without affecting the stability of SecDF complexes with others, suggesting its participation in providing a protective structure for efficient protein folding [1]. Moreover, in *B. subtilis*, topology predictions of SecDF also show besides its 12 transmembranes, two extracytoplasmic loops conserved between TM1-TM2 and TM7-TM8, where studies show SecD and SecF with deletion in the large cytoplasmic

loop, displayed translocation defect without affecting SecD-SecF interaction, reported that it is important for catalyzing protein translocation [1, 20]. The exact function of extracytoplasmic loops is not completely understood, but they may participate as binding sites for lipids [21] or associate with lipids for correct insertion, folding, and proper topology of the protein [22]. In addition, studies of C2 domain structure show surface loops that are required for domain specificity, with their amino acid sequence and conformation varying and mostly involved with lipid binding [23]. Future studies will be performed in order to understand the exact function of the EsaA extracytoplasmic loop and its role with other membrane proteins within the ESAT-6 pathway.

5. CONCLUSION

SDS-PAGE analysis of the expressed soluble stretch of the EsaA membrane protein, a major protein of the ESAT-6 secretion system of *Staphylococcus aureus* that contributes to its virulence showed its stability without its cognate transmembrane domains revealed strong signal, suggesting an independent role for that domain in mediating protein-protein interactions within the ESAT-6 secretion system or Type VII Secretion System (T7SS)

Acknowledgment

We acknowledge the Department of Microbiology and Immunology, Faculty of Pharmacy and Pharmaceutical Industries, Sinai University, El-Arish, North Sinai, Egypt for providing all required Laboratory facilities when required as well as their friendly cooperation.

Conflict of Interest

The author declares no Conflict of Interest

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