Intra- and intermolecular events direct the propeptide-mediated maturation of the Candida albicans secreted aspartic proteinase Sap1p

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Pathogenic yeasts of the genus Candida secrete aspartic proteinases (Sap) which are synthesized as preproenzymes. Expression of the C. albicans SAP1 gene lacking the propeptide-coding region in the methylotrophic yeast Pichia pastoris does not lead to the secretion of the enzyme into the culture supernatant, but results in an accumulation of recombinant protein in the cell. Co-expression in this system of the unattached propeptide from Sap1p, as well as from other Saps, restored Sap1p secretion. A deletion analysis revealed that only a 12 aa sequence in the propeptide, corresponding to a highly conserved region in all Sap propeptides, was necessary and sufficient to produce a large amount of Sap1p in culture supernatant. No Sap1p was secreted when Sap1p was produced with a propeptide carrying an F to D mutation in the identified 12 aa sequence. However, the simultaneous production of equivalent amounts of Sap1p and His-tagged Sap1p (H6-Sap1p) with a mutated and a non-mutated propeptide, respectively, led to the secretion of both proteins in a ratio of approximately 1:2. The restoration of Sap1p secretion occurred at the expense of secretion of H6-Sap1p since the total activity was comparable to that of strains producing only H6-Sap1p with a non-mutated propeptide. In contrast, the proteolytic activity of strains secreting Sap1p and H6-Sap1p both with a functional propeptide was twice that of strains producing either Sap1p or H6-Sap1p alone, and the two enzymes were found in an equivalent amount in the culture supernatant. Altogether, these results show that the propeptide can only function once and that the maturation of recombinant C. albicans secreted aspartic proteinase Sap1p is directed through a combination of intra- and inter-molecular pathways.

Keywords: SAP1 gene, protein secretion, signal peptides, recombinant proteins, Pichia pastoris

INTRODUCTION

The secreted aspartic proteinases (Saps) of the opportunistic pathogens of the genus Candida are of particular interest because of their role as virulence factors (Fallon et al., 1997; Borg-von Zepelin et al., 1998; Schaller et al., 1998, 1999). In C. albicans, they are encoded by a gene family with at least nine members, SAP1 to SAP9 (Monod et al., 1994, 1998). To date, only one gene from C. tropicalis (SAP1; Togni et al., 1991) and two from C. parapsilosis (SAP1 and SAP2; de Viragh et al., 1993) have been cloned and sequenced, although preliminary investigations support the existence of more SAP genes in both species (Monod et al., 1994). The Candida Saps, like many other microbial secreted proteinases, are synthesized as precursors in a preproprotein form. The prepeptide, or signal peptide, of 16–18 aa residues is necessary for entry into the secretory pathway by transporting the protein across the membrane of the endoplasmic reticulum (Pfeffer &
Rothman, 1987). With the exception of Sap7p, which has a putative 195 aa propeptide, the Candida Saps have a relatively short propeptide (32–58 aa) which contains one to four KR sequences, one of which is located immediately before the N-terminus of the mature part of the proenzyme.

The propeptide of many secreted proteinases has been found to be essential and specific for assisting correct folding as well as secretion of the mature domain of the enzyme (for a review, see Eder & Fersht, 1995). Upon completion of folding, the propeptide is cleaved and removed to generate the active enzyme through an autoproteolytic reaction or, as in the Candida Saps, through an exogenous proteolytic reaction in the Golgi apparatus via the membrane-bound proteinase Kex2p through an exogenous proteolytic reaction in the Golgi autoproteolytic reaction or, as in the enzyme (for a review, see Eder & Fersht, 1995). Upon folding as well as secretion of the mature domain of the enzyme. It has been clearly established by many examples in vitro and/or in vivo that the propeptide can also mediate the folding of the proteinase when added as a separate polypeptide chain (Ohita et al., 1991; Fabre et al., 1992; Fukuda et al., 1994; McIver et al., 1995; van den Hazel et al., 1994). In vitro, the propeptide covalently attached to the mature domain of prosubtilisin E has been found to function intermolecularly as folding catalyst, although a parallel intramolecular pathway has not been ruled out (Zhu et al., 1989; Hu et al., 1996).

However, whether the maturation of secreted proteinases occurs through inter- or intramolecular events has never been investigated in vivo. This question instigated our detailed examination of the function of the Sap propeptide in the secretion of the mature enzyme. Our study on C. albicans Sap1p was carried out using the pathogenic yeast Pichia pastoris expression system, which allows the production of substantial amounts of recombinant fungal proteinase (Borg-von Zepelin et al., 1998). We show here that recombinant Sap1p maturation is guided by the propeptide in a combination of intra- and intermolecular processes, and that only a 12 aa sequence in the propeptide is necessary and sufficient to completely ensure the secretion of active enzyme.

**METHODS**

**Organisms and plasmids.** Escherichia coli DH5α was used for transformation and propagation of the recombinant plasmids. The E. coli- P. pastoris shuttle vector pPICZαA was provided with the P. pastoris expression system from Invitrogen. The low-copy-number plasmid pKJ113 has been previously described (Borg-von Zepelin et al., 1998). The P. pastoris strains GS115 and KM71 (Invitrogen) were used as transformation hosts.

**Construction of the expression plasmids.** Expression plasmids were constructed by cloning a SAP PCR product in the multiple cloning site of pKJ113 and pPICZαA. Custom-made primers were provided by Microsynth (Balchach, Switzerland). PCR buffers and AmpliTaq polymerase were from Perkin Elmer Applied Biosystems. The buffer composition was 10 mM Tris/HCl (pH 8.3), 50 mM KCl with 1.5 mM MgCl₂, containing 0.2 mM each dNTP and 2.5 U polymerase per reaction. The PCR was carried out in a GeneAmp PCR system 2400 (Perkin Elmer Applied Biosystems) with a first denaturation step of 5 min at 94 °C followed by 25 cycles of annealing at 55 °C for 30 s, elongation at 72 °C for 30 s and denaturation at 94 °C for 30 s. PCR was completed by a final elongation step at 72 °C for 10 min. DNA from plasmids containing individual SAP genes was used as template. The PCR products were purified using a PCR purification kit (Roche Diagnostics) and were digested by restriction enzymes for which a site (XhoI, BamHI or NotI) was previously designed at the 5’ extremity of the primers. The digested PCR products were then cloned into the appropriate sites of the multiple cloning site of the E. coli- P. pastoris shuttle vector.

The following pairs of sense and antisense primers were used to construct plasmids pSB10, pSB11, pSB42 and pSB249 (Tables 1 and 2), respectively, restriction sites underlined: GATTGCTGAGGAGCTCTAAAAGATCCCAGGT and CA- AAAGATCTCATAGGTTAAGAGCAGCATGTGT; GACGTCGAGAGGATCCTACGGATTTAAATT and CAA- AGATCTAGGTAAAGACGCAATGT; TCTCGAGAGAAGATCTCATCTCAGTTAAGAGCAGCATGTGT; GTCTCGAGAAGATCTCATCTCAGTTAAGAGCAGCATGTGT; TTGGACTAAGAGCAGCATGTGT.

Plasmids expressing N- or C-terminally truncated Sap1p propeptide were constructed after PCR amplification of pSB42 fragments with appropriate homologous primers. The PCR products were used to replace the corresponding part encoding the full-length propeptide in pSB42. Subsequently, plasmids expressing Sap1p propeptide truncated at both extremities were constructed in a similar way using previously constructed plasmids encoding Sap1p propeptide truncated at one extremity (see Fig. 4). Plasmids expressing other Sap propeptides were constructed as pSB42 using different SAP DNA as a template (Togni et al., 1991; de Viragh et al., 1993; Monod et al., 1994). All cloned fragments were further sequenced to confirm the absence of possible PCR induced errors.

**In vitro mutagenesis.** For construction of pSB269, the following pairs of sense and antisense primers were used to amplify two contiguous fragments of pSB10: TTGGAGCTC- GCTATTCTCAAT and AGTGACGTACCTGGGGAT- TTTAGCTGG; GGTGATCTCATTATAGGTTAAGAGCAGCATGTGT; TTGGACTAAGAGCAGCATGTGT. Subsequently, the 0.8 kb SacI-NotI fragment of pKJ113 was excised and replaced by adding end-to-end the two PCR products digested by SacI/AatII and AatII/NotI.

**P. pastoris transformation.** P. pastoris was transformed by electroporation with 5–10 μg linearized plasmid DNA digested by Smal using pKJ113 constructs, and SacI or BstXI using pPICZαA constructs. The His’ Mut’ P. pastoris strain GS115 was transformed with pKJ113 constructs, and transformants...
were selected on histidine-deficient medium [1 M sorbitol, 1% (w/v) glucose, 1:34% (w/v) yeast nitrogen base (YNB) without amino acids, 4 × 10⁻³% (w/v) biotin, amino acids (0:005%, w/v, each of l-glutamic acid, l-methionine, l-lysine, l-leucine and l-isoleucine), 2% (w/v) agarose]. Subsequently, the transformants were screened for insertion of the construct at the AOXI site on minimal methanol plates [1:34% (w/v) YNB without amino acids, 4 × 10⁻³% (w/v) biotin, 0:5% (v/v) methanol, 2% (w/v) agarose]. Transformants unable to grow on media containing methanol as a carbon source were retained for further investigations. They were assumed to contain the construct at the correct yeast genomic location by integration events in the AOXI locus displacing the AOXI coding region. The prototrophic P. pastoris strains were transformed with pPICZαA constructs and transformants selected on YPD medium containing 100 µg zeocin ml⁻¹ (Invitrogen).

Production of Sap1p in P. pastoris. All selected transformants were grown to near saturation (OD₆₀₀ 10) at 30 °C in 10 ml glycerol-based yeast medium [0:1 M potassium phosphate buffer pH 6:0, containing 1% (w/v) yeast extract, 2% (w/v) peptone, 1:34% (w/v) YNB without amino acids, 1% (v/v) glycerol and 4 × 10⁻³% (w/v) biotin]. Cells were harvested by centrifugation and resuspended in 2 ml of the same medium with 0:5% (v/v) methanol instead of glycerol and incubated for 2 d. Thereafter, the supernatants and the cell pellets were separated by centrifugation and retained for protein analysis.

Protein extract analysis. The proteins in 5 or 10 µl of P. pastoris culture supernatant were loaded without further treatment onto SDS-PAGE gels (Laemmli, 1970). Cell protein extracts of 0:1 ml culture were prepared following the method of Yaffe & Schatz (1984). The pellets cells were resuspended in 200 µl 1:8 M NaOH, 1-2 M β-mercaptoethanol, and incubated for 5 min on ice. After addition of 200 µl 10% TCA, the mixture was incubated for 5 min, centrifuged, and the pellet was resuspended in 50 µl 2 × SDS-PAGE loading buffer and neutralized at pH 7-0 by adding 5–10 µl 1 M Tris base. The samples were heated to 95 °C for 3 min before loading onto SDS-PAGE gels (12% acrylamide).

Untagged Sap1p was separated from His₆-tagged Sap1p (H₆-Sap1p) by filtration of the P. pastoris culture supernatant through a nickel chelating resin column (ProBond, Invitrogen) equilibrated with 10 mM phosphate buffer, pH 7.0. After washing the column with the same buffer, adsorbed H₆-Sap1p was eluted with 300 mM imidazole buffer, pH 6.0. All fractions with detectable enzymic activity were retained and pooled. Sap1p was purified from filtered P. pastoris culture supernatant as previously described (Borg-von Zepelin et al., 1998). Protein concentrations were measured by the method of Bradford (1976).

SDS-PAGE gels were stained with Coomassie brilliant blue R-250 (Bio-Rad). Immunodetection of Western blots was performed using an antiserum (z-Sap2) raised in rabbits and cross-reacting with Sap1p (Borg-von Zepelin et al., 1998), and alkaline-phosphatase-conjugated goat anti-rabbit IgG (Bio-Rad), or using an anti-polystyidine peroxidase conjugate monoclonal antibody (Sigma).

Proteolytic assays. The proteolytic activity of Sap isoenzymes was measured with 0.02% (w/v) resorufin-labelled cascin as a substrate (Roche Diagnostics) in sodium citrate buffer (50 mM; pH 4.5) in a total volume of 0.25 ml. After incubation at 37 °C for 60 min, the undigested substrate was precipitated by TCA acid (5% final concentration) and separated from the supernatant by centrifugation. The absorbance of the supernatant was measured in the alkaline range at 574 nm after adding 250 µl 1 M Tris/HCl at pH 10. For practical purposes, one unit of Sap1p activity was defined as that producing an absorbance of 0.001 per min.

RESULTS

Importance of the propeptide for secretion of recombinant Sap1p by P. pastoris

In a preliminary experiment, P. pastoris was used to express the gene SAP1 with and without the nucleotide sequence encoding the propeptide. SAP1 DNA fragments (bp 59–1176 and bp 152–1176 of the open reading frame) were cloned into pKJ113 downstream of the pH0I signal peptide sequence, generating the plasmids pSB10 and pSB11, respectively (Table 1). Plasmid pKJ113 is a low-copy-number E. coli vector which contains the P. pastoris pHIL-S1 expression cassette. The pHIL-S1 constructs insert into the P. pastoris genome via homologous recombination at the AOXI site and carry, in addition to the cloned coding sequence of interest, the HIS4 gene for selection after transformation of P. pastoris GS115 with linearized plasmid DNA. The resulting His⁺ Mut⁻ transformants were selected and screened for proteinase production. C. albicans Sap1p was recovered as a single protein and an active enzyme in the P. pastoris culture supernatants in all transformants with a yield of approximately 350 µm⁻¹ (50 µg ml⁻¹) when its encoding nucleotide sequence was expressed behind that of its propeptide. Direct expression of the SAP1 gene without its propeptide did not lead to secretion of recombinant protein into the culture supernatant of P. pastoris transformants. An accumulation of Sap1p in the cell was demonstrated by Western blotting of total protein extracts immunodeveloped with anti-Sap antibody (Fig. 1). One strain secreting and one strain accumulating Sap1p were selected, and labelled SB10 and SB11, respectively, in subsequent investigations (Table 1).

Rescue of Sap1p secretion by unlinked propeptides

The nucleotide sequence encoding the Sap1p propeptide was cloned into plasmid pPICZαA downstream from the α-factor signal peptide sequence generating the plasmid pSB42 (Table 2). This construction was performed in order to provide an access into the secretory pathway for the propeptide as a separate protein. P. pastoris SB11, which does not secrete Sap1p, was transformed with linearized pSB42 DNA, and clones resistant to zeocin were further tested for proteinase production. C. albicans Sap1p was recovered as a single protein and an active enzyme in all transformant culture supernatants with a yield comparable to that of strain SB10. Therefore, co-expression in P. pastoris of a non-covalently attached propeptide could rescue the lack of cis-expressed propeptide.

Five transformants were retained for restriction fragment analysis. Genomic DNA of the yeasts was digested with SacI, DraI and HindIII, electrophoresed and probed
Table 1. *P. pastoris* GS115 and KM71 transformants generated in this study

<table>
<thead>
<tr>
<th>Strain</th>
<th>Transformed original strain</th>
<th>Plasmid</th>
<th>SAP1 translation product</th>
<th>Secreted proteolytic activity (U ml⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SB10</td>
<td>GS115</td>
<td>pSB10</td>
<td>Pre-Pro-Mat</td>
<td>352</td>
</tr>
<tr>
<td>SB11</td>
<td>GS115</td>
<td>pSB11</td>
<td>Pre-Mat</td>
<td>0</td>
</tr>
<tr>
<td>SB269</td>
<td>GS115</td>
<td>pSB269</td>
<td>Pre-Pro*-Mat</td>
<td>0</td>
</tr>
<tr>
<td>SB249</td>
<td>KM71</td>
<td>pSB249</td>
<td>Pre-Pro-Mat-H₃</td>
<td>324</td>
</tr>
</tbody>
</table>

The SAP1 translation products produced with a full length propeptide, without propeptide, with a non-functional propeptide (F27→D mutation) and with a C-terminal His₈ tag are abbreviated as Pre-Pro-Mat, Pre-Mat, Pre-Pro*-Mat and Pre-Pro-Mat-H₃, respectively.

Fig. 1. Detection of Sap1p from *P. pastoris* SB10, SB11 and SB269. The proteins in 5 µl of supernatant (S) without further treatment and in cell extracts (C) from 5 µl of methanol-induced culture were separated by SDS-PAGE and analysed by Western blotting.

In a similar experiment, we tested whether the secretion defect of Sap1p in *P. pastoris* SB11 could also be rescued by providing other secreted proteinase propeptides as a separate protein (Table 2, Fig. 3). Sap1p was recovered as a single protein and an active enzyme in the *P. pastoris* culture supernatants with a yield varying from 128 to 320 units, corresponding to 16 to 40 µg ml⁻¹, using all Sap propeptides tested. The specific activities of the Sap1p enzymes secreted by the different strains were the same.

Using pSB42, we constructed a series of plasmids encoding the Sap1p propeptide deleted at the N- or the C-termi

Table 2. Sap propeptides used as a separate protein to rescue Sap1p secretion in strain SB11, and Sap1p activity of SB11 transformants

<table>
<thead>
<tr>
<th>Plasmid (encoded propeptide)</th>
<th>Sap propeptide as sequence</th>
<th>Sap1p activity of SB11 transformants (%)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>pSB42 (Sap1p propeptide)</td>
<td>...SPAEK...SGFVTIDPGVTRK...</td>
<td>100</td>
</tr>
<tr>
<td>pSB104 (Sap2p propeptide)</td>
<td>...TPPTYKR...SGFVTIDPGVTRK...</td>
<td>82 ± 4</td>
</tr>
<tr>
<td>pSB105 (Sap3p propeptide)</td>
<td>...TPPTYKR...SGFVTIDPGVTRK...</td>
<td>61 ± 5</td>
</tr>
<tr>
<td>pSB106 (Sap6p propeptide)</td>
<td>...APVEER...SGFVTIDPGVTRK...</td>
<td>78 ± 4</td>
</tr>
<tr>
<td>pSB107 (Sap22p propeptide)</td>
<td>...PGLPKFPGDPGIDPGVTRK...</td>
<td>40 ± 1</td>
</tr>
<tr>
<td>pSB108 (Sap1p propeptide)</td>
<td>...GTYFGIDPGIDPGVTRK...</td>
<td>44 ± 4</td>
</tr>
</tbody>
</table>

* The percentage values refer to the activity of SB11 transformed with pSB42 encoding the Sap1p propeptide. The value of 100% corresponds to 320 U ml⁻¹. The values shown are mean activities with standard deviation obtained from three experiments. No proteolytic activity was detected in SB11 culture supernatant.

The arrow indicates the F residue substituted by a D in strain SB269.
active part necessary for maturation and secretion of the enzyme. All SB11 transformants producing a peptide containing the 12 aa sequence GFVTLDFDVIKT secreted 80–100% of Sap1p activity in comparison to the action of the entire propeptide. Removing the G residue from the N-terminus or the T residue from the C-terminus of this 12 aa sequence reduced the amount of secreted activity by approximately one-third and one-half, respectively. Removing 2 or 3 aa from the N-terminus or from the C-terminus, respectively, prevented Sap1p secretion.

Maturation of Sap1p molecules in an intermolecular process

Following the results of the propeptide deletion analysis, a mutation was introduced in plasmid pSB10 to replace the phenylalanine (F) residue 27 of the SAP1 translation product, which is conserved in different Sap propeptides (Table 2), by an aspartic acid (D) residue. The newly generated plasmid was called pSB269 (Table 1). Direct expression of the mutated SAP1 gene in P. pastoris GS115 did not result in the secretion of Sap1p into the culture supernatant. However, a Sap1p product with an electrophoretic mobility higher than that of the mature form of Sap1p was detected in all transformants, from among which strain SB269 was selected (Table 1, Fig. 1).

The question whether the maturation of Sap1p occurs in vivo through inter- or intramolecular events was investigated by attempting to rescue the secretion of the enzyme produced in SB11 without propeptide, or in SB269 with a mutated non-functional propeptide, by a co-production of Sap1p with non-mutated propeptide. For such an experiment, a DNA fragment encoding Sap1p with a non-mutated prosequence and with a C-terminal His'-tag was cloned into pPICZαA. P. pastoris KM71 transformed with the newly generated plasmid pSB249 produced His'-tagged Sap1p (H₆-Sap1p; Fig. 5, lane 1) with a yield comparable to that of Sap1p produced by strain SB10 (Table 1). The specific activity of purified H₆-Sap1p was equal to that of Sap1p (9.5 U µg⁻¹). When SB10 was transformed by pSB249, both Sap1p and H₆-Sap1p were found, in a ratio close to 1:1 (90:100), in culture supernatant of all transformants (Fig. 5, lane 10). The total secreted proteolytic activity of

![Fig. 2. Chromosomal integration of pSB42 in SB11 transformants. pAOX1, P. pastoris alcohol oxidase gene (AOX1) promoter; proSAP1, part of SAP1 encoding the Sap1p propeptide; mSAP1, part of SAP1 encoding the mature part of Sap1p; BLE, S. hindustanus BLE gene for resistance to zeocin; HIS4, P. pastoris histidinol dehydrogenase gene; 3'AOX1, 3'AOX1 downstream sequence; B, BamHI; BglII, D, DraI; H, HindIII; N, NotI; S, SacI; X, XhoI.](image)
SB10 transformants was twice that of the SB10 strain. Untagged Sap1p could be separated from H<sub>e</sub>-Sap1p by filtration of the <i>P. pastoris</i> culture supernatant through a nickel chelating resin column (Fig. 5, lanes 11 and 12). No H<sub>e</sub>-Sap1p was detected in the culture supernatant after filtration through the column (Fig. 5, lane 12). Adsorbed H<sub>e</sub>-Sap1p could be subsequently released with imidazole for quantification.

When <i>P. pastoris</i> SB11 producing Sap1p without propeptide and SB269 producing Sap1p with a mutated propeptide were transformed by plasmid pSB249, Sap1p and H<sub>e</sub>-Sap1p were again both secreted in the culture supernatants of all transformants, but in a ratio close to 1:2 (54:100) (Fig. 5, lanes 4–9). Therefore, H<sub>e</sub>-Sap1p encoded by pSB249 permitted a restoration of secretion of Sap1p made without propeptide and with a mutated propeptide in the SB11 and SB269 transformants, respectively. However, this restoration occurred at the expense of secretion of H<sub>e</sub>-Sap1p since the proteolytic activity of SB11 and SB269 transformants was similar to that of <i>P. pastoris</i> KM71 transformed with pSB249, but not higher. Furthermore, an accumulation of H<sub>e</sub>-Sap1p product with an electrophoretic mobility higher than that of Sap1p was observed in cell extracts of SB11 and SB269 transformants but not in those of KM71 and SB10 transformants (Fig. 6).

**DISCUSSION**

When not covalently attached to the mature domain, only a 12 aa segment of the propeptide was necessary and sufficient to ensure complete Sap1p secretion. The
Yarrowia lipolytica prosequence was previously investigated with the fate of eukaryotic proproteinases mutated in their subtilisin molecules with unique point mutations distributed throughout the whole propeptide (in positions 76 to 13) did not yield active subtilisin (Lerner et al., 1991; Inouye, 1991).

The fate of eukaryotic proproteinases mutated in their prosequence was previously investigated with the Yarrowia lipolytica alkaline proteinase and the Rhizopus niveus aspartic proteinase I (Fabre et al., 1991; Fukuda et al., 1996). All deletions affecting the proregion of the Y. lipolytica alkaline proteinase precursor resulted in the intracellular accumulation of unprocessed protein. The rates of synthesis of one mutated and one prodeleted precursor of the R. niveus aspartic proteinase I were comparable in Saccharomyces cerevisiae, but both precursors were degraded in the endoplasmic reticulum. As a general rule, misfolded or unfolded proteins are selectively retained in the endoplasmic reticulum (Helenius et al., 1992), where a more or less rapid degradation occurs depending on the protein. The amount of Sap1p product detected intracellularly when Sap1p was produced with a mutated propeptide or without a propeptide was reduced, which suggests protein degradation (Fig. 1). It is likely that the non-functional folding of Sap1p results in the degradation of its mature domain, which consequently causes the lack of secretion of the Sap1p molecules. When SAP1 was expressed without its propeptide sequence an accumulation of Sap1p product with the same electrophoretic mobility as secreted Sap1p was observed in the cell. In contrast, an accumulation of a Sap1p product with an electrophoretic mobility higher than Sap1p was observed when SAP1 was expressed with a sequence encoding a mutated propeptide (Fig. 1). A similar His6-tagged product was detected in cell extracts of pSB249 transformants of SB11 and SB269 in parallel with the rescue of Sap1p secretion (Fig. 6, lanes 2 and 3). It appears that Sap1p produced without a propeptide and Sap1p produced with a mutated propeptide are not degraded in the same way.

The proteolytic activity of the SB11 and SB269 transformants secreting two Sap1ps produced with and without a functional propeptide under the control of two identical promoters was similar to that of KM71 transformed with pSB249 (Fig. 5). The lack of increased proteolytic activity in these transformants producing two different proteins can only be explained by a single-turnover catalytic property of the propeptide, meaning that each propeptide only participates in a single protein-folding reaction (Ohta et al., 1991; Baker et al., 1992).

If folding of the Sap1p molecules were obligatorily assisted by their propeptide in cis (first-order reaction, Fig. 7a), only H₆-Sap1p would have been secreted by the SB11 and SB269 transformants. Therefore, as our experiments demonstrate, there must be an intramolecular pathway to mature Sap1p without a priori exclusion of a possible intramolecular one. It was previously shown that prosubtilisin exists as dimers under non-denaturing conditions in vitro (Hu et al., 1996). A model of enzyme maturation was postulated where the prosequence of one prosubtilisin molecule is C. albicans Sap1p propeptide.

![Fig. 6. Detection of Sap1p and H₆-Sap1p intracellular products from strains KM71 (1), SB11 (2), SB269 (3) and SB10 (4) transformed with pSB249. The proteins in cell extracts from 5 µl of methanol-induced culture were separated by SDS-PAGE and analysed by Western blotting using anti-Sap polyclonal antibody (a) and anti-polyhistidine monoclonal antibody (b). Cell extracts of SB10 (5) and SB11 (6) strains and 5 µl of SB249 culture supernatant (*) were added on the gels for comparison of molecular masses of Sap1p products.](https://www.microbiologyresearch.org)
the template for the refolding of the mature sequence of the second one (second-order reaction). A strict intermolecular mechanism would predict a ratio of H$_6$-Sap1p to Sap1p of 1:1 in the culture supernatant of SB11 and SB269 transformants assuming that Sap1p and H$_6$-Sap1p are synthesized in a ratio of 1:1 (Fig. 7b). In fact, the ratio of H$_6$-Sap1p to Sap1p was experimentally determined to be close to 2:1. This result leads to the conclusion that maturation of Sap1p occurs in P. pastoris through both intra- and intermolecular pathways (Fig. 7). The observation of an intracellular accumulation of a H$_6$-Sap1p product in the SB11 and SB269 transformants (Fig. 6) is consistent with the prediction that H$_6$-Sap1p molecules do not mature without a propeptide or with a non-functional propeptide in heterodimers (Fig. 7).

This is the first report demonstrating that the maturation of an enzyme in vivo can be guided by the propeptide in an intermolecular process. The strategy developed in this work for Sap1p secretion could be used to test whether maturation of other proteinases via a combination of intra- and intermolecular pathways is a general pattern.

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