Characterization of proteinase–adhesin complexes of *Porphyromonas gingivalis*

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Proteinase–adhesin complexes of *Porphyromonas gingivalis* wild-type and RgpA and Kgp mutants were extracted using a Triton X-114 procedure and purified using arginine-affinity chromatography. The complexes were then characterized by peptide mass fingerprinting (PMF) and their equilibrium binding constants, immunogenicity and ability to induce protection as vaccines in the murine lesion model determined. The Triton X-114 procedure resulted in consistently higher yield and specific activity of the wild-type (wt) complex compared with that produced by the previously published sonication method. PMF and N-terminal sequencing of the purified wt complex showed that it consisted of the previously identified Arg-specific proteinase RgpA_{cat}, the Lys-specific proteinase Kgp_{cat} and adhesin domains RgpA_{A1}, RgpA_{A2}, RgpA_{A3}, KgpA_{A1} and KgpA_{A2}. However, analysis of the 30 kDa band in the wt complex, previously suggested to be RgpA_{A4}, indicated that this band contained C-terminally truncated KgpA_{A1} (which has an identical N-terminus to RgpA_{A4}) as well as the HagA_{A1}* adhesin. Analysis of the Triton X-114 extracted complexes from the *P. gingivalis* isogenic mutants *kgp* (RgpA complex) and *rgpA* (Kgp complex) suggested that the Kgp complex consisted of Kgp_{cat}, KgpA_{A1} and KgpA_{A2}/HagA_{A2} and that the RgpA complex consisted of RgpA_{cat}, RgpA_{A1}, HagA_{A1}*, RgpA_{A2} and RgpA_{A3}. Each of the complexes was found to have equilibrium binding constants (K_D) in the nanomolar range for fibrinogen, fibronectin, haemoglobin, collagen type V and laminin. However, the Triton-wt complex exhibited significantly lower K_D values for binding to each host protein compared with the sonication-wt complex, or the Triton-RgpA complex and Triton-Kgp complex. Furthermore, the Triton-wt complex induced a stronger antibody response to the A1 adhesins and tended to be more effective in providing protection in the mouse lesion model compared with the sonication-wt complex.

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INTRODUCTION

Chronic periodontitis is a bacterial-associated inflammatory disease characterized by the destruction of the supporting tissues of the teeth. This disease is a major public health problem, with an estimated 30 % of the adult dentate population affected, with severe forms affecting 5–10 %. The annual economic burden of the disease was estimated at US\$14.3 billion in the USA in 1999 (Brown *et al.*, 2000). Furthermore, chronic periodontitis may have other health implications, as several epidemiological and animal studies indicate an association between periodontitis and increased risk of cardiovascular diseases (Beck *et al.*, 1998; Jain *et al.*, 2003; Miyakawa *et al.*, 2004; Wu *et al.*, 2000), diabetes

(Pucher & Stewart, 2004), rheumatoid arthritis (Mercado *et al.*, 2001) and spontaneous preterm birth and preterm low birth weight (Lin *et al.*, 2003; Offenbacher *et al.*, 1998).

Colonization of the subgingival crevice by a consortium of Gram-negative bacteria with the development of a biofilm has been strongly associated with the onset and progression of chronic periodontitis (Socransky & Haffajee, 2000). Three bacterial species within the biofilm have been strongly associated with chronic periodontitis: *Porphyromonas gingivalis*, *Bacteroides forsythus* and *Treponema denticola* (Socransky *et al.*, 1998). Among these bacteria, *P. gingivalis* has received considerable attention as it has been more closely associated with disease severity (Socransky *et al.*, 1998). Furthermore, *P. gingivalis* has been shown to represent up to 50 % of the anaerobically cultivatable subgingival bacteria from active disease sites (Slots, 1982) and is rarely cultured from subjects without periodontitis (van Winkelhoff *et al.*, 2002). Furthermore, in animal models, subgingival implantation of *P. gingivalis* in mice (Baker

Abbreviations: ABM, adhesin-binding motif; Bz-L-Arg-pNA, benzoyl-L-Arg-p-nitroanilide; CBB, Coomassie brilliant blue; IFA, incomplete Freund's adjuvant; K_D , dissociation equilibrium constant; PMF, peptide mass fingerprinting; TFA, trifluoroacetic acid; TLCK, *N*- α -p-tosyl-L-lysine chloromethylketone; wt, wild-type; z-L-Lys-pNA, benzyloxycarbonyl-L-Lys-p-nitroanilide.

et al., 1994), primates (Persson *et al.*, 1994) and rats (Evans *et al.*, 1992; Rajapakse *et al.*, 2002) induces periodontal bone loss.

P. gingivalis has a number of putative virulence factors that contribute to its pathogenicity. Among them, the Arg- and Lys-specific cysteine proteinases, designated RgpA and Kgp, respectively, are considered major virulence factors as they have been suggested to play a key role in disease pathogenesis (reviewed by O'Brien-Simpson *et al.*, 2003). RgpA and Kgp are encoded by two genes, namely *rgpA* and *kgp* (Curtis *et al.*, 1999). The *rgpA* gene encodes a polyprotein consisting of an N-terminal preprofragment followed by a 45 kDa Arg-specific, calcium-stabilized cysteine proteinase, RgpA_{cat}, and four sequence-related adhesin domains: RgpA_{A1}, RgpA_{A2}, RgpA_{A3} and RgpA_{A4} (O'Brien-Simpson *et al.*, 2003). Similarly, the *kgp* gene encodes a polyprotein with a N-terminal preprofragment followed by a 48 kDa Lys-specific cysteine proteinase, Kgp_{cat}, and five C-terminal adhesin domains: Kgp_{A1}, Kgp_{A2}, Kgp_{A3}, Kgp_{A4} and Kgp_{A5} (O'Brien-Simpson *et al.*, 2003). Although the proteolytic domains of *rgpA* and *kgp* are divergent, with only 25 % identity, their C-terminal adhesin domains share high sequence similarities (Slakeski *et al.*, 1998). In fact, the A₂ adhesins RgpA_{A2} and Kgp_{A2} are identical in both gene products (Slakeski *et al.*, 1999).

Depending on the *P. gingivalis* strain, age of culture and purification method, several forms of RgpA and Kgp, both cell-associated and soluble, have been purified and characterized (reviewed by Potempa *et al.*, 2003). The Arg- and Lys-specific proteinases have been purified as monomeric proteins (Bedi & Williams, 1994; Chen *et al.*, 1992) and as multimeric complexes where the proteinase domains are non-covalently associated with a number of adhesin/haemagglutinin domains (Bhagal *et al.*, 1997; Pike *et al.*, 1994). Using sonication of exponentially growing *P. gingivalis* W50 cells and three chromatographic procedures, Bhagal *et al.* (1997) purified and characterized the major cell-associated Arg- and Lys-specific proteinases as a 300 kDa protein complex containing the Arg and Lys proteinases and their associated adhesins. It was proposed that, after the proteolytic processing of the RgpA and Kgp polyproteins into individual domains, they aggregate via adhesin-binding motifs (ABMs) to form the non-covalently associated cell-surface complex designated the RgpA-Kgp proteinase-adhesin complex (Bhagal *et al.*, 1997; Slakeski *et al.*, 1998). Recently, Takii *et al.* (2005), using sucrose monolaurate to extract *P. gingivalis* cells, confirmed the presence of this proteinase-adhesin complex and its component proteins from RgpA and Kgp, identified by Bhagal *et al.* (1997), but suggested that it existed as larger aggregates (660 kDa) in the form of 10 nm diameter particles on the cell surface. However, Takii *et al.* (2005) used antibodies to recombinant RgpA domains to identify the subunits of the complex and not mass spectrometry. The high sequence similarity of the adhesin domains requires a mass spectrometric approach for definitive identification. Soluble forms

of the Arg- and Lys-specific proteinases complexed with adhesin domains have been purified and characterized from culture supernatants, mainly from *P. gingivalis* strain HG66, which reportedly secretes the entire *rgpA* and *kgp* gene products into the culture media (reviewed by Potempa *et al.*, 2003).

The RgpA and Kgp proteinases and their associated adhesins are recognized as major virulence factors for *P. gingivalis* and therefore are of major interest to study in a purified form (Lamont & Jenkinson, 2000; O'Brien-Simpson *et al.*, 2003). However, the reported methodologies for extracting and purifying RgpA and Kgp as complexes from the cell involve a range of extraction methods and three or more chromatographic purification steps, with many of the purified products being poorly defined. Thus, the major objective of this study was to develop a simple and effective procedure for the extraction and purification of the RgpA and Kgp proteinase-adhesin complexes from *P. gingivalis* cells using Triton X-114 and then to comprehensively characterize the complexes for their subunits using mass spectrometry, as vaccines in an animal model and for their binding properties to a range of host proteins. The non-ionic detergent Triton X-114 was used to extract the complexes as it is a 'mild' detergent that solubilizes membrane-associated proteins while retaining native protein structure (Bordier, 1981) and has been used effectively to extract outer-membrane proteins from a range of bacteria (Brooks *et al.*, 2006; Cullen *et al.*, 2002; Radolf *et al.*, 1988).

METHODS

Bacterial strains and growth condition. *Porphyromonas gingivalis* W50 isogenic mutants lacking either the *rgpA* gene product (W501) or the *kgp* gene product (K1A) as well as *P. gingivalis* W50 wild-type were grown in an anaerobic chamber (MK3 anaerobic workstation; Don Whitley Scientific) at 37 °C on horse blood agar plates supplemented with 10 % (v/v) lysed horse blood and 10 µg erythromycin ml⁻¹ (for the mutant strains). Bacterial colonies were used to inoculate brain heart infusion medium containing 5 µg haemin ml⁻¹ and 0.5 µg cysteine ml⁻¹. Batch culture growth was monitored at 650 nm using a spectrophotometer (Perkin-Elmer model 295E). Culture purity was routinely checked by Gram stain, microscopic examination and using a variety of biochemical tests according to Slots (1982).

Extraction of the proteinase-adhesin complexes. *P. gingivalis* strains W50 wild-type, W501 and K1A were grown to late exponential phase (in 2 l cultures), harvested by centrifugation (7500 g, 30 min, 4 °C) and washed twice with PG buffer (50 mM Tris/HCl, 150 mM NaCl, 5 mM CaCl₂ and 5 mM cysteine-HCl, pH 8.0) in the anaerobic workstation. Cells were either (1) resuspended in PG buffer, total volume 30 ml, containing 0.5 % (v/v) Triton X114 and gently mixed at room temperature for 45 min (Triton complex) or (2) resuspended in 30 ml PG buffer and subjected to mild sonication using a Branson sonifier 250 with an output control of 3 and a 50 % duty cycle (sonication complex). The cell extract was centrifuged (7500 g, 30 min, 4 °C) and the collected supernatant centrifuged (40000 g, 30 min, 4 °C). The supernatant was then filtered (0.2 µm) and the complexes were purified by arginine (Arg)-affinity chromatography. FPLC was performed at room temperature at a flow rate of 1.0 ml min⁻¹. The *P. gingivalis* extracts were applied to

an Arg-Sepharose column (Hiload XK 16/10 Q, Pharmacia), installed in a Pharmacia GP-250 FPLC system, in TC 50 buffer (buffer A) (50 mM Tris/HCl, 50 mM NaCl, 5 mM CaCl₂, pH 7.4) at a flow rate of 1 ml min⁻¹. Non-specifically bound proteins were eluted with a linear gradient of 0–40% TC 50 buffer containing, 500 mM NaCl, 50 mM Tris/HCl, 5 mM CaCl₂, pH 7.4 (buffer B) at a flow rate of 1.0 ml min⁻¹. The column was reequilibrated with buffer A and bound proteins eluted with TC 50 buffer containing 500 mM arginine, pH 7.4 at a flow rate of 1 ml min⁻¹. The eluate was monitored at 280 nm. All fractions were collected at 4 °C and stored at -70 °C before further processing.

Arginine-eluted FPLC fractions were concentrated using Vivaspin 20 concentrators (Vivascience) by centrifugation at 3000 g for 15 min periods at 4 °C until the eluate was reduced to a volume of approximately 1 ml. The filter membrane of the Vivaspin 20 concentrator was then rinsed with 1 ml TC 50 buffer. Concentrated fractions were then desalted using a PD-10 column (Amersham Pharmacia Biotech), equilibrated with TC 50 buffer and the proteins were eluted with the same buffer as per the manufacturer's instructions.

Arg-specific and Lys-specific proteinase assays. Benzoyl-L-Arg-*p*-nitroanilide (Bz-L-Arg-*p*NA) (Sigma) and benzyloxycarbonyl-L-Lys-*p*-nitroanilide (z-L-Lys-*p*NA) (Novabiochem) were used to assay FPLC fractions for Arg- and Lys-specific proteolytic activity, respectively. Samples of each chromatographic fraction were diluted in TC 50 buffer (total volume 360 µl) and incubated for 10 min at 37 °C with 40 µl 100 mM cysteine, pH 8. After incubation, 400 µl of either Bz-L-Arg-*p*NA or z-L-Lys-*p*NA substrate (2 mM Bz-L-Arg-*p*NA or 2 mM z-L-Lys-*p*NA) dissolved in 3 ml propan-2-ol and mixed with 7 ml enzyme buffer (400 mM Tris/HCl, 100 mM NaCl and 20 mM cysteine, pH 8) was added. The proteolytic activity was determined by measuring the absorbance at 410 nm for 3 min at 10 s intervals using a diode array spectrophotometer model 8452A (Hewlett Packard).

Size-exclusion chromatography. This was performed using a macrosphere GPC 300Å column (7 µm, 250 × 4.6 mm, with exclusion limits of 7500–1 200 000 Da; Alltech) installed in a Waters Delta 600 HPLC system. Chromatography was performed at a flow rate of 0.5 ml min⁻¹ in 0.05 M KH₂PO₄ containing 0.15 M Na₂SO₄ (pH 7.0). Material eluted from the column was detected by determining absorbance at 280 nm. A standard curve of molecular mass gel filtration standards (Amersham Pharmacia Biotech) was used to determine the molecular mass of the eluted fractions.

SDS-PAGE. SDS-PAGE was performed on FPLC fractions by using a Novex electrophoresis system with Novex 12% (w/v) Tris/glycine precast mini gels (Invitrogen). Protein samples (10 or 30 µg) were precipitated by addition of trichloroacetic acid (TCA) to a final concentration of 10% (v/v) and incubated 20 min at 4 °C. Precipitated proteins were collected by centrifugation (10 min, 16 000 g), resuspended in 20 µl reducing sample buffer [10% (w/v) SDS, 0.05% (w/v) bromophenol blue, 25% (v/v) glycerol and 0.05% (v/v) 2-mercaptoethanol] and the pH adjusted with the addition of 10 µl of 1.5 M Tris/HCl, pH 8.0, and then heated for 5 min at 100 °C. Samples were loaded onto the precast gels and electrophoresis was performed using a current of 30–50 mA and a potential difference of 125 V. After completion of electrophoresis, the gels were fixed in destain [45% (v/v) methanol, 10% (v/v) acetic acid] for 3 min at room temperature. For Coomassie blue staining, gels were placed in Coomassie brilliant blue (CBB) [0.1% (w/v) CBB R-250, 15% (v/v) ethanol, 2.5% (v/v) acetic acid], heated in a microwave until boiling and then allowed to cool for 5 min. The stain was removed and destain was added and heated in a microwave until boiling and allowed to cool for 5 min. Protein bands were visualized by rinsing gels in Milli Q water overnight.

N-terminal sequence analysis. For sequence analysis, proteins were transferred onto a PVDF membrane (Problott, Applied Biosystems) using a transblot cell (Bio-Rad). The PVDF membrane was wetted in 100% methanol and soaked in transfer buffer [10 mM CAPS, 10% (v/v) methanol, pH 11.5] for 1 min. Transfer was performed using a potential difference of 60 V for 90 min. Membranes were stained with CBB for 30 s and destained in 50% (v/v) aqueous methanol. Protein bands were excised and N-terminal sequences determined using a Hewlett Packard 10005A protein sequencer as previously described (Bhogal *et al.*, 1997).

In-gel digestion of proteins separated by SDS-PAGE and peptide mass fingerprinting. CBB-stained protein bands from SDS-PAGE were excised and subjected to in-gel trypsin digestion and subsequent peptide extraction. Protein bands were excised from the CBB-stained SDS-PAGE gel and gel pieces were washed in 50 mM NH₄HCO₃/ethanol 1:1 (v/v), reduced and alkylated with DTT and iodoacetamide, respectively, and digested with sequencing-grade modified trypsin (10 ng µl⁻¹) (Promega) overnight at 37 °C as previously published (Mortz *et al.*, 1994). The peptide extract containing 25 mM NH₄HCO₃ was then analysed by mass spectrometry using an Ultraflex TOF/TOF instrument (Bruker Daltonics) in positive ion and reflectron mode. A saturated solution of 4-hydroxy- α -cyanocinnamic acid was prepared in 97:3 (v/v) acetone/0.1% aqueous trifluoroacetic acid (TFA). A thin layer was prepared by pipetting and immediately removing 2 µl of this solution onto the 600 µm anchochips of the target plate. Sample (0.5 µl) was deposited on the thin layers with 2.5 µl of 0.1% (v/v) TFA, and allowed to adsorb for 5 min, after which the sample solution was removed, and the thin layers washed once with 10 µl ice-cold 0.1% (v/v) TFA for 1 min. Spectra were calibrated by close external calibration using a standard peptide mix (Bruker Daltonics). MS/MS spectra were acquired using the 'Lift' technique (Suckau *et al.*, 2003). Proteins were identified by peptide mass fingerprinting (PMF) and MS/MS ions searches against the *P. gingivalis* database (available from <http://www.tigr.org>) using an in-house Mascot search engine (Matrix Science).

Immunization and mouse lesion model. The mouse lesion model protocols were approved by the University of Melbourne Ethics Committee for Animal Experimentation. BALB/c mice 6–8 weeks old (10 mice per group) were immunized subcutaneously (s.c., 100 µl) with 25 µg complex in phosphate-buffered saline (PBS, pH 7.4) emulsified in incomplete Freund's adjuvant (IFA). After 30 days, mice were boosted with 25 µg antigen (s.c., 100 µl, emulsified in IFA) and then 12 days later bled from the retrobulbar plexus. Two days after bleeding, mice were challenged with 7.5 × 10⁹ viable cells of *P. gingivalis* strain ATCC 33277 by s.c. injection (100 µl) in the abdomen, and lesion development measured over 14 days. The *P. gingivalis* inocula were prepared in PG buffer in the anaerobic workstation as described above. The number of viable cells in the inocula was verified by enumeration on horse blood agar plates. Lesion sizes were statistically analysed using the Kruskal–Wallis test and the Mann–Whitney *U*-Wilcoxon rank sum test with a Bonferroni correction for type 1 error. Effect sizes, represented as Cohen's *d*, were calculated using the effect size calculator provided on-line by Evidence-Based Education UK web site at <http://www.cemcentre.org/>. According to Cohen (1969) a small effect size is 0.2 ≤ *d* < 0.05, moderate 0.5 ≤ *d* < 0.8 and large *d* ≥ 0.8.

ELISA. ELISAs were performed in triplicate in wells of flat-bottom polyvinyl microtitre plates (Dynatech Laboratories) coated with 10 µg ml⁻¹ Triton-wt complex or sonication-wt complex in PBS, pH 7.4, overnight at 4 °C. After removal of the coating solution, 2% (w/v) skim milk powder in PBS, pH 7.4, containing 0.1% (v/v) Tween 20 was added to wells to block the uncoated plastic for 1 h at room temperature. After washing four times with washing buffer [PBS containing 0.1% (v/v) Tween 20, pH 7.4 or (PBST)], serial

dilutions of mouse sera in PBST containing 0.5% (w/v) skim milk (SK-PBST) were added to each well and incubated for 16 h at room temperature. After washing six times with PBST, a 1/2000 dilution of goat antisera to mouse IgM, IgA, IgG1, IgG2a, IgG2b, or IgG3 (Sigma) was added in SK-PBST and allowed to bind for 2 h at room temperature. Plates were washed six times in PBST and a 1/5000 dilution of horseradish peroxidase-conjugated rabbit anti-goat immunoglobulin (Sigma) in SK-PBST was added to each well and incubated for 1 h at room temperature. After washing (six times, PBST), bound antibody was detected by the addition of 100 µl ABTS substrate [0.9 mM 2,2'-azino-bis (3-ethylbenz-thiazoline-6) sulfonic acid in 80 mM citric acid containing 0.005% (v/v) hydrogen peroxide, pH 4.0] to each well. The A_{415} was measured using a Bio-Rad microplate reader model 450.

Western blot analysis using mouse protective sera. Triton-wt complex and sonication-wt complex (15 µg) were subjected to SDS-PAGE and electrophoretically transferred onto a PVDF membrane as described above. The PVDF membrane was blocked with skim milk powder in TN buffer (50 mM Tris/HCl, 100 mM NaCl, pH 7.4) for 2 h at room temperature. The membrane was then incubated with anti-wt complex sera at a dilution of 1/25 in TN buffer, overnight at room temperature. After the incubation, the membrane was washed four times in TN buffer containing 0.05% (v/v) Tween 20 and incubated with a 1/200 dilution of rabbit anti-mouse horseradish peroxidase-conjugated IgG (Sigma) for 2 h at room temperature. The membrane was washed four times with TN buffer containing 0.05% (v/v) Tween 20 and the bound antibodies were detected by 0.005% (w/v) 4-chloro-1-naphthol in TN buffer containing 16.6% (v/v) methanol and 0.015% (v/v) hydrogen peroxide (30% solution).

Analysis of wt complex, RgpA complex and Kgp complex binding to host proteins. Prior to the ELISA, the proteolytic activity of each proteinase–adhesin complex was inhibited by incubation with 5 mM *N*- α -tosyl-lysine chloromethylketone (TLCK; Sigma) for 30 min at room temperature. Excess TLCK was removed by using a PD-10 column equilibrated with TC 50 buffer with the proteins eluted according to the manufacturer's instructions in the same buffer (Amersham Pharmacia Biotech). ELISAs were performed in triplicate using a solution of 5 µg ml⁻¹ of either fibrinogen, fibronectin, collagen type V, laminin or haemoglobin in PBS, pH 7.4, to coat wells of flat-bottomed polyvinyl microtitre plates (Microtitre, Dynatech Laboratories) overnight at room temperature. After removing the coating solution, remaining uncoated plastic was blocked with 2% (w/v) skim milk powder in PBST for 1 h at room temperature. After washing (4 × PBST), equimolar amounts of TLCK-treated Triton-wt complex, sonication-wt complex, RgpA

complex or Kgp complex in SK-PBST were added to the wells and incubated at room temperature for 2 h. After washing (6 × PBST) a 1/5000 dilution of anti-complex sera in SK-PBST was added to each well and incubated 2 h at room temperature. After washing (6 × PBST), bound antibody was detected by incubation with a 1/5000 dilution of horseradish peroxidase-conjugated anti-mouse immunoglobulin developed in goat (Sigma) in SK-PBST for 1 h at room temperature. Wells were then washed (6 × PBST) and the plates were developed as described above.

ELISA results were used to derive the dissociation equilibrium constant (K_D) based on an analogue of the Michaelis–Menten equation (Qiu *et al.*, 1996): $A = A_{\max}[\text{protein}]/K_D + [\text{protein}]$, where A is the absorbance at a given protein concentration, $A_{\max}$ is the maximum absorbance for the ELISA plate reader, $[\text{protein}]$ is the protein concentration of the analyte and K_D is the dissociation equilibrium constant for a given absorbance at a given protein concentration (ELISA data point). K_D was calculated by a Scatchard analysis of $1/[\text{protein}]$ vs $1/K_D$ (ELISA data point): $1/K_D = m(1/c)$, where m is the slope of the line and c is the intercept at the y axis, with an R^2 value > 0.99.

RESULTS

Characterization of the proteinase–adhesin complexes extracted using Triton X-114 or sonication and purified by Arg-affinity chromatography

The proteinase–adhesin complex was extracted from *P. gingivalis* strain W50 using Triton X-114 (Triton-wt complex) and the previously described method of sonication (sonication-wt complex) (Bhagal *et al.*, 1997). Triton X-114 extraction was consistently more effective in releasing cell-surface protein compared with the sonication method, as the Triton X-114 extraction method produced 70.0 ± 9.2 mg protein from 1 l culture. Sonication produced only 10.3 ± 2.5 mg protein (Table 1). Furthermore, the crude Triton X-114 extract had consistently higher (~threefold) Arg and Lys proteolytic activity compared with the sonication extract (Table 1). The crude Triton X-114 and sonication extracts were subjected to Arg-affinity chromatography and a representative chromatogram is shown in Fig. 1. Non-specifically bound proteins were removed

Table 1. Purification of the wt-complex from Triton and sonication extracts.

	Total protein (mg)	Arg proteolytic activity (U)*	Lys proteolytic activity (U)†	Arg-specific activity (U mg ⁻¹)	Lys-specific activity (U mg ⁻¹)	Purification (-fold)	
						Arg	Lys
Crude cell sonicate‡	10.34 ± 2.52	13.60 ± 2.30	1.89 ± 0.78	1.30 ± 0.62	0.18 ± 0.09	1.00	1.00
Sonication-wt complex	0.72 ± 0.15	1.23 ± 0.24	0.09 ± 0.02	1.72 ± 0.84	0.12 ± 0.06	1.32	0.69
Crude Triton X-114 extract†	70.00 ± 9.23	35.00 ± 4.32	4.56 ± 1.35	0.50 ± 0.22	0.06 ± 0.01	1.00	1.00
Triton-wt complex	0.60 ± 0.12	2.45 ± 0.68	1.19 ± 0.86	4.11 ± 1.24	1.97 ± 0.56	8.22	32.83

*Amidolytic activity using 2.0 mM Bz-L-Arg-pNA: 1 unit (U) = 1 µmol min⁻¹ at 37 °C.

†Amidolytic activity using 2.0 mM z-L-Lys-pNA: 1 unit (U) = 1 µmol min⁻¹ at 37 °C.

‡*P. gingivalis* W50 Triton X-114 extract or cell sonicate was subjected to FPLC and the proteinase–adhesin complexes purified as described in Methods.

Fig. 1. Typical Arg-Sepharose FPLC chromatogram of *P. gingivalis* TX-114 extract and cell sonication. *P. gingivalis* Triton X-114 and sonication extracts were applied to an Arg-Sepharose column installed in a Pharmacia FPLC system. Unbound proteins (peak A) were eluted in TC50 buffer at a flow rate of 1 ml min⁻¹. Non-specifically bound proteins (peak B) were eluted with a linear gradient of 0–40% TC 500 buffer (500 mM NaCl, 50 mM Tris/HCl, 5 mM CaCl₂, pH 7.4) at a flow rate of 1.0 ml min⁻¹. The RgpA-Kgp-HagA complex (peak C) was eluted with TC 50-Arg buffer (500 mM arginine, 50 mM NaCl, 50 mM Tris/HCl, 5 mM CaCl₂, pH 7.4) at a flow rate of 1 ml min⁻¹. The arrows indicate the start of each step gradient.

with a linear gradient of 0–250 mM NaCl (peak B). The proteinase–adhesin complex was then eluted with 500 mM arginine (peak C) which was collected, concentrated using a VivaSpin-10 concentrator (Sartorius), de-salted using a PD-10 column and then subjected to SDS-PAGE. Fig. 2 shows the SDS-PAGE analysis of the 500 mM arginine eluate (peak C) from the Triton X-114 extract (lane 1), with the typical banding pattern associated with the Arg and Lys proteinase–adhesin complex purified using sonication (Bhogal *et al.*, 1997). The N-terminal sequences of the 48, 45, 44, 39, 30, 17 and 15 kDa bands of the Triton-wt complex were determined by Edman sequence analysis of electrophoretically transferred proteins onto a PVDF blot (Table 2). This analysis confirmed the presence of the Kgp_{cat} (48 kDa band), RgpA_{cat} (45 kDa band), RgpA_{A1} (44 kDa band), Kgp_{A1} (39 kDa band), RgpA_{A2}/Kgp_{A2}/HagA_{A2} (17 kDa band) and the RgpA_{A3} (15 kDa band). The 30 kDa band contained two distinct N-terminal sequences; the predominant sequence was ANEAKVVLADN, which is consistent with the N-terminal sequences of RgpA_{A4} and Kgp_{A1}, and APAPYQER, which is consistent with the N-terminal sequence of HagA_{A1}* (Table 2). To further characterize the proteins in the Triton-wt complex, the protein bands were excised from a Tris-glycine SDS-PAGE gel, trypsin digested and identified by PMF. Protein bands running at approximately 48, 45, 44, 39, 17 and 15 kDa were confirmed as Kgp_{cat}, RgpA_{cat}, RgpA_{A1}, Kgp_{A1}, RgpA_{A2}/Kgp_{A2}/HagA_{A2} and RgpA_{A3}, respectively, by PMF analysis

Fig. 2. SDS-PAGE of wt complex, RgpA complex and Kgp complex. The Triton X-114 extracted wt complex, RgpA complex and Kgp complex were subjected to SDS-PAGE and protein bands visualized by Coomassie blue staining. Lanes: 1, wt complex; 2, RgpA complex; 3, Kgp complex. Molecular mass markers (Pharmacia) are indicated in kDa. Protein bands were transferred onto a PVDF membrane and identified by N-terminal sequence analysis, using a Hewlett Packard 10005A protein sequencer. The designation of each protein band identified by N-terminal sequencing is shown.

(Table 3). The band at 30 kDa was found to consist of two proteins. Seven peptides obtained from the band corresponded to Kgp_{A1} and four obtained corresponded to HagA_{A1}* with the sequence of the 1094.56 Da peptide confirmed by MS/MS fragmentation analysis as the HagA_{A1}* sequence HFGCTGIFR (Table 3). In addition, higher molecular mass bands (55 kDa and 97 kDa) were obtained in the SDS-PAGE analysis of the Triton X-114 complex (Fig. 2) that were demonstrated to be partially processed forms of RgpA and Kgp by PMF analysis (data not shown). The Triton X-114 extracted wt complex (Triton-wt complex) had >twofold Arg-specific and >15-fold Lys-specific proteolytic activity compared with the sonication-wt complex (Table 1). N-terminal and PMF analysis of the sonication-wt complex, which produced the same pattern of bands as the Triton-wt complex upon SDS-PAGE, confirmed the presence of the same subunits of the complex except that HagA_{A1}* could not be found in the 30 kDa band. Furthermore, the bands for Kgp_{cat} and RgpA_{cat} were more intense in the Triton-wt complex, consistent with the higher specific Arg and Lys activity.

Characterization of the Kgp complex and RgpA complex extracted using Triton X-114

The individual Kgp complex and RgpA complex were purified by Arg-affinity chromatography from Triton X-114 extracts of *P. gingivalis* W50 *rgpA* mutant (W501) and *kgp* mutant (KIA), respectively. Both the Kgp complex and RgpA complex were subjected to SDS-PAGE (Fig. 2), transferred onto a PVDF membrane and N-terminal sequences of the protein bands determined. In the RgpA complex, bands at 45, 44, 17 and 15 kDa were identified as RgpA_{cat},

Table 2. N-terminal processing of the Triton-wt complex, RgpA complex and Kgp complex protein domains

Cleavage sites are indicated by vertical arrows.

Mol. mass (kDa) determined by SDS-PAGE	Protein domain	N-terminal sequence		
		Triton-wt complex	RgpA complex	Kgp complex
48	Kgp _{cat}	↓ R-DVYTDHGDLYN-	—*	↓ R-DVYTDHGDLYN-
45	RgpA _{cat}	↓ R-YTPVEEKQNGRM-	↓ R-YTPVEEKQNGRM-	—*
44	RgpA _{A1}	↓ R-SGQAEIVLEAHDVWN-	↓ R-SGQAEIVLEAHDVWN-	—*
39	Kgp _{A1}	↓ R-ANEAKVVLAADN-	—*	↓ R-ANEAKVVLAADN-
30	Kgp _{A1}	↓ R-ANEAKVVLAADN-	—*	†
30	HagA _{A1} *	↓ K-APAPYQER-	†	—*
17	RgpA _{A2} /KgpA ₂ /HagA _{A2}	↓ R-ADFTETFESSTH-	↓ R-ADFTETFESSTH-	↓ R-ADFTETFESSTH-
15	RgpA _{A3}	↓ K-PQSVWIERT-	↓ R-EADGAKPQSVWIERT-	—*

*Sequence not present.

†Protein identified by PMF.

RgpA_{A1}, RgpA_{A2} and RgpA_{A3}, respectively (Table 2). In the Kgp complex, protein bands at 48, 39 and 17 kDa were identified as Kgp_{cat}, Kgp_{A1} and Kgp_{A2}/HagA_{A2}, respectively. The band at 30 kDa in the RgpA complex was identified by PMF analysis as HagA_{A1}*. The N-terminal sequences of the proteinases and adhesins in the RgpA complex and Kgp complex, with the exception of RgpA_{A3}, were the same as found in the wt complex and all were preceded by an arginine residue in the original polyprotein sequence from which they were processed (Table 2). The N-terminal sequence of the RgpA_{A3} adhesin, which in the wt complex is PQSVWIERT, is preceded by a lysine residue, whereas the N-terminus of the RgpA_{A3} adhesin in the RgpA complex extracted from the *kgp* mutant was extended by six residues, with the sequence EADGAKPQSVWIERT and is preceded by an arginine residue (Table 2).

The Arg- and Lys-specific proteinase activities of the purified RgpA complex and Kgp complex were measured. The RgpA complex displayed 1.85 units mg⁻¹ of Arg-specific proteolytic activity but no Lys-specific proteolytic activity (Table 4). The Kgp complex exhibited 0.66 units mg⁻¹ of Lys-specific proteolytic activity and no Arg-specific activity (Table 4).

Size-exclusion chromatography of the wt complex, RgpA complex and Kgp complex

The purified Triton-wt complex, Kgp complex and RgpA complex were further characterized by size-exclusion chromatography using a Macrosphere 300Å column (Alltech). The Triton-wt complex eluted as two products, with the major product eluting in the void volume corresponding to a molecular mass of $>1.2 \times 10^6$ and a

second peak eluted at 5.63 min corresponding to a molecular mass of 260 kDa (Fig. 3a). The chromatographic profiles attained for purified Triton-wt complex and sonication-wt complex were similar, with the same retention times and area distribution (data not shown).

The RgpA complex eluted as two products, with the major product eluting in the void volume corresponding to high molecular mass of $>1.2 \times 10^6$ and a second peak eluted at 5.68 min corresponding to a molecular mass of approximately 150 kDa (Fig. 3b). Similarly, the Kgp complex eluted as two products, with the major product eluting in the void volume corresponding to a molecular mass of $>1.2 \times 10^6$ and a second peak eluted at 5.69 min corresponding to a molecular mass of approximately 120 kDa (Fig. 3c).

Immunogenicity of the wt complex

To characterize the immunogenicity of the Triton-wt complex and the sonication-wt complex, BALB/c mice were immunized (day 0 and day 30) with either sonication-wt complex or Triton-wt complex emulsified in incomplete IFA or with adjuvant alone. Twelve days after the second immunization, mice were challenged subcutaneously with *P. gingivalis* strain ATCC 33277. The development of lesions was monitored for the next 14 days for each group. Fig. 4 shows that the mean lesion size of mice immunized with purified Triton-wt complex or sonication-wt complex was significantly ($P < 0.01$; $P < 0.05$, respectively) smaller than that of the PBS/IFA control group. The Triton-wt complex, when used as a vaccine, tended to be more effective ($d = 1.85$, 99% CI: 3.07, 0.47) in providing protection as indicated by a larger effect size compared with the sonication-wt complex ($d = 1.32$, 95% CI: 2.28, 0.07).

Table 3. Peptide mass fingerprinting identification of the proteins in the Arg-affinity purified Triton X114 extracted wt complex

Protein band (k)	Designation	Identifying peptide*	Observed mass (Da)	Calculated mass (Da)
48	Kgp _{cat}	²²⁹ DVYTDHGDLYNTPVR ²⁴³	1764·02	1763·81
		²⁵⁴ EALKPWLTWK ²⁶³	1270·86	1270·71
		²⁶⁷ GFYLDVHYTDEAEVGTTNASIK ²⁸⁸	2429·44	2429·14
		²⁹⁵ YNDGLAASAAPVFLALVGDTDVISGEK ³²¹	2692·71	2692·36
		³²⁸ VTDLYYSAVDGDYFPEMYTFR ³⁴⁸	2551·43	2551·13
		³⁴⁹ MSASSPEELTNIIDK ³⁶³	1634·00	1633·79
		³⁸¹ VLLIAGADYSWNSQVGQPTIK ⁴⁰¹	2259·47	2259·19
		⁴⁰² YGMQYYNQEHGYTDVYNLYK ⁴²²	2711·57	2711·16
		⁶⁰² TNTYTLPASLPQNQASYSIQASAGSYVAISK ⁶³²	3231·08	3230·61
		⁶³³ DGVLYGTGVANASGVATVSMTK ⁶⁵⁴	2097·30	2097·04
		⁶⁵⁵ QITENGNYDVVITR ⁶⁶⁸	1620·99	1620·81
		⁶⁷⁷ QIQVGEPSPYQPVSNLTTATTQGGQK ⁷⁰⁰	2570·62	2570·30
45	RgpA _{cat}	²²⁸ YTPVEEK ²³⁴	864·44	864·42
		²⁶⁹ VAEDIASPVTANAIQQFVK ²⁸⁷	2000·19	2000·06
		²⁹³ EGNDLTYVLLIGDHK ³⁰⁷	1685·98	1685·86
		³¹⁹ SDQVYQGIVGNDHYNEVFIGR ³³⁹	2409·24	2409·13
		⁴¹² CYDPGVTPK ⁴²⁰	1035·53	1035·47
		⁴²¹ NIIDAFNGGISLANYTGHGSETAWGTSHFGTTHVK ⁴⁵⁵	3660·11	3659·74
		⁴⁹³ DGKPTGTVAIIASTINQSWASPMR ⁵¹⁶	2500·40	2500·27
		⁵¹⁷ GQDEMNEILCEK ⁵²⁸	1464·73	1464·62
		⁵³⁶ TFGGVTMNGMFAMVEK ⁵⁵¹	1718·90	1718·78
		⁵⁵⁹ MLDTWTVFGDPSLLVR ⁵⁷⁴	1849·06	1848·94
44	RgpA _{A1}	⁸²⁰ IWIAGQGPTK ⁸²⁹	1069·69	1069·59
		⁸³⁰ EDDYVFEAGK ⁸³⁹	1171·62	1171·61
		⁸⁴¹ YHFLMKK ⁸⁴⁷	966·59	966·22
		⁸⁴⁸ MGSGDGTTELITSEGGGSDYTYTVYR ⁸⁷²	2615·31	2615·30
		⁸⁷⁹ EGLTATTFEEDGVAAGNHEYCVEVK ⁹⁰³	2725·43	2725·42
		⁹¹⁵ DVTVEGSNEFAPVQNLTGSAVGQK ⁹³⁸	2446·39	2446·38
39	Kgp _{A1}	⁸³⁸ MWIAGDGGNQPAP ⁸⁵⁰	1372·80	1372·53
		⁸⁵¹ YDDFTFEAGK ⁸⁶⁰	1192·65	1192·25
		⁸⁵¹ YDDFTFEAGKK ⁸⁶¹	1320·77	1320·42
		⁸⁶¹ KYTFTMR ⁸⁶⁷	946·57	946·14
		⁸⁶⁹ AGMGDGTDMEEVDDSPASYTYTVYR ⁸⁹³	2730·00	2730·88
		⁸⁹⁸ IKEGLTATTFEEDGVAAGNHEYCVEVK ⁹²⁴	2967·70	2968·25
		⁹⁰⁰ EGLTATTFEEDGVAAGNHEYCVEVK ⁹²⁴	2726·54	2726·91
		⁹³⁶ DVTVEGSNEFAPVQNLTGSSVGQK ⁹⁵⁸	2463·48	2463·64
30	HagA _{A1} *	¹¹² HFGCTGIFR ¹²⁰ (confirmed by LIFT ms/ms)	1094·56	1094·26
		⁹⁵ TIDLSAYAGQQVYLAFR ¹¹¹	1916·57	1916·16
		¹⁸⁶ DVTVEGSNEFAPVQNLTGSAVGQK ²⁰⁹	2447·92	2447·64
		¹²¹ LYLDDVAVSGEGSSNDYTYTVYR ¹⁴³	2587·85	2587·74
30	Kgp _{A1}	⁸³⁸ MWIAGDGGNQPAP ⁸⁵⁰	1372·63	1372·53
		⁸⁵¹ YDDFTFEAGKKYTFTMR ⁸⁶⁷	2121·86	2120·37
		⁸⁵¹ YDDFTFEAGK ⁸⁶⁰	1192·65	1192·25
		⁸⁵¹ YDDFTFEAGKK ⁸⁶¹	1320·77	1320·42
		⁸⁶⁹ RAGMGDGTDMEEVDDSPASYTYTVYRDGK ⁸⁹⁷	3287·55	3288·49
		⁹⁰⁰ EGLTATTFEEDGVAAGNHEYCVEVK ⁹²⁴	2726·22	2726·91
		¹¹²⁰ TVDLPAGTKYVAFR ¹¹³³	1536·67	1537·78
17	RgpA _{A2} /KgpA ₂ /HagA _{A2}	¹²³⁸ TGTNAGDFTVVFEEPTNGIN ¹²⁵⁷	2083·18	2082·96
		¹²¹⁸ YYYAVNDGFPGDHYAVMISK ¹²³⁷	2310·23	2310·05
15	RgpA _{A3}	¹²⁷⁴ PQSVWIER ¹²⁸¹	1014·59	1014·15
		¹³³⁴ IKEGLTETTFEEDGVATGNHEYCVEVK ¹³⁶⁰	3055·65	3056·31
		¹³⁷⁰ CVNVTVNSTQFNVPK ¹³⁸⁴ (confirmed by LIFT ms/ms)	1706·97	1706·94
		¹³⁶⁹ KCVNVTVNSTQFNVPK ¹³⁸⁴	1835·09	1835·11

Peptide fragments confirmed by PMF.

Table 4. Purification of the RgpA complex and Kgp complex

	Total protein (mg)	Arg proteolytic activity (U)*	Lys proteolytic activity (U)†	Arg-specific activity (U mg ⁻¹)	Lys-specific activity (U mg ⁻¹)	Purification (-fold)	
						Arg	Lys
Crude <i>kgp</i> mutant Triton extract‡	57.9±5.95	71.40±9.40	0	1.23±0.19	0	1.00	0
RgpA complex purified from the <i>kgp</i> mutant	2.54±0.56	4.68±1.89	0	1.85±0.89	0	1.50	–
Crude <i>rgpA</i> mutant Triton extract‡	40.2±8.60	10.20±2.40	6.00±1.40	0.26±0.04	0.16±0.09	–	1.00
Kgp complex purified from the <i>rgpA</i> mutant	1.11±0.63	0	0.73±0.09	0	0.66±0.23	–	4.13

*Amidolytic activity using 2.0 mM Bz-L-Arg-pNA: 1 unit (U)=1 µmol min⁻¹ at 37 °C.
†Amidolytic activity using 2.0 mM z-L-Lys-pNA: 1 unit (U)=1 µmol min⁻¹ at 37 °C.
‡*P. gingivalis kgp* mutant and *rgpA* mutant crude extracts were prepared as described in Methods.

Fig. 3. Gel filtration chromatography of the Triton-wt complex, RgpA complex and Kgp complex. Arginine-affinity purified Triton X-114 extracted wt complex (a), RgpA complex (b) and Kgp complex (c) were applied to a size-exclusion column (macro-sphere 300Å, 7 µm, 250×4.6 mm, Alltech) installed in a Waters Delta 600 HPLC system and the chromatogram developed at a flow rate of 0.5 ml min⁻¹ in 0.05 M KH₂PO₄ containing 0.15 M Na₂SO₄, pH 7.0. The column eluate was monitored at 280 nm to detect tryptophan and tyrosine residues. V₀ indicates the void volume of the column. The elution volumes of the standard proteins – i, thyroglobulin (669 kDa); ii, ferritin (312 kDa); iii, catalase (232 kDa) – are marked.

Moreover, only 50 % of the mice immunized with the Triton-wt complex developed *P. gingivalis*-induced lesions, whereas 70 % of the mice immunized with the sonication-wt complex developed lesions.

The subclass antibody response induced by the wt complex was determined by ELISA. The results, shown in Fig. 5, demonstrate that immunization with the Triton-wt complex elicited a higher antibody titre compared with the sonication-wt complex. The Triton-wt complex induced higher levels of IgG, IgG1, IgG2a, IgG2b and IgG3 antibodies, with the predominant antibody subclass being IgG1.

Western blot analysis of the Triton-wt complex and sonication-wt complex was performed using protective

Fig. 4. Maximum lesion size of mice immunized with wt complex extracted by sonication or by Triton X-114. BALB/C mice (10 mice per group) were immunized s.c. with Triton complex or sonication complex (emulsified in incomplete Freund's adjuvant, 25 µg per mouse) for the primary (day 0) and secondary (day 30) immunizations and challenged s.c. (day 42) with *P. gingivalis* ATCC 33277 (7.5×10⁹ viable cells). Animals were monitored over a period of 14 days for the development and size of lesions. Lesion sizes were statistically analysed using the Kruskal–Wallis test and the Mann–Whitney U–Wilcoxon rank sum test with a Bonferroni correction for type 1 error and effect size (Cohen's *d*). *, ** group significantly different (*P*<0.05, *P*<0.01, respectively) from the control (PBS/IFA) group.

Fig. 5. Serum antibody subclass responses to the wt complexes extracted using the Triton X-114 and sonication procedures. Sera from mice immunized with the Triton-wt complex (black bars) and sonication complex (white bars) were used in the ELISA with the complex as the absorbed antigen. Antibody responses are expressed as the ELISA titre determined as the reciprocal of the dilution at which absorbance was double the background level, with each titre representing the mean $\pm$ SD of three values.

anti-complex mouse sera. The Triton-wt complex antisera exhibited strong immunoreactive bands at molecular masses 44, 39 and 30 kDa, corresponding to the RgpA_{A1}, Kgp_{A1} and HagA_{A1}* adhesins (Fig. 6). The sonication-wt complex also displayed immunoreactive bands of molecular masses 44 and 39 kDa, corresponding to the complex proteins RgpA_{A1} and Kgp_{A1} (Fig. 6). The immunoreactive 45 kDa band of the wt complex was found not to be RgpA_{cat}, as the complex antisera did not recognize RgpB, which has 97 % sequence identity to the RgpA proteinase, suggesting that the immunoreactive band detected at 45 kDa was also derived from the adhesins (data not shown). These data suggest that the Triton wt complex produces a stronger antibody response directed towards the A1 adhesins of RgpA, Kgp

Fig. 6. Western blot analysis of the wt complex probed with antisera from mice immunized with the sonication-wt complex (A) or Triton-wt complex (B). The Triton-wt complex and sonication-wt complex were separated by SDS-PAGE, transferred onto PVDF membrane and probed with Triton-wt complex (1:50 TN buffer) and sonication-wt complex antisera (1:50 TN buffer), respectively. Molecular mass markers are shown in kDa.

and HagA polyproteins compared with the sonication-wt complex.

K_D for the binding of the wt complex, RgpA complex and Kgp complex to host proteins

The binding of purified Triton-wt complex, sonication-wt complex, RgpA complex and Kgp complex to fibrinogen, fibronectin, haemoglobin, laminin, and collagen types I, III, IV and V was determined by ELISA. K_D for each interaction was determined by Scatchard analysis of the ELISA data (Table 5). The Triton-wt complex bound in the nanomolar and subnanomolar range to the proteins tested, with the strongest binding to fibrinogen followed by fibronectin,

Table 5. Equilibrium constants (K_D) for binding of the wt complex, RgpA complex and Kgp complex to host proteins

Host protein	K _D $\pm$ SD (nM)			
	Triton-wt complex	Sonication-wt complex	RgpA complex	Kgp complex
Fibrinogen	0.90 $\pm$ 0.04†,\$	5.9 $\pm$ 0.8	2.07 $\pm$ 0.11‡	4.90 $\pm$ 0.20
Fibronectin	0.18 $\pm$ 0.01†,\$	3.0 $\pm$ 1.0	1.66 $\pm$ 0.06‡	4.37 $\pm$ 0.10
Haemoglobin	1.81 $\pm$ 0.07†,\$	10.7 $\pm$ 2.3	3.72 $\pm$ 0.14	3.92 $\pm$ 0.14
Collagen types I, II, IV	*	*	*	*
Collagen type V	2.70 $\pm$ 0.17†,\$	5.2 $\pm$ 0.6	4.72 $\pm$ 0.11‡	5.90 $\pm$ 0.12
Laminin	4.49 $\pm$ 0.09	5.3 $\pm$ 0.9	4.70 $\pm$ 0.17‡	*

*Binding not detected.

†Significantly ($P < 0.01$) different to sonication-wt complex.

‡Significantly ($P < 0.01$) different to Kgp complex.

§Significantly ($P < 0.01$) different to Kgp complex and RgpA complex.

haemoglobin, collagen type V and laminin, but no binding was detected for collagen types I, III and IV. Although the sonication-wt complex also bound to host proteins in the nanomolar range, the Triton-wt complex was significantly ($P < 0.01$) more effective at binding to fibrinogen ($d = -6.25$, 99 % CI: -12.69 , -1.4), fibronectin ($d = -2.82$, 99 % CI: -6.35 , -0.02), haemoglobin ($d = -3.87$, 99 % CI: -8.22 , -0.5) and collagen type V ($d = -4.17$, 99 % CI: -8.49 , -0.56). The individual Kgp complex and RgpA complex had significantly ($P < 0.01$) lower binding affinities to all the proteins tested with the exception of laminin (Table 5). Although, the Kgp complex had binding affinities in the nanomolar range, it exhibited significantly ($P < 0.01$) weaker binding to fibrinogen, fibronectin and collagen type V compared with the Triton-wt complex and RgpA complex (Table 5). Furthermore, the Kgp complex, unlike the wt complex and the RgpA complex, did not bind to laminin (Table 5). Both the Kgp complex and RgpA complex had similar binding affinities to haemoglobin (Table 5).

DISCUSSION

The Arg- and Lys-specific proteinases and their associated adhesins (proteinase–adhesin complexes) are major virulence factors of *P. gingivalis* (O'Brien-Simpson *et al.*, 2004). In this study, we describe a simple method of extraction and purification of the major cell-associated proteinase–adhesin complexes from *P. gingivalis* using Triton X-114 and a single Arg-affinity chromatography procedure. Although other authors have used detergents such as CHAPS or Triton X-100 in the extraction step for the proteinase–adhesin complexes from *P. gingivalis*, these detergents were used to solubilize proteins after sonication (Fujimura *et al.*, 1998; Kontani *et al.*, 1996). Takii *et al.* (2005) used sucrose-monomylaurate to extract outer-membrane proteins from *P. gingivalis*, but as with all other reports these authors used multiple chromatography procedures to purify the proteinase–adhesin complexes. Furthermore, Takii *et al.* (2005) did not characterize their purified complex using mass spectrometry. The Triton X-114 extraction method described here was significantly more effective than the previously described sonication method for the extraction of the proteinase–adhesin complexes from the cell surface, as demonstrated by a higher yield and the presence of substantially more Arg- and Lys-specific proteolytic activity. Interestingly, the Lys-specific activity of the Triton-wt complex was comparable to the Lys-specific activity of the Kgp proteinase–adhesin complex purified from culture supernatants of *P. gingivalis* strain HG66 by Pike *et al.* (1994). The higher specific Arg and Lys proteolytic activity of the Triton-wt complex and the presence of the HagA_{A1}* domain are consistent with the Triton X-114 procedure extracting and retaining the native structure of the complex better than the sonication procedure.

Using N-terminal sequence analysis and peptide mass fingerprinting, the extracted and purified Triton-wt complex was found to contain RgpA_{cat}, RgpA_{A1}, RgpA_{A2},

RgpA_{A3}, Kgp_{cat}, Kgp_{A1}, Kgp_{A2}, HagA_{A2} and HagA_{A1}*, corresponding to the molecular masses, as determined by SDS-PAGE and deduced from the gene sequence (Veith *et al.*, 2002) in parentheses of 45 (51), 44 (38), 17 (13), 15 (15), 48 (53), 39 (38), 17 (13), 17 (13) and 30 (28), respectively, which equates to the Triton-wt complex having a combined molecular mass of 272 (262) kDa, assuming stoichiometric amounts of each domain. This molecular mass is very similar to the relative molecular mass (260 kDa) of the smaller molecular mass peak of the Triton-wt complex determined by size-exclusion chromatography. However, the majority of the purified proteinase–adhesin complexes eluted in the void volume of the size-exclusion column, suggesting that the individual complexes may form larger aggregates on the cell surface as suggested by Takii *et al.* (2005) or that the complexes further aggregate after they are released/extracted from the cell surface. Furthermore, for both the Kgp complex and RgpA complex, gel filtration peaks that correspond to the relative molecular masses of 120 kDa and 150 kDa, respectively, fitted well with the stoichiometric amounts of the domains identified for each complex; Kgp_{cat} 48 (53), Kgp_{A1} 39 (38), Kgp_{A2} 17 (13) and HagA_{A2} 17 (13) producing 121 (117) kDa and RgpA_{cat} 45 (51), RgpA_{A1} 44 (38), RgpA_{A2} 17 (13), RgpA_{A3} 15 (15) and HagA_{A1}* 30 (28) producing 151 (145) kDa, respectively.

The N-terminal sequence and cleavage sites of the protein domains from the Triton-Kgp complex and Triton-RgpA complex were compared with the corresponding domains of the Triton-wt complex. The N-terminal sequence and cleavage sites of Kgp_{cat}, Kgp_{A1} and Kgp_{A2} from the *P. gingivalis* *rgpA* mutant and *P. gingivalis* W50 (wild-type) were identical, with an Arg residue preceding all the cleavage sites. This indicates that in the absence of RgpA proteinase activity, the N-terminal processing of Kgp_{cat}, Kgp_{A1} and Kgp_{A2} domains of the Kgp complex is performed by another Arg-X proteinase, probably RgpB. Recently, Veith *et al.* (2002) demonstrated that either RgpA or RgpB proteinase activity had a role in the N-terminal processing of Kgp proteins, where the N-termini of Kgp_{cat}, Kgp_{A1} and Kgp_{A2} from an outer-membrane preparation of a *rgpA/rgpB* double mutant had extended and often ragged N-termini. Similarly, RgpA_{cat}, RgpA_{A1} and RgpA_{A2} domains, but not the RgpA_{A3} adhesin domain, from the *P. gingivalis* *kgp* mutant and *P. gingivalis* W50 (wild-type) had identical N-termini with an Arg residue preceding the cleavage site, consistent with either RgpA and/or RgpB proteolytic activity being involved in the processing of these domains. The RgpA_{A3} adhesin domain from *P. gingivalis* W50 (wild-type) is preceded by a lysine residue, and the RgpA_{A3} adhesin of the RgpA complex extracted from the *P. gingivalis* *kgp* mutant was extended by six residues and was preceded by an arginine residue. This suggests that the Kgp proteinase normally participates in the proteolytic processing of the RgpA_{A3} domain in wild-type *P. gingivalis* and that in the absence of the Kgp proteinase, either RgpA or RgpB proteinase processes the RgpA_{A3} adhesin.

The processing of RgpA and Kgp polypeptides into fully active mature proteins on the cell surface is not yet fully understood. Recently, Mikolajczyk *et al.* (2003) reported a possible activation pathway for RgpB, which involves the removal of N- and C-terminal extensions in three sequential autoproteolytic steps to generate fully active RgpB. Similarly, it has been suggested that prior to becoming fully processed and active proteins on the cell surface, the RgpA or Kgp polypeptides undergo a succession of proteolytic cleavages and these proteolytically processed domains become tightly associated and form the RgpA-Kgp proteinase–adhesin complex on the cell surface (Bhagal *et al.*, 1997). The N-terminal sequencing and PMF data shown in the present study demonstrate that the Kgp proteinase is involved in processing of the RgpA_{A3} adhesin domain and the majority of the RgpA and Kgp domains are processed by either the RgpA or RgpB proteinases. Using proteinase inhibitors to eliminate proteolytic processing during extraction, Veith *et al.* (2002) showed that the RgpA and Kgp proteinase and adhesins are fully processed domains on the cell surface of *P. gingivalis*. Our data suggest that processed domains of the three polyproteins RgpA, Kgp and HagA combine non-covalently to form complexes either on the cell surface or immediately upon release from the cell surface.

It is interesting that in this study the C-terminal domains of both RgpA and Kgp were not found in the extracted complexes. RgpA-Kgp complexes characterized by Pike *et al.* (1994) and Bhagal *et al.* (1997) were reported to contain the C-terminal domain RgpA_{A4} as a 27–30 kDa protein based on SDS-PAGE. Both Pike *et al.* (1994) and Bhagal *et al.* (1997) used N-terminal sequence analysis to identify RgpA_{A4}. However, in the current study we used PMF analysis, which conclusively identified the protein domains in the 30 kDa band of the RgpA-Kgp complexes as Kgp_{A1}/HagA_{A1}*. Kgp_{A1} has an identical N-terminal sequence to RgpA_{A4}, which would explain why Pike *et al.* (1994) and Bhagal *et al.* (1997) both assigned this band to RgpA_{A4}. The pattern of protein bands obtained by SDS-PAGE analysis of the RgpA-Kgp complexes purified by Pike *et al.* (1994) and Bhagal *et al.* (1997) were the same as those obtained in this study with the Triton-wt complex, where PMF analysis confirmed the absence of the C-terminal domains of both RgpA and Kgp in the purified complex. Veith *et al.* (2002) showed using 2D-PAGE analysis of *P. gingivalis* W50 outer membranes that the C-terminal domain of RgpA, RgpA_{A4} was present as a highly post-translationally modified form that reacted with mAb 1B5 whereas the other RgpA and Kgp domains obtained did not react with the mAb. This mAb has recently been reported to react with a phosphorylated branched mannan that has been suggested to be part of the outermost layer of the *P. gingivalis* cell wall (Paramonov *et al.*, 2005). These results together suggest that the C-terminal domain of RgpA_{A4} may be attached to the cell wall and hence may serve to anchor the complex to the cell. Treatment of the cell with Triton X-114 may have dissociated the RgpA-Kgp complex from its C-terminal anchor or may have triggered proteolytic processing,

releasing the complex from the attached C-terminal domain.

As RgpA and Kgp are reported to be major virulence factors of *P. gingivalis* (reviewed by O'Brien-Simpson *et al.*, 2004), we evaluated the protective efficacy of the complexes as vaccines in the murine lesion model. Both the Triton-wt complex and sonication-wt complex provided protection against challenge with *P. gingivalis*. However, the Triton-wt complex, when used as a vaccine, tended to be more effective in providing protection (as indicated by a larger effect size) compared with the sonication-wt complex. Moreover, immunization with Triton-wt complex induced higher levels of IgG, IgG1, IgG2a, IgG2b and IgG3 antibodies compared with the sonication-wt complex. The predominant antibody subclass induced was IgG1 (equivalent to IgG4 in humans), which is a Th2 cytokine-induced antibody that has previously been shown to be involved in a protective immune response against *P. gingivalis*-induced lesions in the mouse (O'Brien-Simpson *et al.*, 2000). Furthermore, Western blot analysis suggested that antibodies to the Triton-wt complex bound strongly to RgpA_{A1}, Kgp_{A1} and HagA_{A1}* adhesins, whereas antibodies to the sonication-wt complex bound to RgpA_{A1} and Kgp_{A1}. These data suggest that the complex extracted using the Triton X-114 method produces a strong antibody response directed towards the A1 adhesins of RgpA, Kgp and HagA. The RgpA_{A1}, Kgp_{A1} and HagA_{A1}* adhesins share a high degree of sequence similarity and interestingly, each contain the previously described protective peptide epitopes ABM1, ABM2 and ABM3 (O'Brien-Simpson *et al.*, 2000). Taken together, our results suggest that proteinase–adhesin complexes extracted from *P. gingivalis* are composed of processed domains of all three polyproteins, RgpA, Kgp and HagA, and that the superiority of the Triton X-114-extracted complex in protection may relate to the Triton complex more closely resembling the form of the proteins on the cell surface.

Binding studies demonstrated that the Triton-wt complex, sonication-wt complex, RgpA complex and Kgp complex bind at nanomolar levels to fibrinogen, fibronectin, collagen type V and haemoglobin. Interestingly, the Triton-wt complex demonstrated a higher affinity to all the proteins tested compared with the sonication-wt complex. The RgpA complex exhibited higher binding for all the proteins tested compared with the Kgp complex with the exception of binding to haemoglobin, where the K_D for each complex was similar. In an earlier study, Pike *et al.* (1996) reported that the RgpA complex and Kgp complex purified from the culture supernatant of *P. gingivalis* strain HG66 bound to fibrinogen, fibronectin and laminin. The K_D of binding to fibrinogen was reported to be 8.5 nM and 4.0 nM for the RgpA complex and Kgp complex, respectively. This K_D of binding to fibrinogen is in good agreement with our findings, where the K_D of the Kgp complex binding to fibrinogen was 4.9 ± 0.2 nM. However, in the present study the RgpA complex had a stronger binding to fibrinogen, with a K_D of 2.07 ± 0.11 nM. This could be due to differences in strain,

age of culture and purification method used between the two studies. Knibbs *et al.* (1998) reported that the affinity of a protein to its ligand can be enhanced with an increase in ligand valency. Therefore, the higher binding to host proteins of the Triton-wt complex and RgpA complex compared with the Kgp complex may be due to the valency of ABMs in each complex. O'Brien-Simpson *et al.* (2005) have shown that antibodies directed to ABM2 (SYTYTVYRDGT-KIKEGLTATTFEEDGVAA) block binding of the complex to fibrinogen, fibronectin and collagen type V, and antibodies to ABM3 (VTLKWDAPNGTPNPNPNPNPNGTTTLESEF) block binding to haemoglobin. ABM2 is repeated three times in the adhesin domains of the RgpA complex and twice in the adhesin domains of Kgp complex, whereas ABM3 is repeated only once in the RgpA complex and Kgp complex. Thus, the stronger binding of the RgpA complex to fibrinogen, fibronectin and collagen type V may be due to the presence of a greater number of ABM2 sequences in its adhesin domains. In contrast, the similar levels of binding to haemoglobin observed with both the RgpA and Kgp complex may be explained by the presence of the same number of ABM3 sequences in their adhesin domains. The Triton-wt complex, which has the highest valency of ABM2 and ABM3 sequences, was found to have the strongest levels of binding to all the proteins tested. However, the higher affinity of the Triton-wt complex for host proteins may also relate to the unique conformation formed upon complex formation. This ability to bind to a variety of extracellular matrix (ECM) proteins may play an important role in *P. gingivalis* virulence as adherence to ECM proteins has been reported to be a crucial step in the pathogenic process associated with many bacteria (Schwarz-Linek *et al.*, 2004). Similar to other microbial surface proteins that bind ECM proteins, the RgpA-Kgp complex can also be considered to be part of the family of cell-surface adhesin molecules called MSCRAMMs (microbial surface components recognizing adhesive matrix molecules) (Patti *et al.*, 1994; Schwarz-Linek *et al.*, 2004). The adherence of the proteinase–adhesin complexes to collagen type V may be an important factor in the reported links between periodontitis and cardiovascular diseases and pre-term birth of infants (Beck *et al.*, 1998; Offenbacher *et al.*, 1998; Spahr *et al.*, 2006). Collagen type V is located in the basement membrane of human aortas, arteries and atherosclerotic plaques. It is also found in the endothelial basement membranes of the placenta and in the amniotic basement membrane (Merker *et al.*, 1987; Polzin *et al.*, 1997). The relative concentration of collagen type V has been reported to increase during the development and ageing of the placenta (Iwahashi *et al.*, 1996).

In conclusion, we have developed a simple and efficient method for the extraction and purification of the complexes of the processed domains of RgpA, Kgp and HagA from *P. gingivalis* using Triton X-114 and Arg-affinity chromatography. The Triton-extracted complexes bound at high affinity to a range of host proteins and when used as an

immunogen provided protection against challenge with *P. gingivalis* in a murine model of infection.

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