Cytolytic DNA vaccine encoding lytic perforin augments the maturation of- and antigen presentation by- dendritic cells in a time-dependent manner

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The use of cost-effective vaccines capable of inducing robust CD8+ T cell immunity will contribute significantly towards the elimination of persistent viral infections and cancers worldwide. We have previously reported that a cytolytic DNA vaccine encoding an immunogen and a truncated mouse perforin (PRF) protein significantly augments anti-viral T cell (including CD8+ T cell) immunity. Thus, the current study investigated whether this vaccine enhances activation of dendritic cells (DCs) resulting in greater priming of CD8+ T cell immunity. In vitro data showed that transfection of HEK293T cells with the cytolytic DNA resulted in the release of lactate dehydrogenase, indicative of necrotic/lytic cell death. In vitro exposure of this lytic cell debris to purified DCs from naïve C57BL/6 mice resulted in maturation of DCs as determined by up-regulation of CD80/CD86. Using activation/proliferation of adoptively transferred OT-I CD8+ T cells to measure antigen presentation by DCs in vivo, it was determined that cytolytic DNA immunisation resulted in a time-dependent increase in the proliferation of OT-I CD8+ T cells compared to canonical DNA immunisation. Overall, the data suggest that the cytolytic DNA vaccine increases the activity of DCs which has important implications for the design of DNA vaccines to improve their translational prospects.

DNA is an ideal platform to develop global vaccines because it is safe, cheap to manufacture rapidly and stable even in ambient temperatures. Furthermore, a landmark phase IIb clinical trial demonstrated that electroproportion of a DNA vaccine (VGX-3100) was capable of eliciting immunity to clear human papillomavirus (HPV) and completely regress HPV-induced cervical intrapithelial neoplasia in humans1. Although the protective mechanism of this vaccine is still not clear, it is likely that cell death and inflammation resulting from electroporation at the vaccination site are contributing factors to its efficacy2.

Several studies from our laboratory suggest that DNA vaccines designed to induce necrosis of vaccine-targeted cells elicit greater T cell immunity and protection than canonical DNA vaccines following intradermal (ID) immunisation3-6. In these studies, expression of the immunogen was driven by the cytomegalovirus (CMV) promoter while the simian virus 40 (SV40) promoter drove expression of a cytolytic protein, truncated mouse perforin (PRF). The CMV promoter is ~10-fold more efficient in driving gene expression than the SV40 promoter7 and the encoded PRF contains a 12 amino acid deletion (12del PRF) at the C-terminus, resulting in cell
death after expression\(^7\). This bicistronic plasmid design allows the immunogen to accumulate significantly within vaccine-targeted cells before necrosis occurs due to 12del PRF expression. Necrotic cell death results in release of damage associated molecular patterns (DAMPs) which act as natural adjuvants and result in inflammation and activation of antigen presenting cells (APCs)\(^8\). Dendritic cells (DCs) are APCs that are pivotal to initiate CD\(^8\)\(^+\) T cell immunity and are the most efficient APC at cross-presenting antigens including antigens derived from dead cells\(^9\). Consequently, in this study, we investigated the hypothesis that the cytolytic DNA vaccine will activate DCs to cross-present antigens to CD\(^8\)\(^+\) T cells more efficiently than canonical DNA vaccines which are non-cytolytic.

Results and Discussion
DCs exposed to necrotic or lytic cell environments mature and cross-present antigens to CD\(^8\)\(^+\) T cells more efficiently than DCs exposed to healthy or apoptotic cells\(^10–12\) and the release of lactate dehydrogenase (LDH) from cells is indicative of lytic/necrotic cell death\(^13\). Consequently, to investigate whether expression of the truncated mouse perforin, 12del PRF, encoded by the bicistronic DNA vaccine resulted in lytic cell death, the release of LDH from transfected HEK293T cells was measured (Fig. 1A). In this experiment, the bicistronic DNA encoded the non-structural protein 3 (NS3) from hepatitis C virus as the immunogen. A mutant form of 12del PRF with a D483A mutation which renders PRF non-cytolytic\(^7, 14\) served as a control for these experiments. Transfection of HEK293T cells with the bicistronic DNA encoding 12del PRF, but not wild-type (WT) PRF, significantly increased LDH release from transfected cells compared to the control (NS3-12del D483A PRF) (Fig. 1A). In contrast, treatment of HEK293T cells with a pro-apoptotic drug, doxorubicin, failed to result in LDH release (Fig. 1A). Thus, only expression of 12del PRF from the bicistronic DNA vaccine resulted in significant necrotic cell death. These results are consistent with previous reports demonstrating the lytic activity...
of 12del PRF and with the general understanding that apoptosis which fails to result in the release of DAMPs is non-inflammatory whereas necrosis not only results in the release of DAMPs, but is inflammatory leading to increased immunogenicity.

Next, to examine if DCs are activated following exposure of lytic cell debris resulting from 12del PRF expression, HEK293T cells were transfected with cytolytic- or control- DNA encoding the model antigen ovalbumin (OVA). These cells were then exposed to purified DCs from naïve C57BL/6 mice. This experiment showed that DCs (gated based on the expression of CD11c and Major Histocompatibility Complex Class (MHC-C) II (CD11c+I-Ab+)) including the cross-presenting CD8α+ subset, which were exposed to HEK293T cells transfected with the cytolytic DNA expressed higher levels of the costimulatory molecules CD80 and CD86, indicative of maturation, compared to DCs exposed to HEK293T cells transfected with control DNA (Fig. 1B and C). However, expression of the immunodominant peptide, SIINFEKL-H2-Kb, was not detected in these DC-HEK293T co-cultures (Fig. 1D). This was not surprising because the 25-D1.16 antibody fails to detect low levels of SIINFEKL-H2-Kb expression and is the only antibody available to quantify this peptide-MHC-CI complex on murine cells. Staining for SIINFEKL-H2-Kb was observed only for DC subsets analysed in co-cultures involving heat-treated necrotic cells (as described in Materials and Methods) in the presence of 0.1 μg/ml SIINFEKL peptide (positive control) (Fig. 1D). Nevertheless, the in vitro data (Fig. 1B,C) suggest that exposure of lytic cell debris resulting from cytolytic DNA expression can trigger maturation of DCs.

Proliferation of adoptively transferred carboxyfluorescein succinimidyl ester (CFSE)-labelled OT-I CD8+ T cells which present OVA epitopes, e.g. SIINFEKL, in recipient mice in the context of MHC-CI is commonly used to quantify antigen presentation by DCs. Thus, this method was used to determine whether cytolytic DNA immunisation could augment antigen presentation by DCs to CD8+ T cells in vivo (Fig. 2). The data show that immunisation with cytolytic DNA failed to increase the number of proliferating C57BL/6 Rag-/- OT-I CD8+ T cells in the draining cervical lymph nodes (CLN) compared to the control when early phases of antigen presentation (days 4-8) were examined (Fig. 2). Similar results were generated when C57BL/6 OT-I CD8+ T cells (as opposed to C57BL/6 Rag-/- OT-I cells (CD45.2+)) were used for adoptive transfer and proliferation of these cells was examined 0–7 days following DNA immunisation (data not shown). However, we have demonstrated previously that cell death in vivo was detected 14 days after ID immunisation with cytolytic DNA and antigen presentation following ID DNA immunisation can be detected for at least 21 days post-immunisation.
Consequently, we examined antigen presentation occurring from 7–22 days following ID immunisation with cytolytic or control DNA. As a positive control, one day after the adoptive transfer of OT-I cells into naïve recipient mice (n = 2), 20 μg SIINFEKL peptide + 5 μg of LPS were injected ID per mouse prior to analysis 14 days later. LPS promotes maturation of antigen presenting cells and SIINFEKL injection is commonly used in adoptive transfer experiments to ensure that OT-I CD8+ T cells in recipients retain the capacity to proliferate and to delineate proliferating cells during flow cytometric analysis. The results of these experiments show that cytolytic DNA immunisation increased the number of proliferating OT-I CD8+ T cells by ∼2-fold in the CLN as well as the spleen compared to the 12del D483A PRF control (Fig. 3). These findings are consistent with our previous report which demonstrated that a higher proportion of mature cross-presenting (i.e. CD8α+ CD80+) DCs emerge in the CLN 14 days following ID immunisation with cytolytic DNA compared to canonical DNA. Furthermore, the increase in antigen presentation due to cytolytic DNA vaccination is consistent with our earlier reports which demonstrated that cytolytic DNA vaccination greatly enhances anti-viral effector CD8+ T cell immunity. Thus, the data suggest that cytolytic DNA more efficiently activates DCs to present vaccine-encoded antigens to CD8+ T cells in vivo, compared to canonical DNA vaccination.

DCs are crucial to initiate CD8+ T cell immunity, but, being a rare cell population, direct targeting of DCs is difficult unless proteins/antibodies recognizing cell surface molecules (e.g. DEC205) are tagged on the DNA or lipid molecules (e.g. liposomes) are used to encapsulate DNA. However, the use of protein or antibody tags increases the manufacturing and storage costs of the vaccine, and despite the potential of these strategies, none has resulted in widespread use. Immunity against the protein tags can also develop resulting in reduced efficacy of the vaccine following booster vaccinations. Consequently, alternative strategies to target DNA-encoded antigens to DCs are required and the findings from the current study suggest that cytolytic DNA encoding 12del PRF...
can be used to deliver vaccine-encoded antigens effectively to DCs in an indirect manner and enhance antigen presentation to naïve CD8+ T cells in a time-dependent manner (Fig. 4).

Following ID delivery of canonical DNA, antigen delivery to DCs is crucial to prime naïve CD8+ T cells; this may occur after direct transfection of DCs although antigen transfer/shedding from transfected fibroblasts/keratinocytes or DCs uptake of antigen derived from dead cells may also contribute to some degree19–21 (Fig. 4B). However, the vast majority of the cells in the dermis are fibroblasts and keratinocytes, and consequently, the bulk of the DNA delivered to this site will likely transfect these cells (Fig. 4). Consequently, cross-presentation is necessary to efficiently prime naïve CD8+ T cells and it is most likely that this mechanism of antigen presentation is exploited more effectively following cytolytic DNA vaccination compared to canonical DNA vaccination (Fig. 4).

A caveat for the model described in Fig. 4 is that it is based on ID delivery of canonical or cytolytic DNA into the ear pinnae. It is possible that the method of antigen presentation to CD8+ T cells will vary depending on the route or mode of delivery of DNA22–25.

It is unclear which DC subsets cross-present antigen to CD8+ T cells following cytolytic DNA immunisation, but based on previous reports it is unlikely that Langerhan cells are required for cross-presentation26. It is likely that langerin+ dermal DCs27, CD8α+ and/or CD103+ DC subsets28,29 are involved, but this requires further investigation.

DNA vaccines have historically failed to show efficacy in large animal models and humans owing mainly to poor immunogenicity and delivery of DNA into cells. The recent success of using DNA vaccines to treat cervical cancer patients30 has reinvigorated the DNA vaccine research field. Furthermore, it is likely that a DNA vaccine will also emerge to protect against Zika virus infection in humans given that a plasmid DNA vaccine conferred protection in mice and macaques against Zika virus challenge31,32. The cytolytic DNA vaccine encoding 12del PRF is also more immunogenic than canonical DNA vaccines in outbred pigs33. Consequently, these findings highlight exciting
translational prospects for the cytolytic DNA to be used as a platform for designing vaccines against infectious diseases and cancer.

**Materials and Methods**

**Mice.** All experiments were approved and performed in accordance with the guidelines and regulations of the University of Adelaide and the SA Pathology animal ethics committees. C57BL/6 and C57BL/6.SJL mice were of the 6J substrain and purchased from the Australian Resource Centre at Perth. C57BL/6 OT-I mice were purchased from the Australian Phenomics Facility at Canberra. Rag\(^{-/-}\) C57BL/6 OT-I mice were bred in specific pathogen free conditions at the Sansom Institute for Health Research, Adelaide. All mice were housed in individually ventilated HEPA-filtered cages at the Queen Elizabeth Hospital, Woodville.

**DNA preparation and immunisations.** The cytolytic DNA was constructed using the pVAX plasmid (Life Technologies) as described previously\(^4\). The genes for *Gallus gallus* ovalbumin (OVA) or codon-optimised NS3\(^4\) were inserted downstream of the CMV promoter, and those for 12del PRF, 12del D483A PRF or WT PRF inserted downstream of the SV40 promoter. The different PRF sequences used are described in Brennan et al\(^7\). All DNA vaccines were prepared using a well-established alkaline lysis method\(^34\) and endotoxins removed with an Endotoxin Removal Solution (Sigma-Aldrich) following the manufacturer’s guidelines. A schematic diagram of the plasmid DNA constructs used in this study is shown in Fig. 5.

Aged matched (6–8 weeks old during initial vaccination) mice were under isofluorane anesthesia during ID DNA immunisations into the ear pinnae using a 29 G needle and syringe. Each mouse received 100 \(\mu\)g of endotoxin-free DNA in phosphate buffer saline (PBS) (50 \(\mu\)g in 10 \(\mu\)l/ear). In the experiment described in Fig. 3, a proportion of the mice were immunised via the ID route with 20 \(\mu\)g SIINFEKL peptide + 5 \(\mu\)g of lipopolysaccharide (LPS) in PBS (Sigma-Aldrich) as a positive control.

**LDH cytotoxicity assay.** HEK293T cells were transfected with 200 ng of DNA in 96-well flat-bottom plates and LDH activity in the culture supernatant measured 48 hours later using the manufacturer’s protocol (Thermo Scientific Pierce) as we described\(^5\). Eight hours after transfection the Dulbecco’s Modified Eagle Medium (DMEM; Life Technologies) + 10% heat inactivated Fetal Bovine Serum (FBS) which was used as cell culture medium during transfection was replaced with DMEM + 2% FBS. To induce apoptosis of HEK293T cells, cells were treated with 2 \(\mu\)M doxorubicin for 24 hours as described\(^5\). The percentage of maximum (specific LDH release) was calculated according to the manufacturer’s protocol (Thermo Scientific Pierce).

**Adoptive transfer.** Pooled splenocytes and lymph node cells from Rag\(^{-/-}\) C57BL/6 OT-I or C57BL/6 OT-I mice were labelled with 10 or 30 \(\mu\)M CFSE (Molecular Probes) respectively as demonstrated\(^17\). After washing, red blood cells (RBC) were depleted in RBC lysis buffer (155 mM NH\(_4\)Cl + 0.01 M Tris-HCl in Milli-Q water, pH 7.65), the labelled cells resuspended in PBS and injected intravenously (i.v.) into the lateral tail vein of C57BL/6. SJL vaccinated mice.

**DC-HEK293T cell coculture.** DCs were purified using splenocytes from naïve C57BL/6 mice using CD11c magnetic MicroBeads according to the manufacturer’s protocol (Miltenyi Biotec). Prior to CD11c enrichment, the spleens were digested for 45 mins at room temperature using 100 mg/ml collagenase type 3 (worthington) + 10 mg/ml DNase I (Sigma-Aldrich) in Roswell Park Memorial Institute Medium (RPMI; Life Technologies) + 2% FBS.

HEK293T cells from 3 wells transfected with DNA as described above were cocultured for 8 hours at 37°C + 5% CO\(_2\) with 1 x 10\(^5\) purified DCs. As a positive control, naïve HEK293T cells (cultured for 48 hours) were incubated at 56°C for 45 minutes to induce necrosis and cultured with 1 x 10\(^5\) purified DCs in the presence of 0.1 \(\mu\)g/ml of SIINFEKL peptide. Following the coculture, the DCs were stained with antibodies that react

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**Figure 5.** Schematic of the plasmid DNA constructs used in the study.
Flow cytometry. Cell surface and intracellular staining was performed similar to that described in Holmes et al. Antibodies against mouse antigens CD8α, T cell receptor (TCR) Vβ3.1/5.2, CD80 and CD11c were purchased from eBioscience. Antibodies against mouse antigen I-Aβ (MHC-II) was from BD Pharmingen. All the other antibodies were purchased from BioLegend.

Statistical analysis. All graphs were constructed using GraphPad Prism 6 software and the p values of the data were calculated using a paired t-test or Kruskal-Wallis H test of the IBM SPSS Statistics software (version 24).

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Acknowledgements
Grants APP1026293, 543139 and 543143 from the National Health and Medical Research Council (NHMRC) of Australia, grant BF040005 from the Australian-Indian biotechnology fund and a grant from the Hospital Research Foundation (THRF) supported the research in this study. THRF supports an early career fellowship for Danushka Wijesundara.

Author Contributions
W.Y., B.Q., P.E., J.H., K.D. and I.V. helped execute the experiments conducted in the manuscript. E.G. and B.G. contributed with ideas necessary to develop the manuscript hypothesis and helped with the write-up of the manuscript. D.W. executed the experiments, analysed the data and wrote the manuscript.

Additional Information
Competing Interests: The authors declare that they have no competing interests.

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