Identification of Two Lpp20 CD4⁺ T Cell Epitopes in Helicobacter pylori-Infected Subjects

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Antigen-specific CD4⁺ T cells play an essential role in effective immunity against Helicobacter pylori (H. pylori) infection. Lpp20, a conserved lipoprotein of H. pylori, has been investigated as one of major protective antigens for vaccination strategies. Our previous study identified two H-2d-restricted CD4⁺ T cell epitopes within Lpp20 and an epitope vaccine based on these epitopes was constructed, which protected mice in prophylactic and therapeutic vaccination against H. pylori infection. Immunodominant CD4⁺ T cell response is an important feature of antiviral, antibacterial, and antitumor cellular immunity. However, while many immunodominant HLA-restricted CD4⁺ T cell epitopes of H. pylori protective antigens have been identified, immunodominant HLA-restricted Lpp20 CD4⁺ T cell epitope has not been elucidated. In this study, a systematic method was used to comprehensively evaluate the immunodominant Lpp20-specific CD4⁺ T cell response in H. pylori-infected patients. Using in vitro recombinant Lpp20 (rLpp20)-specific expanded T cell lines from H. pylori-infected subjects and 27 18mer overlapping synthetic peptides spanning the whole Lpp20 protein, we have shown that L55–72 and L79–96 harbored dominant epitopes for CD4⁺ T cell responses. Then the core sequence within these two 18mer dominant epitopes was screened by various extended or truncated 13mer peptides. The immunodominant epitope was mapped to L57–69 and L83–95. Various Epstein-Barr virus (EBV) transformed B lymphoblastoid cell lines (B-LCLs) with different HLA alleles were used as antigen presenting cell (APC) to present peptides to CD4⁺ T cells. The restriction molecules were determined by HLA class-antibody blocking. L57–69 was restricted by DRB1-1501 and L83–95 by DRB1-1602. The epitopes were recognized on autologous dendritic cells (DCs) loaded with rLpp20 but also those pulsed with whole cell lysates of H. pylori (HP-WCL), suggesting that these epitopes are naturally processed and presented by APC. CD4⁺ T cells were isolated from H. pylori-infected patients and stimulated with L57–69 and L83–95. These two epitopes were able to stimulate CD4⁺ T cell proliferation. This study may be of value for the future development of potential H. pylori vaccine.

Keywords: Helicobacter pylori, Lpp20, CD4⁺ T cell, immunodominant epitope, HLA restriction
INTRODUCTION

Helicobacter pylori (H. pylori) infects more than half of the population in the world. The infection is causally associated with gastritis, peptic ulcer, gastric adenocarcinoma and gastric mucosa-associated lymphoid tissue lymphoma. A growing body of evidence shows that CD4+ T cell responses play a vital role in protective immunity against H. pylori. An increased number of T cells were infiltrating in human stomach with a typical Th1 phenotype during H. pylori infection (Bamford et al., 1998). A predominant Th1-type response was also elicited early during H. pylori infection in rhesus macaques (Mattapallil et al., 2000). Either natural infection or vaccine-induced immunity to H. pylori depends on a strong Th1-type cellular adaptive immune response in mice (Ermark et al., 1998; Eaton et al., 2001; Akhiani et al., 2002). These studies demonstrate that the protective adaptive immunity against H. pylori involves in Th1 response.

Immunodominance is the phenomenon in which the cellular immune response tends to focus on only a few of antigenic epitopes even during responses to complex antigens or pathogens in infected or immunized individuals. Generally, immunodominant T cells are more prevalent and protective in immune response compared with subdominant ones. Therefore, the immunodominant T cells often provide effective immune response and play a pivotal role in the adaptive immunity against pathogens, and this has been well-demonstrated in many bacterial, viral, and tumor systems (Jackson et al., 2006; Wu et al., 2011). Many scientists in the vaccine field think that immunodominant CD4+ T cell epitopes seem to be critical for H. pylori vaccine development (Ermark et al., 1998; Akhiani et al., 2002; Nyström and Svennerholm, 2007). Although several immunodominant epitopes of H. pylori protective antigens were identified (Chen et al., 2013; Yang et al., 2013; Hu et al., 2016), few are known to immunodominant epitope-specific CD4+ T cells response of other H. pylori antigen and immunodominant CD4+ T cell epitope has yet to be elucidated.

Lpp20, an outer membrane lipoprotein on H. pylori, has been considered to be one of potential vaccine candidates (Kostrzynska et al., 1994; Keenan et al., 2000; Peter and Beglinger, 2007; Li et al., 2012). Our previous study identified one Lpp20 B cell epitope (L108–119) which induced mouse anti-Lpp20 serum (Li et al., 2007) and two H-2d-restricted Lpp20 CD4+ T cell epitopes (L58–72 and L83–97) which elicited Th1-type immune response in BALB/c mice (Li et al., 2012). Additionally, the epitope vaccine composed of above mentioned three epitopes could stimulate the production of high level of Lpp20-specific antibodies and Th1-type cytokines and significantly reduced H. pylori colonization in H. pylori-challenged mice (Li et al., 2013), suggesting that Lpp20-derived epitope-based vaccine could be a promising alternative to eradicate H. pylori infection. Due to the MHC molecule differences in mice and humans, H-2d-restricted CD4+ T cell epitopes cannot sometimes induce effective immune responses in humans. Thus, the identification of such epitopes might be important for the future development of potential H. pylori vaccine.

In the present study, we conducted a systematic mapping analysis to screen H. pylori Lpp20 immunodominant CD4+ T cell epitopes using in vitro expanded recombinant Lpp20 (rLpp20)-specific T cell lines from H. pylori-infected subjects and 27 18mer overlapping synthetic peptides covered the sequence of Lpp20 protein. We observed that CD4+ T cell responses to Lpp20 varied remarkably with broad epitope specificity and the main responses focused on L55–72 and L79–96. Further, the core sequence of 18mer immunodominant epitopes was identified by 13mer overlapping peptides and various truncated or extended peptides based on the 13mer peptide. The immunodominant epitope was mapped into L57–69 and L83–95. The former epitope was restricted by HLA-DRB1*1501 and the latter one was restricted by HLA-DRB1*1602 and they were both naturally processed and presented by APC.

MATERIALS AND METHODS

Subjects and Blood Samples

This study was approved by the Institutional Human Ethics Review Board of Nanfang Hospital, Southern Medical University, Guangzhou, China and carried out in accordance with the recommendations. All subjects gave written informed consent in accordance with the Declaration of Helsinki. Gastric diseases were diagnosed by both endoscopic and histopathologic examination. 13C Urea breath test and serum anti-H. pylori antibody ELISA were performed to screen H. pylori-infected subjects. Blood samples from these subjects who donated more than 200 mL blood were collected and peripheral blood mononuclear cells (PBMCs) were isolated by Ficoll-Paque™ (GE Healthcare) gradient and then stored in liquid nitrogen until use. The HLA genotype of PBMC was determined by polymerase chain reaction (PCR) with sequencing-based typing at Beijing Genomics Institute (BGI), Shenzhen, China.

rLpp20 Antigen and Lpp20 Synthetic Peptides

rLpp20 was expressed in Escherichia coli and purified as we described previously (Li et al., 2007) and stored at −70°C. Amino acid sequence of Lpp20 has been submitted to NCBI by us (No. AAZ13599). 18mer synthetic peptides that covered the whole Lpp20 protein and overlapped by 11 amino acids (Figure 1A) and 13mer peptides that covered the initially identified dominant 18mer sequence and overlapped by 11 amino acids (Figures 1B,C) were synthesized and purified (purity >95%) by GL Biochem (Shanghai, China). All synthetic peptides were dissolved in dimethyl sulfoxide (DMSO, Sigma, Shanghai, China) and stored at −80°C.

The Expansion of Lpp20-Specific CD4+ T Cells From H. pylori-Infected Subjects in Vitro

PBMCs (1–2 × 10^6) were pulsed with 0.2 μmol/L Lpp20 (or 5 μmol/L immunodominant peptide) and cultured in RPMI 1640 medium (Gibco) supplemented with 5% human AB
FIGURE 1 | The schematic diagram of 18mer and 13mer overlapping peptides covering Lpp20 protein. (A) 18mer synthetic peptides that covered the whole Lpp20 protein and overlapped by 12 amino acids. (B) Various N- and C-terminus extended or truncated 13mer peptides were based on initially identified dominant 18mer L55–72 and overlapped by 11 amino acids. (C) Various N- and C-terminus extended or truncated 13mer peptides were based on initially identified dominant 18mer L79–96 and overlapped by 11 amino acids.

Sera, 2-mercaptoethanol (5 × 10^{-5} mol/L), L-glutamine (2 mmol/L), and antibiotics (penicillin 100 U/mL and streptomycin 100 µg/mL) in 48-well cell culture plates. The culture medium was 50% replaced by the above-mentioned medium containing 10 U/mL recombinant human interleukin-2 (rh IL-2) on day 5 and then 50% replaced by above medium containing 25 U/mL rhIL-2 when the color of the medium was yellow. The cells were split on day 10 and harvested on day 13.

Systematic Identification of Immunodominant Lpp20 CD4^{+} T Cell Epitopes

PBMCs were thawed and stimulated with rLpp20. After 13 days, T cells were screened against 27 18mer overlapping peptides individually. The intracellular cytokine staining (ICS) and flow cytometry were used to analyze the frequency of IFN-γ-secreting CD4^{+} T cells. Further, the same method was used to determine the core sequence of 18mer immunodominant CD4^{+} T cell epitopes using 13mer overlapping peptides.

Establishment of Epstein-Barr Virus (EBV) Transformed B Lymphoblastoid Cell Lines (B-LCLs)

Using the culture supernatant from EBV-producing B95-8 cells, B lymphoblastoid cell lines (B-LCLs) were established from autologous PBMCs and cultured in RPMI-1640 (GIBCO) supplemented with 10% fetal calf serum (GIBCO), 2-ME (5 × 10^{-5} M), L-glutamine (2 mM), and antibiotics (penicillin 100 U/ml and streptomycin 100 µg/ml).

Intracellular Cytokine Staining (ICS)

In the assays for screening immunodominant CD4^{+} T cell epitopes by 18mer and 13mer synthetic peptide, bulk cultured Lpp20-specific T cells were incubated with 5 µmol/L peptide at 37°C for 5 h in the presence of monensin (Becton Dickinson, Shanghai, China). In the assays for determining HLA-restricting alleles, B-LCLs were incubated with 5 µmol/L peptide for 1 h and then the free peptides were washed out. After that, the B-LCLs were co-cultured with Lpp20-specific T cells at a ratio of 1:10 (B-LCLs/T cells) for 5 h in the presence of monensin. In the antibody-blocking assay, APCs were incubated with 5 µg/mL antibodies against HLA-DP (Abcam), HLA-DR (Biolegend), and HLA-DQ (Biolegend) for 30 min before addition of peptide and monensin. After then, T cell activation was assessed by ICS. The cells were harvested and stained firstly with anti-CD3-PE and anti-CD4-APC (Biolegend, Beijing, China) in 50 µl staining buffer (PBS containing 1% heat inactivated FCS) at 4°C for 30 min, washed, fixed with Fixation/Permeabilization solution (BD Cat. No.554715) at 4°C for 20 min and stained with anti-IFN-γ-FITC (Biolegend, Beijing, China) according to the instruction of the kit protocol. 1 × 10^6 cells were acquired on a FACSCanto II flow cytometer (Becton Dickinson). Lymphocytes were gated by forward scatter (FSC) and side scatter (SSC). CD4^{+}
T cells subsets were further identified by CD3-PE and CD4-APC staining. Finally, gated CD4+ T cells were analyzed for IFN-γ-FITC. FACS data were analyzed using FlowJo software (Tree Star, Inc. Ashland, OR, USA).

Preparation of *H. pylori* Whole Cell Lysates (HP-WCL)

As previously described (Taylor et al., 2006), *H. pylori* NCTC11637 strain was grown on brain-heart infusion (BHI) plates containing 7% goat blood, trimethoprim (5 µg/mL), polymyxin B (5 µg/mL), and vancomycin (10 µg/mL) and propagated in Brucella broth with 5% fetal bovine serum with gentle shaking at 37°C under microaerobic conditions (85% N2, 10% CO2, 5% O2). After being cultured for 1 day, bacteria were collected, washed, and lysed. The lysates were centrifuged at 10,000 g for 20 min at 4°C to remove intact cells and large debris. The supernatant was sonicated as previously described, HP-WCL as previously described (Chen et al., 2004). The protein content was measured using the BCA Protein Assay Kit (Beyotime, Shanghai, China). All lysates were aliquoted and stored at −20°C until future use.

Generation of Dendritic Cells (DCs) and Co-culture With Immunodominant Epitope-Specific T Cells

PBMCs were thawed and CD14+ cells were isolated from PBMCs using CD14 immunomagnetic beads (Miltenyi Biotec, Shanghai, China). Then the cells were cultured in RPMI 1640 medium (Gibco) supplemented with 5% AB human sera, 10 ng/mL interleukin-4, 20 ng/mL granulocyte-macrophage colony-stimulating factor (GM-CSF) at 37°C in a 5% CO2 incubator. On day 6, dendritic cells (DCs) were harvested and pulsed with immunodominant peptides, rLpp20, HP-WCL, and bovine serum albumin (BSA) as a negative control at a final concentration of 50 µg/mL for 24 h. Then, DCs and epitope-specific T cells were co-cultured at a ratio of 1:10 for 5 h in the presence of monensin.

**CD4+ T Cell Proliferation Assay**

CD4+ T cells (2 × 10^5) isolated from *H. pylori*-infected patients were incubated in medium alone or in the presence of phytohemagglutinin (PHA, 5 µg/mL) to assess cell vitality, or in the presence of peptides (20 µg/mL). Cultures were incubated in a total volume of 200 µl for 3 days (37°C, 5% CO2) and pulsed during the last 18 h with tritiated thymidine [3H] (1 µCi/well). [3H] thymidine incorporation was measured in a liquid scintillation counter after collecting cells onto glass fiber filters. The stimulation index (SI) was determined by comparing [3H] thymidine incorporation in the peptide-stimulated wells with unstimulated wells using the following equation: SI = mean cpm of peptide wells/mean cpm of no peptide wells. Experiments were independently repeated three times.

**Statistical Analysis**

The Student’s t-test was used to analyze the differences between two groups. Welch’s correction was applied when the variances of two compared groups were not equal. Differences were considered to be significant when the P-value was < 0.05.

**RESULTS**

The Frequency of Lpp20-Specific CD4+ T Cells in *H. pylori*-Infected Subjects Was Higher Than That in Uninfected-Subjects

To assess Lpp20-specific CD4+ T cell responses in *H. pylori*-infected subjects, PBMCs were isolated from *H. pylori*-infected subjects and the frequency of Lpp20-specific CD4+ T cells was determined by ICS after incubation with pooled 18mer peptides spanning the entire Lpp20 protein. However, the frequency of Lpp20-specific CD4+ T cells was too low to be detected ex vivo (Figure 2A). Specific cell expansion is typically required to low frequency responses, especially in the epitope mapping (Savi et al., 2009). Therefore, Lpp20-specific CD4+ T cells were expanded by pulsing PBMCs with rLpp20. As a result, the frequency of the Lpp20-specific CD4+ T cells became higher after in vitro stimulation (Figure 2B). To further confirm the presence of Lpp20-specific CD4+ T cells in *H. pylori*-infected subjects, PBMCs from 30 *H. pylori*-infected and 30 uninfected subjects were expanded and then the frequency of Lpp20-specific CD4+ T cells were measured as mentioned above. It was shown that the frequency of Lpp20-specific CD4+ T cells in *H. pylori*-infected subjects was significantly higher than that in uninfected- subjects (Figure 2C). These results indicate that the stimulation with rLpp20 was able to reactivate antigen-experienced CD4+ T cells in vivo.

**L55–72 and L79–96 Were Two Dominant Regions Recognized by Lpp20-Specific CD4+ T Cells**

To systematically investigate the fine specificity and full breadth of Lpp20-specific CD4+ T cell responses, PBMCs from several *H. pylori*-infected subjects were stimulated with rLpp20 and T cells were screened against 27 18mer overlapping peptides individually after 13 days. As shown in Figure 3, the Lpp20-specific CD4+ T cells recognized two dominant regions (L55–72 and L79–96). Lpp20-specific CD4+ T cells from subjects H6, H11, and H26 mainly focused on L55–72 (Figures 3B,C,F), whereas T cells from subjects H1, H16, and H21 recognized L79–96 (Figures 3A,D,E). Taken together, L55–72 and L79–96 contain immunodominant CD4+ T cell epitopes of Lpp20.

**L57–69 and L83–95 Were the Core Sequences of Immunodominant Epitope Within L55–72 and L79–96, Respectively**

To further and systematically characterize the core sequence within the immunodominant Lpp20 CD4+ T cell epitope-containing peptides L55–72 and L79–96, T cells reactive to these two peptides were expanded as mentioned above and assessed using a set of 13mer overlapping peptides covering the initially detected 18mer peptides and various N- and C-terminus extended or truncated peptides based on the dominant 18mer sequence (Figures 1B,C). For subject H6, the L55–72-specific CD4+ T cells mainly recognized L57–69 and L59–71 (Figure 4A). L57–69 stimulated more T cells equivalent to the L55–72 response, suggesting that this 13mer peptide was the most...
potent core sequence of the immunodominant Lpp20 CD4⁺ T cell epitope in subject H6 (Figure 3A). In addition, this was further confirmed by titration of L₅⁷–₇₂, L₇₉–₆₉, and L₆₉–₉₁ at a concentration of 5 × 10⁻⁵ mol/L–5 × 10⁻⁹ mol/L (Figure 4B). For subject H1, the L₇₉–₉₆⁺ specific CD4⁺ T cells mainly focused on L₈₃–₉₅ and L₈₅–₉₇ (Figure 4C). L₈₃–₉₅ stimulated more T cells equivalent to the L₇₉–₉₆ response, suggesting that this 13mer peptide was the most potent core sequence of the immunodominant Lpp20 CD4⁺ T cell epitope in subject H1 (Figure 4C). Moreover, this was further confirmed by titration of L₇₉–₉₆, L₈₃–₉₅ and L₈₅–₉₇ at a concentration of 5 × 10⁻⁹ mol/L–5 × 10⁻⁵ mol/L (Figure 4D). Taken together, the most potent immunodominant CD4⁺ T cell epitopes of Lpp20 were L₅⁷–₆₉ and L₈₃–₉₅.

**L₅⁷–₆₉ and L₈₃–₉₅-Responding CD4⁺ T Cells Were Restricted by HLA-DRB1*1501 and HLA-DRB1*1602, Respectively.**

To determine the restricting HLA molecule of L₅⁷–₆₉ and L₈₃–₉₅, a MHC Class-II antibody-blocking assay was operated. PBMCs obtained from subject H6 were pulsed with L₅⁷–₆₉ and cultured in the presence of HLA-DR, HLA-DP, or HLA-DQ. L₅⁷–₆₉ + T cells were naturally presented by APCs, autologous B-LCL from subject H1 efficiently induced L₅⁷–₆₉⁺ specific T cells. In contrast, the B-LCL from subject H17 and H22 expressed the same HLA-DRB1*0901 but did not present this peptide. Therefore, L₅⁷–₆₉ must be restricted by HLA-DRB1*1602.

**L₅⁷–₆₉ and L₈₃–₉₅ Were Naturally Processed and Presented by APCs**

To evaluate whether L₅⁷–₆₉ and L₈₃–₉₅ were naturally presented by APCs, autologous DCs were loaded with rLpp20, HP-WCL, BSA or an immunodominant epitope peptide for 24 h and then co-cultured with epitope-specific T cells for 5 h in the presence of monensin. ICS was carried out to evaluate whether the corresponding epitope-specific T cells could recognize these APCs. As shown in Figure 6B, pulsing the DCs of subject H6 with L₅⁷–₆₉, Lpp20, HP-WCL vigorously stimulated the L₅⁷–₆₉-expanded T cell line. However, the DCs pulsed with BSA or DMSO only elicited the L₅₅–₇₂-specific CD4⁺ T cells.
FIGURE 3 | The mapping and identification of the immunodominant Lpp20 CD4+ T cell epitope. Lpp20-specific T cells from H. pylori-infected subject were expanded in vitro as described above and further screened for their specific response to the 27 Lpp20 18mer overlapping peptides at a final concentration of 5 µmol/L in an ICS assay. The identified 18mer sequences are shown (A–F) corresponding to subjects H1, H6, H11, H16, H21, and H26, respectively.

**DISCUSSION**

Increasing evidence shows that antigen-specific CD4+ T cell response plays an essential role in anti-H. pylori protective immunity (D'Elios et al., 1997a; Sayi et al., 2009; Sjokvist Ottsjo et al., 2015). More importantly, the immunodominant CD4+ T cell response is considered to be more effective than subdominant response in the host adaptive immune response to H. pylori infection (Ermak et al., 1998; Akhiani et al., 2002; Nyström and Svennerholm, 2007). The type of CD4+ T cell response against H. pylori may vary according to the antigen involved (D'Elios et al., 1997b). Although some previous studies on...
FIGURE 4 | The identification and characterization of the core sequence of L55–72 and L79–96 immunodominant epitope. Lpp20-specific T cells from subject H1 and H6 were expanded in vitro as described above. The cells were screened for their specific response to the 5 Lpp20 13mer overlapping peptides (final concentration ∼5 µmol/L) in an ICS assay. (A) The 13mer overlapping peptides within the L55–72 18mer epitope were screened with L55–72-specific T cells. (B) The 13mer overlapping peptides within the L79–96 18mer epitope were screened with L79–96-specific T cells. (C) Three of the reactive peptide L55–72, L57–69, and L59–71 were titrated to compare their activity. (D) Three of the reactive peptide L79–96, L83–95, and L85–97 were titrated to compare their activity.

H. pylori-specific Th1 cell responses focused on specific antigens, they did not identify individual epitopes and therefore were not aware of the influence of involved HLA (D’Ellios et al., 1997b, 2003). Recently, many scientists began to pay attention to identifying HLA-restricted CD4+ T cell epitopes of H. pylori protective antigens using overlapping synthetic peptides, such as UreB373–385 and UreB438–452 (Yang et al., 2013), HpaA88–100 and HpaA142–159 (Chen et al., 2013; Hu et al., 2016). Chen et al. found that the HpaA88–100-specific Th1-polarized response in H. pylori-infected subjects was linked with resistance to severe H. pylori-associated gastric diseases (Chen et al., 2013). This kind of protection was also supported by epidemiologic and genetic studies. These findings suggest that efficient presentation and recognition of some H. pylori antigens by T cell receptor (TCR) could influence the outcome of associated pathologies. Thus, further studies on antigen processing, presentation, and TCR recognition during H. pylori infection are needed to better understand the immune responses against this bacteria.

Although several immunodominant epitopes of H. pylori protective antigens were identified (Chen et al., 2013; Yang et al., 2013; Li et al., 2015; Hu et al., 2016), much remains unclear concerning CD4+ T cell response to many H. pylori antigens. Our previous studies constructed an epitope vaccine based on our identified two Lpp20 H-2d restricted CD4+ T cell epitopes (Li et al., 2012), which stimulated prophylactic and therapeutic responses with Th1-type profile against H. pylori in mice (Li et al., 2015). The MHC loci are genetically variable in mammals so that the highly polymorphic nature of MHC loci causes the immune systems of different subjects/species to focus on different epitopes/antigens within the same antigen/pathogen. This is the reason that many candidate vaccines succeed in murine experiments but fail in human clinical trials. Therefore, we were committed to mapping Lpp20 HLA-restricted epitopes. In the present study, to be able to efficiently identify immunodominant Lpp20 epitopes, a short-term in vitro T cell expansion approach was used to increase the frequency of Lpp20-specific T cells in an unbiased fashion. This identifying analysis also required large arrays of overlapping synthetic peptides. However, the method was accurate and reliable. Moreover, this approach enabled us to evaluate whether these identified epitopes were immunodominant or subdominant.

In this study, the frequency of Lpp20-specific CD4+ T cells in H. pylori-infected subjects were observed higher than that in uninfected subjects, indicating that a systemic immune response can be elicited during H. pylori infection. Indeed, a very low-level Lpp20-specific CD4+ T-cell response was detected in some samples from non-H. pylori-infected patients. We believed that this was because of false-negative diagnosis of subjects who might have had brief or mild H. pylori infection. Then we used a systematic screening approach to evaluate the magnitude and
FIGURE 5 | Peptides alone or in presence of blocking antibodies against HLA-DR, HLA-DP, or HLA-DQ. HLA-II antibodies were used to identify the HLA locus presenting the L<sup>57–69</sup> (A) and L<sup>83–95</sup> (B) 13mer peptide. The results were shown as average frequency ± standard deviation (n = 3). Only anti-HLA DR antibodies abrogated the responses, demonstrating that the responses to L<sup>55–69</sup> and L<sup>83–95</sup> are MHC Class II restricted. (C) A panel of B-LCLs with different DR types was identified. HLA-matched B-LCLs were used to further identify the HLA molecule presenting the 13mer peptide L<sup>57–69</sup> (E) and L<sup>83–95</sup> (D).

FIGURE 6 | The natural processing and presentation of L<sup>57–69</sup> and L<sup>83–95</sup> by DCs. (A) The DCs were pulsed with L<sup>57–69</sup>, rLpp20, HP-WCL, or BSA for 24 h and then co-cultured with L<sup>57–69</sup>-specific T cells from subject H6 for 5 h in the presence of monenisin. The frequency of IFN-γ-secreting CD4<sup>+</sup> T cells was determined by ICS. (B) L<sup>83–95</sup>-specific T cells from subject H1 were used to determine in a manner similar to the peptide described in (A).
It is concluded that Lpp20-specific CD4+ T cell responses in H. pylori-infected individuals. The Lpp20-specific CD4+ T cell response mainly focused on two 18mer peptides (L55–72 and L79–96) on average, consisting with the widely observed immunodominant phenomenon. Our previous studies identified two Lpp20 H-2Kd-restricted CD4+ T cell epitopes (L53–72 and L93–97) response to mice by SYFPEITHI prediction and 15mer overlapping synthetic peptides (Li et al., 2012). It is concluded that Lpp20-specific CD4+ T cell response in humans and mice seems to be almost the same. After that, the most potent core sequence of L55–72 and L79–96 were determined by 13mer overlapping peptides. They were L57–69 and L83–95, which stimulated CD4+ T cell response equivalent to the stimulation of L55–72 and L79–96, respectively.

The MHC loci are variable and highly polymorphic in mammals so that MHC alleles of the same/species generally focus on different peptides let alone MHC alleles from different species. For example, within the 11 reported CD4+ T cell epitopes from H. pylori UreB, including 4 mouse and 7 human epitopes, not a single one is recognized by both murine and human CD4+ T cell responses (Hu et al., 2016). Therefore, identification of a broad spectrum of immunodominant T cell epitopes across different HLA is required for the rational design of T cell epitope-based vaccine against H. pylori. However, only a few HLA-restricted CD4+ T cell epitopes of H. pylori UreB and HpaA antigens have been identified (Chen et al., 2013; Yang et al., 2013; Hu et al., 2016). The immunodominant epitopes were considerably different among individuals with different HLA alleles. The Allele Frequency Net Database (http://www.allelefrequencies.net/default.asp) shows that the frequency of HLA-DRB1*1501 in the Chinese Han population is relatively high (up to 10%) and the frequencies of HLA-DRB1*0803 and HLA-DRB1*1404 in the Chinese Han population are relative low (<1%). However, HLA-DRB1*0803 is more common in some other populations, such as native populations in Papua New Guinea and Taiwan (up to 10%), Australia Aborigine (up to 20%). Chen et al. demonstrated that HpaA88–100-specific CD4+ T cell response was immunodominant in subjects expressing HLA-DRB1*1501 and HLA-DRB1*1501-restricted immunodominant CD4+ T-cell response to HpaA88–100 was related with reduced risk of severe H. pylori-associated gastric diseases (Chen et al., 2013). Interestingly, in the HLA-DRB1*1501 negative subject, HpaA142–159-specific CD4+ T cell response restricted by HLA-DRB1*0901 was the most immunodominant (Hu et al., 2016). In the present study, it was shown that L57–69 and L83–95 could induce dominant CD4+ T cell responses in subjects possessing HLA-DRB1 genotypes and the restriction molecules were detected by antibody blocking. Then, using a panel of B-LCLs with different HLA alleles as APCs to present peptides for CD4+ T cells, we found that L57–69 was restricted by HLA-DRB1*1501 and L83–95 was restricted by HLA-DRB1*1602. Moreover, the CD4+ T cells specific to these epitopes not only recognized autologous DCs pulsed with rLpp20 but also those loaded with HP-WCL, indicating that these epitopes are naturally processed and presented by APCs. Furthermore, L57–69 and L83–95 stimulated the proliferation of CD4+ T cells from H. pylori-infected subjects expressed HLA-DRB1*1501 and HLA-DRB1*1602, respectively. Therefore, HLA-DRB1*1501-restricted L57–69 and HLA-DRB1*1602-restricted L83–95 might be of important value for the development of novel vaccine against H. pylori. To further study the association of L57–69 and L83–95-specific CD4+ T cell responses with gastric diseases induced by H. pylori, HLA-DRB1*1501 and DRB1*1602-expressing subjects with different gastric diseases, including atrophic gastritis, non-atrophic gastritis, antral gastritis pangastritis, peptic ulcer, duodenal ulcer, gastric cancer, will be assessed for L57–69 and L83–95-specific CD4+ T cell responses.

The knowledge gained in our study may not only help us to further understand the mechanism of protective immunity against H. pylori infection, but also help us to improve vaccine design. However, H. pylori has a large genome that encodes many protective antigens, such as UreaseB, UreaseA, Lpp20,
CagA, VacA, and HapA. Thus, it is very important to better understand CD4+ T cell responses to other H. pylori encoded antigens at a population level and to identify other HLA-restricted immunodominant CD4+ T cell epitopes. In the future, we will construct subunit vaccine, which includes CD4+ T cell immunodominant epitopes of many H. pylori protective antigens, to induce Th1-type protective immunity.

ETHICS STATEMENT

The protocol was approved by the Institutional Human Ethics Review Board of Southern Medical Hospital, Southern Medical University, Guangzhou, China.

AUTHOR CONTRIBUTIONS

YN contributed and designed the majority of the work. JY, DW, and JW separated cells and screened overlapping peptides.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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