Phase I Trial of Overlapping Long Peptides from a Tumor Self-Antigen and Poly-ICLC Shows Rapid Induction of Integrated Immune Response in Ovarian Cancer Patients

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Introduction

Cancer vaccines using short-minimal peptides from tumor antigens present limitations because they can only be used in patients with known HLA alleles that present these epitopes in the absence of natural processing; alternatively, full-length protein vaccines often suffer from lack of consistent CD8+ T-cell induction, likely due to inefficient cross-presentation of the exogenous antigen by antigen-presenting cells (APC; ref. 1). In contrast, synthetic long peptides are efficiently presented to both CD4+ and CD8+ T cells and in vivo T cells after intracellular cytokine staining, and tetramer staining).

Experimental Design: Twenty-eight patients with advanced ovarian cancer in second or third remission were enrolled sequentially in three cohorts and received at least one vaccination. Patients in Cohort 1 (n = 4) received 1.0 mg OLP, Cohort 2 (n = 13) received OLP in Montanide-ISA-51, and Cohort 3 (n = 11) received OLP + 1.4 mg Poly-ICLC in Montanide-ISA-51 on weeks 1, 4, 7, 10, and 13. Humoral and cellular responses were evaluated by standardized immunomonitoring techniques (ELISA, ELISPOT assay, intracellular cytokine staining, and tetramer staining).

Results: The vaccine was generally well tolerated with injection site reactions and fatigue that resolved. NY-ESO-1–specific antibody and CD8+ T cells were undetectable after vaccination with OLP alone, but were found in 6 of 13 (46%) and 8 of 13 (62%) patients, respectively, after vaccination with OLP + Montanide, and in 10 of 11 (91%) and 10 of 11 (91%) patients, respectively, after vaccination with OLP + Montanide + Poly-ICLC. NY-ESO-1–specific CD4+ T cells were detected in all patients with greater frequency and polyclonality when Montanide-ISA-51 was used for vaccination. Inclusion of Poly-ICLC as an adjuvant further accelerated the induction of NY-ESO-1–specific immune responses.

Conclusions: The current study shows that NY-ESO-1 OLP vaccine is safe and rapidly induces consistent integrated immune responses (antibody, CD8+ and CD4+) in nearly all vaccinated patients when given with appropriate adjuvants. Clin Cancer Res; 18(23); 6497–508. ©2012 AACR.
Human cancer vaccines have shown limited clinical efficacy to date, and choosing the right combination of antigen and adjuvant requires guidance from in-depth immunomonitoring. Therapeutic vaccines with a tumor self-antigen are achievable in mice, especially in combination with toll-like receptor (TLR) ligands, but questions remain in humans. How to achieve induction of antigen-specific immune responses in all patients? Does prior exposure of the immune system to tumor self-antigens limit the ability to vaccinate? The present study in patients with ovarian cancer shows that overlapping long peptides (OLP) from a tumor-derived self-antigen induce strong immune responses in nearly all patients when combined with Poly-ICLC and Montanide-ISA-51. Inclusion of Poly-ICLC significantly enhanced vaccine immunogenicity compared with OLP alone or OLP in Montanide-ISA-51. Tumor antigen presence or systemic immunosuppression did not negatively affect the vaccine’s immunogenicity. These results provide a scientific rationale for using long peptides and TLR ligands in future human immunotherapeutic developments.

**Translational Relevance**

Human cancer vaccines have shown limited clinical efficacy to date, and choosing the right combination of antigen and adjuvant requires guidance from in-depth immunomonitoring. Therapeutic vaccines with a tumor self-antigen are achievable in mice, especially in combination with toll-like receptor (TLR) ligands, but questions remain in humans. How to achieve induction of antigen-specific immune responses in all patients? Does prior exposure of the immune system to tumor self-antigens limit the ability to vaccinate? The present study in patients with ovarian cancer shows that overlapping long peptides (OLP) from a tumor-derived self-antigen induce strong immune responses in nearly all patients when combined with Poly-ICLC and Montanide-ISA-51. Inclusion of Poly-ICLC significantly enhanced vaccine immunogenicity compared with OLP alone or OLP in Montanide-ISA-51. Tumor antigen presence or systemic immunosuppression did not negatively affect the vaccine’s immunogenicity. These results provide a scientific rationale for using long peptides and TLR ligands in future human immunotherapeutic developments.

**Materials and Methods**

**Eligibility criteria**

Eligible patients in this open-label, single institution phase I study had stage II to IV histologically documented epithelial carcinoma arising in the ovary, fallopian tube, or peritoneum (trial identifiers: NC100616941, MSK07-152, LUD2006-001). Patients initially received cytoreductive surgery and chemotherapy with at least one platinum-based chemotherapy regimen. Following relapse, they returned to a second or third complete clinical remission with additional chemotherapy treatment. Remission was defined as CA125 less than 35 U/mL, unremarkable physical examination, and no definite evidence of disease by computed tomography. Nonspecific lymph nodes or soft tissue abnormalities 1 cm or less were permitted. Cytotoxic chemotherapy was to have concluded at least 4 weeks before the start of the study. Other requirements included Karnofsky Performance Status (KPS) 70% or more, neutrophil count 1.5 × 10^9/L or more, hemoglobin 10 g/dL or more, platelets 80 × 10^9/L or more, and aminotransferases 2.5 × upper limit of the normal range or less. Patients with prior autoimmune diseases, severe allergic reactions, and serious intercurrent illness were excluded. All patients provided written informed consent for this Institutional Review-approved protocol. An Investigational New Drug application was filed by the Ludwig Institute for Cancer Research (LICR, NY).

**Treatment plan**

A total of 28 patients were sequentially enrolled in 3 cohorts. Patients received at least 1 subcutaneous immunization in rotating sites on the upper arms regardless of the expression of NY-ESO-1 in tumor tissues. Patients in Cohort 1 received 1.0 mg NY-ESO-1 OLP in 0.5 mL diluent; Cohort 2 received 1.0 mg NY-ESO-1 OLP in 0.5 mL diluent + 0.5 mL Montanide-ISA-51 VG (total of 1.0 mL); and Cohort 3 received 1.0 mg NY-ESO-1 OLP in 0.5 mL diluent + 0.7 mL Poly-ICLC + 1.0 mL Montanide-ISA-51 VG (total...
of 2.0 mL administered in 2 syringes containing 1.0 mL each. Vaccines were administered on weeks 1, 4, 7, 10, and 13 with final study safety assessment on week 16. Delayed Type Hypersensitivity testing was conducted with 1 mg lyophilized NY-ESO-1 OLP atpretreatment and at week 16.

**Evaluation during study**

Pretreatment evaluations included history, physical and radiologic examination, vital signs, KPS assessment, and laboratory tests including complete blood cell counts, electrolytes and renal function, hepatic profile, thyroid stimulating hormone, prothrombin time, serum CA125 level, and urinalysis. Subsequent visits included vital signs and physical examination, toxicity assessment using CTCAE version 3.0, and serial laboratory studies to include complete blood cell counts, electrolytes and renal function, and hepatic profile. CT imaging was obtained every 2 months while on study, and standard RECIST criteria were used for disease progression. Blood samples for research studies (HLA typing, humoral and cellular immunity) were obtained pretreatment and at weeks 4, 7, 10, 13, and 16.

**Statistical considerations**

The primary objective of the study was to assess the safety of repeated vaccination with NY-ESO-1 OLP with or without the immunoadjuvants Montanide and Poly-ICLC administered every 3 weeks for 5 vaccinations. The secondary endpoint was to characterize the immune response with NY-ESO-1 OLP vaccination with or without the immunoadjuvants. The plan was for Cohort 1 (3 patients), Cohort 2 (9 patients), and Cohort 3 (9 patients) to accrue sequentially, based on safety requirements and availability of Poly-ICLC. Dose adjustment was not permitted. For the evaluation of safety, dose-limiting toxicity (DLT) was defined as ≥grade II allergic reaction, ≥grade III autoimmune reaction, ≥grade III hematologic or nonhematologic toxicity including fever, or ≥grade III injection site reaction. A total of 21 to 24 patients (if DLT was experienced) were planned. If 1 of 3 patients had DLT, an additional 3 patients would be added. If 1 of 6 had DLT, the arm would be considered safe and an additional 3 patients would be added. If an additional patient had DLT, the arm would be discontinued. The immunologic data was to be reported descriptively. Population and individual immune response curves generated by ELISA, tetramer analysis, and IFN-γ-producing T-cell responses were estimated at baseline and at weeks 4, 7, 10, 13, and 16 for all patients. If at least 5 of 9 patients in Cohort 2 or 3 developed immune response (true response range 23%–28%), treatment would be considered worthy of further study. Time to disease progression was not an endpoint of this phase 1 study but would be recorded. Reciprocal antibody titers by ELISA were considered significant if ≥100. Antibody-specific T-cell responses by enzyme-linked immunosorbent spot (ELISPOT) assays were considered significant if spot count was more than 50 out of 50,000 cells and at least 3 times more than the number of spots obtained with control nonpulsed target cells. Remaining data were analyzed using Prism 5 software (GraphPad Software, Inc.) by unpaired 2-tailed t test or Kaplan–Meier method and log-rank test.

**Immunohistochemical evaluation of NY-ESO-1 expression**

Expression of NY-ESO-1 was assessed retrospectively by immunohistochemistry using E978 on archival paraffin blocks with representative areas of the surgical tumor resection specimens (16). Focal expression (<5% of tumor with E978 staining) was considered significant.

**Synthetic peptides**

The following 4 30/32mer peptides (vaccine OLP) were synthesized under cGMP conditions by Multiple Peptide Systems, and formulated as lyophilized powder in 25 mg palmitoyl oleoyl phosphatidyl choline (POPC) and acetic acid by Phares: NY-ESO-179-108; NY-ESO-1100-129; NY-ESO-1121-1150; and NY-ESO-1142-1173. Because most of NY-ESO-1 epitopes recognized by CD4⁺ and CD8⁺ T cells are in the central to C-terminus hydrophilic region of the protein and because of high homology to LAGE-1 (5), the first 78 amino acids of NY-ESO-1 and the last 7 amino acids were excluded from the vaccine design. To enable efficient cross-presentation, the selected length of vaccine OLP was 30 amino acids with an overlap of 9 amino acids to include most of the possible CD4⁺ and CD8⁺ T-cell epitopes. Peptides were designed to avoid any strong anchor HLA-binding motif at the C-terminus to prevent the generation of cryptic epitopes (17). All vaccine peptides were characterized by high-performance liquid chromatography and mass spectrometry. A different set of 20-mer OLP (assay OLP, purchased from Bio-Synthesis or GenScript), covering the entire sequence of NY-ESO-1 with an overlap of approximately 10 amino acids, were used for the analyses of antibody mapping and T-cell responses and to avoid detection of immune responses to potential impurities of the vaccine OLPs.

**Measurement of serum antibody**

Plasma was obtained before and after vaccination by centrifugation of whole blood and stored at −80°C. Recombinant tumor antigen NY-ESO-1 and control dihydrofolate reductase (DHFR) proteins were used to coat plates and measure specific serum antibody levels in ELISA as described previously (18). In some experiments, 20-mer synthetic OLP were used as the coating antigen. A reciprocal titer was estimated from optical density readings of serially diluted plasma samples as described (18). Negative control sera from healthy individual and positive control sera for each antigen from patients with cancer were always included in all assays. The anti-human immunoglobulin antibodies used as secondary reagents were: alkaline phosphatase (AP)-labeled goat-anti-human IgG (polyclonal antiserum; Southern Biotech), biotinylated mouse-anti-human IgG1 (Clone JDC-1; BD Pharmingen), AP-labeled mouse anti-human IgG2 (clone HP6002; Southern Biotech), AP-labeled mouse anti-human IgG3 (clone HP6050; Southern Biotech), AP-labeled mouse anti-human IgG4 (clone HP6023; Southern Biotech), AP-labeled goat-anti-human...
IgA (polyclonal antisera; Southern Biotech), AP-labeled mouse-anti-human IgD-AP (clone IADB6; Southern Biotech), AP-labeled goat-anti-human IgG (Clone HP6029; Southern Biotech), and AP-labeled goat-anti-human IgM (polyclonal antisera; Southern Biotech). To be considered significant, reciprocal titers had to be more than 100.

**Monitoring of T-cell responses**

Peripheral blood mononuclear cells (PBMC) were obtained by centrifugation over lymphocyte separation medium (Mediatech Inc) and aliquots were stored in liquid nitrogen. In vitro stimulation (presensitization) of T cells from patients with cancer was carried out as described (19). Briefly, CD8+ and CD4+ T cells were isolated from PBMC by magnetic beads (Invitrogen Dynal) and independently stimulated with T-cell–depleted cells that were pulsed overnight with NY-ESO-1-1 assay OLP (6 μmol/L/each peptide). In some experiments, cells were pulsed with reduced amount of OLP at 100 nmol/L for each peptide. Presensitization with adenovirus recombinant for NY-ESO-1 (1,000 units/cells) was also conducted in most patients but only for some time points because of the limitation in the number of PBMCs available. CD8+ and CD4+ T cells were cultured in the presence of 10 μg/mL IL-2 (Roche Molecular Biochemicals) and 20 ng/mL IL-7 (R&D systems) in RPMI-1640 medium supplemented with 10% human AB serum (Gemini Bio-Products), 2 mmol/L L-glutamine, 100 U/mL penicillin, 100 μg/mL streptomycin, and 1% nonessential amino acids (RPMI + 10% pooled AB serum). The number of cytokine-producing NY-ESO-1-specific T cells was evaluated by ELISPOT assay and intracellular staining of cytokines at day 9 to 14 for CD8+ T cells and at day 19 to 23 for CD4+ T cells. Each assay was repeated twice. Tetramers were also assayed at day 9 to 14 for CD8+ T cells for patients with appropriate HLA-class I alleles. Autologous Epstein–Barr virus–transformed B-cell lines (EBV-B cells) were generated from a part of CD4+ CD8+ cells by supernatant from B95-8 cells in RPMI + 10% fetal calf serum (FCS) and used as target cells in assays. For 6 patients whose autologous EBV-B cells failed to grow (M06, M11, M15, M24, M26, M28), HLA-compatible allogeneic EBV-B cells were used as target cells in ELISPOT assays. In addition, a part of CD4+ T cells was polyclonally expanded with phytohemagglutinin (PHA, Remel) in RPMI + 10% pooled AB serum in the presence of low-dose IL-2 and IL-7 to be used as target cells (T-APC) of CD4+ T cells in repeat assays.

**ELISPOT assay**

IFN-γ ELISPOT assay was conducted as described previously (19). Briefly, nitrocellulose-coated microtiter plates (Millipore) were coated overnight with anti-IFN-γ monoclonal antibody (2 μg/mL 1-D1K, MABTECH) and blocked with 10% human serum (Sigma-Aldrich) in RPMI-1640 medium. EBV-B cells were pulsed with subpools of assay OLP (10 μmol/L) or infected with vaccinia virus encoding NY-ESO-1 or Influenza virus nucleoprotein (NP) gene overnight at 37°C. Indicated number of effector cells were cocultured with 5 × 10⁴ antigen-pulsed EBV-B cells for 24 hours in RPMI medium without serum. Plates were developed using 0.2 μg/mL biotinylated anti-IFN-γ mAb (7-B6-1-biotin, MABTECH), 1 U/mL streptavidin–alkaline phosphatase conjugate (Roche Diagnostics GmbH), and 5-bromo-4-chloro-3-indolyl phosphate/nitroblue tetrazolium (Sigma-Aldrich). The number of spots was evaluated using C.T.I. Immunospot analyzer and software (Cellular Technologies). Results were shown as the average number of spots from duplicate wells without subtracting the number of background spots against unpulsed target cells. To be considered significant, antigen-specific IFN-γ responses had to have a spot count more than 50 out of 50,000 cells and at least 3 times more than the number of spots obtained with control nonpulsed target cells.

**Tetramer and intracellular cytokine staining assays**

Tetramer assays were conducted using HLA-A2/NY-ESO-1157–165, HLA-B35/NY-ESO-194–102, and HLA-Cw3/NY-ESO-192–100 complexes made by the Lausanne facility of the Ludwig Institute, in patients with relevant HLA alleles, as described (19). Responses were considered significant if tetramer-positive cells were 0.5% or more of CD8+ T cells and were appearing clustered. Intracellular Cytokine Staining assays were conducted using mAb for CD8, CD4, IFN-γ, IL-5, CD45RA, and CCR7, as described (19). Responses were considered significant if more than 0.3% and at least 3 times more than the numbers obtained with control nonpulsed target cells.

**Results**

**Patient characteristics and disposition**

A total of 28 patients with epithelial ovarian cancer, fallopian tube, or primary peritoneal cancer in second or third complete clinical remission were enrolled sequentially in 3 cohorts (Supplementary Table S1), between March 2008 and June 2011. The patient characteristics are listed in Table 1. Of the 28 patients, 26 (93%) had stage III disease at diagnosis, with predominantly serous histology. All

<table>
<thead>
<tr>
<th>Table 1. Patient characteristics</th>
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<tr>
<td><strong>Patient characteristics (N = 28)</strong></td>
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<tr>
<td>Median age (range)</td>
<td>57 (44–73 years)</td>
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<tr>
<td>Median # prior chemotherapy (range)</td>
<td>2.4 (2–4)</td>
</tr>
<tr>
<td>FIGO Stage at diagnosis</td>
<td>N (%)</td>
</tr>
<tr>
<td>II</td>
<td>1 (3.6)</td>
</tr>
<tr>
<td>III</td>
<td>26 (92.2)</td>
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<tr>
<td>IV</td>
<td>1 (3.6)</td>
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<td>Histology</td>
<td></td>
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<tr>
<td>Serous carcinoma</td>
<td>24 (85.8)</td>
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<tr>
<td>Endometrioid carcinoma</td>
<td>2 (7.1)</td>
</tr>
<tr>
<td>Carcinoma not otherwise specified</td>
<td>2 (7.1)</td>
</tr>
<tr>
<td>Second remission at start of study</td>
<td>18 (64.3)</td>
</tr>
<tr>
<td>Third remission at start of study</td>
<td>10 (35.7)</td>
</tr>
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patients had received platinum/taxane-based therapy, and 64% of patients entered the study in second remission and 36% were in third remission. The median time from last chemotherapy to first vaccination was 2.8 months (ranging from 1.2–5.5 months). From 26 patients with available tumor specimens assessed retrospectively, 2 had strong and 7 had focal (<5% sample) but significant NY-ESO-1 expression by immunohistochemistry, consistent with the expected NY-ESO-1 expression frequency in advanced ovarian cancer (Supplementary Table S1). For Cohort 1, 1 out of 4 patients was replaced because of disease progression at the third vaccination. For Cohort 2, 4 of 13 patients were replaced (2 because of disease progression at second and fourth vaccinations, respectively, and 2 due to unrelated medical illness or complication—edema and pneumonia, respectively). For Cohort 3, 2 of 11 patients were replaced (1 because of disease progression at second vaccination and 1 because of significant noncompliance on the part of the patient). The overall number of patients completing study as planned was 16 of 28 (57%). The ability to complete all 5 vaccinations was 3 of 4 in Cohort 1, 8 of 13 in Cohort 2, and 5 of 11 in Cohort 3 (Supplementary Table S1). The main reasons for study discontinuation were progressive disease (4/28 patients) and early study closure (4/11 in Cohort 3).

Safety

The vaccine was generally well tolerated, with mostly grade I (10, 76.9%) and II (1, 7.7%) adverse events unlikely related to the vaccine, except for injection site reactions and fatigue that were definitely or possibly related, respectively, that resolved (Supplementary Table S2). In Cohort 3, 1 patient developed grade II panniculitis at the injection site (Supplementary Fig. S1) and 3 patients had simple grade II injection site reactions. While the injection site reactions were grade II (and technically not DLT), they were more robust than anticipated and persistent (Supplementary Fig. S1), and therefore further vaccinations were held in the last 4 patients from Cohort 3. It was felt prudent to consider a lower dose of Poly-ICLC and/or Montanide before further study. These patients were followed for safety and recurrence as per protocol (Supplementary Table S1). The only grade III (severe) adverse events occurred in 2 patients in Cohort 2: pneumonia (unrelated) and abnormal neutrophil count (possibly related; Supplementary Table S2).

Monitoring of humoral immune responses

Spontaneous and vaccine-induced humoral immune responses were investigated by measuring serum IgG against recombinant full-length NY-ESO-1 protein in ELISA. Results are summarized in Fig. 1A and titers for individual patients are shown in Supplementary Fig. S2. Two patients had significant (reciprocal titer ≥ 100) spontaneous anti-NY-ESO-1 antibody before vaccination (M01 and M28). Patients who received OLP alone (Cohort 1, n = 4) failed to develop detectable NY-ESO-1–specific antibody throughout the vaccination period, except for baseline seropositive patient M01 whose titers gradually increased during vaccination. In contrast, evidence of NY-ESO-1 serum antibody was found in 6 of 13 (46%) patients from Cohort 2 and 10 of 11 (91%) patients in Cohort 3, showing that these peptides are immunogenic. Inclusion of Poly-ICLC in Cohort 3 led to accelerated seroconversions and significantly increased antibody titers compared with Cohort 2 (Fig. 1B). Patients were able to mount similar antibody responses to the vaccine regardless of NY-ESO-1 tumor expression, although the 2 patients with baseline seropositive also had the highest NY-ESO-1 tumor expression. Early discontinuation of vaccination in the last 4 patients (Supplementary Table S1) did not negatively impact antibody responses, as titers had already developed maximally after 2 or 3 injections and remained high during follow-up without additional vaccination.

Characterization of antibody responses

A mapping of epitopes recognized by vaccine-induced NY-ESO-1–specific antibody was determined using 20-mer overlapping peptides as antigen in ELISA. As shown in Fig. 1C and Supplementary Fig. S2, antibody responses seen to NY-ESO-1 protein were confirmed against these peptides and found clustered in 3 areas of NY-ESO-1 centered around amino acids approximately 80, 100 to 110, and approximately 135 to 149. These stretches coincided with hydrophilic areas of the NY-ESO-1 sequences used in the vaccine (Fig. 1C), similarly with what is expected from naturally occurring immune responses. Specific antibody titers were detected to all 4 30-mer overlapping peptides used in vaccination, with no clear evidence of immunodominance, although recognition of the most C-terminus peptide (NY-ESO-1142-173) was comparatively less frequent. Antibody mapping of samples from Cohort 3 was more widespread and showed a greater coverage of NY-ESO-1 epitopes than those from Cohort 2, possibly reflecting differences in overall titers. No clear response was detected against NY-ESO-1 region 1 to 80 outside of the vaccine, expect for patient M28 who had preexisting immunity. Thus, Poly-ICLC enhanced humoral immune responses by not only accelerating the responses and increasing the antibody titer but also by broadening the repertoire of antibodies.

We next investigated isotypes and class switching of NY-ESO-1 antibodies induced by OLP in different formulations. As shown in Fig. 1D, most OLP-induced antibodies were of IgG1 and IgG3 isotypes. NY-ESO-1–specific IgG2, IgG4, IgA, and IgM were also detected in some patients by the end of vaccination. There was no clear evidence of early IgM response when measured at week 4 or 7 (not shown).

Monitoring of CD8+ T-cell responses

All patients were analyzed for CD8+ T-cell responses against NY-ESO-1 at all time points available by IFN-γ ELISPOT assays and whenever possible by staining of intracellular cytokines and tetramers. In patients with appropriate HLA alleles, we also tested purified unstimulated CD8+ T cells for NY-ESO-1 tetramer staining and failed to detect clear evidence of positive cells from any of the samples tested, confirming the difficulty to detect
When patients were retested ex vivo, CD8+ T cells were stimulated once in vitro with a pool of 20-mer peptides from NY-ESO-1 using a protocol standardized to enable the amplification of NY-ESO-1-specific cells from circulating precursors without de novo induction. NY-ESO-1-specific CD8+ T cells were measured against peptide subpools covering C-terminal area 1 to 80 (outside of the vaccine), central area 71 to 130, and N-terminal area 119 to 180, to evaluate polyclonality. Figure 2A shows the magnitude of CD8+ T-cell responses detected by ELISPOT assays. From 4 patients in Cohort 1, CD8+ T-cell responses were detected only in baseline seropositive patient M01. From 8 of 13 (62%) patients in Cohort 2, weak and transient or sporadic CD8+ T-cell responses were detected from week to week, except for patient M16 who showed preexisting sustained responses. In contrast, 10 of 11 (91%) patients from Cohort 3 showed evidence of CD8+ T-cell responses, and 7 of these were consistent and sustained throughout vaccination. Furthermore, Poly-ICLC helped in the rapid induction of NY-ESO-1-specific CD8+ T cells, apparent at week 4 after a single injection of OLP. Responses were polyclonal, as they could recognize at least 2 different areas of NY-ESO-1. Individual minimal epitopes recognized by these CD8+ T-cell responses were confirmed by tetramer assays in several patients with HLA alleles for which prevalidated tetramers were available (Fig. 3A). Corresponding ELISPOT responses against minimal peptide epitopes were also confirmed (not shown).

**Quality of CD8+ T-cell responses**

The ability of vaccine-induced CD8+ T cells to recognize naturally processed NY-ESO-1 was shown by sensitizing and expanding NY-ESO-1-specific CD8+ T cells using APCs transduced with adenovirus recombinant for full-length NY-ESO-1.
NY-ESO-1 (Fig. 3B). In addition, CD8⁺ T cells presensitized with NY-ESO-1 OLP were able to recognize target cells infected with vaccinia virus encoding full-length NY-ESO-1 (vvESO) in 4 of 16 patients tested after vaccination, including baseline NY-ESO-1-seropositive patient M28 (Fig. 3C). The absence of vvESO recognition in some patients could be ascribed to the preferential expansion of low-avidity T cells after in vitro presensitization with the regular 6 μmol/L OLP, as using a lower dose (100 nmol/L) could recover reactivity to vvESO (Fig. 3D). Therefore, OLP vaccine induced CD8⁺ T cells of wide range of avidities, including some high-avidity responses comparable with those observed in patients with spontaneous NY-ESO-1 immunity.

**Monitoring of CD4⁺ T-cell responses**

Monitoring of NY-ESO-1–specific CD4⁺ T cells is shown in Fig. 2B. All patients were analyzed for CD4⁺ T-cell responses against NY-ESO-1 at all time points available by IFN-γ ELISPOT assays and whenever possible by staining of intracellular cytokines. NY-ESO-1–specific CD4⁺ T cells were seen in all patients and were significantly enhanced by Montanide and further by Poly-ICLC. Polyclonal responses to at least 2 areas of NY-ESO-1 were seen as early
as after 1 vaccination in the majority of patients from Cohort 3. Most HLA-DP4+ patients also developed a CD4+ T-cell response to HLA-DP4–restricted epitope NY-ESO-1
\textsuperscript{157–170} (not shown). These vaccine-induced CD4+ T-cell responses were confirmed by intracellular cytokine staining, with mostly IFN-\(\gamma\) and less IL-5 secretion (Supplementary Fig. S3). Interestingly, consistent CD4+ T-cell responses against the region excluded from the vaccine, NY-ESO-1
\textsuperscript{1–76}, were observed in 3 patients from Cohort 3, and transiently in 6 other patients. Detectable presence of NY-ESO-1 in patient tumors did not seem to affect the number or nature of CD4+ T-cell responses generated.

Results of NY-ESO-1–specific antibody, CD4+ and CD8+ T-cell responses are summarized in a heatmap format in Fig. 4. Collectively, occurrence of NY-ESO-1–specific CD4+ T cells together with antibody and CD8+ T-cell responses, that is, integrated immune responses, was seen in 1 of 4 patient from Cohort 1, 4 of 13 patients from Cohort 2, and 10 of 11 patients from Cohort 3, indicating that Poly-ICLC significantly improved the immunization not only by accelerating and broadening immune responses, but also by making them more consistent (higher counts for CD4+ T-cell responses against each NY-ESO-1 peptide subpool at week 4 in Cohort 3 compared with Cohort 2, higher overall maximal CD4 T-cell response and more integrated Ab-CD8-CD4 responses in Cohort 3 than in Cohort 2, \(P < 0.03\) by unpaired \(t\) test).

**Impact of NY-ESO-1 tumor expression on antibody, CD4+ and CD8+ T-cell responses**

The impact of NY-ESO-1 tumor expression on vaccine-induced immune responses was evaluated in Cohort 3 where 5 patients showed NY-ESO-1 tumor expression and 5 did not (Supplementary Table S1). Expression of
NY-ESO-1 antigen by the patient tumors did not affect the induction of NY-ESO-1 antibody, as average maximal titers at the end of vaccination did not significantly differ for patients with and without NY-ESO-1 tumor expression (Fig. 5A, \( P = 0.25 \) at week 16, t test). Similarly, NY-ESO-1 CD8\(^+\) and CD4\(^+\) T-cell responses were very comparable in patients with and without NY-ESO-1 tumor expression (Fig. 5B, \( P = 0.25 \) at week 16, t test).

**Impact of CD4\(^+\)CD25\(^+\) regulatory T cells**

Tumor vaccines have been reported to expand regulatory T cells (Tregs; refs. 4, 20). As reported previously (21, 22), NY-ESO-1–specific CD4\(^+\) T cells that were not detectable from whole CD4\(^+\) T-cell population before vaccination became detectable by removal of CD25\(^+\) T cells in several patients (Fig. 5C). In contrast to the prevaccination specimens, the effect of the removal of CD25\(^+\) T cells at week 4, 13, and 16 was not consistent, increasing some responses in patients while reducing others, or having no effect at all. From the observations that NY-ESO-1–specific IFN-\(\gamma\)–producing T cells were detectable in the presence of Tregs, and that removal of Tregs did not consistently enhance the expansion of effector CD4\(^+\) T cells after vaccination, it is unlikely that vaccination systematically induced NY-ESO-1–specific CD4\(^+\) Tregs able to actively suppress the expansion of NY-ESO-1–specific effector CD4\(^+\) T cells.

**Long-term follow-up and clinical observations**

At the last follow-up in February 2012, 3 of 4 patients had progressive disease (PD) in Cohort 1 (median 9 months, range 1–9), and 1 had no evidence of disease (NED) at 46 months. In Cohort 2, 10 of 13 had PD (median 4.5 months, range 1–15 months), and 3 were NED (median 41 months, range 29–46 months). In Cohort 3, 9 of 11 patients had PD (median 6 months, range 2–22 months) and 2 were NED (median 21 months, range 17–25 months). Although the study was not randomized and the small number of patients in this study does not allow for definitive conclusions, time-to-progression was analyzed posthoc by the Kaplan–Meier method and no significant difference was found between cohorts overall. However, among 10 evaluable patients vaccinated with OLP+Montanide+Poly-ICLC (Cohort 3), those with baseline evidence of NY-ESO-1 tumor expression (\( n = 5 \)) showed a significantly delayed time to recurrence compared with NY-ESO-1–negative patients (Fig. 5D). Such delayed time to recurrence was not observed for the other 2 cohorts when stratified by NY-ESO-1 expression.

**Discussion**

In this study, we showed that NY-ESO-1 OLP were immunogenic when used in combination with adjuvant. Both Montanide and Poly-ICLC significantly influenced the kinetics, magnitude, and quality of responses elicited by vaccination, resulting in integrated responses of antibody, CD8\(^+\) and CD4\(^+\) T cells for NY-ESO-1 in nearly all patients. Patients receiving vaccination with OLP in the presence Montanide and Poly-ICLC developed significantly higher, earlier, and more frequent antibody, CD4\(^+\) and CD8\(^+\) T-cell responses to NY-ESO-1 than those vaccinated with OLP in Montanide alone or without adjuvant. Patients in this study had typical characteristics of second or third ovarian cancer remission, and experienced multiple prior chemotherapies. The rationale for inclusion of patients without NY-ESO-1 expression was to retrospectively evaluate the impact of expression on safety and immune responses. In addition, beside heterogeneity of expression that may lead to false-
negative results, the frequency of NY-ESO-1 expression in cancer is known to increase with stage and grade, so that some patient may express NY-ESO-1 upon recurrence even if their original tumor was considered negative. NY-ESO-1 expression in the tumor was detected in 9 of 26 (35%) of patients, within expectations, and it did not seem to affect the safety or immunogenicity of the vaccine. Treatment with NY-ESO-1 OLP was well tolerated regardless of adjuvant, and was not associated with any clinical systemic toxicities attributable to autoimmunity. An increase in local injection site reactions was seen in Cohort 3 patients receiving both Montanide and Poly-ICLC and led to the discontinuation of vaccinations in this cohort. The skin reactions may have been related to the volume of diluent required for the administration of both adjuvants and seemed to be a local irritant phenomenon, as observed in other studies (23).

Strong antibody responses against vaccine OLP, which excluded the most immunogenic N-terminal region for naturally induced antibodies, were induced in most patients after vaccination with OLP in Montanide, especially when
combined with Poly-ICLC. Titers were sustained even in patients who did not complete all planned immunizations. These results indicate that the antigen in a form of long peptide is a strong immunogen that induces antibody responses even against less immunogenic parts of the protein.

Vaccination with OLP coemulsified in Montanide and Poly-ICLC was also able to induce consistent antigen-specific CD8+ T cells in nearly all patients. This is in contrast to vaccination with whole NY-ESO-1 protein, against which less than 50% of patients developed CD8+ T-cell responses when coemulsified in Montanide with Cpg7909, or even in delivery systems that facilitate cross-presentation of a formulated protein (12-14). While full-length protein vaccines usually result in only a limited repertoire of NY-ESO-1-specific CD8+ T cells in half of the treated patients (9, 12, 24), OLP vaccines emulsified in Montanide with Poly-ICLC generally induced frequent CD8+ T cells recognizing multiple epitopes and in some cases naturally processed NY-ESO-1. Although OLP vaccine induced frequent polyclonal-specific CD8+ T cells, these were found to have a wide range of avidities (Fig. 3), including high-avidity responses which were sometimes obscured by low-avidity cells following in vivo expansion, as observed in short-peptide vaccination (25). Combination vaccination with recombinant DNA or viruses (26), shown to induce high-avidity responses, or using a reduced dose of OLP vaccine should be explored to favor the expansion of high-avidity CD8+ T cells rapidly induced by OLP vaccine.

Vaccination with OLP in Montanide efficiently and consistently induced NY-ESO-1–specific CD4+ T cells recognizing multiple epitopes in vaccine OLP, regardless of NY-ESO-1 tumor expression. The responses were often elicited after a single vaccination, something not previously observed with short-peptide or protein vaccination (12, 27). Surprisingly, CD4+ T-cell responses against the NY-ESO-11-40 region, which is weakly immunogenic for T cells and thus was excluded from the vaccine, were induced in several patients, especially in the presence of Poly-ICLC (Fig. 2B). Potentially, this epitope spreading could be induced by NY-ESO-1 protein released from NY-ESO-1–expressing residual tumors. Another possibility is that proinflammatory cytokines induced by Poly-ICLC allowed the expansion of NY-ESO-1–specific CD4+ T-cell precursors, by interfering with the in vivo suppression by Tregs (28). We are currently investigating whether qualitative differences exist between CD4+ T-cell responses specific for epitopes outside and within the NY-ESO-1 vaccine sequences.

Poly-ICLC is a viral mimic in man, upregulating a broad variety of immune-related genes representing multiple innate immune pathways in a pattern that closely parallels that of a live virus vaccine (29). This innate immune activation is likely playing an important role in the nature and extent of the responses that we show here with OLP. Poly-ICLC increased the kinetics and number of antigen-specific effectors (23, 30).

In summary, vaccination with OLP induced a rapid and strong integrated immune response orchestrated by both CD4+ and CD8+ T cells as well as antibodies. In this study, Montanide was necessary to induce immune responses after peptide vaccination, and Poly-ICLC further accelerated and enhanced these responses. It remains to be determined whether adding Poly-ICLC to a vaccine formulation may render Montanide dispensable, and this question may be answered in an ongoing-related clinical trial of NY-ESO-1 protein and Poly-ICLC with or without Montanide (identifier NCT01079741). No systemic toxicity related to vaccination was seen. Local injection site reactions in some patients who received Montanide and Poly-ICLC warrant the consideration of reducing the volume of administration of Montanide, Poly-ICLC, or both. Future trials should address clinical efficacy in patients with strong NY-ESO-1 tumor expression, with biopsies before and after treatment whenever possible to assess in situ immune responses. Vaccine strategies may be improved further by combining OLP vaccination with modulators of immunosuppression, such as anti-CITLA-4, anti-PD-1, or anti-GITR that are integrally important for the optimization of future cancer vaccines.

Disclosure of Potential Conflicts of Interest

A.M. Salazar is the CEO and Scientific Director of Oncovir Inc, which is developing Poly-ICLC. T. Tsuji, G. Ritter, L.J. Old, and S. Gnjatic are inventors on patents/patent application regarding the NY-ESO-1 antigen, and Ludwig Institute for Cancer Research has licensed such patents. No potential conflicts of interest were disclosed by the other authors.

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