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Abstract

Although checkpoint molecules like CTLA-4 and PD1 have been described several years ago, checkpoint inhibitors such as Nivolumab (an anti-PD-1 antibody) have only recently been used to treat classic Hodgkin lymphoma (cHL). Several studies have shown convincing therapeutic effects of Nivolumab in cHL. However, the mechanism of action of Nivolumab in cHL is not fully understood.

The aim of this study was to monitor changes in cell motility and cell contacts after administration of Nivolumab to an in vitro model of cHL as well as to native hyperplastic lymphoid tissue and native human tissue from cHL. In both tissue and in vitro, CD4+, CD8+, CD30+ and CD20+ cell velocities were unchanged after Nivolumab incubation. In contrast, in primary cHL tissue, the duration of cell contacts between CD4+ T cells and HRS cells was significantly increased after 5 h Nivolumab treatment, and the number of contacts with HRS cells was also slightly increased for CD4+ T cells (not significant), suggesting that CD4+ T cells in particular contribute to the cytotoxicity observed as a result of Nivolumab therapy. There was no change in the duration of cell contacts in the hyperplastic lymphoid tissue after Nivolumab incubation. In conclusion, we show here for the first time by imaging of native lymphoma tissue an enhanced interaction of CD4+ T cells and HRS cells in cHL after Nivolumab administration.
Introduction

The human immune system is a sophisticated structure that controls most types of infectious agents as well as developing malignant cells. However, on relatively rare occasions malignant cells manage to escape this highly efficient control system, leading to the development of a neoplasm. The first immune checkpoints, such as CTLA-4 and PD1, were discovered 20 years ago. It is only in recent years that therapies targeting these specific checkpoints have been developed. Checkpoint inhibitors are very effective, especially in tumors with a high mutational burden and expression of tumor-specific neo-antigens. Among lymphoid-derived tumours, classic Hodgkin lymphoma (cHL) is a neoplasm that often shows a good outcome after checkpoint inhibitor therapy. Patients with de novo early unfavourable cHL showed an overall survival rate of 100% after a median follow-up of 41 months in the NIVAHL trial. However, the exact mechanism that leads to the dissolution of HRS cells remains so far unknown.

To exert their anti-tumour activity, T cells must form a productive conjugate with their targets. In solid tumours, an inability of T cells to migrate and reach cancer cells has been described resulting in an "immune-excluded" tumour profile, which may explain why anti-PD1 antibodies do not always give the expected results. Although T cell motility is a critical factor influencing the success of immunotherapy, there is limited experimental data exploring the potential links between PD1 blockade and T cell migration, particularly in a tumour context.

In vitro co-culture experiments provided the first evidence that PD1 controls T cell motility and interaction time between T cells and antigen-presenting cells. In brief, it has been demonstrated that engagement with PD1 reduces TCR-induced signalling, thereby preventing T cells from establishing stable contact with antigen-presenting cells. Conversely, when PD1 binding to its ligand was blocked, T cells were able to regain stable contact with antigen-presenting cells. Intravital two-photon microscopy of a murine melanoma model showed that PD1 blockade restored stabilized conjugates between T cells and tumour cells.

As the efficacy of Nivolumab in the treatment of cHL has now been demonstrated in multiple studies, the aim of the present study was to further elucidate the direct effect of
Nivolumab on CD4⁺ and CD8⁺ lymphocytes, under reactive/inflammatory conditions and in cHL.

**Methods**

**In vitro model**

For the cHL *in vitro* model, the cHL cell lines L-428 (CVCL_1361) and L-1236 (CVCL_2096) were purchased from DSMZ (Braunschweig, Germany). Cell lines were transduced to express Life-Act GFP as previously described¹⁶ for a better visualization in microfluidics microchannels.

CD4⁺ and CD8⁺ T cells were purified from peripheral blood by magnetic activated cell sorting (MACS) using the untouched CD4⁺ and CD8⁺ T cell Isolation Kits (Miltenyi Biotec, Bergisch-Gladbach, Germany). The purity of the isolated cells was confirmed by FACS to be > 94% for CD4⁺ T cells (CD4-Vioblue antibody, Miltenyi Biotec) and > 89% for CD8⁺ T cells (CD8-PE/Cy7-antibody, Clone SK1, Biolegend, San Diego, CA, USA). T cells were either kept in culture overnight with IL-2 supplementation to recover from the MACS procedure or activated with dynabeads human T-Activator CD3/CD28 (Gibco, Thermofisher Scientific, Waltham, MA, USA) for 48 hours and then seeded at 2 x 10⁵ cells in a microchannel punch. Nivolumab or an IgG4 kappa isotype control antibody (MedChemExpress, Monmouth Junction, NJ, USA) was added at a dose of 10 µg/ml to the cell culture medium that was used to resuspend the cells. The other punch on the opposite side of the 4 x 10 µm microchannels was loaded with 2 x 10⁷ cells of either L-428 or L-1236 cell lines (see Fig. 1A). Time-lapse images were taken every 4 min overnight. The velocity of the cells was measured as previously published.¹⁶ The duration and number of cell-cell interactions was manually analyzed. To assess the expression of immune checkpoint proteins on activated T cells, T cells were cocultured overnight with the L-1236 cell line or an L-1236 PD-L1 knockout cell line at a ratio of 20:1. The expression of PD1, LAG-3, TIM-3 and TIGIT before and after coculture was determined by flow cytometry (Suppl. Methods).

Suggestions for text editing were generated by artificial intelligence (DeepL). All suggestions were checked by the authors for accuracy.
Live cell imaging
For 4D imaging, thick sections from thirteen cases of hyperplastic lymphatic tissue from the pharyngeal tonsil and six cases of cHL were analysed as previously described. CHL cases 1-3 and 6 were nodular sclerosing subtype and EBV-negative, the 4th and 5th case were EBV-positive mixed cellularity (Suppl. Table 1). From the 5th case only movies in the untreated condition could be obtained due to limited tissue size. The study was conducted according to the declaration of Helsinki and informed consent was obtained from all patients. The ethic committee of Goethe University Hospital agreed on the study (Nr20-376aV). The tissue was stained with fluorescent antibodies against CD4 (T helper cells, clone SK3, NovaFluor Blue 585, ThermoFisher Scientific), CD8 (Cytotoxic T lymphocytes, clone SK1 AlexaFluor 647, Biolegend), CD20 (B cells, Clone L26, AlexaFluor 488, ThermoFisher Scientific), and CD30 (Clone BerH2, sc-19658 AF488, Santa Cruz Biotechnologies, Santa Cruz, CA, USA). In each case, Nivolumab was added at a dose of 100 µg/ml on top of the tissue slices and then overlayed with RPMI medium. In the cases of hyperplastic lymphoid tissue an IgG4 kappa isotype antibody was added as negative control in addition to the untreated condition.

Results

CD4+ resting T cells show prolonged cell contacts with cHL cell line L-1236 in microfluidic microchannels after incubation with Nivolumab

To model the effects of Nivolumab on T cells and HRS cells, we first used a model system with microfluidic microchannels, where the different cell types (i.e. T cells vs. HRS cells) can enter the microchannels from opposite sides (Figure 1A). cHL cell lines L-428 and L-1236 were used, which both show expression of PD-L1 (Figure 1B) and loss of expression of major histocompatibility complex (MHC) class I (L-428) or II (L-1236, Suppl. Figure 1) as frequently observed in HRS cells in primary cHL cases. In general, activated T cells moved at a higher velocity (means 3.51 - 5.01 µm/min, Figure 1D) when compared with resting T cells (means of 1.93 - 4.3 µm/min Figure 1C). There
was no systematic difference in velocity when either Nivolumab or the IgG4 kappa isotype control antibody was added to the cell culture medium. The duration of cell-cell contacts with HRS cells of the L-1236 cell line was significantly increased for resting CD4$^+$ T cells in the presence of Nivolumab when compared to the IgG4 kappa isotype control (mean of 96.00 vs. 15.64 min, Figure 1E, G and H, p<0.01, Kruskal-Wallis-test). In addition, contacts with activated CD8$^+$ T cells were significantly longer compared to the IgG4 kappa isotype control (mean of 121.00 vs. 15.84 min, Suppl. Figure 2, p<0.01, Kruskal-Wallis-test). In the L-428 cell line there was no effect of Nivolumab treatment on the duration of cell-cell contacts (Figure 1 F, Suppl. Figure 2 for activated T cells).

Nivolumab treatment of activated T cell cocultures has no effect on the expression of LAG-3, TIM-3 and TIGIT, while PD1 expression becomes undetectable

Expression of the immune checkpoint proteins PD1, LAG-3, TIM-3 and TIGIT was assessed by flow cytometry in activated CD4$^+$ and CD8$^+$ T cells before and after overnight coculture with the cHL cell line L-1236. The aim was to determine if the expression of other checkpoint proteins is altered after Nivolumab treatment. Activated T cells were used because relevant PD1 expression was only observed in activated T cells (Suppl. Figure 3). Both PD1 and LAG-3 expression was reduced in the cocultured CD4$^+$ and CD8$^+$ T cells in the presence of the IgG4 kappa isotype control antibody (Suppl. Figure 4). However, there was no difference in LAG-3 expression between Nivolumab and IgG4 kappa isotype treated cocultures. Using a PD-L1 knockout L-1236 cell line for the T cell cocultures, LAG-3 expression was significantly higher in the activated CD4$^+$ T cells compared to the activated CD4$^+$ T cells cocultured with the L-1236 control cell line (33.88 vs. 18.04% LAG-3 expression, Suppl. Figure 4D and E, p<0.05, Mann-Whitney-test), while all other checkpoint proteins were unchanged when using PD-L1 knockout L-1236 (Suppl. Figure 4).
Duration of T-cell contact with B-cells is not affected in normal lymphoid tissue after Nivolumab treatment

We then examined the motility of CD4⁺ and CD8⁺ T cells and CD20⁺ B cells in native thick sections of native pharyngeal tonsil hyperplastic tissue, as we also observed PD-L1 expression in normal lymphoid tissue (Suppl. Figure 5). Time-lapse movies were taken immediately and after 3 and 5 hours. While the velocity of CD4⁺ and CD8⁺ T cells as well as CD20⁺ B cells was almost unchanged at 3 and 5 h with and without Nivolumab, generally T cells were significantly faster when compared with B cells (means 3.77 and 3.40 µm/min for CD4⁺ and CD8⁺ T cells vs. 2.39 µm/min for B cells, p<0.05, Kruskal-Wallis-test with Dunn´s post-test for multiple comparisons). Similarly, there were no changes in track length or displacement for CD4⁺, CD8⁺ T cells or B cells after Nivolumab incubation (data not shown). After 5 hours of incubation with the IgG4 kappa isotype control antibody, the velocity of CD4⁺ T cells was significantly higher compared to the untreated tissue slices (p<0.05, Kruskal-Wallis test, Figure 2A). This difference may be attributed to the limited sample size and the different areas of imaging, whether within or outside of follicles. In all other settings there was no effect on velocity of either Nivolumab nor IgG4 kappa isotype control antibody.

Cell-cell contacts in untreated tissue were shortest when CD8⁺ cells were involved (CD8-CD20 mean 1.30 min, CD4-CD8 mean 1.65 min), whereas the duration of CD4-CD20 cell contacts under baseline conditions was longer (mean 3.72 min) and had a higher variance, reflecting the physiologic interaction of CD4⁺ T cells and CD20⁺ B cells in follicles and interfollicular areas. Overall, there was no clear effect of Nivolumab on cell contact duration in normal lymphoid tissue (Figure 2D, E and F). However, CD4-CD20 cell contacts were significantly longer after 5h of IgG4 kappa isotype control antibody incubation compared to the Nivolumab treated tissue slices (5.08 vs. 0.74 min, p<0.01, Kruskal-Wallis-test with Dunn's post-test for multiple testing, Figure 2D). This may be due to the limited number of samples and different areas of imaging.

**HRS cells in cHL present enhanced cell-cell contacts with CD4⁺ T cells after Nivolumab treatment**
Next, we incubated thick slices of native tissue from cHL-affected lymph nodes with Nivolumab and took movies at baseline and after 3 and 5 h of Nivolumab incubation. All cases had PD-L1 positive HRS cells and bystander cells (Suppl. Figure 6). The velocity of CD4⁺ T cells (2.87 - 4.94 µm/min), CD8⁺ T cells (3.54 - 4.07 µm/min) and HRS cells (1.94-3.13 µm/min) was largely unchanged after 3 and 5h of Nivolumab incubation (Figure 3A). Similarly, track length and displacement were essentially unchanged in all cell types after 3 and 5h of Nivolumab incubation (data not shown). The number of cell-cell contacts between CD4⁺ T cells and HRS cells was increased after 5h Nivolumab incubation (mean 1 contact/CD4⁺ T cell/movie after 5h Nivolumab vs. mean 0.45 contact/CD4⁺ T cell/movie at baseline, not significant, Figure 3B). No differences were seen in the number of cell contacts between CD8⁺ T cells and HRS cells or between CD4⁺ and CD8⁺ T cells after Nivolumab incubation (Figure 3B). Regarding the duration of cell-cell contacts, a significant increase in the duration of cell-cell contacts between CD4⁺ T cells and CD30⁺ HRS cells was observed after 5h Nivolumab incubation (Figure 3C and E, after 5h Nivolumab mean 5.89 min vs. untreated mean 3.20 min, p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple comparisons). At the same time, the duration of cell contacts between CD8⁺ T cells and CD30⁺ HRS cells was significantly increased after 5h of Nivolumab incubation (mean 3.73 min vs. mean 1.11 min in untreated setting, p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple comparisons). There was no difference in the duration of cell contacts between CD4⁺ and CD8⁺ cells in the untreated vs. Nivolumab-treated groups (Figure 3C).

When only cHL cases with MHC I or II-positive HRS cells were considered (Suppl. Table 1, Suppl. Figure 7-9), the increase in cell-cell contact duration between CD4⁺ T cells and HRS cells remained significant after 5h Nivolumab treatment. Differences in the duration of CD8⁺ and HRS cell contacts were abolished, when only the subgroups of MHC I- or II-positive or -negative cases were considered.

When examining the duration of cell contact between CD4⁺ and CD8⁺ T cells and HRS cells after 5 h of Nivolumab incubation, it was found that there was a significantly longer duration of contact between CD4⁺ T cells and HRS cells in MHC II-negative cHL cases compared to MHC II-positive cHL cases (mean of 8.31 min in MHC II-negative cases vs.
4.51 min in MHC I-positive cases, p<0.05, Mann-Whitney test, Figure 3D). The duration of contacts between CD8⁺ T cells and HRS cells was also longer in cHL cases with MHC I-negative HRS cells (mean 5.15 min in MHC I-negative vs. 1.59 min in MHC I-positive cHL cases, not significant).

Figure 4 shows examples (from case 1) where long tracks with little displacement were seen in most cell types after 5h of Nivolumab treatment.

**Discussion**

This is the first study to demonstrate the dynamic effects of the anti-PD1 antibody Nivolumab on cell motility and cell-cell interactions of T cells, B cells and HRS cells in cHL using live cell imaging. Previously observed differences in the cell dynamics of immune cells¹⁷ were confirmed in this study, with T cells moving significantly faster than B cells in hyperplastic lymphoid tissue.

As PD1 has been described to be expressed on non-malignant CD4⁺, CD8⁺ and CD20⁺ cells¹⁹, we would potentially expect to see effects in all these cell types as PD-L1 was almost ubiquitously expressed in the tissue (Suppl. Figures 5 and 6). However, overall cell motility was not altered after Nivolumab incubation in either microchannels, hyperplastic lymphoid tissue or primary cHL tissue, suggesting that there is no direct effect of Nivolumab on cell motility.

While the duration of cell-cell contacts between CD4⁺ T cells and B cells in in hyperplastic lymphoid tissue was unchanged after Nivolumab treatment, resting CD4⁺ T cells in microchannels showed significantly longer cell-cell contacts with HRS cells of the cHL cell line L-1236 as well as CD4⁺ T cells with HRS cells in primary cHL tissue. Although the microchannel system is more artificial than imaging of primary tissue, this suggests that the situation in cHL is generally different from that in hyperplastic lymphoid tissue. While data from the literature suggests that the IFN-gamma pathway is active in both hyperplastic tonsillar tissue²⁰ and cHL²¹,²², there are several other differences between hyperplastic lymphoid tissue and cHL, such as the composition of the microenvironment, the frequent lack of MHC-I and -II expression in HRS cells, the tumour mutation burden, which has been observed to be on average 3.57/Mb in cHL nodular sclerosing type²³, as well as differences in chemokine secretion in the
neoplastic HRS cells of cHL. In this regard, tumour mutational burden, microsatellite instability, expression of tumour neoantigens, epigenetic silencing, expression of chemokines and receptors such as CCR5, CXCL9 and CXLC13, have been described to be associated with response to checkpoint inhibitor therapy. These findings may explain the divergent observation of cell-cell interaction after Nivolumab treatment in our study and also the large variance between samples, as each patient or tissue has individual factors that influence the efficacy of checkpoint inhibition. HRS cells from cHL strongly secrete various chemokines such as CXCL9, CCL5, CCL17 and CCL22 and thus attract both CD4+ and CD8+ T cells. While an inferior outcome after checkpoint inhibition was observed in patients with loss of MHC I protein expression in HRS cells of cHL, we found rather prolonged cell contacts of CD8+ T cells and HRS cells in cHL with MHC I loss (not significant). Similarly, the duration of cell-cell contacts between CD4+ T cells and HRS cells was significantly longer when MHC II expression was absent in HRS cells. This suggests that the effects of Nivolumab are, to some extent, independent of MHC expression.

The in vitro cHL model can be extended in the future to study T cell activation by visualizing intracellular Ca2+ signalling and measuring T cell cytotoxicity. The importance of CD4+ T cells in mediating the effects of Nivolumab has previously been demonstrated in several studies, including the significant expansion of the CD4+ T cell receptor repertoire in the peripheral blood of patients responding to Nivolumab treatment. In addition, lymph node biopsies taken one week after start of Nivolumab therapy showed little change in the frequency of CD4+ and CD8+ T cells in the cHL microenvironment, despite the frequent disappearance of neoplastic CD30+ HRS cells. Another study found that clonally expanded T cells in untreated cHL lymph nodes were rare, were exclusively CD8+, had a non-naive immune phenotype and only a minority of them expressed classical immune checkpoint molecules. Another study showed that MHC II expression on the HRS cells was predictive of response to Nivolumab treatment whereas no such correlation was observed for MHC I expression on HRS cells. One hypothesis for how Nivolumab might work is that subsets of rosetting CD4+ T cells, which contribute to the growth support of the HRS cells, move away from the HRS
cells, resulting in the removal of growth support. Immunological synapses were observed between HRS cells and rosetting T cells in this regard.\textsuperscript{35} The phenotype of CD4\textsuperscript{+} T cells with regard to LAG-3, TIM-3 or TIGIT expression did not change after Nivolumab incubation in our \textit{in vitro} coculture model of cHL. On the one hand, this model has the limitation that CD4\textsuperscript{+} and CD8\textsuperscript{+} T cells were purified from peripheral blood and did not undergo long-term coculture with HRS cells. On the other hand, the phenotype of CD8\textsuperscript{+} T cells after checkpoint inhibition in a melanoma patient, studied on single cell level in the literature, also did not change with respect to LAG-3, TIM-3 or TIGIT expression.\textsuperscript{36} Co-inhibition of other checkpoint proteins may therefore be an additional therapeutic option.

In general, most studies, including ours, have focussed on the effects of Nivolumab on the activity of CD4\textsuperscript{+} or CD8\textsuperscript{+} T cells. Effects on other cell types such as histiocytes, macrophages and NK cells have not been studied in cHL and may provide explanations for the remaining unanswered questions about the effect of Nivolumab. In a mouse model, it was observed that anti-PD1 antibody can be captured from CD8\textsuperscript{+} T cells by macrophages leading to resistance to checkpoint inhibition.\textsuperscript{37}

In conclusion, Nivolumab altered cell-cell interactions between CD4\textsuperscript{+} T cells and CD30\textsuperscript{+} HRS cells in cHL. The significant prolongation of cell-cell contacts between CD4\textsuperscript{+} T cells and HRS cells may be one component of the successful mode of action of Nivolumab in cHL. In this study, the effects of Nivolumab were found to be independent of the expression of MHC I or II on HRS cells. Future studies, analysing the correlation between Nivolumab-induced changes in motility, cell-cell contacts and outcome after therapy are warranted.
References


Figure Legend

Figure 1. Effects of Nivolumab on T cell motility in microchannels in the presence of Hodgkin-Reed-Sternberg (HRS) cells

A. Schematic overview of experimental setup with 4x10 µm microchannels connecting two reservoirs containing either CD4$^+$ or CD8$^+$ T cells (left side) and classic Hodgkin lymphoma (cHL) cell lines (right side).
B. PD-L1 expression in the classic Hodgkin lymphoma cell lines L-428 and L-1236 (black) compared with isotype control (grey)
C. Velocity of resting CD4$^+$ and CD8$^+$ T cells in microchannels with and without Nivolumab or IgG4 kappa isotype control in presence of classic Hodgkin lymphoma cell lines
D. Velocity of activated CD4$^+$ and CD8$^+$ T cells in microchannels with and without Nivolumab or with IgG4 kappa isotype control in presence of classic Hodgkin lymphoma cell lines
E. Duration of cell-cell contacts between Hodgkin-Reed-Sternberg cells of cell line L-1236 and resting CD4$^+$ or CD8$^+$ T cells with or without Nivolumab or with IgG4 kappa isotype control
F. Duration of cell-cell contacts between Hodgkin-Reed-Sternberg cells of cell line L-428 and resting CD4$^+$ or CD8$^+$ T cells with or without Nivolumab or with IgG4 kappa isotype control
G. Example of several short duration contacts between a resting CD4$^+$ T cell and a Hodgkin-Reed-Sternberg cell without Nivolumab (Hodgkin-Reed-Sternberg cell green, CD4$^+$ T cell red)
H. Example of a long duration contact between a resting CD4$^+$ T cell and a Hodgkin-Reed-Sternberg cell with Nivolumab (Hodgkin-Reed-Sternberg cell green, CD4$^+$ T cell red)

Figure 2. Cell velocity and duration of cell-cell contacts in hyperplastic lymphoid tissue

Each dot represents the mean of a movie. Typically, three movies per condition were obtained.
A. Velocity of CD4$^+$ T cells in hyperplastic lymphoid tissue (from thirteen donors) with and without Nivolumab or IgG4 kappa isotype control antibody; * p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple testing
B. Velocity of CD8$^+$ T cells in hyperplastic lymphoid tissue (from thirteen donors) with and without Nivolumab or IgG4 kappa isotype control antibody
C. Velocity of CD20$^+$ B cells in hyperplastic lymphoid tissue (from thirteen donors) with and without Nivolumab or IgG4 kappa isotype control antibody
D. Duration of cell-cell contacts between CD4$^+$ T cells and CD20$^+$ B cells in hyperplastic lymphoid tissue (from thirteen donors) with and without Nivolumab or IgG4 kappa isotype control antibody; * p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple testing
E. Duration of cell-cell contacts between CD8\(^+\) T cells and CD20\(^+\) B cells in hyperplastic lymphoid tissue (from thirteen donors) with and without Nivolumab or IgG4 kappa isotype control antibody

F. Examples for a short cell contact (left) between a CD8\(^+\) T cell (arrow head) and a CD20\(^+\) B cell (arrow, green cell) and longer duration cell contact (right) between a CD8\(^+\) T cell (arrow head, red cell) and a CD20\(^+\) B cell (arrow, green cell); CD8\(^+\) T cells labelled in red, CD20\(^+\) B cell labelled in green.

Figure 3. Nivolumab incubation extends cell-cell contacts between CD4\(^+\) T cells and Hodgkin-Reed-Sternberg (HRS) cells in classic Hodgkin lymphoma.
The study analysed six cases of classic Hodgkin lymphoma, including five cases in both untreated and Nivolumab-treated conditions, and one case in untreated condition only. Each dot represents the mean of a movie. Typically, three movies per condition were obtained.

A. Velocity of CD4\(^+\) and CD8\(^+\) T cells as well as CD30\(^+\) Hodgkin-Reed-Sternberg cells in classic Hodgkin lymphoma in untreated condition (0h) and after incubation with Nivolumab for 3 or 5 hrs

B. Number of cell-cell contacts between individual CD4\(^+\) T cells/movie and CD30\(^+\) Hodgkin-Reed-Sternberg cells, individual CD8\(^+\) T cells/movie and CD30\(^+\) Hodgkin-Reed-Sternberg cells as well as CD4\(^+\) and CD8\(^+\) T cells in classic Hodgkin lymphoma in untreated condition (0h) and after incubation with Nivolumab (3h and 5h)

C. Duration of cell-cell contacts between CD4\(^+\) T cells and CD30\(^+\) Hodgkin-Reed-Sternberg cells, CD8\(^+\) T cells and CD30\(^+\) Hodgkin-Reed-Sternberg cells as well as CD4\(^+\) and CD8\(^+\) T cells in classic Hodgkin lymphoma in untreated condition (0h) and after incubation with Nivolumab (3h and 5h). * p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple testing

D. Duration of cell-cell contacts between CD4\(^+\) T cells and CD30\(^+\) Hodgkin-Reed-Sternberg cells as well as CD8\(^+\) T cells and CD30\(^+\) Hodgkin-Reed-Sternberg cells after 5h Nivolumab incubation. Cases were stratified based on MHC I and II expression in the Hodgkin-Reed-Sternberg cells. * p<0.05, Kruskal-Wallis-test with Dunn’s post-test for multiple testing

E. Examples for a short cell contact (left) between three CD4\(^+\) T cells (arrow heads) and a CD30\(^+\) Hodgkin-Reed-Sternberg cells (#) in untreated condition and long duration cell contact (right) between a CD4\(^+\) T cell (arrow head) and a CD30\(^+\) Hodgkin-Reed-Sternberg cell (#); CD4\(^+\) T cells labelled in pink, CD30\(^+\) Hodgkin-Reed-Sternberg cell labelled in green.

Figure 4. Cell tracks in classic Hodgkin lymphoma without and with Nivolumab
Top row shows images (merged, CD4 pink, CD8 red, CD30 green) and the respective tracks of the cells in a 15 min movie of untreated classic Hodgkin lymphoma tissue.
Bottom row shows images (merged, CD4 pink, CD8 red, CD30 green) and the respective tracks of the cells in a 15 min movie of classic Hodgkin lymphoma tissue from the same case after incubation with Nivolumab for 5 h. Speed scale bars in µm/sec.
Supplementary Figures and Methods

Supplementary Figures

Suppl. Figure 1. MHC-I and –II expression in classic Hodgkin lymphoma cell lines L-428 and L-1236

Upper row: MHC-I expression as determined by FACS (bold line) versus isotype control (dotted line). The diffuse large B-cell lymphoma cell line TMD-8 was used as positive control.

Bottom row: MHC-II expression as determined by FACS (bold line) versus isotype control (dotted line). The diffuse large B-cell lymphoma cell line TMD-8 was used as positive control.
Suppl. Figure 2. Duration of cell-cell contacts between Hodgkin-Reed-Sternberg (HRS) cells and activated T cells

A. Duration of cell-cell contacts between HRS cells of cell line L-1236 and activated CD4⁺ or CD8⁺ T cells with or without Nivolumab or with IgG4 kappa isotype control
* p<0.05, Kruskal-Wallis test

B. Duration of cell-cell contacts between HRS cells of cell line L-428 and activated CD4⁺ or CD8⁺ T cells with or without Nivolumab or with IgG4 kappa isotype control
Suppl. Figure 3. PD1 expression by flow cytometry in T cells
PD1 expression by flow cytometry in resting and activated CD4+ and CD8+ T cells used in microchannel experiments, compared to isotype control
Suppl. Figure 4. Expression of the immune checkpoint proteins PD1, LAG-3, TIM-3 and TIGIT in an in vitro coculture model of classic Hodgkin lymphoma.
A. Examples of a 20:1 overnight coculture of activated CD4\(^+\) T cells and HRS cells from the L-1236 cell line in presence of Nivolumab or IgG4 kappa isotype control antibody. Gating strategy for the assessment of immune checkpoint protein expression.

B. Expression of PD1, LAG-3, TIM-3 and TIGIT in activated CD4\(^+\) T cells alone and after overnight coculture with HRS cells from the L-1236 cell line in presence of Nivolumab or the IgG4 kappa isotype control antibody.

C. Expression of PD1, LAG-3, TIM-3 and TIGIT in activated CD8\(^+\) T cells alone and after overnight coculture with HRS cells from the L-1236 cell line in presence of Nivolumab or the IgG4 kappa isotype control antibody.

D. Expression of PD1, LAG-3, TIM-3 and TIGIT in activated CD4\(^+\) T cells after overnight coculture with HRS cells from a PD-L1 knockout L-1236 cell line or the L-1236 control cell line in presence of Nivolumab.

E. PD-L1 expression in the PD-L1 knockout L-1236 cell line and the L-1236 control cell line at baseline.
Suppl. Figure 5. PD-L1 expression in hyperplastic lymphatic tissue from the pharyngeal tonsil. Images from four representative of 13 donors. A,C,E,G: 40-fold magnification of four representative examples of hyperplastic lymphoid tissue investigated by live cell imaging. B,D,F,H: 100-fold magnification of the same cases.
Suppl. Figure 6. PD-L1 expression in primary cases of classic Hodgkin lymphoma studied by live cell imaging.
A, C, E, G, I, K: 10-fold magnification of PD-L1 staining of the five cases investigated by live cell imaging.
B, D, F, H, J, L: 300-fold magnification of the same cases.
Suppl. Figure 7. MHC-II (HLA-DPB1) expression in the six classic Hodgkin lymphoma cases investigated by live cell imaging
A.-F. Cases 1-6 25 x magnification. Insert: Representative Hodgkin-Reed-Sternberg cells in 60x magnification.
Suppl. Figure 8. MHC-I (HLA-ABC) expression in the six classic Hodgkin lymphoma cases investigated by live cell imaging
A.-F. Cases 1-6 25 x magnification. Insert: Representative Hodgkin-Reed-Sternberg cell in 60x magnification.
Suppl. Figure 9. Cell-cell contact duration according to MHC expression in Hodgkin-Reed-Sternberg (HRS) cells of the classic Hodgkin lymphoma cases investigated.

A. Cell-cell contact duration between CD4+ T cells and HRS cells, CD8+ T cells and HRS cells and CD4+ and CD8+ T cells in MHC I positive cHL cases (n= 3), each dot represents the mean of one movie

B. Cell-cell contact duration between CD4+ T cells and HRS cells, CD8+ T cells and HRS cells and CD4+ and CD8+ T cells in MHC I negative cHL cases (n= 3), each dot represents the mean of one movie

C. Cell-cell contact duration between CD4+ T cells and HRS cells, CD8+ T cells and HRS cells and CD4+ and CD8+ T cells in MHC II positive cHL cases (n= 3), each dot represents the mean of one movie

D. Cell-cell contact duration between CD4+ T cells and HRS cells, CD8+ T cells and HRS cells and CD4+ and CD8+ T cells in MHC II negative HRS cells
D. Cell-cell contact duration between CD4$^+$ T cells and HRS cells, CD8$^+$ T cells and HRS cells and CD4$^+$ and CD8$^+$ T cells in MHC II negative cHL cases (n= 3), each dot represents the mean of one movie.

Suppl. Table 1. MHC I and II expression by immunohistochemistry in classic Hodgkin lymphoma cases investigated by live cell microscopy
NS: Nodular sclerosing subtype, MC: Mixed cellularity subtype

<table>
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<tr>
<th>Case number</th>
<th>Subtype</th>
<th>MHC-I (HLA-ABC)</th>
<th>MHC-II (HLA-DPB1)</th>
</tr>
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<tbody>
<tr>
<td>1</td>
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<td>+</td>
<td>+</td>
</tr>
<tr>
<td>2</td>
<td>NS, EBV-</td>
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</tr>
<tr>
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<td>+</td>
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</tr>
<tr>
<td>6</td>
<td>NS, EBV-</td>
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<td>+</td>
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Supplementary Methods

Flow cytometry of classic Hodgkin lymphoma (cHL) cell lines
Non-transduced cHL cell lines were stained with antibodies against PD-L1 (Clone 29E.2A3 PE, Biolegend, San Diego, CA, USA), HLA-ABC (Clone G46-2.6, APC, ThermoFisher Scientific, Waltham, MA, USA) HLA-DR, DP, DQ (Clone Tu39 FITC, BD Biosciences, Heidelberg, Germany), and isotype controls and analyzed by fluorescence activated cell sorting (FACS) in a MACSQuant (Miltenyi Biotec, Bergisch Gladbach, Germany).

Cocultures of activated T cells with L-1236
After 48h of activation with dynabeads human T-Activator CD3/CD28 (Gibco, Thermofisher Scientific), activated CD4⁺ or CD8⁺ T cells were seeded in a 24-well plate with L-1236 cells (or L-1236 PD-L1 knockout cells) at a ratio of 20:1. Overnight culture was performed in the presence of Nivolumab or IgG4 isotype control antibody (10 µg/ml). After 24 h, cells were stained with either CD4-Vioblue or anti-CD8-Vioblue (both Miltenyi Biotec) and the following antibodies:
CD223 (LAG-3) Antibody, anti-human, APC, REAfinity (130-119-567), CD366 (TIM-3) Antibody, anti-human, APC, REAfinity (130-119-781), TIGIT Antibody, anti-human, APC, REAfinity (130-116-815), REA Control Antibody (S), human IgG1, APC, REAfinity (130-113-434, all Miltenyi), APC anti-human CD279 (PD-1) Antibody, clone EH12.2H7 (Biolegend) and Isotype APC Mouse IgG1, κ Isotype Ctrl Antibody MOPC-21 (both Biolegend).

Cells were gated on live population and the respective CD4⁺ or CD8⁺ T cell population and expression of the respective checkpoint proteins was assessed (Suppl. Figure 4A).

PD-L1 knockout L-1236 cell line
The PD-L1 knockout L-1236 cell line was obtained from Prof. Roland Schmitz, Gießen:
Vector construction
pLVX-TRE3G-Cas9-Hygro was created by removing the puromycin resistance gene from pLVX-TRE3G (Clontech Laboratories) with BfuAI/KpnI and replacing it with a synthetic Hygro construct. Cas9 was isolated using PCR from LentiCrispr v2 (Addgene #52961) and cloned into pLVX-TRE3G-Hygro digested with BamHI using Gibson cloning. SgRNA sequences targeting CD274 (ACATGTCAGTTCATGTTCAG) or AAVS1 control (GGGGCCACTAGGGACAGGAT) were cloned into pLenti guide dsRed (Addgene #128055) using BsmBI.

**Generation of Doxycycline-inducible Cas9 expressing L-1236 clones**

L-1236 cells were transduced with pLVX-Tet3G (Clontech Laboratories). 72 hours post-transduction, cells underwent selection with G418. Selected cells were transduced with pLVX-Tre3G-Cas9-Hygro. After 3 days of infection, transduced cells were split and incubated with hygromycin. A fraction of the selected pLVX-Tre3G-Cas9-Hygro cells were transduced with a sgRNA vector targeting nonessential surface markers such as ICAM1. 250 ng/mL of doxycycline was added to induce Cas9 and initiate genetic ablations for 8-11 days. The surface expression of the target gene was measured by FACS. After functional validation of Cas9, single-cell clones were generated by using limiting dilution from the pool of Cas9-expressing cells.

**Lentivirus production and transduction**

To generate lentivirus, HEK293FT cells (ThermoFisher) were cultured in DMEM medium supplemented with 10% FBS, 2 mM L-glutamine, and 1% penicillin-streptomycin. One day before transfection, HEK293FT cells were seeded in a 10 cm tissue culture dish at 60% confluency. Cells were transfected using TransIT-293 transfection reagent (Mirus). For each dish, 50 µl serum-free OptiMEM (Gibco) was mixed with 6 µg sgRNA vector, 4.5 µg psPax2 (Addgene #12260) and 1.5 µg pMD2.G (Addgene #12259). 27 µl TransIT-293 transfection reagent was diluted with 950µl OptiMEM, incubated for 5 min at room temperature, and combined with the mixture of plasmids. This transfection mixture was incubated for 30 min at room temperature and then added dropwise to the cells. Virus-containing supernatant was collected 72h post-transfection. Cell debris was removed by
centrifugation at 500xg for 10 min followed by filtration of the supernatant through a 0.45µm low-protein binding membrane (Millipore Steriflip HV/PVDF). Viral supernatant was concentrated by incubation with Lent-X concentrator (Takara) for at least 3h at 4°C. Virus was concentrated 100x, resuspended in sterile PBS, aliquoted, and frozen at -80°C. L-1236 doxycycline-inducible Cas9 cell line was transduced with lentivirus, and after 3 days of transduction, dsRed was measured using FACS. 250 ng/mL doxycycline was added for 8-11 days to induce Cas9. Cells were passaged every three days with fresh medium containing doxycycline and the surface expression of the targeted gene (CD274, clone 29E.2A3) was measured by FACS. Transduced post-doxycycline cells underwent two rounds of FACS sorting.