Development of a novel human phage display-derived anti-LAG3 scFv antibody targeting CD8⁺ T lymphocyte exhaustion

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Abstract

Background: Lymphocyte-activation gene (LAG3) is a 498 aa transmembrane type I protein acting as an immune inhibitory receptor. It is expressed on activated lymphocytes, natural killer cells and plasmacytoid dendritic cells. In activated lymphocytes, LAG3 expression is involved in negative control of cell activation/proliferation to ensure modulation and control of immune responses. In view of its deregulated expression in tumor-infiltrating lymphocytes, LAG3, together with the additional immune checkpoint inhibitors CTLA4 and PD1, is considered a major target in order to reverse the immunosuppression typically mounting in oncologic diseases. Since many patients still fail to respond to current immune checkpoints-based therapies, the identification of new effective immune inhibitors is a priority in the ongoing fight against cancer.

Results: We identified a novel human single-chain variable fragment (scFv) Ab against a conformational epitope of LAG3 by in vitro phage display technology using the recombinant antigen as a bait. This scFv (referred to as F7) was characterized in terms of binding specificity to both recombinant antigen and human LAG3-expressing cells. It was then rebuilt into an IgG format pre-optimized for clinical usage, and the resulting bivalent construct was shown to preserve its ability to bind LAG3 on human cells.

Next, we analyzed the activity of the anti-LAG3 scFvF7 using two different antigen-specific CD8⁺ T lymphocyte clones as target cells. We proved that the reconstituted anti-LAG3 F7 Ab efficiently binds the cell membrane of both cell clones after peptide-activation. Still more significantly, we observed a striking increase in the peptide-dependent cell activation upon Ab treatment as measured in terms of IFN-γ release by both ELISA and ELISPOT assays.

Conclusions: Overall, the biotechnological strategy described herein represents a guiding development model for the search of novel useful immune checkpoint inhibitors. In addition, our functional data propose a novel candidate reagent for consideration as a cancer treatment.

Keywords: LAG3, Single-chain variable fragment, Phage display, Reconstituted scFv, CD8⁺ T lymphocytes, Lymphocyte exhaustion
Background

Lymphocyte activate gene (LAG3) (also known as CD223) is a transmembrane protein showing about 20% identity with CD4 (for a recent review, see [1]). Like CD4, LAG3 binds MHC Class II molecules, albeit at a distinct site and with a greater affinity [2, 3]. LAG3 is not expressed by resting T cells, although it is upregulated on activated T lymphocytes (both CD4+ and CD8+ subpopulations), B lymphocytes, natural killer cells, and plasmacytoid dendritic cells (DCs). Besides MHC Class II molecules, LAG3 has been found to bind Galectin-3 [4, 5], LSECtin [6] and, in central nervous system, α-synuclein [7]. LAG3 engagement leads to a negative regulation of T-cell activation, despite the LAG3 signal transduction mechanism being still largely unknown. A couple of unique, yet largely conserved domains, i.e., the EP [8] and KIEELE [9] motifs, seem important for the transmission of intracellular inhibitory signals. In addition, LAG3 shows a bidirectional signaling activity, as inferred by the modulation of DC activation/maturation observed upon interaction with LAG3 expressing lymphocytes [10, 11].

LAG3 has been identified as one of the major inhibitory receptors (IR) involved in the T lymphocyte exhaustion typically occurring in oncologic diseases [12]. For this reason, specific anti-LAG3 mAbs are currently under investigation in the attempt to reverse cancer-associated immunosuppression. Together with other IRs (i.e. CTLA4 and PD1) LAG3 is now considered a major target of the so called immune checkpoint blockade (ICB) antitumor strategy based on delivery of mAbs against IRs [13]. This therapeutic design has been positively adopted in the field of oncology, providing persistent clinical benefits for a significant number of patients with advanced cancer. Unfortunately, however, many patients fail to respond to ICB-based therapies, and the hyper-immune activation can associate with immune-related adverse events affecting several organs, including skin, gut, heart, lungs, and bone [14]. Therefore, nowadays the emphasis is on the evaluation of new immune checkpoint inhibitors and a combination thereof possibly increasing efficacy while also reducing toxicity [15], all resulting in a considerable interest in LAG3.

IRs maintain self-tolerance and modulate the duration and amplitude of physiological immune responses. It is now clear that tumors could co-opt certain regulatory pathways as a major mechanism of immune resistance, particularly in CD8+ T lymphocytes, and that this dysfunction could be reversed by anti-IRs antagonist mAbs [16]. Meanwhile, awareness is emerging that multiple IRs can be involved in lymphocyte exhaustion [17–19], so paving the way for more effective combinatorial approaches [20, 21]. A first confirmation of this hypothesis came from the fact that the combination of two IR blockers, namely anti-PD1 and anti-CTLA4 mAbs, resulted in a higher rate of response than that observed with single mAb treatments. In solid tumors, LAG3 is upregulated on tumor-infiltrating lymphocytes (TILs), while blockade of LAG3 can enhance antitumor T cell responses. On this basis, several reagents targeting LAG3 are currently being tested in various stages of preclinical and clinical development, thus attesting to the broad interest of this IR in the field.

Monoclonal Abs (mAbs) are molecules largely used in therapy, diagnostic and biotechnology. Among the various recombinant mAb formats, the single chain variable fragment (scFv) is one of the simplest and most versatile constructs that retains the antigen-binding activity [22]. This molecule is formed by the variable regions of both heavy and light chains joined by a peptide linker. While scFvs have the significant advantage that they can be expressed by a single open reading frame, their low stability and short pharmacokinetic may limit their applications. The scFv fusion with CH2 and CH3 domains of the Ig Fc region can overcome such drawbacks [23].

To avoid undesirable immune responses, human Abs represent the most suitable reagents for Ab-based therapeutic purposes. Human scFvs can be readily isolated through the well consolidated phage display technique. In this paper we describe the identification of a novel human anti-LAG3 scFv Ab recovered from a fully human naive phage library by means of an in vitro bio-panning strategy. Upon Fc-rebuilding, this Ab was shown to maintain its binding activity, while strongly increasing the activation of antigen-specific CD8+ T lymphocytes. We propose our anti-LAG3 Ab, in both scFv and reconstituted forms, as a candidate for novel biotechnology and clinic applications.

Results

Isolation of a novel human anti-LAG3 scFv ab by phage display technology

Human recombinant scFvs against LAG3 were obtained by in vitro affinity enrichment process (bio-panning) as previously described [24], and depicted in Fig. 1a. In essence, an aliquot of phage-displayed scFvs representing the complexity of the naive IORISS1 library [25] was subjected to cycles of binding to recombinant LAG3 protein used as a bait and immobilized on immunotubes, followed by elution. Outputs after each round of selection were characterized by a progressive increase in phage titers (Table 1). Five rounds of selection were necessary to isolate a phage population detectable in anti-LAG3 ELISA. Soluble scFvs derived from single colonies were obtained after plating bacteria TG1 infected with the selected phages. Several of these scFvs were demonstrated to bind the recombinant LAG3 protein (Fig. 1b), whereas they were negative against an irrelevant antigen (GO, glucose oxidase) used as a negative control (data not shown). The scFv coding DNA was isolated from the positive clones, and the sequencing revealed a common genetic identity (Fig. 2).
A representative clone, referred to as scFvF7, was produced in bacteria and purified by immobilized metal affinity chromatography using 6× histidine-tag located at its C-terminus. SDS-PAGE analysis showed a single 31 kilodalton (kDa) band, at the expected size of the Ab fragment, with a negligible presence of both contaminant and degraded products (Fig. 3a).

The purified scFv was then tested for binding with recombinant LAG3 antigen in ELISA. The binding assay generated a specific dose-dependent association with LAG3 and a lack of binding against GO protein used as an irrelevant antigen (Fig. 3b).

Concerning the binding activity of scFvF7 on LAG3-expressing eukaryotic cells, flow cytometric analysis on CD4+ T lymphocytes carried out following a 3-day stimulation with phytohaemagglutinin (PHA) showed that scFvF7 binds native LAG3 with a comparable profile to that of a commercial murine anti-LAG3 Ab used as a control (Fig. 3c). Therefore, scFvF7 is capable of recognizing LAG3 antigen when it retains its native conformation, namely in ELISA on recombinant protein and in cytometry on live LAG3-expressing cells. Conversely, scFvF7 failed to react against its target in SDS-PAGE western blot assay (Fig. 3d), whose reducing/denaturing conditions stretch the amino acid structure of protein; this suggests a conformation-dependent epitope. An ELISA conducted on the heat-denatured LAG3 protein (by boiling) and a western blotting assay performed in non-reducing conditions strengthen this hypothesis (see Additional file 1: Figure S3D15).

**Engineering of divalent Fc-scFvF7 Ab**

The use of single-domain Abs overcomes some limitations typical of tetrameric Abs, e.g., unspecific uptake, restricted access to tissues of interest. On the other hand, scFvs have been shown to have low in vivo stability as a consequence of their efficient blood clearance. Hence, to increase the expected in vivo stability of scFvF7 in view of possible clinical applications, a divalent Fc-scFv was generated and assessed for its performance in an ELISA assay and a Western blot.

**Table 1** Selection of native IORISS1 Ab phage library against LAG3 recombinant protein

| Panning cycle | Ag µg/mL | Input | Output | Ratio (O/I) | Enrichment |
|---------------|----------|-------|--------|-------------|------------|
| 1             | 20       | 5×10¹² | 2×10⁴  | 0.4×10⁻⁴    | 1          |
| 2             | 10       | 1×10¹² | 6×10⁶  | 1.6×10⁻⁷    | 3×10       |
| 3             | 5        | 1×10¹² | 1.4×10⁷ | 1.4×10⁻⁵    | 7×10²      |
| 4             | 5        | 1×10¹² | 4×10⁹  | 4×10⁻⁵      | 2×10²      |
| 5             | 2.5      | 1×10¹² | 7.5×10¹⁰ | 7.5×10⁻²     | 3.7×10⁶    |

*both Phage input (I) and output (O) were determined by titration in terms of colony forming units in infected E. coli TGI cells before and after each round of selection. Enrichment was calculated as ratio between outputs from each cycle and the output from the first one.*
use, its fusion with CH2 and CH3 Fc domains was carried out. We generated a divalent scFvF7-Fc construct resulting in a recombinant Ab incorporating two scFvF7 molecules, each of them joined with CH2 and CH3 constant immunoglobulin domains. Similar to classic Abs, their spontaneous dimerization results in two anti-LAG3 binding sites on the same molecule, with the important advantage that the divalent molecule can be expressed by a single open reading frame (Fig. 4a). The presence of a hinge region between the scFv and the constant domains guarantees a flexibility similar to that of natural immunoglobulins. In addition, LALA mutations within the CH2 constant regions would be particularly useful for a possible use in vivo since they are expected to lower the risk of unwanted Fc-dependent effects, e.g., Ab-dependent cell cytotoxicity. The scFvF7 open reading frame was cloned in frame with CH2 and CH3 Fc sequences in the context of the pFUSEss-CHIghGI vector which also expresses the human IgG1 secretory leader peptide sequence (Fig. 4a). The resulting molecular construct was transfected in CHO cells in order to verify the proper synthesis and secretion of the construct in an eukaryotic system. The western blot assay performed under non-reducing conditions on the supernatants of the transfected cells using anti-human Fc as detector showed a dominant signal compatible with the expected molecular weight (~140 kDa) of the reconstituted anti-LAG3 Ab (Fig. 4b). These supernatants efficiently and specifically reacted in LAG3 ELISA (Fig. 4c).

Next, the Fc-reconstituted F7 Ab was produced on a large scale by transient transfection in CHO cells, purified, and formulated in sterile PBS. The purified divalent Ab construct preserved its specificity as evidenced by both its reactivity in ELISA against recombinant LAG-3 (Fig. 5a), and its binding on human HEK 293 cells stably expressing LAG3 (Fig. 5b). Finally, FACS assays proved that the reconstituted F7 Ab binds human PHA-activated CD8+ T cells specifically, and in a concentration-dependent manner (Fig. 5c).

Fc-reconstituted F7 Ab binds peptide-activated, antigen-specific CD8+ T lymphocytes

Besides CTLA4 and PD1, LAG3 is now considered a major target for the mAb-based cancer immunotherapy.
ICBs are selected on the basis of their capacity to reverse the lymphocyte-dependent immune exhaustion, i.e., a typical signature affecting TILs infiltrating solid tumors, and particularly the CD8+ T sub-population [12]. In this context, the F7 Ab’s potential as a novel ICB candidate for new antitumor combination therapies relies on the assessment of its activity on antigen-specific CD8+ T lymphocytes. To this end, we carried out functional assays on a couple of antigen-specific CD8+ T cell clones, i.e., an HLA-B7 restricted clone specific for HIV-1 Nef, and an HLA-A.02-restricted clone specific for the human melanoma antigen Mart-1.

Initially, we were interested in assessing the detection LAG3 expression with the reconstituted F7 Ab following the activation of the CD8+ T cell clones. The two clones were co-cultivated with HLA-matched B-lymphoblastoid cell lines (B-LCLs) previously treated with 100 ng/mL of each specific peptide. As a positive control, CD8+ T lymphocytes isolated from healthy donor peripheral blood mononuclear cells (PBMCs) were treated with PHA for three days. Stimulated cells were incubated respectively with the anti-Flag antibody and the HRP-conjugated anti-mouse antibody, in order to monitor eventual non-specific signals due to the secondary antibodies used to detect scFv antibodies. The ELISA shown below (performed with the same secondary Abs) is a check for the reactivity of the scFv-containing supernatants (scFvGO and scFvF7) used in the assay. Arrows indicate relevant signals. Molecular markers in kilodaltons (kDa) are reported on the right. A representative of three independent assays is shown.
anti-LAG3 Abs (either the reconstituted scFv or a murine mAb) in 100 μL of total volume. After an overnight (o.n.) incubation, supernatants were harvested, and IFN-γ production was evaluated by ELISA. The results reported in Fig. 7a (and Additional file 2: Figure S7AII, and Additional file 3: Figure S7AIII) clearly demonstrate that the treatment with the Fc-reconstituted scFvF7 increased the IFN-γ release even more efficiently than the murine anti-LAG3 mAb validated for functional activity. The effect is more pronounced in the Nef-specific CD8+ T cells, this finding being consistent with the apparently higher level of LAG3 detected by the divalent scFvF7 Fc compared to that of Mart-1 specific CD8+ T lymphocytes (Fig. 6).

The specificity of the functional effect of scFvF7-Fc is supported by the data presented in Fig. 7b (and Additional file 4: Figure S7BII), where the divalent Ab was incubated with cells in the presence or absence of increasing amounts of recombinant LAG3 protein. The F7 Ab induced IFN-γ production in a concentration dependent manner that was effectively inhibited with the addition of the soluble LAG3 antigen. The specific effect of the divalent F7 Ab on the CD8+ T cell activation was further confirmed by IFN-γ ELISpot assays (Fig. 7c and Additional file 5: Figure S7CII).

Taken together, our functional data indicate that the human divalent scFvF7-Fc Ab efficiently binds LAG3 on the surface of antigen-specific CD8+ T lymphocytes, thereby potently increasing their activation state. On the basis of its ability to counteract the LAG3 mediated inhibitory effect on antigen-activated CD8+ T lymphocytes, scFvF7-Fc can be proposed as a novel candidate for the treatment of pathologies characterized by LAG3-driven lymphocyte exhaustion.

Discussion
Here we describe the experimental process aimed at the identification of novel anti-LAG3 human scFvs based on the previously described IORISS1 library [25]. This procedure allowed us to isolate the anti-LAG3 scFvF7. The combination of the results we obtained from western blot assay and ELISA are consistent with scFvF7 binding a conformational epitope of LAG3. Furthermore, functional assays carried out on antigen-specific CD8+ T lymphocytes, scFvF7-Fc biological activity is comparable to the functionally characterized murine anti-LAG3 17B4 mAb. Considering the human origin of scFvF7, these results are of particular relevance in view of possible in vivo uses of
this scFv and derivatives thereof. In fact, monoclonal antibodies obtained with the classic hybridoma technology are extremely valuable instruments in many fields of application, but their use in the clinic is compromised by their xenogenic origin. Conversely, antibodies generated from human antibody libraries are entirely human, with the innate advantage of being only slightly or not at all immunogenic. Nevertheless, some drawbacks remain with the phage display-derived antibody fragments, such as their very short blood clearance. Therefore, the anti-LAG3 antibody here described was engineered in a recombinant ‘scFv-Fc’ format, in order to recover some important requirements for a potential drug compound, namely a structure with the typical avidity of natural antibodies (two binding sites), a prolonged half-life and a reduced ability to recall unwanted cell mediated responses [22, 23].

The interest in the identification and characterization of new ICBs is continually growing as witnessed by the varied clinical studies as well as by the quite recent publication of works describing the efforts to identify new anti-IR Abs. In particular, Everett et coll. Reported the isolation of
novel anti-LAG3 Abs able to target both human and murine LAG3 by means of an additional engineered binding domain within the CH3 Fc region [27]. The in vitro activity of these Ab constructs resulted in an increased release of IL-2 from stimulated PBMCs. In addition, Sasso and colleagues described an Ab selection procedure based on binding phagemid particles on activated human lymphocytes to generate a collection of scFvs against different IRs including LAG3 [28]. Upon reconstitution into human IgG4 Fc, the respective biological activities were evaluated through cytometry-based lymphocyte proliferation assays and cytokine ELISA.

The biologic activity of scFvF7 was evaluated by means of a functional evaluation of CD8+ T lymphocytes specific for either viral or tumor antigens. We focused our interest on CD8+ T cells for the efficacy of scFvF7 as a new ICB candidate since CD8+ T-driven immune reactivation is expected to have a significant therapeutic effect to counteract the exhaustion of this lymphocyte subpopulation in cancer [29] and even chronic infectious diseases [30].

Overall, it is now clear that the broad success of ICB-based immunotherapies will depend on the combination of different treatment options. In a rapidly evolving research landscape, the generation of a repertoire of different immune-modulatory Ab-based constructs could play an essential role in identifying novel therapeutic agents and LAG3 certainly represents one of the most appealing emerging targets. The great interest in LAG3 has been fueled by its sustained expression in dysfunctional T-cells associated with many human tumors [31–33]. In addition, there is strong evidence that LAG3 upregulation could mediate an escape mechanism from PD1 therapy, whose acquired resistance might be overcome by the combined treatment with anti-LAG3 ICBs [34–37]. Another more recent study, for its part, highlights how LAG-3 blockade could result in a superior T cell activation compared to other inhibitory pathways, including PD1/PDL-1 axis [38]. Furthermore, LAG3, together with TIGIT and Tim-3, has been predicted to have a better safety profile in the clinic compared to CTLA4 and PD1 combinations [1, 15]. Also, LAG3 blockade inhibits the suppressive activity of Treg cells.
Fig. 7 (See legend on next page.)
Methods

Ab phage library

The IORISS1 naive human Ab library [24] consists in a large array (i.e., more than \(10^9\) Ab combinations) of scFv polypeptides displayed on the surface of M13 phage. It was constructed by recovering light and heavy Ab chains from a group of healthy donors, then randomly joining them by means of an appropriate peptide linker, and cloning the resulting scFvs into a phagic vector as previously described [25]. It is noteworthy that the scFv inserts have been cloned upstream of two tag sequences, Flag tag and 6-his tag, respectively suitable to detect scFv in immunochemical assays and to purify them by nickel chromatography.

Biopanning strategy for selection LAG3 specific scFvs

LAG3-specific scFv Abs were selected from IORISS1 phage library as described elsewhere [25]. Briefly, a number of phages (~ \(10^{13}\) colony forming units, cfu), proper to represent the complexity of the library, was placed in an immunotube (Nunc Maxisorp) previously coated o.n. at room temperature (r.t.) with 4 mL of phosphate buffered saline (PBS) containing the recombinant LAG3 antigen (Acrobiosystem) at a concentration of 5 \(\mu\)g/mL. After 2 h of incubation, the specific phages were detached by an elution carried out with a solution of 100 mM triethylamine (1 mL), ten minutes later neutralized with 0.5 mL of 1 M Tris–HCl pH 7.4. Afterwards, the phages were amplified in TG1 \(E.\ coli\) bacteria [supE hsd15 thi 1(lac-proAB) F'(traD36 proAB+ lacIq lacZ1M15)], as previously reported [25]; in detail, phages were used to infect TG1 bacteria in exponential growth phase, and the latter distributed on a plate with solid selective medium consisting of 2xTY (Sigma), 2% glucose and ampicillin (100 \(\mu\)g/mL). The day after, 50 mL of liquid selective medium was inoculated with the bacteria grown on the selective plate in order to obtain a 600 nm optical density (O.D.) of 0.05–0.1; when the culture reached the O.D. of 0.4–0.5, the bacteria were re-infected with the M13 K07 helper phage in a ratio phages/bacteria of about 20:1, and incubated at 30°C o.n.. Finally, the phages released into the supernatant were precipitated by means of PEG (Poly-Ethylene Glycol) 6000 before being used for a new cycle of selection on the LAG3 antigen.
An aliquot of TG1 bacteria infected with the phages from each panning cycle were also induced with isopropyl β-D-thiogalactopyranoside (IPTG, Sigma) (final concentration >1 mM) and used to perform a polyclonal ELISA in order to monitor the enrichment of the LAG3-specific population (Table 1). In addition, bacteria infected with phages from the last panning was properly diluted and spread on selective plates in order to obtain single bacterial clones to be tested for their ability to secrete LAG3-specific scFvs.

For ELISA monoclonal screening, one hundred individual clones were inoculated in 96-well plates (Nunc) in a selective medium with 0.1% glucose; after 2 h of growth at 37 °C, clones were induced as above described and incubated for additional 12 h at 30 °C with shaking. Therefore, the scFv-containing supernatants were recovered, spun down (1800×g for 10 min) and used as primary antibodies against LAG3 protein.

DNA characterization and sequences
Plasmid DNAs from individual bacterial colonies were amplified and analyzed by restriction enzyme digestion using Nco I and Not I (i.e., the restriction sites used to clone the antibody fragments in IORISS1 library) to verify the presence of the scFv-encoding inserts. Afterwards, the scFv ORFs from the positive clones were sequenced with an automated DNA sequencer (Eurofins Genomics) using the following primers: scFv forward: 5′-atgaaacacaagcactattgcact-3′; scFv reverse: 5′-tgatatccacaacaagaaggg-3′. The sequences were finally analyzed using the IMGT database.

ELISA on recombinant LAG3 protein
For screening procedure, ninety-six-well ELISA plates were coated with 50 μL/well of recombinant LAG3 in PBS at 4 °C. The following day, a blocking solution consisting of 2% non-fat milk in PBS (MPBS) was added, and after 2 h the plates were washed with PBS containing 0.1% Tween 20 (TPBS). Plates were incubated for 2 h at r.t. with 50 μL of supernatants containing soluble scFv Abs, anti-flag M2 Ab (2.5 μg/mL, Sigma), and HRP-conjugated anti-mouse Ab (5 μg/mL, Dako). All Abs were resuspended in 2% MPBS. The reaction was visualized with 3,3′,5,5′-tetramethylbenzidine (BM blue, POD substrate, Roche), and stopped by adding 50 μL of 1 M sulfidric acid. The reaction was detected by an ELISA reader (Model 680 microplate reader, BioRad), and results expressed in terms of absorbance (A) = A_{450} nm - A_{620} nm.

The same ELISA procedure described above was performed to verify the reactivity of the scFvs-containing bacterial supernatants that were used for the Western blot assays.

The coating step for the assays to test the transfected CHO-derived supernatants was performed, as previously with 0.5 μg of recombinant LAG3/well, while for the dose-response binding analysis of the purified Fc-reconstituted scFv, the antigen was lowered to 0.1 μg/well. In both cases, the reconstituted Ab was incubated for one hour at r.t., followed by the incubation with a HRP-conjugated anti-human Ab (Thermo Fisher Scientific). The remaining procedures were performed as above described.

Soluble scFv purification
For the production of soluble scFvs, specific phage-TG1 E. coli infected cells were cultured at 37 °C in 2 × TY medium containing 100 μg/mL ampicillin and 0.1% glucose up to 0.5 O.D. After induction of Ab expression by adding IPTG 1 mM, cells were incubated o.n. at 30 °C. Then, the bacterial cultures were centrifuged, and scFv-containing supernatants collected. ScFv Abs were precipitated with ammonium sulfate, and then dialyzed in PBS. 6 × His-tagged scFv Abs were purified by immobilized metal affinity chromatography using Ni$_2^+$-nitroloacetic acid agarose (Qiagen). ScFv fragments were eluted with 250 mM imidazole in PBS, dialyzed, aliquoted, and stored at −80 °C.

Eukaryotic cell cultures
PBMCs were obtained by Ficoll–Hypaque density gradient centrifugation from heparinized blood samples of healthy donors. Purified T cells were obtained by PBMC negative selection using a kit for human T cell selection (Miltenyi) according to the manufacturer’s protocol, and cultivated in RPMI medium supplemented with 10% of heat-inactivated fetal calf serum (FCS)/HyClone. Both CHO and HEK293T cells were cultivated in DMEM plus 10% FCS. Both HLA-A02 and HLA-B7 B-LCLs were grown in RPMI medium supplemented with 10% FCS. B-LCLs were generated by in vitro EBV infection of ex vivo B lymphocyte cultures. Both isolation and expansion of CD8+ T cell clones specific for Mart (melanoma-associated recognized by T cells)-1 and HIV-1 Nef have been previously described [41, 42]. Mart1-specific CD8+ T cells recognize the HLA-A02-restricted AAGIGILTV$_{27-35}$ peptide sequence, whereas Nef-specific CD8+ T cells recognize the HLA-B7-restricted TGPFPVRPL$_{128-137}$ peptide. Both antigen-specific CD8+ T-cell clones were cultivated in RPMI plus 10% AB human serum (Gibco), and regularly monitored for their specificity.

Flow cytometry assays on human T cells
T cells isolated from PBMCs were seeded at the concentration of 2 × 10⁶/mL, and activated with 5 μg/mL of PHA. During the screening phase, scFvF7 was preliminarily tested on TCD4 cells at a one shot concentration...
(50 μg/mL), just to have a ‘yes/no’ response about its ability to recognize the antigen on the cell surface. About 1 × 10^6 cells were resuspended in PBS containing primary scFv antibody and incubated for 1 h at r.t. After washing, cells were resuspended for 30 min at 4 °C in PBS containing 25 μg/mL of an anti-FLAG M2 mouse antibody (Sigma) and then incubated for another 30 min at 4 °C with 6 μg/mL of an FITC-goat anti-mouse IgG (Pierce, IL, USA). An irrelevant antibody directed to an irrelevant antigen (glucose oxidase, GO) was used as negative control. After staining, the cell samples were washed, maintained at 4 °C and immediately analyzed by FACScan (Becton-Dickinson, NJ, USA) equipped with 15 mW, 488-nm, argon laser. Fluorescence compensation was determined using samples stained with anti-gluco oxidase scFv and goat FITC-conjugated anti-mouse secondary antibody.

For the cytofluorimetric analysis on HEK293 T cells (both LAG3-negative parental and the derivative LAG3 expressing cell lines, kindly provided by Pantaleo’s team, Lausanne), 2.5 × 10^5 cells were incubated with Fc-reconstituted scFvF7 Ab (10 μg/mL) for 1 h at r.t., and after washing with an goat FITC-conjugated anti-human IgG1 (Pierce) for 30 min at 4 °C.

Similarly, both PBMC-derived T cells and antigen-specific CD8+ T lymphocytes (1 × 10^6 cells/test) were incubated with Fc-reconstituted anti-LAG3 scFv, at different concentrations as specified in each experimental assay (comprised between 0.1 and 20 μg/mL) for 1 h at r.t.; the murine 17B4 anti-LAG3 mAb (EnzoLab, 10 μg/mL) was used as control. Then, cells were washed and incubated with FITC-conjugated secondary either anti-human (for detecting Fc-reconstituted scFvF7) or anti-mouse (for detecting mouse 17B4 mAb) goat IgG1 (Pierce). In all cases, stained cells were analyzed on FACscan platform (BD Biosciences).

**Western blotting**

For the analysis of the scFv binding in Western blot assay, 0.5 μg of both recombinant LAG3 and, as control, glucose oxidase (GO), were loaded in multiple wells, run on 12% SDS-PAGE (under reducing or non-reducing conditions, as specified in each assay) and transferred to nitrocellulose membrane. Separate filter strips were then incubated for 2 h with bacterial supernatants from the anti-LAG3 clone scFvF7 and, as control, from a previously characterized anti-GO clone (scFvGO) [24], both of them previously tested in ELISA to verify their reactivities. Afterwards, the strips were washed and incubated again with 5 μg/mL anti-flag M2 mouse antibody (Sigma) in 2% MPBS. After an additional incubation for 1 h at r.t. in the presence of 5 μg/mL of a goat anti-mouse antibody HRP-conjugated (Dako), the reaction was developed and visualized using a chemiluminescence reagent (NEN Life Science Products). A mouse anti-6 × His mAb (Serotec, 1 μg/mL diluted in 2%MPBS) was used as positive control to detect the recombinant LAG3 protein (which is provided with a 6 × His-tag at its C-terminus).

For monitoring transfection efficiency of CHO cells with the vector encoding the Fc-reconstituted anti-LAG3 antibody, supernatants from transfected cells (50 μL/lane) were separated by 10% SDS-PAGE, and then transferred to nitrocellulose membrane. After blocking with 5% MPBS, the membranes were washed and incubated for 1 h at r.t. with HRP-conjugated, goat anti-human IgG (1:2000 dilution, (Thermo Fisher Scientific)) and developed as described.

**Genetic construction and production of the divalent anti-LAG3 scFvF7-Fc Ab**

The anti-LAG3 scFv coding sequence was amplified from IORISS1 phagemide vector [24] by PCR with tailored primers. The amplified fragment was then digested with Xho I and Eco RI enzymes, gel purified, and ligated in frame to the Xho I/Eco RI double digested pFUSEss-Chlg-HG1 vector (Invivogen) expressing the sequences of both human Fc CH2, including LALA mutations [26], and CH3 Fc domains. The resulting construct was transfected into CHO cells for pilot production of bivalent scFvF7-Fc Ab. Briefly, 10^6 cells were transfected with 1 μg of plasmid DNA per well in a six-well tissue culture plate using Lipofectamine Plus reagent (Life Technologies) according to the manufacturer’s instructions. Supernatants were collected 48 h later to evaluate Ab production through both western blot analysis and ELISA.

**Large-scale production of the divalent scFvF7-Fc Ab**

Divalent scFvF7-Fc Ab was produced through transient transfection of CHO DG44 cells followed by incubation for 6 days in ProCHO5 serum free medium (Lonzon). The scFvF7-Fc Ab was purified from the cell medium using a protein A column (Thermo Fisher) with Abs eluted with 100 mM glycine buffer pH 3.0 into a 1 M Tris-HCl eluate at pH 8.0. Abs were dialyzed twice against PBS, concentrated using a JumboSep centrifuge filter with 3 kDa molecular weight cut-off (Pall Laboratories), and sterile filtered with Millex GP 0.22 μm pore size (Millipore). Prior to evaluation of scFvF7-Fc in the in vitro functional recovery assays, Abs were tested with the FDA-licensed Endosafe-PTS kit (Charles River Laboratory) which showed < 5 EU of endotoxins per mg of protein.

**IFN-γ ELISA and ELISPOT assays**

CD8+ T cells were pre-incubated with different concentrations of the reconstituted anti-LAG3 scFv for 2 h, and then co-cultured at a 2:1 ratio with HLA-matched B-LCLs previously pulsed with 100 ng/mL of the appropriate peptide. The co-cultures were incubated o.n., and
finally the supernatants were clarified and assayed for IFN-γ content by ELISA (Immunological Sciences). For ELISPOT assays, the co-cultures were carried out for 16 h in ELISPOT microwells previously coated with a mAb against human IFN-γ (clone D1K, Mabtech). Afterwards, the cells were removed, and a biotinylated Ab against human IFN-γ added, followed by the addition of a streptavidin-alkaline phosphatase. The plate was then developed using BCIP/NBT substrate (Sigma). Spot-forming cells were analyzed and counted using an ELISPOT reader (Amplimedical Bioline A-EL-VIS GmbH).

**Statistical analysis**

Statistical analyses were performed by using GraphPad Prism version 5.01 software. Data are expressed as means ± standard error of the mean (SEM). The student’s t test was used to determine significance. A p value of < 0.05 was considered statistically significant.

### Supplementary information

**Supplementary information** accompanies this paper at https://doi.org/10.1186/s12896-019-0559-x.

- **Additional file 1:** Figure S3D
- **Additional file 2:** Figure S7A
- **Additional file 3:** Figure S7A
- **Additional file 4:** Figure S7B

### Abbreviations

B-LCLs: B-lymphoblastoid cell lines; DCs: Dendritic cells; ELISA: Enzyme-linked immunosorbent assay; ELISPOT: Enzyme-linked immunospot; Fc: Fragment crystallizable; FCS: Fetal calf serum; GO: Glucose oxidase; HLA: Human leukocyte antigen; IC: Immune checkpoint blockers; IFN: Interferon; Ig: Immunoglobulin; IPTG: Isopropyl β-D-thiogalactopyranoside; IR: Inhibitory receptor; kDa: Kilodaltons; LAG3: Lymphocyte activate gene-3; mAb: Monoclonal antibody; O.D.: Optical density; o.n.: Overnight; PBMCs: Peripheral blood mononuclear cells; PBS: Phosphate buffered saline; PHA: Phytohaemagglutinin; r.t.: Room temperature; scFv: Single-chain variable fragment; SE: Standard error; TILs: Tumor-infiltrating lymphocytes

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### Authors’ contributions

AA isolated and characterized the scFvF7, participated in the design of the research and drafted the manuscript; CA performed the experiments with CD8+ T cell clones; AM worked on the design and genetic construction of the diabody scFvF7-Fc; MFL carried out biochemical assays and binding studies with scFvs; MG performed western blot analyses; MA analyzed data from flow cytometry analyses and interpreted the results; CF and GP produced and characterized the divalent scFvF7; SV coordinated and supervised the study design on the basis of the scFv isolation and divalent antibody construction; MFE wrote the paper and supervised the cell immunology experiments. All authors have seen and agreed to the final version of the paper. All authors read and approved the final manuscript.

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### Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

### Ethics approval and consent to participate

Human specimens used in some experiments described in our ms are PBMCs from healthy donors. Multiple stocks of human peripheral blood mononuclear cells (PBMCs) isolated from a large number of healthy donors (n > 20), have been stored in liquid nitrogen in our laboratory. The PBMCs have been purified from buffy coats obtained from the Transfusion Center of the University of Rome “La Sapienza” as waste material resulting from the procedures of plasma/platelet/red blood isolation from whole blood of healthy donors. They were volunteers and consented for their donated blood to be used for research. In line with the Italian law on this matter (Legislative Decree of the Italian Ministry of Health of January 25, 2001, published on the Official Gazette of April 3, 2001) donor ages ranged between 21 and 60 years for women and 21–65 years for men. There have been no restrictions for blood donors as to sex and ethnic background, except for children, mentally disabled individuals and pregnant women. All donors were healthy, non-medicated consulting adults, and were pre-screened for exposure to infectious agents. Blood was obtained by venipuncture following standard medical procedures using sterile technique and a compression bandage in the area of blood removal. A record of the donor and hematocrit result is stored on file at the Transfusion Center. Research investigators do not have any access to personal data or other information concerning the donors in agreement with the Italian law Number 675 of December 31, 1996.

### Consent for publication

Not applicable.
Competing interests
Alessandro Ascione (AA), Michela Pieggo (MFL), Alessandra Mallano (AM), Stefano Vella (SV) and Maurizio Federico (MFE) are co-inventors of a patent recently filed, regarding the human monoclonal antibody against LAG3 described in the present manuscript (Date of patent filing: 12/07/2019. Title: Humoral recombinant antibody against LAG3 membrane receptor, medical and diagnostic uses thereof). Apart from that the authors declare that they have no competing interests.

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