Construction, characterisation and kinetics of a single chain antibody recognising the tumour associated antigen placental alkaline phosphatase

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Summary The murine monoclonal antibody H17E2 recognises placental alkaline phosphatase (PLAP), an antigen present in the human term placenta and also expressed by many tumours. The antibody is of value in both immunoscintigraphy and radioimmunotherapy in testicular and ovarian cancer.

Radiolabelled monoclonal antibodies have been investigated by many centres for the experimental diagnosis and treatment of malignant disease (Maraveyas & Epenetos, 1991). However the properties of murine IgG result in several diagnostic and therapeutic problems. The relatively large size of an IgG antibody (~150 kDa) can be a factor in limiting penetration into tumours, leading to areas of sub-optimal uptake (Jain, 1990). The long serum half life (3–5 days, Stewart et al., 1989) of radiolabelled intravenously administered monoclonal antibody can result in nontarget tissues, particularly the bone marrow, receiving toxic doses of radiation (Epenetos et al., 1986; Vaughan et al., 1987). Prolonged blood residence also reduces the tumour: blood ratio thus limiting the imaging information available. A further clinical problem is that of immunogenicity. Repeated administrations of murine monoclonal antibody often lead to patients making an antibody response, generating human anti-mouse antibodies (HAMA) (Schroff et al., 1985). This HAMA response leads to rapid clearance of subsequent antibody administrations, with immune complex formation and the risk of anaphylaxis (Courtenay-Luck et al., 1986).

Univalent Fab and bivalent F(ab')2 antibody fragments produced by proteolysis of antibody show increased tumour penetration, clear more rapidly (Kennel et al., 1991) but remain immunogenic, can accumulate in the kidney and are difficult to produce (Milenic et al., 1989). Single chain antibodies (SCAs), each comprising one variable heavy- and one variable light-chain domain of immunoglobulin joined by a polypeptide linker, provide an alternative approach to antibody miniaturisation (Huston et al., 1988). In addition to their small size (25–30 kDa) SCAs have a number of advantages over IgG. They can be produced economically in bacteria, their specificity can be selected by in vitro ‘immunisation’ (Marks et al., 1991) and they can be manipulated by genetic engineering to form anti-tumour fusion proteins incorporating additional effector functions (Chaudary et al., 1989; Savage et al., 1993).

The anti-PLAP murine monoclonal antibody H17E2 recognises the tumour associated antigen placental alkaline phosphatase which is expressed in a large number of cases of testicular and ovarian cancers (Travers & Bodmer, 1984; Epenetos et al., 1984). It shows no cross-reactivity with human liver and intestinal alkaline phosphatases, allowing it to be used effectively in both immunoscintigraphy and radioimmunotherapy (Epenetos et al., 1985, 1986). In an effort to avoid the HAMA response H17E2 has been ‘reshaped’ by grafting its CDRs onto human framework regions (Verhoeyen et al., 1991). This antibody, Hu2PLAP, has demonstrated clinical benefits in early trials (Hird et al., 1991).

We describe here the construction, expression, and characterisation of a single chain antibody based on the murine monoclonal antibody H17E2 and report early preclinical in vivo results in nude mice bearing human xenografts.

Materials and methods

Plasmid construction

The cDNA sequences encoding the VH and VK domains were cloned from the hybridoma secreting monoclonal antibody H17E2, and were used to replace the corresponding VH and VK sequences in pSW2, a plasmid designed for expression of the Fv derivative of the anti-hen egg lysozyme antibody D1.3 (Ward et al., 1989), yielding the plasmid pFvPLAP. A bacterial colony transformed with pFvPLAP was boiled in 500 μl of water for 5 min. A sample (5 μl) of the cleared supernatant was subjected to 30 rounds (94°C, 1 min; 55°C, 1 min; and 72°C, 1 min) of polymerase chain reaction (PCR)-mediated amplification using VH1 and VK1 oligonucleotide primers (Orlandi et al., 1989), using a PEC ampliqa kit. Reaction products were digested with appropriate restriction enzymes, gel-purified and the VH and VK fragments were cloned into the appropriate sites of the plasmid pSWsFvD1.3myc (McCafferty et al., 1990). These regions were sequenced in a number of the progeny plasmids, and one with the expected sequence was designated pSH17E2.2myc. This encodes a SCA derived from H17E2 under the transcriptional control of the lac promoter. Since H17E2 SCA retains the c-myc antigenic tag derived from pSWsFvD1.3myc, it can be detected with the antibody 9E10 (Evan et al., 1985).
Expression and purification of H17E2 SCA

A 500 ml culture of E. coli KS476 (Stauch et al., 1989) transformed with plasmid pSH17E2.2myc was grown and expression of H17E2 SCA was induced as previously described (Ward et al., 1989). No biological activity of H17E2 SCA was detected in cell supernatants, and the c-myc antigenic tag used for SCA recognition could only be detected in the cell pellets (not shown). To produce functional SCA a refolding protocol was employed (George et al., 1993). Briefly, after pelleting, cells were disrupted by sonication and resuspended in 50 ml of 8 M urea. Insoluble material was removed by centrifugation and the soluble material dialysed against 0.1 M Tris base, 2 mM EDTA, 0.4 M arginine, pH 8.0. Insoluble material produced during dialysis was removed by centrifugation and the solution was then dialysed exhaustively against PBS.

Affinity purification of the refolded material was performed on PLAP (Calzyme, San Luis Calif.) immobilised on a Carbolink (Pierce, Rockford IL, USA) column. Crude refolded material was passed down the column under gravity at 4°C. After washing with PBS, the bound protein was eluted with 50 mM unbuffered diethylamine. Elution fractions were dialysed against PBS, concentrated in terms 1/10 volume 1 M Tris-HCl pH 7.5, and then dialysed against PBS and stored at 4°C.

PLAP binding assay, ELISA, SDS-PAGE and Western blot

Biotinylated H17E2 antibodies were immobilised on streptavidin-coated nylon pegs by incubation of the pegs in biotin-H17E2 solution for 1 h at room temperature. The pegs were rinsed in water and then placed in wells of a microtitre plate (Dynatech Immulon) predosed with PLAP and either competing SCA preparation or intact unlabelled H17E2 IgG. In the absence of these competing molecules, PLAP bound to the pegs via immobilised biotin-H17E2. The level of this binding could be assayed by the enzymatic activity of the bound enzyme. With either the SCA or unlabelled H17E2 IgG present, the PLAP binding was reduced in proportion to the concentration of the competing species. The competitive binding step was continued for 1 h at room temperature, then the pegs were washed to remove unbound PLAP. Bound PLAP was assayed by incubating the pegs for 1 h in microtitre plates containing 200 µl volumes of pNPP (1 mg ml⁻¹) in 2 M diethanolamine adjusted to pH 10.2. Substrate conversion was determined in terms of absorbance at 408 nm measured in Titertek MCC/340 plate reader.

Specific in vitro binding of the SCA to immobilised antigen was also confirmed by ELISA. Performed essentially as described previously (Savage et al., 1993) the plates were coated with either 100 µg ml⁻¹ PLAP, or lysozyme, BSA, FCS, insulin or milk powder at appropriate dilutions. Bound SCA was detected via its myc peptide tail with mAb 9E10 (Evant et al., 1985) and with HRP-conjugated anti-mouse antibody.

Analysis of SCA preparations was carried out by SDS-PAGE (Laemmli, 1970) using a 15% gel with a 3% stack. Coomassie blue staining was used for direct visualisation. For western blotting, proteins were transferred electrophoretically to a nitrocellulose membrane and probed with mAb 9E10 using AP-conjugated anti-mouse antibodies (Promega) for visualisation.

Radiolabelling and solid phase radioimmunoassay (RIA)

Samples of the SCAs H17E2 and TEL9 (gift of Dr T. Bonnert), (Marks et al., 1992) were labelled with 125I and 131I respectively using the Iodogen method of Fraker and Speck (1978). Unincorporated iodine was removed by gel filtration on a G25 Sephadex column (Pharmacia). SCA-containing fractions were pooled and the protein concentration and specific activity were measured.

To estimate the antigen binding affinity of the labelled SCA, a RIA and Scatchard analysis were used. Using 100 µl volumes throughout, flexible Titertek Elisa plates were coated overnight at room temperature with 100 µg ml⁻¹ PLAP in 50 mM bicarbonate buffer, pH 9.6. After washing in PBS, non-specific binding sites were blocked by incubation with a 1% solution of milk powder in PBS for 30 min at room temperature. Dilutions of radiolabelled SCA were applied and incubated for 1 h at room temperature. After further washes in PBS, bound radioactivity was measured by cutting out the individual wells and counting in a Minaxi 5550 gamma counter (Canberra Packard). Samples of the orginal dilutions were also counted to determine the total of bound and unbound counts at each dilution. To determine the binding affinity calculations were performed by Scatchard (1949) were performed.

In vivo pharmacokinetics

Tumour xenografts of the H.Ep-2 human epidermoid tumour cell line (Toolan, 1954) were produced by subcutaneous inoculation of female nude mice with 5 x 10⁶ cells. After 3 weeks growth, when the tumours measured 6–8 mm in diameter, 0.5 µCi (0.5 µg) of 125I-H17E2 SCA and 131I-TEL9 SCA were administered concurrently in 100 µl of PBS via a lateral tail vein. In parallel, 5 µCi of radiolabelled H17E2 IgG was injected into a similar group of mice. Mice were sacrificed at 1, 3, 5, 24 and 48 h post-injection by cardiac puncture and exsanguination. Samples of tumour and non-target tissues were weighed and incorporated radioactivity measured for both the 125I and 131I content in a gamma counter. Results are expressed as the percentage of injected dose per gram of wet tissue (percentage ID g⁻¹) and as tumour: blood ratios.

Results

Expression, refolding and purification of SCA H17E2

Since H17E2 SCA was expressed as an insoluble protein in E. coli, cells were sonicated, the sonicate was dialysed in 8 M guanidine hydrochloride and after dialysis against 0.1 M Tris base, 2 mM EDTA, 0.4 M arginine, pH 8.0 and then PBS, the soluble fraction was applied to an affinity column containing immobilised PLAP. Gel electrophoresis and Western blotting showed this simple refolding protocol to be very inefficient (Figure 1). Monoclonal antibody 9E10 (which recognises the polypeptide myc tag fused at the carboxyl end of the SCA) identified a single band at 30 kDa as expected. The material eluted from the column was subjected to SDS-PAGE and reduced conditions and then stained with Coomassie blue. Figure 2 shows again the presence of a single band at 30 kDa. In this initial series of experiments no attempt was made to optimise the yield of functional SCA and the total yield of purified material was only 100 µg.

Specificity and affinity of H17E2 SCA

Specificity of H17E2 SCA was demonstrated by a competition assay in which affinity-purified refolded SCA was able to inhibit competitively the binding of PLAP to immobilised biotinylated parent antibody H17E2. A dose-dependent decrease in signal occurred with increasing concentration of H17E2 SCA and with increasing concentrations of IgG H17E2 (Figure 3). Neither non-refolded crude H17E2 SCA (not shown) nor refolded anti-lysozyme SCA D1.3 had any ability to inhibit the binding of PLAP. In binding studies with a panel of different immobilised antigens (PLAP, hen egg lysozyme, BSA, FCS, insulin and milk powder), PLAP was the only material to which there was any significant binding of the H17E2 SCA with no evidence of any non-specific sticking to the panel of other antigens tested (not shown).

When the results of the RIA (Figure 4) for the estimation of the antigen binding affinity were subjected to Scatchard analysis, they give a value for the association constant of at least 10⁻⁸ M.
The studies described here confirm that a functional SCA has been generated from the variable domains of the anti-PLAP monoclonal antibody H17E2. The SCA was successfully expressed in bacteria, but the biologically active form was only produced after the use of a simple refolding protocol. The ability of this SCA to specifically bind to PLAP is demonstrated through the competitive binding studies with the parent IgG. The comparative binding studies with the panel of antigens indicates that this SCA is free of the non-specific stickiness that can undermine the usefulness of some recombinant antibody fragments (Ward et al., 1989).

The antigen binding affinity of this SCA (>10⁻⁸ M) is similar to those of other SCAs (Glockshuber et al., 1990; Colcher et al., 1990) but is reduced from the measured affinity of the parent IgG (M. Verhoeyen, unpub). This may result from constraints of the SCA linker which may alter slightly the native configuration of the antigen binding site. In this initial study the total yield of purified material was low, perhaps owing to the inefficiency of the simple refolding protocol used. This figure should be greatly improved by optimising the refolding protocol (McCartney et al., 1991). It should be possible to increase the final yield, as experience with Fab fragment produced by bacterial fermentation indicates that expression levels of over 500 mg L⁻¹ can be achieved (Better et al., 1990).

**Discussion**

In addition to their low cost of production and ease of manipulation, single chain antibodies offer a number of theoretical advances over IgG for *in vivo* applications. These advantages result from the smaller size of the SCA that lead to improved tumour penetration (Yokota et al., 1992) and more rapid clearance from the circulation (Colcher et al., 1990; Milenic et al., 1991). The studies described here confirm that a functional SCA has been generated from the variable domains of the anti-PLAP monoclonal antibody H17E2. The SCA was successfully expressed in bacteria, but the biologically active form was only produced after the use of a simple refolding protocol. The ability of this SCA to specifically bind to PLAP is demonstrated through the competitive binding studies with the parent IgG. The comparative binding studies with the panel of antigens indicates that this SCA is free of the non-specific stickiness that can undermine the usefulness of some recombinant antibody fragments (Ward et al., 1989).

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H17E2 SCA

TEL9 SCA

H17E2 IgG

Time post-injection (h)

Tumour: blood ratio

Figure 4 Scatchard plot of the radioimmunoassay measuring antigen binding affinity of H17E2 SCA.

Figure 5 Pharmacokinetics of H17E2 SCA and IgG and the non-specific SCA Tel9 in tumour bearing nude mice. Results are expressed as tumour:blood ratios. Means and standard deviations of replicate data-points are given.

In vivo, H17E2 SCA localised to the tumour bearing tumour associated antigens in marked contrast to non-specific TEL9 SCA, a result in agreement with previous reports of successful in vivo localisation of anti-tumour SCAs (Colcher et al., 1990).

This current work extends previous studies; here the observed binding is shown to be through specific uptake as there is no significant tumour binding of the non-specific SCA TEL9. This study is the first to compare the binding of concurrently administered specific and non-specific SCAs. Tumour:blood ratios achieved with use of a specific SCA are significantly improved over those from use of whole IgG and are observed much earlier. These results with the H17E2 SCA given tumour:blood values less than those reported with some other SCA preclinical studies. The potential reason for these discrepancies include; the relatively poor vascularity of H-ep 2 tumours in mice, the presence of circulating PLAP shed from the tumour and the fact that this SCA has not been optimised regarding its charge for optimal tumour penetration. However these early good ratios should be of considerable benefit in imaging and potential therapy, with improved and earlier definition occurring. The rapid blood clearance noted in previous reports on SCA kinetics is confirmed here. Although tumour:blood ratios obtained with specific SCAs are superior to those seen with IgG, the levels of absolute uptake are lower. This poses a potential difficulty for the use of radioimmunotherapy with SCAs. Optimisation of administration methods may overcome this. A steady state produced by continuous infusion over a longer period might result in higher absolute tumour levels. With the rapid renal clearance of the iodinated SCA it should be possible to control the levels very accurately to achieve maximal tumour uptake, while keeping the serum level below the threshold for non-target toxicity. As shown here, administration by bolus injection does not seem the optimal way to achieve therapeutic doses to a tumour. Another difficulty relates to the potential immunogenicity of the SCA, particularly of the linker sequence. This might be minimised by sequential use of SCAs with different linker sequences, a number of which have already been shown to be equally effective in producing functional molecules.

In summary the H17E2 SCA selectively binds antigen-positive tumour cells in vivo, and shows favourable pharmacokinetic behaviour for therapy and localisation. It is therefore an attractive candidate for a SCA-based fusion proteins (Chaudary et al., 1989; Savage et al., 1993) to target other effector functions to tumour cells.

Table 1 Tumour uptake and pharmacokinetics of specific (H17E2) and control (TEL9) single chain antibodies and H17E2 IgG. Means and standard deviations of replicate data-points are given.

| Time (h) | H17E2 SCA | TEL9 SCA | H17E2 IgG |
|---------|-----------|----------|-----------|
| 1       | 2.2 ± 0.4 | 2.2 ± 0.4 | ND        |
| 3       | 1.4 ± 0.2 | 1.6 ± 0.4 | ND        |
| 5       | 0.7 ± 0.2 | 0.9 ± 0.4 | ND        |
| 24      | 0.15 ± 0.04 | 0.08 ± 0.02 | 7.7 ± 1.1 |
| 48      | 0.18 ± 0.07 | 0.08 ± 0.02 | 9.3 ± 1.0 |
| T1 2 β (h) | 1.8 | 2.1 | 141 |
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