

Cellular processing in the SW1222 cell line of mAb A33 directly and indirectly radiohalogenated

JOHANNA HÖGLUND^{1,2}, ANNA ORLOVA², ANDERS SUNDIN¹,
HANS LUNDQVIST² and VLADIMIR TOLMACHEV²

Divisions of ¹Radiology and ²Biomedical Radiation Sciences, Department of Oncology,
Radiology and Clinical Immunology, Uppsala University, Uppsala, Sweden

Received December 20, 2005; Accepted March 2, 2006

Abstract. Investigations into the cellular processing of radio-labeled monoclonal antibodies (mAbs) for their further use in radioimmunodiagnosis and cancer therapy are needed in order to understand the fate of internalized and catabolized mAbs. The anti-colorectal cancer mAb, A33, was labelled with ⁷⁶Br and ¹²⁵I using the direct Chloramine-T method, or by labelling *N*-succinimidyl *para*-(tri-methylstannyl) benzoate and its further conjugation to the mAb. The cellular processing of the four conjugates was investigated in SW1222 cells *in vitro*. Uptake of mAb was rapid, peaking after 14-16 h. Intracellular degradation was slow and the early loss of radioactivity was due to dissociation of cell-surface bound mAb. The indirect labelling resulted in stronger binding of the mAb as well as prolonged intracellular retention of the radiolabel. Direct and indirect halogen radiolabelling results in different cell-processing patterns of radiolabels, and radioactive catabolic products follow different routes of cellular excretion. The results of this cellular study indicate that indirect labelling is preferable to the direct Chloramine-T method.

Introduction

Positron emission tomography (PET) is increasingly used for tumour visualization. Detection sensitivity is higher for PET in comparison with single photon emission computerized tomography. It was shown that the use of tumour-specific agents such as monoclonal antibodies (mAbs) directed against tumour-associated antigens and radiolabelled with positron emitters may result in more specific detection than with widely spread non-tumour-specific PET targeting agents such as [¹⁸F]FDG

and [¹¹C]methionine (1). Labelling of mAbs for PET radio-immuno diagnosis requires positron-emitting nuclides with half-lives longer than the conventional ¹¹C and ¹⁸F (*T*_{1/2}=20 and 110 min) that can readily be attached to proteins. The radiohalogens ⁷⁶Br and ¹²⁴I have been suggested as suitable candidates for this purpose (2-4) by virtue of its decay half-life and reasonable positron yield (*T*_{1/2}=16.2 h and 55% for ⁷⁶Br and *T*_{1/2}=4.2 days and 55% for ¹²⁴I) as well as the established and simple direct and indirect radiohalogenation of proteins (5-8). Indirect radiohalogenation of proteins seems preferable because it was found that catabolism of directly radiohalogenated mAbs resulted in high amounts of free radiohalogene, which had a slow excretion rate (bromine) (5) or accumulated in normal tissue (iodine) (9).

We are planning to introduce A33 mAb for radioimmuno diagnosis and therapy of colorectal cancers (10,11). The expression of the colon-cancer antigen, A33, appears to be organ specific, with the large and small intestinal mucosa as the principal sites of expression. An almost uniform tumour expression of the A33 antigen has been found in 95% of primary and metastatic colorectal cancers, while other epithelial cancers, sarcomas, neuroectodermal cancers and lymphoid neoplasms are generally A33-negative (12). The antigen is not secreted or shed and some human colon cancer cell lines have shown high levels of antigen expression, binding up to 370,000 A33 antibodies per cell (4). For this reason, this antibody might be a useful targeting vector in radioimmuno diagnosis. However, there are indications of non-typical cellular processing of the mAb A33 (13,14), possibly invalidating general considerations about tracer design and processing.

Introduction of the new radiolabelled protein in clinical practice requires detailed study (15) including labelling chemistry, cellular processing, radiocatabolites, and pharmacokinetic. The aim of the present study was to further characterize the cellular processing of mAb A33 labelled with radiohalogens ⁷⁶Br and ¹²⁵I by using the conventional direct Chloramine-T (CAT) method and the indirect procedure with *N*-succinimidyl *para*-(tri-methylstannyl) benzoate (SPMB) as the precursor molecule (5,7). Studies were carried out on the A33-antigen positive SW1222 cell line. The four conjugates were analyzed regarding cellular uptake and retention. Furthermore, excreted catabolites were analyzed to identify the distribution of high and low molecular weight components.

Correspondence to: Dr Vladimir Tolmachev, The Rudbeck Laboratory, Division of Biomedical Radiation Sciences, S-751 85 Uppsala, Sweden
E-mail: vladimir.tolmachev@bms.uu.se

Key words: ⁷⁶Br, indirect labeling, antibody internalization, cellular processing

Materials and methods

Antibodies and cells. The CDR-grafted huA33 antibody (9 mg/ml in phosphate buffer (50 mM, pH 7.0) with 154 mM NaCl) (16) and the colorectal cancer cell line, SW1222, expressing the A33 antigen (17) were kindly donated by the Ludwig Institute (New York, NY, USA). The cells were counted with an electronic cell counter (Beckman Coulter, Bromma, Sweden). Cells were plated in culture dishes (\varnothing 3.5 cm); at the time of the experiments, the number of cells per dish was $0.7\text{--}1 \times 10^6$. Cells were washed once with complete culture medium before the radioactive medium (1 ml per dish) was applied. The radioactive medium contained $0.2 \mu\text{g}$ mAb/ml (30–60 kBq ^{76}Br or 15–30 kBq ^{125}I). For each data-point three cell dishes were used, and experiments were repeated three times.

Chemicals. All chemicals and reagents were of analytical grade. CAT and sodium metabisulfite (Sigma Chemical Co., St Louis, MO, USA) were used as 10 mg/ml aqueous solutions. The precursor, SPMB, was synthesized according to Kozirowski *et al.* (18).

Radioactivity measurements, radionuclides and radiolabelling of mAb A33. The radioactivity was measured using either an ultra-pure germanium detector with multi-channel analyzer (ORTEC, The Nucleus, Inc., Oak Ridge, TN, USA) working on-line with a PC, or an automatic gamma counter (Wallac, Upplands-Väsby, Sweden) using a ^{76}Br measurement protocol with an energy window of 300–2047 keV and a ^{125}I standard protocol.

The production of ^{76}Br was carried out at a low-energy cyclotron (MC17, Scanditronix, Uppsala, Sweden) at the Uppsala University PET Centre according to a method described previously (19). [^{125}I]NaI was purchased from Amersham Pharmacia Biotech UK Limited (Buckinghamshire, UK).

Direct and indirect radiohalogenation was performed as described elsewhere (5–7,10). Directly (^{125}I -A33, ^{76}Br -A33) and indirectly labelled mAb ([^{125}I]PIB-A33 and [^{76}Br]PBrB-A33) were purified on Sephadex G-25 NAP-5 columns (Pharmacia, Uppsala, Sweden) equilibrated with PBS.

Analysis of cell culture medium. After cell incubation, the cell culture medium was analyzed regarding the amount of radioactivity released and size of degradation products. Half-ml samples of the cell culture medium were aspirated and separated into high and low molecular weight (HMW and LMW) fractions on NAP-5 columns. The radioactivity of the fractions was then determined using the gamma well-counter.

Continuous incubation with radioactive media. Radioactive medium was added to the dishes and the cells were incubated at 37°C from 30 min up to 48 h. At various time-points, cell-dishes were taken to analyze membrane-bound and internalized radioactivity as described elsewhere (20). Additional dishes were incubated 1–48 h with an excess of unlabeled mAb, $12 \mu\text{g}$ per dish, to determine unspecific binding.

Interrupted incubation with radioactive medium. Radioactive medium was added to the dishes and the cells were incubated

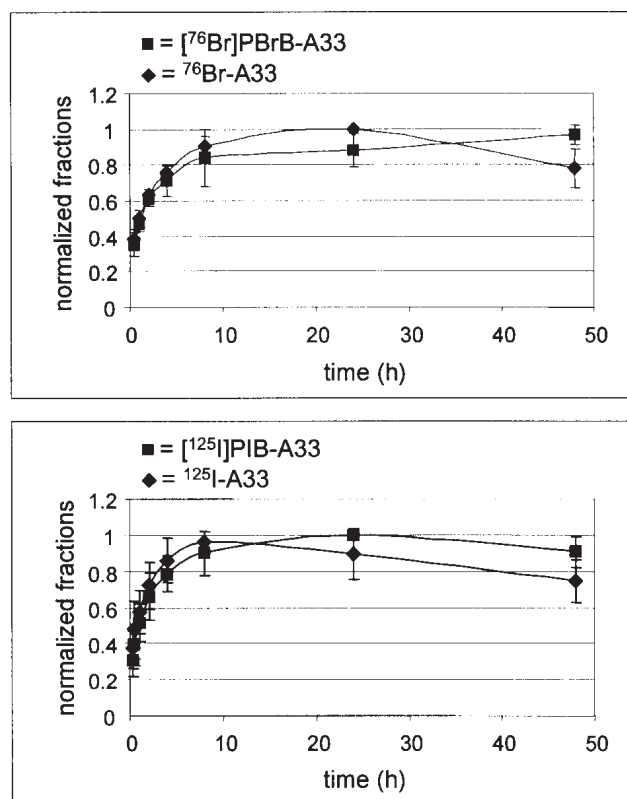


Figure 1. Uptake as a function of time for the four conjugates: (A) ^{76}Br -A33 (\blacklozenge) and [^{76}Br]PBrB-A33 (\blacksquare); (B) ^{125}I -A33 (\blacklozenge) and [^{125}I]PIB-A33 (\blacksquare). Means and maximum errors are shown for 3 or 4 experiments, each performed in triplicate.

at 37°C for 12–13 h. Incubation was interrupted by washing the dishes six times with ice-cold serum-free medium. Fresh, complete medium at 37°C was added, 1 ml per dish, and the cells were again incubated at 37°C . At various time-points, from 0–24 h, the incubation medium was aspirated from cell-dishes for size exclusion chromatography. The cells were then analyzed for membrane-bound and internalized radioactivity.

Cellularly processed mAb. In order to evaluate the processing of internalized radioactivity, cells after interrupted incubation with radiolabelled mAb were incubated with $12 \mu\text{g}$ non-labelled mAb at 4°C for 4 h in order to displace membrane-bound radioactive mAb. Cells were washed with cold media and, after addition of 1 ml fresh media, were incubated at 37°C . At various time-points, 0–24 h, incubation medium was aspirated from the cell-dishes and analyzed by size exclusion chromatography. The cells were then washed six times with ice-cold serum-free culture medium and analyzed for membrane-bound and internalized radioactivity.

Results

Continuous incubation with radioactive media. The cellular uptake of the four conjugates as a function of time was studied; the results are shown in Fig. 1, where 1.0 represents maximum uptake. The uptake pattern was rather similar for the four conjugates although the absolute uptake was higher for indirectly labelled mAb ([^{76}Br]PBrB-A33 and [^{125}I]PIB-A33) than for

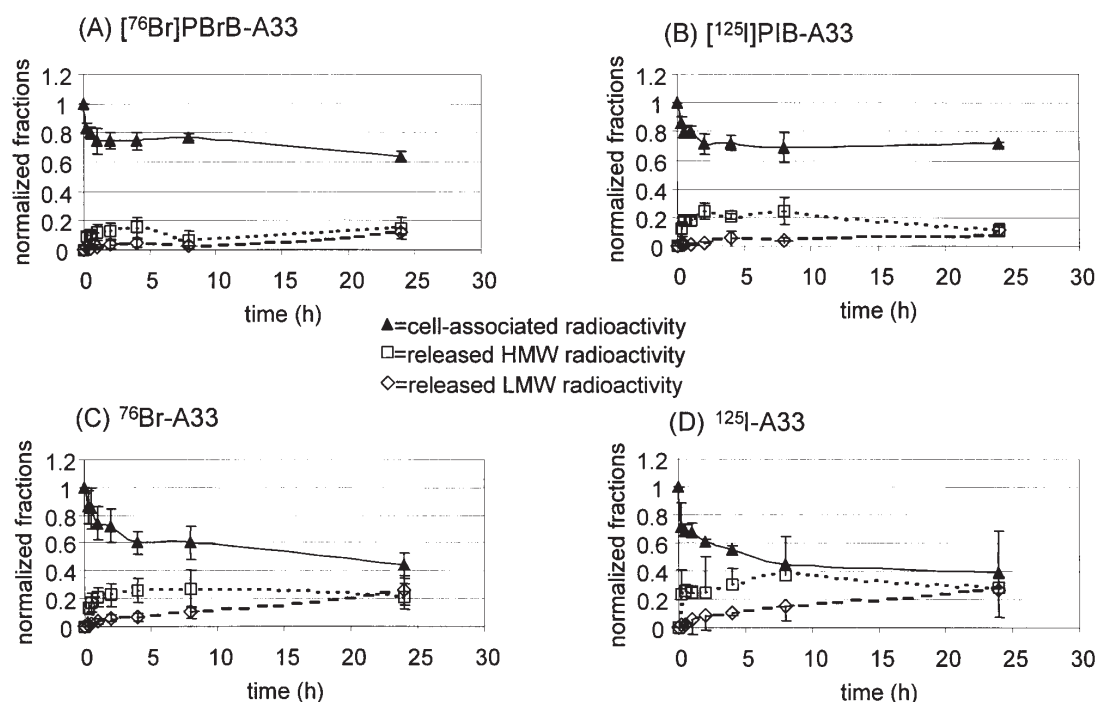


Figure 2. Release of radioactivity from cells as a function of time for the four conjugates [^{76}Br]PBrB-A33 (A), [^{125}I]PIB-A33 (B), ^{76}Br -A33 (C) and ^{125}I -A33 (D). \blacktriangle — \blacktriangle , total cell-associated radioactivity; \diamond — \diamond , low molecular weight; and \square — \square , high molecular weight released radioactive products. Means and maximum errors are shown for 3 experiments, each performed in triplicate.

directly labelled mAb (^{76}Br -A33 and ^{125}I -A33) (20-60% and 10-20% from added radioactivity, respectively). A tendency toward a decreased cell-associated radioactivity at the latest time-point (48 h) was seen for both brominated and iodinated conjugates in the case of directly labelled mAb. Furthermore, the membrane-bound radioactivity corresponded to approximately two-thirds and internalized approximately one-third of the total cell-associated radioactivity throughout the study (data not shown). The background invariably accounted for less than 10% of the total cell-associated radioactivity.

Interrupted incubation with radioactive media. The cellular retention of radioactivity after interrupted incubation with the four conjugates was studied; the results are shown in Fig. 2. Retention of indirectly labelled mAb (Fig. 2A and B) was more pronounced than that of directly labelled mAb (Fig. 2C and D), 64-72% vis-à-vis 39-44% cell-associated radioactivity after 24 h for indirectly and directly radiolabelled mAb respectively. Initially, a quick release of radioactivity was observed, 25-28% and 40-45% of initially bound, for indirectly and directly radiolabelled mAb, respectively. This early decrease in cell-associated radioactivity (completed within 4 h) was almost entirely due to a release of membrane-bound radioactivity (data not shown). The release of radioactivity from the cells showed a bi-exponential pattern, with a rapid early and a slower late phase. The $T_{1/2}$ (early) was less than 20 min for all four conjugates, whereas the $T_{1/2}$ (late) was several days. Furthermore, the $T_{1/2}$ (late) was longer for the indirectly labelled conjugates, 5-48 days, compared with the directly labelled mAbs, with a $T_{1/2}$ (late) of 1-2 days. The released radioactivity as measured in the cell-culture medium was, for all four conjugates, to the greatest extent in HMW form until the latest

time-point when the amounts of HMW and LMW radioactivity were alike. Furthermore, the release of LMW radioactivity was slower in the case of indirectly vs. directly labelled mAb.

Cellularly processed mAb. The excretion pattern of degradation products was assessed after interrupted incubation, with subsequent displacement of cell-surface bound mAb. Results of the cellular excretion of radioactivity and of the analysis of degradation products are shown in Fig. 3. In the case of direct labelling, the retained intracellular radioactivity was 29-31% after 24 h, whereas for indirect labelling it was 49-61%. The two indirectly labelled conjugates (Fig. 3A and B) revealed similar excretion patterns, with a somewhat higher fraction of HMW radioactivity at the start of incubation but with augmented release of LMW radioactivity over time. The two directly labelled conjugates (Fig. 3C and D) also showed similar excretion patterns, but LMW radioactivity was released to a significantly greater extent in the case of directly vis-à-vis indirectly labelled mAb. A part of the excreted radioactivity was in HMW form also in the case of directly labelled mAb.

Discussion

The studies presented were carried out on the A33-antigen positive SW1222 cell line. We investigated the cellular processing of the four radiolabelled conjugates, mAb A33 directly or indirectly radiolabelled with ^{76}Br and ^{125}I . The fate of the radiolabels was analyzed as regards cellular uptake and retention. The excreted catabolites were further analyzed to identify the distribution of high and low molecular weight components.

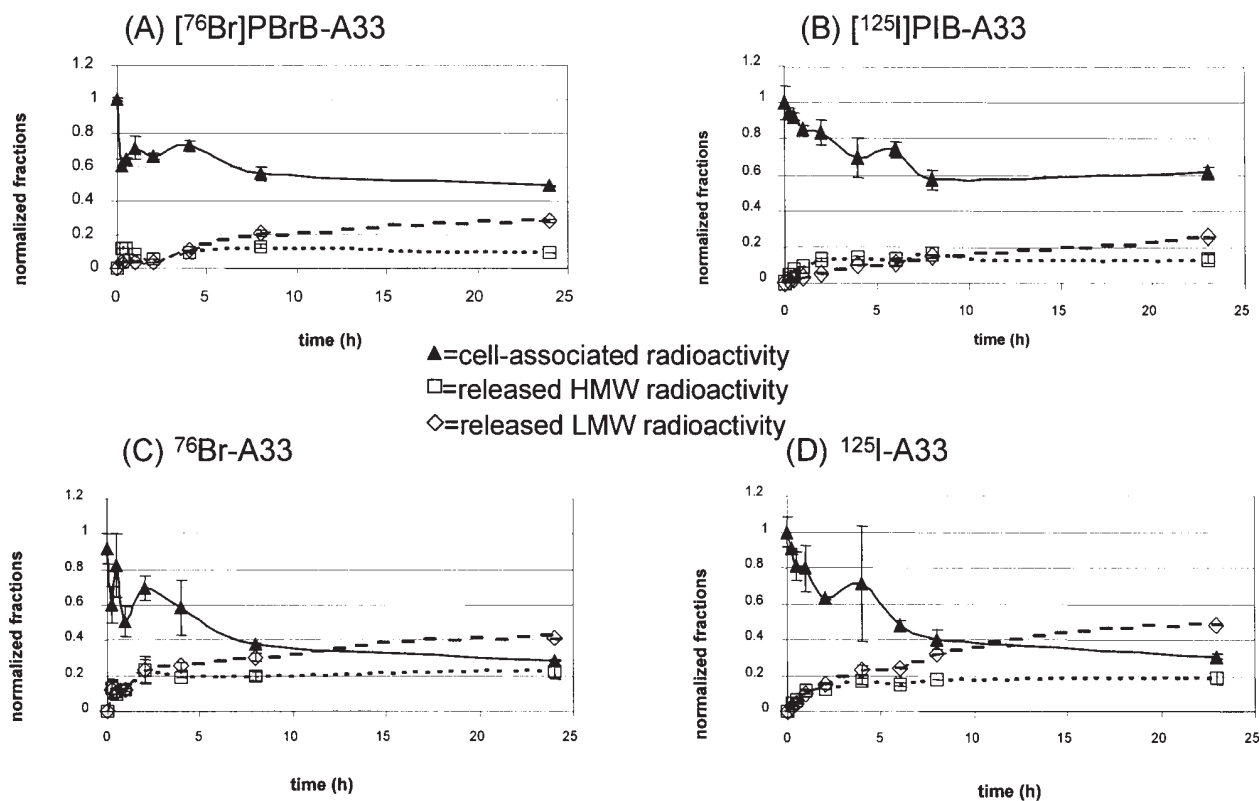


Figure 3. Excretion of radioactivity from cells as a function of time for the four conjugates [^{76}Br]PBrB-A33 (A), [^{125}I]PIB-A33 (B), ^{76}Br -A33 (C) and ^{125}I -A33 (D). \blacktriangle — \blacktriangle , total cell-associated radioactivity; \diamond — \diamond , low molecular weight; and \square — \square , high molecular weight released radioactive products. Means and maximum errors are shown for 1 experiment, performed in triplicate.

As indicated in Fig. 1 the cellular uptake pattern of the four conjugates was fairly similar. However, the absolute uptake of the radiolabelled mAb depended on the labelling method. A possible explanation for the greater absolute uptake of indirectly vs. directly labelled mAb in the present study is that indirect labelling might cause less damage to the protein. The indirect labelling used in the present study conjugates the radiohalogenated linker molecule to lysine residues of the protein, while direct labelling introduces the radiohalogen into the tyrosine moieties of the protein. Many antibodies, including A33, are known to have an increased fraction of tyrosine residues at the antigen-binding site while the lysine residues are more evenly distributed throughout the antibody (21). Several earlier studies (22–24) have actually shown that the affinity of radiolabelled mAb may depend on the experimental protocol and experimental conditions rather than the properties of the mAb itself.

The difference in retention between directly and indirectly halogenated mAb A33, as demonstrated in Fig. 2, could have several explanations. It might be caused partly by different time-patterns in the release of membrane-bound mAb. The bi-exponential shape of the dissociation curves with a rapid early and a slow late phase agrees closely with previously published data (22,23,25). The reason for the bi-phase dissociation curves is not known definitely, but one explanation, suggested by Mattes (23) may be that a fraction of the mAb is monovalently bound and represents the first rapid dissociation. Monoclonal Abs bind predominantly bivalently to the cell surface and wobble at a significant rate to generate a transient, monovalently bound form. Antibodies labelled by different methods

differ in their binding behaviour in that the fraction of antibody released intact can differ (26) possibly as a result of complete destruction or hampered function of one binding site by a particular labelling, as mentioned above. It has been shown (25) that Fab fragments dissociated much more rapidly than the intact mAbs, indicating the need of bivalent attachment for irreversible binding.

Another explanation for the differing retention of radioactivity after indirect vs. direct radiohalogenation may be differences in the cellular processing of radiolabelled degradation products. The cellular processing of internalized mAb, shown in Fig. 3, elucidates the differing intracellular retention of degradation products between the two labelling methods. The results show a pronounced retention of internalized radioactivity in the case of indirect labelling as compared with direct labelling. Furthermore, the results in Figs. 2 and 3 demonstrate a difference in release of HMW and LMW radioactivity for the two labelling methods. In Fig. 2, the results reflect the sum of dissociated membrane bound and intracellularly processed mAb whereas, in Fig. 3, the results reflect intracellularly processed mAb.

mAb A33 is thought to be internalized, upon binding to the A33 antigen, into a not yet fully-characterized large cytoplasmic vesicular compartment, a so-called micropinosome (13,14), in contrast with the majority of mAbs which, after internalisation, are transported to lysosomes where they are metabolised (27). The subcellular distribution of mAb A33 upon internalization into A33 antigen-positive cells has been examined (13), revealing that up to 40% of total bound mAb A33 accumulates in these cytoplasmic vesicles, which traverse

the cytoplasm and come into close proximity with the cell nucleus (13). In the present study, the fraction of internalized radioactivity (approximately one-third of total bound) tallies closely with previously published data.

The fact that HMW radioactivity was released from cells that were treated with excess unlabeled mAb at 4°C to displace membrane-bound mAb (Fig. 3) indicates an excretion of internalized intact mAb. It has been claimed (28) that part of the internalized mAb A33 can be exteriorized in an intact form and thus bind back to the cell surface so that the process of uptake and internalization can be repeated.

Indirect labelling resulted in greater absolute uptake and prolonged intracellular retention of the radiolabel, when compared with direct labelling. Furthermore, no difference in cellular uptake and retention was observed between the two halogens, bromine and iodine. The cellular processing of the radiohalogens thus depends on the labelling method used rather than on the halogen itself. The more favourable retention of the indirectly radiohalogenated mAb compared with directly labelled may be beneficial *in vivo* for some antibodies.

Acknowledgements

The authors would like to thank the staff at the Uppsala University PET Center for their assistance with the radiobromine production and the The Svedberg Laboratory for allowing us to use their facilities. This study was supported by the Swedish Cancer Society.

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