## Supplemental Information

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## Method: Biodistribution Study of SAAC $\left({ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6 4 hours postinjection

B16/F10 melanoma and 38C13 lymphoma tumor models were used in this study. When the tumors had grown to $6-8 \mathrm{~mm}$ in diameter, the mice were randomly divided into treated group (TG) and non-treated group (N-TG), with each group consisting of 4-5 mice. In the TG groups, mice bearing murine B16/F10 melanoma received a single dose of intravenous polymeric paclitaxel (equivalent dose $=80 \mathrm{mg} / \mathrm{kg}$ ); and mice bearing 38C13 xenografts received intraperitoneal injection of cyclophosphamide ( $100 \mathrm{mg} / \mathrm{kg}$ ). Twenty-four hours after treatment, all mice in the TG received intravenous injection of SAAC $\left({ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6 ( $20 \mu \mathrm{Ci} /$ mouse $)$. Mice in the N-TG also received intravenous injection of SAAC $\left({ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6 (20 $\mu \mathrm{Ci} /$ mouse $)$. Mice were killed 4 hours after radiotracer injection. Blood, heart, liver, spleen, kidney, lung, stomach, intestine, muscle, bone, and tumor tissues were removed and weighed. Radioactivity was measured with a gamma counter. Uptake of SAAC( $\left.{ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6 in various tissues was calculated as the percentage of the injected dose per gram of tissue (\% ID/g).


Supplemental Figure 1. Structure of SAAC ( $\left.{ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6.

A


B


SAAC ( ${ }^{99 \mathrm{~m} T \mathrm{Tc}) \text {-PSBP-6 uptake increase rate: }}$ $142.4 \pm 36.9 \%$

Supplemental Figure 2. Changes in tumor uptake values of radiotracers in 38C13 tumors after therapy. (A) Changes of ${ }^{18}$ F-FDG uptake, expressed as percentages of injection dose per gram tumor (\%ID/g). (B) Changes of SAAC ( ${ }^{99 \mathrm{~m}} \mathrm{Tc}$ )-PSBP-6 uptake, expressed as percentages of injection dose (\% ID). An1-An5, animals 1-5.


Supplemental Figure 3. Changes in tumor uptake values of radiotracers in B16/F10 tumors after therapy. (A) Changes of ${ }^{18} \mathrm{~F}$-FDG uptake, expressed as percentages of injection dose per gram tumor (\%ID/g). (B) Changes of SAAC $\left({ }^{99 m} \mathrm{Tc}\right)$-PSBP-6 uptake, expressed as percentages of injection dose (\% ID). An1-An5, animals 1-5.


Supplemental Figure 4. Representative autoradiographs and photographs of H\&E-, TUNEL-, and active-caspase 3-stained slides of B16/F10 tumors from a non-treated mouse (A) and a PG-TXL-treated mouse (B). Tumors were removed at 4 hours following i.v. injection of SAAC( ${ }^{99 m}$ Tc)-PSBP-6. Red in fluorescent microphotographs shows TUNEL-positive apoptotic cells; blue represents DAPI-stained cells.


Supplemental Figure 5. Biodistribution of SAAC( $\left.{ }^{99 m} \mathrm{Tc}\right)-\mathrm{PSBP}-6$ in $\mathrm{C} 3 \mathrm{H} / \mathrm{HeJ}$ mice bearing 38C13 lymphoma. Mice in treated group received i.p. injection of cyclophosphamide.


Supplemental Figure 6. Biodistribution of SAAC( $\left.{ }^{99 \mathrm{~m}} \mathrm{Tc}\right)$-PSBP-6 in nude mice bearing B16/F10 melanoma. Mice in treated group received i.v. injection of PG-paclitaxel.

