## Supplementary Information (SI)

Article title: Modelling DTPA therapy following Am contamination in rats

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Table A1 Kinetics of Am-citrate in blood of rats. Data from Turner and Taylor (1968) and from CEA/LRT studies. Data are expressed as the percentage of the administered activity (% IA; mean ± SD).

Exp.	n <sup>a</sup>		Time after administration of Am-citrate (d)									
		6.94E-04	0.01	0.021	0.042	0.21	1	7	17			
Turner and Taylor (1968)		14.9 ± 0.5		5.0 ± 0.5	2.2 ± 0.3							
CEA/LRT	4		10.3 ± 0.5			0.35 ± 0.05	$0.13 \pm 0.03$	0.03 ± 0.03	0.013 ± 0.05			

<sup>a</sup> n = number of rats per group; this information is not available for the study of Turner and Taylor.

Table A2 Urinary excretion of Am-citrate in control rats. Data are expressed as 24 hours cumulated percentage of injected activity (% IA; mean ± SD).

nª		Time after administration of Am-citrate (d)									
	1	2	3	4	5	6	7	8	9	10	
4	10 ± 2	1.2 ± 0.3	0.55 ± 0.07	0.37 ± 0.04	0.27 ± 0.05	0.23 ± 0.05	0.25 ± 0.22	$0.21 \pm 0.10$	$0.16 \pm 0.08$	$0.11 \pm 0.04$	

<sup>a</sup> n = number of rats per group

Table A3 Faecal excretion of Am-citrate in control rats. Data are expressed as 24 hours cumulated percentage of injected activity (% IA; mean ± SD).

nª		Time after administration of Am-citrate (d)									
	1	2	3	4	5	6	7	8	9	10	
4	1.3 ± 0.2	1.8± 0.8	2.3 ± 0.9	2.9 ± 0.9	2.9 ± 0.8	3.2 ± 0.3	2.7 ± 0.9	2.6 ± 0.5	2.5 ± 0.7	$1.8 \pm 0.4$	

<sup>a</sup> n = number of rats per group

Table A4: Pu-DTPA in blood after injection of 12.2 kBq Pu-DTPA. Data are expressed as the percentage of the administered activity (% IA). The uncertainty was calculated as 30 % of the measured value.

Exp.	nª	Time after administration of Pu-DTPA (min.)									
		10	20	40	60	120	240	1440			
CEA/LRT	1	7.14±2.14	4.03± 1.21	2.77± 0.83	0.91± 0.27	$0.14 \pm 0.04$	$0.03 \pm 0.01$	0.02 ± 0.005			

<sup>a</sup> n = number of rats per time point

Table A5: Pu-DTPA in urine after injection of 12.2 kBq Pu-DTPA. Data are expressed as the percentage of the administered activity (% IA). The uncertainties correspond to the measurement error, or to 3 % of the measured value (whichever the highest).

Ехр	nª	Time after administration of Pu-DTPA (min.)								
		10	20	40	60	120	240	1440		
Pu-DTPA from CEA/LRT	1	19.6 ± 0.6	20.7 ± 0.6	68 ± 4	88 ± 6	92 ± 8	91 ± 7	93 ± 16		

<sup>a</sup> n = number of rats per time point

Table A6: Measurements of hepatic and skeletal Am in rats on Day 14 for different DTPA treatment times. Data are expressed as the mean percentage of the injected activity (% IA; mean ± SD).

Exp.	nª	dose	Т	Liver	Skeleton
		(µmol /kg)			
А	4	300	1d	1.05±0.31	28±2
В	4	300	3d	1.16±0.19	30±4
С	4	300	7d	1.25±0.25	35±3

<sup>a</sup> n = number of rats per group

d = dose of DTPA treatment

T = time of DTPA administration

Table A7: Urinary excretion of Am in DTPA-treated rats. Data are expressed as 24 hours cumulated percentage of injected activity (% IA; mean ± SD).

Exp.	nª	dose	Т	Time after administration of Am-citrate (d)											
		(µmol /kg)	(d)	2	3	4	5	6	7	8	9	10			
А	4	300	1	12.0 ± 1.2	3.7±0.4	2.2±0.2	1.41±0.05	1.05±0.09	0.78±0.14	0.76±0.08					
В	4	300	3			10.2 ± 0.1	3.1 ± 0.3	1.9 ± 0.9							
С	4	300	7							8.2 ± 1.2	2.4 ± 0.3	$1.6 \pm 0.2$			

<sup>a</sup> n = number of rats per group

d = dose of DTPA treatment

T = time of DTPA administration

Table A8: Faecal excretion of Am in DTPA-treated rats. Data are expressed as 24 hours cumulated percentage of injected activity (% IA; mean ± SD).

Exp.	nª	dose	Т		Time after administration of Am-citrate (d)									
		(µmol /kg)	(d)	2	3	4	5	6	7	8	9	10		
А	4	300	1	9.4±1.1	9.7±3.0	3.9±1.6	1.8±0.2	1.11±0.13	0.70±0.12	0.51±0.10	0.34±0.09	0.21±0.01		
В	4	300	3			3.8±1.3	11±3	4.7±1.1						
С	4	300	7							5.0±1.9	8.4±0.8	3.3±0.8		

<sup>a</sup> n = number of rats per group

d = dose of DTPA treatment

T = time of DTPA administration



**Figure S1** comparison of model prediction (lines) and data (circles) of plasma uptake and clearance of Am in control rats; Data from Turner and Taylor (1968) plus additional unpublished data from studies conducted at CEA/LRT are expressed as the percentage of the injected activity (% IA; mean ± SD).