

INTERNAL DOSIMETRY AT RESEARCH AND MEDICAL FACILITIES

The University Five
Case Studies

Learning Objectives

- Identify radionuclides of concern at research and medical facilities
- Analyze two case studies

The University Five

- C-14: radiolabeling, radiocarbon dating
- P-32: radiolabeling of biomolecules
 - (also H-3, e.g., tritiated thymidine)
- I-125: diagnostic nuclear medicine
- I-131: therapeutic nuclear medicine
- Cf-252: neutron sources
 - Not an internal hazard

H-3, C-14 and P-32

- Low-,Medium- and High-energy beta emitters
- C-14 and P-32 detectable with pancake GM
- Normally quantified by smears and BLS
- Organically bound tritium (OBT): 50 % has a T(b) of 10 d and (50%) has a T(b) of 40 d (same as carbon)
- ALIs: C-14 = 2 mCi, P-32 = 0.9 mCi

Iodines

- All ALI's (except I-120m, I-128, I-134) are based on non-stochastic limit of 500 mSv CODE to thyroid ($w_T = 0.03$)
- Volatile, so easily dispersed; treatment is usually blocking with stable KI (325 mg); unblocked thyroid deposition = 25%; KI administered within 1-2 minutes lowers to 2%, after 6 hr lowers to 10%, no effect more than 12 hr after intake; however, daily dose prevents re-uptake of circulating iodide
- MDA in thyroid for photon-emitting isotopes is less than 0.4 Bq

Medical Uses of Radioiodines

- Thyroid uptake studies:
 - 15 MBq I-123 NaI (3.4 mGy/MBq)
 - or 0.4 MBq I-131 NaI (340 mGy/MBq)
- Thyroid scanning:
 - 4 MBq I-131 NaI p.o. for routine
 - 75 MBq I-131 NaI for post-operative carcinoma scanning
 - Tc-99m O_4 actually more common
- Thyroid ablation:
 - 1.1 GBq I-131 NaI for hyperthyroidism;
 - 7.4 GBq I-131 NaI for carcinoma
 - pregnancy testing always indicated

Estimates of Intakes and Internal Doses from Ingestion of P-32 at MIT and NIH

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Introduction

- June 29, 1995 - individual at NIH reportedly internally contaminated with P-32
- August 14, 1995 - individual at MIT also reported to have intake of P-32
- Internal Dose Center at Oak Ridge contacted for technical support
 - General internal dose information support center
 - Subcontractor to the USNRC - role in assessment of accidental intakes of radionuclides
 - Association with REAC/TS

Introduction (cont'd)

MIT Intake Incident

- Researcher working with P-32 - labeling of DNA components
- Contamination discovered early - external contamination
- Whole body counting
 - Shielded chair type NaI detector system
 - Started within ~5 days post intake
 - Continued to ~50 days post intake

Introduction (cont'd)

MIT Intake Incident

- Urine samples (mostly 24-hr)
 - Liquid scintillation counting
 - Started within ~10 days post intake
 - Continued to ~60 days post intake
- Single set of whole body counting data (N=35)
- Three separate sets of urine data analyzed
 - Supplied by MIT (N = 40)
 - Provided by the researcher (N = 48)
 - Independently analyzed by laboratories at ORISE (N = 51) (2 duplicate samples)

Methods

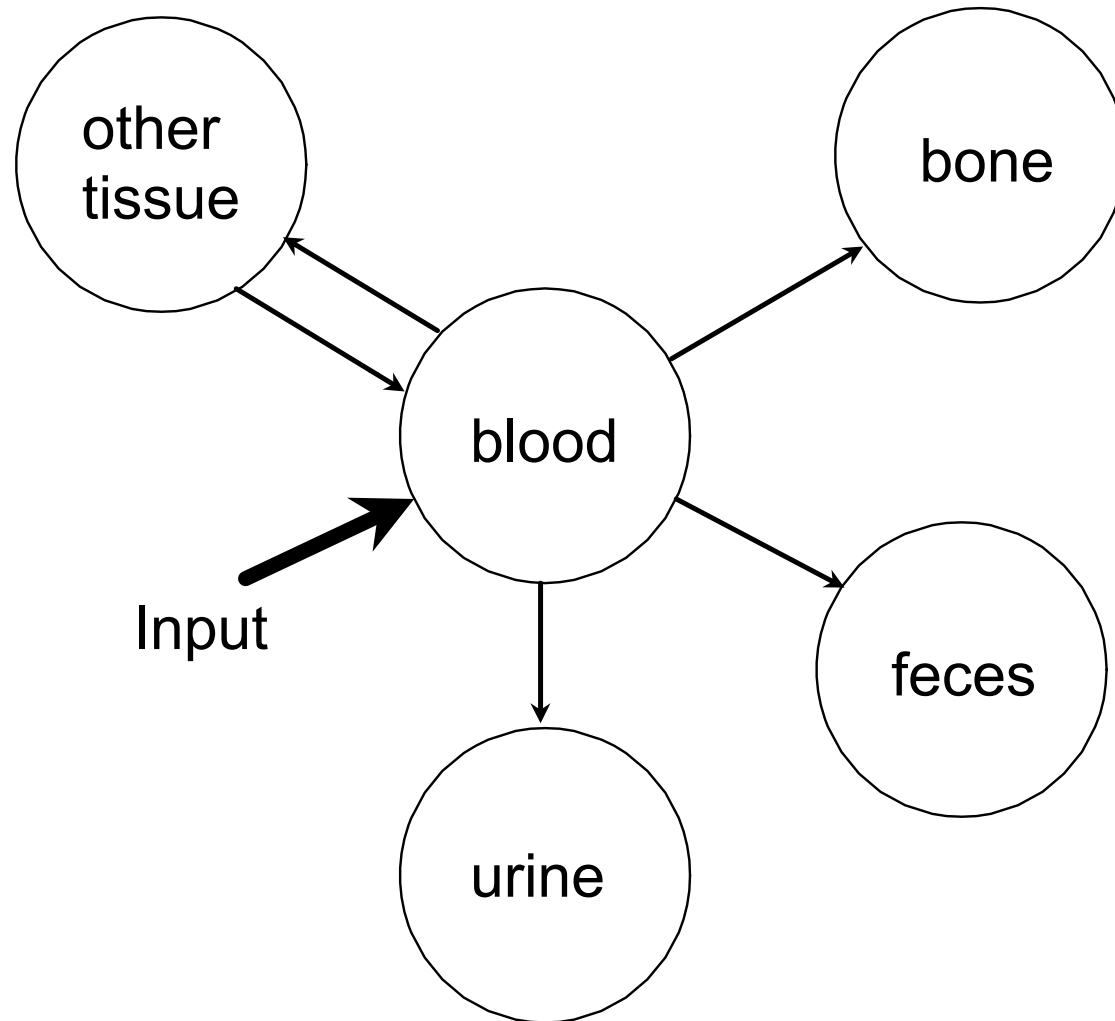
MIT Intake Incident

- Dose Assessment
 - Based on data from site, intake assumed to be ingestion
 - Only one f_1 category - licensee suggested that this was reasonable
 - Intake estimates using both weighted and unweighted least squares techniques for single intake/multiple bioassay data points
 - NUREG/CR-4884 used as source of IRFs for whole body retention, 24-hr urinary excretion

Methods (cont'd) -MIT Intake Incident

- Dose conversion factors from ICRP 30 applied to intake estimates
- Because of slight divergence of model from data at late times, an individual-specific model was also developed
 - Compartmental model developed
 - Whole body and urine data fitted - all model parameters allowed to vary
 - Intake estimated
 - Time integrals of activity in compartments used with ICRP 30 SEEs to obtain dose estimates

Individual-Specific Kinetic Model



Methods (cont'd) -MIT Intake Incident

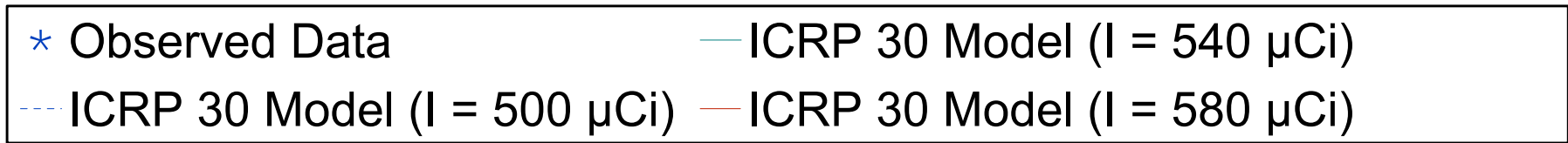
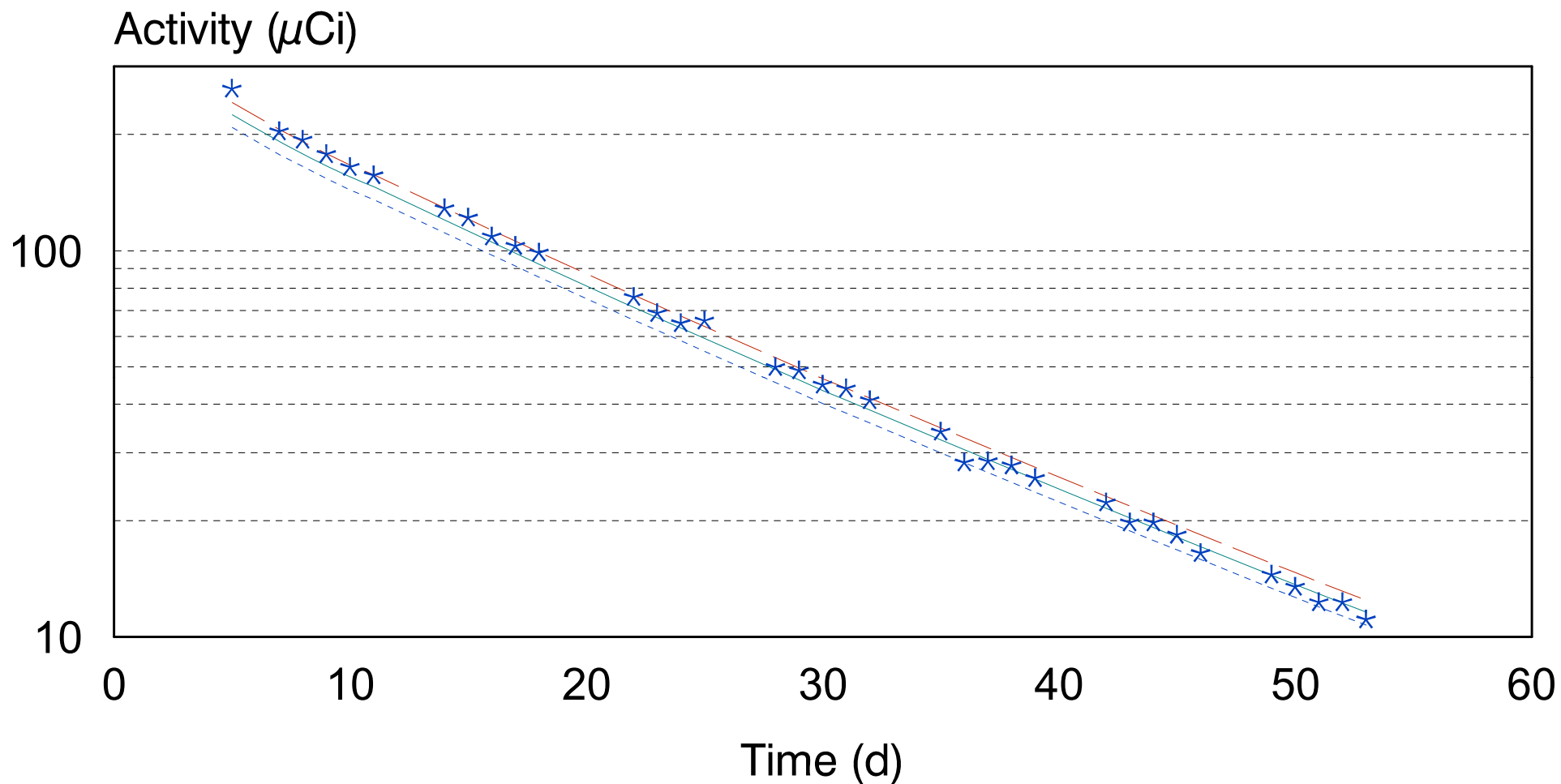
- Individual-specific modeling (cont'd)
 - Doses from intake with standard ICRP 30 model compared with those from the individual-specific model
 - Doses also compared with those predicted by the MIRDOSE 3.1 software program - less conservative marrow dose model
 - Organ doses, Effective Dose Equivalent, Effective Dose calculated

Results - Intake Estimates (μCi)

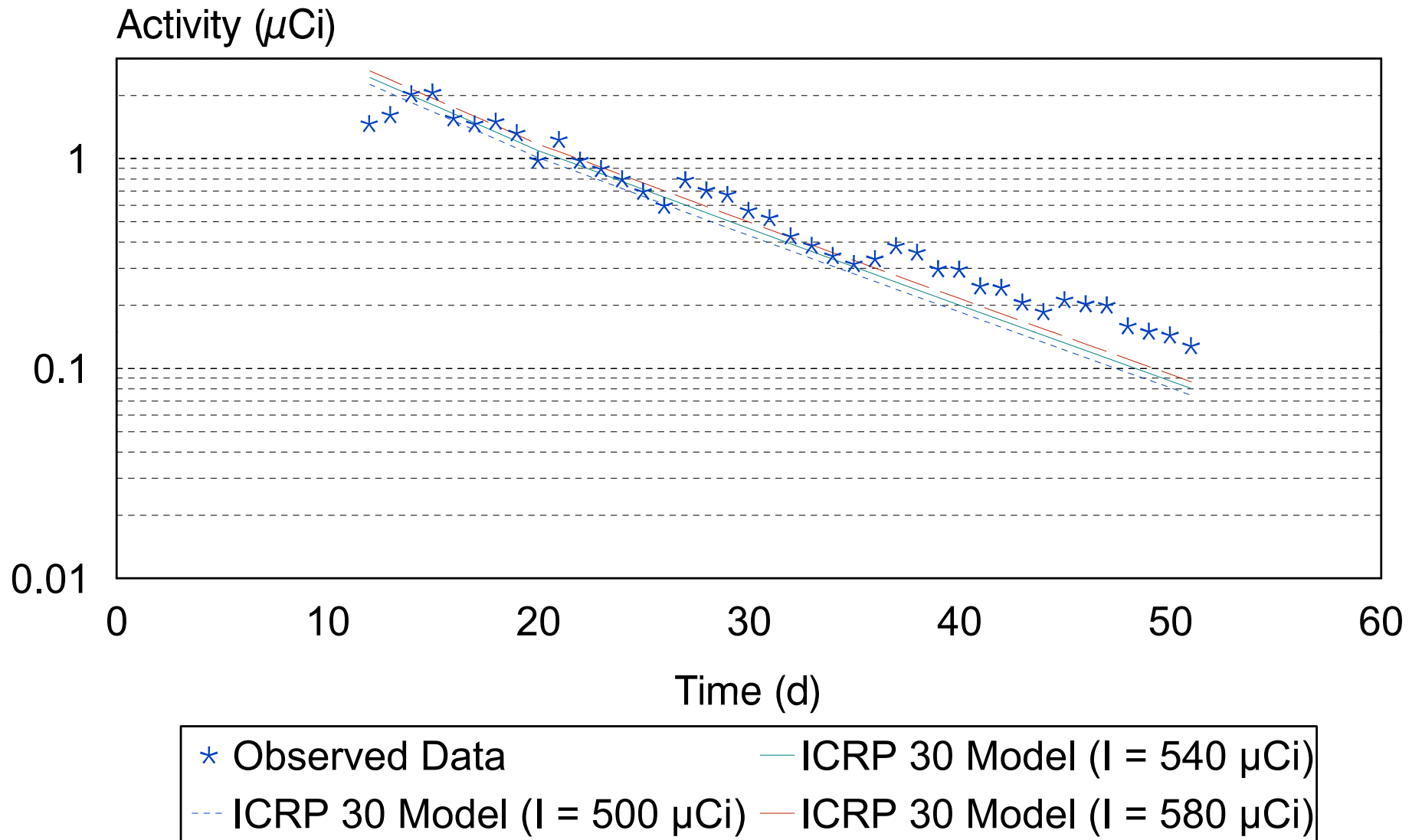
MIT Intake Incident

	Unweighted	Weighted
Whole Body Counting	586	576
Urine - MIT	505	561
Urine - Researcher	522	538
Urine - ORISE	484	537

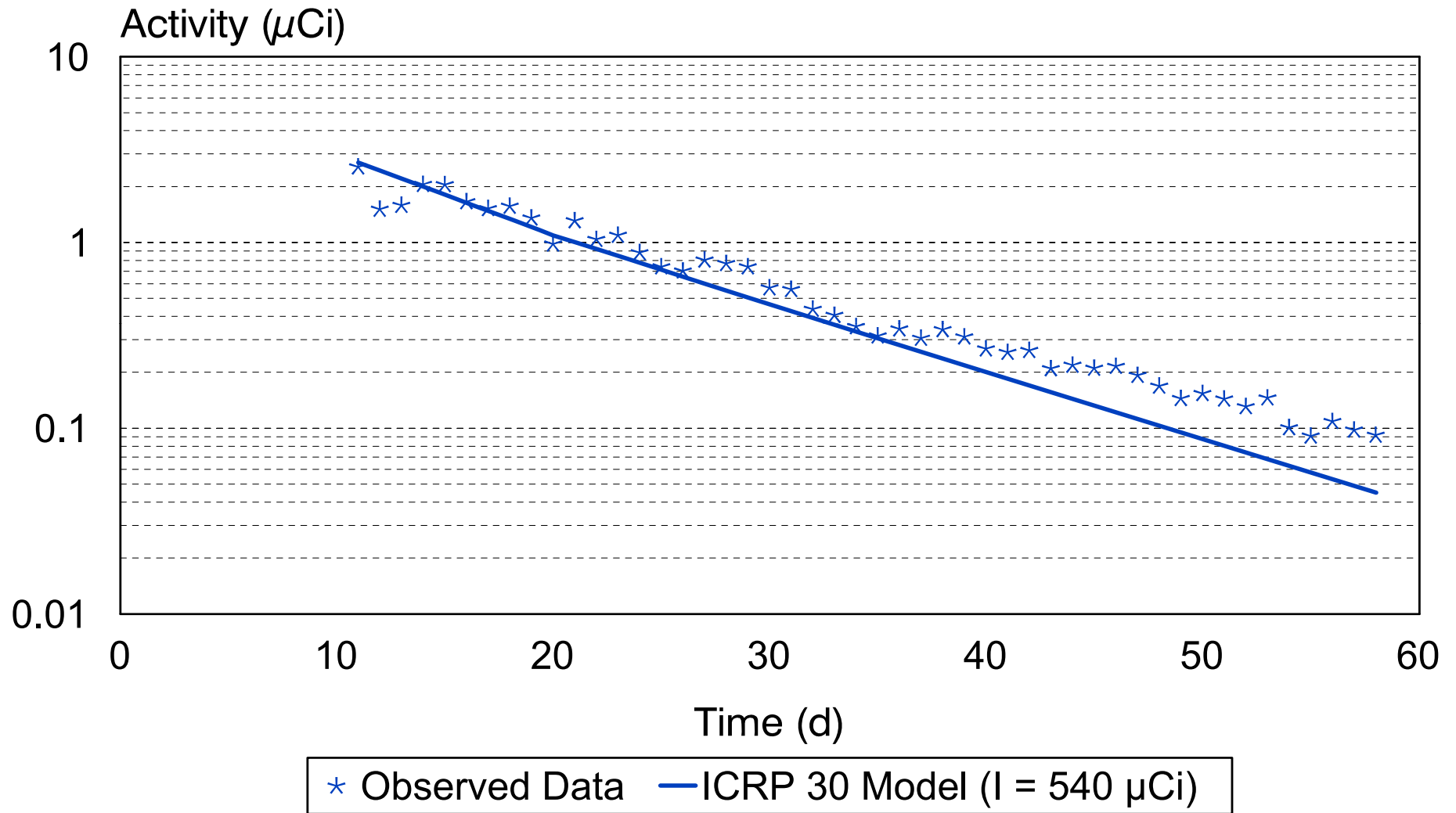
MIT P-32 Intake - WB Counting Data



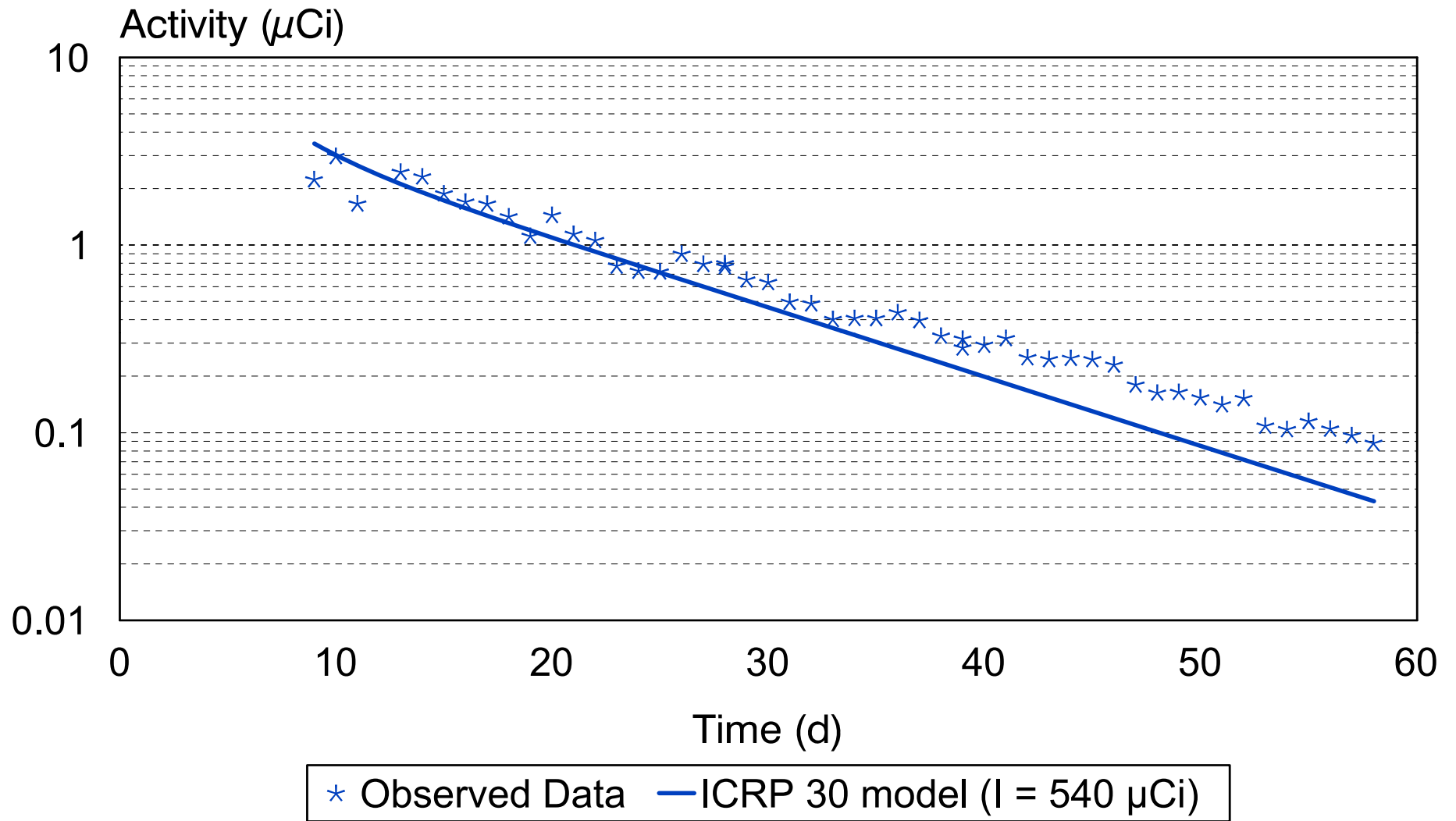
MIT P-32 Intake - Urine Data (MIT)



MIT P-32 Intake - Urine Data (Contaminated Individual)



MIT P-32 Intake - Urine Data (ORAU Analysis)



Results - Individual-Specific Kinetic Model

MIT Intake Incident

	T = 2.2 days	T = 25.5 days	T = Infinity
Blood	0.882	0.118	0
Other Tissue	-0.495	0.495	0
Bone	-0.077	-0.122	0.199
Total	0.31	0.491	0.199

Results - Committed Doses
MIT Incident - Effect of Change in Kinetic Model

	CDE (Sv/Bq)	CDE (Sv)	CDE (Sv/Bq)	CDE (Sv)
Gonads	6.5E-10	0.013975	9.6E-10	0.02064
Breast	6.5E-10	0.013975	9.6E-10	0.02064
Red Marrow	8.1E-9	0.17415	3.9E-9	0.08385
Bone Surfaces	7.9E-9	0.16985	3.8E-9	0.0817
ULI Wall	3E-9	0.0645	4.5E-9	0.09675
LLI Wall	7.2E-9	0.1548	1.1E-8	0.2365
EDE	2.1E-9	0.04515	1.9E-9	0.04085
ED	2.1E-9	0.04515	2E-9	0.043

Results - Committed Doses

MIT Intake Incident - Effect of Change in DCFs

	CDE (Sv/Bq)	CDE (Sv)	CDE (Sv/Bq)	CDE (Sv)
Gonads	9.6E-10	0.02064	9.9E-10	0.021285
Breast	9.6E-10	0.02064	9.9E-10	0.021285
Red Marrow	3.9E-9	0.08385	3E-9	0.0645
Bone Surfaces	3.8E-9	0.0817	3.5E-9	0.07525
ULI Wall	4.5E-9	0.09675	3.8E-9	0.0817
LLI Wall	1.1E-8	0.2365	1E-8	0.215
EDE	1.9E-9	0.04085	1.7E-9	0.03655
ED	2E-9	0.043	1.9E-9	0.04085

Discussion

MIT Intake Incident

- A large amount of data was available for analysis.
- All of the urine data were in good agreement.
- In general, the agreement of the data with the ICRP model was good.
- The range of estimates of intake from all sources (whole body counting data and urine data) was very small - good confidence in results.
- At late times, the ICRP model appeared to slightly overpredict whole body retention and underpredict excretion

Discussion (cont'd)

MIT Intake Incident

- With the individual-specific model, the long term biological half-time appeared to be around 26 d.
- The bone uptake was only about 20%, instead of 30%.
- The predictions of the internal workings of the compartment model are not very reliable, as no data on bone or soft tissue uptake per se was available, only whole body retention and excretion.
- In any case, the most likely intake appears to be around 15 kBq (540 μ Ci) and the dose about 40-45 mSv (4-4.5 rem).

Introduction

NIH P-32 Contamination Incident

- Pregnant researcher (~17 weeks gestation) discovered to be contaminated - internal contamination suspected.
- Conditions of intake difficult to determine.
- Spot urine sample taken immediately, follow-up with many urine samples (mostly 24-hr).

Introduction

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Introduction

NIH P-32 Contamination Incident

- Liquid scintillation counting
 - Started on the day that the contamination was discovered
 - Continued for 29 days
- Two whole body images taken with a nuclear medicine camera and quantitated.

Introduction

NIH P-32 Contamination Incident

- 26 other workers contaminated, but at much lower level.
 - 1-3 urine samples taken, liquid scintillation counting
 - Intakes assumed to come from contaminated water cooler.

Introduction

NIH P-32 Contamination Incident

- Single researcher - intake assumed to be ingestion
 - Intake from water cooler?
 - Contaminated food?
 - Intentional/accidental?

Methods

NIH P-32 Contamination Incident

- Intake form also assumed to be phosphate, due to lack of knowledge of true form. First few samples' kinetics not inconsistent with this model.
- Intake estimates using weighted, unweighted least squares, using NUREG/CR-4884 IRFs for urinary excretion.

Methods

NIH P-32 Contamination Incident

- Most samples assumed to be 24-hr values.
 - Highly variable volumes. Some suggestion that fluids were being forced.
 - Times of collection not always 24 hour periods. Some adjustments made to make samples as close as possible to 24-hr samples.
- ICRP 30 DCFs.

Results

NIH P-32 Contamination Incident

- In vivo images - 32 MBq (862 μ Ci) on 6/30, 13 MBq (342 μ Ci) on 7/7.
- Early estimate of intake, from first 4 data points, was 9.6 MBq (260 μ Ci), but final estimate, with 10 data points, was 27-30 MBq (740-820 μ Ci).
- Using a 70 kg adult, this implies a CEDE of 58-64 mSv (5.8-6.4 rem).
- Using a 57 kg adult, this implies a CEDE of 72-80 mSv (7.2-8.0 rem).

Results

NIH P-32 Contamination Incident

- Biological half-time from data apparently about 18.3 days. Good agreement with ICRP 30 phosphate model.
- Based on model in NUREG/CR-5631, Rev. 2, dose to fetus at 120 days' gestation about 2.1 mSv/MBq (7.8 rem/mCi). Intake of 27 MBq implies a fetal dose of 57 mSv (5.7 rem).