

Transfer Standard Alanine Study

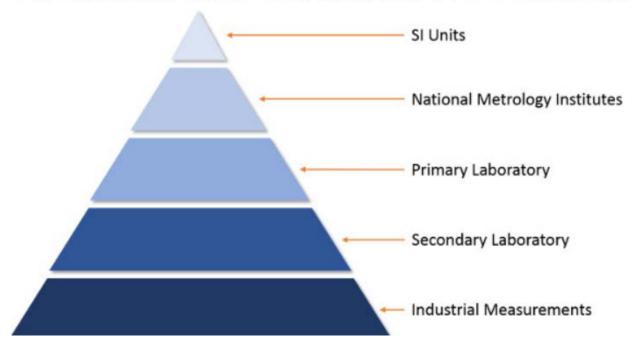
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Background

Ionizing radiation is used to produce various desired effects in products. Examples of applications include; the sterilization of medical products, microbial reduction, modification of polymers and electronic devices, and curing of inks, coatings, and adhesives. Radiation dose measurements are necessary to ensure the effectiveness of the radiation process. Dosimetry systems used for these dose measurements must be calibrated traceable to a national standard with a stated level of dose measurement uncertainty.

Dose traceability requires evidence of an unbroken chain of calibrations from the realization of the Gray (Gy) as determined from a primary national standard of dose. Certified traceable doses can only be provided by approved high dose reference laboratories (HDRLs). Approved labs are either a primary national standard HDRL or a traceable ISO 17025 accredited secondary standard HDRL lab.

MEASUREMENT TRACEABILITY PYRAMID



The ISO/ASTM 51261 Standard Practice for Calibration of Routine Dosimetry Systems for Radiation Processing specifies the requirements for calibrating routine dosimetry systems for use in radiation processing, including establishing measurement traceability. There are two methods for irradiating dosimeters that may be used for calibration of a dosimetry system.

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- Perform a full dosimetry system dosimeter batch specific calibration by irradiating representative dosimeter batch samples alongside reference transfer standard dosimeters, provided by an approved laboratory.
- 2. Irradiating representative dosimeter batch samples to known doses under fixed dose rate and temperature conditions, at an approved laboratory, followed by a calibration verification exercise with targeted dose irradiations performed in-situ or in the user's plant; the user's representative batch sample routine dosimeters are placed alongside reference transfer standard dosimeters, provided by an approved laboratory.

In either case, the use of reference transfer standard dosimeters is required.

Reference Transfer Standard Alanine Dosimetry

The successor to liquid chemical and radiochromic film reference transfer standard dosimetry is alanine dosimetry, which has demonstrated exceptional post-irradiation response stability that is key to postal-based transfer dosimetry services offered by calibration laboratories. The dosimetry system is based on irradiated crystalline alanine measured by electron paramagnetic resonance (EPR) spectrometry. Alanine is highly respected for its wide calibration dose range and environmental and time stable properties, and has become the transfer dosimetry system of choice for high-dose dosimetry services for National Metrology Institutes (NMI).

A large portion of the transfer standard alanine dosimeters being provided today have calibration traceability to either the NIST or NPL primary standards. NIST and NPL use somewhat different approaches in realizing the Gy to establish their primary standard of dose. These labs also use one or more types of internal transfer dosimetry systems, including alanine EPR dosimetry systems that use different sources of alanine pellets for their transfer standard alanine dosimetry. In addition, the two labs apply a different temperature factor (0.15%/°C vs 0.11%/°C) to adjust for transfer alanine irradiation temperatures that differ for their calibration temperature condition.

However, with all these differences, one should expect doses reported by these laboratories to be approximately "equivalent", meaning their results should be expected to be normally distributed within the overall combined uncertainty limits of the laboratories. These laboratories and other NMIs along with other ISO 17025 accredited laboratories participate in periodic formal inter-lab comparisons conducted by the BIPM, and typically evidence agreement within 1.0% under controlled temperature and dose rate irradiation conditions at 1.0, 10.0, and 30.0 kGy doses.

Inter-comparison Study

GEX Corporation (Centennial, Colorado USA) conducted an independent blind study of transfer national standard traceable alanine dosimetry systems during August 2016. The primary purpose of the study was to evaluate the level of agreement (equivalency) of the NIST and NPL calibration dose chains. The study involved the co-location of NIST and NPL transfer alanine dosimeters, along with GEX (NIST traceable) and DTU Risø (NPL traceable) transfer alanine dosimeters that were irradiated to a range of targeted doses, in order to assess the equivalency of dose results of the two popular, primary standard traceability chains under "typical" industry usage conditions.

This four lab inter-comparison study included in-situ site irradiations carried out under both a "fixed" radiation Gammacell 220 field where dose rate and temperature were held constant, and the "variable" radiation field of a largescale industrial electron beam irradiator where dose rate and temperature were uncontrolled.

Transfer standard alanine dosimeters were obtained from the two primary standard NIST and NPL HDRLs, along with two ISO 17025 certified secondary HDRLs, GEX and DTU / Risø. The alanine pellet transfer dosimeter sets were prepared in individually labeled target dose point packages by GEX along with irreversible temperature indicators, and followed detailed irradiation, shipping, and handling instructions, then sent to the two irradiation sites.

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The four sets of transfer alanine dosimeters for each dose point were irradiated by the individual sites to the targeted doses, and returned to GEX along with process records and irreversible temperature labels. GEX returned the transfer alanine packets to their respective labs with a specified average dose point temperature to be used to make appropriate temperature adjustments. The HDRLs were only instructed to prepare and report their doses in accordance with their established transfer alanine practices, and make appropriate temperature adjustments based on using the "average" irradiation temperatures as stipulated by GEX.

Study Results

The table below provides the dose results as reported by the four laboratories.

		Result	s of the Fo	ur Lab Inte	rcomparis	on Testin	g	
Cal	ID#	Irradiated	Target kGy	°C Temp _{corr}	DTU Risø	NIST	NPL	GEX
34	51	8/17/2016	7.5	32.5	7.55	7.45	7.38	7.39
34	51	8/18/2016	25.0	32.5	25.1	24.8	24.6	25.0
34	51	8/18/2016	45.0	32.5	45.3	45.1	44.6	45.2
34	52	8/16/2016	0.5	27.5	0.81	0.807	0.83	0.84
34	52	8/16/2016	2.5	27.5	2.30	2.30	2.27	2.29
34	52	8/16/2016	6.0	30.0	5.53	5.56	5.44	5.48
34	52	8/16/2016	12.0	32.5	11.1	11.0	10.9	11.1
34	52	8/16/2016	20.0	36.0	18.6	18.7	18.3	19.3
34	52	8/16/2016	45.0	47.5	47.4	47.7	46.6	47.9
34	52	8/16/2016	70.0	57.5	70.0	72.5	69.4	70.8

The table below shows the associated uncertainties used in equivalency analysis calculations.

Associated Dose Measurement Uncertainty (k=1)						
Lab "fixed" field "variable" fie						
NIST	0.900	0.090				
NPL	1.200	1.200				
GEX	1.065	1.350				
DTU Risø	1.750	1.750				

NOTE 1 - The temperature rise observed in the irreversible temperature indicators for the "variable" electron beam Cal 3452 dose points was not inconsistent with a normally expected 0.65 °C temperature rise in alanine and the 0.80 °C temperature rise in polystyrene per kGy of absorbed dose. The four HDRLs did apply different alanine response temperature adjustment factors ranging from a low of 0.11%/°C for NIST to a high of 0.15%/°C for NPL with GEX and DRU Risø each using a 0.14%/°C adjustment factor.

NOTE 2 - GEX assigned a higher average level of uncertainty associated with the doses for "variable" radiation conditions based on experience in providing in-situ irradiation transfer doses to industrial users. NIST, NPL and Risø reported the same levels of associated uncertainty for both "fixed" and "variable" irradiation doses with NPL, noting that 'no consideration is given to response differences arising from differences in-user site irradiation conditions or the additional uncertainty differences in the event actual temperature may differ from the user stated temperatures'. NIST also noted that their stated uncertainty 'does not attempt to account for uncertainties arising from customer reported temperatures or radiation field variability', and provided a memo with usage instructions and conditional usage information along with their alanine shipment. DTU Risø provided both an average stated uncertainty value along with individual dose point uncertainty estimates as did GEX.

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Analysis of Results

Analysis of results for the comparison and the degree of equivalence was determined by calculation of En-values as follows:

$$E_n = X_{lab} - X_{ref} / \sqrt{U_{lab^2} + U_{ref^2}}$$

Where:

Xlab = the result of the specific HDRL transfer alanine dose

Xref = the value of the reference (NIST, NPL GEX or Risø)

Ulab = the measurement uncertainty of the specific HDRL transfer alanine dosimetry system

Uref = the measurement uncertainty of the other lab or reference (NIST, NPL, GEX or DTU Risø transfer alanine dosimetry system)

The uncertainties were based on the NIST, NPL, DTU Risø, and GEX stated calibration uncertainty as calculated by GEX at k = 1, 2 and 3. E_n results of less than 1.0 are considered to evidence equivalency. Results are expected to be normally distributed and follow the so called "empirical rule" with 68% of the results falling within 1-sigma, 95% within 2-sigma, and 99.7% within 3-sigma. The distribution of E_n results may be plotted and evaluated for potential bias.

Results from the NIST and NPL primary standards labs for the "fixed" Gammacell 200 radiation field target irradiations are shown in the table below. These results evidence close agreement with all dose points shown equivalent and in agreement within ±1-sigma of the overall combined uncertainty limits.

	Fixed Dose/						
	Cal ID#3451 NIST/NPL Lab Intercomparison Results						
Dose							
Point ID	NIST kGy	NPL kGy	Difference kGy	% Difference			
1	7.45	7.38	-0.07	-0.94%			
2	24.8	24.6	-0.20	-0.81%			
3	45.1	44.6	-0.50	-1.11%			
NIST U	NIST Uncertainty (k=1) 0.90 Transfer Alanine						
NPL (Jncertainty (k=1)	1.20	Transfer Alanine				
	<u>Equivale</u>	ncy Calculation					
Dose							
Point ID	En @ k=1	En @ k=2	En @ k=3				
1	-0.05	-0.02	-0.02				
2	-0.13	-0.07	-0.04				
3	-0.33	-0.18	-0.11				

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The NIST and NPL dose results from the "variable" industrial 10 MeV electron beam radiation process field are shown in the table below.

Variable Dose/Temp Radiation Conditions						
Dose						
Point ID	NIST kGy	GEX kGy	Difference kGy	% Difference		
1	0.807	0.84	0.03	4.09%		
2	2.30	2.29	-0.01	-0.43%		
3	5.56	5.48	-0.08	-1.44%		
4	11.0	11.1	0.10	0.91%		
5	18.7	19.3	0.60	3.21%		
6	47.7	47.9	0.20	0.42%		
7	72.5	70.8	-1.70	-2.34%		
NIST Un	NIST Uncertainty (k=1) 0.90 Transfer Alanine					
GEX Un	certainty (k=1)	1.35	Transfer Alanine	ansfer Alanine		
	<u>Equivale</u>	ncy Calculation	<u>1</u>			
Dose						
Point ID	En @ k=1	En @ k=2	En @ k=3			
1	0.02	0.01	0.01			
2	-0.01	0.00	0.00			
3	-0.05	-0.02	-0.02			
4	0.06	0.03	0.02			
5	0.37	0.18	0.12			
6	0.12	0.06	0.04			
7	-1.05	-0.52	-0.35			

Dose results are shown to be equivalent and in agreement within the combined limits of the stated measurement uncertainty of the laboratories. Higher point-to-point variability is observable compared with the highly uniform results obtained from the "fixed" radiation field irradiations. However, these results also carry an expectation of higher variability associated with the "variable" irradiation conditions of a large-scale industrial field where temperature and dose rate variability come into play as well as higher dosimeter handling and placement uncertainty components.

For example, the variability of irradiation temperature measurements made in an industrial irradiator using irreversible temperature indicators with 2.5°C increments is larger than temperature measurements obtained using the high precision thermistor or thermocouple used to maintain and control fixed temperature Gammacell 220 irradiations.

In summary, the dose results obtained from the "fixed" and "variable" radiation field experiments indicate agreement with all but one dose point (70 kGy target) being in agreement within the ±1-sigma combined laboratory uncertainty limits.

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The NIST-traceable calibration chain of dose used in the GEX dose estimates for the "fixed" irradiation conditions, as shown below, reflected close agreement, with all dose points found within ±1-sigma of the combined uncertainty limits of the labs.

	Fixed Dose/				
	Lab Intercom	ts			
Dose					
Point ID	NIST kGy	GEX kGy	Difference kGy	% Difference	
1	7.45	7.39	-0.06	-0.81%	
2	24.8	25.0	0.20	0.81%	
3	45.1	45.2	0.10	0.22%	
NIST U	Jncertainty (k=1)	0.600	Transfer Alanine		
GEX (Jncertainty (k=1)	1.065	Transfer Alanine		
	Equivalency Calculation				
Dose					
Point ID	En @ k=1	En @ k=2	En @ k=3		
1	-0.05	-0.02	-0.02		
2	0.16	0.08	0.05		
3	0.08	0.04	0.03		

NIST / GEX comparison results obtained from the "variable" radiation field experiment were also shown to be in agreement, with all dose points shown equivalent within the ±2-sigma combined uncertainty limits of the labs.

The same equivalency analysis was also performed on the NPL calibration traceability chain by comparing the NPL results with the DTU Risø data as shown below.

	Fixed Dose/						
Cal ID#3451 DTU Risø/NPL Lab Intercomparison Results							
Dose							
Point ID	DTU Risø kGy	NPL kGy	Difference kGy	% Difference			
1	7.55	7.38	0.17	2.25%			
2	25.1	24.6	0.50	1.99%			
3	45.3	44.6	0.70	1.55%			
DTU Risø U	DTU Risø Uncertainty (k=1) 1.75 Transfer Alanine						
NPL U	Incertainty (k=1)	1.20	Transfer Alanine	er Alanine			
	<u>Equivale</u>	ncy Calculation					
Dose							
Point ID	En @ k=1	En @ k=2	En @ k=3				
1	-0.08	-0.04	-0.03				
2	-0.24	-0.12	-0.08				
3	-0.33	-0.17	-0.11				

The DTU and NPL reported doses are shown to be equivalent with all dose points shown in agreement within ±1-sigma of the overall combined uncertainty limits stated by the labs.

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Below are the DTU Risø/NPL comparison results for the "variable" irradiation conditions experiment.

<u> </u>	/ariable Dose	Temp Radia	tion Conditions					
Cal ID#	Cal ID#3452 DTU Risø/NPL Lab Intercomparison Results							
Dose								
Point ID	DTU Risø kGy	NPL kGy	Difference kGy	% Difference				
1	0.81	0.83	-0.02	-2.47%				
2	2.30	2.27	0.03	1.30%				
3	5.33	5.44	-0.11	-2.06%				
4	11.1	10.9	0.20	1.80%				
5	18.6	18.3	0.30	1.61%				
6	47.4	46.6	0.80	1.69%				
7	70.0	69.4	0.60	0.86%				
J Risø Un	certainty (k=1)	1.75	Transfer Alanin	2				
NPL Un	certainty (k=1)	1.20	Transfer Alanin	nsfer Alanine				
	<u>Equivale</u>	ncy Calculatio	<u>n</u>					
Dose								
Point ID	En @ k=1	En @ k=2	En @ k=3					
1	0.01	0.00	0.00					
2	-0.01	-0.01	0.00					
3	0.05	0.03	0.02					
4	-0.09	-0.05	-0.03					
5	-0.14	-0.07	-0.05					
6	-0.38	-0.19	-0.13					
7	-0.28	-0.14	-0.09					

The DTU and NPL reported doses are shown to be equivalent with all dose points in agreement within ±1-sigma of the overall combined uncertainty limits stated by the labs.

The equivalency of the NIST and NPL primary standard traceability chains at the secondary standard lab level was also evaluated by comparing results between GEX that utilized a NIST-traceable calibration and DTU Risø which used an NPL-traceable calibration as shown below.

<u>Fi</u>							
	Lab Intercomparison Results						
Dose Point ID	DTU Risø	GEX kGy	Difference kGy	% Difference			
1	7.55	7.39	0.16	2.12%			
2	25.1	25.0	0.10	0.40%			
3	45.3	45.2	0.10	0.22%			
DTU Risø Unce	rtainty (k=1)	1.750	Transfer Alanin	e			
GEX Unce	rtainty (k=1)	1.065	Transfer Alanine				
	Equivalenc	y Calculation					
Dose Point ID	En @ k=1	En @ k=2	En @ k=3				
1	-0.08	-0.04	-0.03				
2	-0.05	-0.02	-0.02				
3	-0.05	-0.02	-0.02				

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The results of the "fixed" radiation field conditions show the DTU Risø NPL-traceable doses and the GEX NIST-traceable equivalency demonstrated of all dose points within ±1-sigma limits.

Below are the comparison results for the "variable" radiation conditions of the 10 MeV industrial irradiator experiment.

Variable Dose/Temp Radiation Conditions								
Lab Intercomparison Results								
Dose Point ID	DTU Risø	GEX kGy	Difference kGy	% Difference				
1	0.81	0.84	-0.03	-3.70%				
2	2.30	2.29	0.01	0.43%				
3	5.33	5.48	-0.15	-2.81%				
4	11.1	11.1	0.00	0.00%				
5	18.6	19.3	-0.70	-3.76%				
6	47.4	47.9	-0.50	-1.05%				
7	70.0	70.8	-0.80	-1.14%				
DTU Risø Uncertainty (k=1) 1.75 Transfer Alanine								
	, , ,							
	ertainty (k=1)	1.35	Transfer Alanin	ie				
		1.35	Transfer Alanir	ie				
				ie				
	ertainty (k=1)			ie				
	ertainty (k=1)			ne				
GEX Unc	ertainty (k=1) Equivalency (Calculation		ne				
GEX Unco	Equivalency of En @ k=1	Calculation En @ k=2	En @ k=3	ne				
Dose Point ID	Equivalency (En @ k=1	Calculation En @ k=2 0.01	En @ k=3 0.00	ne				
Dose Point ID	Equivalency (Fig. 1) Equivalency (Fig. 2) En @ k=1 0.01 0.00	En @ k=2 0.01 0.00	En @ k=3 0.00 0.00	ne				
Dose Point ID 1 2 3	En @ k=1 0.01 0.00 0.07	En @ k=2 0.01 0.00 0.04	En @ k=3 0.00 0.00 0.02	ne				
Dose Point ID 1 2 3 4	En @ k=1 0.01 0.00 0.07 0.00	En @ k=2 0.01 0.00 0.04 0.00	En @ k=3 0.00 0.00 0.02 0.00	ne				

The results of the "variable" irradiation experiment show the DTU Risø and GEX reported doses to be equivalent within ±1 sigma of the overall combined limits of the labs.

Conclusion

Analysis of the data sets of doses reported by the four HDRL laboratory dose inter-comparison provides evidence that the traceable doses reported by the laboratories were shown to be equivalent within the normally expected overall combined uncertainty limits of the comparing laboratories.

This study, although limited to only a single set of "fixed" and "variable" radiation field experiments, does provide evidence that doses, as currently being realized and reported within the two major Primary National Standards laboratory dose chains, NIST and NPL, are demonstrated equivalent under actual "field" conditions when the stated combined uncertainties of the laboratories are taken into account.

The dose data from this study also provides an example of normally expected magnitude dose differences that may be expected to occur from a single calibration exercise. Multiple calibration exercises and calibration audit exercises would be expected to yield results consistent with this study and provide evidence of reproducibility.

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Although formal HDRL inter-comparisons are performed and monitored by BIPM and reported in the literature periodically, field study experiments such as this demonstrate actual results obtained under normal industry usage conditions. GEX hopes to be able to continue to fund and conduct these types of actual industry usage studies to demonstrate the continued effectiveness among international dose standards. We are also hopeful that future funding of the national standard labs will be sufficient to sustain existing national dose standards, and support their continued staff participation in important international industry association activities.

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Recent BIPM International High Dose Inter-comparisons:

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REVISION CONTROL HISTORY

DATE	CHANGE DESCRIPTION	REVISION
09/01/16	Initial release.	Α

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