#### Deriving Correction Factors for a Primary Standard for Radiation Dosimetry

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Presented at the 2011 COMSOL Conference Oct 13, 2011



### **The NIST Water Calorimeter**





#### Vessel with thermistors

Vessel with an ion chamber



### Application Note: chamber calibration





- NIST water calorimeter terminates calibration chain for QA protocols in clinics/hospitals.
- Calibration transferred to referencequality standards used in secondary calibration labs.

### The "Bureau International des Poids et Mesures" (BIPM)

- BIPM = International laboratory created by the metre convention; has an lonizing Radiation Division
- Role: "The task of the BIPM is to ensure world-wide uniformity of measurements and their traceability to the International System of Units (SI)."
- www.bipm.org



### **BIPM Intercomparison Program:**

- Enables NMIs (like NIST) to declare calibration and measurement capabilities (CMCs)
- Key comparisons and database (http://kcdb.bipm.org)



## Water Calorimetry at NIST





## Heat Transport Corrections

Questions regarding existing remedies:

- Would the system exhibit a stable, steady-state behavior? Interruptions to reestablish thermal equilibrium might not be necessary.
- If so, to what extent would convection contribute? *Possibility to operate at room temperature instead of 3.98 °C.*



## Heat Transport



- Beam: distributed heat source with semiempirical depth and transverse dose gradients.
- Ext stirring: simulated by elevated thermal conductivity in unstirred water.
- Heat transport: conduction only; no convection.

### Response to chopped <sup>60</sup>Co beam -time domain



### Response to chopped <sup>60</sup>Co beam -frequency domain



## k<sub>ht</sub> vs. shutter time (sec)

#### - derived from simulation



 $k_{ht}$  (70 sec) = 1.0009

# Conduction-like behavior obtained at low duty cycle (70s on, 1940 s off)



### Heat Transport cont'd

Variation of "apparent" dose rate with shutter freq/period demonstrates agreement with heat equation (i.e. conduction only) *except* at higher shutter periods.

 $\rightarrow$  steady-state operation at room temperature appears feasible at shorter exposure periods.



### Heat Defect

Table 1. Model IIIR: reactions and rate constants (4 °C)

Reactions*				Rate constar
1	e + e	$\rightarrow$	$H_2 + OH^- + OH^-$	$3.48 \times 10$
2	e_ + H	$\rightarrow$	$H_2 + OH^-$	$1.73 \times 10$
3	e <sub>au</sub> + OH	$\rightarrow$	OH-	$2.38 \times 10$
4	$e_{aa}$ + H <sub>2</sub> O <sub>2</sub>	$\rightarrow$	OH <sup>-</sup> + OH	$8.84 \times 10$
5	$e_{aq} + O_2$	$\rightarrow$	Ož	$1.16 \times 10$
6	$e_{aq} + O_2$	$\rightarrow$	$HO_2 + OH^-$	$8.48 \times 10$
7	$e_{aq}^-$ + HO <sub>2</sub>	$\rightarrow$	HO <sub>2</sub>	$8.48 \times 10$
8	H+H	$\rightarrow$	H <sub>2</sub>	$3.44 \times 10$
9	H + OH	$\rightarrow$	H <sub>2</sub> O	$1.21 \times 10$
10	$H + H_2O_2$	$\rightarrow$	$OH + H_2O$	$3.18 \times 10$
11	$H + O_2$	$\rightarrow$	HO <sub>2</sub>	$9.58 \times 10$
12	$H + HO_2$	$\rightarrow$	H <sub>2</sub> O <sub>2</sub>	$7.24 \times 10$
13	$H + O_2^-$	$\rightarrow$	HO <sub>2</sub>	$7.24 \times 10$
14	OH + OH	$\rightarrow$	H <sub>2</sub> O <sub>2</sub>	$3.76 \times 10$
15	$OH + H_2$	$\rightarrow$	$H + H_2O$	$2.40 \times 10$
16	$OH + H_2O_2$	$\rightarrow$	$H_2O + H_2O$	$1.79 \times 10$
17	$OH + HO_2$	$\rightarrow$	$H_2O + O_2$	$9.08 \times 10$
18	$OH + O_2^-$	$\rightarrow$	$OH^- + O_2$	$7.89 \times 10$
19	$HO_2 + HO_2$	$\rightarrow$	$H_2O_2 + O_2$	$3.72 \times 10$
20	$HO_2 + O_2^-$	$\rightarrow$	$H_2O_2 + O_2 + OH^-$	$5.84 \times 10$
21	H <sub>2</sub> O	$\rightarrow$	$H^* + OH^-$	$2.22 \times 10$
22	$H^+ + OH^-$	$\rightarrow$	H <sub>2</sub> O	$7.23 \times 10$
23	H <sub>2</sub> O <sub>2</sub>	$\rightarrow$	$H^* + HO_2^-$	$1.34 \times 10$
24	$H^{+} + HO_{2}$	$\rightarrow$	H <sub>2</sub> O <sub>2</sub>	$3.13 \times 10$
25	$H_2O_2 + OH^-$	$\rightarrow$	$HO_2^- + H_2O$	$7.56 \times 10$
26	$HO_2 + H_2O$	$\rightarrow$	$H_2O_2 + OH^-$	$5.45 \times 10$
27	н	$\rightarrow$	$e_{aq}^{-}$ + H <sup>+</sup>	8.83 × 10
28	e <sub>aq</sub> + H <sup>*</sup>	$\rightarrow$	Н	1.88 × 10
29	$e_{aq} + H_2O$	$\rightarrow$	H+OH	$5.08 \times 10$
30	H+OH-	$\rightarrow$	e <sub>aq</sub> + H <sub>2</sub> O	$7.77 \times 10$
31	OH UN OF	$\rightarrow$	H'+0	1.34 × 10
32	H'+0	$\rightarrow$	OH U O	3.13 × 10
33	OH + OH	$\rightarrow$	$O + H_2O$	7.56 × 10
24	0 + H <sub>2</sub> 0	$\rightarrow$	OH + OH	5.45 × 10
30	HO <sub>2</sub>		02 + H	4.21 × 10
27	U2 + H	-	0.102	5.15 × 10
29	0-100	-	UQ + H20	1.04 × 10
20	$O_2 + H_2O$		HU2 + 0H	1.94 × 10 7.05 × 10
40	0-140		0-140	2.44 × 10
40	0 + H <sub>2</sub> O <sub>2</sub>	-	01 + H0	5.44 × 10
42	01+102		HO:	6.02 × 10
42			0-+04-	$2.10 \times 10$
43	$e_{aq} + 10_2$		OH-+ OH-	1.82 × 10
45	0-0		0	$2.62 \times 10$
46	0:	_	0.+0	670 × 10
40	0" + H05	$\rightarrow$	05 + 0H	$2.84 \times 10$
48	0 + 05		OH-+OH-+O-	4.26 × 10
49	$HO_{2} + H_{2}O_{2}$	$\rightarrow$	$OH + H_0 + O_2$	$2.90 \times 10$
50	$05 + H_{2}O_{2}$	$\rightarrow$	$OH^- + OH + O$	9 30 × 10
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N.V. Klassen and Carl K. Ross, J. Res. Natl. Inst. Stand. Techol. **107**, 171-178 (2002).

#### Causes:

its<sup>b</sup>

 Chemical reactions involving products of incident radiation and various dissolved species within the water

#### Effect on signals:

- Transient can be huge (~100%).
- Steady state depends on dissolved species (0 to few %).

#### Remedy:

 $H_2$  – saturated, high-purity water in a sealed glass vessel.



## Summary

We convert from temperature rise to absorbed dose as follows:

