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**Appendix F**

**Reproducibility Analyses for LLNA: BrdU-ELISA with Decision  
Criterion of SI  $\geq$  1.5**

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## 30 **1.0 Test Method Reliability**

31 **Section 7** provides the reproducibility analyses for the LLNA: BrdU-ELISA using  $SI \geq 2.0$  to  
32 classify substances as sensitizers. The decision criterion of  $SI \geq 2.0$  was used in the JSAAE  
33 interlaboratory validation study. The  $SI \geq 2.0$  criterion produced an accuracy of 87% (27/31),  
34 a false positive rate of 0% (0/9), and a false negative rate of 18% (4/22) when LLNA: BrdU-  
35 ELISA results were compared to the results of the traditional LLNA (**Table 6-6**). This  
36 appendix provides the reproducibility analyses using  $SI \geq 1.5$  to classify substances as  
37 sensitizers. This was one of the alternate SI criterion evaluated in **Section 6.5**. The  $SI \geq 1.5$   
38 criterion produced an accuracy of 84% (26/31), a false positive rate of 33% (3/9), and a false  
39 negative rate of 9% (2/22) when LLNA: BrdU-ELISA results were compared to the results of  
40 the traditional LLNA (**Table 6-6**).

### 41 **1.1 Intralaboratory Reproducibility**

42 The test results for the LLNA: BrdU-ELISA were amenable to intralaboratory reproducibility  
43 analyses for three endpoints: sensitizer or nonsensitizer classification, SI values, and EC1.5  
44 values. Analyses of intralaboratory reproducibility were performed using a concordance  
45 analysis for the qualitative results (sensitizer vs. nonsensitizer) (**Section 1.1.1**) and a CV  
46 analysis for the quantitative results (SI values and EC3 values) (**Sections 1.1.2 and 1.1.3**,  
47 respectively).

#### 48 1.1.1 *Intralaboratory Reproducibility – Qualitative Results*

49 The dataset available for an intralaboratory concordance analysis of the qualitative test  
50 results for the LLNA: BrdU-ELISA included eight substances that were tested multiple times  
51 and classified as sensitizers or nonsensitizers. Hexyl cinnamic aldehyde was tested six times,  
52 eugenol was tested five times, and isoeugenol was tested three times, and 2,4-  
53 dinitrochlorobenzene, glutaraldehyde, hexane, 4-phenylenediamine, and propylene glycol  
54 were each tested twice (Takeyoshi et al. 2003, 2004a, 2005, 2006, 2007a; unpublished data)  
55 (**Table F-1**). All substances were sensitizers in the traditional LLNA except for propylene  
56 glycol and hexane. The multiple test results for 7/8 substances were 100% concordant when  
57  $SI \geq 1.5$  was used to classify substances as sensitizers. Discordant test results were noted for  
58 propylene glycol tested at a maximum concentration of 50%. The test result from Takeyoshi

59 et al. (2005) was positive (SI = 1.6) while the result from Takeyoshi et al. (2006) produced a  
60 negative result (SI = 0.9). Both tests used AOO as the vehicle.

61 By comparison, the qualitative intralaboratory concordance analysis for the traditional LLNA  
62 (ICCVAM 1999) was based on a dataset of six substances that included six results each for  
63 benzocaine and hexyl cinnamic aldehyde, five results for eugenol, four results each for  
64 isoeugenol and methyl salicylate, and three results for 2,4-dinitrochlorobenzene.

65 Intralaboratory results for each substance were 100% concordant with the exception of  
66 benzocaine. One of the six benzocaine (5/6 or 83% concordance) results in the traditional  
67 LLNA was reported as equivocal because SI increased with dose, but did not reach the  
68 criterion of  $SI \geq 3.0$ . Thus, the proportion of substances for which intralaboratory  
69 concordance of qualitative results was 100% was similar for LLNA: BrdU-ELISA (7/8) and  
70 the traditional LLNA (5/6).

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71 **Table F-1 Intralaboratory Reproducibility for the LLNA: BrdU-ELISA Outcome of**  
 72 **Substances Tested Multiple Times**

Substance	Highest Concentration Tested (%)	Highest SI	Outcome <sup>1</sup>	Takeyoshi et al. Reference
2,4-Dinitro-chlorobenzene	2	17.9	+	2005
	2	6.8	+	2006, 2007b
Eugenol	30	3.3	+	2004a
	30	3.8	+	2007a
	50	12.3	+	2005
	50	3.1	+	2006
	50	17.7	+	2007b
Glutaraldehyde	2	14.6	+	2005, 2007b
	10	15.5	+	2005, 2007b
Hexane	50	1.9	+	2005
	100	1.8	+	Unpublished data
Hexyl cinnamic aldehyde	25	2.4	+	2003
	50	3.6	+	2003
	50	5.9	+	2005
	50	3.6	+	2006
	50	2.7	+	2006
	50	3.0	+	2007b
Isoeugenol	10	8.4	+	2005
	10	2.4	+	2006, 2007b
	30	6.7	+	
4-Phenylenediamine	2	11.7	+	2005, 2007b
	10	14.7	+	2005, 2007b
Propylene glycol	50	1.6	+	2005
	50	0.9	-	2006, 2007b

73 Abbreviations: LLNA: BrdU-ELISA = Murine local lymph node assay (LLNA) with enzyme-linked  
 74 immunosorbent assay (ELISA) detection of bromodeoxyuridine (BrdU); SI = Stimulation index.

75 <sup>1</sup>(+) = Sensitizer; (-) = nonsensitizer

76 1.1.2 *Intralaboratory Reproducibility – SI*

77 There were six substances that were tested multiple times by Takeyoshi et al. (2003, 2004a,  
78 2005, 2006, 2007a, 2007b, unpublished data). Because two substances had multiple tests for  
79 more than one concentration, there were nine substance/concentration combinations that  
80 were tested two to five times in separate experiments. The multiple SI values for each  
81 substance/concentration were used to calculate a CV for the assessment of intralaboratory  
82 variability. As shown by **Table F-2**, the CVs ranged from 1% (25% hexyl cinnamic  
83 aldehyde) to 79% (10% isoeugenol). The intralaboratory reproducibility of the traditional  
84 LLNA was not assessed by CV analysis of SI values (ICCVAM 1999).

85 1.1.3 *Intralaboratory Reproducibility – EC1.5*

86 CV values were also calculated for the EC1.5 values for the three sensitizers that were tested  
87 more than once using multiple doses by Takeyoshi et al. (2003; 2004a, 2005, 2006, 2007a,  
88 2007b). The individual animal data for eugenol, hexyl cinnamic aldehyde, and isoeugenol,  
89 were used to calculate EC1.5 values for the LLNA: BrdU-ELISA. The methods for  
90 calculating EC1.5 values for each sensitizer were modified from those used by Ryan et al.  
91 (2007) to calculate EC3 values. Linear interpolation was used to calculate EC1.5 values for  
92 each test with SI values higher or lower than two and extrapolation was used to calculate  
93 EC1.5 values for tests with no SI values below two. The equation for linear interpolation  
94 was:

$$EC1.5 = c + \left[ \frac{(1.5 - d)}{(b - d)} \right] \times (a - c)$$

95 The linear interpolation equation uses the points immediately above and below SI = 2, with  
96 the (dose, SI) coordinates of (a, b) immediately above SI = 2 and (c, d) immediately below SI  
97 = 2. The equation for extrapolation was:

98

$$EC1.5_{ex} = 2 \left\{ \log_2(c) + \frac{(1.5-d)}{(b-d)} \times [\log_2(a) - \log_2(c)] \right\}$$

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100 The extrapolation equation uses the two points immediately above SI = 2, with the  
101 coordinates of (a, b) for the point closest to SI = 2, and (c, d) for the higher point.

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103**Table F-2 Intralaboratory Reproducibility for the SI of Tested Substances in LLNA: BrdU-ELISA - Coefficient of Variation**

Substance	Concentration Tested (%)	SI	Mean	SD	CV (%)	Takeyoshi et al. Reference
2,4-Dinitrochlorobenzene	2	17.9	12.4	7.8	64	2005
	2	6.8				2006, 2007b
Eugenol	30	3.3	3.6	0.4	10	2004a
	30	3.8				2007a
Eugenol	50	12.3	11.0	7.4	67	2005
	50	3.1				2006
	50	17.7				2007b
Hexane	50	1.9	1.8	0.07	4	2005
	50	1.8				Unpublished
Hexyl cinnamic aldehyde	12.5	1.87	1.73	0.21	12	2003
	12.5	1.58				2003
Hexyl cinnamic aldehyde	25	2.42	2.4	0.01	1	2003
	25	2.40				2003
Hexyl cinnamic aldehyde	50	3.6	3.8	1.3	34	2003
	50	5.9				2005
	50	3.6				2006
	50	2.7				2006
	50	3.0				2007b
Isoeugenol	10	8.4	5.4	4.2	79	2005
	10	2.4				2006, 2007b
Propylene glycol	50	1.6	1.1	0.6	55	2005
	50	0.7				2006, 2007b

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Abbreviations: LLNA: BrdU-ELISA = Murine local lymph node assay (LLNA) with enzyme-linked immunosorbent assay (ELISA) detection of bromodeoxyuridine (BrdU); CV = Coefficient of variation; SD = Standard deviation; SI = Stimulation index.

108 As shown in **Table F-3**, there were five EC1.5 values for hexyl cinnamic aldehyde, four  
 109 EC1.5 values for eugenol, and two EC1.5 values for isoeugenol. The CV values were 37%  
 110 for hexyl cinnamic aldehyde, 66% for eugenol, and 52% for isoeugenol. The ICCVAM  
 111 LLNA *Performance Standards* criteria for demonstrating adequate intralaboratory  
 112 reproducibility is based on results from at least four independent tests of hexyl cinnamic  
 113 aldehyde (ICCVAM 2009). Intralaboratory reproducibility is considered adequate when each  
 114 test yields an ECt value (i.e., the estimated concentration needed to produce an SI of a  
 115 specific threshold value, 1.5, in this case) within 5% to 20% (ICCVAM 2009). All five  
 116 EC1.5 values for hexyl cinnamic aldehyde were within the acceptable range for  
 117 intralaboratory reproducibility.

118 **Table F-3 Intralaboratory Reproducibility for the EC1.5 of Tested Substances in**  
 119 **LLNA: BrdU-ELISA - Coefficient of Variation**

Substance	EC1.5	Mean	SD	CV (%)	Takeyoshi et al. Reference
Eugenol	5.9	7.2	4.7	66	2004a
	11.0				2006
	10.7				2007a
	1.0				2007b
Hexyl cinnamic aldehyde	11.6	12.9	4.8	37	2003
	5.5				2003
	15.9				2006
	18.1				2006
	13.5				2007b
Isoeugenol	6.3	4.6	2.4	52	2006, 2007b
	2.9				2007a

120 Abbreviations: LLNA: BrdU-ELISA = Murine local lymph node assay (LLNA) with enzyme-linked  
 121 immunosorbent assay (ELISA) detection of bromodeoxyuridine (BrdU); CV = Coefficient of variation; EC1.5 =  
 122 Estimated concentration needed to produce a stimulation index of two; SD = Standard deviation.

123

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124 The intralaboratory reproducibility of the traditional LLNA was assessed by CV analysis of  
 125 EC3 values using a larger dataset (ICCVAM 1999) than that available for the LLNA: BrdU-  
 126 ELISA analysis. Two EC3 values were reported by each of five laboratories for 2, 4-dinitro-  
 127 chlorobenzene, five EC3 values were reported by one laboratory for isoeugenol, six EC3  
 128 values were reported for hexyl cinnamic aldehyde by two laboratories, and five EC3 values  
 129 were reported for eugenol by one laboratory (Table F-4).

130 **Table F-4 Intralaboratory Reproducibility for the EC3 of Tested Substances in the**  
 131 **Traditional LLNA<sup>1</sup>**

Substance	Number of Laboratories	Number of Tests per Laboratory	CV (%)
2, 4-Dinitrochlorobenzene	5	2	13 – 47
Isoeugenol	1	5	26
Hexyl cinnamic aldehyde	2	6	19-27
Eugenol	1	5	18

132 Abbreviations: LLNA = Murine local lymph node assay (LLNA) with enzyme-linked immunosorbent assay  
 133 (ELISA) detection of bromodeoxyuridine (BrdU); CV = Coefficient of variation; EC3 = Estimated  
 134 concentration needed to produce a stimulation index of three.

135 <sup>1</sup>From ICCVAM (1999).

136 For all three substances in common, the intralaboratory CV values for the EC1.5 values from  
 137 LLNA: BrdU-ELISA tests were higher than those reported in ICCVAM (1999) for EC3  
 138 values from the traditional LLNA. The intralaboratory EC1.5 CV for the LLNA: BrdU-  
 139 ELISA tests of eugenol was 66% vs. 18% for the CV of EC3 values reported by ICCVAM  
 140 (1999). The intralaboratory EC1.5 CV for isoeugenol was 52% vs. 26% for the CV of EC3  
 141 values from ICCVAM (1999), and the intralaboratory EC1.5 CV for hexyl cinnamic  
 142 aldehyde was 37% vs. 19 to 27% for the CV reported by ICCVAM (1999) for EC3 values.

## 143 1.2 Interlaboratory Reproducibility

144 The interlaboratory reproducibility of the LLNA: BrdU-ELISA was assessed using the  
 145 individual animal data from the multi-laboratory validation study organized by the JSAAE  
 146 (Kojima et al. 2008). The study design is described in Section 7.2. The LLNA: BrdU-ELISA  
 147 test results from the study are amenable to interlaboratory reproducibility analyses for two  
 148 endpoints: sensitizer or nonsensitizer classification and EC2 values. Analyses of  
 149 interlaboratory reproducibility were performed using a concordance analysis for the

150 qualitative results (sensitizer vs. nonsensitizer) (**Section 1.2.1**) and a CV analysis for the  
151 quantitative results (EC1.5 values) (**Section 1.2.2**).

152 1.2.1 *Interlaboratory Reproducibility – Qualitative Results*

153 The available quantitative absorbance data for interlaboratory reproducibility analysis were  
154 used to calculate SI values for each substance and dose tested. Substances with  $SI \geq 1.5$  at  
155 any dose were classified as sensitizers. The qualitative (i.e., sensitizer vs. nonsensitizer)  
156 interlaboratory concordance analysis for the 10 substances tested during Phase II of the  
157 JSAAE interlaboratory validation study is shown in **Table F-6**. The qualitative comparison  
158 of LLNA: BrdU-ELISA results for nine substances tested in up to seven laboratories show  
159 that interlaboratory concordance was 100% (3/3, 6/6, or 7/7). However, one of these  
160 substances, lactic acid, was misclassified as a nonsensitizer in all three laboratories. The  
161 concordance for isopropanol, the substance that produced discordant results among  
162 laboratories, the concordance was 50% (3/6). The test of isopropanol at Laboratory 2 failed  
163 ( $SI = 1.09$ ) because the concurrent positive control ( $SI = 1.29$ ) failed the acceptance criterion  
164 of  $SI \geq 2$ . The other six laboratories reported maximum SI values of 2.22, 0.98, 1.57, 0.94,  
165 2.04, and 1.01 for isopropanol. Isopropanol produces a nonsensitizer result in the traditional  
166 LLNA.

167 The Validation Management Team, which evaluated the reproducibility using  $SI \geq 2$  to  
168 identify sensitizers, considered the interlaboratory reproducibility to be acceptable (Kojima et  
169 al. 2008). Because the evaluation of interlaboratory reproducibility for the traditional LLNA  
170 did not include an evaluation of qualitative results (ICCVAM 1999), there were no traditional  
171 LLNA concordance data for comparison with the LLNA: BrdU-ELISA concordance.

172

172 **Table F-6 Qualitative Results for the Phase II Interlaboratory Validation Study on**  
 173 **the LLNA: BrdU-ELISA<sup>1</sup>**

Substance	Laboratory							Concordance
	1	2	3	4	5	6	7	
2,4-Dinitrochlorobenzene	+	+	+	+	+	+	+	7/7
Glutaraldehyde	+				+	+		3/3
Nickel sulfate			+	+			+	3/3
<i>trans</i> -Cinnamic aldehyde		+		+	+			3/3
Formaldehyde	+				+	+		3/3
Eugenol		+				+	+	3/3
Hexyl cinnamic aldehyde	+	- <sup>3</sup>	+	+	+	+ <sup>5</sup>	+	6/6
Isopropanol	+	- <sup>3</sup>	-	+	-	+ <sup>4</sup>	-	3/6
Lactic acid			+	+			+	3/3
Methyl salicylate	-	-	-					3/3

174 Abbreviation: LLNA: BrdU-ELISA = Murine local lymph node assay (LLNA) with enzyme-linked immunosorbent assay  
 175 (ELISA) detection of bromodeoxyuridine (BrdU).

176 <sup>1</sup>(+) indicates sensitizer result; (-) indicates nonsensitizer result using  $SI \geq 1.5$  to classify sensitizers.

177 <sup>2</sup>Test failed because concurrent positive control ( $SI = 1.29$ ) failed the acceptance criterion (i.e.,  $SI < 2$ ). The positive control  
 178 would have also failed if the acceptance criterion was  $SI \geq 1.5$ . This isopropanol result was not included in the concordance  
 179 analysis.

180 <sup>3</sup>Three mice tested at highest dose.

181 <sup>4</sup>Three mice per dose group.

### 182 1.2.2 Interlaboratory Reproducibility – EC1.5 Values

183 The SI values for each test used to calculate EC1.5 values for each sensitizer according to the  
 184 methods reported in Section 1.1.3. The EC1.5 values from each laboratory were used to  
 185 calculate CV values for each substance. The resulting values are shown in **Table F-7**. CV  
 186 values ranged from 31% (*trans*-cinnamic aldehyde) to 95% (glutaraldehyde). The mean CV  
 187 was 63%.

188 The ICCVAM LLNA *Performance Standards* indicate that interlaboratory reproducibility  
 189 should be evaluated with at least two sensitizing chemicals with well-characterized activity in  
 190 the traditional LLNA (ICCVAM 2009). Acceptable reproducibility is attained when each  
 191 laboratory obtains ECt values within 0.025% to 0.1% for 2,4-dinitrochlorobenzene and  
 192 within 5% to 20% for hexyl cinnamic aldehyde (ICCVAM 2009). For 2,4-dinitrochloro-  
 193 benzene, the EC1.5 values from four laboratories were outside the acceptable range, and for  
 194 hexyl cinnamic aldehyde, the EC1.5 values from four laboratories were outside the  
 195 acceptable range. All values outside the acceptable ranges were below the low end of the  
 196 range. This indicates that the discordance was due to the LLNA: BrdU-ELISA producing a  
 197 more sensitive result.

198 **Table F-7 EC1.5 Values from the Phase II Interlaboratory Validation Study of the LLNA: BrdU-ELISA<sup>1</sup>**

Substance	Laboratory							Mean	% CV
	1	2	3	4	5	6	7		
Glutaraldehyde	0.064	NT	NT	NT	0.031	0.21	NT	0.10	95
Nickel sulfate	NT	NT	1.5	0.5	NT	NT	0.6	0.8	65
<i>trans</i> -Cinnamic aldehyde	NT	1.7	NT	1.0	1.8	NT	NT	1.5	31
Formaldehyde	0.3	NT	NT	NT	0.2	0.6	NT	0.3	66
Eugenol	NT	12.5	NT	NT	NT	10.5	3.5	8.8	54
<b>2,4-Dinitro-chlorobenzene</b>	0.058 (4.3 @ 1%)	0.010 (8.37 @ 1%)	0.022 (5.99 @ 0.3%)	0.022 (5.50 @ 1%)	0.0022 (18.80 @ 0.3%)	0.015 (4.83 @ 0.3%)	0.049 (12.18 @ 1%)	0.025	81
<b>Hexyl cinnamic aldehyde</b>	9.4 (3.4 @ 50%)	- <sup>1</sup> (1.83 @ 50%)	15.2 (2.87 @ 50%)	4.1 (3.34 @ 50%)	3.5 (13.5 @ 50%)	7.9 <sup>2</sup> (3.27 @ 50%)	9.5 (3.84 @ 50%)	8.3	52

199 Note: Bolded font indicates substances recommended for assessing interlaboratory reproducibility in *Recommended Performance Standards* (ICCVAM 2009). Shading shows  
 200 EC1.5 values that are outside of the acceptable range from the ICCVAM *LLNA Performance Standards*: 5 - 20% for hexyl cinnamic aldehyde and 0.025 - 0.1% for 2,4-  
 201 dinitrochlorobenzene. Values in parentheses are highest SI values achieved.

202 Abbreviations: CV =Coefficient of variation; LLNA: BrdU-ELISA = Murine local lymph node assay (LLNA) with enzyme-linked immunosorbent assay (ELISA) detection  
 203 of bromodeoxyuridine (BrdU); NT = Not tested; SI = Stimulation index.

204 <sup>1</sup>Test failed because associated positive control failed acceptance criterion (i.e., SI < 2; vehicle control absorbance was unusually high). At SI= 1.29, the positive control  
 205 would have failed even if the acceptance criterion was SI ≥ 1.5. Result not included in the mean EC1.5 and CV.

206 <sup>2</sup>Three mice tested at highest dose.  
 207

208 The interlaboratory CV values for the LLNA: BrdU-ELISA EC1.5 values were higher than  
 209 that for the traditional LLNA EC3 values. The analysis of interlaboratory variation of EC3  
 210 values for the traditional LLNA reported CV values of 7 to 84% for five substances tested in  
 211 five laboratories (**Table F-8**; ICCVAM 1999). Three of the same substances were evaluated  
 212 in the traditional LLNA and the LLNA: BrdU-ELISA. All interlaboratory CV values for the  
 213 EC1.5 from LLNA: BrdU-ELISA tests were greater than that for EC3 values from the  
 214 traditional LLNA. The CV of 81% for EC1.5 values for 2,4-dinitrochlorobenzene was greater  
 215 than the two CV values of 37% and 27%, calculated from five EC3 values each, reported by  
 216 ICCVAM (1999). The CV of 52% for EC1.5 values for hexyl cinnamic aldehyde tested in  
 217 the LLNA: BrdU-ELISA was greater than the CV for EC3 values reported by ICCVAM  
 218 (1999). The CV of 54% for EC1.5 values for eugenol tested in the LLNA: BrdU-ELISA was  
 219 greater than the CV of 42% for EC3 values reported by ICCVAM (1999).

220 **Table F-8 Interlaboratory Reproducibility of the EC3 for Substances Tested in the**  
 221 **Traditional LLNA<sup>1</sup>**

Substance	Laboratory					CV (%)
	1	2	3	4	5	
2, 4-Dinitrochlorobenzene	0.3	0.5	0.6	0.9	0.6	37
	0.5	0.6	0.4	0.6	0.3	27
Hexyl cinnamic aldehyde	7.9	7.6	8.4	7.0	8.1	7
Isoeugenol	1.3	3.3	1.8	3.1	1.6	41
Eugenol	5.8	14.5	8.9	13.8	6.0	42
SLS	13.4	4.4	1.5	17.1	4.0	84

222 Abbreviations: CV = Coefficient of variation; EC3 = Estimated concentration needed to produce a  
 223 stimulation index of three; LLNA = Murine local lymph node assay; SLS = Sodium lauryl sulfate.  
 224 <sup>1</sup>From ICCVAM (1999).

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