

# MHLW to ISO Comparison

	Extraction/Sample Preparation		Methodology		TAT (DAYS)		Sample Requirements	
	MHLW	ISO	MHLW	ISO	MHLW	ISO	MHLW	ISO
<b>Cytotoxicity</b>	24 hrs in EMEM + 5% FBS	24 hrs in EMEM + 5% FBS	Dilute suspension of cells is concurrently plated with extract and incubated for 6-7 days.	Extract is added to cell monolayer and observed up to 72 hrs.	31	18	2.5g or 150cm <sup>2</sup>	1g, 30cm <sup>2</sup> or 15cm <sup>2</sup>
<b>Genotoxicology Bacterial Reverse Mutation Test</b>	Exhaustive extract in methanol/acetone with terminal evaporation	50°C / 72 hrs extract in saline & DMSO or PEG	Extracts combined with agar and bacteria. Plates incubated for 72 hrs and colonies enumerated.		Pretest + 35	28	10 grams (Pretest) Method 1: TBD Method 2: 1g	0.8g, 24cm <sup>2</sup> or 12 cm <sup>2</sup> (x2)
<b>Genotoxicology – Chromosome Aberration</b>	Exhaustive extract in methanol/acetone with terminal evaporation	50°C / 72 hrs extract in saline & DMSO or PEG	Extracts applied to a cell monolayer for up to 20 hrs. Cells harvested and prepared for microscopic examination of gross chromosomal damage.		Pretest + 69	62	10 grams (Pretest) Method 1: TBD Method 2: 2g	2g, 60cm <sup>2</sup> or 30cm <sup>2</sup> (x2)
<b>Genotoxicology – Mouse Micronucleus</b>	Exhaustive extract in methanol/acetone with terminal evaporation	50°C / 72 hrs extract in saline & sesame oil	The test article extract used to evaluate potential to induce micronuclei formation in bone marrow of CD-1 mice.		Pretest + 62	55	10 grams (Pretest) Method 1: TBD Method 2: 3g	4g, 120cm <sup>2</sup> or 60 cm <sup>2</sup> (x2)
<b>Genotoxicology – Mouse Lymphoma</b>	Exhaustive extract in methanol/acetone with terminal evaporation	50°C / 72 hrs extract in saline & DMSO or PEG	Extract evaluated for ability to induce forward mutations at thymidine kinase locus as assayed by colony growth of L5178Y mouse lymphoma cells.		Pretest + 46	39	10 grams (Pretest) Method 1: TBD Method 2: 2g	2g, 60cm <sup>2</sup> or 30cm <sup>2</sup> (x2)
<b>Hemolysis</b>	Extract exposure for 1, 2 & 4 hrs	Extraction in PBS: 37°C / 72 hrs; 50°C / 72 hrs 70°C / 24 hrs; 121°C / 1 hr	Extract placed in dilute solution of rabbit blood for 1, 2 & 4 hrs. Optical density is scanned.	Test article or extract exposed to dilute solution of rabbit blood for 3 hrs. Optical density is scanned.	32	21	8g, 240cm <sup>2</sup> or 120cm <sup>2</sup> (x3)	4g, 120cm <sup>2</sup> or 60cm <sup>2</sup> (x3)
<b>Complement Activation</b>	Direct contact with NHS for 1 hr. (Sponsor supplied comparison recommended.)		Test and control NHS tested via ELISA plate		25		4g, 120cm <sup>2</sup> or 60cm <sup>2</sup>	
<b>Irritation – Intracutaneous</b>	Extraction methods identical		3 rabbits dosed intracutaneously with each extract. Observations & scoring up to 72 hrs. MHLW requires site photos at dosing & scoring periods. Terminal weights recorded.		28		1.2g, 36cm <sup>2</sup> or 18cm <sup>2</sup> (x2)	
<b>Irritation – Primary Skin</b>	Standard extraction	One-inch-square pieces of test article	3 rabbits per extract, at least 2 extracts. Patched 24 hrs, scratched & unscratched skin. Site photos during scoring. Terminal weights recorded.	3 rabbits, patches applied for 4 or 24 hrs, removed and scored.	28		4g, 120cm <sup>2</sup> or 60cm <sup>2</sup> (x2)	2.5cm <sup>2</sup> x 2.5cm <sup>2</sup> (x7)
<b>Sensitization – Guinea Pig Maximization</b>	Exhaustive extract in methanol/acetone with terminal evaporation	50°C / 72 hrs extract in saline & sesame oil	Per extract: 11 test animals plus 6 negative & 6 positive controls run concurrently. Terminal weights recorded.	Per extract: 11 test animals plus 6 negative controls. Positive control validation with 11 animals run every 3 months.	Pretest + 54	47	10 grams (Pretest) Method 1: TBD Method 2: 20g	2g, 60cm <sup>2</sup> or 30cm <sup>2</sup> (x6)
<b>Sensitization – Murine Local Lymph Node</b>	Three options: • standard extraction • exhaustive extract in methanol/acetone • concentrations/dilutions prepared in saline or DMSO	50°C / 72 hrs extract in saline & DMSO or PEG	15 mice per extract or 25 mice for liquids or dissolved solids	15 mice per extract	32	Pretest + 32	Contact lab.	1g, 30cm <sup>2</sup> or 15cm <sup>2</sup> (x6)
<b>Implantation</b>	Preparation methods identical		3 rabbits, either sex. 4-6 test articles can be implanted. Photos at necropsy & pathology evaluation.	3 rabbits, either sex. 4-6 test articles can be implanted.	Same implant duration		~10mm x 3mm (x15)	
<b>Acute Systemic Toxicity</b>	50°C / 72 hrs or 121°C / 1 hr extract in saline & sesame oil	50°C / 72 hrs extract in saline & sesame oil	Necropsy required at study termination. Body weight loss indicative of toxicity if statistically significant between test and control.	Necropsy not required. Body weight loss indicative of toxicity if 3 or more lose greater than 10%.	27		1.6g, 48cm <sup>2</sup> or 24cm <sup>2</sup> (x2)	
<b>Subacute Toxicity (MHLW) Subacute/Subchronic Toxicity (ISO)</b>	Mentions only use of saline as extraction vehicle.	Saline and/or sesame oil can be used.	Rats only. IV dosing for 14 days. Dose of 20 mL/kg.	Mice or rats. Study length and number of doses not outlined. Daily dosing regimen.	105	14-Day: 82 28-Day: 98	20g, 600cm <sup>2</sup> or 300cm <sup>2</sup> (x28)	Mice: 1.6g, 48cm <sup>2</sup> or 24cm <sup>2</sup> Rats: 9g, 270cm <sup>2</sup> or 135cm <sup>2</sup> (x14 or x28)
<b>Pyrogen</b>	Extraction methods identical		Initial test method identical to ISO. 3 rabbits for each continued test based on summation of temperature rise for all animals (up to 2).	Initial test method identical to MHLW. 5 rabbits used for (only 1) continued test, based on ≥ 0.5°C temperature rise in any one rabbit.	26		30g, 900cm <sup>2</sup> or 450cm <sup>2</sup>	

Talk to us and find out how our biocompatibility testing services can work for you.

Contact a WuXi AppTec Account Manager at 651-675-2000 or email: [info@wuxiapptec.com](mailto:info@wuxiapptec.com)

