

FINAL REPORT

MEM ELUTION

PROCEDURE NO. STP0032 REV 03

LABORATORY NO. 439112.1 AMENDED

PREPARED FOR:

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LABORATORY NUMBER: PROCEDURE NUMBER:

SAMPLE SOURCE:

TYPE OF TEST:

SAMPLE IDENTIFICATION:

DEVIATIONS:

CELL LINE:

INCUBATION PERIOD:

METHOD OF SCORING:

AMOUNT TESTED/SAMPLE EXTRACT: SAMPLE RECEIVED DATE: LAB PHASE START DATE:

LAB PHASE COMPLETION DATE:

REPORT ISSUE DATE: STUDY COMPLETION DATE:

AMENDED REPORT ISSUE DATE:

439112.1 Amended STP0032 REV 03

American Biotech Labs

Solid, Gel

Refer to Table 1

None

Mouse Heteroploid Connective Tissue (L-929)

72 ± 3 hours at 37 ± 1°C Cytopathic Effect (0-4)

3 g / 15 mL 20 Aug 2008

21 Aug 2008 26 Aug 2008

27 Aug 2008

27 Aug 2008

03 Nov 2008

AMENDMENT JUSTIFICATION:

At the request of the sponsor, the final report was changed from short format to long format.

INTRODUCTION:

The MEM Elution test was designed to determine the cytotoxicity of extractable substances. An extract of the sample was added to cell monolayers and incubated. The cell monolayers were examined and scored based on the degree of cellular destruction.

ACCEPTANCE CRITERIA:

The United States Pharmacopeia & National Formulary states that the sample meets the requirements if the reactivity grade is not greater than grade 2 or a mild reactivity. The AAMI/ISO 10993-5 standard states that the overall assessment of the results shall be made by capable persons based upon the data and results. Nelson Laboratories acceptance criteria was based upon the negative and media controls receiving "0" reactivity grades and positive results receiving a 3-4 reactivity grades (moderate to severe).



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PROCEDURE:

The amount of test material extracted was based on ANSI/AAMI/ISO and USP surface area recommendations or weight (0.20 g/mL extract fluid for polymers and plastic). The sample was extracted for 24-25 hours at $37 \pm 1^{\circ}$ C in 1X Minimal Essential Media with 5% calf serum. Positive (Latex Natural Rubber) and negative (Polypropylene Pellets) controls were extracted and included in the assay. A blank of extraction media (media control) was also included in the assay.

Multiple well cell culture plates were seeded with a verified quantity of L-929 cells and incubated until 80-90% confluent. The cell culture media was removed from the plates. The test extracts were filtered and the appropriate amount of extract was added to each well on the cell culture plates. Each extract was tested on three wells of cells. The cells were incubated at $37 \pm 1^{\circ}$ C with $5 \pm 1\%$ CO₂ for 72 ± 3 hours.

The cell monolayers were examined microscopically. The wells were scored as to the degree of discernable morphological cytotoxicity on a relative scale of 0 to 4:

CONDITIONS OF ALL CULTURES	REACTIVITY	GRADE
No cell lysis, intracytoplasmic granules. Not more than 20% rounding, occasional lysed cells. Not more than 50% rounding, no extensive cell lysis. Not more than 70% rounding and lysed cells. Nearly complete cell destruction.	NONE SLIGHT MILD MODERATE SEVERE	0 1 2 3 4

The results from the three wells were averaged to give a final cytotoxicity score.

RESULTS:

The results are summarized in Table 1. The test is acceptable if all three of the negative control and medium control test wells have a score of 0 and all three of the positive control test wells have a score of 3 or higher.

The sample meets USP requirements if none of the cell culture exposed to the sample shows greater than a mild reactivity (grade 2).



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STATEMENT OF UNCERTAINTY:

If applicable, the statement of uncertainty is available to sponsors upon request.

Technical Reviewer

Bobbi Rushton-Castro

Study Director

Amended Report Date

jzw



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TABLE 1. Results

IDENTIFICATION	SCORE #1	SCORE #2	SCORE #3	AVERAGE
Negative Control Media Control Positive Control 429606 Lot #060908	0	0	0	0
	0	0	0	0
	4	4	4	4
	1	1	1	1



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