

MRL/MpJ

Stock No: 000486 | MRL

 Inbred Strain

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Also Known As: Murphy Roths Large, MRL

The MRL/MpJ mice are the parent and control strain for for MRL/MpJ-*Fas*^{*lpr*} (Stock Nos. 000485, 006825). Despite carrying the normal *Fas* gene, MRL/MpJ mice also exhibit autoimmune disorders, but symptoms are manifested much later in life compared to those the MRL/MpJ-*Fas*^{*lpr*} mice. As a strain developed as the control for MRL/MpJ-*Fas*^{*lpr*}, MRL/MpJ mice are useful in the study of their comparable defects and diseases, including systemic lupus erythematosus and Sjogren syndrome.

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GENETIC OVERVIEW

Genetic Background

Generation

[Contact Technical Support](#)
(2018-07-27 00:00:00)

VIEW GENETICS

RESEARCH APPLICATIONS

Internal/Organ Research
Immunology, Inflammation and Autoimmunity Research
Neurobiology Research
Sensorineural Research

[VIEW ALL RESEARCH APPLICATIONS](#)

BASE PRICE

Starting at:

\$100.93 Domestic price for female 3-week

[VIEW PRICE LIST](#)

Details

Detailed Description

The MRL/MpJ mice are large but docile to the point that males rarely fight. They are the parent and control strain for for MRL/MpJ-*Fas*^{lpr} (Stock Nos. 000485, 006825). Despite carrying the normal *Fas* gene, MRL/MpJ mice also exhibit autoimmune disorders, but symptoms are manifested much later in life compared to those the MRL/MpJ-*Fas*^{lpr} mice. Starting at about three months of age, levels of circulating immune complexes rise greatly in the MRL-*Fas*^{lpr} mouse but not in the wildtype control, MRL/MpJ. Also beginning at 3 months *Fas*^{lpr} mice exhibit very severe proliferative glomerulonephritis, whereas in the MRL/MpJ controls usually only mild glomerular lesions are detected. MRL/MpJ inbred female typically die at 73 weeks of age and males die at 93 weeks. This compares to a lifespan of 17 weeks in the female and 22 weeks for males in the mouse homozygous for *Fas*^{lpr}. See MRL/MpJ-*Fas*^{lpr} (Stock No. [000485](#)) for additional information. As a strain developed as the control for MRL/MpJ-*Fas*^{lpr}, MRL/MpJ mice are useful in the study of their comparable defects and diseases.

MRL/MpJ, and one of its ancestral strains LG/J, display heightened wound healing relative to a panel of other inbred strains. At 4 weeks post-injury, 2mm ear punch wounds heal to 0-0.4mm in MRL/MpJ mice but are still 1.2-1.6mm in C57BL/6 mice. At 15 days post-injury C57BL/6 show a maximal closure of 30% reduction in ear hole size while MRL show 85% reduction. The process of healing in MRL/MpJ mice is faster, more complete, showed increased swelling, angiogenesis, fibroblast migration, extracellular matrix deposition, and decreased scarring and fibrosis. Additionally, hair follicles and accompanying sebaceous glands regenerate to a much greater degree. The other ancestral strains of MRL/MpJ (C3H, C57BL/6, and AKR) do not display this enhanced healing. Bone marrow transplantation shows that the MRL/MpJ healing phenotype does not readily transfer with bone marrow and remains in the irradiated host tissues. Enhanced healing of cardiac wounds has also been reported in MRL/MpJ mice. In this model, a very high mitotic index (10-20%) is found, similar to that seen in non-mammalian tissue regeneration. Using F2 and backcross mapping of MRL/MpJ-*Fas*^{lpr} x B6 progeny McBrearty *et al.* identified multiple wound healing QTLs, *Heal2* and *Heal3*, on MRL/MpJ chromosome 13 in the region of D13Mit115 and D13Mit129 respectively; *Heal5* on MRL/MpJ chromosome 12 in the region of D12Mit233; *Heal1* on chromosome 8 of C57BL/6 in the region of D8Mit211; and a highly suggestive locus on MRL/MpJ chromosome 7 in the region of D7Mit220. In crosses between MRL/MpJ x SJL/J, Masinde *et al.* identified 10 QTL for wound healing, confirming and extending the

findings of McBrearty et al. Chromosomes 1, 3, 6, and 13 each have a single QTL with that on chromosome 13 being statistically suggestive but not significant, while chromosomes 4, 7, and 9 each have two statistically significant QTLs. (Clark et al., 1998; Leferovich et al., 2001; Kench et al., 1999; McBrearty et al., 1998; Masinde et al., 2001.)

Microarray analysis and SELDI ProteinChip analysis identified multiple genes and proteins that have varied expression in the ear punch wounds of MRL/MpJ-*Fas*^{lpr} versus C57BL/6. The changes in expression patterns suggest that in MRL/MpJ mice there is less of an inflammatory response and an earlier transition into tissue repair than is seen in C57BL/6. (Li et al., 2000 and 2001.)

Blankenhorn et al. found that MRL/MpJ females heal faster and more completely than males. Some *Heal* QTLs are sexually dimorphic with *Heal2*, 3, 7, 8, 10, and 11 having greater effects in males and *Heal4*, 5, and 9 having greater effects in females. Castration improves wound healing in MRL/MpJ males to nearly the degree seen in females, but ovariectomy does not improve the degree of healing seen in MRL/MpJ females. (Blankenhorn et al., 2003)

Relative to B10.D2nSnJ mice, MRL/MpJ mice have decreased Neutrophil accumulation in the bronchiolar lavage in response to LPS infusion, and tests using bone marrow chimeras reveal that the pulmonary inflammatory response transfers with bone marrow. Transforming growth factor beta 1 autologous induction is reduced in MRL/MpJ splenocytes while macrophages show a reduction in the transforming growth factor beta 1 induction of interleukin 1 beta and tumor necrosis factor alpha production but no significant reduction in transforming growth factor beta 1 production. (Kench et al., 1999.)

+ Development

+ Control Suggestions

+ Selected References

- Genetics

+ *I12^{m1}*

+ *Gpr84^{del}*

- Disease/Phenotype

+ Disease Terms

+ Research Areas By Phenotype

+ Mammalian Phenotype Terms by Genotype

+ Phenotype Information

Technical Support

C O N T A C T T E C H N I C A L S U P P O R T

Genotyping Protocols

End Point Analysis: [H2rs8249979 Alternate 1-EP](#)

[Genotyping resources and troubleshooting](#)

Inbred mouse strains are maintained through sibling (sister x brother) matings; no genotyping required.

Dietary Information

LabDiet® 5K52 formulation (6% fat)

Breeding Considerations

[This strain is a good breeder.](#)

Due to the heightened healing which occurs in mice with the MRL genetic background, ear punch is not a good method for individual mouse identification in this strain.

[Additional Breeding and Husbandry Support](#)

Mating System

Sibling x Sibling

Appearance

albino, unaffected

Related Genotype: *a/a Tyr^f/Tyr^f Fas⁺/Fas⁺*

Citation

When using the MRL mouse strain in a publication, please include JAX stock #000486 in your Materials and Methods section.

Animal Health Reports

[Facility Barrier Level Descriptions](#)

 [AX28 \(Maximum\)](#)

 [MP14 \(Maximum\)](#)

 [RB08 \(Maximum\)](#)

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4 weeks	Female	Not Applicable	\$100.93
	Male	Not Applicable	\$100.93
5 weeks	Female	Not Applicable	\$100.93
	Male	Not Applicable	\$100.93
6 weeks	Female	Not Applicable	\$104.38
	Male	Not Applicable	\$104.38
7 weeks	Female	Not Applicable	\$107.83
	Male	Not Applicable	\$107.83
8 weeks	Female	Not Applicable	\$111.28
	Male	Not Applicable	\$111.28
9 weeks	Female	Not Applicable	\$114.73
	Male	Not Applicable	\$114.73
10 weeks	Female	Not Applicable	\$118.18
	Male	Not Applicable	\$118.18
11 weeks	Female	Not Applicable	\$121.63
	Male	Not Applicable	\$121.63

RELATED PRODUCTS AND SERVICES		
Mouse ES Cells	MRL/MpJ mES cells	\$1095.00

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