Extracting Mouse Genetic Project raw phenotyping data

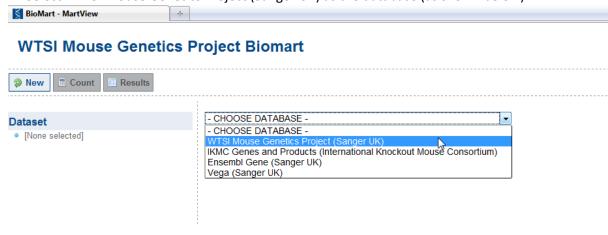
Introduction:

Raw phenotyping data is available through a publicly available BioMart¹ interface. This can be accessed at the following web address http://www.sanger.ac.uk/htgt/biomart. Mouse Genetics Project data which has passed all necessary internal quality control checks is automatically published to this BioMart within 24 hours. As of December 2011 this data set comprises approximately 9 x 10⁶ data points and hence it is not recommended that users attempt to extract all data in one bolus. Rather, the query made to the BioMart should be carefully phrased to extract only the data pertinent to answering your specific scientific question. Below is a step-wise example of how to extract the data using the standard BioMart user interface. Further information on using a BioMart can be found at: http://bowtie-bio.sourceforge.net/recount/biomaRt.pdf.

Demonstration Example:

The following example, demonstrates the extraction of data associated with the Mysm1 knockout line.

- 1. Open the BioMart by following the link: http://www.sanger.ac.uk/htgt/biomart.
- 2. Select "WTSI Mouse Genetics Project (Sanger UK) as the database (as shown below).

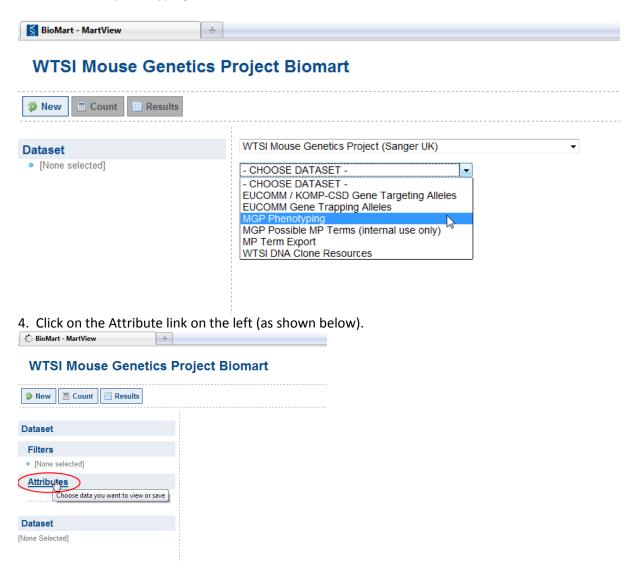


¹ BioMart is a generic data management system [Smedley et al "BioMart – biological queries made easy" (2009) BMC Genomics 10 (22)].





3. Select "MGP phenotyping" as the dataset (as shown below).





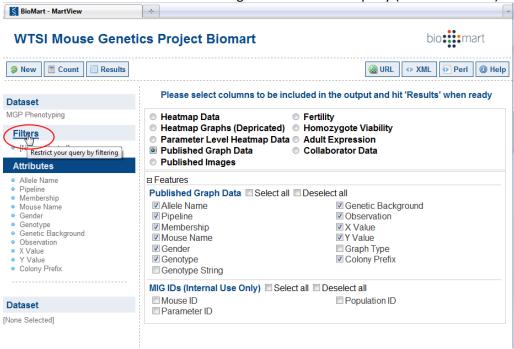


5. Select Published Graph Data (as shown below). S BioMart - MartView **WTSI Mouse Genetics Project Biomart** WILL <> XML New Count Perl Help Results Please select columns to be included in the output and hit 'Results' when ready **Dataset** MGP Phenotyping Fertility Meatmap Data Heatmap Graphs (Depricated)
Homozygote Viability **Filters** Parameter Level Heatmap Data Adult Expression [None selected] R Published Graph Data Collaborator Data ு Published Images **Attributes** Marker Symbol □ Features Allele Name Allele/Colony Details Select all Deselect all Colony Prefix Marker Symbol ■ Finalised Popuation Count Strain Adult LacZ Expression Significant Call Count Allele Name Auditory Brainstem Response Colony Prefix ■ Mouse Line Summary Body Composition (DEXA) Strain Brain Histopathology Citrobacter Challenge Phenotyping Tests Select all Deselect all Dysmorphology Adult LacZ Expression Micronuclei Embryo LacZ Expression Eye Histopathology Auditory Brainstem Response Modified SHIRPA Eye Morphology Body Composition (DEXA) ■ Non-Invasive Blood Pressure Fertility Brain Histopathology Open Field General Observations Citrobacter Challenge Peripheral Blood Lymphocytes Glucose Tolerance (ip) Plasma Chemistry Grip Strength Dysmorphology Haematology (CBC) ■ Embryo LacZ Expression Plasma Immunoglobulins Heart Histology Eye Histopathology Prepulse Inhibition Heart Weight Eye Morphology Recessive Lethal Study Hot Plate Fertility Rotarod Indirect Calorimetry MicroCT & Quantitative Faxitron General Observations ☑ Salmonella Challenge Micronuclei ☑ Glucose Tolerance (ip) Skin Histopathology Modified SHIRPA Grip Strength Stress Induced Hyperthermia Non-Invasive Blood Pressure ■ Haematology (CBC) ☑ Tail Epidermis Wholemount Open Field





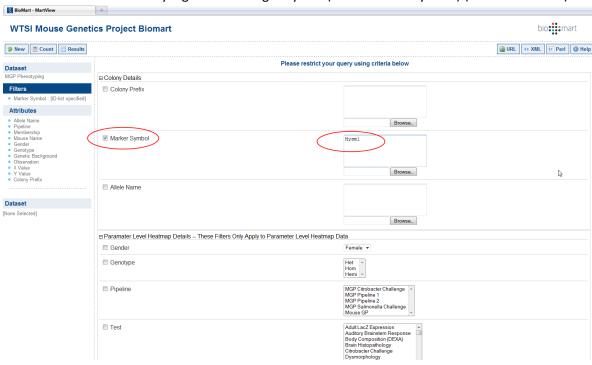
6. Now click on Filters to access a dialog box to restrict the query (as shown below).







7. We can filter the data by a gene name e.g. Mysm1 (a.k.a. Marker symbol) (as shown below).

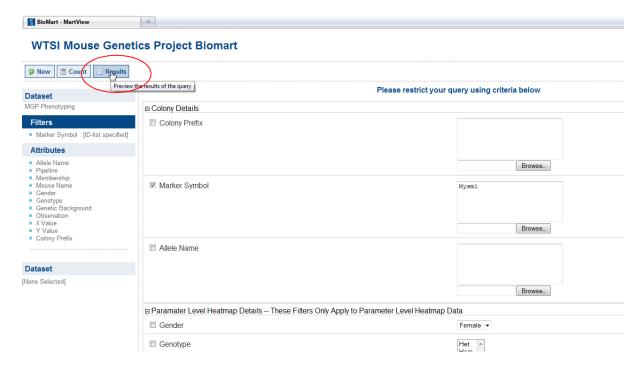






8. Click on the Results button (as shown below) to filter as specified.

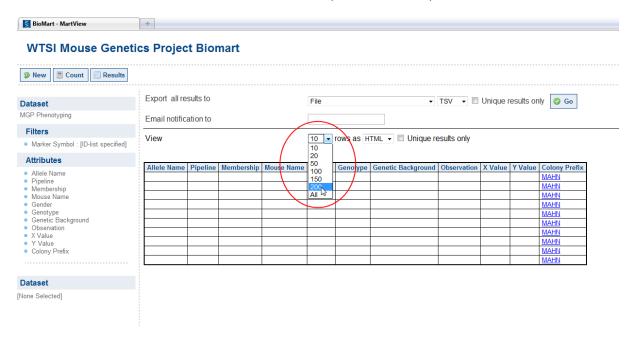
Note if the program is running for a long time and no results appear, it is likely you have asked for too big a slice of the data in one query. Please close and reconsidered your filters. A poorly formulated query can cause so much data to be sent to your computer that the web browser (and potentially your computer) will become overwhelmed and stop responding. It should also be noted that the volume of data available through this interface is roughly two orders of magnitude greater than supported by Excel. As such careful pre-planning of queries is advised.







9. Edit the number of rows shown to view the data (as shown below).







10. Scroll down the dataset. Empty rows can appear at the top if entries in the database have the correct colony prefix but no information to populate the remaining columns of the table. This is a limitation of the BioMart system and, while inelegant, does not present a challenge to the analysis of the data.

You may notice large numbers of digits appearing after a decimal place in the values displayed in X or Y value columns. These are a result of displaying the numerical values at the maximum available computer precision. NB. Software, such as Excel, also internally store values in this format but truncate them before display. To allow for the significant differences in the numerical precision of different assays we have not applied truncation. Users requiring improved formatting should extract the data (see step 11) and use the number formatting capabilities within, for example, excel to choose their desired reporting precision and format.

es	Allele Name	Pipeline	Membership	Mouse Name	Gender	Genotype	Genetic Background	Observation	X Value	Y Value	Co P
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											MAH
rship											MAH
Name											MAH
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Background											MAH
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	Mysm1 ^{tm1a(KOMP)Wtsi}	MGP Salmonella Challenge	Baseline	NTAC246.2a	Male	+/+	B6	Caecal Contents Count	20	1750	MAH
	Mysm1 ^{tm1a(KOMP)Wsi}	MGP Salmonella Challenge	Baseline	MACW61.2a	Male	+/+	B6	Caecal Contents Count	12	250	MAH
	Mysm1 ^{tm1a(KOMP)Wisi}	MGP Salmonella Challenge	Baseline	NTAC273.5a	Male	+/+	B6	Caecal Contents Count	7		MAH
	Mysm1 ^{tm1a(KOMP)Wtsi}	MGP Salmonella Challenge	Baseline	MADC46.1a	Male	+/+	B6	Caecal Contents Count	28	2250	MAH
	Mysm1 ^{tm1a(KOMP)Wtsi}	MGP Salmonella Challenge	Baseline	MAED34.2a	Male	+/+	B6	Caecal Contents Count	14	42500	MAH
	Mysm1 ^{tm1a(KOMP)Wtsi}	MGP Salmonella Challenge	Baseline	MADC60.1c	Male	+/+	B6	Caecal Contents Count	28		MAH





11. Results can be exported using the export facility (as shown below).

