

Finding Genetic Reagents

We have made changes to the **Alleles, Insertions, and Transgenic Constructs** section of our **Gene Reports** and to the **QuickSearch "GAL4 etc"** tab to make it easier for you to find alleles and transgenes which might be useful for your experiments. In both cases, we have added experimental tool information and have introduced "dynamic" tables which allow you to filter and sort these genetic reagents to find the ones with the particular characteristics that you need.

Gene Reports

NEW New "dynamic" tables in **Alleles, Insertions, and Transgenic Constructs** section

Download your list or send it to a FlyBase HitList

QuickSearch "GAL4 etc" tab

NEW Search has been expanded to include **ANY** transgenic construct/insertion that encodes an experimental tool, not just a limited set of drivers/reporters

Human Disease Associations on Gene Reports

Potential Disease Models based on Orthology

We have implemented a new automated pipeline to annotate *D. melanogaster* genes that may be relevant to human diseases based on their orthology to human "disease-related" genes.

- The pipeline uses three sets of data:
- **DIOPT** *D. melanogaster*-to-human orthology relationships
 - human gene-to-phenotype relationships within **OMIM**
 - **OMIM** phenotype cross-references within the **Disease Ontology (DO)**

The annotations are shown as **Potential Models Based on Orthology** in the **Human Disease Associations** section of **Gene Reports**

Disease Ontology (DO) Annotations			
Models Based on Experimental Evidence (7)			
Allele	Disease	Evidence	References
LamC ^{G489V} UAS	model of myopathy	CEA	(Dialynas et al., 2012)
LamC ^{N496I} UAS	model of muscular dystrophy	CEA	(Dialynas et al., 2015)
LamC ^{V528P} UAS	model of myopathy	CEA	(Dialynas et al., 2012)
LamC ^{M553R} UAS	model of muscular dystrophy	CEA	(Dialynas et al., 2015)
LamC ^{ΔN} UAS.c8a	model of muscular dystrophy	CEA	(Dialynas et al., 2015)
LamC ^{E174K} UAS	model of progeria	CEA with LamC ^{rspecified}	(Li et al., 2016)
LamC ^{rspecified}	model of progeria	CEA with LamC ^{E174K} UAS	(Li et al., 2016)
Potential Models Based on Orthology (9)			
Human Ortholog	Disease	Evidence	References
LMNA; lamin A/C	model of dilated cardiomyopathy 1A	inferred from electronic annotation	(FlyBase, 2019-)
	model of progeria	inferred from electronic annotation	(FlyBase, 2019-)
	model of lethal restrictive dermopathy	inferred from electronic annotation	(FlyBase, 2019-)
	model of autosomal dominant Emery-Dreifuss muscular dystrophy 2	inferred from electronic annotation	(FlyBase, 2019-)
	model of familial partial lipodystrophy type 2	inferred from electronic annotation	(FlyBase, 2019-)
	model of autosomal recessive Emery-Dreifuss muscular dystrophy 3	inferred from electronic annotation	(FlyBase, 2019-)
	model of congenital muscular dystrophy due to LMNA mutation	inferred from electronic annotation	(FlyBase, 2019-)
	model of Charcot-Marie-Tooth disease type 2B1	inferred from electronic annotation	(FlyBase, 2019-)
LMNB1; lamin B1	model of adult-onset autosomal dominant demyelinating leukodystrophy	inferred from electronic annotation	(FlyBase, 2019-)

Disease Model Summary Ribbon

We have added a graphical **Summary Ribbon** to the **Human Disease Associations** section of **Gene Reports**:

- it summarises Disease Ontology data into broad categories
- the same categories are used on all gene pages, so it easy to compare between genes
- both experimentally determined models and *potential* models identified by our new automated pipeline are included in the ribbon

Finding out about new features

This poster highlights just two recent improvements to FlyBase. There are a number of ways you can keep up-to-date with the latest improvements and new features.

From the Navigation Bar at the top of each page:

- Sign up for our **Newsletter**
- Follow us on twitter **@FlyBaseDotOrg**
- Check out the **New in This Release** and **Release Notes** pages
- From the Home Page:
- Check out the **Commentaries**

Got Questions ?

Help menu on the Navigation bar