

SPECIAL ARTICLE

Predicting human disease mutations and identifying drug targets from mouse gene knockout phenotyping campaigns

Robert Brommage^{1,*}, David R. Powell¹ and Peter Vogel²

ABSTRACT

Two large-scale mouse gene knockout phenotyping campaigns have provided extensive data on the functions of thousands of mammalian genes. The ongoing International Mouse Phenotyping Consortium (IMPC), with the goal of examining all ~20,000 mouse genes, has examined 5115 genes since 2011, and phenotypic data from several analyses are available on the IMPC website (www.mousephenotype.org). Mutant mice having at least one human genetic disease-associated phenotype are available for 185 IMPC genes. Lexicon Pharmaceuticals' Genome5000™ campaign performed similar analyses between 2000 and the end of 2008 focusing on the druggable genome, including enzymes, receptors, transporters, channels and secreted proteins. Mutants (4654 genes, with 3762 viable adult homozygous lines) with therapeutically interesting phenotypes were studied extensively. Importantly, phenotypes for 29 Lexicon mouse gene knockouts were published prior to observations of similar phenotypes resulting from homologous mutations in human genetic disorders. Knockout mouse phenotypes for an additional 30 genes mimicked previously published human genetic disorders. Several of these models have helped develop effective treatments for human diseases. For example, studying *Tph1* knockout mice (lacking peripheral serotonin) aided the development of telotristat ethyl, an approved treatment for carcinoid syndrome. *Sglt1* (also known as *Slc5a1*) and *Sglt2* (also known as *Slc5a2*) knockout mice were employed to develop sotagliflozin, a dual SGLT1/SGLT2 inhibitor having success in clinical trials for diabetes. Clinical trials evaluating inhibitors of AAK1 (neuropathic pain) and SGLT1 (diabetes) are underway. The research community can take advantage of these unbiased analyses of gene function in mice, including the minimally studied 'ignorome' genes.

KEY WORDS: Knockout mice, Mouse models, Phenotyping, Phenomics, Translational medicine

Introduction

Understanding gene function can explain the disease phenotypes observed in carriers of common genetic variants and deleterious mutations. Great progress is being made, deciphering the functions of the ~20,000 human genes, but the actions of many genes remain

poorly understood. For example, the Undiagnosed Diseases Network and other DNA sequencing efforts can typically identify gene mutations for one-third of patients with unknown rare genetic diseases (Splinter et al., 2018). The genes, and their actions, responsible for the remaining patients remain unknown. Identifying the actions and biochemical pathways of disease genes provides insights for potential therapies. Although imperfect, mice are the best-established models for human disease (Justice and Dhillon, 2016; Perlman, 2016; Sundberg and Schofield, 2018; Nadeau and Auwerx, 2019). This article summarizes data from two large-scale mouse gene knockout phenotyping campaigns: the International Mouse Phenotyping Consortium (IMPC) and Lexicon Pharmaceuticals' Genome5000™ program.

Both campaigns employed reverse genetics, the approach that relies on analyzing the phenotypes that result from the inactivation of specific genes to provide information on the physiological functions of these genes, to generate knockout mouse strains. Forward genetics approaches, involving the identification of the genes responsible for mouse phenotypes resulting from spontaneous mutations (Davisson et al., 2012) or chemical mutagenesis (Probst and Justice, 2010; Arnold et al., 2012; Sabrautzki et al., 2012; Potter et al., 2016; Wang et al., 2018), have also made major contributions to our understanding of genetic disease. Besides identifying inactivating gene mutations, forward genetics approaches often identify hypomorphic, gain-of-function and dominant-negative mutations. For example, The Jackson Laboratory (JAX) employed whole-exome sequencing to decipher spontaneous pathogenic mutations in 124 mouse strains (Fairfield et al., 2015; Palmer et al., 2016).


Mouse gene knockout phenotyping

Although examining mutant mice in individual laboratories has uncovered the functions of many genes, such piecemeal studies have several limitations. First, individual research groups often focus on the systems in which they have interest, hypotheses and experimental expertise. As a result, they can miss or ignore additional phenotypes. For example, a behavior laboratory can easily overlook concurrent immune disorders. Second, since research groups tend to individualize experimental techniques, comparisons among different laboratories can be difficult. Mouse strains, sex and age, along with assays and computational analyses, also vary. Third, there is a strong bias in the community to repeatedly study well-characterized genes, leaving thousands of genes, known as the 'ignorome' or the 'dark genome', unexplored (Edwards et al., 2011; Pandey et al., 2014; Oprea et al., 2018; Stoeger et al., 2018). The Mouse Genome Informatics database (Eppig, 2017) includes 13,924 genes with published mutant alleles in mice (data correct as of 19 February 2019), indicating that 6000 mouse genes remain unexplored and are therefore part of the ignorome.

Large-scale mutant mouse phenotyping campaigns that employ a panel of assays covering a wide range of phenotypes and apply

¹Department of Metabolism Research, Lexicon Pharmaceuticals, 8800 Technology Forest Place, The Woodlands, TX 77381, USA. ²St. Jude Children's Research Hospital, Pathology, MS 250, Room C5036A, 262 Danny Thomas Place, Memphis, TN 38105, USA.

*Author for correspondence (rbrommage@lexpharma.com)

 R.B., 0000-0002-9947-3822; D.R.P., 0000-0001-8072-388X; P.V., 0000-0002-7535-0545

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<https://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution and reproduction in any medium provided that the original work is properly attributed.

standardized experimental protocols and statistical analyses can address these limitations. The thousands of genes examined in these projects include both ignored and previously characterized genes. Technical staff generally have no knowledge of purported gene functions, which minimizes subconscious bias, and the large amounts of data collected from wild-type mice allow tracking of possible variations from normal values over time (Moore et al., 2018a).

Two large-scale mouse gene knockout phenotyping campaigns have been undertaken: Lexicon Pharmaceuticals' Genome5000™ campaign, designed to identify novel drug targets, and the International Mouse Phenotyping Consortium (IMPC), which aims to characterize mutant phenotypes for all ~20,000 mammalian genes. As summarized in Table 1, these two campaigns have many similarities but also differences.

Lexicon's high-throughput phenotyping analyses were performed between 2001 and the end of 2008, and included alliances with Bristol-Myers Squibb (Toyn et al., 2010; Kostich et al., 2016), Genentech (Tang et al., 2010) and Organon/Merck. The ongoing IMPC effort evolved from and includes data for 449 genes obtained during earlier European Mouse Disease Clinic (EUMODIC) and Sanger Mouse Genetics Program (MGP) mutant mouse phenotyping campaigns (Ayadi et al., 2012; White et al., 2013; de Angelis et al., 2015). Individual IMPC phenotyping centers select the genes they examine based on institutional investigator interests. Two focused mouse gene knockout phenotyping campaigns included examinations of 36 genes coding for glycan-binding proteins and glycosyltransferases (Orr et al., 2013) and 54 testes-expressed genes for male fertility (Miyata et al., 2016). In the early 2000s, Deltagen generated 750 mouse gene knockout lines using standard homologous recombination techniques (Moore, 2005) and phenotypic data are publicly available for 134 of these knockout lines (Table S1).

The IMPC effort utilizes murine embryonic stem (ES) cells generated by the International Knockout Mouse Consortium (IKMC) (Bradley et al., 2012; Brown et al., 2018). The IMPC phenotyping screen generally examines seven male and seven female mutant mice, with comparisons to phenotyping center-specific male and female historical control wild-type mice, which are shared among all genes examined (Meehan et al., 2017; Brown et al., 2018). An example of IMPC control data for body bone mineral density (BMD) is provided in Fig. S1. Lexicon's effort utilized ES cells generated by gene-trap mutagenesis using the

OmniBank® I library (Abuin et al., 2007; Hansen et al., 2008) or homologous recombination involving a λ -phage knockout shuttle system (Wattler et al., 1999). Phenotypes of *Adipor1*, *Angptl4*, *Ptprg*, *Rpn13* (also known as *Adrm1*) and *Tph1* mouse knockout lines generated independently via both ES cell technologies were identical. The Lexicon primary phenotyping screen generally examined four male and four female mutant mice, with comparisons to both littermate/cagemate and historical control wild-type mice. The parents of the mutant mice examined initially were subsequently mated a second time to provide a second cohort of mice for possible replication studies. The primary screen clearly identified dramatic phenotypes (*Alpk3*, *Brs3*, *Ksr2*, *Lrrk1*, *Mc4r* and *Sost*), with milder phenotypes confirmed or refuted with the second cohort. This approach follows the Bayesian statistical paradigm. If phenotypic replication was successful and the gene encoded a potential drug target, multiple additional cohorts of mutant mice were generated for sophisticated analyses. For example, more than 700 homozygous mutant mice were generated for *Aak1*, *Dagla*, *Ksr2*, *Ptprg*, *Sglt1* (also known as *Slc5a1*), *Sglt2* (also known as *Slc5a2*), *Stk4* and *Tph1* genes.

Both Lexicon and the IMPC employ similar phenotyping screens for audiology, behavior, blood cell counts, cardiology, body BMD and composition, immunology, metabolism, ophthalmology, radiology and serum chemistry. When gene knockout was lethal, yielding no adult homozygous mice, both campaigns examined mutant heterozygous mice. Beyond the common screening assays discussed above, Lexicon examined cortical and trabecular bone architecture by micro computed tomography (microCT) (Brommage et al., 2014), pain sensitivity by hot plate and formalin skin responses (Kostich et al., 2016), neuronal amyloid- β levels (Toyn et al., 2010) and comprehensive histopathology (Schofield et al., 2012). Metabolic responses to feeding a high-fat diet were analyzed in a second cohort (Brommage et al., 2008). Whereas IMPC extends the embryonic lethal analysis to time of death and high-throughput optical projection and microCT imaging (Dickinson et al., 2016), Lexicon did not examine the developmental abnormalities responsible for embryonic lethality.

The IMPC publishes detailed mutant mouse phenotype data. These publications include histopathology for 50 genes (Adissu et al., 2014); plasma metabolic profiling for 62 genes (Probert et al., 2015); skin, hair and nail abnormalities for 169 genes (Sundberg et al., 2017); developmental abnormalities for 401 embryonic-lethal knockout lines (Dickinson et al., 2016); skin data from 500+ genes

Table 1. Overview of the International Mouse Phenotyping Consortium (IMPC) and Lexicon Pharmaceuticals' Genome5000™ gene knockout mouse phenotyping campaigns

Campaign	IMPC	Lexicon Genome 5000™
Funding	Multiple government agencies	Publicly traded stock+corporate alliances (Bristol-Myers Squibb, Genentech, Organon/Merck)
Goals	Characterize all 20,000 genes	Identify novel drug targets
Locations	Phenotyping centers worldwide	Single facility in Texas, USA
Genes examined	Selected by phenotyping centers	Druggable genome
Mouse strain	C57BL/6N	C57BL/6J X 129SvEvBrd F2 hybrids
Gene knockouts	International Knockout Mouse Consortium (IKMC) ES cells	Homologous recombination (64%) and OmniBank® gene trap (36%)
Active	2007 start, ongoing (Bradley et al., 2012)	1998 through 2008 (high-throughput screen), 2014 (full phenotyping) (Zambrowicz et al., 1998)
Mutant mice examined	7 males and 7 females	4 males and 4 females+additional cohorts for interesting phenotypes
Control wild-type mice	Historical data for each phenotyping center	Littermates/cagemates+historical data
Mutant genes phenotyped	5057 (October 2018)	~4654
Mice/ES cells available	IMPC website (www.mousephenotype.org)	See 'Web resources for obtaining ES cells, mice and phenotypic data' section

(DiTommaso et al., 2014; Liakath-Ali et al., 2014); whole-mount LacZ reporter tissue expression profiles (Armit, 2015) in adult mice for 313 (West et al., 2015) and 424 (Tuck et al., 2015) genes; hearing data for 3006 genes (Bowl et al., 2017); metabolic phenotyping for 2016 genes (Rozman et al., 2018); and ophthalmic data for 4364 genes (Moore et al., 2018b). A manuscript summarizing IMPC bone data and relationships to human skeletal diseases is in preparation. The IMPC website (www.mousephenotype.org) provides comprehensive mutant mouse phenotype data in a readily searchable format (Koscielny et al., 2014). Updates of ongoing progress in IMPC mouse phenotyping continue, with Release 9.2 (5614 phenotyped genes) published in January 2019.

All high-throughput screens have false positives and false negatives (Karp et al., 2010) and ‘...it has never been easier to generate high-impact false positives than in the genomic era’ (MacArthur, 2012). The occurrence of false negatives can be estimated by the ability to identify the expected phenotypes arising from knockouts of benchmark genes, which are associated with well-established human and mouse mutant phenotypes. Examples of successful benchmark gene confirmation include *Brs3*, *Cnr1* and *Mcr4* in Lexicon’s obesity screen (Brommage et al., 2008), and *Crtap*, *Lrp5*, *Ostm1*, *Src* and *Sost* in Lexicon’s bone screen (Brommage et al., 2014). Conversely, researchers can detect false positives by phenotyping additional cohorts of mutant mice. The IMPC campaign provides data for the primary screen only, and statistical modeling calculations (Karp et al., 2010) estimate an 11.4% false-positive rate averaged among all IMPC phenotyping assays. Lexicon’s primary screen included fewer mice than that of the IMPC, and many false positives, subsequently identified with secondary screens, were observed.

Complete and variably penetrant lethality are common in gene knockout mice (Wilson et al., 2017). The IMPC defines subviable mutant lines as having fewer (<12.5% of the litter) than the expected 25% surviving homozygous mice resulting from heterozygous crosses (<http://www.mousephenotype.org/data/embryo>). The latest IMPC data for 4969 mutant lines show 24% preweaning lethality and 10% subviability. Lexicon observed ~16% preweaning lethality among 4654 mutant lines (Brommage et al., 2014).

Two IMPC phenotyping centers (Freudenthal et al., 2016; Dymont et al., 2016; Rowe et al., 2018) perform specialized skeletal analyses beyond the body BMD and radiology data obtained as part of the high-throughput screen (Table S1). Combined bone quantitative X-ray microradiography (Butterfield et al., 2019) and bone breaking strength data are available for 100 genes, with skeletal phenotypes observed for nine genes (Bassett et al., 2012). Gene knockout of the murine *Slc20a2* phosphate transporter (Beck-Cormier et al., 2019) results in skeletal defects and brain calcification, mimicking the homologous human genetic disease. Integration of IMPC mouse bone data and human genome-wide association study (GWAS) of heel bone BMD and fracture data from the UK Biobank identified variants in *GPC6* (Kemp et al., 2017) and *DAAM2* (Morris et al., 2019) as key determinants of skeletal health.

A summary of Lexicon’s phenotyping campaign (~4654 genes, with 3762 viable adult homozygous gene knockout lines undergoing bone phenotyping) was published in 2014 (Brommage et al., 2014). Published phenotypes involving multiple cohorts of knockout mice are available for 100 genes summarized below.

Modeling human Mendelian genetic disorders

Mutant mice contribute to our understanding of the mechanisms responsible for human genetic disorders. The IMPC performs an automated comparison of mutant mouse phenotypes to over 7000

rare human diseases in the Online Mendelian Inheritance in Man (OMIM) and Orphanet databases. The comprehensive 2017 update (Meehan et al., 2017) summarizes IMPC disease model discovery findings. Briefly, of the 3328 IMPC mouse genes examined, 621 had previous MGI mouse model annotations, with 385 genes (62%) having common observed phenotypes. Importantly, 90% (8984 of 9942) of the gene-phenotype annotations described by the IMPC had not previously been described in the literature. From the OMIM or Orphanet databases, 889 known rare disease-gene associations have an orthologous IMPC mouse mutant displaying at least one phenotype. These 889 associations involve 185 IMPC genes for which mutant mice showed at least one human disease-associated phenotype. Details on these data are available in supplementary tables 1-4 in Meehan et al. (2017). Updates to these analyses are provided within the ‘Human Diseases’ section of the IMPC website.

Lexicon published mouse knockout phenotypes for 100 genes (Fig. 1; Table S2) in both focused papers ($N=81$) and summaries ($N=19$) on obesity (Brommage et al., 2008) and bone phenotypes (Brommage et al., 2014), and the Genentech Secreted Protein Discovery Initiative (SPDI) gene alliance (Tang et al., 2010). Manual annotation of the PubMed database (www.ncbi.nlm.nih.gov/pubmed) identified human Mendelian disease phenotypes for 66 of these 100 mouse genes, with the remaining 34 having no known associated human Mendelian genetic disorder. Table 2 lists 30 genes for which Lexicon’s mutant mouse data support previously identified human phenotypes. All 30 genes have an OMIM disease designation.

Importantly, 29 mutant mouse phenotypes mimicking human disease phenotypes were characterized and published prior to the identification of their orthologous human disease genes (Table 3). Eighteen of these 29 genes have an OMIM disease designation, and OMIM summaries for many of the remaining 11 genes are outdated. At the time of Lexicon’s mouse phenotypic analyses, most of these 29 genes were minimally studied ignormeg genes.

ADIPOR1 (Rice et al., 2015) and *HDAC4* (Rajan et al., 2009) are classified as variants of unknown significance in OMIM, as subsequent human studies did not confirm the initial disease phenotype-gene associations observed in humans (Zhang et al., 2016) and mice. *Hdac4* knockout mice are presently in the IMPC phenotyping queue. *Adipor1* mice showed abnormal retinal morphology in both Lexicon and IMPC screens. Diverging human and mouse phenotypes have been described for five genes [*PTPRG* (Zhang et al., 2012), *RRM2B* (Powell et al., 2005), and *SLC25A1*, *SLC30A5* and *SLC30A10* (Brommage et al., 2014)], which can result from incomplete human and/or mouse phenotypic evaluations (Table 4). For example, human *SLC30A5* mutations affecting a zinc transporter reduce human breast milk zinc content without other clinical observations (Kumar et al., 2015), whereas *Slc30a5* mutant mice have low bone mass (Inoue et al., 2002; Brommage et al., 2014), but mouse milk composition was not examined. In the IMPC campaign, *Slc30a10* mice are currently in the phenotyping queue, *Slc25a1* mice exhibited preweaning lethality and the other three genes have not been examined yet.

The 34 Lexicon mouse phenotypes described without corresponding published human Mendelian genetic disorders fall into several categories (Table 5). Mouse mutant phenotypes for *Slc6a4* [selective serotonin reuptake inhibitor (SSRI) drug target] and *Tph1* (carcinoid syndrome drug target) have been examined by independent laboratories, but not by the IMPC. Human GWAS data exist for *EPHA6*, *FADS1*, *KCNK16*, *NOTUM*, *TPH2* and *WNT16*, with the IMPC having examined *Epha6*, *Fads1*, *Notum* and *Wnt16* mutant mice and observing preweaning lethality in *Notum* and *Wnt16* knockout mice. Multiple studies indicate that *ATG4B*, *CLDN18*,

Flow chart categorizing 100 published Lexicon mouse gene knockout phenotypes

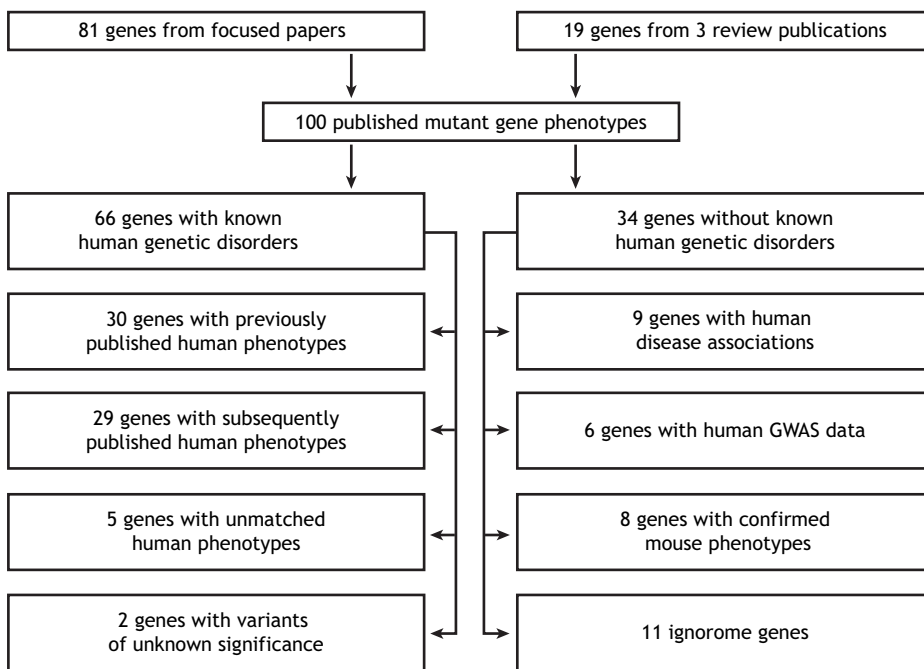


Fig. 1. Flow chart categorizing 100 published Lexicon mouse gene knockout phenotypes. We grouped these based on known or unknown human-mouse gene associations.

LIMK2, *MDM4*, *MKP1* (also known as *DUSP1*), *RPN13* and *Uchl5* are human oncogenes, with the IMPC having examined *Atg4b*, *Cldn18*, *Limk2*, *Mkp1* and *Uchl5* mutant mice and observing preweaning lethality in *Limk2* and *Uchl5* knockout animals. There is minimal published information for 11 ignorome genes (*Aak1*, *Ak8*, *Dpcd*, *Itfg2*, *Kif27*, *Kirrel1*, *Nme5*, *Tmem218*, *Tmub1*, *Tomm5* and *Till1*), and the IMPC examined only preweaning lethal *Till1* mutant mice. Independently published mouse knockout data exist for eight genes (*Brs3*, *Fam20b*, *Pik3c2a*, *Rock1*, *Rock2*, *Sh2d3c*, *Slc30a7* and *Spns2*), with the IMPC having examined *Pik3c2a*, *Rock1* and *Spns2* mutant mice, observing preweaning lethality in *Pik3c2a* and *Rock1* mutant mice.

Of the 100 Lexicon genes summarized in Fig. 1, embryonic lethality was observed in *Fam20b*, *Mdmx* and *Wnk1* knockout mice, perinatal lethality in *Kirrel1* knockout mice, and juvenile lethality in *Arl3*, *Cln7*, *Fgf23*, *Klotho*, *Npr2*, *Ostm1*, *Slc4a1* and *Sumf1* knockout mice. Subviability, defined as a deviation from the expected 1-2-1 Mendelian ratio of wild-type, heterozygous and homozygous mice from heterozygous crosses at $P < 0.001$ by Chi-squared testing, was observed in *Angptl4* (754, 17%), *Notum* (931, 19%), *Pkd11l1* (61, 10%), *Pomk* (395, 10%), *Rock1* (197, 15%), *Rock2* (227, 4%), *Rpn13* (39, 9%) and *Uchl5* (644, 14%) mice. Numbers in parentheses indicate observed numbers of wild-type mice and percentages of homozygous mutant mice, respectively.

Studies in mutant mice can also provide guidance for treating human genetic diseases. For example, Lexicon (Iwaniec et al., 2007) and others (Sawakami et al., 2006) showed that teriparatide treatment increases bone mass in *Lrp5* gene knockout mice with low bone mass. Similarly, teriparatide treatment increased BMD in a patient with osteoporosis pseudoglioma syndrome resulting from an inactivating *LRP5* mutation (Arantes et al., 2011).

IMPC – Lexicon comparisons

These two successful phenotyping campaigns had different objectives, funding and approaches to phenotypic screening

(Table 1), and comprehensive comparisons are beyond the scope of this article. Of the 100 Lexicon genes discussed here, 36 were also examined by the IMPC. Preweaning lethality was observed for 15 genes (*Fam20c*, *Fzd4*, *Limk2*, *Mboat7*, *Notum*, *Pik3c2a*, *Rock1*, *Sgpl1*, *Slc25a1*, *Slc46a1*, *Stk36*, *Sumf1*, *Till1*, *Uchl5* and *Wnt16*) in the IMPC, but not the Lexicon, phenotypic analyses. The OMIM autosomal-recessive disease genes *FAM20C*, *MBOAT7*, *SGPL1*, *SLC46A1* and *SUMF1* are not expected to exhibit disease phenotypes in the heterozygous mutant mice examined by the IMPC. Lexicon examined F2 hybrid C56BL/6J X 129SvEv-Brd mice, and hybrid vigor presumably contributed to better viability compared to the purebred C57BL/6N mice examined by the IMPC. The lower rate of lethality across all genes examined (~16% for Lexicon versus 25% for IMPC) is consistent with this hypothesis. Incomplete penetrance is common in human inherited diseases (Cooper et al., 2013) and variations in modifier genes likely contribute to this variable penetrance (Riordan and Nadeau, 2017).

Of the 36 genes examined in both phenotyping campaigns, 17 genes model human Mendelian disease. Both campaigns provided robust mouse data consistent with human genetic disorders involving mutations of *ALPK3*, *ANGPT4*, *DAGLA*, *DUOXA2*, *LRRK1* and *SLC24A5*. *Lrrk1* mice have the highest body volumetric BMD and BMD values in the Lexicon and IMPC screens, respectively (Fig. S1). Preweaning lethality and/or subviability of homozygous mice in the IMPC screen for *Fam20c*, *Fzd4*, *Grem2*, *Mboat7*, *Sgpl1*, *Slc25a1*, *Slc46a1* and *Sumf1* precluded the evaluation of homozygous knockout phenotypes for these genes. In contrast to observations by Lexicon, the IMPC did not observe soft tissue calcification in *Fam20a* mice, situs inversus in *Nme7* mice, nor any phenotypes in *Sgtl2* (only immunological parameters were examined) or *Slc30a8* mice. Human gene mutation phenotypes for *FAM20A* (enamel renal syndrome), *NME7* (situs inversus), *SGLT2* (familial renal glycosuria) and *SLC30A8* (resistance to Type 2 diabetes) are consistent with Lexicon's mutant mouse phenotypes.

Table 2. Lexicon mouse gene knockout phenotypes mimicking human genetic disorders

Gene ID	OMIM	Human disease	Knockout mouse phenotype	Lexicon publication	Human publication
<i>AGPAT2</i>	608594	lipodystrophy	lipodystrophy	Vogel et al., 2011	Agarwal et al., 2002
<i>ANGPTL3</i>	605019	hypotriglyceridemia	hypotriglyceridemia	Lee et al., 2009	Romeo et al., 2009
<i>CHSY1</i>	605282	brachydactyly	chondrodysplasia	Tang et al., 2010*	Li et al., 2010; Tian et al., 2010
<i>CLCN7</i>	607649	neurological defects	neurodegeneration	Rajan et al., 2011	Frattini et al., 2003
<i>CRTAP</i>	610682	osteogenesis imperfecta	low bone mass	Brommage et al., 2014	Morello et al., 2006
<i>DUOXA2</i>	274900	hypothyroidism	hypothyroidism	Brommage et al., 2014	Zamproni et al., 2008
<i>ENPP1</i>	208000	arterial calcification	calcinosis	Brommage et al., 2014	Rutsch et al., 2003
<i>FGF23</i>	193100	hypophosphatemic rickets	skeletal abnormalities	Brommage et al., 2014	ADHR Consortium, 2000
<i>FMR1</i>	300624	fragile X syndrome	learning, memory defects	Baker et al., 2010	1990s
<i>FZD4</i>	133780	exudative vitreoretinopathy	retinal angiogenesis	Paes et al., 2011	Robitaille et al., 2002
<i>GNPTAB</i>	252500	mucopolipidosis II	mucopolipidosis	Gelfman et al., 2007; Vogel et al., 2009	4 papers (2005)
<i>GNPTG</i>	252605	mucopolipidosis IIIC	mucopolipidosis	Vogel et al., 2009	Raas-Rothschild et al., 2000
<i>KL</i>	617994	tumoral calcinosis	skeletal abnormalities	Brommage et al., 2014	Ichikawa et al., 2007
<i>LRP5</i>	259770	osteoporosis-pseudoglioma	osteopenia	Iwaniec et al., 2007; Cui et al., 2011	Gong et al., 2001
<i>MC3R</i>	602025	obesity	obesity	Brommage et al., 2008	Lee et al., 2002; Feng et al., 2005
<i>MC4R</i>	601665	obesity	obesity	Brommage et al., 2008	Vaisse et al., 1998; Yeo et al., 1998
<i>MIP</i>	615274	cataracts	cataracts	Shiels et al., 2001	Berry et al., 2000; Francis et al., 2000
<i>MSTN</i>	614160	muscle hypertrophy	muscle hypertrophy	Brommage et al., 2014	Schuelke et al., 2004
<i>NPR2</i>	616255	acromesomelic dysplasia	achondroplastic dwarfism	Brommage et al., 2014	Bartels et al., 2004
<i>OSTM1</i>	259720	osteopetrosis	osteopetrosis	Brommage et al., 2014	Chalhoub et al., 2003
<i>SGLT1</i>	606824	glucose malabsorption	glucose malabsorption	Powell et al., 2013	Turk et al., 1991
<i>SGLT2</i>	233100	familial renal glycosuria	glucosuria	Powell et al., 2013	van den Heuvel et al., 2002
<i>SLC29A3</i>	602782	familial histiocytosis	histiocytosis	Tang et al., 2010 [‡]	Morgan et al., 2010
<i>SLC30A8</i>	125853	type 2 diabetes protection	insulin secretion	Syring et al., 2016	Flannick et al., 2014
<i>SLC39A13</i>	612350	spondylocheiro dysplasia	growth plate dysplasia	Brommage et al., 2014	Giunta et al., 2008
<i>SLC46A1</i>	229050	folate malabsorption	folate malabsorption	Salojin et al., 2011	Qiu et al., 2006
<i>SOST</i>	269500	sclerosteosis	sclerosteosis	Tang et al., 2010 [§]	Balemans et al., 2001; Brunkow et al., 2001
<i>STK4</i>	614868	immunodeficiency	immunodeficiency	Salojin et al., 2014	Abdollahpour et al., 2012; Nehme et al., 2012
<i>SUMF1</i>	272200	multiple sulfatase deficiency	lethality, cartilage defects	Brommage et al., 2014	Cosma et al., 2003
<i>WNK1</i>	614492	pseudohypoadosteronism	low blood pressure	Zambrowicz et al., 2003	Wilson et al., 2001

*Lexicon – Genentech 2005 SPDI gene alliance. Comprehensive analyses performed at Genentech (Wilson et al., 2012).

‡Lexicon – Genentech 2005 SPDI gene alliance. Comprehensive analyses performed at Genentech (Hsu et al., 2012).

§Lexicon – Genentech 2005 SPDI gene alliance.

Lexicon's extended phenotyping allowed characterization of human disease phenotypes not measured in the initial high-throughput screening assays. For example, human *DUOXA2* (Zamproni et al., 2008) and *SLC26A7* (Zou et al., 2018) mutations result in hypothyroidism, and Lexicon observed abnormal thyroid gland histology in both gene knockouts. Moreover, Lexicon's *Slc26a7* mice had reduced circulating thyroxine levels (Brommage et al., 2014).

Identifying novel drug targets

Studying human genetic disorders (Plenge et al., 2013; Nelson et al., 2015; Williams, 2016) in conjunction with knockout mice (Zambrowicz and Sands, 2003) can identify previously unknown tractable targets and lead to effective drugs. *PCKS9* is an example of this strategy, as knowledge that human inactivating mutations result in hypocholesterolemia led to the development of neutralizing antibodies to treat this condition (Jaworski et al., 2017). *Pcsk9* knockout mice are also hypocholesterolemic (Rashid et al., 2005) and blood cholesterol levels are halved in IMPC mice.

Unlike the genome-wide effort of the IMPC, Lexicon's choice of genes for knockout mouse analyses emphasized the druggable genome (Plewczynski and Rychlewski, 2009; Finan et al., 2017; Santos et al., 2017), which includes enzymes, receptors, ligands,

channels and secreted proteins. Ideally, drugs should influence disease processes without adversely affecting healthy tissues. In addition to identifying novel drug targets from beneficial mutant phenotypes, human genetic diseases and global gene knockout mice quickly identify or, preferably rule out, the possible adverse phenotypes that are likely to contribute to secondary drug target effects. For example, hypocholesterolemic subjects with inactivating *PCKS9* mutations and IMPC *Pcsk9* mutant mice have no unexpected health problems related to this mutation, suggesting that therapeutic inhibition of *PCKS9* activity should be safe. Generally, once this approach identifies novel drug targets, the preclinical drug development pipeline involves establishing robust enzymatic or binding assays, screening chemical libraries, optimization of chemical structures for potency and pharmacokinetic properties, followed by increasingly sophisticated animal pharmacology and toxicology studies. Thus, Lexicon stopped examining new mouse gene knockouts after December 2008 and stopped all basic research after January 2014 to focus on clinical development of small molecule drugs against selected targets previously identified in its gene knockout phenotyping campaign.

Lexicon's preclinical drug development program included the generation of neutralizing antibodies against *ANGPTL3* (Lee et al., 2009), *ANGPTL4* (Desai et al., 2007), *DKK1* (Brommage et al.,

Table 3. Lexicon mouse gene knockout phenotypes characterized prior to identification of human gene mutations

Gene ID	OMIM	Human disease	Knockout mouse phenotype	Lexicon publication	Human publication
<i>AK7</i>	617965	male infertility	male infertility	Vogel et al., 2012a	Lorès et al., 2018
<i>ALPK3</i>	618052	cardiomyopathy	cardiomyopathy	van Sligtenhorst et al., 2012	Phelan et al., 2016; Almomani et al., 2016
<i>ANGPTL4</i>	615881	hypotriglyceridemia	hypotriglyceridemia	Desai et al., 2007	Romeo et al., 2009
<i>ARL3</i>	(2013)	retinitis pigmentosa	retinitis pigmentosa	Schrick et al., 2006	Strom et al., 2016; Alkanderi et al., 2018
<i>CELSR2</i>	(2005)	Joubert syndrome	neurological defects	Vogel et al., 2012a	Vilboux et al., 2017a,b
<i>DAGLA</i>	(2011)	neurological phenotypes	neurological phenotypes	Powell et al., 2015	Smith et al., 2017
<i>ENTPD5</i>	(2011)	oncogene (mutation)	liver tumors	Read et al., 2009	MacCarthy and Notario, 2013; Bracco et al., 2014
<i>FAM20A</i>	204690	enamel renal syndrome	renal calcinosis	Vogel et al., 2012b	Jaureguiberry et al., 2012
<i>FAM20C</i>	259775	non-lethal Raine syndrome	hypophosphatemic rickets	Vogel et al., 2012b*	Whyte et al., 2017
<i>GPIHBP1</i>	615947	hypertriglyceridemia	hypertriglyceridemia	Beigneux et al., 2007 [‡] ; Sonnenburg et al., 2009 [‡]	Beigneux et al., 2009
<i>GREM2</i>	617275	tooth malformations	incisor tooth shape	Vogel et al., 2015a	Kantaputra et al., 2015
<i>KSR2</i>	(2011)	hyperphagic obesity	hyperphagic obesity	Revelli et al., 2011	Pearce et al., 2013
<i>LRRC4C</i>	(2008)	sensory processing	auditory response	Zhang et al., 2008	Maussion et al., 2017
<i>LRKK1</i>	615198	osteosclerotic metaphyseal dysplasia	osteopetrosis	Xing et al., 2013	Iida et al., 2016; Guo et al., 2017
<i>MBOAT7</i>	617188	intellectual disability	hydrocephalus	Vogel et al., 2012a	Johansen et al., 2016
<i>NME7</i>	(2018)	situs inversus	situs inversus	Vogel et al., 2010a, 2012a	Reish et al., 2016
<i>PAPPA2</i>	no gene	short stature	growth retardation	Conover et al., 2011	Dauber et al., 2016
<i>PKD1L1</i>	617205	situs inversus	situs inversus	Vogel et al., 2010a	Vetrini et al., 2016
<i>PLIN1</i>	613877	lipodystrophy	leanness	Brommage et al., 2008	Gandotra et al., 2011
<i>POMK</i>	615249	neurological deficits	cerebellar dysplasia	Vogel et al., 2012a	Di Costanzo et al., 2014; von Renesse et al., 2014
<i>SEMA4A</i>	610283	retinal degeneration	retinal degeneration	Rice et al., 2004	Abid et al., 2006
<i>SFRP4</i>	265900	Pyle's bone disease	cortical and trabecular bone phenotypes	Kiper et al., 2016	Kiper et al., 2016; Galada et al., 2017
<i>SGPL1</i>	617575	nephrotic syndrome with adrenal calcification	multiple immune phenotypes	Vogel et al., 2009	Janecke et al., 2017; Lovric et al., 2017; Prasad et al., 2017
<i>SLC10A7</i>	(2007)	skeletal dysplasia	osteochondrodysplasia	Brommage et al., 2014	Ashikov et al., 2018; Dubail et al., 2018
<i>SLC24A5</i>	113750	oculocutaneous albinism	oculocutaneous albinism	Vogel et al., 2008	Wei et al., 2013
<i>SLC26A7</i>	(2004)	hypothyroidism	hypothyroidism	Brommage et al., 2014	Zou et al., 2018
<i>STK36</i>	(2009)	ciliary dyskinesia	ciliary dyskinesia	Vogel et al., 2012a	Edelbusch et al., 2017
<i>TSPAN12</i>	613310	familial exudative vitreoretinopathy	retinal vasculature development	Junge et al., 2009 [‡]	Nikopoulos et al., 2010; Poulter et al., 2010
<i>ULK4</i>	615010	neurological deficits	neural development	Vogel et al., 2012a	Lang et al., 2014; Tassano et al., 2018

Underlined values in parentheses indicate the year of the latest OMIM update. The *PAPPA2* gene is not listed in the OMIM database.

*Lexicon – Organon/Merck 2005 alliance.

[‡]Lexicon – Genentech 2005 SPDI gene alliance (Tang et al., 2010).

2014), *FZD4* (Paes et al., 2011) and *NOTUM* (Brommage et al., 2019). Treating wild-type mice with each antibody successfully replicated the phenotypes observed in knockout mice. Subsequent work by Regeneron Pharmaceuticals demonstrated the efficacy of anti-*ANGPTL3* antibodies for the treatment of hypercholesterolemia in human patients (Dewey et al., 2017). In addition to providing phenotypic information, gene knockout mice provide two advantages in antibody generation and characterization. First, producing antibodies should, theoretically, be more efficient in specific gene knockout compared to wild-type mice, as the knockout mouse immune systems have never been exposed to the immunizing

proteins. Second, lack of antibody specificity is a major experimental problem, and the ‘... most stringent control for antibody specificity requires comparison of antibody reactivity in wild-type tissues or cells to reactivity in knockout animals...’ (Schonbrunn, 2014). Lexicon demonstrated the specificities of its anti-*ANGPTL3* and anti-*ANGPTL4* antibodies by showing lack of reactivity to tissues from the corresponding gene knockout mice.

In addition to antibodies, Lexicon developed small-molecule chemical inhibitors to 12 targets and information on these targets is provided in Table S3. Orally active inhibitors of *AAK1*, *SGLT1*, *SGLT2*, *SGPL1*, *SLC6A7* and *TPH1* entered human clinical trials.

Table 4. Genes with unmatched human and mouse mutant phenotypes

Gene ID	OMIM	Human disease	Knockout mouse phenotype	Lexicon publication	Human publication
<i>PTPRG</i>	(2004)	psychosis, schizophrenia	depressive behaviors	Zhang et al., 2012	Kranz et al., 2015; Kranz et al., 2016
<i>RRM2B</i>	612075	mitochondrial DNA depletion	renal failure	Powell et al., 2005	Bourdon et al., 2007; Kollberg et al., 2009
<i>SLC25A1</i>	615182	hydroxyglutaric aciduria	neonatal lethality	Brommage et al., 2014	Nota et al., 2013
<i>SLC30A5</i>	(2007)	reduced milk zinc content	severe osteopenia	Brommage et al., 2014	Kumar et al., 2015
<i>SLC30A10</i>	613280	hypermanganesemia	juvenile lethality	Brommage et al., 2014	Quadri et al., 2012; Tuschl et al., 2012

Underlined values in parentheses indicate the year of the latest OMIM update.

Table 5. Mouse phenotypes without known human Mendelian genetic disorders

Human gene	Function or pathway	IMPC mouse	Non-IMPC mouse	Lexicon phenotype	Human association	Lexicon publication
<i>AAK1</i>	protein kinase	yes	no	pain perception	ignorome	Kostich et al., 2016
<i>AK8</i>	adenylate kinase	no	no	hydrocephalus	ignorome	Vogel et al., 2012a
<i>ATG4B</i>	autophagy pathway	yes	2010	motor performance	oncogene	Read et al., 2011
<i>BRS3</i>	feeding behavior	no	1997	obesity	none	Brommage et al., 2008
<i>CLDN18</i>	osteoclast function	yes	2014	osteopenia	oncogene	Linares et al., 2012
<i>DPCD</i>	ciliary function	no	no	hydrocephalus	ignorome	Vogel et al., 2010a; Vogel et al., 2012a
<i>EPHA6</i>	ephrin receptor kinase	yes	2014	learning/memory	GWAS	Savelieva et al., 2008a
<i>FADS1</i>	fatty acid desaturase	yes	2012	lean/plaque resistance	GWAS	Powell et al., 2016
<i>FAM20B</i>	glycosaminoglycan synthesis	no	yes	lethality	none	Vogel et al., 2012b
<i>ITFG2</i>	integrin	yes	no	B-cell differentiation	ignorome	Al-Shami et al., 2013
<i>KCNK16</i>	potassium channel	yes	no	insulin secretion	GWAS	Vierra et al., 2015
<i>KIF27</i>	hedgehog signaling	no	no	hydrocephalus	ignorome	Vogel et al., 2012a
<i>KIRREL1</i>	podocyte formation	no	no	proteinuria	ignorome	Donoviel et al., 2001
<i>LIMK2</i>	actin dynamics	yes	2007	eyelid opening	oncogene	Rice et al., 2012
<i>MDM4</i>	p53 (also known as TP53) oncogene regulator	no	2001	fibroblast	oncogene	Finch et al., 2002
<i>MKP1</i>	MAPK phosphatase	yes	2006	immune responses	oncogene	Salojin et al., 2006
<i>NME5</i>	nucleoside diphosphate kinase	no	no	hydrocephalus high cortical	ignorome	Vogel et al., 2012a; Vogel et al., 2016
<i>NOTUM</i>	WNT inactivation	yes	2018	bone thickness; dental dysplasia	GWAS	Brommage et al., 2019; Nilsson et al., 2018
<i>PIK3C2A</i>	phosphoinositide kinase	yes	2014	glomerular structure	none	Harris et al., 2011
<i>ROCK1</i>	Rho-activated kinase	yes	2005	intraocular pressure	pleiotropy	Whitlock et al., 2009
<i>ROCK2</i>	Rho-activated kinase	no	2005	intraocular pressure	pleiotropy	Whitlock et al., 2009
<i>RPN13</i>	proteasome pathway	no	no	mouse pleiotropy	oncogene	Al-Shami et al., 2010a
<i>SH2D3C</i>	integrin pathway	no	2010	B-cell development	none	Al-Shami et al., 2010b
<i>SLC6A4</i>	serotonin transporter	no	1998	behavioral	SSRI drug target	Zhao et al., 2006
<i>SLC30A7</i>	zinc transporter	no	2007	insulin secretion	none	Syring et al., 2016
<i>SPNS2</i>	sphingosine-1-PO ₄ transporter	yes	2010	immune defects	none	Donoviel et al., 2015
<i>TMEM218</i>	membrane protein	no	no	cystic kidney disease	ignorome	Vogel et al., 2015b
<i>TMUB1</i>	CNS membrane protein	no	no	locomotor activity	ignorome	Zhang et al., 2010
<i>TOMM5</i>	mitochondrial translocase	no	no	pneumonia	ignorome	Vogel et al., 2015c
<i>TPH1</i>	peripheral serotonin synthesis	no	2003	gut motility syndrome	carcinoid	Savelieva et al., 2008b; Cui et al., 2011
<i>TPH2</i>	CNS serotonin synthesis	no	2004	behavior	GWAS	Savelieva et al., 2008b; Brommage et al., 2015
<i>TLL1</i>	tubulin ligase	yes	2010	male infertility	ignorome	Vogel et al., 2010b
<i>UCHL5</i>	proteasome pathway	yes	2015	mouse pleiotropy	oncogene	Al-Shami et al., 2010a
<i>WNT16</i>	WNT pathway	yes	2015	low cortical bone thickness	GWAS	Zheng et al., 2012; Wergedal et al., 2015

The Non-IMPC mouse column indicates the year of the first gene knockout publication. Gene expression data exist for many of the genes characterized as belonging to the ignorome.

Lexicon's peripheral serotonin synthesis inhibitors LX1031 and telotristat ethyl (both acting on tryptophan hydroxylase 1 encoded for by the *TPH1* gene) showed efficacy in subjects with irritable bowel syndrome (Brown et al., 2011) and carcinoid syndrome (Kulke et al., 2017), respectively. Telotristat ethyl was approved for the treatment of carcinoid syndrome in 2017. Neither drug crosses the blood-brain barrier to inhibit the neuronal TPH2 serotonin-synthesizing enzyme. Sotagliflozin, a dual SGLT1/SGLT2 glucose transport inhibitor, showed efficacy in Phase 3 trials for Type 1 diabetes (Garg et al., 2017) and in Phase 2 trials for Type 2 diabetes (Rosenstock et al., 2015), and is currently being developed, in collaboration with Sanofi, for both indications. Early clinical development is underway examining inhibitors of SGLT1 for Type 2 diabetes (Goodwin et al., 2017) and *AAK1* for neuropathic pain (Kostich et al., 2016).

Although drug development is not formally part of its mission or funding, the IMPC generates important knowledge for drug target identification and precision medicine initiatives (Lloyd et al., 2015).

Unlike Lexicon, the main goal of the IMPC is not drug development. However, we believe that its data and collaborative nature are an unmatched resource for future downstream work, both aimed at improving our fundamental understanding of mammalian gene function and at applying this knowledge to treatment of human genetic diseases.

Conclusions

The IMPC and Lexicon mouse gene knockout phenotyping campaigns provide key data for scientists studying mouse and human genomics. By continually updating its online database, the IMPC increasingly characterizes ignorome genes. The future success of the IMPC in identifying gene functions of significance to human health can be expected based on the results of Lexicon's successful mutant mouse phenotyping efforts. Lexicon's clinical drug development efforts, aiming for approval of SGLT1, SGLT2 and *AAK1* inhibitors, continue and their success should help patients with diabetes and neuropathic pain. We anticipate that future work

will develop additional drugs from Lexicon's knowledge base and, with adequate support, that of the IMPC. Both campaigns are expected to continue to contribute key mouse data for researchers studying ignomine genes associated with human genetic diseases.

Although this article focuses on published results, we stress that networking and presenting preliminary mouse data at conferences facilitates interactions with scientists working in human genomics and can contribute to collaborations ultimately resulting in publication of newly identified human genetic data. Successful examples of this process include *FAM20A*, *GREM2* (Kantaputra et al., 2015), *KSR2* (Pearce et al., 2013) and *SFRP4* (Kiper et al., 2016), and GWAS data for *WNT16* (Medina-Gomez et al., 2012; Zheng et al., 2012; Wergedal et al., 2015). Lexicon collaborated with academic scientists on many projects, and IMPC collaborations with academia and pharma should be encouraged. Recent publications involving IMPC mouse bone data and human data from the UK Biobank (Kemp et al., 2017; Morris et al., 2019) should stimulate additional collaborations in the future.

We encourage scientists to visit the IMPC website for further understanding of the actions of genes of interest. As Francis Collins, Director of the US National Institutes of Health, stated in 2006, 'A graduate student shouldn't spend a year making a knockout that's already been made. It's not a good use of resources' (Grimm, 2006). IMPC data showing either lethality, lack of a specific phenotype of interest, presence of this phenotype, and/or presence of additional phenotypes can guide research decisions for individual laboratories and optimize the use of limited resources.

Web resources for obtaining ES cells, mice and phenotypic data

The Mouse Genome Informatics (MGI) website (www.informatics.jax.org) is an excellent source of information on the availability of genetically modified mice. The IMPC website provides information on obtaining ES cells and cryopreserved sperm made available through the IKMC. The Monash University Embryonic Stem Cell (ES Cell)-to-Mouse Service group has published their experiences, from a 'client' perspective, using IKMC ES cells obtained worldwide (Cotton et al., 2015).

Information from individual IMPC phenotyping centers and the publicly available data from Deltagen and Lexicon are available in Table S1.

Note added in proof

Sotagliflozin has been approved within the European Union for use as an adjunct to insulin therapy to improve glycemic control in adults with Type 1 diabetes and a body mass index ≥ 27 kg/m², who could not achieve adequate glycemic control despite optimal insulin therapy.

Acknowledgements

The Lexicon and IMPC phenotyping campaigns involved contributions of hundreds of scientists, who are imperfectly captured in publications. R.B. worked for 18 months at the German Mouse Clinic (<https://www.mouseclinic.de>), an IMPC phenotyping center, and had many positive interactions with IMPC scientists. Special thanks to DMM Reviews Editor Julija Hmeljak for many helpful comments.

Competing interests

D.R.P. is currently employed at Lexicon Pharmaceuticals and has stock shares and stock options. R.B. and P.V. were previously employed at Lexicon Pharmaceuticals. R.B. owns Lexicon stock shares. P.V. has no financial interests.

Author contributions

All authors contributed equally to writing and reviewing the manuscript.

Funding

This work was supported by Lexicon Pharmaceuticals.

Supplementary information

Supplementary information available online at <http://dmm.biologists.org/lookup/doi/10.1242/dmm.038224.supplemental>

References

- Abdollahpour, H., Appaswamy, G., Kotlarz, D., Diestelhorst, J., Beier, R., Schäffer, A. A., Gertz, E. M., Schambach, A., Kreipe, H. H., Pfeifer, D. et al. (2012). The phenotype of human STK4 deficiency. *Blood* **119**, 3450-3457. doi:10.1182/blood-2011-09-378158
- Abid, A., Ismail, M., Mehdi, S. Q. and Khaliq, S. (2006). Identification of novel mutations in the SEMA4A gene associated with retinal degenerative diseases. *J. Med. Genet.* **43**, 378-381. doi:10.1136/jmg.2005.035055
- Abu, A., Hansen, G. M. and Zambrowicz, B. (2007). Gene trap mutagenesis. *Handb. Exp. Pharmacol.* **178**, 129-147. doi:10.1007/978-3-540-35109-2_6
- ADHR Consortium (2000). Autosomal dominant hypophosphataemic rickets is associated with mutations in FGF23. *Nat. Genet.* **26**, 345-348. doi:10.1038/81664
- Adisu, H. A., Estabel, J., Sunter, D., Tuck, E., Hooks, Y., Carragher, D. M., Clarke, K., Karp, N. A., Sanger Mouse Genetic Project, Newbigging, S. et al. (2014). Histopathology reveals correlative and unique phenotypes in a high-throughput mouse phenotyping screen. *Dis. Model. Mech.* **7**, 515-524. doi:10.1242/dmm.015263
- Agarwal, A. K., Arioglu, E., De Almeida, S., Akkoc, N., Taylor, S. I., Bowcock, A. M., Barnes, R. I. and Garg, A. (2002). AGPAT2 is mutated in congenital generalized lipodystrophy linked to chromosome 9q34. *Nat. Genet.* **31**, 21-23. doi:10.1038/ng880
- Al-Shami, A., Jhaver, K. G., Vogel, P., Wilkins, C., Humphries, J., Davis, J. J., Xu, N., Potter, D. G., Gerhardt, B., Mullinax, R. et al. (2010a). Regulators of the proteasome pathway, Uch37 and Rpn13, play distinct roles in mouse development. *PLoS ONE* **5**, e13654. doi:10.1371/journal.pone.0013654
- Al-Shami, A., Wilkins, C., Crisostomo, J., Seshasayee, D., Martin, F., Xu, N., Suwanichkul, A., Anderson, S. J. and Oravec, T. (2010b). The adaptor protein Sh2d3c is critical for marginal zone B cell development and function. *J. Immunol.* **185**, 327-334. doi:10.4049/jimmunol.1000096
- Al-Shami, A., Crisostomo, J., Wilkins, C., Xu, N., Humphries, J., Chang, W. C., Anderson, S. J. and Oravec, T. (2013). Integrin- α FG-GAP repeat-containing protein 2 is critical for normal B cell differentiation and controls disease development in a lupus model. *J. Immunol.* **191**, 3789-3798. doi:10.4049/jimmunol.1203534
- Alkanderi, S., Molinari, E., Shaheen, R., Elmaghloob, Y., Stephen, L. A., Sammut, V., Ramsbottom, S. A., Srivastava, S., Cairns, G., Edwards, N. et al. (2018). ARL3 mutations cause Joubert Syndrome by disrupting ciliary protein composition. *Am. J. Hum. Genet.* **103**, 612-620. doi:10.1016/j.ajhg.2018.08.015
- Almomani, R., Verhagen, J. M. A., Herkert, J. C., Brosens, E., van Spaendonck-Zwarts, K. Y., Asimaki, A., van der Zwaag, P. A., Frohn-Mulder, I. M. E., Bertoli-Avella, A. M., Boven, L. G. et al. (2016). Biallelic truncating mutations in ALPK3 cause severe pediatric cardiomyopathy. *J. Am. Coll. Cardiol.* **67**, 515-525. doi:10.1016/j.jacc.2015.10.093
- Arantes, H. P., Barros, E. R., Kunii, I., Bilezikian, J. P. and Lazaretti-Castro, M. (2011). Teriparatide increases bone mineral density in a man with osteoporosis pseudoglioma. *J. Bone Miner. Res.* **26**, 2823-2826. doi:10.1002/jbmr.530
- Armit, C. (2015). Into the blue: the importance of murine lacZ gene expression profiling in understanding and treating human disease. *Dis. Model. Mech.* **8**, 1341-1343. doi:10.1242/dmm.023606
- Arnold, C. N., Barnes, M. J., Berger, M., Blasius, A. L., Brandl, K., Croker, B., Crozat, K., Du, X., Eidenschien, C., Georgel, P. et al. (2012). ENU-induced phenovariance in mice: inferences from 587 mutations. *BMC Res Notes.* **5**, 577. doi:10.1186/1756-0500-5-577
- Ashikov, A., Abu Bakar, N., Wen, X.-Y., Niemeijer, M., Rodrigues Pinto Osorio, G., Brand-Arzamendi, K., Hasadsri, L., Hansikova, H., Raymond, K., Vicogne, D. et al. (2018). Integrating glycomics and genomics uncovers SLC10A7 as essential factor for bone mineralization by regulating post-Golgi protein transport and glycosylation. *Hum. Mol. Genet.* **27**, 3029-3045. doi:10.1093/hmg/ddy213
- Ayadi, A., Birling, M.-C., Bottomley, J., Bussell, J., Fuchs, H., Fray, M., Gailus-Durner, V., Greenaway, S., Houghton, R., Karp, N. et al. (2012). Mouse large-scale phenotyping initiatives: overview of the European Mouse Disease Clinic (EUMODIC) and of the Wellcome Trust Sanger Institute Mouse Genetics Project. *Mamm. Genome* **23**, 600-610. doi:10.1007/s00335-012-9418-y
- Baker, K. B., Wray, S. P., Ritter, R., Mason, S., Lanthorn, T. H. and Savelieva, K. V. (2010). Male and female Fmr1 knockout mice on C57 albino background exhibit spatial learning and memory impairments. *Genes Brain Behav.* **9**, 562-574. doi:10.1111/j.1601-183X.2010.00585.x
- Balemans, W., Ebeling, M., Patel, N., Van Hul, E., Olson, P., Dioszegi, M., Lacza, C., Wuyts, W., Van Den Ende, J., Willems, P. et al. (2001). Increased bone density in sclerosteosis is due to the deficiency of a novel secreted protein (SOST). *Hum. Mol. Genet.* **10**, 537-543. doi:10.1093/hmg/10.5.537
- Bartels, C. F., Bükülmez, H., Padayatti, P., Rhee, D. K., van Ravenswaaij-Arts, C., Pauli, R. M., Mundlos, S., Chitayat, D., Shih, L.-Y., Al-Gazali, L. I., et al. (2004). Mutations in the transmembrane natriuretic peptide receptor NPR-B

- impair skeletal growth and cause acromesomelic dysplasia, type Maroteaux. *Am. J. Hum. Genet.* **75**, 27-34. doi:10.1086/422013
- Bassett, J. H. D., Gogakos, A., White, J. K., Evans, H., Jacques, R. M., van der Spek, A. H., Ramirez-Solis, R., Ryder, E., Sunter, D., Boyde, A. et al.** (2012). Rapid-throughput skeletal phenotyping of 100 knockout mice identifies 9 new genes that determine bone strength. *PLoS Genet.* **8**, e1002858. doi:10.1371/journal.pgen.1002858
- Beck-Cormier, S., Lelliott, C. J., Logan, J. G., Lafont, D. T., Merametzjian, L., Leitch, V. D., Butterfield, N. C., Protheroe, H. J., Croucher, P. I., Baldock, P. A. et al.** (2019). Slc20a2, encoding the phosphate transporter PiT2, is an important genetic determinant of bone quality and strength. *J. Bone Miner. Res.* e3691. doi:10.1002/jbmr.3691
- Beigneux, A. P., Davies, B. S. J., Gin, P., Weinstein, M. M., Farber, E., Qiao, X., Peale, F., Bunting, S., Walzem, R. L., Wong, J. S. et al.** (2007). Glycosylphosphatidylinositol-anchored high-density lipoprotein-binding protein 1 plays a critical role in the lipolytic processing of chylomicrons. *Cell Metab.* **5**, 279-291. doi:10.1016/j.cmet.2007.02.002
- Beigneux, A. P., Franssen, R., Bensadoun, A., Gin, P., Melford, K., Peter, J., Walzem, R. L., Weinstein, M. M., Davies, B. S. J., Kuivenhoven, J. A. et al.** (2009). Chylomicronemia with a mutant GPIHBP1 (Q115P) that cannot bind lipoprotein lipase. *Arterioscler. Thromb. Vasc. Biol.* **29**, 956-962. doi:10.1161/ATVBAHA.109.186577
- Berry, V., Francis, P., Kaushal, S., Moore, A. and Bhattacharya, S.** (2000). Missense mutations in MIP underlie autosomal dominant 'polymorphic' and lamellar cataracts linked to 12q. *Nat. Genet.* **25**, 15-17. doi:10.1038/175538
- Bourdon, A., Minai, L., Serre, V., Jais, J.-P., Sarzi, E., Aubert, S., Chrétien, D., de Lonlay, P., Paquis-Flucklinger, V., Arakawa, H. et al.** (2007). Mutation of RRM2B, encoding p53-controlled ribonucleotide reductase (p53R2), causes severe mitochondrial DNA depletion. *Nat. Genet.* **39**, 776-780. doi:10.1038/ng2040
- Bowl, M. R., Simon, M. M., Ingham, N. J., Greenaway, S., Santos, L., Cater, H., Taylor, S., Mason, J., Kurbatova, N., Pearson, S. et al.** (2017). A large scale hearing loss screen reveals an extensive unexplored genetic landscape for auditory dysfunction. *Nat. Commun.* **8**, 886. doi:10.1038/s41467-017-00595-4
- Bracco, P. A., Bertoni, A. P. S. and Wink, M. R.** (2014). NTPDase5/PCPH as a new target in highly aggressive tumors: a systematic review. *Biomed. Res. Int.* **2014**, 123010. doi:10.1155/2014/123010
- Bradley, A., Anastassiadis, K., Ayadi, A., Battey, J. F., Bell, C., Birling, M.-C., Bottomley, J., Brown, S. D., Bürger, A., Bult, C. J. et al.** (2012). The mammalian gene function resource: the International Knockout Mouse Consortium. *Mamm. Genome* **23**, 580-586. doi:10.1007/s00335-012-9422-2
- Brommage, R., Desai, U., Revelli, J.-P., Donoviel, D. B., Fontenot, G. K., Dacosta, C. M., Smith, D. D., Kirkpatrick, L. L., Coker, K. J., Donoviel, M. S. et al.** (2008). High-throughput screening of mouse knockout lines identifies true lean and obese phenotypes. *Obesity* **16**, 2362-2367. doi:10.1038/oby.2008.361
- Brommage, R., Liu, J., Hansen, G. M., Kirkpatrick, L. L., Potter, D. G., Sands, A. T., Zambrowicz, B., Powell, D. R. and Vogel, P.** (2014). High-throughput screening of mouse gene knockouts identifies established and novel skeletal phenotypes. *Bone Res.* **2**, 14034. doi:10.1038/boneres.2014.34
- Brommage, R., Liu, J., Doree, D., Yu, W., Powell, D. R. and Yang, M. Q.** (2015). Adult Tph2 knockout mice without brain serotonin have moderately elevated spine trabecular bone but moderately low cortical bone thickness. *Bonekey Rep.* **4**, 718. doi:10.1038/bonekey.2015.87
- Brommage, R., Liu, J., Vogel, P., Mseeh, F., Thompson, A. Y., Potter, D. G., Shadoan, M. K., Hansen, G. M., Jeter-Jones, S., Cui, J. et al.** (2019). NOTUM inhibition increases endocortical bone formation and bone strength. *Bone Res.* **7**, 2. doi:10.1038/s41413-018-0038-3
- Brown, P. M., Drossman, D. A., Wood, A. J. J., Cline, G. A., Frazier, K. S., Jackson, J. I., Bronner, J., Freiman, J., Zambrowicz, B., Sands, A. et al.** (2011). The tryptophan hydroxylase inhibitor LX1031 shows clinical benefit in patients with nonconstipating irritable bowel syndrome. *Gastroenterology* **141**, 507-516. doi:10.1053/j.gastro.2011.05.005
- Brown, S. D. M., Holmes, C. C., Mallon, A.-M., Meehan, T. F., Smedley, D. and Wells, S.** (2018). High-throughput mouse phenomics for characterizing mammalian gene function. *Nat. Rev. Genet.* **19**, 357-370. doi:10.1038/s41576-018-0005-2
- Brunkow, M. E., Gardner, J. C., Van Ness, J., Paepfer, B. W., Kovacevich, B. R., Proll, S., Skonier, J. E., Zhao, L., Sabo, P. J., Fu, Y.-H. et al.** (2001). Bone dysplasia sclerosteosis results from loss of the SOST gene product, a novel cysteine knot-containing protein. *Am. J. Hum. Genet.* **68**, 577-589. doi:10.1086/318811
- Butterfield, N. C., Logan, J. G., Waung, J., Williams, G. R. and Bassett, J. H. D.** (2019). Quantitative X-Ray imaging of mouse bone by Faxitron. *Methods Mol. Biol.* **1914**, 559-569. doi:10.1007/978-1-4939-8997-3_30
- Chalhoub, N., Benachenhou, N., Rajapurohitam, V., Pata, M., Ferron, M., Frattini, A., Villa, A. and Vacher, J.** (2003). Grey-lethal mutation induces severe malignant autosomal recessive osteopetrosis in mouse and human. *Nat. Med.* **9**, 399-406. doi:10.1038/nm842
- Conover, C. A., Boldt, H. B., Bale, L. K., Clifton, K. B., Grell, J. A., Mader, J. R., Mason, E. J. and Powell, D. R.** (2011). Pregnancy-associated plasma protein-A2 (PAPP-A2): tissue expression and biological consequences of gene knockout in mice. *Endocrinology* **152**, 2837-2844. doi:10.1210/en.2011-0036
- Cooper, D. N., Krawczak, M., Polychronakos, C., Tyler-Smith, C. and Kehrer-Sawatzki, H.** (2013). Where genotype is not predictive of phenotype: towards an understanding of the molecular basis of reduced penetrance in human inherited disease. *Hum. Genet.* **132**, 1077-1130. doi:10.1007/s00439-013-1331-2
- Cosma, M. P., Pepe, S., Annunziata, I., Newbold, R. F., Grompe, M., Parenti, G. and Ballabio, A.** (2003). The multiple sulfatase deficiency gene encodes an essential and limiting factor for the activity of sulfatases. *Cell* **113**, 445-456. doi:10.1016/S0092-8674(03)00348-9
- Cotton, L. M., Meilak, M. L., Templeton, T., Gonzales, J. G., Nenci, A., Cooney, M., Truman, D., Rodda, F., Lynas, A., Viney, E. et al.** (2015). Utilising the resources of the International Knockout Mouse Consortium: the Australian experience. *Mamm. Genome* **26**, 142-153. doi:10.1007/s00335-015-9555-1
- Cui, Y., Niziolek, P. J., MacDonald, B. T., Zylstra, C. R., Alenina, N., Robinson, D. R., Zhong, Z., Matthes, S., Jacobsen, C. M., Conlon, R. A. et al.** (2011). Lrp5 functions in bone to regulate bone mass. *Nat. Med.* **17**, 684-691. doi:10.1038/nm.2388
- Dauber, A., Muñoz-Calvo, M. T., Barrios, V., Domené, H. M., Kloverpris, S., Serra-Juhé, C., Desikan, V., Pozo, J., Muzumdar, R., Martos-Moreno, G. Á. et al.** (2016). Mutations in pregnancy-associated plasma protein A2 cause short stature due to low IGF-I availability. *EMBO Mol. Med.* **8**, 363-374. doi:10.15252/emmm.201506106
- Davissou, M. T., Bergstrom, D. E., Reinholdt, L. G. and Donahue, L. R.** (2012). Discover genetics: the history and future of spontaneous mutation research. *Curr. Protoc. Mouse Biol.* **2**, 103-118. doi:10.1002/9780470942390.mo110200
- de Angelis, M. H., Nicholson, G., Selloum, M., White, J. K., Morgan, H., Ramirez-Solis, R., Sorg, T., Wells, S., Fuchs, H., Fray, M. et al.** (2015). Analysis of mammalian gene function through broad-based phenotypic screens across a consortium of mouse clinics. *Nat. Genet.* **47**, 969-978. doi:10.1038/ng.3360
- Desai, U., Lee, E.-C., Chung, K., Gao, C., Gay, J., Key, B., Hansen, G., Machajewski, D., Platt, K. A., Sands, A. T. et al.** (2007). Lipid-lowering effects of anti-angiopoietin-like 4 antibody recapitulate the lipid phenotype found in angiopoietin-like 4 knockout mice. *Proc. Natl. Acad. Sci. USA* **104**, 11766-11771. doi:10.1073/pnas.0705041104
- Dewey, F. E., Gusarova, V., Dunbar, R. L., O'Dushlaine, C., Schurmann, C., Gottesman, O., McCarthy, S., Van Hout, C. V., Bruse, S., Dansky, H. M. et al.** (2017). Genetic and pharmacologic inactivation of ANGPTL3 and cardiovascular disease. *N. Engl. J. Med.* **377**, 211-221. doi:10.1056/NEJMoa1612790
- Di Costanzo, S., Balasubramanian, A., Pond, H. L., Rozkalne, A., Pantaleoni, C., Saredi, S., Gupta, V. A., Sunu, C. M., Yu, T. W., Kang, P. B. et al.** (2014). POMK mutations disrupt muscle development leading to a spectrum of neuromuscular presentations. *Hum. Mol. Genet.* **23**, 5781-5792. doi:10.1093/hmg/ddu296
- Dickinson, M. E., Flenniken, A. M., Ji, X., Teboul, L., Wong, M. D., White, J. K., Meehan, T. F., Weninger, W. J., Westerberg, H., Adissu, H. et al.** (2016). High-throughput discovery of novel developmental phenotypes. *Nature* **537**, 508-514. doi:10.1038/nature19356
- DiTommaso, T., Jones, L. K., Cottle, D. L., Gerdin, A.-K., Vancollie, V. E., Watt, F. M., Ramirez-Solis, R., Bradley, A., Steel, K. P., Sundberg, J. P. et al.** (2014). Identification of genes important for cutaneous function revealed by a large scale reverse genetic screen in the mouse. *PLoS Genet.* **10**, e1004705. doi:10.1371/journal.pgen.1004705
- Donoviel, D. B., Freed, D. D., Vogel, H., Potter, D. G., Hawkins, E., Barrish, J. P., Mathur, B. N., Turner, C. A., Geske, R., Montgomery, C. A. et al.** (2001). Proteinuria and perinatal lethality in mice lacking NEPH1, a novel protein with homology to NEPHRIN. *Mol. Cell. Biol.* **21**, 4829-4836. doi:10.1128/MCB.21.14.4829-4836.2001
- Donoviel, M. S., Hait, N. C., Ramachandran, S., Maceyka, M., Takabe, K., Milstien, S., Oravec, T. and Spiegel, S.** (2015). Spinster 2, a sphingosine-1-phosphate transporter, plays a critical role in inflammation and autoimmune diseases. *FASEB J.* **29**, 5018-5028. doi:10.1096/fj.15-274936
- Dubail, J., Huber, C., Chantepie, S., Sonntag, S., Tüysüz, B., Mihci, E., Gordon, C. T., Steichen-Gersdorf, E., Amiel, J., Nur, B. et al.** (2018). SLC10A7 mutations cause a skeletal dysplasia with amelogenesis imperfecta mediated by GAG biosynthesis defects. *Nat. Commun.* **9**, 3087. doi:10.1038/s41467-018-05191-8
- Dyment, N. A., Jiang, X., Chen, L., Hong, S.-H., Adams, D. J., Ackert-Bicknell, C., Shin, D.-G. and Rowe, D. W.** (2016). High-throughput, multi-image cryohistology of mineralized tissues. *J. Vis. Exp.* e54468. doi:10.3791/54468
- Edelbusch, C., Cindrić, S., Dougherty, G. W., Loges, N. T., Olbrich, H., Rivlin, J., Wallmeier, J., Pennekamp, P., Amirav, I. and Omran, H.** (2017). Mutation of serine/threonine protein kinase 36 (STK36) causes primary ciliary dyskinesia with a central pair defect. *Hum. Mutat.* **38**, 964-969. doi:10.1002/humu.23261
- Edwards, A. M., Isserlin, R., Bader, G. D., Frye, S. V., Willson, T. M. and Yu, F. H.** (2011). Too many roads not taken. *Nature* **470**, 163-165. doi:10.1038/470163a
- Eppig, J. T.** (2017). Mouse Genome Informatics (MGI) Resource: genetic, genomic, and biological knowledgebase for the laboratory mouse. *ILAR J.* **58**, 17-41. doi:10.1093/ilar/ilx013
- Fairfield, H., Srivastava, A., Ananda, G., Liu, R., Kircher, M., Lakshminarayana, A., Harris, B. S., Karst, S. Y., Dionne, L. A., Kane, C. C. et al.** (2015). Exome

- sequencing reveals pathogenic mutations in 91 strains of mice with Mendelian disorders. *Genome Res.* **25**, 948-957. doi:10.1101/gr.186882.114
- Feng, N., Young, S. F., Aguilera, G., Puricelli, E., Adler-Wailles, D. C., Sebring, N. G. and Yanovski, J. A. (2005). Co-occurrence of two partially inactivating polymorphisms of MC3R is associated with pediatric-onset obesity. *Diabetes* **54**, 2663-2667. doi:10.2337/diabetes.54.9.2663
- Finan, C., Gaulton, A., Kruger, F. A., Lumbers, R. T., Shah, T., Engmann, J., Galver, L., Kelley, R., Karlsson, A., Santos, R. et al. (2017). The druggable genome and support for target identification and validation in drug development. *Sci. Transl. Med.* **9**, eaag1166. doi:10.1126/scitranslmed.aag1166
- Finch, R. A., Donoviel, D. B., Potter, D., Shi, M., Fan, A., Freed, D. D., Wang, C. Y., Zambrowicz, B. P., Ramirez-Solis, R., Sands, A. T. et al. (2002). Mdmx is a negative regulator of p53 activity in vivo. *Cancer Res.* **62**, 3221-3225.
- Flannick, J., Thorleifsson, G., Beer, N. L., Jacobs, S. B. R., Grarup, N., Burtt, N. P., Mahajan, A., Fuchsberger, C., Atzmun, G., Benediktsson, R. et al. (2014). Loss-of-function mutations in SLC30A8 protect against type 2 diabetes. *Nat. Genet.* **46**, 357-363. doi:10.1038/ng.2915
- Francis, P., Chung, J.-J., Yasui, M., Berry, V., Moore, A., Wyatt, M. K., Wistow, G., Bhattacharya, S. S. and Agre, P. (2000). Functional impairment of lens aquaporin in two families with dominantly inherited cataracts. *Hum. Mol. Genet.* **9**, 2329-2334. doi:10.1093/oxfordjournals.hmg.a018925
- Frattoni, A., Pangrazio, A., Susani, L., Sobacchi, C., Mirolo, M., Abinun, M., Andolina, M., Flanagan, A., Horwitz, E. M., Mihci, E. et al. (2003). Chloride channel CICN7 mutations are responsible for severe recessive, dominant, and intermediate osteopetrosis. *J. Bone Miner. Res.* **18**, 1740-1747. doi:10.1359/jbmr.2003.18.10.1740
- Freudenthal, B., Logan, J., Sanger Institute Mouse Pipelines, Croucher, P. I., Williams, G. R. and Bassett, J. H. D. (2016). Rapid phenotyping of knockout mice to identify genetic determinants of bone strength. *J. Endocrinol.* **231**, R31-R46. doi:10.1530/JOE-16-0258
- Galada, C., Shah, H., Shukla, A. and Girisha, K. M. (2017). A novel sequence variant in SFRP4 causing Pyle disease. *J. Hum. Genet.* **62**, 575-576. doi:10.1038/jhg.2016.166
- Gandotra, S., Le Dour, C., Bottomley, W., Cervera, P., Giral, P., Reznik, Y., Charpentier, G., Auclair, M., Delépine, M., Barroso, I. et al. (2011). Perilipin deficiency and autosomal dominant partial lipodystrophy. *N. Engl. J. Med.* **364**, 740-748. doi:10.1056/NEJMoa1007487
- Garg, S. K., Henry, R. R., Banks, P., Buse, J. B., Davies, M. J., Fulcher, G. R., Pozzilli, P., Gestly-Palmer, D., Lapuerta, P., Simó, R. et al. (2017). Effects of Sotagliflozin added to Insulin in patients with type 1 diabetes. *N. Engl. J. Med.* **377**, 2337-2348. doi:10.1056/NEJMoa1708337
- Gelfman, C. M., Vogel, P., Issa, T. M., Turner, C. A., Lee, W.-S., Kornfeld, S. and Rice, D. S. (2007). Mice lacking alpha/beta subunits of GlcNAc-1-phosphotransferase exhibit growth retardation, retinal degeneration, and secretory cell lesions. *Invest. Ophthalmol. Vis. Sci.* **48**, 5221-5228. doi:10.1167/iovs.07-0452
- Giunta, C., Elcioglu, N. H., Albrecht, B., Eich, G., Chambaz, C., Janecke, A. R., Yeowell, H., Weis, M. A., Eyre, D. R., Kraenzlin, M. et al. (2008). Spondylocheiro dysplastic form of the Ehlers-Danlos syndrome—an autosomal-recessive entity caused by mutations in the zinc transporter gene SLC39A13. *Am. J. Hum. Genet.* **82**, 1290-1305. doi:10.1016/j.ajhg.2008.05.001
- Gong, Y., Slee, R. B., Fukui, N., Rawadi, G., Roman-Roman, S., Reginato, A. M., Wang, H., Cundy, T., Glorieux, F. H., Lev, D. et al. (2001). LDL receptor-related protein 5 (LRP5) affects bone accrual and eye development. *Cell.* **107**, 513-523. doi:10.1016/S0092-8674(01)00571-2
- Goodwin, N. C., Ding, Z.-M., Harrison, B. A., Strobel, E. D., Harris, A. L., Smith, M., Thompson, A. Y., Xiong, W., Mseeh, F., Bruce, D. J. et al. (2017). Discovery of LX2761, a sodium-dependent glucose cotransporter 1 (SGLT1) inhibitor restricted to the intestinal lumen, for the treatment of diabetes. *J. Med. Chem.* **60**, 710-721. doi:10.1021/acs.jmedchem.6b01541
- Grimm, D. (2006). Mouse genetics. A mouse for every gene. *Science* **312**, 1862-1866. doi:10.1126/science.312.5782.1862
- Guo, L., Girisha, K. M., Iida, A., Hebbar, M., Shukla, A., Shah, H., Nishimura, G., Matsumoto, N., Nismath, S., Miyake, N. et al. (2017). Identification of a novel LRRK1 mutation in a family with osteosclerotic metaphyseal dysplasia. *J. Hum. Genet.* **62**, 437-441. doi:10.1038/jhg.2016.136
- Harris, D. P., Vogel, P., Wims, M., Moberg, K., Humphries, J., Jhaver, K. G., DaCosta, C. M., Shadoan, M. K., Xu, N., Hansen, G. M. et al. (2011). Requirement for class II phosphoinositide 3-kinase C2alpha in maintenance of glomerular structure and function. *Mol. Cell. Biol.* **31**, 63-80. doi:10.1128/MCB.00468-10
- Hsu, C.-L., Lin, W., Seshasayee, D., Chen, Y.-H., Ding, X., Lin, Z., Suto, E., Huang, Z., Lee, W. P., Park, H. et al. (2012). Equilibrative nucleoside transporter 3 deficiency perturbs lysosome function and macrophage homeostasis. *Science* **335**, 89-92. doi:10.1126/science.1213682
- Ichikawa, S., Imel, E. A., Kreiter, M. L., Yu, X., Mackenzie, D. S., Sorenson, A. H., Goetz, R., Mohammadi, M., White, K. E. and Econs, M. J. (2007). A homozygous missense mutation in human KLOTHO causes severe tumoral calcinosis. *J. Clin. Invest.* **117**, 2684-2691. doi:10.1172/JCI31330
- Iida, A., Xing, W., Docx, M. K. F., Nakashima, T., Wang, Z., Kimizuka, M., Van Hul, W., Rating, D., Spranger, J., Ohashi, H. et al. (2016). Identification of biallelic LRRK1 mutations in osteosclerotic metaphyseal dysplasia and evidence for locus heterogeneity. *J. Med. Genet.* **53**, 568-574. doi:10.1136/jmedgenet-2016-103756
- Inoue, K., Matsuda, K., Itoh, M., Kawaguchi, H., Tomoike, H., Aoyagi, T., Nagai, R., Hori, M., Nakamura, Y. and Tanaka, T. (2002). Osteopenia and male-specific sudden cardiac death in mice lacking a zinc transporter gene, Znt5. *Hum. Mol. Genet.* **11**, 1775-1784. doi:10.1093/hmg/11.15.1775
- Iwaniec, U. T., Wronski, T. J., Liu, J., Rivera, M. F., Arzaga, R. R., Hansen, G. and Brommage, R. (2007). PTH stimulates bone formation in mice deficient in Lrp5. *J. Bone Miner. Res.* **22**, 394-402. doi:10.1359/jbmr.061118
- Janecke, A. R., Xu, R., Steichen-Gersdorf, E., Waldegger, S., Entenmann, A., Giner, T., Krainer, I., Huber, L. A., Hess, M. W., Frisberg, Y. et al. (2017). Deficiency of the sphingosine-1-phosphate lyase SGPL1 is associated with congenital nephrotic syndrome and congenital adrenal calcifications. *Hum. Mutat.* **38**, 365-372. doi:10.1002/humu.23192
- Jaureguiberry, G., De la Dure-Molla, M., Parry, D., Quentric, M., Himmerkus, N., Koike, T., Poulter, J., Klootwijk, E., Robinette, S. L., Howie, A. J. et al. (2012). Nephrocalcinosis (enamel renal syndrome) caused by autosomal recessive FAM20A mutations. *Nephron Physiol.* **122**, 1-6. doi:10.1159/000349989
- Jaworski, K., Jankowski, P. and Kosior, D. A. (2017). PCSK9 inhibitors - from discovery of a single mutation to a groundbreaking therapy of lipid disorders in one decade. *Arch. Med. Sci.* **13**, 914-929. doi:10.5114/aoms.2017.65239
- Johansen, A., Rosti, R. O., Musaev, D., Sticca, E., Harripaul, R., Zaki, M., Çağlayan, A. O., Azam, M., Sultan, T., Froukh, T. et al. (2016). Mutations in MBOAT7, encoding lysophosphatidylinositol acyltransferase I, lead to intellectual disability accompanied by epilepsy and autistic features. *Am. J. Hum. Genet.* **99**, 912-916. doi:10.1016/j.ajhg.2016.07.019
- Junge, H. J., Yang, S., Burton, J. B., Paes, K., Shu, X., French, D. M., Costa, M., Rice, D. S. and Ye, W. (2009). TSPAN12 regulates retinal vascular development by promoting Norrin- but not Wnt-induced FZD4/beta-catenin signaling. *Cell* **139**, 299-311. doi:10.1016/j.cell.2009.07.048
- Justice, M. J. and Dhillon, P. (2016). Using the mouse to model human disease: increasing validity and reproducibility. *Dis. Model. Mech.* **9**, 101-103. doi:10.1242/dmm.024547
- Kantaputra, P. N., Kaewgahya, M., Hatsadaloi, A., Vogel, P., Kawasaki, K., Ohazama, A. and Ketudat Cairns, J. R. (2015). GREMLIN 2 mutations and dental anomalies. *J. Dent. Res.* **94**, 1646-1652. doi:10.1177/0022034515608168
- Karp, N. A., Baker, L. A., Gerdin, A.-K. B., Adams, N. C., Ramirez-Solis, R. and White, J. K. (2010). Optimising experimental design for high-throughput phenotyping in mice: a case study. *Mamm. Genome* **21**, 467-476. doi:10.1007/s00335-010-9279-1
- Kemp, J. P., Morris, J. A., Medina-Gomez, C., Forgetta, V., Warrington, N. M., Youtlen, S. E., Zheng, J., Gregson, C. L., Grundberg, E., Trajanoska, K. et al. (2017). Identification of 153 new loci associated with heel bone mineral density and functional involvement of GPC6 in osteoporosis. *Nat. Genet.* **49**, 1468-1475. doi:10.1038/ng.3949
- Kiper, P. O. S., Saito, H., Gori, F., Unger, S., Hesse, E., Yamana, K., Kiviranta, R., Solban, N., Liu, J., Brommage, R. et al. (2016). Cortical-bone fragility - insights from sFRP4 deficiency in Pyle's disease. *N. Engl. J. Med.* **374**, 2553-2562. doi:10.1056/NEJMoa1509342
- Kollberg, G., Darin, N., Benan, K., Moslemi, A.-R., Lindal, S., Tulinius, M., Oldfors, A. and Holme, E. (2009). A novel homozygous RRM2B missense mutation in association with severe mtDNA depletion. *Neuromuscul. Disord.* **19**, 147-150. doi:10.1016/j.nmd.2008.11.014
- Koscielny, G., Yaikhom, G., Iyer, V., Meehan, T. F., Morgan, H., Aienza-Herrero, J., Blake, A., Chen, C.-K., Easty, R., Di Fenza, A. et al. (2014). The International Mouse Phenotyping Consortium Web Portal, a unified point of access for knockout mice and related phenotyping data. *Nucleic Acids Res.* **42**, D802-D809. doi:10.1093/nar/gkt977
- Kostich, W., Hamman, B. D., Li, Y.-W., Naidu, S., Dandapani, K., Feng, J., Easton, A., Bourin, C., Baker, K., Allen, J. et al. (2016). Inhibition of AAK1 kinase as a novel therapeutic approach to treat neuropathic pain. *J. Pharmacol. Exp. Ther.* **358**, 371-386. doi:10.1124/jpet.116.235333
- Kranz, T. M., Harroch, S., Manor, O., Lichtenberg, P., Friedlander, Y., Seandel, M., Harkavy-Friedman, J., Walsh-Messinger, J., Dolgalev, I., Heguy, A. et al. (2015). De novo mutations from sporadic schizophrenia cases highlight important signaling genes in an independent sample. *Schizophr. Res.* **166**, 119-124. doi:10.1016/j.schres.2015.05.042
- Kranz, T. M., Berns, A., Shields, J., Rothman, K., Walsh-Messinger, J., Goetz, R. R., Chao, M. V. and Malaspina, D. (2016). Phenotypically distinct subtypes of psychosis accompany novel or rare variants in four different signaling genes. *EBioMedicine* **6**, 206-214. doi:10.1016/j.ebiom.2016.03.008
- Kulke, M. H., Hörsch, D., Caplin, M. E., Anthony, L. B., Bergsland, E., Öberg, K., Welin, S., Warner, R. R. P., Lombard-Bohas, C., Kunz, P. L. et al. (2017). Telotristat Ethyl, a tryptophan hydroxylase inhibitor for the treatment of carcinoid syndrome. *J. Clin. Oncol.* **35**, 14-23. doi:10.1200/JCO.2016.69.2780
- Kumar, L., Michalczyk, A., McKay, J., Ford, D., Kambe, T., Hudek, L., Varigos, G., Taylor, P. E. and Ackland, M. L. (2015). Altered expression of two zinc transporters, SLC30A5 and SLC30A6, underlies a mammary gland disorder of

- reduced zinc secretion into milk. *Genes Nutr.* **10**, 487. doi:10.1007/s12263-015-0487-x
- Lang, B., Pu, J., Hunter, I., Liu, M., Martin-Granados, C., Reilly, T. J., Gao, G.-D., Guan, Z.-L., Li, W.-D., Shi, Y.-Y. et al. (2014). Recurrent deletions of ULK4 in schizophrenia: a gene crucial for neurogenesis and neuronal motility. *J. Cell Sci.* **127**, 630-640. doi:10.1242/jcs.137604
- Lee, Y.-S., Poh, L. K.-S. and Loke, K.-Y. (2002). A novel melanocortin 3 receptor gene (MC3R) mutation associated with severe obesity. *J. Clin. Endocrinol. Metab.* **87**, 1423-1426. doi:10.1210/jcem.87.3.8461
- Lee, E.-C., Desai, U., Gololobov, G., Hong, S., Feng, X., Yu, X.-C., Gay, J., Wilganowski, N., Gao, C., Du, L.-L. et al. (2009). Identification of a new functional domain in angiopoietin-like 3 (ANGPTL3) and angiopoietin-like 4 (ANGPTL4) involved in binding and inhibition of lipoprotein lipase (LPL). *J. Biol. Chem.* **284**, 13735-13745. doi:10.1074/jbc.M807899200
- Li, Y., Laue, K., Temtamy, S., Aglan, M., Kotan, L. D., Yigit, G., Canan, H., Pawlik, B., Nürnberg, G., Wakeling, E. L. et al. (2010). Temtamy preaxial brachydactyly syndrome is caused by loss-of-function mutations in chondroitin synthase 1, a potential target of BMP signaling. *Am. J. Hum. Genet.* **87**, 757-767. doi:10.1016/j.ajhg.2010.10.003
- Liakath-Ali, K., Vancollie, V. E., Heath, E., Smedley, D. P., Estabel, J., Sunter, D., Ditommaso, T., White, J. K., Ramirez-Solis, R., Smyth, I. et al. (2014). Novel skin phenotypes revealed by a genome-wide mouse reverse genetic screen. *Nat. Commun.* **5**, 3540. doi:10.1038/ncomms4540
- Linares, G. R., Brommage, R., Powell, D. R., Xing, W., Chen, S.-T., Alshbool, F. Z., Lau, K.-H. W., Wergedal, J. E. and Mohan, S. (2012). Claudin 18 is a novel negative regulator of bone resorption and osteoclast differentiation. *J. Bone Miner. Res.* **27**, 1553-1565. doi:10.1002/jbmr.1600
- Lloyd, K. C. K., Meehan, T., Beaudet, A., Murray, S., Svenson, K., McKeirle, C., West, D., Morse, I., Parkinson, H., Brown, S. et al. (2015). Precision medicine: look to the mice. *Science* **349**, 390. doi:10.1126/science.349.6246.390-a
- Lorès, P., Coutton, C., El Khouri, E., Stouvenel, L., Givellet, M., Thomas, L., Rode, B., Schmitt, A., Louis, B., Sakheli, Z. et al. (2018). Homozygous missense mutation L673P in adenylate kinase 7 (AK7) leads to primary male infertility and multiple morphological anomalies of the flagella but not to primary ciliary dyskinesia. *Hum. Mol. Genet.* **27**, 1196-1211. doi:10.1093/hmg/ddy034
- Lovric, S., Goncalves, S., Gee, H. Y., Oskouian, B., Srinivas, H., Choi, W.-I., Shril, S., Ashraf, S., Tan, W., Rao, J. et al. (2017). Mutations in sphingosine-1-phosphate lyase cause nephrosis with ichthyosis and adrenal insufficiency. *J. Clin. Invest.* **127**, 912-928. doi:10.1172/JCI89626
- MacArthur, D. (2012). Methods: face up to false positives. *Nature* **487**, 427-428. doi:10.1038/487427a
- MacCarthy, C. M. and Notario, V. (2013). The ENTPD5/mt-PCPH oncoprotein is a catalytically inactive member of the ectonucleoside triphosphate diphosphohydrolase family. *Int. J. Oncol.* **43**, 1244-1252. doi:10.3892/ijco.2013.2052
- Maussion, G., Cruceanu, C., Rosenfeld, J. A., Bell, S. C., Jollant, F., Szatkiewicz, J., Collins, R. L., Hanscom, C., Kolobova, I., de Champfleury, N. M. et al. (2017). Implication of LRRC4C and DPP6 in neurodevelopmental disorders. *Am. J. Med. Genet. A* **173**, 395-406. doi:10.1002/ajmg.a.38021
- Medina-Gomez, C., Kemp, J. P., Estrada, K., Eriksson, J., Liu, J., Reppe, S., Evans, D. M., Heppe, D. H. M., Vandenput, L., Herrera, L. et al. (2012). Meta-analysis of genome-wide scans for total body BMD in children and adults reveals allelic heterogeneity and age-specific effects at the WNT16 locus. *PLoS Genet.* **8**, e1002718. doi:10.1371/journal.pgen.1002718
- Meehan, T. F., Conte, N., West, D. B., Jacobsen, J. O., Mason, J., Warren, J., Chen, C.-K., Tudose, I., Relac, M., Matthews, P. et al. (2017). Disease model discovery from 3,328 gene knockouts by The International Mouse Phenotyping Consortium. *Nat. Genet.* **49**, 1231-1238. doi:10.1038/ng.3901
- Miyata, H., Castaneda, J. M., Fujihara, Y., Yu, Z., Archambeault, D. R., Isotani, A., Kiyozumi, D., Kriseman, M. L., Mashiko, D., Matsumura, T. et al. (2016). Genome engineering uncovers 54 evolutionarily conserved and testis-enriched genes that are not required for male fertility in mice. *Proc. Natl. Acad. Sci. USA* **113**, 7704-7710. doi:10.1073/pnas.1608458113
- Moore, M. W. (2005). High-throughput gene knockouts and phenotyping in mice. *Ernst Schering Res. Found. Workshop* **50**, 27-44. doi:10.1007/s12263-015-0487-x
- Moore, B. A., Roux, M. J., Sebbag, L., Cooper, A., Edwards, S. G., Leonard, B. C., Imai, D. M., Griffey, S., Bower, L., Clary, D. et al. (2018a). A population study of common ocular abnormalities in C57BL/6N rd8 mice. *Invest. Ophthalmol. Vis. Sci.* **59**, 2252-2261. doi:10.1167/iovs.17-23513
- Moore, B. A., Leonard, B. C., Sebbag, L., Edwards, S. G., Cooper, A., Imai, D. M., Straiton, E., Santos, L., Reilly, C., Griffey, S. M. et al. (2018b). Identification of genes required for eye development by high-throughput screening of mouse knockouts. *Commun. Biol.* **1**, 236. doi:10.1038/s42003-018-0226-0
- Morello, R., Bertin, T. K., Chen, Y., Hicks, J., Tonachini, L., Monticone, M., Castagnola, P., Rauch, F., Glorieux, F. H., Vranka, J. et al. (2006). CRTAP is required for prolyl 3-hydroxylation and mutations cause recessive osteogenesis imperfecta. *Cell* **127**, 291-304. doi:10.1016/j.cell.2006.08.039
- Morgan, N. V., Morris, M. R., Cangul, H., Gleeson, D., Straatman-Iwanowska, A., Davies, N., Keenan, S., Pasha, S., Rahman, F., Gentle, D. et al. (2010). Mutations in SLC29A3, encoding an equilibrative nucleoside transporter ENT3, cause a familial histiocytosis syndrome (Faisalabad histiocytosis) and familial Rosai-Dorfman disease. *PLoS Genet.* **6**, e1000833. doi:10.1371/journal.pgen.1000833
- Morris, J. A., Kemp, J. P., Youlten, S. E., Laurent, L., Logan, J. G., Chai, R. C., Vulpesu, N. A., Forgetta, V., Kleinman, A., Mohanty, S. T. et al. (2019). An atlas of genetic influences on osteoporosis in humans and mice. *Nat. Genet.* **51**, 258-266. doi:10.1038/s41588-018-0302-x
- Nadeau, J. H. and Auwerx, J. (2019). The virtuous cycle of human genetics and mouse models in drug discovery. *Nat. Rev. Drug Discov.* **18**, 255-272. doi:10.1038/s41573-018-0009-9
- Nehme, N. T., Schmid, J. P., Debeurme, F., André-Schmutz, I., Lim, A., Nitschke, P., Rioux-Laucat, F., Lutz, P., Picard, C., Mahlaoui, N. et al. (2012). MST1 mutations in autosomal recessive primary immunodeficiency characterized by defective naive T-cell survival. *Blood* **119**, 3458-3468. doi:10.1182/blood-2011-09-378364
- Nelson, M. R., Tipney, H., Painter, J. L., Shen, J., Nicoletti, P., Shen, Y., Floratos, A., Sham, P. C., Li, M. J., Wang, J. et al. (2015). The support of human genetic evidence for approved drug indications. *Nat. Genet.* **47**, 856-860. doi:10.1038/ng.3314
- Nikopoulos, K., Gilissen, C., Hoischen, A., van Nouhuys, C. E., Boonstra, F. N., Blokland, E. A. W., Arts, P., Wieskamp, N., Strom, T. M., Ayuso, C. et al. (2010). Next-generation sequencing of a 40 Mb linkage interval reveals TSPAN12 mutations in patients with familial exudative vitreoretinopathy. *Am. J. Hum. Genet.* **86**, 240-247. doi:10.1016/j.ajhg.2009.12.016
- Nilsson, K., Mönväre-Skrčić, S., Henning, P., Funck-Brentano, T., Nethander, M., Rivadeneira, F., Koskela, A., Tuukkanen, J., Tuckermann, J., Perret, C. et al. (2018). Osteoblast-derived NOTUM reduces cortical bone mass in mice and the NOTUM locus is associated with bone mineral density in humans. *J. Bone Miner. Res.* **32** Suppl. 1. Available at: <http://www.asbmr.org/education/AbstractDetail?aid=addeafd3-36ee-4813-bc0f-fc8701deb7de>.
- Nota, B., Struys, E. A., Pop, A., Jansen, E. E., Fernandez Ojeda, M. R., Kanhai, W. A., Kranendijk, M., van Dooren, S. J., Bevova, M. R. et al. (2013). Deficiency in SLC25A1, encoding the mitochondrial citrate carrier, causes combined D-2- and L-2-hydroxyglutaric aciduria. *Am. J. Hum. Genet.* **92**, 627-631. doi:10.1016/j.ajhg.2013.03.009
- Oprea, T. I., Bologa, C. G., Brunak, S., Campbell, A., Gan, G. N., Gaulton, A., Gomez, S. M., Guha, R., Hersey, A., Holmes, J. et al. (2018). Unexplored therapeutic opportunities in the human genome. *Nat. Rev. Drug Discov.* **17**, 377. doi:10.1038/nrd.2018.52
- Orr, S. L., Le, D., Long, J. M., Sobieszczyk, P., Ma, B., Tian, H., Fang, X., Paulson, J. C., Marth, J. D. and Varki, N. (2013). A phenotype survey of 36 mutant mouse strains with gene-targeted defects in glycosyltransferases or glycan-binding proteins. *Glycobiology* **23**, 363-380. doi:10.1093/glycob/cws150
- Paes, K. T., Wang, E., Henze, K., Vogel, P., Read, R., Suwanichkul, A., Kirkpatrick, L. L., Potter, D., Newhouse, M. M. and Rice, D. S. (2011). Frizzled 4 is required for retinal angiogenesis and maintenance of the blood-retina barrier. *Invest. Ophthalmol. Vis. Sci.* **52**, 6452-6461. doi:10.1167/iovs.10-7146
- Palmer, K., Fairfield, H., Borgeia, S., Curtain, M., Hassan, M. G., Dionne, L., Yong Karst, S., Coombs, H., Bronson, R. T., Reinholdt, L. G. et al. (2016). Discovery and characterization of spontaneous mouse models of craniofacial dysmorphology. *Dev. Biol.* **415**, 216-227. doi:10.1016/j.ydbio.2015.07.023
- Pandey, A. K., Lu, L., Wang, X., Homayouni, R. and Williams, R. W. (2014). Functionally enigmatic genes: a case study of the brain ignorome. *PLoS ONE* **9**, e88889. doi:10.1371/journal.pone.0088889
- Pearce, L. R., Atanassova, N., Banton, M. C., Bottomley, B., van der Klaauw, A. A., Revelli, J.-P., Hendricks, A., Keogh, J. M., Henning, E., Doree, D. et al. (2013). KSR2 mutations are associated with obesity, insulin resistance, and impaired cellular fuel oxidation. *Cell* **155**, 765-777. doi:10.1016/j.cell.2013.09.058
- Perlman, R. L. (2016). Mouse models of human disease: an evolutionary perspective. *Evol. Med. Public Health* **2016**, 170-176. doi:10.1093/emph/eow014
- Phelan, D. G., Anderson, D. J., Howden, S. E., Wong, R. C. B., Hickey, P. F., Pope, K., Wilson, G. R., Pébay, A., Davis, A. M., Petrou, S. et al. (2016). ALPK3-deficient cardiomyocytes generated from patient-derived induced pluripotent stem cells and mutant human embryonic stem cells display abnormal calcium handling and establish that ALPK3 deficiency underlies familial cardiomyopathy. *Eur. Heart J.* **37**, 2586-2590. doi:10.1093/eurheartj/ehw160
- Plenge, R. M., Scolnick, E. M. and Altshuler, D. (2013). Validating therapeutic targets through human genetics. *Nat. Rev. Drug Discov.* **12**, 581-594. doi:10.1038/nrd4051
- Plewczynski, D. and Rychlewski, L. (2009). Meta-basic estimates the size of druggable human genome. *J. Mol. Model.* **15**, 695-699. doi:10.1007/s00894-008-0353-5
- Potter, P. K., Bowl, M. R., Jeyarajan, P., Wisby, L., Bleas, A., Goldsworthy, M. E., Simon, M. M., Greenaway, S., Michel, V., Barnard, A. et al. (2016). Novel gene function revealed by mouse mutagenesis screens for models of age-related disease. *Nat. Commun.* **7**, 12444. doi:10.1038/ncomms12444
- Poulter, J. A., Ali, M., Gilmour, D. F., Rice, A., Kondo, H., Hayashi, K., Mackey, D. A., Kearns, L. S., Ruddle, J. B., Craig, J. E. et al. (2010). Mutations in TSPAN12 cause autosomal-dominant familial exudative vitreoretinopathy. *Am. J. Hum. Genet.* **86**, 248-253. doi:10.1016/j.ajhg.2010.01.012

- Powell, D. R., Desai, U., Sparks, M. J., Hansen, G., Gay, J., Schrick, J., Shi, Z.-Z., Hicks, J. and Vogel, P. (2005). Rapid development of glomerular injury and renal failure in mice lacking p53R2. *Pediatr. Nephrol.* **20**, 432-440. doi:10.1007/s00467-004-1696-5
- Powell, D. R., DaCosta, C. M., Gay, J., Ding, Z.-M., Smith, M., Greer, J., Doree, D., Jeter-Jones, S., Mseeh, F., Rodriguez, L. A. et al. (2013). Improved glycaemic control in mice lacking SglT1 and SglT2. *Am. J. Physiol. Endocrinol. Metab.* **304**, E117-E130. doi:10.1152/ajpendo.00439.2012
- Powell, D. R., Gay, J. P., Wilganowski, N., Doree, D., Savelieva, K. V., Lanthorn, T. H., Read, R., Vogel, P., Hansen, G. M., Brommage, R. et al. (2015). Diacylglycerol lipase α knockout mice demonstrate metabolic and behavioral phenotypes similar to those of cannabinoid receptor 1 knockout mice. *Front. Endocrinol.* **6**, 86. doi:10.3389/fendo.2015.00086
- Powell, D. R., Gay, J. P., Smith, M., Wilganowski, N., Harris, A., Holland, A., Reyes, M., Kirkham, L., Kirkpatrick, L. L., Zambrowicz, B. et al. (2016). Fatty acid desaturase 1 knockout mice are lean with improved glycaemic control and decreased development of atheromatous plaque. *Diabetes Metab. Syndr. Obes.* **9**, 185-199. doi:10.2147/DMSO.S106653
- Prasad, R., Hadjidemetriou, I., Maharaj, A., Meimaridou, E., Buonocore, F., Saleem, M., Hurcombe, J., Bierzynska, A., Barbagelata, E., Bergadá, I. et al. (2017). Sphingosine-1-phosphate lyase mutations cause primary adrenal insufficiency and steroid-resistant nephrotic syndrome. *J. Clin. Invest.* **127**, 942-953. doi:10.1172/JCI90171
- Probert, F., Rice, P., Scudamore, C. L., Wells, S., Williams, R., Hough, T. A. and Cox, I. J. (2015). ^1H NMR metabolic profiling of plasma reveals additional phenotypes in knockout mouse models. *J. Proteome Res.* **14**, 2036-2045. doi:10.1021/pr501039k
- Probst, F. J. and Justice, M. J. (2010). Mouse mutagenesis with the chemical supermutagen ENU. *Methods Enzymol.* **477**, 297-312. doi:10.1016/S0076-6879(10)77015-4
- Qiu, A., Jansen, M., Sakaris, A., Min, S. H., Chattopadhyay, S., Tsai, E., Sandoval, C., Zhao, R., Akabas, M. H. and Goldman, I. D. (2006). Identification of an intestinal folate transporter and the molecular basis for hereditary folate malabsorption. *Cell* **127**, 917-928. doi:10.1016/j.cell.2006.09.041
- Quadri, M., Federico, A., Zhao, T., Breedveld, G. J., Battisti, C., Delnooz, C., Severijnen, L.-A., Di Toro Mammarella, L., Mignarri, A., Monti, L. et al. (2012). Mutations in SLC30A10 cause Parkinsonism and dystonia with hypermanganesemia, polycythemia, and chronic liver disease. *Am. J. Hum. Genet.* **90**, 467-477. doi:10.1016/j.ajhg.2012.01.017
- Raas-Rothschild, A., Cormier-Daire, V., Bao, M., Genin, E., Salomon, R., Brewer, K., Zeigler, M., Mandel, H., Toth, S., Roe, B. et al. (2000). Molecular basis of variant pseudo-Hurler polydystrophy (mucopolidosis IIIC). *J. Clin. Invest.* **105**, 673-681. doi:10.1172/JCI5826
- Rajan, I., Savelieva, K. V., Ye, G.-L., Wang, C.-Y., Malbari, M. M., Friddle, C., Lanthorn, T. H. and Zhang, W. (2009). Loss of the putative catalytic domain of HDAC4 leads to reduced thermal nociception and seizures while allowing normal bone development. *PLoS ONE* **4**, e6612. doi:10.1371/journal.pone.0006612
- Rajan, I., Read, R., Small, D. L., Perrard, J. and Vogel, P. (2011). An alternative splicing variant in *Clcn7*—/— mice prevents osteopetrosis but not neural and retinal degeneration. *Vet. Pathol.* **48**, 663-675. doi:10.1177/0300985810370164
- Rashid, S., Curtis, D. E., Garuti, R., Anderson, N. N., Bashmakov, Y., Ho, Y. K., Hamner, R. E., Moon, Y.-A. and Horton, J. D. (2005). Decreased plasma cholesterol and hypersensitivity to statins in mice lacking *Pcsk9*. *Proc. Natl. Acad. Sci. USA.* **102**, 5374-5379. doi:10.1073/pnas.0501652102
- Read, R., Hansen, G., Kramer, J., Finch, R., Li, L. and Vogel, P. (2009). Ectonucleoside triphosphate diphosphohydrolase type 5 (*Entpd5*)-deficient mice develop progressive hepatopathy, hepatocellular tumors, and spermatogenic arrest. *Vet. Pathol.* **46**, 491-504. doi:10.1354/vp.08-VP-0201-R-AM
- Read, R., Savelieva, K., Baker, K., Hansen, G. and Vogel, P. (2011). Histopathological and neurological features of *Atg4b* knockout mice. *Vet. Pathol.* **48**, 486-494. doi:10.1177/0300985810375810
- Reish, O., Aspit, L., Zouella, A., Roth, Y., Polak-Charcon, S., Baboushkin, T., Benyamini, L., Scheetz, T. E., Mussaffi, H., Sheffield, V. C. et al. (2016). A homozygous *Nme7* mutation is associated with situs inversus totalis. *Hum. Mutat.* **37**, 727-731. doi:10.1002/humu.22998
- Revelli, J.-P., Smith, D., Allen, J., Jeter-Jones, S., Shadoan, M. K., Desai, U., Schneider, M., van Slijtenhorst, I., Kirkpatrick, L., Platt, K. A. et al. (2011). Profound obesity secondary to hyperphagia in mice lacking kinase suppressor of Ras 2. *Obesity* **19**, 1010-1018. doi:10.1038/oby.2010.282
- Rice, D. S., Huang, W., Jones, H. A., Hansen, G., Ye, G.-L., Xu, N., Wilson, E. A., Troughton, K., Vaddi, K., Newton, R. C. et al. (2004). Severe retinal degeneration associated with disruption of semaphorin 4A. *Invest. Ophthalmol. Vis. Sci.* **45**, 2767-2777. doi:10.1167/iovs.04-0020
- Rice, D. S., Hansen, G. M., Liu, F., Crist, M. J., Newhouse, M. M., Potter, D., Xu, N., Abuin, A., Vogel, P. J. and Zambrowicz, B. P. (2012). Keratinocyte migration in the developing eyelid requires LIMK2. *PLoS ONE* **7**, e47168. doi:10.1371/journal.pone.0047168
- Rice, D. S., Calandria, J. M., Gordon, W. C., Jun, B., Zhou, Y., Gelfman, C. M., Li, S., Jin, M., Knott, E. J., Chang, B. et al. (2015). Adiponectin receptor 1 conserves docosahexaenoic acid and promotes photoreceptor cell survival. *Nat. Commun.* **6**, 6228. doi:10.1038/ncomms7228
- Riordan, J. D. and Nadeau, J. H. (2017). From peas to disease: modifier genes, network resilience, and the genetics of health. *Am. J. Hum. Genet.* **101**, 177-191. doi:10.1016/j.ajhg.2017.06.004
- Robitaille, J., MacDonald, M. L. E., Kaykas, A., Sheldahl, L. C., Zeisler, J., Dubé, M.-P., Zhang, L.-H., Singaraja, R. R., Guernsey, D. L., Zheng, B. et al. (2002). Mutant frizzled-4 disrupts retinal angiogenesis in familial exudative vitreoretinopathy. *Nat. Genet.* **32**, 326-330. doi:10.1038/ng957
- Romeo, S., Yin, W., Kozlitina, J., Pennacchio, L. A., Boerwinkle, E., Hobbs, H. H. and Cohen, J. C. (2009). Rare loss-of-function mutations in *ANGPTL* family members contribute to plasma triglyceride levels in humans. *J. Clin. Invest.* **119**, 70-79. doi:10.1172/JCI37118
- Rosenstock, J., Cefalu, W. T., Lapuerta, P., Zambrowicz, B., Ogbaa, I., Banks, P. and Sands, A. (2015). Greater dose-ranging effects on A1C levels than on glucosuria with LX4211, a dual inhibitor of SGLT1 and SGLT2, in patients with type 2 diabetes on metformin monotherapy. *Diabetes Care* **38**, 431-438. doi:10.2337/dc14-0890
- Rowe, D. W., Adams, D. J., Hong, S.-H., Zhang, C., Shin, D.-G., Renata Rydzik, C., Chen, L., Wu, Z., Garland, G., Godfrey, D. A. et al. (2018). Screening gene knockout mice for variation in bone mass: analysis by μCT and histomorphometry. *Curr. Osteoporos. Rep.* **16**, 77-94. doi:10.1007/s11914-018-0421-4
- Rozman, J., Rathkolb, B., Oestereicher, M. A., Schütt, C., Ravindranath, A. C., Leuchtenberger, S., Sharma, S., Kistler, M., Willershäuser, M., Brommage, R. et al. (2018). Identification of genetic elements in metabolism by high-throughput mouse phenotyping. *Nat. Commun.* **9**, 288. doi:10.1038/s41467-017-01995-2
- Rutsch, F., Ruf, N., Vaingankar, S., Toliat, M. R., Suk, A., Höhne, W., Schauer, G., Lehmann, M., Roscioli, T., Schnabel, D. et al. (2003). Mutations in *ENPP1* are associated with 'idiopathic' infantile arterial calcification. *Nat. Genet.* **34**, 379-381. doi:10.1038/ng1221
- Sabrautski, S., Rubio-Aliaga, I., Hans, W., Fuchs, H., Rathkolb, B., Calzada-Wack, J., Cohrs, C. M., Klafien, M., Seedorf, H., Eck, S. et al. (2012). New mouse models for metabolic bone diseases generated by genome-wide ENU mutagenesis. *Mamm. Genome* **23**, 416-430. doi:10.1007/s00335-012-9397-z
- Salojin, K. V., Owusu, I. B., Millerchip, K. A., Potter, M., Platt, K. A. and Oravec, T. (2006). Essential role of MAPK phosphatase-1 in the negative control of innate immune responses. *J. Immunol.* **176**, 1899-1907. doi:10.4049/jimmunol.176.3.1899
- Salojin, K. V., Cabrera, R. M., Sun, W., Chang, W. C., Lin, C., Duncan, L., Platt, K. A., Read, R., Vogel, P., Liu, Q. et al. (2011). A mouse model of hereditary folate malabsorption: deletion of the PCFT gene leads to systemic folate deficiency. *Blood* **117**, 4895-4904. doi:10.1182/blood-2010-04-279653
- Salojin, K. V., Hamman, B. D., Chang, W. C., Jhaver, K. G., Al-Shami, A., Crisostomo, J., Wilkins, C., Digeorge-Foushee, A. M., Allen, J., Patel, N. et al. (2014). Genetic deletion of *Mst1* alters T cell function and protects against autoimmunity. *PLoS ONE* **9**, e98151. doi:10.1371/journal.pone.0098151
- Santos, R., Ursu, O., Gaulton, A., Bento, A. P., Donadi, R. S., Bologa, C. G., Karlsson, A., Al-Lazikani, B., Hersey, A., Oprea, T. I. et al. (2017). A comprehensive map of molecular drug targets. *Nat. Rev. Drug Discov.* **16**, 19-34. doi:10.1038/nrd.2016.230
- Savelieva, K. V., Rajan, I., Baker, K. B., Vogel, P., Jarman, W., Allen, M. and Lanthorn, T. H. (2008a). Learning and memory impairment in Eph receptor A6 knockout mice. *Neurosci. Lett.* **438**, 205-209. doi:10.1016/j.neulet.2008.04.013
- Savelieva, K. V., Zhao, S., Pogorelov, V. M., Rajan, I., Yang, Q., Cullinan, E. and Lanthorn, T. H. (2008b). Genetic disruption of both tryptophan hydroxylase genes dramatically reduces serotonin and affects behavior in models sensitive to antidepressants. *PLoS ONE* **3**, e33301. doi:10.1371/journal.pone.0003301
- Sawakami, K., Robling, A. G., Ai, M., Pitner, N. D., Liu, D., Warden, S. J., Li, J., Maye, P., Rowe, D. W., Duncan, R. L. et al. (2006). The Wnt co-receptor LRP5 is essential for skeletal mechanotransduction but not for the anabolic bone response to parathyroid hormone treatment. *J. Biol. Chem.* **281**, 23698-23711. doi:10.1074/jbc.M601002000
- Schofield, P. N., Vogel, P., Gkoutos, G. V. and Sundberg, J. P. (2012). Exploring the elephant: histopathology in high-throughput phenotyping of mutant mice. *Dis. Model. Mech.* **5**, 19-25. doi:10.1242/dmm.008334
- Schonbrunn, A. (2014). Editorial: antibody can get it right: confronting problems of antibody specificity and irreproducibility. *Mol. Endocrinol.* **28**, 1403-1407. doi:10.1210/me.2014-1230
- Schrick, J. J., Vogel, P., Abuin, A., Hampton, B. and Rice, D. S. (2006). ADP-ribosylation factor-like 3 is involved in kidney and photoreceptor development. *Am. J. Pathol.* **168**, 1288-1298. doi:10.2353/ajpath.2006.050941
- Schuelke, M., Wagner, K. R., Stolz, L. E., Hübner, C., Riebel, T., Kömen, W., Braun, T., Tobin, J. F. and Lee, S.-J. (2004). Myostatin mutation associated with gross muscle hypertrophy in a child. *N. Engl. J. Med.* **350**, 2682-2688. doi:10.1056/NEJMoa040933
- Shiels, A., Bassnett, S., Varadaraj, K., Mathias, R., Al-Ghoul, K., Kuszak, J., Donoviel, D., Lilleberg, S., Friedrich, G. and Zambrowicz, B. (2001). Optical dysfunction of the crystalline lens in aquaporin-0-deficient mice. *Physiol. Genomics* **7**, 179-186. doi:10.1152/physiolgenomics.00078.2001

- Smith, D. R., Stanley, C. M., Foss, T., Boles, R. G. and McKernan, K. (2017). Rare genetic variants in the endocannabinoid system genes CNR1 and DAGLA are associated with neurological phenotypes in humans. *PLoS ONE* **12**, e0187926. doi:10.1371/journal.pone.0187926
- Sonnenburg, W. K., Yu, D., Lee, E. C., Xiong, W., Gololobov, G., Key, B., Gay, J., Wilganowski, N., Hu, Y., Zhao, S. et al. (2009). GPIHBP1 stabilizes lipoprotein lipase and prevents its inhibition by angiotensin-like 3 and angiotensin-like 4. *J. Lipid. Res.* **50**, 2421-2429. doi:10.1194/jlr.M900145-JLR200
- Splinter, K., Adams, D. R., Bacino, C. A., Bellen, H. J., Bernstein, J. A., Cheatele-Jarvela, A. M., Eng, C. M., Esteves, C., Gahl, W. A., Hamid, R. et al. (2018). Effect of genetic diagnosis on patients with previously undiagnosed disease. *N. Engl. J. Med.* **379**, 2131-2139. doi:10.1056/NEJMoa1714458
- Stoeger, T., Gerlach, M., Morimoto, R. I. and Nunes Amaral, L. A. (2018). Large-scale investigation of the reasons why potentially important genes are ignored. *PLoS Biol.* **16**, e2006643. doi:10.1371/journal.pbio.2006643
- Strom, S. P., Clark, M. J., Martinez, A., Garcia, S., Abelazeem, A. A., Matynia, A., Parikh, S., Sullivan, L. S., Bowne, S. J., Daiger, S. P. et al. (2016). De novo occurrence of a variant in ARL3 and apparent autosomal dominant transmission of retinitis pigmentosa. *PLoS ONE* **11**, e0150944. doi:10.1371/journal.pone.0150944
- Sundberg, J. P. and Schofield, P. N. (2018). Living inside the box: environmental effects on mouse models of human disease. *Dis. Model. Mech.* **11**, dmm035360. doi:10.1242/dmm.035360
- Sundberg, J. P., Dadras, S. S., Silva, K. A., Kennedy, V. E., Garland, G., Murray, S. A., Sundberg, B. A., Schofield, P. N. and Pratt, C. H. (2017). Systematic screening for skin, hair, and nail abnormalities in a large-scale knockout mouse program. *PLoS ONE* **12**, e0180682. doi:10.1371/journal.pone.0180682
- Syring, K. E., Boortz, K. A., Oeser, J. K., Ustione, A., Platt, K. A., Shadoan, M. K., McGuinness, O. P., Piston, D. W., Powell, D. R. and O'Brien, R. M. (2016). Combined deletion of Slc30a7 and Slc30a8 unmasks a critical role for ZnT8 in glucose-stimulated insulin secretion. *Endocrinology* **157**, 4534-4541. doi:10.1210/en.2016-1573
- Tang, T., Li, L., Tang, J., Li, Y., Lin, W. Y., Martin, F., Grant, D., Solloway, M., Parker, L., Ye, W. et al. (2010). A mouse knockout library for secreted and transmembrane proteins. *Nat. Biotechnol.* **28**, 749-755. doi:10.1038/nbt.1644
- Tassano, E., Uccella, S., Giacomini, T., Striano, P., Severino, M., Porta, S., Gimelli, G. and Ronchetto, P. (2018). Intragenic microdeletion of ULK4 and partial microduplication of BRWD3 in siblings with neuropsychiatric features and obesity. *Cytogenet Genome Res.* **156**, 14-21. [Epub ahead of print]. doi:10.1159/000491871
- Tian, J., Ling, L., Shboul, M., Lee, H., O'Connor, B., Merriman, B., Nelson, S. F., Cool, S., Ababneh, O. H., Al-Hadidy, A. et al. (2010). Loss of CHSY1, a secreted FRINGE enzyme, causes syndromic brachydactyly in humans via increased NOTCH signaling. *Am. J. Hum. Genet.* **87**, 768-778. doi:10.1016/j.ajhg.2010.11.005
- Toyn, J. H., Lin, X.-A., Thompson, M. W., Guss, V., Meredith, J. E., Jr, Sankaranarayanan, S., Barrezueta, N., Corradi, J., Majumdar, A., Small, D. L. et al. (2010). Viable mouse gene ablations that robustly alter brain A β levels are rare. *BMC Neurosci.* **11**, 143. doi:10.1186/1471-2202-11-143
- Tuck, E., Estabel, J., Oellrich, A., Maguire, A. K., Adissu, H. A., Souter, L., Siragher, E., Lillistone, C., Green, A. L., Wardle-Jones, H. et al. (2015). A gene expression resource generated by genome-wide lacZ profiling in the mouse. *Dis. Model. Mech.* **8**, 1467-1478. doi:10.1242/dmm.021238
- Turk, E., Zabel, B., Mundlos, S., Dyer, J. and Wright, E. M. (1991). Glucose/galactose malabsorption caused by a defect in the Na⁺/glucose cotransporter. *Nature* **350**, 354-356. doi:10.1038/350354a0
- Tuschl, K., Clayton, P. T., Gospe, S. M., Jr, Gulab, S., Ibrahim, S., Singhi, P., Aulakh, R., Ribeiro, R. T., Barsottini, O. G., Zaki, M. S. et al. (2012). Syndrome of hepatic cirrhosis, dystonia, polycythemia, and hypermanganesemia caused by mutations in SLC30A10, a manganese transporter in man. *Am. J. Hum. Genet.* **90**, 457-466. doi:10.1016/j.ajhg.2012.01.018
- Vaisse, C., Clement, K., Guy-Grand, B. and Froguel, P. (1998). A frameshift mutation in human MC4R is associated with a dominant form of obesity. *Nat. Genet.* **20**, 113-114. doi:10.1038/2407
- van den Heuvel, L. P., Assink, K., Willemsen, M. and Monnens, L. (2002). Autosomal recessive renal glucosuria attributable to a mutation in the sodium glucose cotransporter (SGLT2). *Hum. Genet.* **111**, 544-547. doi:10.1007/s00439-002-0820-5
- van Sligtenhorst, I., Ding, Z.-M., Shi, Z.-Z., Read, R. W., Hansen, G. and Vogel, P. (2012). Cardiomyopathy in α -kinase 3 (ALPK3)-deficient mice. *Vet. Pathol.* **49**, 131-141. doi:10.1177/0300985811402841
- Vetrini, F., D'Alessandro, L. C., Akdemir, Z. C., Braxton, A., Azamian, M. S., Eldomery, M. K., Miller, K., Kois, C., Sack, V., Shur, N. et al. (2016). Bi-allelic mutations in PKD1L1 are associated with laterality defects in humans. *Am. J. Hum. Genet.* **99**, 886-893. doi:10.1016/j.ajhg.2016.07.011
- Vierra, N. C., Dadi, P. K., Jeong, I., Dickerson, M., Powell, D. R. and Jacobson, D. A. (2015). Type 2 diabetes-associated K⁺ channel TALK-1 modulates β -cell electrical excitability, second-phase insulin secretion, and glucose Homeostasis. *Diabetes* **64**, 3818-3828. doi:10.2337/db15-0280
- Vilboux, T., Doherty, D. A., Glass, I. A., Parisi, M. A., Phelps, I. G., Cullinane, A. R., Zein, W., Brooks, B. P., Heller, T., Soldatos, A. et al. (2017a). Molecular genetic findings and clinical correlations in 100 patients with Joubert syndrome and related disorders prospectively evaluated at a single center. *Genet. Med.* **19**, 875-882. doi:10.1038/gim.2016.204
- Vilboux, T., Malicdan, M. C. V., Roney, J. C., Cullinane, A. R., Stephen, J., Yildirimli, D., Bryant, J., Fischer, R., Vemulapalli, M., Mullikin, J. C. et al. (2017b). CELSR2, encoding a planar cell polarity protein, is a putative gene in Joubert syndrome with cortical heterotopia, microphthalmia, and growth hormone deficiency. *Am. J. Med. Genet. A* **173**, 661-666. doi:10.1002/ajmg.a.38005
- Vogel, P., Read, R. W., Vance, R. B., Platt, K. A., Troughton, K. and Rice, D. S. (2008). Ocular albinism and hypopigmentation defects in Slc24a5^{-/-} mice. *Vet. Pathol.* **45**, 264-279. doi:10.1354/vp.45-2-264
- Vogel, P., Donoviel, M. S., Read, R., Hansen, G. M., Hazlewood, J., Anderson, S. J., Sun, W., Swaffield, J. and Oravec, T. (2009). Incomplete inhibition of sphingosine 1-phosphate lyase modulates immune system function yet prevents early lethality and non-lymphoid lesions. *PLoS ONE* **4**, e4112. doi:10.1371/journal.pone.0004112
- Vogel, P., Read, R., Hansen, G. M., Freay, L. C., Zambrowicz, B. P. and Sands, A. T. (2010a). Situs inversus in Dpcc/Poll^{-/-}, Nme7^{-/-}, and Pkd11^{-/-} mice. *Vet. Pathol.* **47**, 120-131. doi:10.1177/0300985810353553
- Vogel, P., Hansen, G., Fontenot, G. and Read, R. (2010b). Tubulin tyrosine ligase-like 1 deficiency results in chronic rhinosinusitis and abnormal development of spermatid flagella in mice. *Vet. Pathol.* **47**, 703-712. doi:10.1177/0300985810363485
- Vogel, P., Read, R., Hansen, G., Wingert, J., Dacosta, C. M., Buhning, L. M. and Shadoan, M. (2011). Pathology of congenital generalized lipodystrophy in Agpat2^{-/-} mice. *Vet. Pathol.* **48**, 642-654. doi:10.1177/0300985810383870
- Vogel, P., Read, R. W., Hansen, G. M., Payne, B. J., Small, D., Sands, A. T. and Zambrowicz, B. P. (2012a). Congenital hydrocephalus in genetically engineered mice. *Vet. Pathol.* **49**, 166-181. doi:10.1177/0300985811415708
- Vogel, P., Hansen, G. M., Read, R. W., Vance, R. B., Thiel, M., Liu, J., Wronski, T. J., Smith, D. D., Jeter-Jones, S. and Brommage, R. (2012b). Amelogenesis imperfecta and other biomineralization defects in Fam20a and Fam20c null mice. *Vet. Pathol.* **49**, 998-1017. doi:10.1177/0300985812453177
- Vogel, P., Liu, J., Platt, K. A., Read, R. W., Thiel, M., Vance, R. B. and Brommage, R. (2015a). Malformation of incisor teeth in Grem2^{-/-} mice. *Vet. Pathol.* **52**, 224-229. doi:10.1177/0300985814528218
- Vogel, P., Gelfman, C. M., Issa, T., Payne, B. J., Hansen, G. M., Read, R. W., Jones, C., Pitcher, M. R., Ding, Z.-M., DaCosta, C. M. et al. (2015b). Nephronophthisis and retinal degeneration in Tmem218^{-/-} mice: a novel mouse model for Senior-Løken syndrome? *Vet. Pathol.* **52**, 580-595. doi:10.1177/0300985814547392
- Vogel, P., Read, R. W., Reh, J. E. and Hansen, G. M. (2015c). Cryptogenic organizing pneumonia in Tomm5^{-/-} mice. *Vet. Pathol.* **50**, 65-75. doi:10.1177/0300985812450723
- Vogel, P., Read, R. W., Hansen, G. M., Powell, D. R., Kantaputra, P. N., Zambrowicz, B. and Brommage, R. (2016). Dentin dysplasia in Notum knockout mice. *Vet. Pathol.* **53**, 853-862. doi:10.1177/0300985815626778
- von Renesse, A., Petkova, M. V., Lützkendorf, S., Heinemeyer, J., Gill, E., Hübner, C., von Moers, A., Stenzel, W. and Schuelke, M. (2014). POMK mutation in a family with congenital muscular dystrophy with merosin deficiency, hypomyelination, mild hearing deficit and intellectual disability. *J. Med. Genet.* **51**, 275-282. doi:10.1136/jmedgenet-2013-102236
- Wang, T., Bu, C. H., Hildebrand, S., Jia, G., Siggs, O. M., Lyon, S., Pratt, D., Scott, L., Russell, J., Ludwig, S. et al. (2018). Probability of phenotypically detectable protein damage by ENU-induced mutations in the Mutagenetix database. *Nat. Commun.* **9**, 441. doi:10.1038/s41467-017-02806-4
- Wattler, S., Kelly, M. and Nehls, M. (1999). Construction of gene targeting vectors from lambda DOS genomic libraries. *BioTechniques* **26**, 1150-1156. doi:10.2144/99266rr02
- Wei, A.-H., Zang, D.-J., Zhang, Z., Liu, X.-Z., He, X., Yang, L., Wang, Y., Zhou, Z.-Y., Zhang, M.-R., Dai, L.-L. et al. (2013). Exome sequencing identifies SLG2A5 as a candidate gene for nonsyndromic oculocutaneous albinism. *J. Invest. Dermatol.* **133**, 1834-1840. doi:10.1038/jid.2013.49
- Wergedal, J. E., Kesavan, C., Brommage, R., Das, S. and Mohan, S. (2015). Role of WNT16 in the regulation of periosteal bone formation in female mice. *Endocrinology* **156**, 1023-1032. doi:10.1210/en.2014-1702
- West, D. B., Pasumarthi, R. K., Baridon, B., Djan, E., Trainor, A., Griffey, S. M., Willis, B., Rapp, J., de Jong, P. J. and Lloyd, K. C. (2015). A lacZ reporter gene expression atlas for 313 adult KOMP mutant mouse lines. *Genome Res.* **25**, 598-607. doi:10.1101/gr.184184.114
- White, J. K., Gerdin, A.-K., Karp, N. A., Ryder, E., Buljan, M., Bussell, J. N., Salisbury, J., Clare, S., Ingham, N. J., Podrini, C. et al. (2013). Genome-wide generation and systematic phenotyping of knockout mice reveals new roles for many genes. *Cell* **154**, 452-464. doi:10.1016/j.cell.2013.06.022
- Whitlock, N. A., Harrison, B., Mixon, T., Yu, X.-Q., Wilson, A., Gerhardt, B., Eberhart, D. E., Abuin, A. and Rice, D. S. (2009). Decreased intraocular

- pressure in mice following either pharmacological or genetic inhibition of ROCK. *J. Ocul. Pharmacol. Ther.* **25**, 187-194. doi:10.1089/jop.2008.0142
- Whyte, M. P., McAlister, W. H., Fallon, M. D., Pierpont, M. E., Bijanki, V. N., Duan, S., Otaify, G. A., Sly, W. S. and Mumm, S.** (2017). Raine syndrome (OMIM #259775), caused by FAM20C mutation, is congenital sclerosing osteomalacia with cerebral calcification (OMIM 259660). *J. Bone Miner. Res.* **32**, 757-769. doi:10.1002/jbmr.3034
- Williams, S. C. P.** (2016). Genetic mutations you want. *Proc. Natl. Acad. Sci. USA* **113**, 2554-2557. doi:10.1073/pnas.1601663113
- Wilson, F. H., Disse-Nicodème, S., Choate, K. A., Ishikawa, K., Nelson-Williams, C., Desitter, I., Gunel, M., Milford, D. V., Lipkin, G. W., Achard, J. M. et al.** (2001). Human hypertension caused by mutations in WNK kinases. *Science* **293**, 1107-1112. doi:10.1126/science.1062844
- Wilson, D. G., Phamluong, K., Lin, W. Y., Barck, K., Carano, R. A. D., Diehl, L., Peterson, A. S., Martin, F. and Solloway, M. J.** (2012). Chondroitin sulfate synthase 1 (*Chsy1*) is required for bone development and digit patterning. *Dev. Biol.* **363**, 413-425. doi:10.1016/j.ydbio.2012.01.005
- Wilson, R., Geyer, S. H., Reissig, L., Rose, J., Szumska, D., Hardman, E., Prin, F., McGuire, C., Ramirez-Solis, R., White, J. et al.** (2017). Highly variable penetrance of abnormal phenotypes in embryonic lethal knockout mice. *Wellcome Open Res.* **1**, 1. doi:10.12688/wellcomeopenres.9899.2
- Xing, W., Liu, J., Cheng, S., Vogel, P., Mohan, S. and Brommage, R.** (2013). Targeted disruption of leucine-rich repeat kinase 1 but not leucine-rich repeat kinase 2 in mice causes severe osteopetrosis. *J. Bone Miner. Res.* **28**, 1962-1974. doi:10.1002/jbmr.1935
- Yeo, G. S. H., Farooqi, I. S., Aminian, S., Halsall, D. J., Stanhope, R. G. and O'Rahilly, S.** (1998). A frameshift mutation in MC4R associated with dominantly inherited human obesity. *Nat. Genet.* **20**, 111-112. doi:10.1038/2404
- Zambrowicz, B. P. and Sands, A. T.** (2003). Knockouts model the 100 best-selling drugs—will they model the next 100? *Nat. Rev. Drug Discov.* **2**, 38-51. doi:10.1038/nrd987
- Zambrowicz, B. P., Friedrich, G. A., Buxton, E. C., Lilleberg, S. L., Person, C. and Sands, A. T.** (1998). Disruption and sequence identification of 2,000 genes in mouse embryonic stem cells. *Nature* **392**, 608-611. doi:10.1038/33423
- Zambrowicz, B. P., Abuin, A., Ramirez-Solis, R., Richter, L. J., Piggott, J., BeltrandelRio, H., Buxton, E. C., Edwards, J., Finch, R. A., Friddle, C. J. et al.** (2003). Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap screen to identify potential targets for therapeutic intervention. *Proc. Natl. Acad. Sci. USA* **100**, 14109-14114. doi:10.1073/pnas.2336103100
- Zamproni, I., Grasberger, H., Cortinovis, F., Vigone, M. C., Chiumello, G., Mora, S., Onigata, K., Fugazzola, L., Refetoff, S., Persani, L. et al.** (2008). Biallelic inactivation of the dual oxidase maturation factor 2 (DUOXA2) gene as a novel cause of congenital hypothyroidism. *J. Clin. Endocrinol. Metab.* **93**, 605-610. doi:10.1210/jc.2007-2020
- Zhang, W., Rajan, I., Savelieva, K. V., Wang, C.-Y., Vogel, P., Kelly, M., Xu, N., Hasson, B., Jarman, W. and Lanthorn, T. H.** (2008). Netrin-G2 and netrin-G2 ligand are both required for normal auditory responsiveness. *Genes Brain Behav.* **7**, 385-392. doi:10.1111/j.1601-183X.2007.00361.x
- Zhang, W., Savelieva, K. V., Suwanichkul, A., Small, D. L., Kirkpatrick, L. L., Xu, N., Lanthorn, T. H. and Ye, G.-L.** (2010). Transmembrane and ubiquitin-like domain containing 1 (*Tmub1*) regulates locomotor activity and wakefulness in mice and interacts with CAMLG. *PLoS ONE* **5**, e11261. doi:10.1371/journal.pone.0011261
- Zhang, W., Savelieva, K. V., Tran, D. T., Pogorelov, V. M., Cullinan, E. B., Baker, K. B., Platt, K. A., Hu, S., Rajan, I., Xu, N. et al.** (2012). Characterization of PTPRG in knockdown and phosphatase-inactive mutant mice and substrate trapping analysis of PTPRG in mammalian cells. *PLoS ONE* **7**, e45500. doi:10.1371/journal.pone.0045500
- Zhang, J., Wang, C., Shen, Y., Chen, N., Wang, L., Liang, L., Guo, T., Yin, X., Ma, Z., Zhang, B. et al.** (2016). A mutation in ADIPOR1 causes nonsyndromic autosomal dominant retinitis pigmentosa. *Hum. Genet.* **135**, 1375-1387. doi:10.1007/s00439-016-1730-2
- Zhao, S., Edwards, J., Carroll, J., Wiedholz, L., Millstein, R. A., Jaing, C., Murphy, D. L., Lanthorn, T. H. and Holmes, A.** (2006). Insertion mutation at the C-terminus of the serotonin transporter disrupts brain serotonin function and emotion-related behaviors in mice. *Neuroscience* **140**, 321-334. doi:10.1016/j.neuroscience.2006.01.049
- Zheng, H.-F., Tobias, J. H., Duncan, E., Evans, D. M., Eriksson, J., Paternoster, L., Yerges-Armstrong, L. M., Lehtimäki, T., Bergström, U., Kähönen, M. et al.** (2012). WNT16 influences bone mineral density, cortical bone thickness, bone strength, and osteoporotic fracture risk. *PLoS Genet.* **8**, e1002745. doi:10.1371/journal.pgen.1002745
- Zou, M., Alzahrani, A. S., Al-Odaib, A., Alqahtani, M. A., Babiker, O., Al-Rijjal, R. A., BinEssa, H. A., Kattan, W. E., Al-Enezi, A. F., Al Qarni, A. et al.** (2018). Molecular analysis of congenital hypothyroidism in Saudi Arabia: SLC26A7 mutation is a novel defect in thyroid dysmorphogenesis. *J. Clin. Endocrinol. Metab.* **103**, 1889-1898. doi:10.1210/jc.2017-02202