

Maternal–Fetal Conflict: Rapidly Evolving Proteins in the Rodent Placenta

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Abstract

Conflicting evolutionary interests between mother and offspring are hypothesized to drive an evolutionary arms race during mammalian pregnancy, and thus, positive selection may cause the rapid divergence of placental proteins that affect maternal or fetal fitness. We investigated the genomic consequences of placental expression in rodents and report that a substantial proportion (20.5%) of genes specifically expressed in the mature placenta are rapidly evolving. Moreover, we found that most rapidly evolving genes belong to just three pregnancy-related gene families: placental cathepsins, prolactins, and placental carcinoembryonic antigens. We then sequenced the most rapidly evolving gene, trophoblast-specific protein alpha (*Tpbpa*), in nine different *Mus* species/subspecies and found evidence of positive selection within the *Mus* lineage, with an excess of nonsynonymous changes clustering near a functionally important interaction site. Together, these results suggest that placental proteins, which mediate interactions between mother and offspring, often may be the targets of evolutionary conflict.

Key words: adaptive evolution, evolutionary conflict, placenta, positive selection, reproduction.

Comparative genomic studies across many diverse taxa consistently have found that those genes related to fertilization and immune responses are among the most rapidly evolving in the genome (e.g., Swanson and Vacquier 2002; Nielsen et al. 2005; Gibbs et al. 2007). Many of these genes are probably involved in evolutionary “arms races,” driven by host–pathogen conflict or male–female sexual conflict, in which they experience strong positive selection to maintain optimal fitness (Dawkins and Krebs 1979; Chapman et al. 2003). Another arena for evolutionary conflict, which is less well studied, is antagonism between mother and offspring during pregnancy. The maternal–fetal conflict hypothesis proposes that there is a selective advantage to offspring who can selfishly maximize their own growth at the expense of the fitness of their mother and siblings, whereas there is an opposing advantage to mothers who can counter these adaptations by maintaining equal investment in all offspring and surviving to breed in future generations (Haig 1993; Zeh and Zeh 2000).

The primary site of physiological exchange between mother and offspring during mammalian pregnancy is the placenta—a temporary organ formed by the fusion of fetal extraembryonic membrane with maternal uterine tissue. At this maternal–fetal interface, a complex interplay of proteins from these two distinct individuals mediates nutrient and waste exchange, immunoregulation, and other pregnancy-related physiological processes. These interactions are likely to result in an evolutionary conflict between maternal and fetal genes (Haig 2008) in which genes under conflict may either experience balancing selection and reach a stable resolution (McVean and Hurst 1997) or experience repeated bouts of positive selection in an

arms race to maintain maximal fitness. In this study, we asked whether we could identify placental genes that show strong signatures of continuous positive selection and might therefore be involved in an evolutionary arms race between mother and offspring.

Early comparative genomic screens across mammalian orders (e.g., human vs. mouse) did not identify placental genes to be rapidly evolving as a class (e.g., Clark et al. 2003; Lindblad-Toh et al. 2005), which is unsurprising because many placental genes are lineage-specific and would therefore be missed in comparisons of divergent taxa (Rawn and Cross 2008). As more mammalian genomic data have become available, several reports have revealed signatures of positive selection in placental genes within lineages (e.g., primates and artiodactyls; Hughes et al. 2000; Hou et al. 2009). Here, we investigated placental gene evolution within the rodent lineage, which offers several advantages: 1) rodents have short gestation times and are a widely used experimental model, so we can readily examine subsets of placental genes from each stage of development rather than focusing on the term placenta, which has a distinct expression profile compared with a developing placenta (Knox and Baker 2008), 2) rodents generally have large and multisired litters, which likely intensifies the potential maternal–fetal conflict (Zeh and Zeh 2000), and 3) there are several examples of interfertile monogamous/promiscuous sister species (e.g., *Peromyscus maniculatus* and *P. polionotus*, respectively), which allows for the role of mating system, and of particular candidate genes, in maternal–fetal conflict to be tested directly.

For this study, we confined our analysis to placental genes in two rodents, mouse (*Mus musculus*) and rat

Table 1. Pairwise Comparisons of Placentally Expressed Genes between *Mus* and *Rattus*.

Category	N	Total			Nonsecreted $\bar{\omega}$ (N)	Secreted $\bar{\omega}$ (N)	$\omega \geq 0.5$ (%)
		\bar{d}_N	\bar{d}_S	$\bar{\omega}$			
Genome	18,708	0.03	0.21	0.17	0.15 (14,683)	0.24 (4,025)	2,438 (13.1)
Placental	704	0.08	0.24	0.25	0.20 (469)	0.35 (235)	95 (13.5)
Developing	519	0.06	0.24	0.23	0.20 (388)	0.31 (131)	54 (10.4)
Fetal	342	0.06	0.24	0.22	0.19 (290)	0.36 (52)	32 (9.4)
Maternal	200	0.06	0.23	0.25	0.20 (111)	0.31 (89)	29 (14.5)
Mature	191	0.09	0.26	0.33	0.23 (85)	0.41 (106)	41 (20.5)
Fetal	61	0.11	0.25	0.44	0.31 (31)	0.56 (30)	21 (34.4)
Maternal	150	0.09	0.26	0.33	0.23 (62)	0.40 (88)	35 (23.3)

NOTES.—Categories following Knox and Baker (2008). “Placental genes” exhibit maximal expression in the placenta. Genes classified as “developing” are those overexpressed from E8.5 to 13.5, whereas “mature” genes are overexpressed from E13.5 to term. “Fetal” and “maternal” refer to the tissue of expression and are not mutually exclusive (e.g., a gene may be overexpressed in both fetal and maternal mature placental tissue). N, number of unique *Mus/Rattus* coding sequence alignments (obtained from Ensembl build 49); \bar{d}_N , mean nonsynonymous substitution rate; \bar{d}_S , mean synonymous substitution rate; $\bar{\omega}$, mean ω for all genes estimated using PAML v4 (Yang 2007); nonsecreted and secreted, secreted proteins predicted using Phobius, values are significantly different for all secreted/nonsecreted comparisons within a gene class (one-way analysis of variance, $P < 0.01$); percentage and number of rapidly evolving genes with $\omega \geq 0.5$ are given.

(*Rattus norvegicus*), which share the same type of placenta (Mess 2003) and are the only rodents with well-annotated genomes. Because the timing of expression during development has been demonstrated to correlate with the rate of evolution (e.g., in testes; Good and Nachman 2005), we utilized a comprehensive time course microarray over mouse placental development (Knox and Baker 2008) to group placental genes into “developing” (specifically expressed between E8.5 and 13.5) and “mature” (E13.5 to term) as well as “maternal” (decidua) and “fetal” (fetal chorion) based on the tissue of origin. Finally, we examined whether secreted proteins, which can traverse the maternal–fetal boundary, are more likely to be targets of positive selection.

Our specific goals in examining the molecular evolution of rodent placental genes were to identify the most rapidly evolving genes and to determine whether specific functional classes of genes and/or their developmental timing affect their rate of evolution. By comparing the nonsynonymous substitution rate to the synonymous substitution rate, we estimated the rate of evolution for each gene ($d_N/d_S = \omega$) in *Mus* and *Rattus*. High values of ω ($\omega \geq 0.5$ for pairwise comparisons) indicate elevated rates of amino acid sequence divergence and suggest that some sites are likely to be targets of positive selection (Swanson et al. 2004).

From a total of 786 genes predominantly expressed in the placenta (following Knox and Baker 2008), we aligned 704 coding sequences between mouse and rat based on the Ensembl peptide alignments. We next performed pairwise estimates of ω using maximum likelihood (PAML v4, runmode = -2; Yang 2007); table 1 summarizes these results. When all placental genes are considered together, $\bar{\omega}$ is significantly higher than the genome-wide average ($\bar{\omega}$: 0.25 vs. 0.17, one-way analysis of variance, $P < 0.0001$), although evolutionary rate is often elevated for genes with tissue-specific expression (Winter et al. 2004). Secreted proteins had a higher $\bar{\omega}$ than did nonsecreted proteins (table 1) within all gene classes (genome wide and for each placental subclass). When we considered the temporal pattern of expression, we found that genes highly expressed in the mature placenta exhibit significantly faster rates of evolution than genes expressed earlier in placental development

($\bar{\omega}$: 0.33 vs. 0.23, $P < 0.0001$). In fact, secreted proteins from the mature fetal component of the placenta are the most rapidly evolving class of placental genes even compared with other placental secreted genes ($N = 30$, $\bar{\omega}$: 0.56 vs. 0.31, $P < 0.005$).

Table 2 lists the 15 most rapidly evolving genes from our analysis. Notably, we identified rapidly evolving genes from three major rodent placental gene families: carcinoembryonic antigens (CEAs), prolactins, and placentally expressed cathepsins (PECs). All three of these families are comprised of secreted proteins (Zebhauser et al. 2005; Soares et al. 2007; Mason 2008), and the majority of family members represented in our analysis are expressed in the fetal portion of the mature placenta (CEAs: 7 of 8 exhibit fetal expression and 8 of 8 expressed in mature placenta; prolactins: 14 of 16 and 9 of 16; and PECs: 10 of 10 and 8 of 10).

The CEA family is located on mouse chromosome 7 and comprises a large group of pregnancy-specific glycoproteins (PSGs) and carcinoembryonic antigen-related cell-adhesion molecules (CEACAMs), many of which are rodent specific and exhibit placenta-specific expression (Zebhauser et al. 2005). Although their precise function is unknown, they probably help modulate the maternal immune system (Wynne et al. 2006). In our analysis, the most rapidly evolving placental CEAs are *Ceacam11* ($\omega = 1.04$), *Ceacam12* ($\omega = 1.11$), *Ceacam13* ($\omega = 0.95$), and *Ceacam14* ($\omega = 1.06$). In addition, our data set includes CEAs *Ceacam18*, *Psg17*, *Psg18*, and *Psg28*, which also exhibit elevated rates of evolution ($N = 4$, $\bar{\omega} = 0.62 \pm 0.09$).

Placental prolactin hormones are located in a cluster on mouse chromosome 13 and play extremely diverse roles in pregnancy; members of this family are important for placental development, vascularization, stress response, and a variety of other physiological functions (Haig 2008). The most rapidly evolving prolactins we identified were *Prl3c1* ($\omega = 0.93$), a maternally expressed prolactin, and *Prl7a2* ($\omega = 0.92$), an important stimulator of hematopoiesis in trophoblast (Soares et al. 2007). Additional placental prolactins identified in our analysis also exhibit high rates of evolution ($N = 14$, $\bar{\omega} = 0.62 \pm 0.10$).

Finally, the PECs are a rodent-specific family of placental cathepsins that are also located in a cluster on mouse

Table 2. Top 15 Rapidly Evolving Placental Genes Based on Pairwise ω Estimated Using Maximum Likelihood.

Gene	Chr	d_N	d_S	ω	Fetal	Maternal	Developing	Mature	Secreted
<i>Tpbpa</i>	13qB2	0.30	0.20	1.50	X	—	—	X	X
<i>Pdgfrl</i>	8qA4	0.09	0.06	1.44	—	X	X	—	—
<i>2610036L11Rik</i>	10qA4	0.14	0.10	1.43	X	—	X	—	—
<i>Tpbpb</i>	13qB2	0.28	0.20	1.39	X	X	—	X	X
<i>Ctla2a</i>	13qB2	0.17	0.14	1.22	X	—	—	X	X
<i>Ceacam12</i>	7qA2	0.27	0.24	1.12	X	X	—	X	X
<i>Ceacam14</i>	7qA2	0.26	0.24	1.07	X	X	—	X	—
<i>Ccl9</i>	11qC	0.41	0.40	1.05	—	X	—	X	X
<i>Ceacam11</i>	7qA2	0.25	0.24	1.04	X	X	—	X	X ^a
<i>Pvr</i>	7qA3	0.36	0.38	0.97	—	X	—	X	X
<i>Glrx</i>	13qC1	0.09	0.09	0.97	X	—	X	—	—
<i>Spink3</i>	18qB3	0.29	0.30	0.96	X	—	X	—	X
<i>Ceacam13</i>	7qA2	0.22	0.24	0.95	X	X	—	X	X
<i>Prl3c1</i>	13qA3	0.21	0.23	0.93	—	X	X	—	X
<i>Prl7a2</i>	13qA3	0.18	0.19	0.92	X	X	—	X	X

Chr, chromosomal position in mouse; d_N , nonsynonymous substitution rate; d_S , synonymous substitution rate; ω , pairwise ω estimated with PAML; fetal, expressed in the fetal portion of the placenta; maternal, expressed in maternal portion; developing, high placental expression during E8.5 to E13.5; mature, high placental expression during E13.5 to term; secreted, predicted signaling molecule (identified with Phobius; Käll et al. 2004). Expression data annotation obtained from Knox and Baker (2008).

^a From Kataoka et al. (2000).

chromosome 13 and include mature proteases (cathepsins 3, 6, 7, 8, M, Q, R) and predicted propeptide inhibitors (*Tpbpa*, *Tpbpb*, and *Ctla2a*). Together, the PECs function to structurally remodel placental tissue during development (Mason 2008). The most rapidly evolving genes from this family are *Tpbpa* ($\omega = 1.50$), *Tpbpb* ($\omega = 1.39$), and *Ctla2a* ($\omega = 1.22$). In addition, the mature proteases as a group are also rapidly evolving ($N = 7$, $\bar{\omega} = 0.48 \pm 0.09$), potentially as a consequence of an evolutionary conflict between the placental proteases and their inhibitors.

To test for positive selection within a lineage, we sequenced the most rapidly evolving gene, trophoblast-specific protein alpha (*Tpbpa*), in nine different *Mus* species/subspecies (fig. 1; Supplementary Material, Supplementary Material online). Under the M8a-M8 model comparison in PAML, we found that *Tpbpa* was indeed evolving rapidly within the *Mus* lineage (Likelihood ratio test = 37.5, $P < 0.001$). The peptide encoded by *Tpbpa* contains an interaction domain predicted to block to the substrate-binding site of mature cathepsin to prevent premature

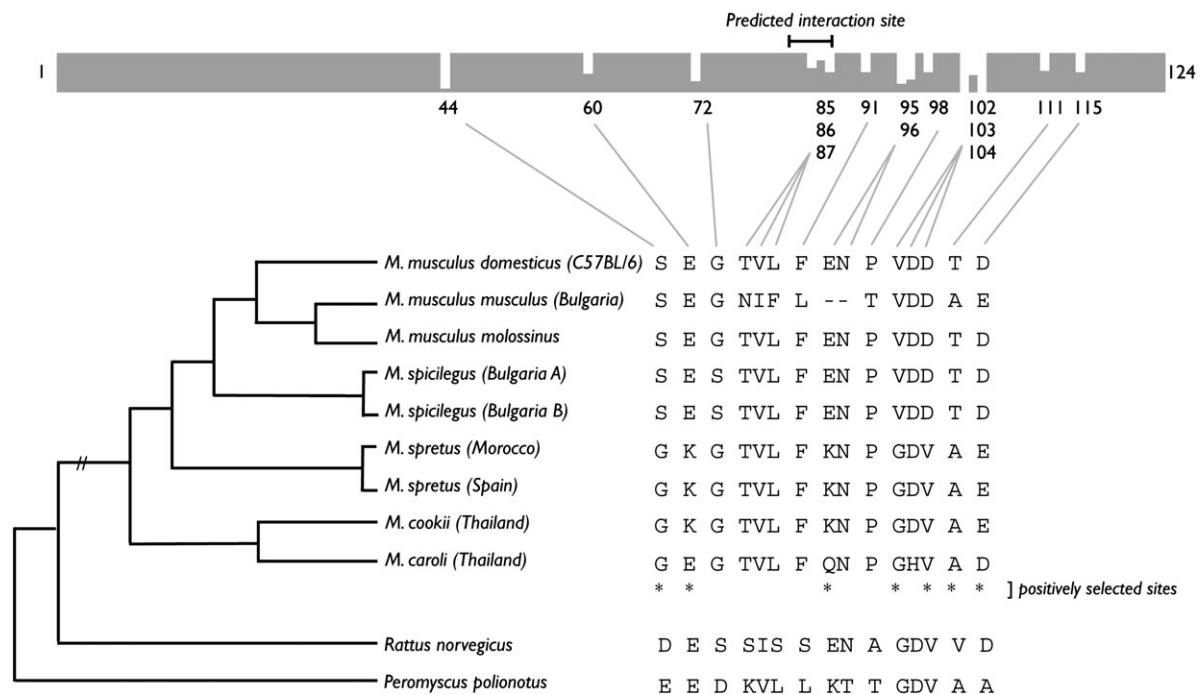


Fig. 1. Rapid evolution of *Tpbpa* in *Mus*. The conservation plot from the amino acid alignment of *Tpbpa* from nine *Mus* species/subspecies in which gray indicates conservation and white shows variation. Divergent amino acid sites are given and displayed next to a species phylogeny (Tucker et al. 2005). Sites marked with "*" are positively selected with $P > 99\%$ using the PAML Bayes Empirical Bayes analysis under M8. The corresponding amino acid in *Rattus norvegicus* and *Peromyscus polionotus* is shown below each site.

proteolytic activity (Coulombe et al. 1996) and shows high sequence variation when compared with both placental and nonplacental cathepsins, possibly reflecting different substrate specificities (Deussing et al. 2002). We observed that positively selected sites in *Tpbpa* clustered significantly near this predicted interaction site (permutation test, $P = 0.001$); although the sites closest to the interaction site do not appear to be under positive selection, their proximity is suggestive of functional consequences. Furthermore, we observed that 12 of 15 amino acid sites that are variable among *Mus* species are also variant in either *Rattus* or deer mice (genus *Peromyscus*; GenBank accession number EV469066.1) compared with the *Mus* consensus, which suggests that these may be functionally significant sites and undergo repeated adaptive substitutions.

Overall, we show that a subset (~13.5%) of placental proteins, especially those that are secreted and expressed late in development, probably experience positive selection based on their pattern of sequence divergence. Although a pairwise comparison is a generally low power and conservative test for positive selection, we find supporting evidence that the most rapidly evolving gene *Tpbpa* is experiencing positive selection in a functionally relevant domain even within the *Mus* lineage. It is important to note that many genes we identified are part of large gene families in which there may be some functional redundancy; hence, relaxed functional constraint may also contribute to their rapid evolution (Hughes 2007). We avoided the inclusion of pseudogenes by only considering full-length coding sequences that were in-frame and known to be highly expressed in the mouse placenta. Future comparative genomic data from promiscuous and monogamous rodents (e.g., *P. maniculatus* and *P. polionotus*, respectively) can test whether the rapidly evolving genes are under positive or relaxed selection because maternal–fetal conflict is hypothesized to be most severe in polyandrous species with multisired litters (Zeh and Zeh 2000).

Several lines of evidence suggest that many of these placental proteins may be involved in an evolutionary conflict between mother and offspring. First, the most rapidly evolving placental genes are expressed in the mature placenta in which the majority of genes are involved in physiological exchange between mother and offspring (Knox and Baker 2008). Second, among genes expressed in the mature placenta, fetal-secreted proteins are the most rapidly evolving class, consistent with their role in directly interacting with maternal physiology or immune response. Although these observations are also consistent with maternal–offspring coadaptation (Wolf and Brodie 1998), we also find evidence for positive selection acting broadly in the placental gene families, CEAs, prolactins, and PECs. These three gene families share characteristics with sexual conflict genes first identified in *Drosophila* seminal fluid; the most rapidly evolving seminal proteins include secreted proteins such as glycoproteins, proteins with enzymatic functions including proteases and their inhibitors, and proteins involved in the immune response (Wolfner 2002; Ravi Ram and Wolfner 2007). We suggest, then, that evolution-

ary arms races between mother and offspring may be an important selective force in the evolution of viviparous species, especially those with large and multisired litters.

Supplementary Material

Supplementary file is available at *Molecular Biology and Evolution* online (<http://www.mbe.oxfordjournals.org>).

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