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Maya Belghazi, Benoît Guyonnet, Xavier Druart, Jean Luc Gatti, Françoise
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Mammalian Epididymal Proteome

Jean-Louis Dacheux⁽¹⁾, Clémence Belleannée, Russell Jones⁽²⁾, Valérie Labas⁽³⁾, Maya Belghazi⁽³⁾, Benoît Guyonnet, Xavier Druart, Jean Luc Gatti, Françoise Dacheux

INRA-CNRS 6175, Nouzilly, 37380, France ;

(2) The University of Newcastle, Newcastle, NSW, Australia.

(3) Service de Spectrométrie de Masse pour la Protéomique, INRA, Nouzilly, F.

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(1) **Corresponding author:** Jean Louis Dacheux, Tel: 33 2 47 42 79 49, Fax: 33 2 47 42 77 43

jdacheux@tours.inra.fr,

Summary

20 In all mammalian species, the final differentiation of the male germ cell occurs in the epididymal duct where the spermatozoa develop the ability to be motile and fertilize an ovum. Understanding of these biological processes is the key to understanding and controlling male fertility. Comparative studies between several mammals could be an informative approach to finding common sperm modifications which are not species-specific. The new global biological approaches such the transcriptomes and proteomes provide considerable information which can be used for such comparative approaches. This report summarizes our proteomic studies of the epididymis of several mammals, including humans.

Introduction:

30 Epididymal function is essential for the fertility of male mammals because their sperm are infertile when they leave the testes and only acquire the ability to fertilize an ovum during passage through the epididymides. It is also essential that the epididymides accumulate and store sperm as, depending upon species, it takes 0.5 to 2 days for the testes to produce the number of sperm in a normal ejaculate, and in a competitive mating system males may inseminate up to 50 females in a day (Jones, 1999; Jones et al., 2007). The efficacy of sperm storage in the epididymis is so high that fertile sperm can survive in an isolated epididymis for several days at 4°C (Guérin et al., 2003)

40 The epididymis is a very long duct which receives testicular sperm via the efferent ducts. In eutherian and marsupial mammals the duct is differentiated into about 6 structurally distinct segments, indicating a well developed division of labor. Although all mammals have an initial segment of the epididymis with distinctive characteristics, there is variation between species in the structure and extent of the different segments, suggesting some variation in post-testicular sperm maturation and storage (Jones, 2002:). The division of labor through the epididymis has been confirmed in studies of epididymal physiology and sperm modifications along the duct, and these findings have lead to the paradigm that the maturational changes in sperm in the epididymis are the result of sequential changes in their milieu, particularly the proteins secreted by the epididymal epithelium (Dacheux et al., 2003). In view of the variations in epididymides between species mentioned above, it is considered that there must be conserved aspects and variations between species in the changes in protein composition throughout the epididymis, presumably reflecting the occurrence and relative significance of sperm maturation and storage between species. This

short review therefore compares our findings regarding the proteome and secretome of epididymal fluid in mammalian species: i.e. the stallion, ram, boar and human.

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Characteristics of intra luminal epididymal proteomes

Most of the studies analyzing the proteins of epididymal fluids have been performed on samples obtained either from tubule micropuncture (Turner et al., 1995; Turner et al., 1999) or from microperfusion techniques (Druart et al., 1994; Syntin et al., 1996 ; Fouhécourt et al., 2000 ; Dacheux et al., 2006) The epididymal fluid provides a milieu for the gamete analogous to blood plasma providing a milieu for cell tissues. Due to the presence of the blood-testis and blood-epididymis barriers, most of the blood proteins are not found in epididymal plasma (albumin, transferrin and certain others being exceptions). In contrast to the stable concentration of blood plasma proteins, the protein concentration in epididymal fluid varies greatly along the duct: from 2-4 mg/ml in the initial segment of the epididymis, a maximum of 50-60 mg/ml in the distal caput and 20-30 mg/ml in the more distal regions of the organ (Fig 1C.). For most of the species studied, these variations in protein concentration follow the changes in water content of the fluid as assessed by changes in sperm concentration (Fig.1A).

Several hundred epididymal proteins have been described electrophoretically and some have been identified. There is a wide range of dynamics in the abundance of these proteins (probably around 10 orders of magnitude). About 15-20 proteins make up more than 60 to 80% of the total protein concentration. The most common proteins found are lactoferrin, procathepsin D, NCP2 (HE1, CTP, cholesterol transfer protein), GPX (glutathione peroxidase), beta-N-acetyl-hexosaminidase, mannosidase, galactosidase, PGDS (Prostaglandin D2 synthase), clusterin, CRISP (Cystein-rich secretory protein) and E-RAPB (epididymal retinoic acid-binding protein).

The protein composition changes continuously throughout the duct, independently of the protein concentration in the fluid. The concentrations of the major common proteins cited above vary between species (Fig. 2). Lactoferrin, mannosidase, PGDS and albumin are present in high concentrations in the stallion, boar, ram and human, respectively, but GPX and PGDS are virtually absent in humans and boar, respectively.

Most of the epididymal proteins are characterized by their numerous isoforms which result from their high degree of glycosylation. The pI of these multi isoforms can vary widely, ranging from pH 3 to 8 for the same protein (i.e. RNase 10 in the boar, Fig. 3). The degree of glycosylation for the same protein can be different according to the epididymal

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region eg. clusterin and PGDS in the horse, where the number of isoforms is different between the anterior and the posterior part of the organ (Fig 3.), or RNase 10 (Train A) which is different between species for the boar and the ram (Fig 3).

Dynamics of epididymal fluid proteomes

The spatial changes in the composition of luminal proteins are the result of two opposite activities of the epithelium: protein secretion and protein absorption throughout the epididymal duct. In the anterior part of the epididymis, the epididymal fluid is composed of a mixture of testicular and epididymal proteins. Most of the proteins originating from the testis, such as albumin, transferrin, testicular clusterin and PGDS (Fouchecourt et al., 2000) are reabsorbed in the efferent ducts (Clulow et al., 1994) The rapidity of their absorption is species-specific and generally almost all are absent in the posterior part of the epididymis, except in humans in which albumin and transferrin are still present in large quantities.

The epididymal epithelium has high protein synthesis and secretion activity, activity being high both in the rates of protein synthesis and secretion and in the variety of proteins secreted. The anterior part of the epididymis is the most active (Figs. 1B, 2). As for the protein concentration, from 70 to 80% of the secretome is composed of 10 to 20 of the major secreted proteins present in the luminal fluid (Fig. 4). Most of the luminal proteins are secreted by the epithelium, but some, such as ACE, are released from the sperm surface by an unknown sheddase in an anterior part of the epididymis (Fig. 6) (Gatti et al., 1999; Metayer et al., 2002; Thimon et al., 2005).

Among the different proteins secreted, the same protein can be secreted in the same region of the epididymis in different species, for example PGDS, GPX and clusterin in the anterior part, and glucosidases in the middle part. Clusterin is secreted at a greater rate than the other proteins and can represent 30-40% of the all the proteins secreted. This clusterin can be sequentially secreted under different isoforms in different parts of the epididymis as in the stallion and the ram (Figs. 2, 3). Some highly secreted proteins are characteristic of a species, for example, lactoferrin in the stallion, PGDS and GPX in the ram and RNase 10, mannosidase and hexosaminidase in the boar (Fig. 2).

In humans, in contrast to other species, few changes in pattern of protein secretion occur throughout the epididymis, a finding which correlates with the low degree of structural differentiation of the epididymal epithelium along the duct (Holstein, 1969).

Variations in luminal protein concentrations, controlled by secretory and absorption activities, are modulated for each species by the length of the epididymal duct involved in

the two activities, i.e. the flow rate of the luminal fluid, enzyme degradation or the binding of protein on the sperm membrane. All these parameters differentially affect the concentrations of almost all proteins. Some are reabsorbed as soon as they are secreted and never reach high luminal concentrations, for example RNase 10 in the boar (Fig 5B) (Castella et al., 2004), and several isoforms of clusterin in the stallion (Fig.3) (Fouchecourt et al., 2000). For most other proteins, concentrations are high in the region where they are secreted and gradually decrease in the following region, for example GPX and PGDS in different species (Figs.2, 5A). The concentrations of some other proteins increase throughout the epididymal duct, as for example lactoferrin in the stallion and hexosaminidase and mannosidase in the boar (Fig.2).

Relationships between sperm maturation and epididymal proteome among species

The most obvious and easily observed change in sperm directly related to acquisition of fertility is the activation of their flagella machinery. A gradual increase in the coordination of propagation of flagellar bending makes the gamete motile and progressive. In species that have been studied, this activation occurs in sperm in the corpus epididymides, after a transitory phase of uncoordinated beating. In humans, this activation occurs in sperm from the distal caput of the epididymis (Fig 6).

Transit of the sperm through the epididymis must be necessary for sperm maturation since sperm maturation has not been achieved in vitro to date. Sperm maturation is probably related to sequential modifications which occur mainly on the sperm surface. During the maturation process, the immature gametes progressively lose or modify most of their testicular surface proteins and gain new transient or permanent proteins in well organized membrane protein domains. The pattern of sperm surface modification is species-specific, related to the nature of the surface proteins involved or the epididymal regions where these modifications occur. However, common changes in several identified sperm surface proteins have been described, e.g. such as fertilin (Primakoff et al., 1988), CRISP1 (Roberts et al., 2006), P34H (Sullivan et al., 2006) and ACE (Gatti et al., 2002), which represent the most important sperm surface changes (Fig 6). Such common changes can also be illustrated by the global surface changes of sialoglycoproteins on the sperm surface in the last part of the epididymis (Dacheux et al., 1989; Dacheux et al., 1987).

Few relationships have been observed between the major sequential modifications of the epididymal proteome and the sequential changes on the sperm surface. Furthermore, there is currently no direct evidence of a specific role for these major proteins, although several of them are known to be enzymes, inhibitors or binding proteins. For several of them, such

150 as clusterin which is the most common epididymal protein found, gene KO in mice does not change the fertility of the animals (McLaughlin et al., 2000) .

It is probable that these major epididymal proteins which surround the gametes are more involved in sperm preservation than in inducing specific and localized modifications on the sperm surface. The high concentrations of proteins involved in the protection of gametes from oxidative stress, such as GPX5, thioredoxin peroxidase, glutathione *S*-transferase P, thioredoxin peroxidase and superoxide dismutase, probably contribute to sperm survival during epididymal storage.

160 However, no more than 10% of all the proteins present in the epididymal fluid have been identified. The wide range of protein concentrations makes identification of the less highly represented proteins by mass spectrometry more difficult. Furthermore, these unidentified proteins contain the most hydrophobic proteins. These proteins are almost never present in a soluble form in the epididymal luminal fluid but partly associated with membrane structures such as epididymosomes (Frenette et al., 2002; Gatti et al., 2005), or directly transferred to the sperm surface or associated with several binding proteins such as clusterin and several lipocalins (Ecroyd et al., 2005).

Conclusion

170 New approaches, including large scale analysis, are now being applied to the study of epididymis physiology and the post-testicular differentiation of spermatozoa. Both transcriptomic (Turner et al., 2006) and proteomic approaches provide a dramatic advance to our understanding of male reproduction in different species. From the first results of these global approaches, it is evident that significant differences exist between species either in the sequential changes in the luminal proteome or sperm surface proteins. Each species appears to have developed its own strategy for sperm maturation and preservation. A general feature of sperm maturation probably exists between mammals but it probably involves different protein combinations. However, there is a need to identify more epididymal and sperm proteins in order to obtain greater understanding of this aspect of the male reproductive system which is fundamental for the survival of all species.

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Legends of the figures:

270 Figure 1: Spermatocrit (A), protein secretion (B) and concentrations of luminal proteins (C) in the epididymal fluid from 9 regions of the epididymis (1-4: caput; 5-6: corpus; 7-9: cauda) for three species (ram, boar, stallion) from (Syntin et al., 1996; Fouchecourt et al., 2000) and unpublished data.

Figure 2: Epididymal secretomes and proteomes of four species. For proteomes, each plate corresponds to 1D gel electrophoresis separation of about the same quantity of epididymal protein from each region of the epididymis. The secretome diagrams represent each secreted protein from the different epididymal regions and are expressed as the percentage of total secretion of the whole organ. The plates illustrating the secretory activities of the different epididymal regions for the
280 four animal species correspond to the autoradiograms of the same four 1D gel separations presented above for the proteome (Syntin et al., 1996; Fouchecourt et al., 2000 ; Druart et al., 1994 ; Druart, unpublished data ; Dacheux et al., 2006).

Figure 3: Immunodetection of 2D electrophoresis gels of isoforms of three proteins (PGDS, clusterin and RNase 10) according to their epididymal and species origins (stallion: Fouchecourt et al., 2000; boar: Castella et al., 2004 and ram: unpublished data).

Figure 4: Distribution of the major proteins secreted by the human and boar epididymis (Dacheux et al., 2006; Syntin et al., 1996)

290 Figure 5: Relationship between secretome and proteome. A) Secretion and protein concentrations for five major proteins present in the epididymal fluid of the stallion. B) Epididymal localisation of the secretion, luminal protein and corresponding RNA of RNase 10 (Train A) from the initial segment (IS), caput (1-4), corpus (5-6), cauda (7-9) in the boar (Castella et al., 2004).

Figure 6: Sperm epididymal maturation according to epididymal region in the human (Dacheux et al., 1987) and porcine (Dacheux et al., 1984) related to ACE, sialoprotein modifications on the sperm surface and percentage of progressive motility along the epididymis.

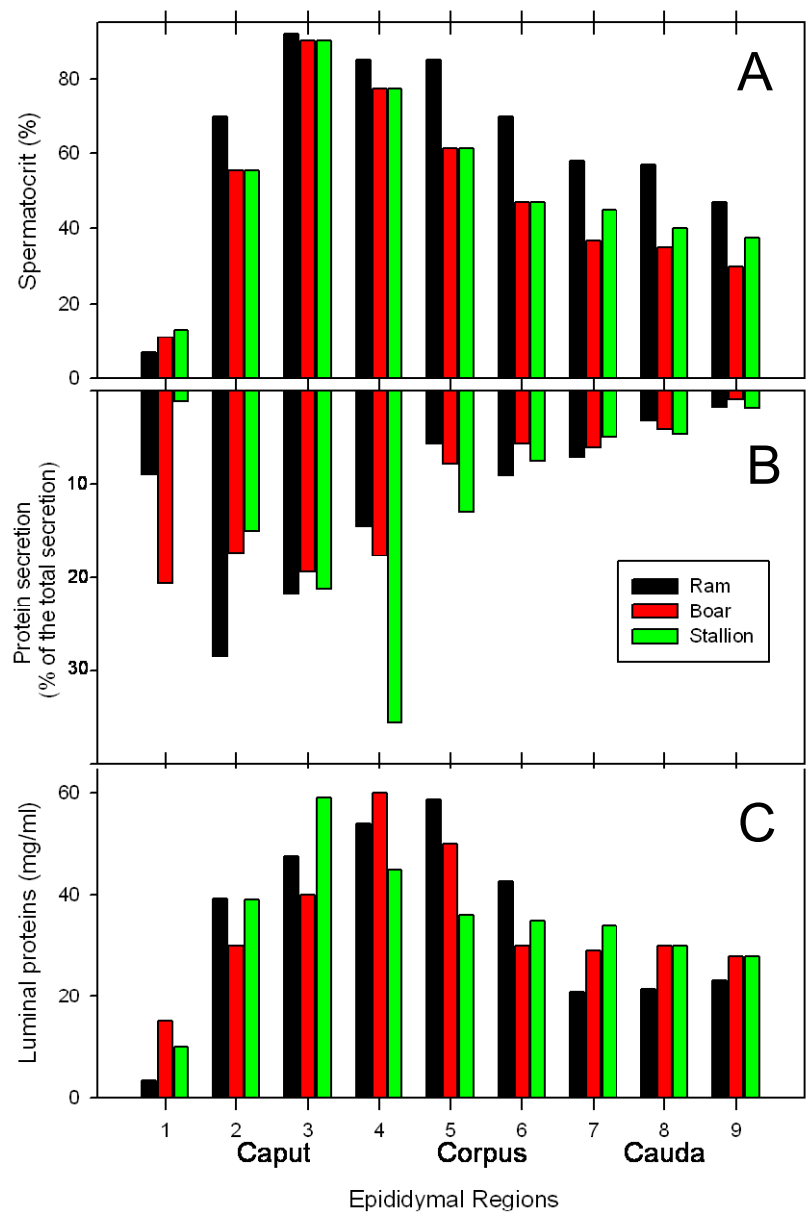


Figure 1

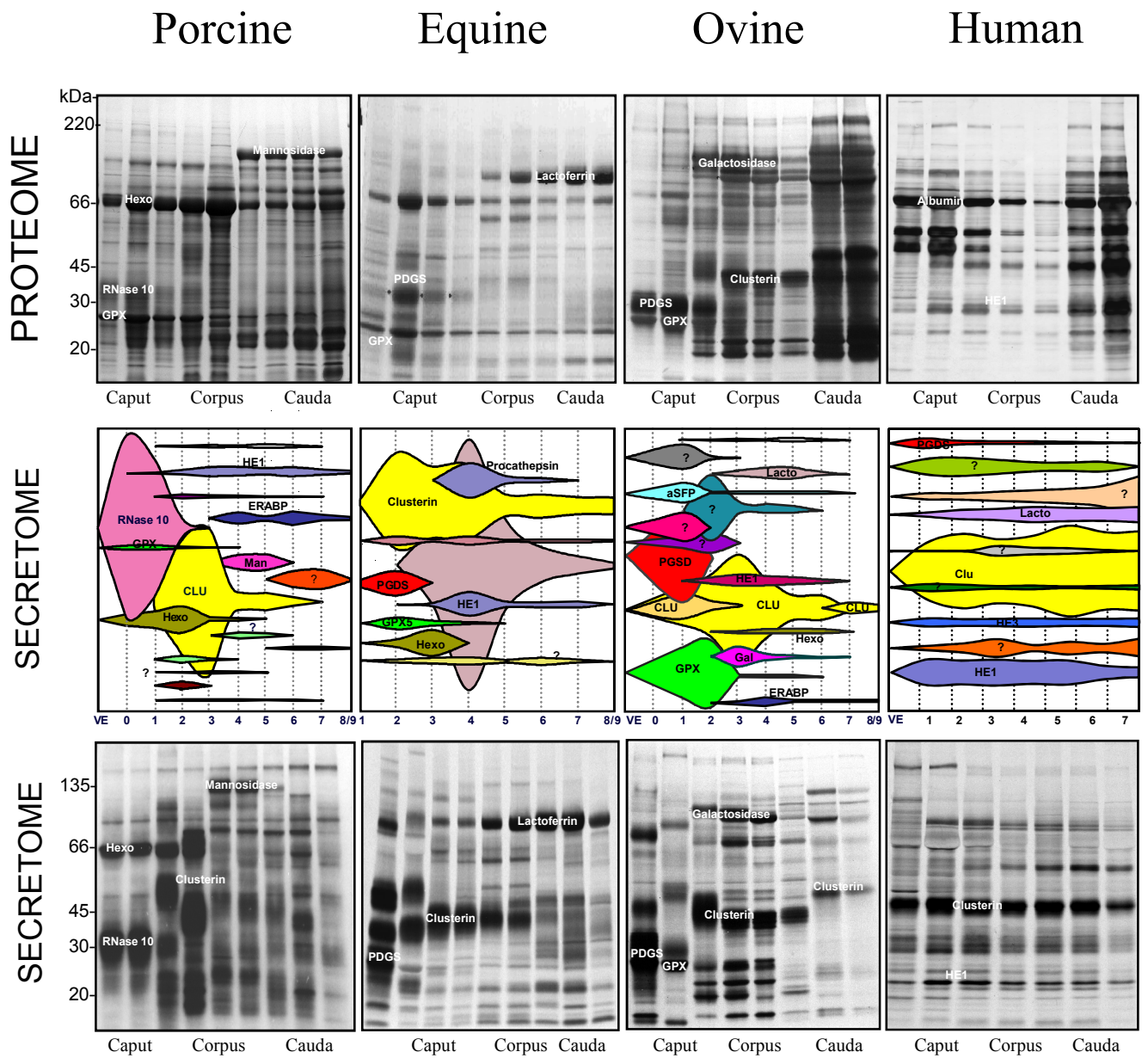


Figure 2

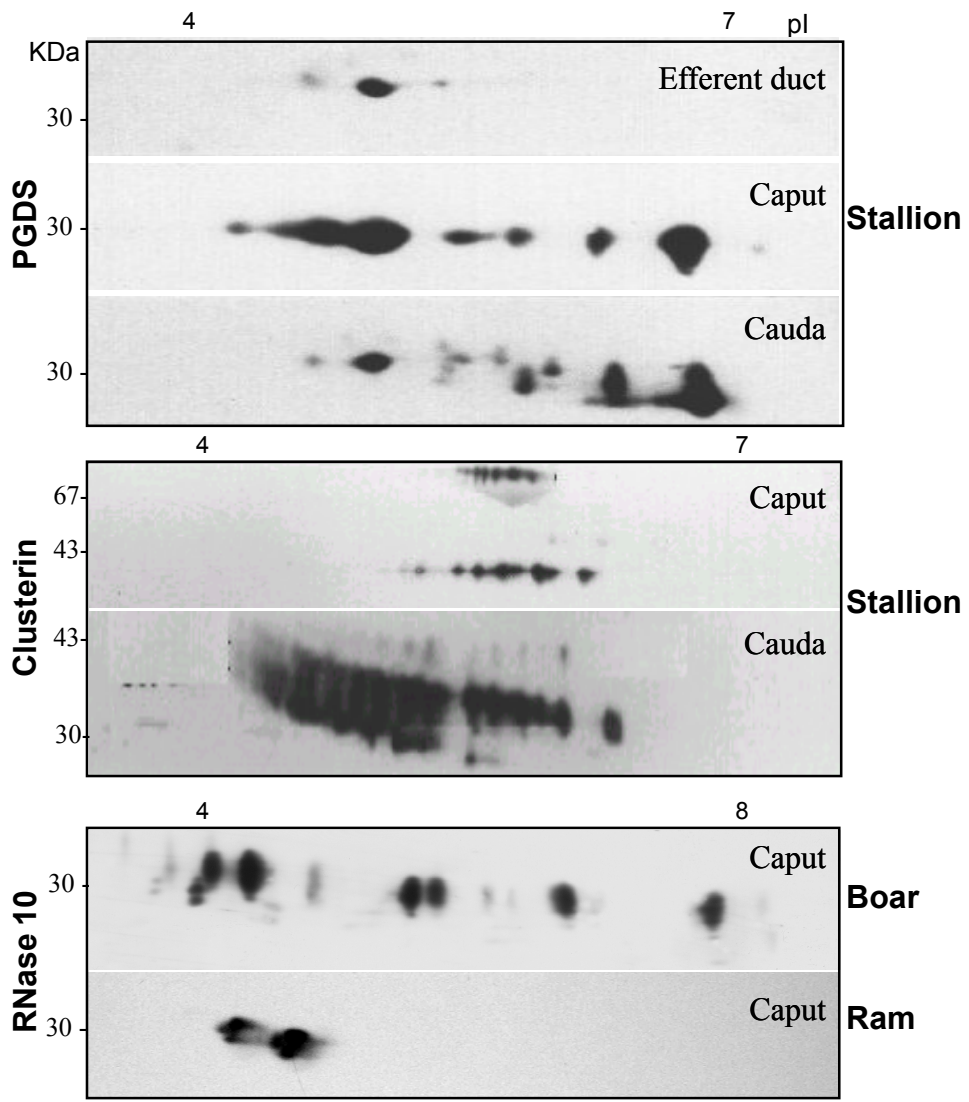
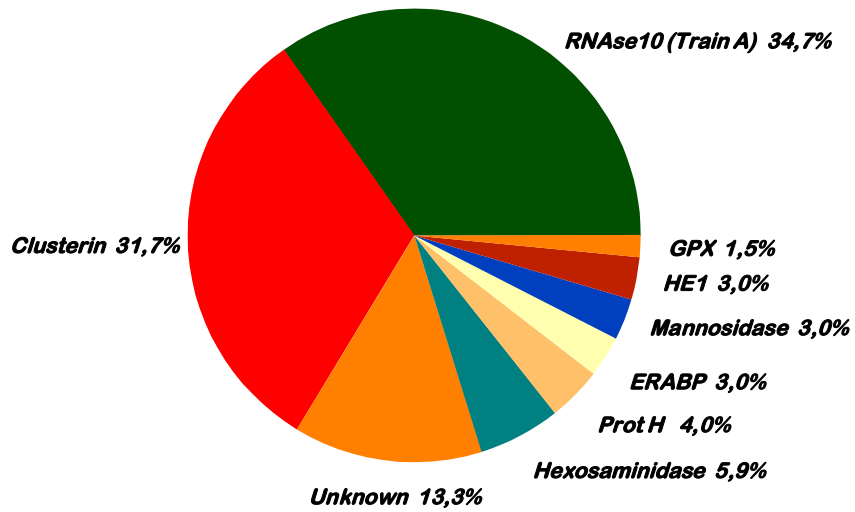
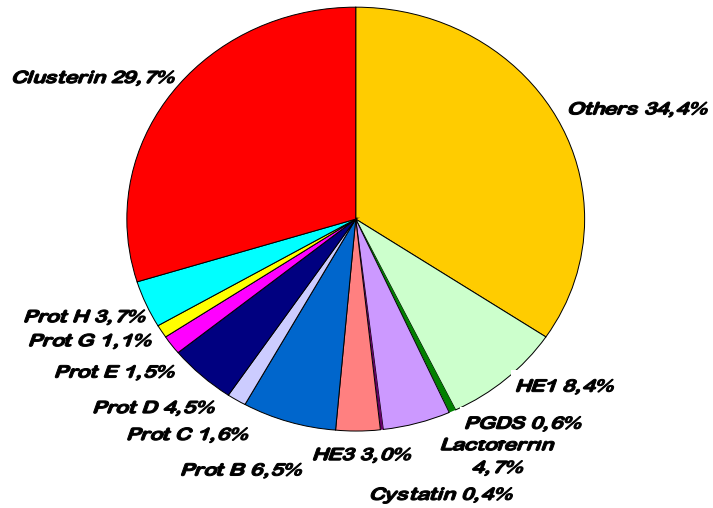


Figure 3



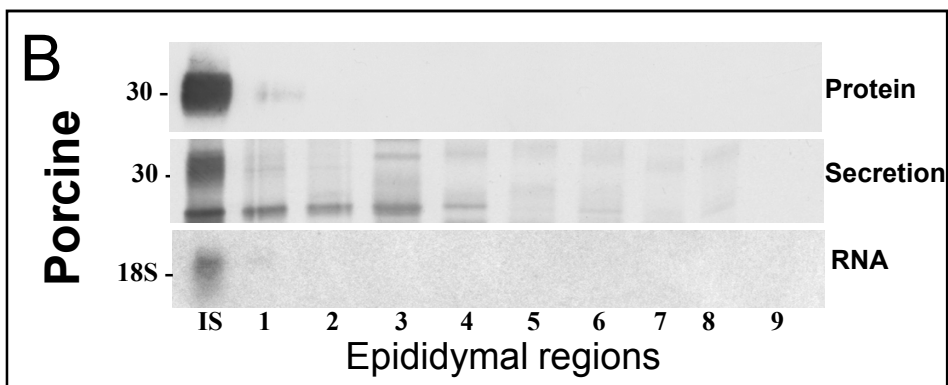
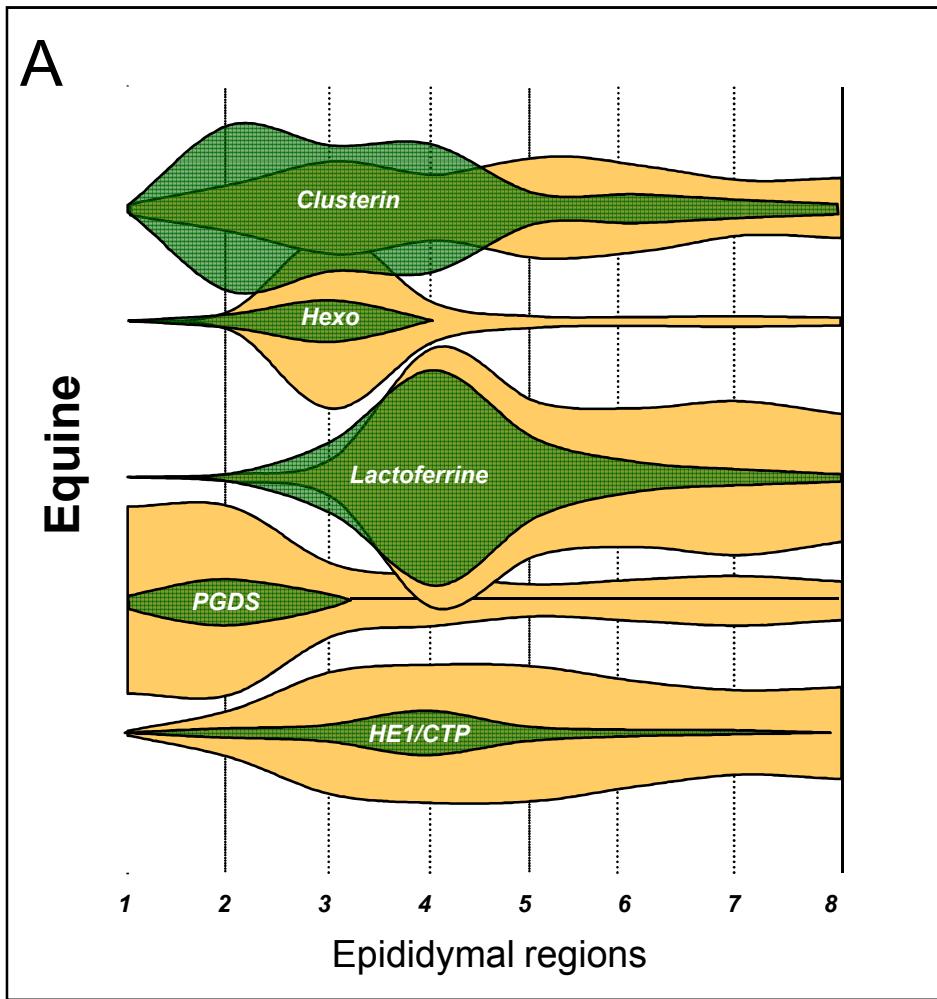


Figure 5

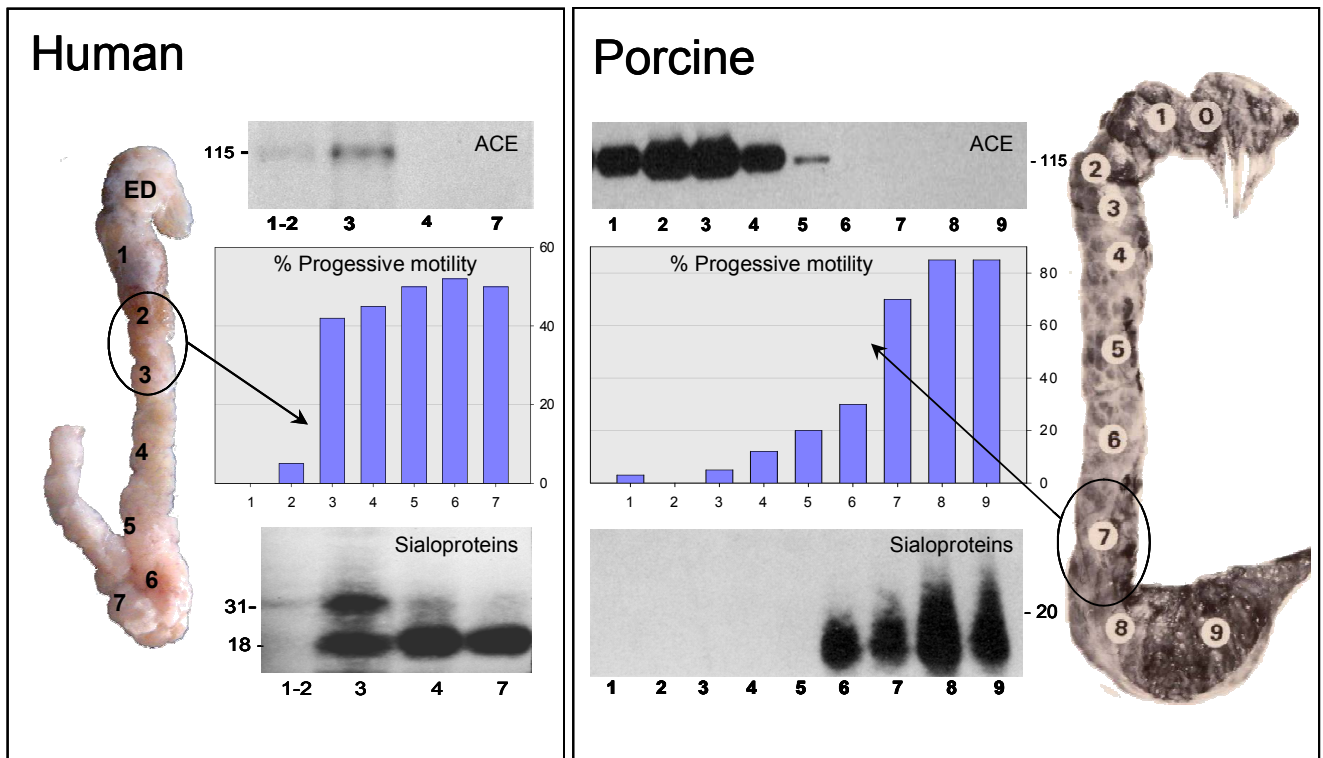


Figure 6