Loss of genes implicated in gastric function during platypus evolution

Gonzalo R Ordoñez*, LaDeana W Hillier†, Wesley C Warren†, Frank Grützner‡, Carlos López-Otín* and Xose S Puente*

Addresses: *Departamento de Bioquímica y Biología Molecular, Facultad de Medicina, Instituto Universitario de Oncología, Universidad de Oviedo, C/Fernando Bongera s/n, 33006 Oviedo, Spain. †Genome Sequencing Center, Washington University School of Medicine, Campus Box 8501, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA. ‡Discipline of Genetics, School of Molecular & Biomedical Science, The University of Adelaide, 5005 South Australia, Adelaide, Australia.

Correspondence: Xose S Puente. Email: xspuente@uniovi.es

Published: 15 May 2008

Genome Biology 2008, 9:R81 (doi:10.1186/gb-2008-9-5-r81)

The electronic version of this article is the complete one and can be found online at http://genomeweb.com/2008/9/5/R81

© 2008 Ordoñez et al; licensee BioMed Central Ltd.
This is an open access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: The duck-billed platypus (Ornithorhynchus anatinus) belongs to the mammalian subclass Prototheria, which diverged from the Theria line early in mammalian evolution. The platypus genome sequence provides a unique opportunity to illuminate some aspects of the biology and evolution of these animals.

Results: We show that several genes implicated in food digestion in the stomach have been deleted or inactivated in platypus. Comparison with other vertebrate genomes revealed that the main genes implicated in the formation and activity of gastric juice have been lost in platypus. These include the aspartyl proteases pepsinogen A and pepsinogens B/C, the hydrochloric acid secretion stimulatory hormone gastrin, and the α subunit of the gastric H⁺/K⁺-ATPase. Other genes implicated in gastric functions, such as the β subunit of the H⁺/K⁺-ATPase and the aspartyl protease cathepsin E, have been inactivated because of the acquisition of loss-of-function mutations. All of these genes are highly conserved in vertebrates, reflecting a unique pattern of evolution in the platypus genome not previously seen in other mammalian genomes.

Conclusion: The observed loss of genes involved in gastric functions might be responsible for the anatomical and physiological differences in gastrointestinal tract between monotremes and other vertebrates, including small size, lack of glands, and high pH of the monotreme stomach. This study contributes to a better understanding of the mechanisms that underlie the evolution of the platypus genome, might extend the less-is-more evolutionary model to monotremes, and provides novel insights into the importance of gene loss events during mammalian evolution.
Background

A major goal in the sequencing of different genomes is to identify the genetic changes that are responsible for the physiological differences between these organisms. In this regard, the comparison between human and rodent genomes has identified an expansion in rodents of genes that are implicated in fertilization and sperm maturation, host defense, odor perception, or detoxification [1-3], confirming at the genetic level the physiological differences in these processes between humans and rodents. Additionally, the development of specific biological processes during evolution, for example the production of milk in mammals, has been accompanied by the appearance of novel genes that are implicated in these novel functions, such as casein and α-lactalbumin [4]. Therefore, it appears that the acquisition of novel physiological functions during vertebrate evolution has been driven by the generation of novel genes adapted to these newer functions. However, although gene gains constitute an intuitive mechanism for the development of novel biological functions, gene losses have also been important during evolution, both quantitatively and qualitatively [5-9]. The recent availability of numerous vertebrate genomes has opened the possibility to perform large-scale evolutionary analysis in order to identify differential genes responsible for the specific differences in particular biological processes.

The duck-billed platypus (Ornithorhynchus anatinus) represents a valuable resource for unraveling the molecular mechanisms that have been active during mammalian evolution, due both to its phylogenetic position and to the presence of unique biological characteristics [10]. Together with the echidnas, platypus constitutes the Monotremata subclass (prototherians); this is one of the two subclasses into which mammals are divided, together with therians, which are further subdivided into marsupials (metatherians) and placentals (eutherians) [11]. The appearance of mammal-specific characteristics such as homeothermy, presence of fur, and mammary glands makes this organism a key element in elucidating the genetic factors that are implicated in the appearance of these biological functions. Nevertheless, since the last mammalian common ancestor, more than 166 million years ago (MYA) [12,13], other characteristics have emerged, such as the presence of venom glands or electroreception, and some vertebrate characteristics have been lost, resulting in the absence of adult teeth or a functional stomach [14,15].

In this work, we show that there has been a selective deletion and inactivation in the platypus genome of several genes that are implicated in the activity of the stomach, including all genes encoding pepsin proteases, which are involved in the initial digestion of proteins in the acidic pH of the stomach, as well as the genes required for the secretion of acid in this organ (Figure 1). The loss and inactivation of these genes provide a molecular basis for understanding the mechanisms that are responsible for the absence in platypus of a functional stomach, and expand our knowledge of the evolution of mammalian genomes.

Results and discussion

Loss of pepsin genes in the platypus genome

During the initial annotation and characterization of the platypus genome, we noticed the absence of several protease genes in this organism that were present in other mammalian species [2,10]. Most of these lost protease genes encode members of rapidly evolving protease families, including proteases that are implicated in immunological functions, spermatogenesis, or fertilization [2,16]. However, when we performed a further detailed analysis of all of these protease genes lost in platypus, we observed that those encoding three major gastric aspartyl proteases (pepsinogen A, pepsinogen B, and gastricsin/pepsinogen C) were also absent from the platypus genome assembly. These proteases are responsible for the proteolytic cleavage of dietary proteins at the acidic pH of the stomach, and have been highly conserved through evolution, from fish to mammals and birds [17]. The genes encoding these proteases (PGA, PGB, and PGC) are located in different chromosomal loci, whose overall structure has also been well conserved in most vertebrate genomes, including platypus (Figure 2). Therefore, it appeared unlikely that their absence in platypus could be due to the incompleteness of the genome assembly in a specific chromosomal region. Moreover, analysis of more than 2 million trace sequences not present in the assembly and expressed sequence tag (EST) sequences from different platypus tissues [10] also failed to reveal the existence of any of these pepsinogen genes, reinforcing the hypothesis that they had been specifically deleted in the genome of this mammal.

To investigate this possibility further, we first compared the genomic organization of these three aspartyl protease genes - PGA, PGB and PGC - in the genomes of human, dog, opossum, chicken, lizard, and frog [18-21]. It is well established that the genes encoding pepsinogens have undergone several expansions during vertebrate evolution, leading to the presence of at least three to six distinct functional members in the genomes of these organisms (Figure 2a). Additionally, a duplication event in PGC in the therian lineage has resulted in the formation of PGB, which appears to be functional in opossum and dog, and in the latter has probably replaced the function of PGC, which has been inactivated by pseudogenization. The loci containing these pepsinogen genes have been highly preserved through evolution, and their flanking genes are also perfectly conserved in both order and nucleotide sequence in vertebrate genomes (Figure 2a).

Analysis of platypus bacterial artificial chromosomes (BACs) and/or fosmids corresponding to these regions revealed that the genes flanking the pepsinogen genes in other species are conserved and map to the corresponding syntenic region of the platypus genome (Figure 2). However, a DNA probe cor-
responding to murine pepsinogen A failed to hybridize with the analyzed platypus BACs or fosmids spanning the regions of interest (see Additional data file 1). Moreover, complete sequencing of the platypus genomic regions flanked by TFEB and FRS3 as well as by Ctorf88 and CHIA2 failed to detect any genes encoding pepsinogen C or pepsinogen B, respectively. Additionally, and in order to test the possibility that pepsinogen genes have been transposed to other loci during platypus evolution, a Southern blot analysis with the same probe was performed using total genomic DNA. This analysis resulted in the absence of hybridization when genomic DNA from platypus and one echidna species (Tachyglossus aculeatus) were used, whereas the same probe readily detected two hybridization bands in more evolutionary distant species such as lizard (Podarcis hispanica) and chicken (data not shown).

Together, these data indicate that the genes encoding these gastric proteases have been specifically deleted in the genome of monotremes, probably resulting in important differences in the digestion of dietary proteins in these species when compared with other vertebrates.

**Loss or inactivation of platypus genes implicated in stomach acid secretion**

Pepsinogens are synthesized by chief cells in the oxyntic glands of the stomach as inactive precursors that become activated when they are exposed to the low pH of the gastric fluid [22]. The secretion of hydrochloric acid is stimulated by the gastric hormone gastrin, which is released by enteroendocrine G cells that are present in pyloric glands in response to amino acids and digested proteins. To try to extend the above findings on the absence of pepsinogen genes in platypus, we next evaluated the possibility that the gene encoding gastrin (GAST) could also be absent from the platypus genome.
After comparative genomic analysis following the same strategy as in the case of pepsinogen genes, we failed to detect any evidence of the presence of GAST in platypus (see Additional data file 1), which suggests that acid secretion might also be impaired in this species. Consistent with this observation, parallel genomic analysis also showed that the α subunit of the H⁺/K⁺-ATPase (ATP4A), which is responsible for the acidification of the stomach content by parietal cells, has also been deleted from the platypus genome. This gene, which is present from fish to amniotes, has been highly conserved through evolution but is absent from the platypus genome assembly (Figure 3a). Also similar to the case of pepsinogen genes, the ATP4A-flanking genes (TMEM147 and KIAA0841), which are present in fish, therians, and chicken, were readily identified in platypus. Thus, analysis of a fosmid clone corresponding to this region with a probe for the most proximal gene (TMEM147) resulted in detection of a specific hybridization band in platypus (see Additional data file 1). However, no hybridization bands could be detected in platypus fosmid KAAG-0404B19, or total genomic DNA from platypus and T. aculeatus when using a human derived ATP4A probe, which otherwise recognized specific bands in mouse, chicken, and lizard (Additional data file 1 and data not shown). These results extend the above findings on gastric
protease genes and demonstrate that other genes involved in the digestive activity of gastric juice have also been selectively deleted from the genomes of monotremes.

We next examined the possibility that mechanisms distinct from those involving the specific deletion of gastric genes could also contribute to the apparent loss in platypus of evolutionarily conserved digestive functions. This analysis led us to conclude that two well known gastric genes - namely CTSE and ATP4B [23-25], which encode the aspartyl protease cathepsin E and the β subunit of the H+/K+-ATPase, respectively - have been inactivated by pseudogenization. Thus, we first observed that the platypus genome contains sequences with high similarity to both gastric genes in the corresponding syntenic regions, suggesting that CTSE and ATP4B could indeed be functional genes in platypus. However, further detailed analysis of their nucleotide sequence revealed that CTSE is nonfunctional in this species due both to the presence of a premature stop codon in exon 7 (Lys295Ter) and to the loss of six of its nine exons. Similarly, the gene encoding ATP4B has been pseudogenized in platypus because of the presence of premature stop codons in exons 3, 4, and 7 showing the presence of premature stop codons and a frameshift (red arrow). MYA, million years ago.

Loss of gastric genes during platypus evolution

The mammalian stomach is lined with a glandular epithelium that contains four major cell types [26]: mucous, parietal, chief, and enteroendocrine cells. The data presented above show that the genes encoding different products of these four major cell types of the gastric glandular epithelium have been selectively deleted or inactivated during monotreme evolution (Figure 1 and Table 1). Although the genes encoding proteases have been shown to be subjected to processes of gene gain/loss events in both vertebrate and invertebrate genomes [5,16,27], we have determined that these gene loss events observed in platypus gastric genes do not represent a general process affecting all proteins that are involved in food digestion, because analysis of genes implicated in gastrointestinal functions revealed that those encoding proteases and hormones expressed in the intestine or exocrine pancreas from eutherians are perfectly conserved in platypus (Figure 1). It therefore appears that there has been a selective loss of platypus genes responsible for the biological activity of gastric juice.

To address this question further, we next performed a detailed search for the putative occurrence in the platypus genome of functional genes encoding proteins secreted by gastric glands. This search led us to the identification of two genes with interesting characteristics in this regard. The gene encoding gastric intrinsic factor (GIF), which is necessary for the absorption of vitamin B12, is perfectly conserved in platypus. This protein is secreted by chief or parietal cells in most eutherians, but it is mainly produced by pancreatic cells in dogs as well as in opossum, in which no gastric expression can be detected [28,29]. It is therefore likely that the expression...
of this gene was pancreatic before the prototherian-therian split, and the intrinsic factor might still be secreted by the pancreas in platypus, where it can exert its physiological function.

To investigate this possibility, we conducted RT-PCR analysis using specific primers for GIF and RNA from different tissues from either platypus or echidna (T. aculeatus). This allowed us to find that GIF expression can be detected in pancreas, and lower expression could be also detected in liver as well as in echidna brain, whereas no expression was detected in muscle or brain from platypus (see Additional data file 2). Therefore, these findings indicate that, similar to the case of marsupials, the GIF gene is also expressed by the pancreas in monotremes. A similar situation could occur in the case of chymosin, an aspartyl protease that participates in milk clotting by limited proteolysis of κ casein [30]. Chymosin is present in chicken and in most mammalian species, although it has been inactivated by pseudogenization in humans and other primates [2,31]. Our genomic analysis also detected a gene containing a complete open reading frame that might constitute a functional chymosin gene in the platypus genome. This finding, together with the absence of soluble pepsins and cathepsin E in platypus, suggests that chymosin might be the only aspartyl protease with ability to contribute to food digestion in the stomach of platypus. Nevertheless, it is very unlikely that chymosin could compensate for the lack of pepsin activity in platypus stomach because of its much lower proteolytic activity when compared with that of pepsins [30]. Additionally, the high pH of platypus stomach might prevent the zymogen activation and proteolytic activity of this peptidase. Finally, it is possible that, similar to the case of the intrinsic factor, platypus chymosin might be also produced by other tissues. In this regard, we have been unable to detect the expression of this gene in any of the tissues analyzed above (data not shown), although its putative participation in the digestion of dietary proteins should be further characterized.

The loss of stomach function in prototherians is unique among vertebrates, because this organ has been functional for more than 400 million years, from fish to therians and birds, and it has been adapted to specific dietary habits, resulting in the formation of multiple chambers in birds and ruminants [32]. In contrast, the stomach of platypus is completely aglindrical and has been reduced to a simple dilatation of the lower esophagus [14,15]. It is remarkable that some fish species such as zebrafish (Danio rerio) and pufferfish (Takifugu rubripes) have also lost their gastric glands during evolution, although this fact has not apparently resulted in the loss of so many gastric genes in these teleosts as in platypus [33,34]. On the other hand, the small stomach, high pH of gastric fluid, and lack of gastric glands in echidna, together with the finding that some of the gastric genes lost in platypus are also absent in T. aculeatus, suggest that the loss of the stomach function and gastric genes in monotremes occurred before the platypus-echidna split, more than 21 MYA [10]. However, it is difficult to determine whether the loss of gastric genes in platypus has conferred a selective advantage during evolution, or whether they have been lost as a result of a relaxed constraint due to additional changes in this species.

In this regard, it is possible that the loss of gastric genes in monotremes might have conferred a selective advantage to this population against parasites or pathogens that rely on the presence of an acidic pH in the stomach for their infection or propagation, or the use of cell surface proteins such as ATP4A, ATP4B, or CTSE as receptors for the infection. Should this be the case, then this would represent a clear example of the 'less-is-more' hypothesis [35,36], which postulates that the loss of a gene might confer a selective advantage under specific conditions. Nevertheless, in the absence of additional data, it cannot be ruled out that additional changes in the digestive system of monotremes made irrelevant the function of the genes described in this work, and they were subjected to the accumulation of deleterious mutations because of a relaxed constraint. However, an interesting question at this point is whether additional strategies have

### Table 1

| Protein                                      | Gene   | Status in platypus genome | Confirmatory evidence |
|----------------------------------------------|--------|---------------------------|-----------------------|
| ATPase, H⁺/K⁺ exchanging, α polypeptide      | ATP4A  | Absent                    | Southern blot         |
| ATPase, H⁺/K⁺ exchanging, β polypeptide      | ATP4B  | Pseudogene                | PCR/direct sequencing |
| Cathepsin E                                  | CTSE   | Pseudogene                | PCR/direct sequencing |
| Gastrin                                      | GAST   | Absent                    | Southern blot         |
| Neurogenin 3                                 | NGN3   | Absent                    | Southern blot         |
| Pepsin A                                     | PGA    | Absent                    | Southern blot/sequencing |
| Pepsin C                                     | PGC    | Absent                    | Southern blot/sequencing |
| Gastric intrinsic factor                     | GIF    | Present (expression pancreatic) | RT-PCR |
| Chymosin                                     | CYMP   | Present (expression not detected) | Sequencing/RT-PCR |

RT-PCR, reverse transcription polymerase chain reaction.

Summary of genes implicated in gastric function in platypus.
been adopted by platypus to accomplish efficient protein digestion in the absence of a number of gastric enzymes. Changes in dietary habits, such as feeding on insect larvae, which are easily digested; the presence of specific anatomical structures, such as grinding plates or cheek-pouches, which allow food trituration and storage; and the putative occurrence of a characteristic gastrointestinal flora in platypus might constitute mechanisms by which this species has overcome the loss of a functional stomach.

Another question raised by this comparative genome analysis is whether the loss of all of the above discussed genes is cause or consequence of this particular platypus gastric phenotype. Deletion of the gene encoding gastrin might have contributed to this process, because mice deficient in gastrin exhibit an atrophy of the oxyntic mucosa, with a reduced number of parietal and enteroendocrine cells, achlorhydria, and decreased mucosa thickness [37-39]. Additionally, inactivation of ATP4B has been shown to produce a significant decrease in pepsin-producing chief cells and alterations in the structure of parietal cells [25]. Moreover, loss of PGA might also contribute to the gastric atrophy observed in platypus, because this protease was recently shown to be required for the processing and activation of the morphogen sonic hedgehog (Shh) in the stomach [40]. Therefore, deletion or inactivation of gastrin, the acid-secreting ATPase, and pepsinogen A could have contributed to a substantial reduction in the formation of gastric glands in monotremes. Nevertheless, we cannot discard the possibility that the stomach function was lost by some other unrelated mechanism, and - in the absence of a selective pressure to maintain the genes encoding proteins implicated in the gastric function - these genes were lost by pseudogenization and/or deletion events. However, the exclusive absence of these genes cannot explain the significant reduction in size observed in the stomach of platypus, suggesting that other factors might be responsible for this characteristic feature.

To evaluate this possibility, we first selected a series of genes previously described to influence stomach size in mice and examined its putative presence and sequence conservation in the platypus genome (Additional data file 3). This analysis allowed us to determine that the gene encoding neurogenin-3 has been lost in platypus (Additional data file 1 and Table 1).

Neurogenin-3 is a transcription factor whose activity is required for the specification of gastric epithelial cell identity, and deficiency of this factor results in considerably smaller stomachs and absence of gastrin-secreting G cells, somatostatin-secreting D cells and glucagon-secreting A cells [41]. Therefore, it is tempting to speculate that neurogenin-3 could be a candidate gene to explain, at least in part, the morphological differences between platypus stomach and that of other vertebrates. Nevertheless, further studies of the role of neurogenin-3 in different species will be required to ascribe a role to this transcription factor in defining structural or functional differences in stomach during mammalian evolution.

**Mechanisms involved in the loss of gastric genes in platypus**

Finally, in this work we have also examined putative mechanisms responsible for the loss of gastric genes in the platypus genome. A first possibility in this regard should be the occurrence of directed gene losses specifically occurring in platypus and the two extant echidna species Zaglossus and Tachyglossus. As a first step in this analysis, and based on recent studies of specific gene losses during hominoid evolution [42], we examined the hypothesis that gastric genes were independently deleted in platypus by nonallelic homologous recombination or by insertion of repetitive sequences. Consistent with this possibility, and in agreement with the increased activity of interspersed elements in the platypus genome [10,43], we have found that the CTSE gene has been disrupted in platypus by the insertion of long interspersed elements (LINEs) and short interspersed elements (SINEs) in exons 7 and 9, disrupting the protein coding region (Figure 4). Interestingly, exon 9 was disrupted by the insertion of a LINE2 Platym element, which was further disrupted by the insertion of a SINE Mon1f3 element (Figure 4). In this regard, analysis of different interspersed elements in the platypus genome has revealed that the main period of activity of Mon1f3 elements was between 88 and 159 MYA [10], indicating that pseudogenization of CTSE might have occurred within this period, and suggesting that the inactivation of gastric genes in monotremes started at least 88 MYA. Furthermore, the high abundance of repetitive elements in the CTSE region (more than 3.8 interspersed elements per kilobase as compared with 2 for the genome average [10]) might have contributed to the deletion of six out of the nine exons of CTSE by nonallelic homologous recombination between these repetitive elements. The variable density of interspersed elements in the regions examined in this study raises the possibility that similar mechanisms to that observed in CTSE might have been responsible for the complete deletion of other gastric genes, although the participation of other mechanisms in this process cannot be ruled out.

**Conclusion**

In summary, detailed analysis of the platypus genome sequence has allowed us to demonstrate that a number of genes that are implicated in food digestion in the stomach have specifically been deleted or inactivated in this species, as well as in echidna. It is remarkable that the results presented here may constitute an exceptional example of the less-is-more evolutionary model [35,36], both for the number of genes involved as well as for the physiological consequences derived from these genetic losses. In fact, the loss of the gastric genes reported in this study appears to be responsible for the specific characteristics of the platypus gastrointestinal system, although it cannot be ruled out that the loss of the
stomach by other unrelated events might have resulted in the neutral evolution of these genes. The gastric genes lost in the platypus genome include those encoding the aspartyl proteases pepsinogen A, pepsinogens B/C and cathepsin E, the hydrochloric acid secretion stimulatory hormone gastrin, and both subunits of the gastric H⁺/K⁺-ATPase. Likewise, genes encoding proteins implicated in stomach development, such as the neurogenin-3 transcription factor, are also absent in the platypus genome. All of these genes have been highly conserved in vertebrates for more than 400 million years, reflecting a unique pattern of evolution in the platypus genome when compared with other mammalian genomes. On the basis of these findings, we propose that loss of genes involved in gastric functions might be responsible for the remarkable anatomical and physiological differences of the gastrointestinal tract between monotremes and other vertebrates, and underscores the importance of gene loss for mammalian evolution.

Materials and methods

Bioinformatic analysis

The identification of protease-coding genes in the platypus genome was carried out as previously described [27], using a 6X assembly (version 5.0) generated with the PCAP assembly program, with an estimated coverage of 90% to 93% [10]. Briefly, protein sequences corresponding to human proteases were searched in the platypus assembly using the TBLASTN algorithm with an expected threshold of 10. In most cases this was sufficient to identify individual contigs containing exons with high sequence identity to the queried protease, which were further analyzed to obtain the full-length coding sequence. In those cases in which no clear ortholog was found in the platypus genome assembly, the following procedure was used. First, the traces and the EST sequences were analyzed using BLASTN and TBLASTN, increasing the expected threshold up to 1,000, which was sufficient to detect the orthologous genes in the assembly and traces of more evolutionary distant vertebrates such as lizard, chicken, or frog. Second, to exclude the possibility that these results arose simply because the human gene was too divergent from the platypus one, the query sequence was replaced by the corresponding ortholog in mouse, dog, opossum, chicken, lizard, frog, or fish (when available), and the search was performed in the platypus assembly, traces, and ESTs using BLASTN and TBLASTN. Third, if the previous strategies failed, then the 5'- and 3'-flanking genes in other vertebrate genomes were used as query to identify platypus contigs corresponding to the locus in which the candidate gene was supposed to lie. These contigs were then searched with the TBLASTN algorithm with increasing expected threshold to identify potential exons of the gene or pseudogene, and the contigs were analyzed for the presence of large gaps. When large gaps were found, BACs and/or fosmids corresponding to those regions were obtained and analyzed by Southern blot and/or sequencing.

Southern blot and sequencing

Platypus BACs were obtained from Children’s Hospital Oakland Research Institute, and fosmids and genomic DNA were provided by the platypus genome sequencing project [10]. DNA was digested with the indicated enzymes, separated in a 0.7% agarose gel, and transferred to a nylon membrane. Southern blot hybridization was performed using specific oli-
gonucleotides corresponding to platypus genes present in the assembly (Additional data file 4) or using human or mouse-derived cDNA probes for ATP4A (corresponding to nucleotides 1,899 to 2,503 of sequence NM_000704), PGA (corresponding to nucleotides 867 to 1,259 of sequence NM_021453), and NGN3 (corresponding to nucleotides 387 to 593 of sequence NM_020999). DNA probes were PCR-amplified using Taq Platinum (Invitrogen, Carlsbad, CA) and purified. All PCRs were performed in a Veriti 96-well thermal cycler (Applied Biosystems, Foster City, CA) for 35 cycles of denaturation (95°C for 15 seconds), annealing (60°C for 15 seconds), and extension (72°C for 30 seconds). Double-stranded DNA probes were radiolabeled with [α-32P]dCTP (3,000 Ci/mmole) from GE Healthcare (Uppsala, Sweden), using a commercial random priming kit purchased from the same company. When specific oligonucleotides were used for hybridization, they were labeled with [γ-32P]ATP (3,000 Ci/mmole) from GE Healthcare using T4 Polynucleotide Kinase (USB, Cleveland, OH). Hybridization was performed at 42°C or 60°C for oligonucleotides or cDNA probes, respectively, using a Rapid-Hyb hybridization solution (GE Healthcare).

Additionally, the regions corresponding to the PGC and PGB loci in platypus were cloned from the indicated BACs and fosmids, and subjected to direct sequencing using the kit DR terminator TaqFS and the automatic DNA sequencer ABI-PRISM 310 (Applied Biosystems), with specific oligonucleotides as primers. Mutations in gastric genes were confirmed by amplification of the corresponding exons with specific primers (Additional data file 4) using platypus genomic DNA as template, and the amplified product was subjected to nucleotide sequencing.

Analysis of GIF expression in platypus and echidna tissues
Total RNA from platypus and echidna (T. aculeatus) tissues was reverse-transcribed using oligo-dT and the RNA-PCR Core kit from Perkin Elmer Life Sciences (Foster City, CA) and subjected to PCR amplification using specific primers for GIF (5'-TGCGCTCTGACCTGTATGTA and 5'-GGTTTG-GCTTTCCAGG GAAGG) and GAPDH (5'-AAGGCTGT-GGGCAAGGTCAT and 5'-CTGTGGAATGCACAGGAC).

Abbreviations
BAC, bacterial artificial chromosome; EST, expressed sequence tag; LINE, long interspersed element; MYA, million years ago; RT-PCR, reverse transcription polymerase chain reaction; SINE, short interspersed element.

Authors’ contributions
GRO, CLO, and XSP conceived of the study, carried out the data analysis and interpretation, and contributed to the writing of the manuscript. LWH and WCW performed the analysis of BAC and Fosmid ends, and provided individual clones for the indicated loci. FG provided platypus and echidna samples. All authors read and approved the final manuscript.

Additional data files
The following additional data files are available. Additional data file 1 is a figure showing the following: Southern blot analysis of platypus fosmids KAAG-0287H03, KAAG-0109P06, and BAC KAAG-711F22; synteny map of the gastrin locus in the indicated species; synteny map of the ATP4A locus in different vertebrates and platypus fosmid KAAG-0404B19 corresponding to this region. Additional data file 2 is a figure showing the analysis of GIF expression in platypus and echidna tissues. Additional data file 3 is a table listing genes implicated in stomach size and development and their status in the platypus genome. Additional data file 4 is a table listing the oligonucleotides used for amplification, sequencing, and hybridization of the indicated platypus genes.

Acknowledgements
We thank T Graves for help with fosmid clones; A Fuego, V Quesada, and A Smit for helpful discussions; and F Rodríguez for technical assistance. This work was supported by grants from the European Union (CancerDegrome-FP6), Ministerio de Educación y Ciencia-Spain, Ministerio de Sanidad-Spain; Fundación La Caixa, Fundación M Botín, Fundación Lilly, and Ramón y Cajal Program (XSP). The Instituto Universitario de Oncología is supported by Obra Social Cajastur.

References
1. Gibbs RA, Weinstock GM, Metzker ML, Muzny DM, Sodergren EJ, Scherer S, Scott G, Steffen D, Worley KC, Burch PE, Owuomo G, Hines S, Lewis L, DeRamo C, Delgado O, Dugan-Rocha S, Miner G, Morgan M, Hawes A, Gill R, Celera, Holt RA, Adams MD, Amanatides PG, Baden-Tillson H, Barnstead M, Chin S, Evans CA, Ferreria S, Fusler C, et al.: Genome sequence of the Brown Norway rat yields insights into mammalian evolution. Nature 2004, 428:493-521.
2. Puente XS, Sánchez LM, Overall CM, López-Otín C: Human and mouse proteases: a comparative genomic approach. Nat Rev Genet 2003, 4:544-558.
3. Godfrey PA, Malnic B, Buck LB: The mouse olfactory receptor gene family. Proc Natl Acad Sci USA 2004, 101:2156-2161.
4. Kawasaki K, Weiss KM: Mineralized tissue and vertebrae evolution: the secretory calcium-binding phosphoprotein gene cluster. Proc Natl Acad Sci USA 2003, 100:4060-4065.
5. Hahn MW, Han MV, Han SG: Gene family evolution across 12 Drosophila genomes. PLoS Genet 2007, 3:e197.
6. Smedman HH, Kozjak BW, Nelson A, Theiler DM, Su LT, Low DW, Bridges CR, Shragger JB, Minugh-Purvis N, Mitchell MA: Myosin gene mutation correlates with anatomical changes in the human lineage. Nature 2004, 428:415-418.
7. Krylov DM, Wolf YI, Rogozin IB, Koonin EV: Gene loss, protein sequence divergence, gene dispensability, expression evolution, and interactivity are correlated in euarchontian evolution. Genome Res 2003, 13:2229-2235.
8. Blomme T, Vandepoele K, De Bodt S, Simillion C, Maere S, Peer Y Van de: The gain and loss of genes during 600 million years of vertebrate evolution. Genome Biol 2006, 7:R43.
9. Wang X, Grus WE, Zhang J: Gene losses during human origins. PLoS Biol 2006, 4:e52.
10. Warren WC, Hillier LW, Marshall-Graves JA, Birney E, Ponting CP, Grutzner F, Belov K, Miller W, Clarke L, Chinwalla AT, Yang SP, Heager A, Clarke D, Mietheke P, Waters P, Veyrunes F, Fulton L, Graves T, Puente XS, López-Otín C, Ordoñez GR, Eichler EE, Deakin
JE, Thompson K, Kirby P, Papenfuss AT, Wakefield M, Olender T, Lambert J, Zamecnik P, Lander ES, Gaffney PT, Fuentes C, et al. Gene regulation in mammalian placental development. Science 2004, 304:1364-1368.

27. Fong FY, Yang X, Maniatis T: A novel retrotransposon in mouse brain. PLoS Biol 2005, 3:e54.

28. Van de Water K, Wilkens LF, Bicknell B, et al. Differentiation of colon epithelial cells in 3D culture. Dev Biol 2006, 293:27-41.

29. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

30. Kageyama R, Nakamura M, et al. Gene expression change in mouse brain. Science 2005, 309:1587-1591.

31. Kageyama R, Nakamura M, et al. Gene expression change in mouse brain. Science 2005, 309:1587-1591.

32. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

33. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

34. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

35. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

36. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

37. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

38. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

39. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

40. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

41. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

42. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

43. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

44. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

45. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

46. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

47. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

48. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

49. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

50. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

51. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

52. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

53. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

54. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

55. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

56. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

57. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

58. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

59. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

60. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

61. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

62. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

63. Watanabe Y, Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.

64. Watanabe Y,Komatsu A, et al. Cell cycle regulation in type 1 diabetes. Proc Natl Acad Sci USA 2005, 102:996-1001.
regulate multiple aspects of gastrointestinal development. 
*Development* 2000, 127:2763-2772.

53. Sock E, Rettig SD, Enderich J, Bosl MR, Tamm ER, Wegner M: Gene targeting reveals a widespread role for the high-mobility-group transcription factor Sox11 in tissue remodeling. *Mol Cell Biol* 2004, 24:6635-6644.

54. Kanai-Azuma M, Kanai Y, Gad JM, Tajima Y, Taya C, Kurohmaru M, Sanai Y, Yonekawa H, Yazaki K, Tam PP, Hayashi Y: Depletion of definitive gut endoderm in Sox17-null mutant mice. *Development* 2002, 129:2367-2379.

55. Chiang MK, Liao YC, Kuwabara Y, Lo SH: Inactivation of tensin3 in mice results in growth retardation and postnatal lethality. *Dev Biol* 2005, 279:368-377.