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Regulatory activities of transposable elements: from conflicts to benefits

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Abstract

Transposable elements (TEs) are a prolific source of tightly regulated, biochemically active non-coding elements, such as transcription factor binding sites and non-coding RNAs. A wealth of recent studies reinvigorates the idea that these elements are pervasively co-opted for the regulation of host genes. We argue that the inherent genetic properties of TEs and conflicting relationships with their hosts facilitate their recruitment for regulatory functions in diverse genomes. We review recent findings supporting the long-standing hypothesis that the waves of TE invasions endured by organisms for eons have catalyzed the evolution of gene regulatory networks. We also discuss the challenges of dissecting and interpreting the phenotypic impact of regulatory activities encoded by TEs in health and disease.

Graphical abstract

Table of contents blurb: Transposable elements (TEs) are widely known for the pathological consequences of selfish propagation and mutagenesis. However, as described in this Review, TEs also provide hosts with rich, beneficial gene-regulatory machinery in the form of regulatory DNA elements and TE-derived gene products. The authors highlight the diverse regulatory contributions of TEs to organismal physiology and pathology, provide a framework for responsibly assigning functional roles to TEs and offer visions for the future.

Introduction

Transposable elements are ubiquitous in eukaryotic genomes, and persist through independent replication of their sequences. The idea that TEs play a fundamental role in the evolution of eukaryotic gene regulation reaches back 75 years to the seminal work of Barbara McClintock on ‘controlling elements’ of maize. She regarded these elements as “normal components of the chromosome responsible for controlling, differentially, the time

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and type of activity of individual genes”¹. McClintock's perspective, although scorned by some of her contemporaries, was further developed by other pioneers, most notably Britten and Davidson in the late 1960s. Building on early insights into the complex repetitive nature of eukaryotic genomes², which they correctly attributed to transposition activity, Britten and Davidson envisioned a model where the amplification of diverse repeat families in the genome could spread ‘pre-built’ regulatory elements to drive the evolution of gene regulatory networks³.

Half-a-century forward, we now appreciate that the movement and accumulation of TEs in genomes may be solely explained by their ‘selfish’ replication activities^{4,5} and other non-adaptive forces acting at the level of the host population, such as genetic drift [G], inexorably shaping genome architecture⁶. Many studies have documented the disruptive, and often deleterious impact of these activities, as well as the more ‘constructive’ influence of TEs in the evolution of chromosome structure and gene content. But to what extent the pervasive colonization of genomes by TEs has impacted the evolution of eukaryotic gene regulation remains a matter of speculation and controversy. At the heart of the debate lies a series of recent large-scale analyses of the genetic regulatory landscape of mammalian cells revealing the engagement of an unexpectedly large fraction of TE sequences in a wide range of regulatory processes and molecular interactions. These observations, now reported for a diverse range of organisms, have rejuvenated some of the original ideas proposed by McClintock, Britten and Davidson and repositioned transposition as a potent mechanism underlying the evolution of transcriptional gene networks in eukaryotes.

In this Review, we consider emerging evidence revealing TEs as a genome-wide source of regulatory elements. We also discuss recent advances in our ability to experimentally capture the regulatory activities of TEs, with a primary focus on their contribution as *cis*-regulatory DNA elements. We argue that most of these regulatory activities can be interpreted as relics of strategies used by TEs to spread within genomes and host populations. This simple view bypasses the need to evoke widespread adaptive consequences for the host, while recognizing the occasional long-term repercussions of TEs on genome evolution. We weigh current challenges in deciphering the biological impact and evolutionary relevance of TE-derived regulatory activities, drawing examples mostly from studies performed in mammalian species, but also other eukaryotes including plants and insects. Finally, we propose how unbalanced control of TE-derived functions might result in transcriptional misregulation promoting disease states.

What predisposes TEs to *cis*-regulatory activity?

Evolutionary origins of TE *cis*-regulatory activities

Autonomous TEs encode genes that promote their replication independently of that of host chromosomes. However, as genomic parasites, TEs rely on host cell machinery to express their genes. Thus replication-competent TEs have evolved *cis*-regulatory sequences that function to mimic host promoters (Fig 1a). In long terminal repeat (LTR) elements [G], such as those of endogenous retroviruses (ERVs), *cis*-regulatory sequences [G] and RNA polymerase II (Pol II) promoters are present in duplicate within each of the two LTRs flanking the coding sequence of the elements⁸. By contrast, full-length long interspersed

nuclear elements [G] (LINEs) possess an internal Pol II promoter located in their 5' untranslated region (UTR)⁹ as well as an antisense promoter¹⁰. Other TEs collectively referred to as short interspersed nuclear elements [G] (SINEs) are derived from cellular genes transcribed by Pol III (e.g. tRNA or 7SL RNA). Thus, an intact, full-length SINE copy contains internal sequence motifs (A and B boxes) that are capable of recruiting Pol III⁹.

The diversity of TEs in their abundance, form, and replication mechanism greatly impacts the fate of the promoters and *cis*-regulatory elements they carry. For instance, the mechanism of LTR element replication dictates that each new insertion will introduce two exact copies of the LTR in the host genome⁸. However, following insertion, these elements often undergo ectopic recombination between their LTRs. These events result in the removal of the coding regions of these elements, but leave intact solitary LTRs, which contain the original *cis*-regulatory sequences. ERVs occupy ~8% of the human genome but 90% exist as solitary LTRs¹¹. By contrast, LINEs account for a larger fraction of the human genome (~20%), but the vast majority suffered 5' truncations upon insertion that removed their promoter sequences¹¹. Similarly, DNA transposons [G], which generally transpose via a cut-and-paste mechanism, are mostly propagated as miniature inverted repeat TEs (MITEs), which arise from internal deletion derivatives of autonomous elements¹². Predictably, many MITEs lack the promoter sequences of their parental element. In summary, not all TE insertions are 'created equal' in their ability to retain their original promoters and *cis*-regulatory elements (see also Fig 1a).

As for canonical host gene promoters, TE promoters often show spatially or temporally regulated activity that is dependent on cell type, or in response to environmental clues such as stress or infection^{13,14}. Although relatively few TE promoters have been extensively characterized, it is known that a variety of TEs exhibit regulated patterns of expression that is established by clusters of *cis*-regulatory sequences. Much like canonical promoters and enhancers, these sequences can recruit and 'integrate' specific combinations of host-encoded transcription factors^{9,15-22}.

What evolutionary forces shaped the regulatory features of TE promoters? For new TE insertions to persist through vertical inheritance, transposition events must occur in the germline or prior to germline development, which results in strong selection for TEs to be transcriptionally active in the germline. Indeed, the expression of many TEs appears restricted to various stages of gametogenesis or early embryogenesis in both plants²³ and animals²⁴ (see Box 1). Paradoxically, some TEs also exhibit tightly regulated activity in somatic tissues in a variety of organisms²³⁻²⁶. Given that somatic transposition events are not trans-generationally inherited, activity in these tissues holds no immediately apparent benefit or consequence to the TE. It may be tempting to speculate that regulated somatic TE activity may reflect a symbiotic relationship between host and TE, but an alternative, yet not mutually exclusive explanation is that these somatic regulatory activities have been previously moulded by selection on the TE. For example, all ERVs originate from infectious retroviruses that must have infected germ cells (or their progenitors) but possibly also somatic cells from diverse tissues. The past viral lifestyle of ERVs may thus explain why their LTRs harbour complex regulatory elements that are capable of driving transcription in a variety of tissues and cell types. Regardless of the evolutionary forces driving somatic

activity, it is clear that TEs have the potential to impact host gene regulation well beyond the germline and early embryo.

Non-random integration of TEs in the genome

As TEs evolved regulatory sequences that essentially mimic host *cis*-regulatory elements, it naturally follows that a TE insertion that lands in the vicinity of a host gene has a strong potential to interfere with its expression. Thus, where a TE initially inserts in the genome will often dictate its fate in the population. Accordingly, many TEs appear to have evolved mechanisms that favour integration into genomic regions that maximize their chance of propagation. For instance, some TE families have evolved sophisticated molecular mechanisms to target genomic 'safe havens', such as gene-poor or heterochromatic regions, whereas others favour integration within areas of open, transcriptionally active chromatin²⁷. For instance, *Mutator* elements in maize²⁸, *mPing* MITEs in rice²⁹, Tf1 retroelements in fission yeast³⁰, and P elements in *Drosophila*³¹ all actively target the 5' region of genes. Some retroviruses, including HIV-1, preferentially integrate within highly transcribed regions of the genome³². Presumably, this pattern of propagation ensures that newly integrated elements reside in a favourable environment for expression and transmission. Such elements are likely to have a stronger propensity for modulating adjacent gene expression and, if tolerated by natural selection, may be more prone to *cis*-regulatory co-option — possibly immediately upon insertion²⁹. Finally, it is important to note that as time elapses after TE propagation, forces such as selection and genetic drift are likely to obscure initial integration preferences.

TEs are a rich source of regulatory activities

What is the impact of the widespread dispersal of *cis*-regulatory elements by TEs on genome regulation and cell function? To cope with the parasitic burden of TEs in their genomes, eukaryotes appear to have evolved multi-layered mechanisms to prevent TE transcription^{23,33,34}. Consequently, the bulk of TE-derived DNA is generally assumed to be silenced and biochemically inert in most cells. Yet examples of host genes driven by TE promoters have been documented in diverse species over the past several decades^{15,22,35–37}. Do these cases simply represent rare events where individual TEs have escaped host silencing, or are they hints of pervasive regulatory activity?

A large body of literature has illustrated the myriad mechanisms by which TEs can alter host gene expression both in *cis* and in *trans*, transcriptionally or post-transcriptionally^{15,38} (Fig 1b). Whereas most of these mechanisms were initially discovered through the genetic analysis of mutations caused by *de novo* TE insertions, genomic studies have revealed that these activities permeate throughout the genome and often emanate from TEs that have long been fixed in the host population. In particular, genome-wide assays such as those mapping transcriptional activity, open chromatin, or binding of transcription factors, have provided compelling evidence that a multitude of TE-derived sequences exhibit the biochemical hallmarks of active regulatory elements. Together these studies indicate that TEs are not as robustly or systematically silenced as commonly assumed.

Notably, transcriptome analyses have uncovered TEs as an abundant source of tissue-specific and/or alternative promoters. Two principal methods have allowed the mapping of TE-derived promoters at a genome-wide scale. First, deep sequencing of full-length cDNAs (a form of RNA sequencing (RNA-seq [G])) revealed that chimeric transcripts that initiate within TE-derived promoters constitute a considerable fraction of mammalian transcriptomes, notably during early development³⁹ but also in many other mammalian tissues⁴⁰. Second, approaches mapping sites of transcription initiation, such as cap analysis of gene expression followed by sequencing (CAGE-seq [G]) revealed surprising amounts of Pol II initiation within TEs in a variety of human and mouse cell types and tissues¹³ (e.g. up to 30% of all transcription start sites mapped in human embryonic cells) as well as in *Drosophila melanogaster* embryonic development⁴¹. There is also growing evidence that TEs are a substantial source of promoter activity in plants³⁷. Thus TEs appear to have dispersed vast numbers of developmentally-regulated promoters in a wide range of species that often remain active and drive the transcription of adjacent DNA in a tissue-or stage-specific fashion.

Evidence is mounting that TEs provide a profusion of other types of *cis*-regulatory elements including enhancers, insulators, and repressive elements. In mammals, chromatin immunoprecipitation followed by sequencing (ChIP-seq [G]) studies have revealed that for any given transcription factor (TF) and cell type examined, TEs contribute a substantial fraction of binding sites across the genome (5-40%, average ~20%)^{16,20}. LTR elements tend to contribute more than other TE types^{15,22,42}, probably because LTRs are more likely to possess and retain their ancestral *cis*-regulatory activities as explained above (Fig. 1a). Importantly, for a given TF the majority of TE-derived binding sites are contributed by a relatively small number of specific TE families that are highly enriched for TF binding events relative to what would be expected based on the density of these TE families in the genome^{18,20,43,44}. In most cases for which the origin and sequence of the binding sites have been examined, it could be inferred that a canonical binding motif pre-existed within the ancestral TE sequence and was subsequently dispersed through the genome via transposition – a scenario consistent with a ‘copy-and-paste’ model of TF binding site dispersion¹⁵.

Interestingly, TE-derived TF binding sites tend to be lineage-specific. For instance, in an analysis of binding events for 26 pairs of orthologous TFs across two comparable human and mouse cell lines, >98% of >130,000 TE-derived peaks identified in each species were species-specific²⁰. This result can be partly explained by the fact that the majority of binding events occur within TEs that have amplified after the split of the two species examined, i.e. primate- or rodent-specific TE families. Another factor is that more ancient TEs have accumulated many more neutral substitutions, leading to degradation of their ancestral TF binding sites. The differential decay of these ancestral TE sequences across species may also result in species-specific transcription factor binding events. In any case, these data suggest that transposition represents a common mechanism for the gain of novel TF binding sites during mammalian evolution.

TEs have also been documented to function as insulator and/or boundary elements. Such sequences serve to partition the genome into domains of active or inactive transcription ranging in size from 100 kb to 1 Mb, often by preventing the spread of heterochromatin⁴⁵.

Several studies showed that many TEs, in particular SINEs, harbour binding sites for factors such as CTCF or TFIIC that confer insulator activity and organize nuclear architecture^{46–48}. A subset of these TEs appear to play roles in the three-dimensional organization of the genome by functioning as ‘anchors’ that serve to isolate regions of active transcription. Indeed, studies investigating intra- or inter-chromosomal interactions underlying these topologies at a genome-wide scale using chromatin interaction profiles have found that SINEs are enriched at the borders of these domains^{49–51}. By providing a fertile source of binding sites for architectural factors, TEs may be important contributors to high-order genomic organization, which governs the transcriptional regulation of large chromosomal regions containing many genes.

In addition to providing *cis*-regulatory DNA elements, TEs have also been documented to contribute a wealth of non-coding regulatory RNA transcripts such as microRNAs (miRNAs) and long non-coding RNAs (lncRNAs) that can modulate gene expression in *cis* or in *trans* (see Fig. 1b and Box 2).

Finally, while we primarily focus this Review on regulatory activities encoded within TE sequences, TEs can also alter gene expression through many other effects, including the disruptive effects of their chromosomal integration. An interesting example is the adaptive insertion of a *POGONI* DNA transposon within the 3′ UTR of the gene *CG11699* in *D. melanogaster*⁵². The insertion disrupts the ‘normal’ (ancestral) polyadenylation signal of the gene resulting in a shorter 3′ UTR, elevated mRNA levels, and increased resistance to xenobiotic stress⁵². TEs may also exert a broad influence on genome regulation as a result of their targeting by host silencing pathways³³ (Fig. 1b). In particular, the repressive chromatin nucleated at TE sequences may ‘spread’ to adjacent regions and silence the expression of nearby genes, a phenomenon which has been observed in several organisms including plants⁵³ and mammals⁵⁴.

The biological impact of TE regulatory activities

The work summarized above firmly anchors the idea that TEs are a prolific source of biochemical regulatory activity in host cells. These observations raise the question as to whether this phenomenon has exerted a significant influence on the biology and phenotypic evolution of species. Although it may be tempting to interpret the tightly-controlled regulatory activity originating from TEs as indicative of a widespread role in physiology or development, the biochemical manifestation of this activity does not alone signify that it is important for proper genome function^{55,56} (Fig. 1c). The terms ‘function’, ‘activity’ or ‘co-option’ are widely used yet can have different meanings depending on the field of study, and we caution that imprecise usage of these terms might confuse our understanding of the influence of TE regulatory activity on biological outcomes. Coming from an evolutionary perspective, we use the term “co-option”, which is also referred to as exaptation⁵⁷ or domestication⁵⁸, when a TE-derived sequence acquired a cellular function conferring a selectable phenotype that predictably increases the fitness of the host organism (Fig. 1c). Given that this review focuses on regulatory activity of TEs rather than their protein products, we will use the term ‘co-opted TEs’ to refer specifically to the co-option of DNA sequences within the TE that are responsible for *cis*-regulatory activity.

With the goal of advancing our current understanding of the biological importance of TE regulatory activities, we discuss the major challenges involved in disentangling effects that are simply relics of the selfish ‘behaviour’ of TEs with no benefit to the host from those that reflect bona fide co-option events resulting in adaptive cellular innovations.

Co-option of TEs revealed by evolutionary sequence conservation

How can evolutionary co-opted, or ‘domesticated’ TEs be distinguished from TEs whose regulatory activity is non-adaptive and simply tolerated by host cells? One of the most direct lines of evidence that a DNA sequence has exerted biological function is to show that it has evolved under a regime of purifying selection [G]. Selective constraint on non-coding DNA is most effectively revealed by detecting a signature of evolutionary sequence conservation when compared across distantly related species, relative to DNA sequences that are assumed to be unconstrained and neutrally evolving. Such comparative genomic studies have revealed tens of thousands of non-coding TE fragments in the human genome that are orthologous and highly conserved across species and display clear signatures of purifying selection^{15,56,59}. Although most of these sequences derive from relatively ancient TEs (which are old enough to be compared orthologously across distant species), they are of various ages and types. They are also unevenly distributed in the genome and tend to be enriched near transcription factor genes and other developmental genes. Together these data strongly suggest that TEs have indeed provided a rich source of non-coding sequence material fueling regulatory innovation during vertebrate evolution.

Striking examples of TEs co-opted for gene regulation identified by virtue of evolutionary conservation are those of the so-called ‘living fossil’ SINE (LF-SINE) family⁶⁰. Multiple LF-SINE copies are highly conserved across tetrapods, and one was shown to possess tissue-specific enhancer activity in a mouse reporter assay [G] (Table 1). Another interesting example is a highly conserved mammalian-wide interspersed repeat (MIR) element that in mice serves as an intronic enhancer for the *FOXP3* gene required for extrathymic generation of regulatory T cells, which re-enforces maternal-fetal immunotolerance during pregnancy⁶¹. Additionally, two ancient TEs, a mammalian apparent LTR retrotransposon [G] (MaLR) and a SINE, were independently co-opted during mammalian evolution to serve as hypothalamus-specific enhancers that are required to regulate the proopiomelanocortin (*POMC*) gene, which functions in the brain to control food intake⁶². Examples like these highlight how evolutionary conservation of TEs can guide experiments validating regulatory functions.

Although tens of thousands of TEs in the human genome contain sequences that have evolved under purifying selection⁵⁹, as well as sequence motifs and biochemical signatures that are consistent with some form of *cis*-regulatory activity^{63,64}, it is important to emphasize that, thus far, very few of these elements have been characterized using functional assays. Interestingly, a recent study⁶⁵ identified a DNA segment that is highly conserved across eutherian mammals and derived from three distinct juxtaposed TEs (AmnSINE1, X6b_DNA, and MER117) that appears to function as an enhancer driving frontonasal expression of the crucial development gene *Wnt5* in the mouse embryo. Surprisingly, however, mice carrying a homozygous deletion of the TE-derived enhancer did not show any

obvious phenotypic defects, raising the paradox — not unprecedented⁶⁶ — that even sequences with a strong signature of evolutionary constraint and demonstrated *cis*-regulatory activity may only exert subtle effects on organismal development.

Uncovering recently co-opted TEs

Although sequence conservation is a strong predictor of biological function, there are several caveats that may preclude comparative genomics from identifying TEs that have been co-opted for host gene regulation. First, only relatively ancient TEs that have been co-opted for a sufficiently long evolutionary period will display a robust signature of sequence constraint⁶⁷. These ancient TEs are technically challenging to recognize as such, especially in species with rapid evolution rate such as fruit flies (Box 3). In human and mouse, the vast majority of TEs exhibiting regulatory activity in functional genomics assays are non-orthologous and derive from relatively young insertions^{20,42}. The recent origins of these elements make it difficult to assess whether they have evolved under functional constraint, although some methods have been developed in attempt to address this issue⁶⁸. The matter is worsened by the intrinsic shortness and degeneracy of the sequence motifs underlying regulatory activity, which increases the difficulty of detecting purifying selection acting on these motifs through sequence analysis alone^{69,70}.

Despite these hurdles, there is a growing set of recent TEs for which solid evidence has been gathered to support the idea that they were co-opted for important regulatory innovation³⁶ (some examples are provided in Table 1 and Fig. 2). These are often cases in which the TE serves as a tissue-specific promoter for a gene with crucial function in that tissue. Indeed, these cases can be readily discerned by the detection of a tissue-specific gene transcript initiating within a TE sequence³⁵. A classic example in mammals is the acquisition of salivary expression of the digestive enzyme amylase from a retroviral LTR inserted in the common ancestor of anthropoid primates⁷¹. Another interesting case is the endometrial expression of the hormone prolactin, which is critical for pregnancy in mammals^{72,73} (Box 3).

In contrast to TEs with promoter activity, the function of TEs located distal to genes is more challenging to discern and validate. Therefore, far fewer examples of TE-derived enhancers with clear biological roles have been documented³⁶. A traditional experimental approach to delineate the *cis*-regulatory effects of a TE is a reporter assay. If the reporter expression pattern driven by the TE recapitulates that of the endogenous gene associated with the TE, it may be reasonable to conclude that the TE contributes to the regulation of the endogenous gene *in vivo*^{60,74}. However, these experiments are still limited by the fact that they dissociate the TE from its native chromosomal context and cannot establish a direct causal link between the *cis*-regulatory activity of the TE and endogenous gene expression.

A more conclusive approach to assess the effects of an individual TE on host gene regulation is to perform a loss-of-function experiment. The recent development of precise genome editing technologies has begun to facilitate such experimental knockout studies. In one study, transcription activator-like effector nuclease (TALEN)-mediated deletion of a mouse MT element providing an alternative promoter for the *Dicer1* gene led to oocyte malfunction and female sterility⁷⁵. In another study, CRISPR-mediated deletion of a MER41 ERV-

derived interferon-inducible enhancer associated with the *AIM2* gene in human cells impaired the innate immune response to viral infection⁴⁴ (Fig. 2b). In a third example, a screen for *cis*-elements regulating the pregnancy-specific histocompatibility gene *HLA-G* combining massively parallel reporter assays and CRISPR genome editing identified a LTR71-derived distal enhancer required for *HLA-G* expression in the placenta⁷⁶.

TE-mediated cis-regulatory variation and the domestication of species

TEs were first discovered in maize and have since been characterized as a major component of other crops. Intriguingly, there is mounting evidence that many traits associated with the domestication of crop plants evolved through artificial selection of TEs with *cis*-regulatory effects on adjacent host genes^{14,37}. In maize for instance, a *Hopscotch* retrotransposon insertion serves as an enhancer for the *teosinte branched1* gene underlying increased apical dominance relative to its ancestral wild relative teosinte⁷⁷. Another study examining blood oranges identified a *copla*-like retrotransposon that functions as a cold temperature-inducible enhancer of the *ruby* locus that modulates fruit color⁷⁸ (Table 1).

Furthermore, there is evidence that regulatory variation introduced by TEs have facilitated the domestication of animals. A study investigating the domesticated silkworm found that the insertion of a *Taguchi* LINE enhances expression of the *ecdysone oxidase (EO)* gene, which inhibits premature metamorphosis of silk-producing pupae⁷⁹. Additionally, several TE insertions with *cis*-regulatory effects have been associated with traits selected in domestic dog and cat breeds, such as small body size⁸⁰ and coat color pattern^{81,82}.

A recent report⁸³ identified an intronic insertion of a TE that enhances the expression of the *cortex* gene, and this increased expression was found to underlie industrial melanism in the English peppered moth (Fig. 2c). In this textbook example of adaptation, peppered moths were not domesticated *per se*, but were subject to strong artificial selection by coal pollution during the industrial revolution.

Together these findings suggest that TEs frequently play pivotal roles facilitating plant and animal domestication, which is characterized by artificial selection for specific traits often modulated through changes in gene expression⁸⁴.

TEs and the evolution of gene regulatory networks

Whereas most previous studies focused on individual cases of a particular TE regulating a single host gene, it is theoretically possible that TEs exert a more extensive influence in regulatory evolution by providing the ‘wiring’ connecting large gene batteries. Such regulatory networks coordinate the expression of multiple gene products that act in concert to control entire pathways and complex biological processes. The assembly of new gene regulatory networks is thought to underlie major evolutionary innovations including the emergence of new morphological structures and cell types⁸⁵. There is mounting evidence that modification of the *cis*-regulatory architecture underlying the transcriptional control of genes is an important force driving the evolution of new networks⁸⁶. Transposition has long been proposed as an attractive mechanism to facilitate the concurrent mutational events that are required to deeply remodel *cis*-regulatory architecture^{3,15}.

This model has gained support from recent studies linking the expansion of certain TE families with the dispersal of a regulatory module that is important for the execution of a specific developmental program. For instance, MER20 transposons appear to have deposited numerous cAMP-inducible enhancers specifying endometrial gene expression in the eutherian mammal ancestor, which coincided with the emergence of mammalian pregnancy over 100 million years ago⁶³. Another example is the dispersal of hundreds of enhancers with forebrain-specific activity in the mouse through the ancient expansion of the MER130 family, which may have been associated with the evolution of the mammalian neocortex⁶⁴. There is also evidence supporting a role for lineage-specific TEs in driving more recent adaptive evolution of gene regulatory networks. For example, the murine-specific RLTR13 family of ERVs dispersed hundreds of placenta-specific enhancers within the past 15-25 million years⁸⁷, which may reflect and possibly directly contribute to the rapid morphological diversification of the mammalian placenta⁸⁸.

Functional evidence for network rewiring by TEs

These recent findings support a model whereby waves of TE activity deposited the raw material for large-scale *cis*-regulatory changes underlying major evolutionary innovations. However current evidence is mostly limited to correlative observations based on chromosomal association of TEs with nearby genes encoding evocative functions^{63,64,87,89,90}. Although these associations are tantalizing, they should be interpreted with caution. An alternative, but not mutually exclusive interpretation is that TEs inserted within or near genes that are highly transcribed in a given cell type are more likely to be accessible to regulatory proteins (e.g. TFs) expressed in those cells. Because TEs may have an intrinsic preference for integration within 'open' transcriptionally permissive chromatin^{91,92} and must also replicate in the germline for new insertions to be inherited, these properties may introduce a biased association with genes that are highly expressed in the germline and early embryonic cell types⁹³⁻⁹⁵. Second, there is probably some level of selection against insertions that significantly alter host gene expression patterns upon insertion, meaning that most TEs retained in the genome might cause minimal changes in host gene expression⁹⁶⁻¹⁰². This bias would also favour the retention of TEs near or within genes that share similar expression profiles, as these insertions are less likely to significantly alter existing gene expression patterns. Thus, although genomic analyses often appear to support a causal role for TEs in shaping regulatory networks, these data alone cannot falsify the null hypothesis that a given TE has no regulatory effect on a nearby gene, even when exhibiting biochemical signatures that are consistent with *cis*-regulatory activity⁵⁶.

A few recent studies have gone a step further in testing more directly the role of TE co-option in the evolution of gene regulatory networks by using experimental genetic manipulation. In a study investigating the interferon-inducible gene regulatory network, CRISPR-mediated deletion of four separate primate-specific MER41 elements impaired the expression of adjacent genes with known innate immune functions⁴⁴. Thus the co-option of multiple MER41 elements with the same regulatory properties at several genomic loci appear to have facilitated the establishment of a coordinated transcriptional response to infection during primate evolution.

Another genetic network where TEs have been implicated is the *cis*-regulatory circuit enabling dosage compensation on sex chromosomes. Studies of the emergence of dosage compensation on the newly evolved (<1 million year old) 'neo-X' chromosome of the fruit fly *Drosophila miranda* showed that an ISX transposon was responsible for spreading dozens of binding sites for the dosage compensation machinery¹⁰³ (Box 3). These ISX elements are strikingly enriched on the X chromosome, and experimental insertion of an ISX element into an autosomal chromosome was sufficient to recruit the dosage compensation complex to this ectopic site. Together with data indicative of the involvement of TEs in mammalian dosage compensation^{104,105}, these findings suggest that transposition played a recurring role in the evolution of sex chromosomes by enabling the rapid copying-and-pasting of *cis*-regulatory sequences along entire chromosomes.

A growing number of studies also suggest a potential role for TEs in rewiring gene regulatory networks orchestrating early mammalian development²⁴. Although the biological relevance of these findings remains to be clarified, these reports raise the exciting possibility that ERVs have been repeatedly co-opted during mammalian evolution to remodel the complex genetic circuits underlying early embryonic development (Box 1).

Pathogenic effects of TE regulatory activities

In line with their parasitic origins and selfish behaviour, TEs have long been associated with mutant phenotypes and disease. TEs are well documented to cause disease through two primary mechanisms: insertional mutagenesis and chromosomal rearrangements. For example, *de novo* germline TE insertions disrupting normal gene function have been implicated in over a hundred human inherited diseases¹⁰⁶. Furthermore, both transposition and TE-mediated chromosomal rearrangements in somatic cells have been causally linked to several types of cancer¹⁰⁷. Here we explore how the regulatory activities of TEs may also represent an underappreciated source of disease phenotypes.

TEs as pathogenic regulatory variants

Cis-regulatory variation is increasingly recognized as an important factor influencing disease susceptibility¹⁰⁸. As TE insertions represent a common form of structural variation [G] in human genomes¹⁰⁹, it is plausible that some of these polymorphic insertions contribute to disease risk by modulating the expression of adjacent genes. This idea has yet to be investigated systematically, but a plausible example recently surfaced from the discovery of a link between the complement C4 system and schizophrenia risk¹¹⁰. The authors found that individuals carrying a polymorphic ERV intronic insertion in the C4 gene have elevated C4 expression, which in turn was found to cause synapse over-pruning, which is a phenotype that is associated with schizophrenia¹¹⁰. Although the evidence linking ERV regulatory activity to disease remains indirect, this case is intriguing in light of previous observations of an association between schizophrenia and elevated ERV transcriptional activity¹¹¹.

Causes and consequences of TE reactivation

TEs that are fixed in the human population, but encode dormant regulatory sequences, may also contribute to pathogenesis. It has been widely observed that TE transcript levels are

significantly increased in numerous cancers and other disease states^{112,113}. The reasons why particular TEs appear to be upregulated in certain disease conditions remain poorly understood. Recent data suggest that environmental stimuli, including infection¹¹⁴ and cellular stress¹¹⁵, as well as natural cellular processes such as senescence^{116–118}, destabilize epigenetic marks that normally silence the bulk of TEs in the genome, thereby triggering their sporadic transcriptional activation.

Does the derepression of TEs play a major role in driving disease states, or is it simply a side effect of pathogenesis^{113,119,120}? There are several routes by which inappropriate transcriptional activation of TEs might have pathogenic consequences for the host (Fig. 3a). First, this activity may result in increased rates of propagation through transposition. Indeed, DNA hypomethylation and transcriptional reactivation of replication-competent LINE-1 copies seems to explain why some tumours, particularly epithelial ones, may exhibit an increased rate of transposition relative to matched healthy tissues^{121–123}. The role of somatic insertional mutagenesis and its contribution to tumorigenesis and malignancy is an area of considerable interest¹¹⁹. Indeed, recent studies have demonstrated that *de novo* LINE-1 insertions can activate oncogenic pathways in hepatocellular carcinoma¹²⁴ and colorectal cancer¹²⁵. Activation of the LINE-1 transposition machinery may have additional pathological consequences, such as the infliction of double strand breaks in the genome through LINE-1-encoded endonuclease activity¹²⁶.

It is important to keep in mind that the vast majority of TEs that are transcriptionally activated in disease (such as ERVs) are no longer replication competent, and as such are unable to generate new transposition events. However, there are other potential routes for an excess of TE-encoded transcripts to lead to pathogenesis (Fig. 3a). For example, some of these transcripts may encode peptides with adverse cellular activities. For instance, overexpression of ERV envelope proteins, as seen in the brain of patients affected with some neurodegenerative and autoimmune disorders, can induce a wide range of cellular processes and abnormalities associated with these pathologies, such as neurodegeneration¹²⁷, autoinflammation¹²⁸, demyelination¹²⁹, and superantigen activity¹³⁰. Furthermore, the cytoplasmic accumulation of nucleic acids derived from activated TEs, including double-stranded RNA, reverse transcribed cDNA, or RNA–DNA hybrids, are increasingly regarded as potent immunological ‘adjuvants’ that may trigger autoimmune responses^{131,132} and other toxicities when present in elevated quantities^{133,134}.

Misregulation of host genes by TEs

Perhaps most importantly, and in-line with the major theme of this Review, the reactivation of TEs may promote disease states indirectly by altering host gene expression. TEs that are normally silenced by DNA methylation may exhibit *cis*- or *trans*-regulatory activity that could cause global dysregulation of host genes in *cis* or *trans* (through the various mechanisms illustrated in Fig. 1b). Thus ectopic activation of dormant TE *cis*-regulatory sequences may result in the pathogenic activation of genes or pathways in some cells. Although evidence supporting this model remains limited, there is a growing number of studies showing how derepression of a particular TE copy activates transcription of an adjacent proto-oncogene^{135–137} (Fig. 3b). Whereas the loss of regulatory control at these

specific elements may be a rare stochastic event occurring in a small subset of cells, it is possible that this process could be favoured by selection during tumour evolution. In this model termed ‘onco-exaptation’, individual cells where an oncogenic TE is aberrantly unmasked acquire a fitness advantage over other cells as a result of altered oncogene expression¹³⁸. This process would favour clonal propagation of cells where the TE is unmasked and perpetuate tumour growth.

In addition to altering the expression of adjacent protein-coding genes, reactivated TEs can also drive widespread expression of non-coding RNAs, which themselves are largely derived from TE sequences^{139–141}. Many of these transcripts are likely to be non-functional, but there is evidence that some have oncogenic properties. For example the lncRNA *BANCR* is specifically expressed in melanoma cells and promotes the proliferation and migration of these cells in culture¹⁴². Interestingly, *BANCR* exons are largely derived from a MER41 ERV insertion, and its promoter is derived from the MER41 LTR¹⁴⁰. A similar example is *EVADR*, which is a lncRNA derived from the ERV MER48 that is recurrently overexpressed in adenocarcinomas¹⁴³. Further studies are warranted to determine the mechanisms by which these TE-derived transcripts might contribute to tumorigenesis.

The regulatory activities of TEs are also emerging as drivers of pathogenesis in non-human species. In the oil palm for example, a fixed *Karma* TE insertion within an intron of the gene *MANTLED* is normally methylated, but sporadic demethylation of the region provides an alternative splice site and a premature termination signal for *MANTLED*¹⁴⁴ (Fig. 2d). This epigenetic dysregulation was found to underlie the spontaneous production of deformed oil palm fruits by genetic clones¹⁴⁴.

The double-edged sword of co-opting TEs

How can potentially pathogenic TEs persist in host genomes and not be purged by negative selection? In some cases it might just be a matter of time until TE insertions segregating at low frequency in the population are eventually eliminated. It is also possible that the pathogenic activity of certain TEs is only unmasked in a small number of individuals, or exert only weak or post-reproductive effects, and therefore impose an insufficient fitness cost to be purged by natural selection. Another possibility is that some TEs play an adaptive role, but occasional misregulation acts a negative, disease-causing side effect of co-option. Under such a scenario co-opted TEs might be viewed as ‘double-edged swords’ in host evolution: beneficial in some situations or individuals, but detrimental in others.

Our recent discovery of a MER41 element conferring interferon-inducibility to the *AIM2* gene⁴⁴ (Fig. 2b) may represent an example of a double-edged TE. *AIM2* encodes an important immunity factor that also acts as a potent tumour suppressor¹⁴⁵. Constitutive transcriptional dysregulation of *AIM2* (either up or down) has been recurrently observed in cancer and autoimmune diseases, although the mechanisms underlying this misregulation are not well-understood¹⁴⁶. In one study examining colorectal colon cancer cells, where *AIM2* is constitutively silent and unresponsive to interferon, the upstream promoter region (~700 bp upstream of the *AIM2* gene) was found to be consistently hypermethylated in cancer samples¹⁴⁷. Intriguingly, this region coincides with the location of the MER41

element, suggesting that aberrant methylation of this TE might account for silencing of *AIM2* in the cancer cells.

Conclusions and future perspectives

The past decade was marked by tremendous progress in our understanding of how TEs shape genome evolution. Major headways were propelled in large part by advances in technologies enabling genome-scale analysis. Genome-wide surveys revealed TEs as a substantial source of *cis*-regulatory elements in diverse eukaryotic species, lending credence to ideas pioneered decades ago by McClintock, Britten and Davidson, among others. It is now apparent that TEs evolved many complex mechanisms and biochemical activities that, to various extents, predispose them to transitions from parasitic elements to integral components of host gene regulation; ‘from conflicts to benefits’. A pressing challenge is to obtain more direct assessments of the biological consequences of the regulatory activity of TEs, which in turn will provide a better understanding of the long-term evolutionary implications of TE dispersal. Despite decades of genetic analyses of mutant and disease phenotypes illustrating how TE insertions and rearrangements can alter host gene expression patterns in many ways, there is still relatively little experimental evidence that TEs have promoted evolutionary innovations in organismal development and physiology.

Spurred by new tools for direct genetic manipulation such as the CRISPR-Cas9 system, recent studies add to the small but growing list of TEs that have been unambiguously co-opted for regulating cellular functions. A major outstanding task is to gain a better grasp on the role of TE co-option in driving the evolution of gene regulatory networks on a broader scale. Recent work suggests that TEs can extensively remodel regulatory networks that are involved in specific processes including dosage compensation¹⁰³, immunity⁴⁴, and early embryonic development²⁴. Other comparative studies of enhancer evolution in the liver¹⁴⁸ and the neocortex¹⁴⁹ among mammals found only a minor contribution for host co-option of TE activity. These observations raise the question of whether TE-mediated regulatory evolution may be inherently biased for certain biological processes, or if it is a more general mechanism for large-scale genetic innovation.

Taking phylogenetic and other retrospective approaches for assessing the function of ancient TEs will also provide insights into how more-recent TEs have shaped gene regulatory networks in modern species. Here one of the major hurdles is to reach back to events of the distant past to study specific steps leading to TE co-option, which are obscured by millions of years of evolution (Box 2). Therefore, in addition to these types of retrospective studies, investigations of ongoing adaptation in populations^{150,151} or using protocols of real-time experimental evolution^{152,153} will shed light on how TEs can facilitate regulatory evolution in response to defined selective pressures during the earliest stages of adaptation.

There is also growing evidence linking aberrant TE regulatory activity and disease states. Although only a few examples of pathogenic TE misregulation have been documented so far for contributing to cancer, TE reactivation is clearly associated with other pathological conditions including ageing¹⁵⁴, neurological disorders¹⁵⁵, and autoimmunity¹³³. Our mechanistic understanding of the cause and consequence of such ectopic TE activation for

disease aetiology or progression is still in its infancy. As technological advances continue to increase our ability to functionally test the involvement of non-coding regulatory processes in pathogenesis, it will open new avenues to assess the role of TE-mediated gene dysregulation in disease.

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Glossary

Genetic drift

A process by which mutations become fixed in the population by chance alone

Cis-regulatory sequences

Segments of DNA that regulate the transcription of adjacent genes

Long terminal repeat (LTR) elements

A class of retrotransposons containing direct long terminal repeats flanking the protein-coding sequence

Long interspersed nuclear elements (LINEs)

Class of non-LTR retrotransposons that retrotranspose by target-primed reverse transcription

Short interspersed nuclear elements (SINEs)

Class of non-autonomous retrotransposons that are copied by the LINE replication machinery

Retrotransposon

A type of transposable element that replicates through an RNA intermediate in a ‘copy-and-paste’ mechanism

DNA transposons

Transposable elements that do not generate an RNA intermediate during transposition, which generally occurs through a ‘cut-and-paste’ mechanism

CAGE-seq (Cap Analysis of Gene Expression followed by sequencing)

A method used to precisely map the transcription start sites of capped RNAs genome-wide

RNA-seq

High-throughput sequencing of complementary DNAs derived from RNAs extracted from cells or tissues

ChIP-seq (Chromatin immunoprecipitation followed by sequencing)

A method for identifying protein–DNA interactions genome-wide. Following crosslinking, a protein of interest is immunoprecipitated, and its binding sites in the genome are identified by high-throughput sequencing of the co-purified DNA fragments

Reporter assay

A putative *cis*-regulatory DNA sequence is cloned upstream of a reporter gene (such as luciferase) either in an episomal vector or as a chromosomally integrated construct and tested for its ability to enhance transcription of the reporter gene

Purifying selection

Selection against mutations that are deleterious to the fitness of the individual

Structural variation

Genomic variation resulting from large-scale DNA mutations such as deletions, insertions, or rearrangements

Biographies

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Edward Chuong received his Ph.D. from Stanford University in 2013, where he worked with Julie Baker to study the impact of endogenous retroviruses on mammalian placenta development. Since 2013, he has been a Postdoctoral Fellow at the University of Utah School of Medicine, Salt Lake City, USA, working with Nels Elde and Cédric Feschotte to study how transposable elements influence gene regulatory evolution in host immunity.

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Cédric Feschotte

Cédric Feschotte received his Ph.D. in 2001 from the University of Paris VI, France, during which he characterized mosquito transposable elements with Claude Mouchès. From 2000 to 2004, he was a Postdoctoral Associate in the group of Susan R. Wessler at the University of Georgia, Athens, USA, where he explored the origin and amplification mechanism of 'MITE' transposons in plant genomes. In 2004, he became an Assistant Professor in the Department of Biology at the University of Texas, Arlington, USA, and an Associate Professor in 2009. In June 2012, he joined the Department of Human Genetics at the University of Utah School of Medicine, Salt Lake City, USA, where he currently holds the title of Professor and runs a laboratory investigating the biological impact of mobile genetic elements.

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Box 1**A role for ERVs in regulating early mammalian development?**

Genome-wide epigenetic silencing of transposable elements (TEs) is a pivotal step in early mammalian development, yet recent studies characterizing the transcriptomes of embryonic stem cells and early embryos have revealed surprisingly high transcriptional activity emanating from endogenous retrovirus (ERV) sequences. ERVs are amongst the first sequences to be transcribed during zygotic genome activation in mouse two-cell embryos³⁹. Similarly, in human embryos, distinct families of primate-specific ERVs are expressed at each stage of pre-implantation development, and this activity ceases as cells differentiate into somatic cells^{156,157}. These ERVs are transcriptionally regulated by long terminal repeat (LTR) promoters that contain binding sites for transcription factors governing early development, such as OCT4 and NANOG²⁴. Taken together, these findings indicate that ERVs are seemingly unleashed during pre-implantation development, with potentially widespread regulatory effects on the cellular transcriptome.

A major challenge now lies in understanding the biological consequences of ERV activity in embryonic development. RNA sequencing (RNA-seq) analyses have revealed that hundreds of LTRs drive expression of lncRNAs¹⁵⁸ and protein-coding genes^{39,159}, and chromatin immunoprecipitation followed by sequencing (ChIP-seq) analyses have revealed thousands more solitary LTRs that function as embryonic stem cell (ESC)-specific distal enhancers that may also influence host gene expression¹⁸. The expression of some host genes that are required for pre-implantation development may thus be regulated by ERV-derived promoters or enhancers.

The activity of some of these ERVs, such as the primate-specific HERV-H, is highly correlated with stem cell pluripotency^{139,159–161}, and knockdown of long non-coding RNAs (lncRNAs) driven by HERV-H results in rapid differentiation into somatic cell types^{159,161–164}. Further molecular investigation in human embryonic stem cells suggested that HERV-H transcripts may function as RNA scaffolds recruiting transcriptional activators including OCT4, Mediator, and p300¹⁶⁴. Yet another lncRNA — named human pluripotency-associated transcript 5 (HPAT5) and derived from both a primate-specific HUERS-P1 ERV and an *Alu* element — was also discovered to promote pluripotency by functioning as a molecular sponge for the *Let-7* family of microRNAs (miRNAs)¹⁶⁵ (Fig. 2a). Together these results indicate that ERV-derived lncRNAs are capable of modulating stem cell pluripotency, which may be important for proper development. Another study found that the protein Rec encoded by HERV-K binds to a multitude of host mRNAs and potentiates antiviral resistance in cell culture, raising the provocative hypothesis that HERV-K expression protects pre-implantation embryos from exogenous viral infection¹⁵⁷.

These data provide mounting evidence that ERVs have been co-opted as essential regulators of human development. However, an important limitation to these studies is that they are typically performed in cultured embryonic stem cells, thus it remains possible that functional phenotypes linked to ERV regulatory activity may not be required for organismal development. One pioneering study used small interfering RNA (siRNA)

injections to simultaneously knock down expression of three ERV-derived lncRNAs (*HPAT2*, *HPAT3* and *HPAT5*) in human two-cell embryos, and found that cells depleted for these lncRNAs were no longer capable of contributing to the inner cell mass of the blastocyst¹⁶⁵. This result suggests, but does not prove, that ERVs have been co-opted to regulate blastocyst development. Given the limited sample size and potential for off-target or transcriptional effects of siRNA silencing, further studies are necessary to conclusively establish a developmental role for human ERVs. Indeed it is possible that ERV activity in embryonic stem cells is not essential for development, but instead merely reflect the recent selfish exploitation of this developmental niche by retroviruses¹⁶⁶.

Although the necessary experiments would be challenging with human material, mice similarly exhibit dynamic expression of ERVs during early embryonic development²⁴. Whereas all ERVs with regulatory activity in human embryos are primate-specific, mouse embryos similarly exhibit rodent-specific ERVs that are expressed throughout early development^{39,158}. If ERV activity is truly required for both human and mouse early development, it would imply a remarkable scenario whereby ERVs were independently co-opted in primates and rodents to regulate embryogenesis. Note there are precedents for such convergent TE co-option events^{167,168} (Box 3).

Box 2**TEs as a source of non-coding regulatory RNAs**

In addition to providing *cis*-regulatory DNA elements, transposable elements (TEs) have also been documented to contribute non-coding regulatory sequences that modulate gene expression post-transcriptionally. Best documented are TE-derived regulatory sequences embedded within the untranslated regions (UTRs) of protein-coding genes. For instance, *Alu* elements inserted within 3' UTRs can provide recognition sequences for Staufen 1-mediated mRNA decay¹⁶⁹ or influence mRNA localization¹⁷⁰, whereas *Alu* elements incorporated within 5' UTRs often modulate mRNA translation efficiency¹⁷¹.

TEs are also major contributors to the evolutionary origination and biogenesis of various regulatory RNAs that regulate gene expression either transcriptionally or post-transcriptionally, including microRNAs (miRNAs)¹⁷², long non-coding RNAs (lncRNAs)^{139,140}, and circular RNAs (circRNAs)^{173,174}. In human, mouse and zebrafish, more than two thirds of lncRNAs were found to contain exonic TE sequence, whereas they seldom occur in protein-coding transcripts^{139,140}. Of course, many of these lncRNAs may have no biological function and could merely reflect the pervasive transcriptional activity of TEs. It could also reflect the greater capacity of lncRNAs to tolerate and assimilate TE insertions over evolutionary time^{15,70,175}. Nevertheless, there is a growing number of examples where the sequences conferring regulatory activities to non-coding RNAs are directly derived from tes^{70,164,165,175–177} (Box 1).

Many of the conceptual ideas developed in this Review for TE-derived *cis*-regulatory DNA elements can be applied to those acting at the RNA level. In particular, a *cis*-acting sequence (e.g. a recognition motif for an RNA-binding protein) that is present within an ancestral TE sequence will be seeded as the TE amplifies throughout the genome, opening the door for the co-option of multiple TE copies to assemble complex regulatory circuits modulating the expression of a vast number of genes^{15,176,178}. Further details on the role of TEs in the evolution of regulatory RNAs and post-transcriptional gene regulation have been reviewed elsewhere^{38,70,175}.

Box 3**Evolutionary dynamics of TE regulatory activities**

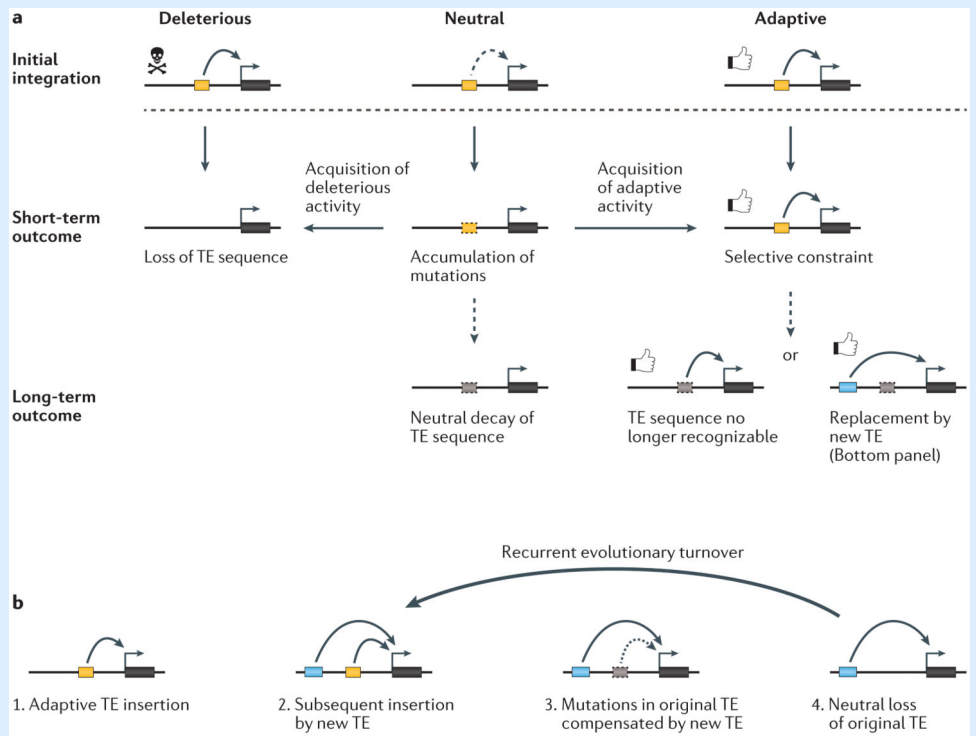
What is the mode and tempo of transposable element (TE) co-option for *cis*-regulatory function? Does TE co-option occur in major bursts that coincide with the invasion of TEs, or is it a more gradual process tapping into a reservoir of decaying TEs with various levels of predisposition?

The most straightforward path to co-option is when a TE confers an adaptive regulatory function immediately upon insertion (See the figure, part a, right). This scenario may conceivably be common because the *cis*-regulatory elements mapped within adaptive TEs are often inferred to have pre-existed at the time of their insertion in the genome⁴⁴. Some of the adaptive TE insertions that recently swept in the *Drosophila melanogaster* population may represent examples of new *cis*-regulatory sequences gained by transposition¹⁷⁹. It could also be that some TEs have immediate *cis*-regulatory effects that are independent of their own sequences, simply through their disruptive⁵² or epigenetic activities.

There is also evidence that a TE may become ‘latently’ co-opted long after it inserted in the genome (See the figure, part a, middle). Although many TEs possess built-in *cis*-regulatory sequences, these may remain silent and/or inconsequential for adjacent gene expression until additional mutations unmask or bolster their regulatory activity. Along those lines, Emera and Wagner proposed a model of ‘epistatic capture’ to describe the series of mutational events by which a TE was transformed into a strong promoter for decidual prolactin⁷². It is also conceivable that such epistatic mutational events occurring outside of the TE sequence but within its genomic neighbourhood could also promote latent co-option of the TE. A variety of mechanisms can be envisioned, such as point mutations introducing or removing other TF binding sites, as well as insertion or deletion events of *cis*-regulatory elements, genes, or even other TEs.

What is the evolutionary fate of a TE that has become co-opted for host function? As discussed in the main text, a recent study in *Drosophila miranda* found that dosage compensation on a newly evolved ‘neo-X’ chromosome (~1 million years old) was mediated by the recent spread of an ISX transposon, which harbours a ‘recognition element’ motif for the male-specific lethal (MSL) dosage compensation complex¹⁰³. Remarkably, the authors also found evidence that dosage compensation on the older XR chromosome (~15 million years old) was partly established by an older expansion of a related TE family, named ISXR¹⁰³. The older ISXR elements were characterized by stronger MSL recognition elements and decaying TE sequence. The authors also found evidence that the MSL recognition elements in the younger ISX elements have already undergone additional evolutionary ‘fine-tuning’ to optimize binding of the MSL dosage complex, facilitated by a gene-conversion-like process between copies¹⁸⁰. These data suggest a two-step model of TE domestication, whereby elements with a regulatory predisposition upon insertion are further refined by post-insertional mutation and selection. Eventually, co-opted TEs will lose signatures of their TE origins as all but the most essential nucleotides are eroded by neutral substitutions¹⁰³.

Another emerging insight is that the co-option of TEs as regulatory elements may be a surprisingly volatile process. Several cases have now been described whereby multiple species have independently, but convergently co-opted lineage-specific TEs to regulate the same gene. In mammals, this is exemplified by the prolactin gene⁷² and also the neuronal apoptosis inhibitory protein (*NAIP*) gene¹⁸¹. One potential explanation is that similar selective pressures drove independent co-option of TEs at the same locus in different species. However, it is also possible that the ancestral locus was also regulated by a TE, which has since been replaced by a new TE in each lineage. Thus, analogous to a gene duplication event, the insertion of a TE with similar regulatory properties might result in relaxed selection and eventual decay of a nearby co-opted TE, and over time this would perpetuate in a cycle of co-opted TEs being replaced by newer TEs (See the figure, part **b**). Such a turnover model could explain the prevalence of lineage-specific TEs associated with cellular processes that have deep evolutionary origins, such as pregnancy⁹⁰ or stem cell development¹⁸.



Key points

- Transposable elements (TEs) are increasingly recognized as a potent source of regulatory sequences in eukaryotic genomes.
- The selfish replication cycle of TEs drove the evolution of finely tuned regulatory activities that favoured their propagation and has predisposed them to be co-opted for the regulation of host genes.
- There is a growing number of examples of TE-derived sequences that have been co-opted to regulate important biological processes in organismal development and physiology.
- Dysfunction of TE-derived regulatory sequences is also emerging as a potential driver of diseases including cancer and autoimmunity.
- Functional genomics and genome editing technologies herald an exciting era for understanding the biological impact of TEs.

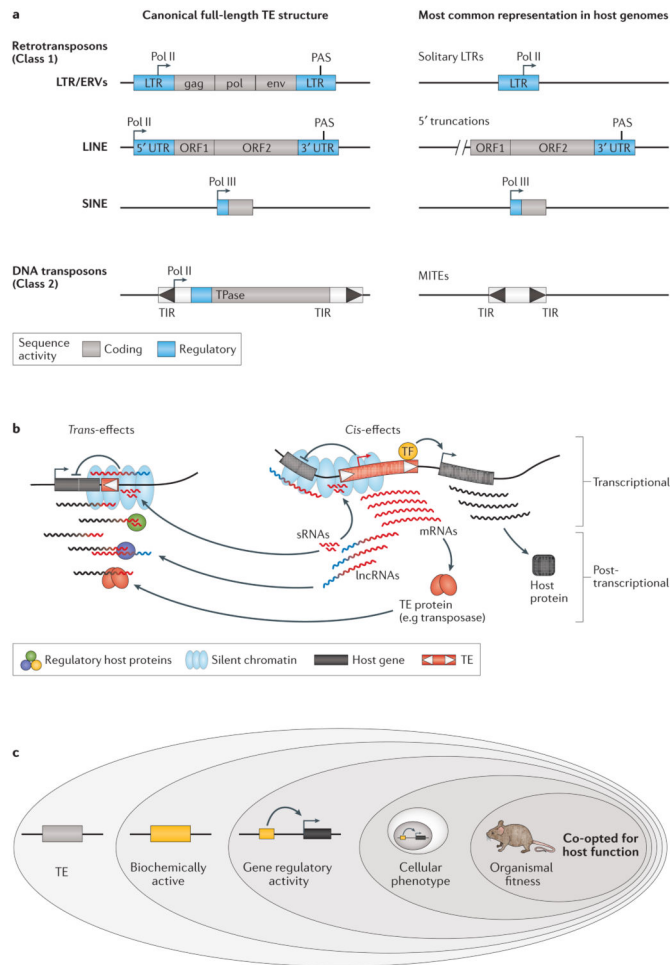


Figure 1. Origins of TE regulatory activities and how they may impact host genes
a | Schematic of major transposable element (TE) classes and their typical genetic organization. The left panel depicts the idealized full-length version of each TE type. Most TEs harbour regulatory sequences that function to promote their own transcription and regulation, such as promoters for RNA polymerase II (Pol II) or RNA polymerase III (Pol III), and polyadenylation signals (PAS). The right panel shows the structures of each type of TEs as they most commonly occur in the genome, which can differ substantially based on the TE (see the main text for details). **b** | Diagram depicting different types of regulatory activities exerted by TEs. These include effects mediated by *cis*-regulatory DNA and RNA elements (right panel) as well as *trans* effects mediated by TE-produced non-coding RNAs and proteins (left panel). **c** | Hierarchy of evidence to consider when determining if a TE has been co-opted for host functions. Many TEs exhibit biochemical hallmarks of regulatory activity based on genome-wide assays. However, additional evidence is required to determine which of these TEs alter the regulation of host genes and affect organismal phenotype and fitness. Abbreviations: LTRs: long terminal repeats; ERVs: endogenous retrovirus; MITEs: miniature inverted-repeat transposable elements; sRNAs: small RNAs; TIR: terminal inverted repeats.

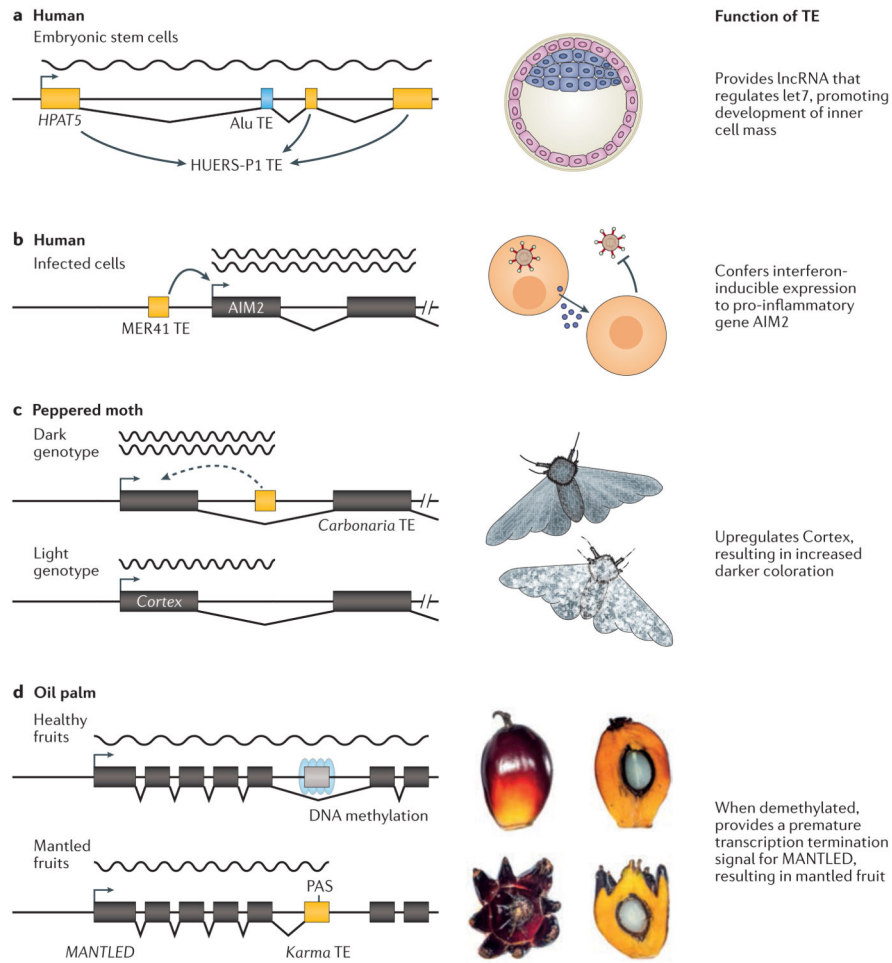


Figure 2. Examples of phenotypes driven by TE regulatory activity

a | The human pluripotency-associated transcript 5 (*HPAT5*) is a long non-coding RNA (lncRNA) that is derived from a composite of a HUERS-P1 endogenous retrovirus (ERV) element and an *Alu* short interspersed nuclear element (SINE). In all figure parts, wavy lines indicate pre-mRNA transcripts and angled lines indicate spliced introns. *HPAT5* regulates the *let7* family of microRNAs through *let7* binding sites carried by the *Alu* element, and was shown to be essential for inner cell mass formation during early embryonic development¹⁶⁵. **b** | A MER41 transposable element (TE) provides an interferon-inducible enhancer upstream of the human *AIM2* gene, which regulates inflammation in response to infection⁴⁴. **c** | In the peppered moth, a polymorphic *Carbonaria* TE insertion within an intron of the *Cortex* gene enhances *Cortex* expression levels (dotted line indicates an uncharacterized regulatory mechanism) which underlay adaptive cryptic colouration during the industrial revolution⁸³. **d** | In oil palm, sporadic demethylation of a *Karma* TE within an intron of the *MANTLED* gene results in unmasking of a cryptic splice acceptor site and premature termination signal, causing the mantled fruit phenotype¹⁴⁴. Left panel depicts genomic locus (not drawn to scale). The fruit images in part d are reproduced from ref¹⁴⁴

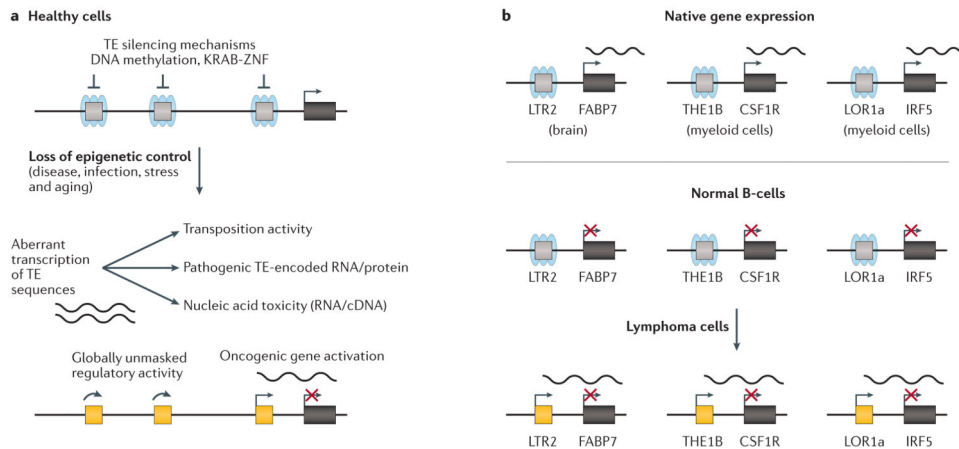


Figure 3. TEs can be aberrantly unmasked to promote disease states

a | Epigenetic perturbations can result in global transposable element (TE) reactivation and pathogenic consequences. The repressive epigenetic marks that normally silence TE transcription include DNA methylation and binding by KRAB zinc finger (ZNF) transcription factors, and these can be depleted upon various stresses. The reactivation of TE sequences across the genome can result in a wide variety of pathogenic consequences, including genome instability via transposition, other pathogenic activities of TE-encoded peptides or non-coding RNAs, and cellular toxicity due to buildup of RNA or cDNA intermediate molecules. **b** | Studies examining B-cell lymphomas have revealed oncogenic activation of colony-stimulating factor 1 receptor (*CSF1R*) (driven by a *THE1B* promoter in Hodgkin Lymphoma)¹³⁵, fatty acid-binding protein gene (*FABP7*) (driven by a *LTR2* promoter in diffuse large B-cell lymphoma)¹³⁶, and interferon regulatory factor 5 (*IRF5*) (driven by a *LOR1a* promoter in Hodgkin Lymphoma)¹³⁷. All these examples involve long terminal repeat (LTR) or endogenous retrovirus (ERV) elements, highlighting again the proclivity of this class of TEs to retain potent but generally repressed *cis*-regulatory activity in the human genome.

Table 1

Case examples of TEs functioning as *cis*-regulatory DNA elements.

Species	Gene product	TE	<i>Cis</i> -regulatory activity	Function	Evidence	References
Tetrapod vertebrates	ISL1	LF-SINE (vertebrate SINE)	Neural enhancer	Brain development (inferred)	Sequence conservation, mouse reporter assay	60
Human	AIM2	MER41 (primate ERV)	Interferon-inducible enhancer	Regulates inflammatory response	CRISPR knockout in human cell culture	44
Human	β -globin	ERV9 (primate ERV)	Erythroid enhancer	Controls developmental switch from fetal to adult globin	Cre-LoxP knockout in transgenic mouse BAC, chromatin looping (3C)	182
Mouse	Dicer	MT-C (rodent LTR retrotransposon)	Oocyte-specific promoter	Necessary for female oocyte function and fertility	TALLEN knockout in mouse	75
Mouse	Growth Hormone	B2 SINE (mouse retrotransposon)	Insulator	Developmental regulation of growth hormone (inferred)	Enhancer blocking assay	183
Human	Prolactin	MER39 (ERV)	Endometrial-specific promoter	Prolactin expression during pregnancy	5' RACE on endometrial tissue	184
Fly (<i>D. melanogaster</i>)	Cyp6g1	Accord (LTR retrotransposon)	Enhancer	Increased resistance to DDT insecticide	Selective sweep, genetic mapping, reporter assay	185
Fly (<i>D. melanogaster</i>)	Jheh1/2	Bari1	Antioxidant response elements (enhancer)	Increased resistance to oxidative stress	Selective sweep, phenotypic assays	186
Fly (<i>D. simulans</i>)	Slowpoke	Sheldler (LTR retrotransposon)	Altered splicing	Courtship song variation	Trait mapping, in vivo CRISPR knockout	187
Maize	TB1	Hopscoich (teosinte LTR retrotransposon)	Enhancer	Apical dominance branching pattern	Trait mapping, reporter assay in maize leaf protoplast	77
Sicilian blood orange	Ruby	Tcs1 (LTR retrotransposon)	Cold-responsive promoter	Required for cold-induced accumulation of anthocyanin	Trait mapping, 5' RACE	78

Further examples can be found in the Catalogue of genes affected by TEs (C-GATE) <https://sites.google.com/site/tecatalog/> (Ref. 36). 3C, chromatin conformation capture; BAC, bacterial artificial chromosome; ERV, endogenous retrovirus; LF-SINE, living-fossil SINE; LTR, long terminal repeat; RACE, rapid amplification of cDNA ends; SINE, short interspersed nuclear element; TALLEN, transcription activator-like effector nuclease.