Transcript assembly and quantification by RNA-Seq reveals unannotated transcripts and isoform switching during cell differentiation

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High-throughput mRNA sequencing (RNA-Seq) promises simultaneous transcript discovery and abundance estimation1–3. However, this would require algorithms that are not restricted by prior gene annotations and that account for alternative transcription and splicing. Here we introduce such algorithms in an open-source software program called Cufflinks. To test Cufflinks, we sequenced and analyzed >430 million paired 75-bp RNA-Seq reads from a mouse myoblast cell line over a differentiation time series. We detected 13,692 known transcripts and 3,724 previously unannotated ones, 62% of which are supported by independent expression data or by homologous genes in other species. Over the time series, 330 genes showed complete switches in the dominant transcription start site (TSS) or splice isoform, and we observed more subtle shifts in 1,304 other genes. These results suggest that Cufflinks can illuminate the substantial regulatory flexibility and complexity in even this well-studied model of muscle development and that it can improve transcriptome-based genome annotation.

Recently, RNA-Seq has revealed tissue-specific alternative splicing4, novel genes and transcripts5 and genomic structural variations6. Deeply sampled RNA-Seq permits measurement of differential gene expression with greater sensitivity than expression7 and tiling8 microarrays. However, the analysis of RNA-Seq data presents major challenges in transcript assembly and abundance estimation, arising from the ambiguous assignment of reads to isoforms9–10.

In earlier RNA-Seq experiments conducted by some of us, we estimated the relative expression for each gene as the fraction of reads mapping to its exons after normalizing for gene length11. We did not attempt to allocate reads to specific alternate isoforms, although we found ample evidence that multiple splice and promoter isoforms are often coexpressed in a given tissue12. This raised biological questions about how the different forms are distributed across cell types and physiological states. In addition, our prior methods relied on annotated gene models that, even in mouse, are incomplete. Longer reads (75 bp in this work versus 25 bp in our previous work) and pairs of reads from both ends of each RNA fragment can reduce uncertainty in assigning reads to alternative splice variants13. To produce useful transcript-level abundance estimates from paired-end RNA-Seq data, we developed a new algorithm that can identify complete novel transcripts and probabilistically assign reads to isoforms.

For our initial demonstration of Cufflinks, we performed a time course of paired-end 75-bp RNA-Seq on a well-studied model of skeletal muscle development, the C2C12 mouse myoblast cell line13 (see Online Methods). Regulated RNA expression of key transcription factors drives myogenesis, and the execution of the differentiation process involves changes in expression of hundreds of genes14,15. Previous studies have not measured global transcript isoform expression; however, there are well-documented expression changes at the whole-gene level for a set of marker genes in this system. We aimed to establish the prevalence of differential promoter use and differential splicing, because such data could reveal much about the model system’s regulatory behavior. A gene with isoforms that code for the same protein may be subject to complex regulation to maintain a certain level of output in the face of changes in expression of its transcription factors. Alternatively, genes with isoforms that encode different proteins could be functionally specialized for different cell types or states. By analyzing changes in the relative abundances of transcripts produced by the alternative splicing of a single primary transcript, we hoped to infer the effects of post-transcriptional processing (for example, splicing) on RNA output separately from rates of primary transcription. Such analysis could identify key genes in the system and suggest experiments to establish how they are regulated.

We first mapped sequenced fragments to the mouse genome using an improved version of TopHat16, which can align reads across splice junctions without relying on gene annotation (Supplementary Methods, section 2). A fragment corresponds to a single cDNA molecule, which can be represented by a pair of reads from each end. Out of 215 million fragments, 171 million (79%) mapped to the genome, and 46 million spanned at least one putative splice
Of the splice junctions spanned by fragment alignments, 70% were present in transcripts annotated by the UCSC, Ensembl or VEGA groups (known genes).

To recover the minimal set of transcripts supported by our fragment alignments, we designed a comparative transcriptome assembly algorithm. Expressed sequence tag (EST) assemblers such as PASA introduced the idea of collapsing alignments to transcripts on the basis of splicing compatibility, and Dilworth’s theorem has been used to assemble a parsimonious set of haplotypes from virus population sequencing reads. Cufflinks extends these ideas, reducing the transcript assembly problem to finding a maximum matching in a weighted bipartite graph that represents compatibilities among fragments (Fig. 1a–c and Supplementary Methods, section 4). Noncoding RNAs and microRNAs have been reported to regulate cell differentiation and development, and coding genes are known to produce noncoding isoforms as a means of regulating protein levels through nonsense-mediated decay. For these biologically motivated reasons, the assembler does not require that assembled transcripts contain an open reading frame (ORF). As Cufflinks does not make use of existing gene annotations during assembly, we validated the transcripts by first comparing individual time point assemblies to existing annotations.

We recovered a total of 13,692 known isoforms and 12,712 new isoforms of known genes. We estimate that 77% of the reads originated from previously known transcripts (Supplementary Table 2). Of the new isoforms, 7,395 (58%) contain novel splice junctions, with the remainder being novel combinations of known splicing outcomes; 11,712 (92%) have an ORF, 8,752 of which end at an annotated stop codon. Although we sequenced deeply by current standards, 73% of the moderately abundant transcripts (15–30 expected fragments per kilobase of transcript per million fragments mapped, abbreviated FPKM; see below for further explanation) detected at the 60-h time point with three lanes of GAII transcriptome sequencing were fully recovered with just a single lane. Because distinguishing a full-length transcript from a partially assembled fragment is difficult, we conservatively excluded from further analyses the novel isoforms that were unique to a single time point. Out of the new isoforms, 3,724 were present in multiple time points, and 581 were present at all time points; 6,518 (51%) of the new isoforms and 2,316 (62%) of the multiple time point novel isoforms were tiled by high-identity paired-end RNA-Seq. Cufflinks treats each pair of fragment reads as a single alignment. The algorithm assembles overlapping ‘bundles’ of fragment alignments separately, which reduces running time and memory use, because each bundle typically contains the fragments from no more than a few genes. Cufflinks then estimates the abundances of the assembled transcripts. The first step in fragment assembly is to identify pairs of ‘incompatible’ fragments that must have originated from distinct spliced mRNA isoforms. Fragments are connected in an ‘overlap graph’ when they are compatible and their alignments overlap in the genome. Each fragment has one node in the graph, and an edge, directed from left to right along the genome, is placed between each pair of compatible fragments. In this example, the yellow, blue and red fragment pairs must have originated from separate isoforms, but any other fragment could have come from the same transcript as one of these three. Isoforms are then assembled from the overlap graph. Paths through the graph correspond to sets of mutually compatible fragments that could be merged into complete isoforms. The overlap graph here can be minimally ‘covered’ by three paths (shaded in yellow, blue and red), each representing a different isoform. Dilworth’s Theorem states that the number of mutually incompatible reads is the same as the minimum number of transcripts needed to ‘explain’ all the fragments. Cufflinks implements a proof of Dilworth’s Theorem that produces a minimal set of paths that cover all the fragments in the overlap graph by finding the largest set of reads with the property that no two could have originated from the same isoform. Next, transcript abundance is estimated. Fragments are matched (denoted here using color) to the transcripts from which they could have originated. The violet fragment could have originated from the blue or red isoform. Gray fragments could have come from any of the three shown. Cufflinks estimates transcript abundances using a statistical model in which the probability of observing each fragment is a linear function of the abundances of the transcripts from which it could have originated. Because only the ends of each fragment are sequenced, the length of each may be unknown. Assigning a fragment to different isoforms often implies a different length for it. Cufflinks incorporates the distribution of fragment lengths to help assign fragments to isoforms. For example, the violet fragment would be much longer, and very improbable according to the Cufflinks model, if it were to come from the red isoform instead of the blue isoform. Last, the program numerically maximizes a function that assigns a likelihood to all possible sets of relative abundances of the yellow, red and blue isoforms, producing the abundances that best explain the observed fragments, shown as a pie chart.
EST alignments or matched RefSeq isoforms from other organisms, and end point RT-PCR experiments confirmed new isoforms in genes of interest (Supplementary Table 3). We concluded that most of the unannotated transcripts we found are in the myogenic transcriptome and that the mouse annotation remains incomplete.

To estimate transcript abundances, we first selected a set of 11,079 genes containing 17,416 high-confidence isoforms (Supplementary Data 1). Of these, 13,692 (79%) were known, and the remaining 3,724 (21%) were novel isoforms of known genes present in multiple time points. We then developed a statistical model specifying the probability of observing an RNA-Seq fragment. The model is parameterized by the abundances of these transcripts (Fig. 1d–f and Supplementary Methods, section 3). Cufflinks’ model allows for the probabilistic deconvolution of RNA-Seq fragment densities to account for cases in which genome alignments of fragments do not uniquely correspond to source transcripts. The model incorporates minimal assumptions about the sequencing experiment, and it extends the unpaired read model of Jiang and Wong to the paired-end case.

Abundances were reported in FPKM. In these units, the relative abundances of transcripts are described in terms of the expected biological objects (fragments) observed from an RNA-Seq experiment, which in the future may not be represented by single or paired reads. Confidence intervals for estimates were obtained using a Bayesian inference method based on importance sampling from the posterior distribution. Abundances of spiked control sequences ($R^2 = 0.99$) and benchmarks with simulated data ($R^2 = 0.96$) revealed that Cufflinks’ abundance estimates are highly accurate. The inclusion of novel isoforms of known genes during abundance estimation had a strong impact on the estimates of known isoforms in many genes ($R^2 = 0.90$), highlighting the importance of coupling transcript discovery together with abundance estimation.

We identified 7,770 genes and 10,480 isoforms undergoing significant abundance changes between some successive pair of time points (expected false discovery rate, abbreviated FDR, of <5%). Many genes show substantial transcript-level dynamics that are not reflected in their overall expression patterns (Supplementary Data 2). For example, Myc (Fig. 2a), a proto-oncogene that is known to be transcriptionally and post-transcriptionally regulated during myogenesis, is downregulated overall during the time course, and, although isoforms A and B follow this pattern, isoform C has a more complex expression pattern (Fig. 2b).

We noted that many genes showed switching between major and minor transcripts; furthermore, some contained isoforms with muscle-specific functions, such as tropomyosin I and II, which display a marked switch in isoform dominance upon differentiation (Supplementary Methods, appendix b). However, many genes featured dynamics involving several isoforms with behavior too complex to be deemed ‘switching’.

In light of these observations, we classified the patterns of expression dynamics for transcripts, assigning them one of four ‘trajectories’ based on their expression curves being flat, increasing, decreasing or mixed (see Online Methods). On the basis of this trajectory classification, a total of 1,634 genes were found to have multiple isoforms with different trajectories in the time course, and we hypothesized that differential promoter preference and differential splicing were responsible for the divergent patterns.

To explore the impact of regulation on mRNA output and to check whether it could explain the variability of trajectories, we grouped transcripts by their TSS instead of just by gene. Changes in the relative abundances of mRNAs spliced from the same pre-mRNA transcript are by definition post-transcriptional, so this grouping effectively discriminated changes in mRNA output associated with differential transcription from changes associated with differential post-transcriptional processing. Of the 3,486 genes in our high-confidence set with isoforms that shared a common TSS, 41% had TSS groups containing different isoform trajectories. Summing the expressions of isoforms...
sharing a TSS produces the trajectory for their primary transcript, and we identified 401 (48%) genes with multiple distinct primary transcript trajectories. However, trajectory classification was not precise enough to prioritize further investigation into individual genes and could not form the basis for statistical significance testing.

We therefore formalized and quantified divergent expression patterns of isoforms within and between TSS groups with an information-theoretic metric derived from the Jensen-Shannon divergence. With this metric, relative transcript abundances are represented as points along a logarithmic spiral in a real Hilbert space, and as a result the distance between points measures the extent of change in relative expression. Quantification of expression change in this way revealed significant (FDR < 5%) differential transcriptional regulation and splicing in 882 of 3,486 (25%) and 273 of 843 (32%) candidate genes, respectively, with 70 genes showing both types of differential regulation (Supplementary Table 4). Myc (Fig. 2a,b) undergoes a shift in transcriptional regulation of transcript abundances to post-transcriptional control of abundances (Fig. 2c) between 60 h and 90 h, as myocytes are beginning to fuse into myotubes.

Focusing on the genes with significant promoter and isoform changes (FDR < 5%), we noted that in many cases changes in relative abundance reflected switch-like events in which there was an inversion of the dominant primary transcript. For example, in the gene encoding FHL3, a transcriptional regulator recently reported to inhibit myogenesis,26 Cufflinks assembled the known isoform and another with a novel start site. We validated the 5′ exon of this isoform along with other novel start sites and splicing events by form-specific RT-PCR (Fig. 3a and Supplementary Methods, section 4). Limiting analysis to known isoforms would have produced an incorrect abundance estimate for the known isoform of FHL3. Moreover, the novel isoform is dominant before differentiation, so this potentially important differentiation–associated promoter switch would have been missed (Fig. 3b). In total, we tested and validated 153 of 185 putative novel TSSs by comparison against TAF1 and RNA polymerase II chromatin immunoprecipitation (ChIP)-Seq peaks.

We also observed switches in the major isoform of alternatively spliced genes. In total, 10% of multi-promoter genes featured a switch in major primary transcript, and 7% of alternatively spliced primary transcripts switched major isoforms. We concluded that not only does promoter switching have a substantial impact on mRNA output, but also many genes show evidence of post-transcriptionally induced expression changes, supporting a role for dynamic splicing regulation in myogenesis. A key question is whether genes that show divergent expression patterns of isoforms are differentially regulated in a particular system because they have isoforms that are functionally specialized for that system. Of the genes undergoing transcriptional or post-transcriptional isoform switches, 26% and 24%, respectively, encode multiple distinct proteins according to annotation.

We excluded genes with novel isoforms from the coding sequence analysis, so this fraction probably underestimates the impact of differential regulation on coding potential. We thus speculate that differential RNA level isoform regulation, whether transcriptional, post-transcriptional or mixed in underlying mechanism, suggests functional specialization of the isoforms in many genes.

Although Cufflinks was designed to investigate transcriptional and splicing regulation in this experiment, it is applicable to a broad range of RNA-Seq studies (Fig. 4). The open-source software runs on commonly available and inexpensive hardware, making it accessible to any researcher using RNA-Seq data. We are currently exploring the use of the Cufflinks assembler to annotate genomes of newly sequenced organisms and to quantify the effect of various mechanisms of gene regulation on expression. When coupled with assays of upstream regulatory activity, such as chromatin-state mapping or promoter occupancy, Cufflinks should help unveil the range of mechanisms governing RNA manufacture and processing.

METHODS

Methods and any associated references are available in the online version of the paper at http://www.nature.com/naturebiotechnology/.

Accession code. NCBI Gene Expression Omnibus: The data discussed in this publication have been deposited with accession number GSE20846.

Note: Supplementary information is available on the Nature Biotechnology website.

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AUTHOR CONTRIBUTIONS

C.T. and L.P. developed the mathematics and statistics and designed the algorithms; B.A.W. and G.K. performed the RNA-Seq and B.A.W. designed and executed experimental validations; C.T. implemented Cufflinks and Cuffdiff; G.P. implemented CuffCompare; M.J.v.B. and A.M. tested the software; C.T., G.P. and A.M. performed the analysis; L.P., A.M. and B.J.W. conceived the project; C.T., L.P., A.M., B. J.W. and S.L.S. wrote the manuscript.

COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

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Preparation of cDNA followed the procedure described previously, with minor modifications as described below. Before fragmentation, a 7-µl aliquot (total mass ~500 pg) containing known concentrations of seven ‘spiked in’ control transcripts from *Arabidopsis thaliana* and the lambda phage genome were added to a 100-ng aliquot of mRNA from each time point. This mixture was then fragmented to an average length of 200 nucleotides by metal-ion and heat-catalyzed hydrolysis. The hydrolysis was performed in a 25-µl volume at 94 °C for 90 s. The 5× hydrolysis buffer components were 200 mM Tris acetate, pH 8.2, 500 mM potassium acetate and 150 mM magnesium acetate. After removal of hydrolysis ions by G50 Sephadex filtration (USA Scientific, catalog no. 1451-1602), the fragmented mRNA was randomly primed with hexamers and reverse-transcribed using the Super Script II cDNA synthesis kit (Invitrogen, catalog no. 11917010). After second-strand synthesis, the cDNA went through end-repair and ligation reactions according to the Illumina ChiP-Seq genomic DNA preparation kit protocol (Illumina, catalog no. IP102-1001), using the paired-end adapters and amplification primers (Illumina, catalog no. PE102-1004). Ligation of the adapters adds 94 bases to the length of the cDNA molecules.

**Size selection.** The cDNA library was size-fractionated on a 2% TAE low-melt agarose gel (Lonza, catalog no. 500080), with a 100-bp ladder (Roche, catalog no. 14703220) run in adjacent lanes. Before loading of the gel, the ligated cDNA library was taken over a G50 Sephadex column to remove excess salts that interfere with loading the sample in the wells. After staining of the gel in ethidium bromide, a narrow slice (~2 mm) of the cDNA lane centered at the 300-bp marker was cut. The slice was extracted using the QiaEx II kit (Qiagen, catalog no. 202021), and the extract was filtered over a Microcon YM-100 microconcentrator (Millipore, catalog no. 42409) to remove DNA fragments shorter than 100 bp. Filtration was performed by pipetting the extract into the upper chamber of a microconcentrator and adding ultra-pure water (Gibco, catalog no. 10977) to a final volume of 500 µl. The filter was spun at 500g until only 50 µl remained in the upper chamber (about 20 min per spin) and then the upper chamber volume was replenished to 500 µl. This procedure was repeated six times. The filtered sample was then recovered from the filter chamber according to the manufacturer’s protocol. Fragment-length distributions obtained after size selection were estimated from the spike-in sequences and are shown in Supplementary Figure 1.

**Amplification.** One-sixth of the filtered sample volume was used as template for 15 cycles of amplification using the paired-end primers and amplification reagents supplied with the IlluminaChiP-Seq genomic DNA prep kit. The amplified product was cleaned up over a Qiaquick PCR column (Qiagen, catalog no. 28104), and then the filtration procedure using the Microcon YM-100 microconcentrators described above was repeated, to remove both amplification primers and amplification products shorter than 100 bp. A final pass over a G50 Sephadex column was performed, and the library was quantified using the Qubit fluorometer and PicoGreen quantification reagents (Invitrogen, catalog no. Q3283). The library was then used to build clusters on the Illumina flow cell according to protocol.

**Mapping cDNA fragments to the genome.** Fragments were mapped to build 37.1 of the mouse genome using TopHat version 1.0.13. We extended our previous algorithms to exploit the longer paired reads used in the study. TopHat version 1.0.7 and later splits a read 75 bp or longer in three or more segments of approximately equal size (25 bp) and maps them independently. Reads with segments that can be mapped to the genome only noncontiguously are marked as possible intron-spanning reads. These ‘contiguously unmappable’ reads are used to build a set of possible introns in the transcriptome. TopHat accumulates an index of potential splice junctions by examining segment mapping for all contiguously unmappable reads. For each junction, the program then concatenates 22 bp pairs upstream of the donor to 22 bp pairs downstream of the acceptor to form a synthetic spliced sequence around the junction. The segments of the contiguously unmappable reads are then aligned against these synthetic sequences with Bowtie. The resulting contiguous and spliced segment alignments for these reads are merged to form complete alignments to the genome, each spanning one or more splice junctions. Further details of how version 1.0.13 of TopHat differs from the published algorithm are provided in section 2 of the Supplementary Methods.

**Transcript abundance estimation.** We estimated transcript abundances using a generative statistical model of RNA-Seq experiments. The model was parameterized by the relative abundances of the set of all transcripts in a sample. For computational convenience, abundances of non-overlapping transcripts in disjoint genomic loci were calculated independently. The parameters of the model were the non-negative abundances $\rho_i$. Denoting the fragment distribution by $f$, we defined the effective length of a transcript to be:

$$ l(t) = \sum_{i=1}^n f(i)l(t)_i $$

where $l(t)_i$ is the length of a transcript. The likelihood function for our model was then given by:

$$ L(R) = \prod_{r \in R} \sum_{u \in T} \rho_u l(u) \left( \frac{F(I(r))}{I(t)_i - I(r)} \right) $$

where the products were over all fragment alignments $R$ and transcripts $T$ in the transcriptome, and $l(t)_i$ was the implied length of a fragment determined by a pair of reads assuming it originated from transcript $t$ (Supplementary Fig. 2). This is the likelihood function for a non-negative linear model, and therefore, the likelihood function had a unique maximum, which our implementation calculated via a numerical optimization procedure. Rather than reporting this estimate, we instead found the maximum *a posteriori* (MAP) estimate using a Bayesian inference procedure based on importance sampling from the posterior distribution. The proposal distribution we used was multivariate normal, with a mean given by the maximum likelihood estimate discussed above, and the variance-covariance matrix given by the inverse of the observed Fisher information matrix. The samples were also used to compute 95% confidence intervals for the MAP estimates. The MAP estimates and (associated confidence intervals) were used for differential expression testing.

Abundances were reported in FPKM (expected fragments per kilobase of transcript per million fragments sequenced). This unit is a scalar multiple of the parameters $\rho_i$, FPKM is conceptually analogous to the reads per kilobase per million reads sequenced (RPKM) measure, but it explicitly accommodates sequencing data with one, two or—if needed for future sequencing platforms—higher numbers of reads from single source molecules.

Abundance estimates were validated using spike-in sequences (Supplementary Fig. 3) and simulations (Supplementary Fig. 4). To confirm that all transcripts of a gene are necessary for accurate abundance estimation, novel transcripts were removed from the analysis (Supplementary Fig. 5), showing that resulting estimates may be biased.

**Transcript assembly.** Transcripts were assembled from the mapped fragments sorted by reference position. Fragments were first divided into non-overlapping loci, and each locus was assembled independently of the others using the Cufflinks assembler. The assembler was designed to find the minimal number of potential splice junctions by examining segment mapping for all contiguously unmappable reads. For each junction, the program then concatenates 22 bp pairs upstream of the donor to 22 bp pairs downstream of the acceptor to form a synthetic spliced sequence around the junction. The segments of the contiguously unmappable reads are then aligned against these synthetic sequences with Bowtie. The resulting contiguous and spliced segment alignments for these reads are merged to form complete alignments to the genome, each spanning one or more splice junctions. Further details of how version 1.0.13 of TopHat differs from the published algorithm are provided in section 2 of the Supplementary Methods.
graph’ $G$. A directed edge $(x,y)$ was placed between nodes $x$ and $y$ when the alignment for $x$ started at a lower coordinate than $y$, the alignments overlapped in the genome and the fragments were ‘compatible’ (Supplementary Fig. 6).

Compatibility was defined for overlapping fragments for which every implied intron in one fragment matched an identical implied intron in the other fragment. The resulting directed, acyclic graph was transitively reduced to produce $G$, to avoid including redundant path information. Cufflinks then found a minimum path cover of $G$, meaning that every fragment node was contained in some path in the cover, and the cover contained as few paths as possible. Each path in the cover corresponded to a set of mutually compatible fragments overlapping each other on the left and right (except initial and terminal fragments on the path). Dijkstra’s theorem implied that this path cover could be constructed by first finding the largest set of fragments with the property that no two are compatible. This set was determined by finding a maximum matching in a bipartite graph constructed from the transitive closure of $G$. The bipartite ‘reachability graph’ had a node in each partition for all fragments in $G$, and nodes were connected if there was a path between them in $G$. Given a maximum cardinality matching $M$, any fragment without an incident edge in $M$ was a member of an ‘antichain’. Each member of this antichain could be extended to a path, and this extension was a minimum path cover of $G$. The minimum cardinality chain decomposition computed using the approach described above was not guaranteed to be unique. To ‘phase’ distant exons, we leveraged the fact that abundance in homogeneities could link distant exons by their coverage. We therefore weighted the edges of the bipartite reachability graph on the basis of the percent-splitted-in metric introduced previously. Cufflinks arbitrated between multiple parsimonious assemblies by choosing the minimum-cost maximum matching in the reachability graph. In our setting, the percent-splitted-in $\psi$, for an alignment $x$ was computed by counting the alignments overlapping $x$ in the genome that were compatible with $x$, dividing by the total number of alignments that overlap $x$, and then normalizing for the length of the $x$. The cost $C(y, z)$ assigned to an edge between alignments $y$ and $z$ reflected the belief that they originated from different transcripts:

$$C(y, z) = -\log(1 - \frac{\psi_y - \psi_z}{2})$$

A useful feature of the Cufflinks assemblies is that they resulted in provably identifiable models. Complete details of the Cufflinks assembler are provided in the Supplementary Methods (section 4), along with proofs of several key theorems.

Structural comparison of time point assemblies. To validate Cufflinks transcripts (assembled transcript fragments) against annotated transcriptomes, and also to find transfrags common to multiple assemblies, we developed a tool called ‘Cuffcompare’ that builds structural equivalence classes of transcripts. We ran Cuffcompare on the assembly from each time point against the combined annotated transcriptomes of UCSC, Ensembl and VEGA (Supplementary Fig. 7). Because of the stochastic nature of sequencing, assembly of the same transcript in two different samples may result in transfrags of slightly different lengths. A Cufflinks transfrag was considered a complete match when there was a transcript with an identical chain of introns in the combined annotation. When no complete match was found between a Cufflinks transfrag and the transcripts in the combined annotation, Cuffcompare determined and reported whether there was another potentially significant relationship with any of the annotation transcripts that could be found in or around the same genomic locus.

Assembly and abundance robustness analysis. A total of 61,787,833 cDNA fragments were sequenced at 60 h. We mapped and assembled subsets of these fragments (at fractions 1/64, 1/32, 1/16, 1/8, 1/4 and 1/2 of the total) using TopHat and Cufflinks. Each assembly of parts of the data was compared to the assembly obtained with the full fragment set using Cuffcompare. We counted transcripts recovered in assemblies from partial data that structurally matched some transcripts in the assembly using all the reads. We assessed robustness of abundance estimation by counting the fraction of assembled transcripts that were assigned abundances within 15% of the FPKM value reported for the full fragment set transcript.

Simulation-based validation. To assess the accuracy of the Cufflinks estimates, we simulated an RNA-Seq experiment using the FluxSimulator, a freely available software package that models whole-transcriptome sequencing experiments with the Illumina Genome Analyzer. The software works by first randomly assigning expression values to the transcripts provided by the user, constructing an amplified, size-selected library, and then sequencing it. Mouse UCSC transcripts were supplied to the software, along with build 37.1 of the genome. FluxSimulator then randomly assigned expression levels to 18,935 UCSC transcripts. From these relative expression levels, the software constructed an in silico RNA-Seq sample, with each transcript assigned a number of library molecules according to its abundance. FluxSimulator produced 13,203,516 75-bp paired-end RNA-Seq reads from 6,601,805 library fragments, which were mapped with TopHat to the mouse genome using identical parameters to those used to map the C2C12 reads. A total of 6,176,961 fragments were mapped (93% of the library). These alignments were supplied along with the exact set of expressed transcripts to Cufflinks, to measure Cufflinks’ abundance estimation accuracy when working with a ‘perfect’ assembly.

Validation of novel transcription start sites. Transcripts with 5’ exons not in the UCSC, Ensembl or VEGA annotations were selected for validation. We excluded transcripts with estimated abundances of <0.5 FPKM at all time points, as well as those with a 5’ exon within 200 bp of an annotated exon. To validate our novel observed 5’ exons, we conducted ChIP-Seq experiments as described previously. Mouse UCSC, Ensembl or VEGA annotations were selected for validation. We also to find transcripts common to multiple assemblies, we developed a tool called ‘Cuffcompare’ that builds structural equivalence classes of transcripts. We ran Cuffcompare on the assembly from each time point against the combined annotated transcriptomes of UCSC, Ensembl and VEGA annotations were selected for validation. We then randomly assigned expression levels to 18,935 UCSC transcripts. From these relative expression levels, the software constructed an in silico RNA-Seq sample, with each transcript assigned a number of library molecules according to its abundance. FluxSimulator produced 13,203,516 75-bp paired-end RNA-Seq reads from 6,601,805 library fragments, which were mapped with TopHat to the mouse genome using identical parameters to those used to map the C2C12 reads. A total of 6,176,961 fragments were mapped (93% of the library). These alignments were supplied along with the exact set of expressed transcripts to Cufflinks, to measure Cufflinks’ abundance estimation accuracy when working with a ‘perfect’ assembly.

End point RT-PCR validation of novel isoforms. Six genes with multiple assembled splice isoforms were chosen as cases for end point PCR validation, including three with novel isoforms (Supplementary Figs. 8 and 9). Amplification primers that cross the Cufflinks-predicted spliced-exon junctions were purchased from Integrated DNA Technologies, Inc. For each time point, 5 μg of total RNA was primed with oligo(dT)20 (Invitrogen, catalog no. 18418020) and reverse transcribed at 50 °C using SuperScript III reverse transcriptase (Invitrogen, catalog no. 18080044) according to the manufacturer’s protocol. One-tenth of the cDNA reaction was used as template for 35 rounds of PCR amplification with each pair of junction-crossing primers. The PCR reactions were cleaned up using the Qiagick PCR cleanup kit (Qiaegen, catalog no. 28104) and quantified using a Nanodrop spectrophotometer. An equal mass (30 ng) of DNA from each reaction was then loaded in each lane of a 2.0% agarose gel, post-stained with Sybr Gold (Invitrogen, catalog no. S11494) and visualized on a UV transilluminator.

Analysis of gene expression and regulation dynamics. To test for divergent expression dynamics among isoforms, we tested all high-confidence isoforms for significant changes between each time point using the variance estimates produced by our statistical model (FDR < 5%). Trajectories were assigned to transcript expression curves on the basis of significant (FDR < 5%) increases or decreases in expression between consecutive time points. To be deemed significant, expression between consecutive time points also had to change by at least 25%. The possible trajectories were therefore reduced to 81 combinatorial possibilities (increasing, decreasing or flat between any of the three pairs of consecutive time points). Trajectories were then classified into four groups: increasing (three consecutive increases), decreasing (three consecutive decreases), flat (no changes) and mixed (presence of both increases and decreases in expression along the time course). To test for significant changes in the relative abundance of a group of transcripts, we calculated the square root of the Jensen-Shannon divergence on the relative abundances in each of two time points. The variance of this metric under the null hypothesis of no change in relative abundance can be estimated using the delta method from the variance-covariance matrix on abundances estimates. Using the estimated variance of the Jensen-Shannon metric, we applied a one-sided t-test for significant changes in relative abundance of transcripts grouped by TSS and also primary transcripts grouped by gene. Type I errors were controlled with the Benjamini-Hochberg correction for multiple testing of differential expression, splicing and promoter preference throughout the analysis.
Figures 10 and 11 show examples of genes with significant changes in relative transcript abundances during the time course.

Software availability. TopHat (http://tophat.cbcb.umd.edu) is freely available as source code. It takes a reference genome (as a Bowtie® index) and RNA-Seq reads as FASTA or FASTQ and produces alignments in SAM® format. TopHat is distributed under the Artistic License and runs on Linux and Mac OS X.

The Cufflinks assembler and abundance estimation algorithms (http://cufflinks.cbcb.umd.edu/) are open-source C++ programs and are freely available in both source and binary. The package includes the assembler along with utilities to structurally compare Cufflinks output between samples (Cuffcompare) and to perform differential expression testing (Cuffdiff). Cufflinks is distributed under the Boost License and runs on Linux and Mac OS X. The source code for Cufflinks version 0.8.0 is provided in Supplementary Data 3.