Recent insights into Groucho co-repressor recruitment and function

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Gene expression is often controlled by transcriptional repressors during development. Many transcription factors lack intrinsic repressive activity but recruit co-factors that inhibit productive transcription. Here we discuss new insights and models for repression mediated by the Groucho/Transducin-Like Enhancer of split (Gro/TLE) family of co-repressor proteins.

**Drosophila** Groucho was the original member of the Groucho/Transducin-Like Enhancer of split (Gro/TLE) family of co-repressor proteins. This family is conserved across metazoans and includes four human (TLE1-4) and mouse Gro-related gene (Grg1-4) orthologues.1,2 Gro/TLE family proteins do not bind DNA directly, but act as repressive co-factors for a numerous and diverse range of sequence-specific DNA-binding transcription factors, including members of the Hes, Runx, Nkx, LEF1/Tcf, Pax, Six, and Fox families. Several transcription factors that recruit Gro/TLE proteins are effectors of cell fate specification pathways, including Notch, Wnt, NFkB and Estrogen Receptor signalling. Thus, Gro/TLE proteins participate in a diverse range of biological processes, including neurogenesis, somitogenesis, osteogenesis, haematopoiesis, stem cell maintenance, and proliferation. Human Gro family members have been linked to cancer pathogenesis. For example, epigenetic inactivation of the human TLE1 gene contributes to the development of hematologic malignancies,3 and Estrogen Receptor binding and transcriptional activity is regulated by TLE1 in breast cancer cells.4

Gro/TLE family members are characterized by highly conserved N-terminal and C-terminal domains.2 The conserved N-terminal domain is glutamine rich and so is known as the Q-domain. The Q-domain has been shown to mediate oligomerization of Gro/TLE proteins into tetramers and to facilitate binding of a subset of recruiting transcription factors. The highly conserved C-terminal domain contains WD-repeats and folds into a seven-blade β-propeller that binds many transcription factors.5

Although it is now over 20 years since it was first established that *Drosophila* Gro acts as a co-repressor,6 the precise molecular mechanisms underlying Gro/TLE-mediated repression have remained enigmatic. To gain new insight into Gro function, we took advantage of recent advances in genome-wide profiling technology and used chromatin immunoprecipitation (ChIP) followed by high-throughput sequencing (ChIP-seq) to determine how Gro is recruited to target genes and to identify factors associated with its binding. We opted to use cell lines (Kc167 and S2) to avoid the complications of interpreting data from mixed cell populations (e.g., embryos) where peaks obtained by ChIP may represent binding to overlapping or adjacent regulatory elements used in different cell types. In addition, use of Kc167 cells facilitated direct comparison of our data with the numerous published genome-wide datasets for transcriptional regulator binding and chromatin modifications in Kc167 cells from the modENCODE project and from the van Steensel group.8,9

**Drosophila** Gro was first described as mediating "long-range" repression (acting over distances of >1 kb) in 1997 in two independent studies that considered Gro
that the Gro binding represented a single continuous peak from the site of recruitment.

Our ChIP-seq analysis generated a series of observations in direct conflict with predictions made by the aforementioned “spreading” model in which Gro oligomerizes along several kilobases of DNA to mediate silencing at distant (>1 kb away) promoters. First, Gro typically bound the genome at discrete peaks spanning less than one kilobase; the median peak width was 708 bp in Kc167 cells, and 425 bp in S2 cells and fewer than 3% of peaks were greater than 2 kb. Second, oligomerization via the Q-domain did not contribute to the peak width of Gro recruitment. The average peak width bound by a GFP-tagged variant of Gro unable to oligomerize was actually slightly broader than that of GFP-(wild-type)Gro. Third, Gro was frequently found bound at annotated transcription start sites (TSSs); approximately 40% of Gro peaks overlapped TSSs in Kc167 cells (25% in S2 cells). Finally, genes that had Gro bound in peaks either upstream, at the TSS, or within the gene, were not completely silenced; these were genes that recruited RNA polymerase II (RNAP II) and expressed mRNA at some level.

From these observations, it is clear that Gro does not spread over multi-kilobase regions of chromatin in either Kc167 or S2 cells. Furthermore, meta-analysis of ChIP-seq data for the human Gro orthologue TLE3 in MCF7 cells revealed that the average peak width for TLE3 was not significantly different to that of Gro, indicating that it is recruited in a similar manner. However, we did notice that discrete peaks of Gro binding were frequently found in clusters (e.g., peaks within the Enhancer of split complex [E(spl)-C], Fig. 2) that could potentially be mistaken for longer continuous binding of Gro by low resolution mapping techniques (e.g., ChIP-qPCR, DamID). While it is clear that oligomerization has a role to play in Gro-mediated repression, our genome-wide binding data indicates that oligomerization does not influence the breadth of peaks of Groucho recruitment as had been previously proposed.

Another long-standing issue we attempted to address was the relationship between Gro and histone acetylation status. Gro has been shown to physically and genetically interact with the histone deacetylase HDAC1, referred to as Rpd3 in Drosophila. These observations led to the model that Gro recruits Rpd3 to deacetylate histones, generating chromatin that is not permissive for transcription. Support for this model came from a series of published reports studying ectopic Gro recruitment, though it had long been acknowledged that Gro repression could, at most, be only partially dependent on Rpd3 in vivo. Embryos with reduced maternal Rpd3 have a segmenta- tion phenotype that may reflect failure of repression by the products of segmentation genes that recruit Gro (e.g., Hairy, Eve, and Runt). However, embryos lacking maternal Rpd3 do not exhibit the characteristic proneural phenotype found

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**Figure 1.** Models for Gro repression. (A) The original “spreading” model (adapted from Chen and Courey) for long range action of Gro predicted the presence of broad domains (kilobases) of continuous Gro recruitment, which we did not observe in ChIP-seq analysis of Gro. In this model Gro recruits Rpd3 that leads to chromatin that is not acetylated. Speculative and simplified models based on our ChIP-seq analysis of how Gro may act both locally (B) and from a distance (C) to repress transcription. In both cases sequence-specific DNA-binding proteins recruit Gro to the chromatin, which somehow interacts (either directly or indirectly) with the RNP II complex to inhibit transcription when bound at the TSS (B), or via chromatin looping from a distance (C). Rpd3 co-localizes with Gro at a small majority of recruitment sites. Peaks of Gro binding are associated with hypacetylated histones, however acetylated histones are frequently observed flanking Gro peaks.
in embryos depleted of maternal Gro, resulting from failure of Gro-mediated repression by E(spl)-HLH proteins. Our ChIP-seq analysis revealed that a small majority (59%) of Gro peaks in Kc167 cells overlapped peaks of Rpd3 binding, largely consistent with the previous supposition that Gro repression involves Rpd3 in some, but not all, contexts. Gro peaks were generally associated with genomic regions where histones are not highly acetylated. However, depleting cells of Gro by RNAi did not lead to detectable changes in the pattern of histone acetylation at the sites where Gro bound in wild-type cells. This result held true at the E(spl)mβ-HLH locus where transcript production was significantly upregulated in gro-RNAi cells, indicating that increased histone acetylation at Gro targets is not absolutely required for derepression. However, we could not formally rule out the possibility that the residual levels of Gro left after RNAi treatment were sufficient to maintain histone hypoacetylation and Gro-mediated repression at most loci. Indeed, we saw relatively few significant changes in gene expression in gro-RNAi treated cells. Recent advances in genome engineering techniques including CRISPR and TALENS should, in the future, expedite the construction of gro<sup>−/−</sup> cells, allowing us to resolve the effect of Gro on Rpd3 recruitment and histone acetylation status at target loci.

ChIP-seq analysis of Gro recruitment in Kc167 cells also allowed us to look for additional factors, including histone modifications associated with Gro binding. Even though Gro acts as a repressor, we found that Gro peaks were most commonly associated with binding of factors and chromatin modifications found in "active chromatin" that is permissive for transcription. This observation was entirely consistent with results from Filion et al., who used the DamID technique to profile the genome-wide binding of 53 transcriptional regulators, including Gro. Moreover, Gro peaks were found at sites associated with DNase I hypersensitivity, a further indication that Gro binds in open, active chromatin.

Peaks of Gro at transcription start sites (TSSs) co-localize with RNAP II; therefore, the presence of Gro does not block RNAP II recruitment. However, we found evidence that Gro can alter RNAP II activity. The transition from transcription initiation to elongation involves a checkpoint that is a key point of control of the expression of many developmentally regulated genes (recently reviewed in refs. 22-24). When this checkpoint is rate-limiting, the amount of RNAP II observed around the TSS is higher than that through the gene body; this phenomenon is often described as promoter proximal RNAP II pausing. The "pause ratio" of a transcript is the ratio of total RNAP II at the TSS to that within the gene body. In Kc167 cells, almost half of transcripts where Gro is bound at the TSS have a very high pause ratio (in the top 10% of all transcripts). Furthermore, a majority of Gro peaks overlap peaks of GAGA Factor (GAF) binding, which has previously been linked to promoter proximal pausing at many genes in Drosophila. Although we did not detect any global effects on RNAP II pausing in gro-RNAi treated cells, loss of Gro led to decreased RNAP II pausing at the E(spl)mβ-HLH locus, a validated target of Gro repression in Kc167 cells.

RNAP II pausing has been proposed as a means of attenuating the expression of genes primed for rapid activation in response to environmental or developmental cues. It is therefore not surprising that Gro is found to associate with paused genes, as Gro is known to repress signal-responsive genes in the absence of both Notch and Wnt signalling. For example, in the absence of induced Notch signalling in Kc167 cells, loss of Gro alone was sufficient to cause reduced RNAP II pausing and increased transcript production at the E(spl)mβ-HLH locus. Thus, one action of Gro at E(spl)mβ-HLH is to promote RNAP II pausing. This raises the possibility that the many different sequence-specific DNA-binding transcription factors that recruit Gro may induce repression, at least in part, by promoting RNAP II pausing.

The current understanding of RNAP II pausing revolves largely around its release; much less is known about its establishment. Briefly, after transcribing approximately 30 nucleotides, RNAP II passes through a checkpoint where positive Transcription Elongation Factor b (P-TEFb) must phosphorylate specific amino acid residues in RNAP II, Spt5 (an RNAP II co-factor) and Negative Elongation Factor (NELF) for further transcription elongation to occur. A number of factors are known to stimulate P-TEFb activity to overcome RNAP II pausing and maximise transcription rates, including the Super Elongation Complexes (SECs) Brd4 and Myc. The factors involved in establishing paused RNAP II at specific genes are not so well understood. The 7SK snRNP inhibits P-TEFb activity, but it is not...
known if its actions are gene-specific or more general in Drosophila.28 GAF is associated with a subset of paused genes in Drosophila and is thought to promote RNAPII pausing through NELF recruitment.29 Interestingly, we found that the sequences recovered from Gro peaks were enriched for GAF binding motifs and that a high proportion of Gro peaks overlapped GAF peaks (82% at TSSs in Kc167 cells) raising the possibility that GAF and Gro collaborate to promote pausing. Further studies are required to determine if Gro inhibits P-TEFb activity by blocking activating factors, or if Gro recruits factors that impede P-TEFb activity and stabilise pausing.

Although we found many Gro peaks at TSSs, we also found Gro peaks further away from promoters in intergenic regions. Due to the failure to detect continuous regions of Gro bound from distant regulatory elements to promoters, we favour a model in which transcription factors by blocking activating factors, or if Gro recruits factors that impede P-TEFb activity and stabilise pausing.

To our initial surprise, the removal of Gro did not lead to a general up-regulation of target genes. However, on reflection, we realised that loss of repression and activation of gene expression are separable processes. We were examining stable populations of cells in culture that were not subject to changes in environmental conditions or cell-cell signalling that had the potential to induce immediate de novo expression and recruitment of activating factors to Gro target genes. Cells in a developing embryo are constantly bombarded with changing signals from neighbouring cells and are likely to be more vulnerable to a loss of repression causing rapid gene activation.

Genome-wide profiling of Gro recruitment in single cell types has led us to re-evaluate the old “spreading” model for Gro recruitment and function, but we have not been able to rule out the possibility that Gro recruits histone deacetylases to attenuate transcription. In addition, we have found that Gro is associated with paused genes and that it facilitates RNAPII pausing at a bona fide target gene. The molecular mechanisms through which Gro acts as a co-repressor for so many different transcription factors remain to be elucidated. However, this study shows that RNAPII pausing and histone acetylation status are likely to be two important aspects of Gro function. Future studies should address how Gro influences RNAPII pausing and histone acetylation status, and whether these two phenomena are linked. It will also be important to expand the set of cell types studied to understand if the patterns we observed in Kc167 and S2 cells are representative, or if the expression of other factors shifts the binding and function of Gro.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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