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research in this proposal addresses three aspects of gene regulation that might be applicable to the physiological control of cocaine addiction. First, the role of transcriptional regulation in heroin addiction will be investigated. An analysis of response elements present in the 5'-flanking regions of rat norepinephrine transporter (NEAT), tyrosine hydroxylase (TH), and dopamine beta-hydroxylase (DBH) genes will be completed. Also, the hypothesis that these genes are coordinately regulated will be tested with an analysis of promoter-reporter gene constructs. The second aspect will investigate transcriptional regulation by the activated glucocorticoid receptor (GR). Full-length and truncated forms of the murine GR gene will be cloned into different expression vectors, and the effect of the different GR genes on transcription of a sensitive tTA-tk promoter-luciferase reporter will be assessed. The third aspect will investigate transcriptional regulation by the oncogene, erbB-2. This will be accomplished by identifying the critical intragenic regions of the human erbB-2 gene with deletion mutagenesis

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