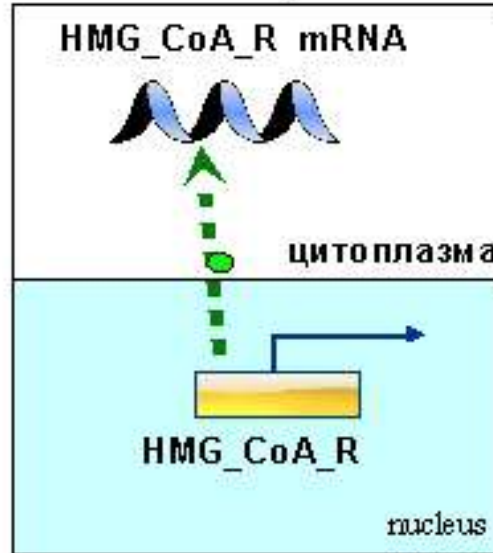
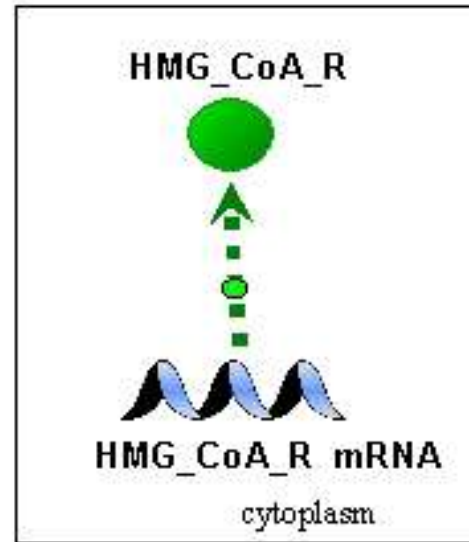


# The elementary processes in GENENET database

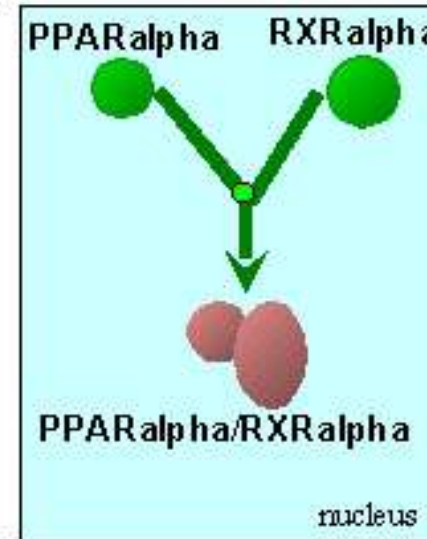
## Transcription



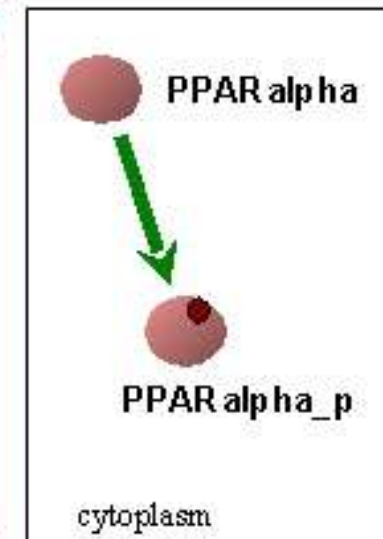
## Translation



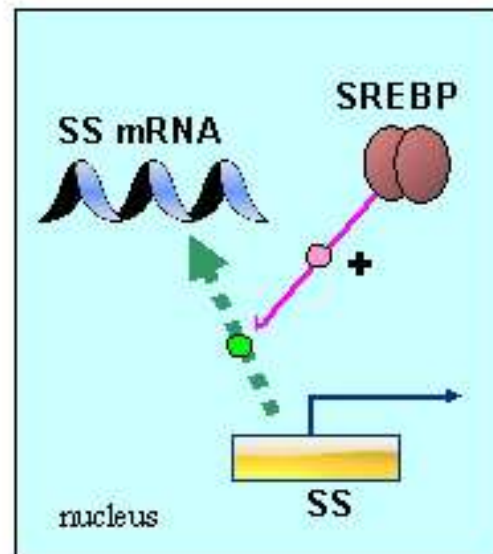
## Multimerization



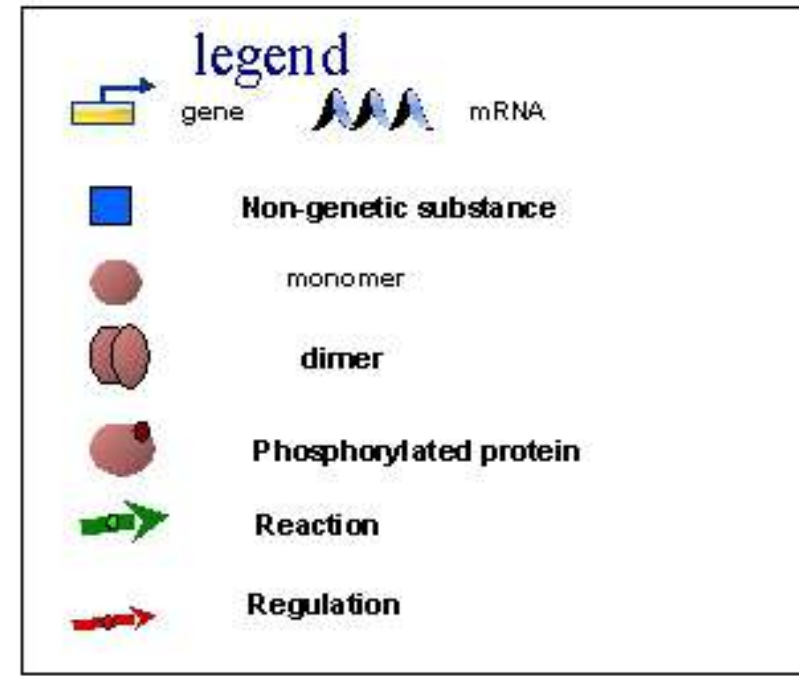
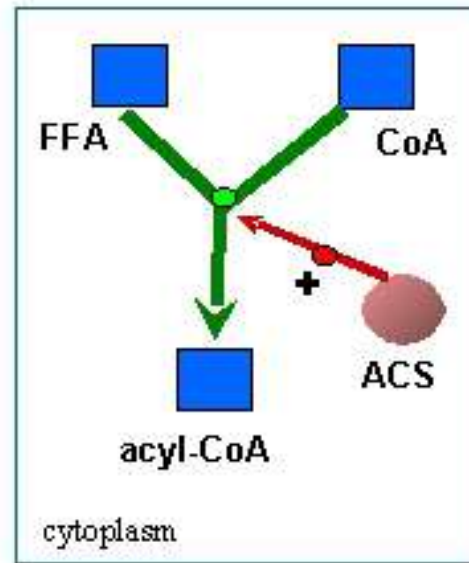
## Phosphorylation



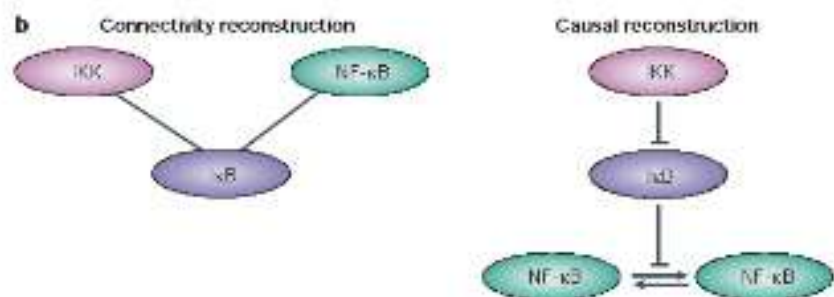
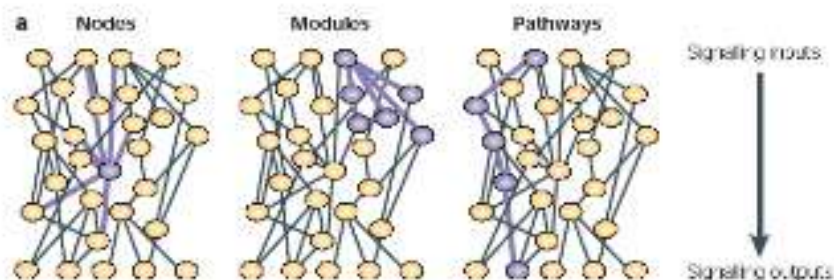
## Transcription activation



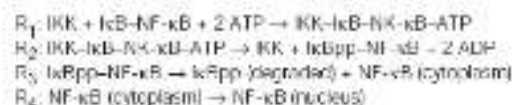
## reaction



# Reconstructing a gene network



**c** Stoichiometric reconstruction



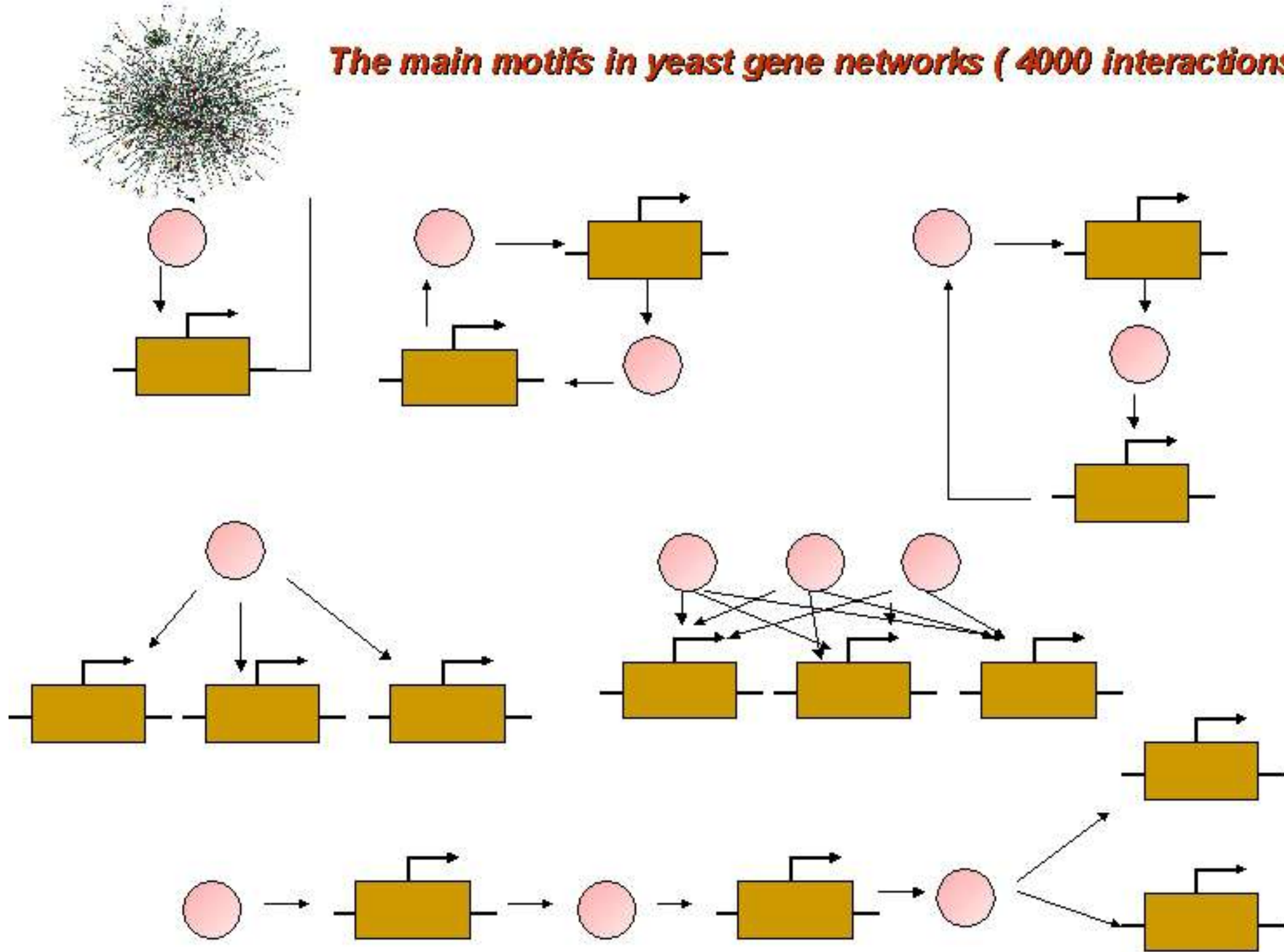
Compounds	Reactions			
	R <sub>1</sub>	R <sub>2</sub>	R <sub>3</sub>	R <sub>4</sub>
IKK	-1	+1	0	0
IKK complex	+1	-1	0	0
IκBpp-NF-κB	0	+1	-1	0
IκBpp (degraded)	0	0	-1	0
NF-κB (cytoplasm)	0	0	-1	-1
IκB-NF-κB	-1	0	0	0
ATP	-2	0	0	0
ADP	0	+2	0	0
NF-κB (nucleus)	0	0	0	+1



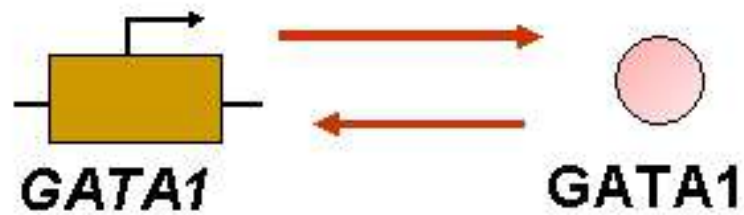
The most signalling network reconstructions focus on particular nodes, modules or pathways in given network. Network “nodes” describe the many interactions that a given compound (for example transcription factor NF-κB) participates in. Network “modules” consist of a group of related reaction that often incorporate feedback mechanisms. Network “pathways” connect a signalling input to signalling output.

There are three levels of resolution in reconstruction of NF-κB activation.

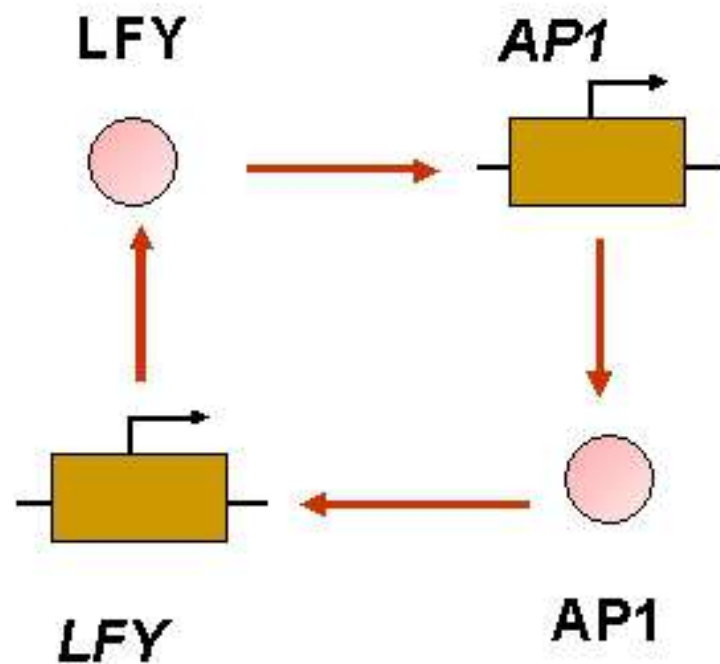
**The main motifs in yeast gene networks ( 4000 interactions)**



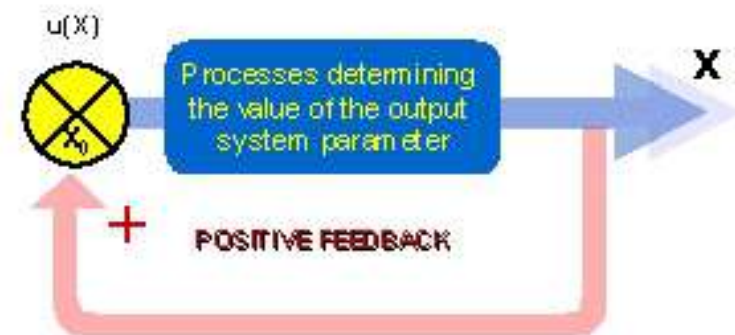
## Positive feedbacks



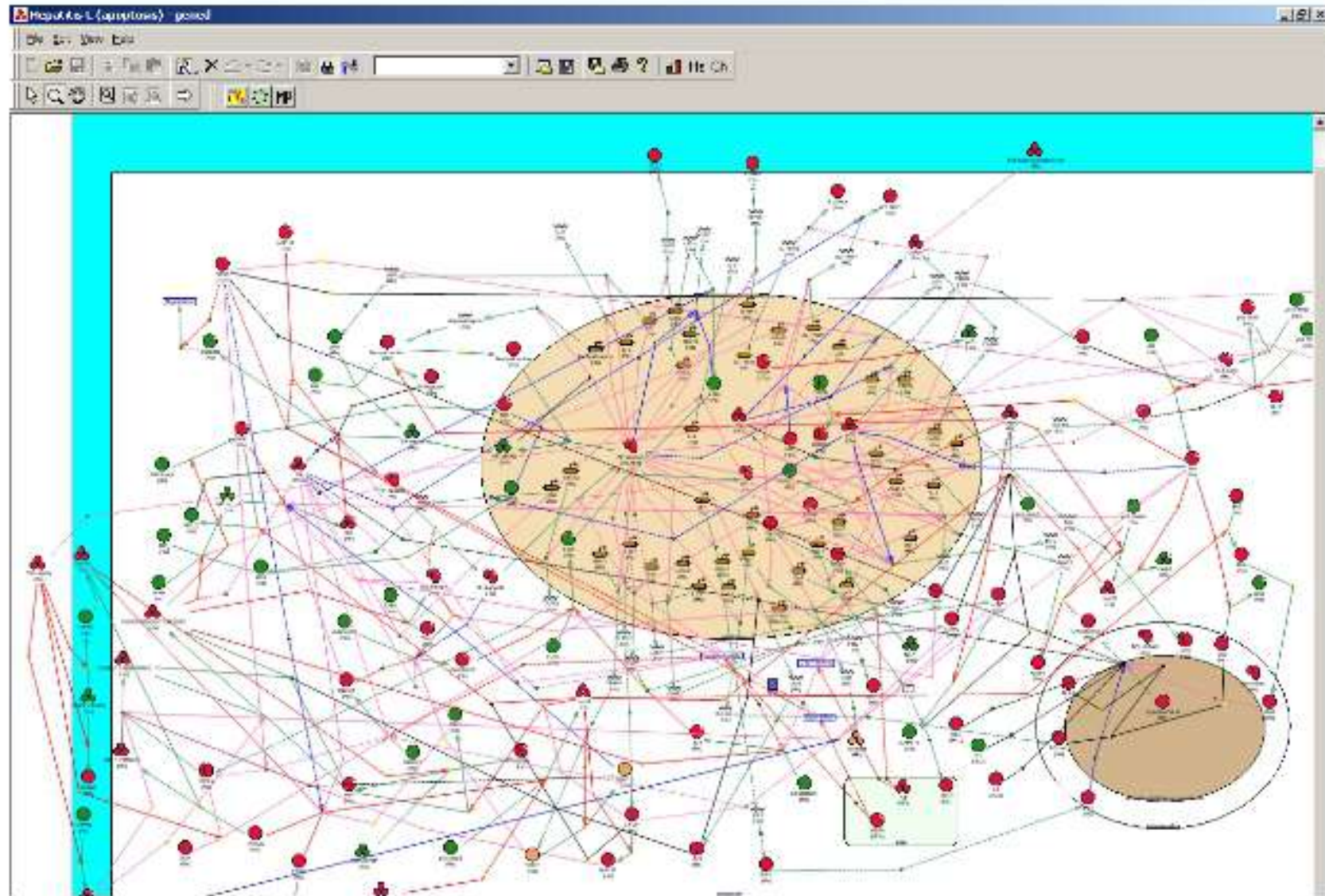
*Transcription factor activates its own gene*



*Cross effect of transcription factors with positive feedback*



## Apoptosis gene network in hepatitis C



Apoptosis plays a significant role in the pathogenesis of hepatitis C. This process may be viewed as a host defense mechanism against viral infection and hepatocarcinogenesis. Hepatitis C virus (HCV)-regulated gene network in apoptosis was reconstructed on the basis of the data extracted from 273 experimental papers. The 157 negative and positive autoregulatory circuits are clustered due to the coalescing nodes NF- $\kappa$ B/I $\kappa$ B, c-FLIP, TRAF2, and caspase 3.