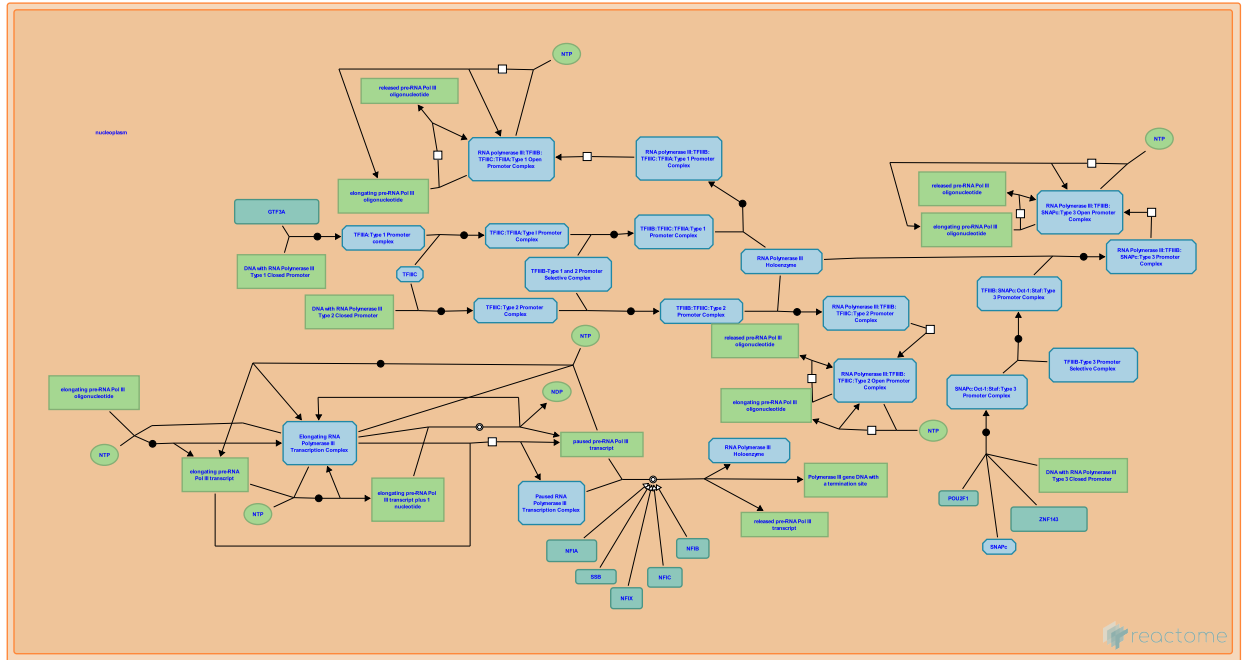


# RNA Polymerase III Transcription



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This is just an excerpt of a full-length report for this pathway. To access the complete report, please download it at the [Reactome Textbook](https://reactome.org/textbook/).

26/04/2024

## Introduction

Reactome is open-source, open access, manually curated and peer-reviewed pathway database. Pathway annotations are authored by expert biologists, in collaboration with Reactome editorial staff and cross-referenced to many bioinformatics databases. A system of evidence tracking ensures that all assertions are backed up by the primary literature. Reactome is used by clinicians, geneticists, genomics researchers, and molecular biologists to interpret the results of high-throughput experimental studies, by bioinformaticians seeking to develop novel algorithms for mining knowledge from genomic studies, and by systems biologists building predictive models of normal and disease variant pathways.

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## Literature references

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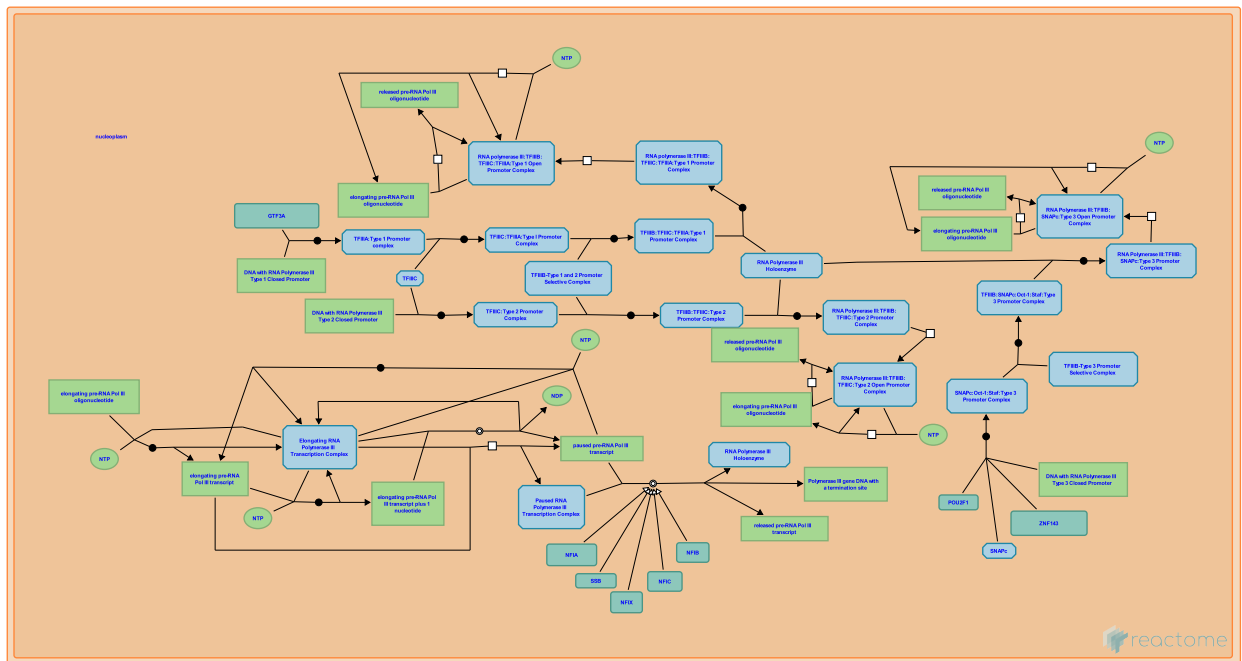
Reactome database release: 88

This document contains 5 pathways ([see Table of Contents](#))

# RNA Polymerase III Transcription ↗

Stable identifier: R-HSA-74158

Compartments: nucleoplasm



RNA polymerase III is one of three types of nuclear RNA polymerases present in eucaryotic cells. About 10% of the total transcription in dividing cells can be attributed to its activity. It synthesizes an eclectic collection of catalytic or structural RNA molecules, some of which are involved in protein synthesis, pre-mRNA splicing, tRNA processing, and the control of RNA polymerase II elongation, whereas some others have still unknown functions. Like other RNA polymerases, RNA polymerase III cannot recognize its target promoters directly. Instead it is recruited to specific promoter sequences through the help of transcription factors. There are three basic types of RNA polymerase III promoters, called types 1, 2, and 3 (Geiduschek and Kassavetis, 1992). Although in vivo, RNA polymerase III may be recruited to these promoters as part of a large complex (holo RNA polymerase III) containing the polymerase and its initiation factors (Wang et al., 1997), in vitro the reaction can be divided into several steps. First, the promoter elements are recognized by DNA binding factors, which then recruit a factor known as TFIIB. TFIIB itself then directly contacts RNA polymerase III. In human cells but not in *S. cerevisiae*, there are at least two versions of TFIIB. One contains TBP, Bdp1, and Brf1 (Brf1-TFIIB), and the other TBP, Bdp1, and Brf2 (Brf2-TFIIB) (Schramm et al., 2000; Teichmann et al., 2000).

## Literature references

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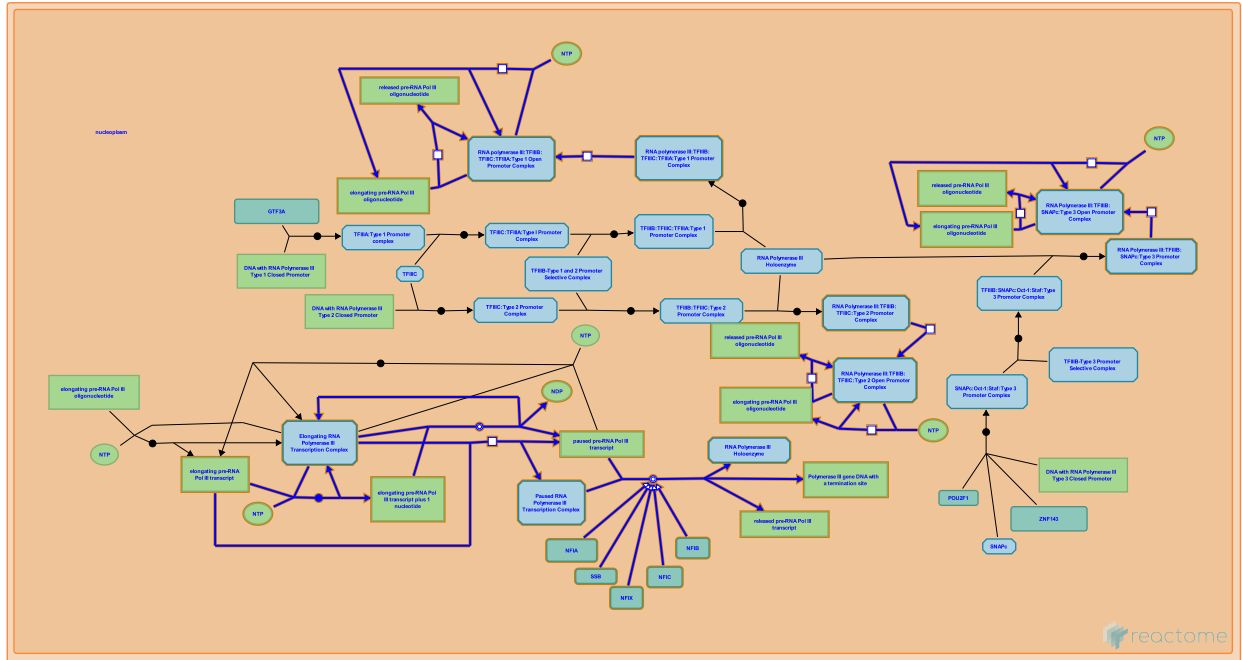
2003-09-11	Authored	Hernandez, N.
2024-03-06	Edited	Joshi-Tope, G.

# RNA Polymerase III Abortive And Retractive Initiation ↗

**Location:** RNA Polymerase III Transcription

**Stable identifier:** R-HSA-749476

**Compartments:** nucleolus



Abortive initiation, the repetitive formation of short oligonucleotides, is a ubiquitous feature of transcriptional initiation. This Pathway contains events inferred from events in *Saccharomyces cerevisiae*.

## Editions

2004-03-29

Authored

Geiduschek, EP.

2024-03-06

Edited

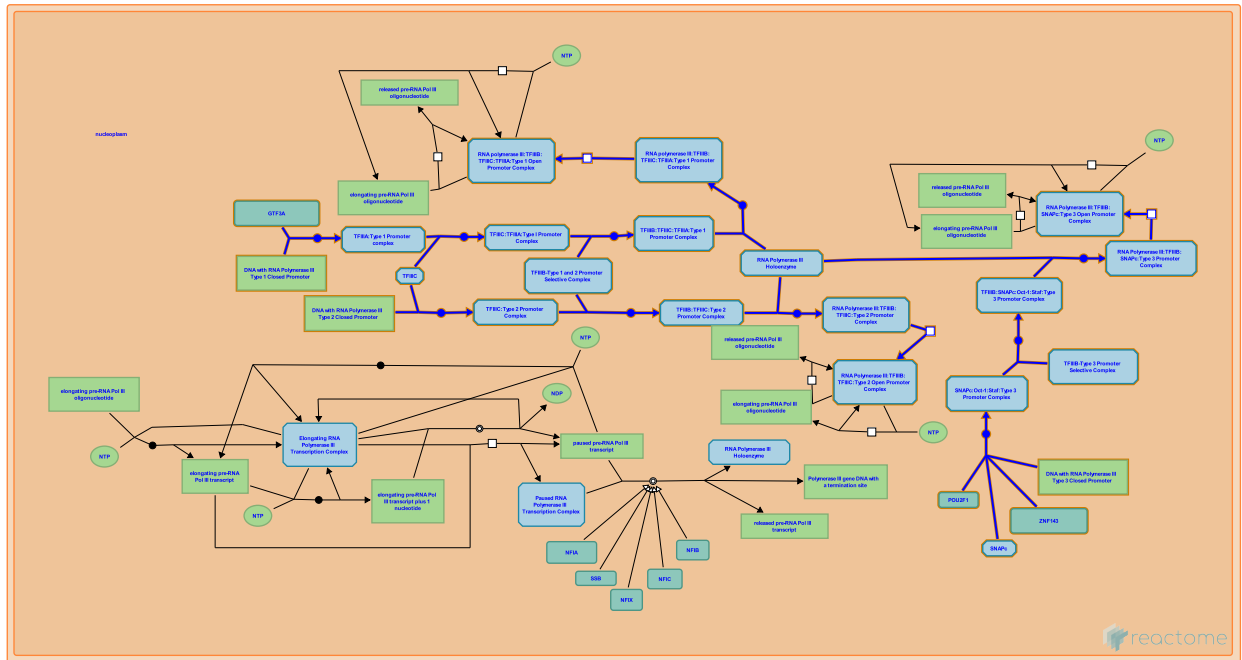
Gillespie, ME.

# RNA Polymerase III Transcription Initiation ↗

**Location:** RNA Polymerase III Transcription

**Stable identifier:** R-HSA-76046

**Compartments:** nucleoplasm



There are three basic types of RNA polymerase III promoters. The three types of RNA polymerase III promoters are known as type 1, type 2, and type 3 promoters. Type 1 promoters are found in the 5S genes and consist of a gene-internal element called the internal control region (ICR), that is subdivided into A block, intermediate element, and C block (Boghenagen, 1985; Sakonju et al., 1980). Type 2 promoters are found in tRNA genes, Adenovirus 2 VAI gene, and other genes (Galli et al., 1981; Sharp et al., 1981). These promoters consists of two gene-internal elements called the A and the B boxes. Type 3 promoters consist of a distal sequence element (DSE) that serves as an enhancer, a proximal sequence element (PSE), and a TATA box (Baer et al., 1989; Lobo and Hernandez, 1989).

Some promoters combine elements from type 2 and 3 promoters. For example, the *S. cerevisiae* U6 promoter, also shown in the figure, contains the TATA box typical of type 3 promoters and the A and B boxes typical of type 2 promoters. Moreover, in *S. pombe*, nearly all tRNA and 5S genes contain a TATA box in addition to gene-internal elements, and the TATA box is required for transcription.

## Literature references

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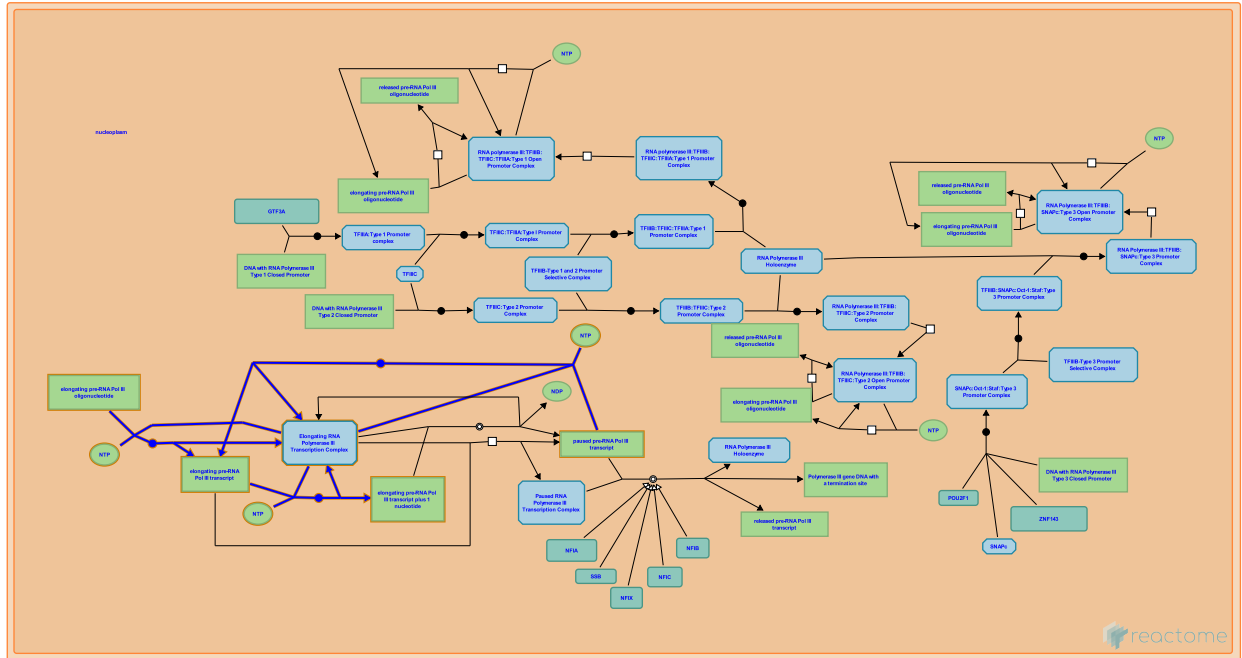
2003-09-11	Authored	Hernandez, N.
2024-03-06	Edited	Joshi-Tope, G.

# RNA Polymerase III Chain Elongation ↗

**Location:** RNA Polymerase III Transcription

**Stable identifier:** R-HSA-73780

**Compartments:** nucleoplasm



Pol III initiation complexes open the promoter spontaneously similar to the mechanism employed in archaeal and bacterial transcription.

## Editions

2004-02-27

Authored

Kassavetis, GA., Geiduschek, EP.

2024-03-06

Edited

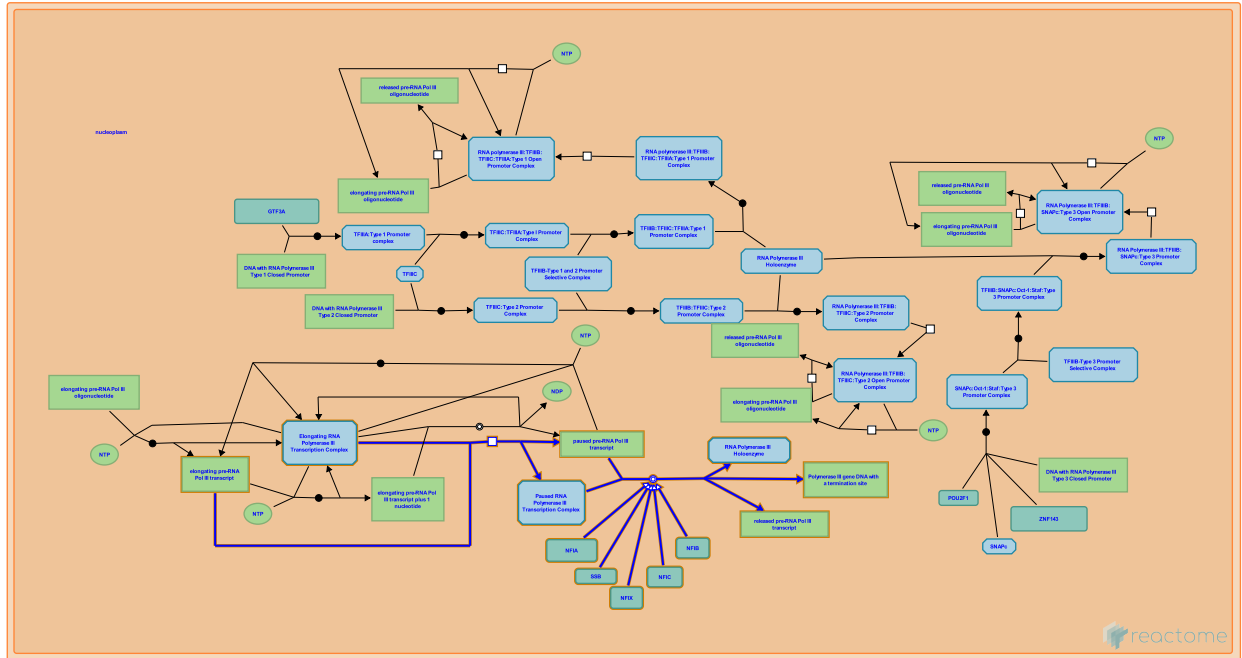
Gillespie, ME.

# RNA Polymerase III Transcription Termination ↗

**Location:** RNA Polymerase III Transcription

**Stable identifier:** R-HSA-73980

**Compartments:** nucleoplasm



At the end of the cycle, the elongation complex (EC) must be destabilized to release its transcript and DNA. Analogous to initiation, cis-signals and protein factors are required to mediate EC destabilization and release of the transcript from the grip of the RNA polymerase (RNAP). RNAP III achieves efficient termination despite the apparent simplicity of its cis-acting DNA terminator element, a stretch of five or more T residues on the non-template (NT) strand, which directs termination within this site without need for additional cis-elements or trans-acting factors.

## Literature references

Maraia, RJ., Arimbasseri, AG. (2015). Mechanism of Transcription Termination by RNA Polymerase III Utilizes a Non-template Strand Sequence-Specific Signal Element. *Mol. Cell*, 58, 1124-32. ↗

## Editions

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