

Download GenBank from NCBI

- Download NG or NC Accession
- Download NT Accession
- Save GenBank

Gene ▾ EGFR

Search

NCBI Home

Resource List (A-Z)

All Resources

Chemicals & Bioassays

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Domains & Structures

Genes & Expression

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Homology

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Sequence Analysis

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Variation

Welcome to NCBI

The National Center for Biotechnology Information advances science and health by providing access to biomedical and genomic information.

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Get Started

- [Tools](#): Analyze data using NCBI software
- [Downloads](#): Get NCBI data or software
- [How-To's](#): Learn how to accomplish specific tasks at NCBI
- [Submissions](#): Submit data to GenBank or other NCBI databases

Genomic Structural Variation

dbVar archives large scale genomic variation data and associates defined variants with phenotypic information.



|| 1 2 3 4 5 6 7 8

Popular Resources

PubMed

Bookshelf

PubMed Central

PubMed Health

BLAST

Nucleotide

Genome

SNP

Gene

Protein

PubChem

NCBI Announcements

NCBI's April Newsletter is on the Bookshelf

04 May 2012

Information about May's Discovery Workshop, the new GTR and Assembly

New Filter Sidebar has been added to PubMed

Gene

Gene

EGFR

Search

Save search Advanced

Help

Show additional filters

Display Settings: Tabular, 20 per page, Sorted by Relevance

Send to:

Hide sidebar >>

Clear all

Gene sources

- Genomic
- Mitochondria
- Organelles
- Plasmids

Categories

- Alternatively spliced
- Annotated genes
- Non-coding
- Protein-coding
- Pseudogene

Sequence content

- CCDS
- Ensembl
- RefSeq
- RefSeqGene

Status

clear

Current only

Did you mean EGFR as a gene symbol?
Search Gene for [EGFR](#) as a symbol.

Results: 1 to 20 of 3600 << First < Prev Page 1 of 180 Next > Last >>

Filters activated: Current only. [Clear all](#) to show 3615 items.

Name/Gene ID	Description	Location	Aliases
<input checked="" type="checkbox"/> EGFR ID: 1956	epidermal growth factor receptor [<i>Homo sapiens</i> (human)]	Chromosome 7, NC_000007.14 (55019032..55207338)	ERBB, ERBB1, H
<input checked="" type="checkbox"/> Egfr ID: 13649	epidermal growth factor receptor [<i>Mus musculus</i> (house mouse)]	Chromosome 11, NC_000077.6 (16752203..16913907)	RP23-295E4.1, Erb1, Errp, Was
<input checked="" type="checkbox"/> Egfr ID: 37455	Epidermal growth factor receptor [<i>Drosophila melanogaster</i>]	Chromosome 2R, NT_033778.4 (21522420..21559977)	Dmel_CG10079, DEGFR, DER, D ball, DER/top, D Der, DmHD-33, [

Filters: [Manage Filters](#)

Top Organisms [\[Tree\]](#)

- Homo sapiens (1140)
- Mus musculus (231)
- Drosophila melanogaster (132)
- Rattus norvegicus (108)
- Listeria monocytogenes (39)
- All other taxa (1980)

More...

Find related data

Database:

Select

Find items

Search details

EGFR[All Fields] AND alive[property]

Scroll down to Genomic regions, select the appropriate assembly

Genomic context

Location: 7p12
Exon count: 30

Download NT Accession → [MapViewer](#)

Annotation release	Status	Assembly	Chr	Location
106	current	GRCh38 (GCF_000001405.26)	7	NC_000007.14 (55019032..55207338)
105	previous assembly	GRCh37.p13 (GCF_000001405.25)	7	NC_000007.13 (55086678..55279262)

Chromosome 7 - NC_000007.14

Genomic regions, transcripts, and products

Download NG or NC Accession to [reference sequence details](#)

Gen: [NC_000007.14 chromosome 7 reference GRCh38 Primary Assembly](#)

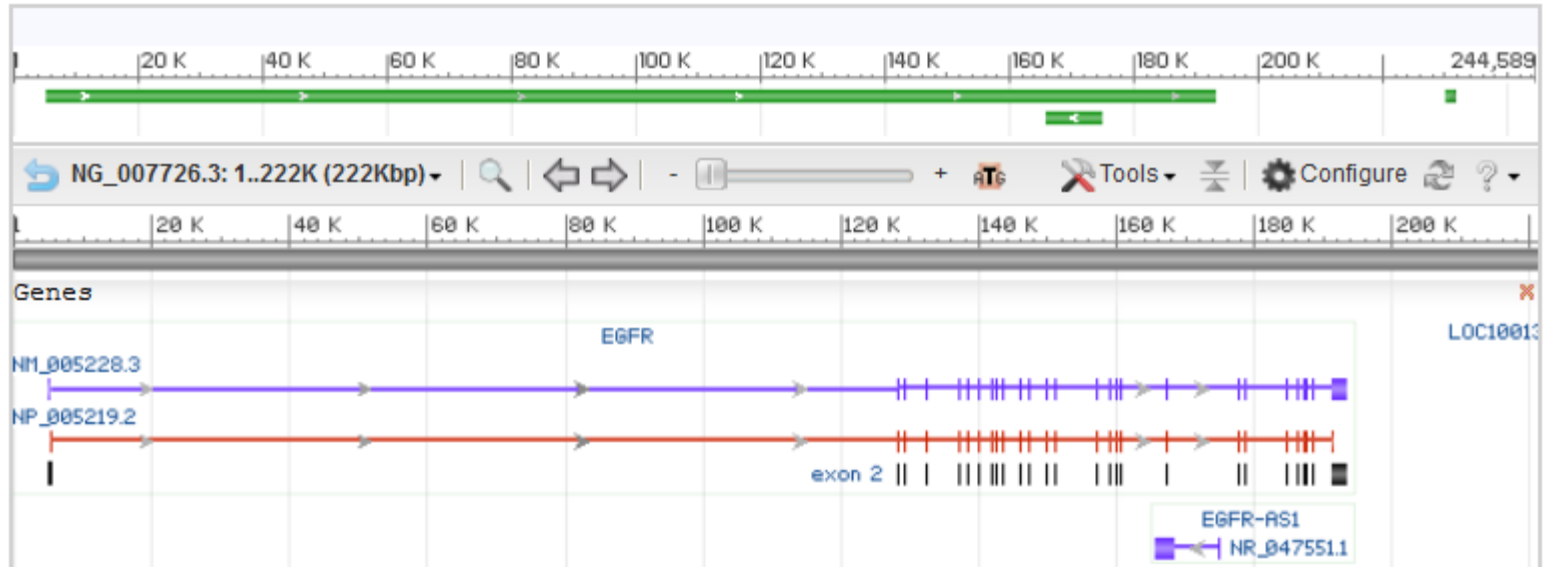
- NC_000007.14 chromosome 7 reference GRCh38 Primary Assembly
- NG_007726.3 RefSeqGene
- NC_018918.2 chromosome 7 alternate CHM1_1.1
- AC_000139.1 chromosome 7 alternate HuRef
- NC_000007.13 Chromosome 7 Reference GRCh37.p13 Primary Assembly

- NG accession is the RefSeq
- Most RefSeq GenBanks contain only a single transcript
- NG accession can show dbSNP
- Nucleotide positions do not relate to chromosome coordinates
- Click “GenBank” link to download NG Accession

Genomic regions, transcripts, and variants

Genomic Sequence:

Go to nucleotide: [Graphics](#) [FASTA](#) [GenBank](#)



- NC Accessions often contains Several Transcripts
- NC Accessions do not have dbSNP
- Nucleotide positions are chromosome coordinates
- Click “GenBank” link to download NC Accession

Genomic regions, transcripts, and products

Go to [reference sequence details](#)

Genomic Sequence: NC_000007.13 Chromosome 7 Reference GRCh37.p13 Primary Assembly ▾

Go to nucleotide: [Graphics](#) [FASTA](#) [GenBank](#)



[LOC643168](#) †

[sv](#) [dlevmm](#)

best RefSeq 7p11.2 Rho GTPase activating protein 5 pseudogene

- NT Accessions often contains Several Transcripts
- NT Accessions can show dbSNP
- Nucleotide positions can be translated to chromosome coordinates
- Click “dl” link to download NT Accession

[EGFR](#) †

[OMIM](#) [HGNC](#) [sv](#) [gr](#) [dlevmm](#) [hm](#) [sts](#) [CCDS](#) [SNP](#) best RefSeq 7p12 OTTHUMP00000209210|avian erythroblastic

[LOC100507500](#) †

[sv](#) [dlevmm](#) [sts](#)

mRNA

hypothetical LOC100507500

[Homo sapiens \(human\)](#) (Build 37.3)

Region to retrieve (in chromosome coordinates):

Chromosome: Strand:

from: adjust by:

to: adjust by:

Sequence Format:

This chromosome region corresponds to the contig region(s):

Contig	start	stop	strand	
NT_033968.6	4676094	4864400		+ Display Save to Disk View Evidence ModelMaker

Save GenBank

NCBI Resources How To Sign in to NCBI

Nucleotide Nucleotide Search Help

Advanced

Display Settings: GenBank Send:

Homo sapiens chromosome 7 genomic scaffold, GRCh38 Primary Assembly HSCHR7_CTG1

NCBI Reference Sequence: NT_007819.18

[FASTA](#) [Graphics](#)

Go to:

LOCUS NT_007819 188307 bp DNA linear CON 03-FEB-2014

DEFINITION Homo sapiens chromosome 7 genomic scaffold, GRCh38 Primary Assembly HSCHR7_CTG1.

ACCESSION [NT_007819](#) REGION: 55009032..55197338 GPS_003205514

VERSION NT_007819.18 GI:568815311

DBLINK BioProject: [PRJNA168](#)
Assembly: [GCF_000001405.26](#)

KEYWORDS RefSeq.

SOURCE Homo sapiens (human)

ORGANISM [Homo sapiens](#)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 188307)

CONSRM International Human Genome Consortium (International Human Genome Consortium)

Change region shown

Whole sequence (abbreviated view)

Selected region

from: 55009032 to: 55197338

Update View

Customize view

Abbreviated view

Customize

Basic Features

Default features

Gene, RNA, and CDS features only

Features added by NCBI

2522401 SNPs

Display options

Show sequence

Show reverse complement

Update View

With NG_ accessions, you have the option to include SNPs - select this and press Update View.

For genes located on minus strand, deselect Reverse complemented strand and press Update View.

Expand the range if you are interested in the gene's promoter region or other information outside of the gene, then press Update View.

Nucleotide Nucleotide

Advanced

Display Settings: GenBank

Homo sapiens chromosome 7 genomic scaffold, GR Assembly HSCHR7_CTG1

NCBI Reference Sequence: NT_007819

[FASTA](#) [Graphics](#)

Go to:

LOCUS NT_007819
 DEFINITION Homo sapiens chromosome 7 genomic scaffold, GR Assembly HSCHR7_CTG1.
 ACCESSION [NT_007819](#) REGION: 55009032..55197338 GPS_003205514
 VERSION NT_007819.18 GI:568815311
 DBLINK BioProject: [PRJNA168](#)
 Assembly: [GCF_000001405.26](#)
 KEYWORDS RefSeq.
 SOURCE Homo sapiens (human)

Choose Send
 Choose "File" Destination
 Choose "GenBank" Format
 Click "Create File"

Send:

- Complete Record
- Coding Sequences
- Gene Features

Choose Destination

- File
- Clipboard
- Collections
- Analysis Tool

Download 1 items.

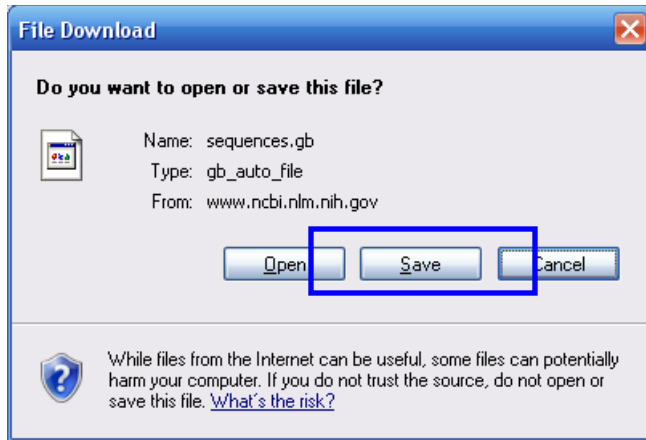
Format

GenBank

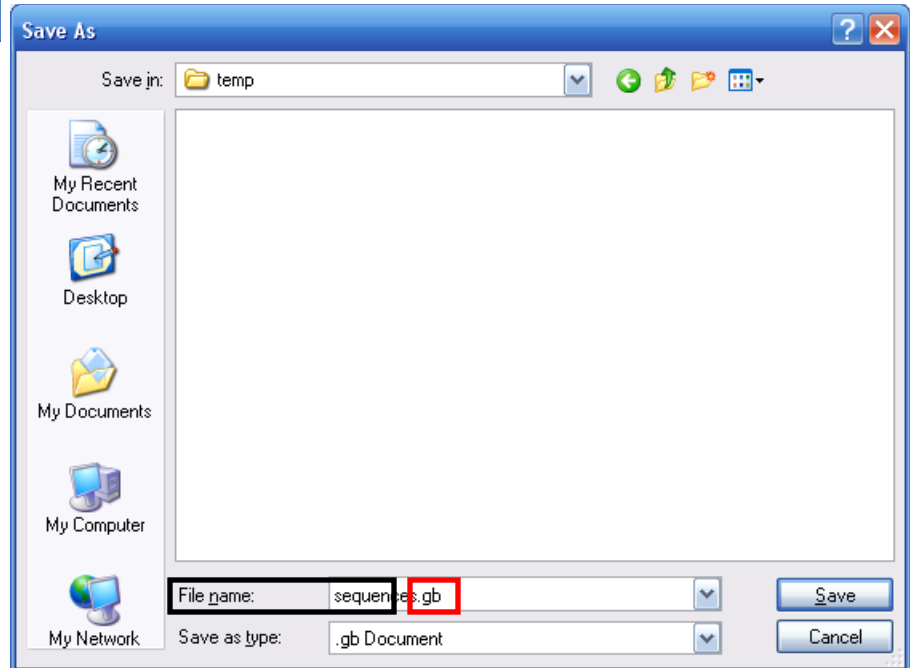
Create File

Features added

2522401 SNPs



Choose to Save File.



Gbk File Editor - C:\Documents and Settings\HP_Administrator\Desktop\EGFR.txt

File Edit Search Tools Help

New GBKFile

- Gene
 - EGFR
 - CDS
 - NP_005219.2**
 - NP_958441.1
 - NP_958439.1
 - NP_958440.1
 - mRNA
 - NM_005228.3
 - NM_201284.1
 - NM_201282.1
 - NM_201283.1
 - Variations

Basic Information | Sequence

Item	Value
Region	247..334,123255..123406,124274..124457,127575..127709,132263..132331,133515..133633,134980..13512
InterestRegion	
CodonStart	1
product	epidermal growth factor receptor isoform a
protein_id	NP_005219.2
note	isoform a is encoded by transcript variant 1;epidermal growth factor receptor (avian erythroblasticleukemia viral
translation	MRPSGTAGAALLALLAALCPASRALEEKVKCGT SNKLTQLGTFEDHFLSLQRMFNNCEWVLGNLEITYVQRNY
Corresponding_mRNA_Region	NM_005228.3
External primer	
Internal primer	

Choose the appropriate protein for your analysis

Advanced GBK File Editor - C:\Documents and Settings\HP_Administrator\My Documents\Downloads\EGFR_NC.txt

File Edit Search Tools

New GBKFile

- Gene
 - EGFR
 - CDS
 - NP_005219.2
 - NP_958441.1
 - NP_958439.1
 - NP_958440.1
 - mRNA
 - NM_005228.3
 - NM_201284.1
 - NM_201282.1
 - NM_201283.1
 - LOC100507500
 - Variations

Basic Information

Item	
CDS	80..135121,136799..136915,137502..137628,137728..137812)
Region of Interest	
Amplicon ID	
External primer	
Internal primer	
CodonStart	
product	
protein_id	
note	derived by automated computational analysis using gene prediction r
translation	MRPSGTAGAALLALLAALCPASRALEEKKVCQGTSNKLTQLGTFEDHFLSLQRMFNNCEVVLGNLEITYVQRNYDLSFLKT
Corresponding_mRNA_Region	NM_201283.1
transl_table	

Save As

Mode

- Selected gene region
- Selected gene region, Selected mRNA region
- Seq files for selected CDS

OK Cancel

- Choose Save As from File menu
- Choose Selected gene region, selected mRNA region
- Then open this GBK file for your project.

Advanced GBK File Editor - C:\Documents and Settings\HP_Administrator\My Documents\Downloads\EGFR_NC.txt

File Edit Search Tools

New GBKFile

- Gene
 - EGFR
 - CDS
 - NP_958440.1
 - mRNA
 - NM_201283.1
 - Variations

Basic Information | Sequence

Item	
LOCUS	
SEQUENCE TYPE	
DEFINITION	
ACCESSION	
VERSION	
KEYWORDS	