

## NCBI News, November 2011

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### Phase One Rollout of the New Genome Site

A completely redesigned Genome site, [www.ncbi.nlm.nih.gov/genome](http://www.ncbi.nlm.nih.gov/genome), is now available. Major improvements include a more natural organization at the level of the organism for prokaryotic, eukaryotic, and viral genomes. Reports include information about the availability of nuclear or prokaryotic primary genomes as well as organelles and plasmids. The new Genome resource provides a summary view of the data from all genome-scale projects including genome maps, assemblies, annotation, and transcriptomes. Genome collects data from primary data resources and provides links to more detailed information. While not new with this release, it is worth noting that the Genome interface has been upgraded to the new NCBI standard with the new search bar, Limits and Advanced Search pages, and NCBI footer. Moreover, search results and record views in Genome are discovery-oriented and feature the Discovery Column with analysis tools and easy access to related data. Figure 1 shows sample pages from the Genome resource and highlights these new features. An [information page](#) accompanying the release provides additional details and help with transitioning to the new service.

The new Genome site is much easier to navigate and provides rapid access to all genome data for a particular organism. The new site will continue to improve as additional displays and features are added in phases. A feature article in the next NCBI News will provide more detailed coverage of Genome with examples illustrating the power of the new system.

### Note: Changes affecting genome identifiers

Because of the reorganization to a natural classification system, older genome identifiers are no longer valid. Typically these genome identifiers were not exposed in the previous system and were used mainly for programmatic access. To aid in the transition to the new system, a [file](#) that maps previous Genome identifiers to the identifier for the genome sequence is available on the FTP site. The sequence identifiers can be used to retrieve the genome sequence from the Nucleotide database.

### New BLAST videos on NCBI's YouTube channel

Two new instructional videos about BLAST statistics are on the NCBI YouTube channel: [An explanation of BLAST E-values \(pt.1\)](#), and [Answers to a few E-value FAQs \(pt.2\)](#). These BLAST videos should help in interpreting BLAST output and designing more effective BLAST search strategies. The BLAST videos join a growing collection of 54 videos on the [NCBI YouTube channel](#).

The figure displays three panels from the NCBI Genome browser. The top-left panel shows the species-level page for *Homo sapiens*, including genome statistics, assembly information, and a list of related bio-projects. The top-right panel shows the species-level page for *Staphylococcus aureus*, featuring a sub-species tree and a table of genome projects. The bottom-left panel shows search results for the query "plants", listing *Zea mays*, *Oryza sativa*, and *Physcomitrella patens*. The bottom-right panel shows the strain-level page for *Staphylococcus aureus* subsp. *aureus* MRSA252, including a graphical sequence viewer of the chromosome.

**Top panel, right: Species-level page for human (*Homo sapiens*)**

Lineage: Eukaryota[3473]; Metazoa[2593]; Chordata[1694]; Craniata[1672]; Vertebrata[1670]; Euteleostomi[1649]; Mammalia[343]; Eutheria[307]; Euarchontoglires[168]; Primates[56]; Haplorhini[41]; Catarrhini[36]; Hominoidea[7]; Homo[2]

The human genome project is the result of an international consortium among many different sequencing and bioinformatics centers. A wealth of data is available including: the annotated assembled genomic sequence, transcript sequence, library resources, expression data, map data, disease and functional information, and more. The result is an unprecedented amount of knowledge concerning human genetics that will eventually result in breakthroughs in understanding human biology as well as significant medical advances. A challenge facing researchers today is that of analyzing and integrating the plethora of data available. The human genome sequence provides a critical foundation for continued advances in medicine, basic research, and clinical diagnostic technologies.

**Chromosomes**

Click on chromosome name to open Map Viewer

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18  
19 20 21 22 X Y

**Assembly and Annotation**

8 other assemblies are available

Default assembly	
Assembly Name	GRCh37.p5
Last sequence update	06-Mar-2009
Highest level of assembly	some chromosomes assembled
Size (total bases)	3,101,788,170
Number of genes	36,036
Number of proteins	32,130

**Mitochondrial Genome**

Last record update	30-Apr-2010
Last sequence update	08-Jul-2009
Size	16,569
Number of genes	37
Number of proteins	13

**Related BioProjects**

Type	Count
RefSeq Genome	4
Assembly	1
Clone ends	3
Epigenomics	1
Exome	31
Genome sequencing	65
Map	3
Other	2
Phenotype or Genotype	3
Random survey	1
Targeted Locus (Loc)	3
Transcriptome or Gene expression	20
Variation	4

**Top panel, right: Species-level page for *Staphylococcus aureus***

Lineage: Bacteria[2656]; Firmicutes[630]; Bacillales[144]; Staphylococcus[22]

**Staphylococci.** The genus *Staphylococcus* are pathogens of humans and other mammals. Traditionally they were divided into two groups based on the coagulase reaction. *Staphylococci* are generally found inhabiting the skin and mucous membranes of mammals and birds. Some members of this genus can be found as human commensals and these are generally believed to have the greatest pathogenic potential in opportunistic infections. *Staphylococcus aureus*. This organism is a major cause of nosocomial (hospital-acquired) and community-acquired infections. Since its discovery as an opportunistic pathogen, *S. aureus* continues to be a major cause of mortality and is responsible for a variety of infections including, boils, furuncles, styes, impetigo and other superficial skin infections in humans. Also known to cause more serious infections particularly in the chronically ill or immunocompromised. These include pneumonia, deep abscesses, osteomyelitis, endocarditis, phlebitis, mastitis and meningitis. The ability to cause invasive disease is associated with persistence in the nasal cavity of a host.

**Sub-species tree**

1%

**Genome Projects**

Highest level of Assembly	Count
Chromosomes	26
Scaffolds or contigs	68
SRAs or Traces	98
No data	27
All	219

**Related BioProjects**

Type	Count
RefSeq Genome	89
Genome sequencing	222

**Bottom panel, right: Strain-level page for *Staphylococcus aureus* subsp. *aureus* MRSA252**

Lineage: Bacteria[2656]; Firmicutes[630]; Bacillales[144]; Staphylococcus[22]

**Staphylococcus aureus subsp. aureus strain 252.** This strain is a hospital-acquired strain isolated in the United Kingdom, representative of the methicillin-resistant (MRSA) strains. It is one of the two major MRSA strains found in British hospitals in the late 1990's.

**Sequence Information (See Genome Summary)**

Type	Name	RefSeq	INSDC	Size (Mb)	GC%	proteins	rRNA	tRNA	other rna	gene	pseudogene
Chr	-	NC_002952.2	BX371856.1	2.9	32.8	2,650	16	60	25	2,839	88

**Bottom panel, left: Sample search results for the query "plants"**

Results: 1 to 20 of 357

- Zea mays***  
Major crop plant and a classic genetic model  
Kingdom: Eukaryotes  
Haplod chromosomes: 10; Organelles: 2  
ID: 12
- Oryza sativa***  
Cultivated rice, used as staple food by majority of world's population  
Kingdom: Eukaryotes  
Haplod chromosomes: 12; Organelles: 2  
ID: 10
- Physcomitrella patens***  
Moss, used as a model plant for functional genomics, evo-devo studies and comparative genomics.  
Kingdom: Eukaryotes  
Haplod chromosomes: 27; Organelles: 2  
ID: 383

Figure 1. Sample Genome pages. *Top panel, right*: Species-level page for human showing the right hand Discovery Column. *Top panel, left*: Species-level page for the bacterium *Staphylococcus aureus*. The subspecies tree has links to strain-level pages. *Bottom panel, right*: Strain-level page for the antibiotic resistant *S. aureus* MRSA252 showing the imbedded graphical sequence viewer display of the chromosome. *Bottom panel, left*: Sample search results for the query "plants" showing the updated search result page.

## BLAST Results: Expect Values, part 1

NCBI 54 videos  [www.youtube.com/ncbinlm](http://www.youtube.com/ncbinlm)

Alignments

Select All [Get selected sequences](#) [Distance tree of results](#) [Multiple alignment](#)

```
> ref|NP\_001008976.1 | UGM apolipoprotein A-I [Pan troglodytes]
Length=100
GENE ID: 449498 APOA2 | apolipoprotein A-I [Pan troglodytes]
(10 or fewer PubMed links)
Score = 177 bits (448), Expect = 2e-61, Method: Compositional matrix adjust.
Identities = 97/100 (97%), Positives = 100/100 (100%), Gaps = 0/100 (0%)
Query 1  MKLLAATVLLLITCSLEGALVRRQAKEPCVESLVSQYFQTVTDYGKDLMEKVKSPQLQAE  60
          MKLLAATVLLLITCSLEGALVRRQAKEPCV++LVSQYFQTVTDYGKDLMEKVKSPQLQAE
Sbjct 1  MKLLAATVLLLITCSLEGALVRRQAKEPCVDNLVSQYFQTVTDYGKDLMEKVKSPQLQAE  60
Query 61  AKSYFEKSKEQLTPLIKKAGTELVNFLSYFVELGTQPATQ  100
          AKSYFEKSKEQLTPLIKKAGTELVNFLSYFVELGTQPATQ
```

Homology?

## BLAST Results: Expect Values, part 2

NCBI 54 videos

**Question #3:**

What is an E-value of 0.0?

E-value < 1e-179

2:55 / 3:08 360p

## Entrez Utility Changes: New EFetch Version and New alternative ESummary XML

An updated EFetch Entrez Utility (version 2.0) is now in production. EFetch retrieves records from the NCBI databases by unique identifier. Additions include support for the BioSample, BioProjects, and SRA databases as



well as defined default values for retrieval mode (retmode) and retrieval type (rettype). Updated retmode and rettype values are given in the [table](#) in the [Entrez Programming Utilities Help Manual](#).

An alternative XML record is now available from the ESummary Entrez Utility. The content in the new record is unique to each Entrez database and has additional content not available in the traditional ESummary record. The new XML can be requested by including &version=2.0 in the ESummary URL. The traditional ESummary record will continue to be supported and will be returned without the version parameter. The [traditional](#) and [version 2.0](#) ESummary for NM\_000240 (gi=33469954) from the nuccore (nucleotide) database are shown below. The [release notes](#) provide details on changes. The [Entrez Programming Utilities Help Manual](#) has complete information on EFetch, ESummary, and the other EUtility programs.

```

<eSummaryResult>
  --<DocSum>
    <Id>33469954</Id>
    <Item Name="Caption" Type="String">NM_000240</Item>
    --<Item Name="Title" Type="String">
      Homo sapiens monoamine oxidase A (MAOA), nuclear gene encoding mitochondrial protein, mRNA
    </Item>
    <Item Name="Extra" Type="String">gi33469954[refNM_000240.2][33469954]</Item>
    <Item Name="Gi" Type="Integer">33469954</Item>
    <Item Name="CreateDate" Type="String">1999/04/01</Item>
    <Item Name="UpdateDate" Type="String">2011/10/23</Item>
    <Item Name="Flags" Type="Integer">512</Item>
    <Item Name="TaxId" Type="Integer">9606</Item>
    <Item Name="Length" Type="Integer">4090</Item>
    <Item Name="Status" Type="String">live</Item>
    <Item Name="ReplacedBy" Type="String"/>
    <Item Name="Comment" Type="String"></Item>
  </DocSum>
</eSummaryResult>

```

Traditional XML

ESummary XML for Nucleotide NM\_000240

```

<eSummaryResult>
  --<DocumentSummarySet status="OK">
    --<DocumentSummary uid="33469954">
      <Caption>NM_000240</Caption>
      --<Title>
        Homo sapiens monoamine oxidase A (MAOA), nuclear gene encoding mitochondrial protein, mRNA
      </Title>
      <Extra>gi33469954[refNM_000240.2]</Extra>
      <Gi>33469954</Gi>
      <CreateDate>1999/04/01</CreateDate>
      <UpdateDate>2011/10/23</UpdateDate>
      <Flags>512</Flags>
      <TaxId>9606</TaxId>
      <Slen>4090</Slen>
      <Biomol>mRNA</Biomol>
      <MolType>rna</MolType>
      <Topology>linear</Topology>
      <SourceDb>refseq</SourceDb>
      <SegSetSize>0</SegSetSize>
      <ProjectId>0</ProjectId>
      <Genome>genomic</Genome>
      <SubType>chromosomeimap</SubType>
      <SubName>XIXp11.3</SubName>
      <AssemblyG>14165523</AssemblyG>
      <AssemblyAcc>X60819.1</AssemblyAcc>
      <Tech>
      <Completeness>has-right</Completeness>
      <GeneticCode>1</GeneticCode>
      <Strand>
      <Organism>Homo sapiens</Organism>
      +<Statistics></Statistics>
      <AccessionVersion>NM_000240.2</AccessionVersion>
      <Properties na="1">1</Properties>
      <Comment>
      <OSLT indexed="yes">NM_000240.2</OSLT>
      <IdGiClass mol="2" repr="2" gi_state="10" sat="4" sat_key="59045920" owner="20" sat_name="NCBI" o
      </DocumentSummary>
    </DocumentSummarySet>
  </eSummaryResult>

```

Version 2.0 XML

```

--<Statistics>
  <Stat type="Length" count="4090"/>
  <Stat type="Length" subtype="literal" count="4090"/>
  <Stat type="all" count="6"/>
  <Stat type="blob_size" count="16469"/>
  <Stat type="cdregion" count="1"/>
  <Stat type="cdregion" subtype="CDS" count="1"/>
  <Stat type="gene" count="1"/>
  <Stat type="gene" subtype="Gene" count="1"/>
  <Stat type="imp" count="3"/>
  <Stat type="imp" subtype="polyA_signal" count="1"/>
  <Stat type="imp" subtype="polyA_site" count="2"/>
  <Stat type="org" count="1"/>
  <Stat type="pub" count="10"/>
  <Stat type="pub" subtype="PubMed" count="5"/>
  <Stat type="pub" subtype="PubMedGene-rif" count="5"/>
  <Stat source="CDS" type="all" count="13"/>
  <Stat source="CDS" type="prot" count="1"/>
  <Stat source="CDS" type="region" count="2"/>
  <Stat source="CDS" type="region" subtype="Region" count="2"/>
  <Stat source="CDS" type="site" count="10"/>
  <Stat source="CDS" type="site" subtype="Site" count="10"/>
  <Stat source="CDS/CDD" type="all" count="6"/>
  <Stat source="CDS/CDD" type="region" count="6"/>
  <Stat source="CDS/CDD" type="region" subtype="Region" count="6"/>
  <Stat source="CDS/SNP" type="all" count="30"/>
  <Stat source="CDS/SNP" type="imp" count="30"/>
  <Stat source="CDS/SNP" type="imp" subtype="variation" count="30"/>
  <Stat source="Exon" type="all" count="15"/>
  <Stat source="Exon" type="evidence" count="15"/>
  <Stat source="Exon" type="imp" count="15"/>
  <Stat source="Exon" type="imp" subtype="exon" count="15"/>
  <Stat source="SNP" type="all" count="42"/>
  <Stat source="SNP" type="imp" count="42"/>
  <Stat source="SNP" type="imp" subtype="variation" count="42"/>
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  <Stat source="all" type="Length" count="4090"/>
  <Stat source="all" type="all" count="121"/>
  <Stat source="all" type="blob_size" count="16469"/>
  <Stat source="all" type="cdregion" count="1"/>
  <Stat source="all" type="evidence" count="15"/>
  <Stat source="all" type="gene" count="1"/>
  <Stat source="all" type="imp" count="99"/>
  <Stat source="all" type="org" count="1"/>
  <Stat source="all" type="prox" count="1"/>
  <Stat source="all" type="pub" count="10"/>
  <Stat source="all" type="region" count="8"/>
  <Stat source="all" type="site" count="10"/>
</Statistics>

```

## Highlight Features Link Now on Sequence Records

A Highlight Features link now appears in the Analyze this Sequence section of protein and nucleotide sequence records displayed at the NCBI site. This link activates the new Feature Highlight function described in the

August 2011 NCBI News. Clicking the link opens the Feature Highlight Bar and highlights the first coding sequence (CDS) feature or the first linked feature if no CDS feature is present.

As pointed out in the original NCBI News article, the Highlight Features function is helpful in visualizing the extent and location of such important features as genes, coding regions, exons, and mRNAs in nucleotide sequences and conserved domains, modification sites, and interaction sites in protein sequences. This function joins the links to BLAST, Primer-BLAST, and Conserved Domain searches as well as the Find-in-Sequence pattern finder as on-the-fly analysis capabilities in the NCBI sequence databases. The image below shows the link and the activated CDS highlight on the ReSeqGene record for the human monoamine oxidase gene (NG\_008957).

**Homo sapiens monoamine oxidase A (MAOA), RefSeqGene on chromosome X**

NCBI Reference Sequence: NG\_008957.1

FASTA Graphics

Go to: [dropdown]

LOCUS NG\_008957 97660 bp  
 DEFINITION Homo sapiens monoamine oxidase X.  
 ACCESSION NG\_008957  
 VERSION NG\_008957.1 GI:212549708  
 KEYWORDS RefSeqGene.  
 SOURCE Homo sapiens (human)  
 ORGANISM [Homo sapiens](#)  
 Eukaryota; Metazoa; Chordata; C  
 Mammalia; Eutheria; Euarchontog  
 Catarrhini; Hominidae; Homo.  
 COMMENT REVIEWED [REFSEQ](#): This record has  
 reference sequence was derived  
[BX530072.B](#) and [BX537148.1](#).  
 This sequence is a reference standard in the [RefSeqGene](#) project.

Change region shown [dropdown]  
 Customize view [dropdown]

Analyze this sequence [dropdown]  
 Run BLAST  
 Pick Primers  
**Highlight Sequence Features**  
 Find in this Sequence

Articles about the MAOA gene [dropdown]  
<sup>3</sup>H kinetic isotope effects and pH dependence of catalysis as mechanistic probe [Biochemistry. 2011]  
 Association of the MAOA promoter uVNTR polymorphism with suicide [BMC Med Genet. 2011]  
 Search for association between suicide and 5-HTT, MAOA and DAT [Arch Med Sadowe] Kryminol. 2010]

92581 accctgccca ccttcccaag taactctgtg taacctcttg gttcccttga aggggtgatcc  
 92641 gtaaacccgt gggcaggatt ttctttgggg gcacagagac tgccacaaag tggagcggct  
 92701 acatggaag ggcagttgag gctggagaac gacagactag ggaggaagc aggaaagccc  
 92761 aggctctctc cctcccgagt cacggcaacg tttttggcat ctggctctgc tagttcttga  
 92821 cactgataga atctgtatgt ccatttctct gccctcact catggggctc  
 92881 agcagggcct tgaatctgta gaaactatac agcctctttt cataataacc  
 92941 cttcaggtc taaatggtc tcgggaaggt gaccgagaaa gatatctgg  
 93001 tgaatcaang gtaagtttgg tgactctggg cactatctct ccttagacc  
 93061 ataaactca catctccctt ctctagcct cggatttaat tatagatgc  
 93121 gggctcccat gcattgatct tgcagtgtt ttgtctctct tgtcagcat  
 93181 tcatgatctg ttttccctca tctaggacgt tcacgggta gaaatcacc  
 93241 ggaagggaac ctgccctctg tttctggcct gctgaagatc attggattc  
 93301 aactgccctg gggtttctgc tgtacaaata caagctcctg ccacggctc  
 93361 tcttatgctc tctgtcact ggttttcaat accaccaaga ggaaatat  
 93421 aaggctgtgt cattgggcca tgtttaagtg tactggattt aactacctt  
 93481 caatcattgt taaagtaaaa acaattcaaa gaatcaccta attaatttc  
 93541 gctccatctt atttgcagt gtagatcaac tcatgttaat tgatagaat  
 93601 atcaactttc gaaattcaca aagttaaacg tgatgtgctc atcagaaaac  
 93661 cctgttttta ttcccttcaa tgcaaaatac atgatgattt cagaaacaa  
 93721 ttctgtctgt ggaggtggag taggtgaagg ccagcctgt aactgtcct  
 93781 taggcaatgg tgaactgtca ttacagagcc tagaggctca cagcctcct  
 93841 cctccacttt ggatcaggaa atagtaaagg aaagcagtg tgggggtag  
 93901 ccctcagacc agaatgggga catcttctgt tctgtctcct caggaatct

join(5182..5254,32353..32447,42130..42267,60711..60815,  
 61544..61635,77012..77153,80080..80229,80533..80692,  
 81538..81634,85066..85119,89520..89577,90789..90886,  
 92633..92744,92948..93010,93206..93352)  
 /gene="MAOA"  
 /EC\_number=" 1.4.3.4 "  
 /note="MAO-A; monoamine oxidase type A"  
 /codon\_start=1  
 /product="amine oxidase [flavin-containing] A"  
 /protein\_id=" NP\_000231.1 "  
 /db\_xref="GI:4557735"  
 /db\_xref="CCDS: CCDS14260.1 "  
 /db\_xref="GeneID: 4128 "  
 /db\_xref="HGNC: 6833 "  
 /db\_xref="MIM: 309850 "

CDS Feature 1 of 1 NG\_008957 : 15 segments

Details [dropdown] Display: FASTA GenBank Help [X]

## New BLAST 16S Prokaryotic Ribosomal RNA Database

A prokaryotic 16S ribosomal RNA database is now available through the database pull-down list on the main nucleotide BLAST service. The 16S database contains both bacterial and archaeal sequences from two RefSeq Targeted Loci projects (BioProjects PRJNA33175 and PRJNA33317). These data represent near full-length 16S



ribosomal RNA sequences from more than 250 archaeal and 7200 bacterial strains. The 16S BLAST database is useful for identifying or establishing the taxonomic affinities of unknown bacterial 16S sequences such as those from environmental or organismal samples or metagenomes. Figure 2 shows how the database can be used to partially classify a 16S sequence (JF340503) obtained from a concrete sewer biofilm (PubMed: 21981064, PopSet: 330372088). The top panel of the figure shows the basic nucleotide BLAST form with the 16S database selected. The center panel shows the BLAST results (RID: CUR81JZY012). The results indicate that the query sequence has the closest affinity to the acetobacteriaceae, particularly *Acidocella facilis*. The [BLAST Distance Tree](#), also shown in the figure provides a useful way to see the results of the analysis at a glance.

The pre-formatted 16S microbial database is also available in the [BLAST db FTP directory](#) as the file [16SMicrobial.tar.gz](#).

## New Phenotype-Genotype Integrator (PheGenI)

The [Phenotype-Genotype Integrator \(PheGenI\)](#) is a new service that integrates genome-wide association study (GWAS) catalog data from NHGRI with molecular and literature databases at the NCBI. PheGenI takes chromosome location, gene, SNP, or phenotype as input and provides annotated tables of SNPs, genes, association results, and gene expression data. A new [tutorial video](#) on YouTube demonstrates how to use PheGenI.

## Eukaryotic Genome Builds and Updates

Twelve new genome assemblies with annotations have recently been released at the NCBI. Nine of the new builds are genomes that make their first appearance (build 1.1) at NCBI. Highlights include the first genome for a sponge (*Amphimedon queenslandica*), the first for a reptile – the green anole (*Anolis carolinensis*), the first for a perciform fish – the Nile tilapia (*Oreochromis niloticus*), and two new rodent genomes – the guinea pig (*Cavia porcellus*) and the Chinese hamster CHO-K1 cell line (*Cricetulus griseus*). In addition updated annotations for six more genomes are also available including human build 37.3 described in the next section. A complete list of new builds and updates is given below. The NCBI [BioProject](#), [Genome](#), [Gene](#), [Nucleotide](#), [Protein](#), [BLAST](#) and [Map Viewer](#) services provide access to these data. The assemblies and annotations may also be downloaded from the [genomes area](#) of the FTP site.

### First NCBI Builds (build 1.1)

Sponge (*Amphimedon queenslandica*) [[BioProject](#), [Map Viewer](#)]

Buff-tailed bumblebee (*Bombus terrestris*) [[BioProject](#), [Map Viewer](#)]

Nile tilapia (*Oreochromis niloticus*) [[BioProject](#), [Map Viewer](#)]

Domestic turkey (*Meleagris gallopavo*) [[BioProject](#), [Map Viewer](#)]

Green Anole (*Anolis carolinensis*) [[BioProject](#), [Map Viewer](#)]

Guinea pig (*Cavia porcellus*) [[BioProject](#), [Map Viewer](#)]

African savannah elephant (*Loxodonta africana*) [[BioProject](#), [Map Viewer](#)]

White-faced gibbon (*Nomascus leucogenys*) [[BioProject](#), [Map Viewer](#)]

Chinese hamster (CHO-K1 cell line) (*Cricetulus griseus*) [[BioProject](#), [Map Viewer](#)]

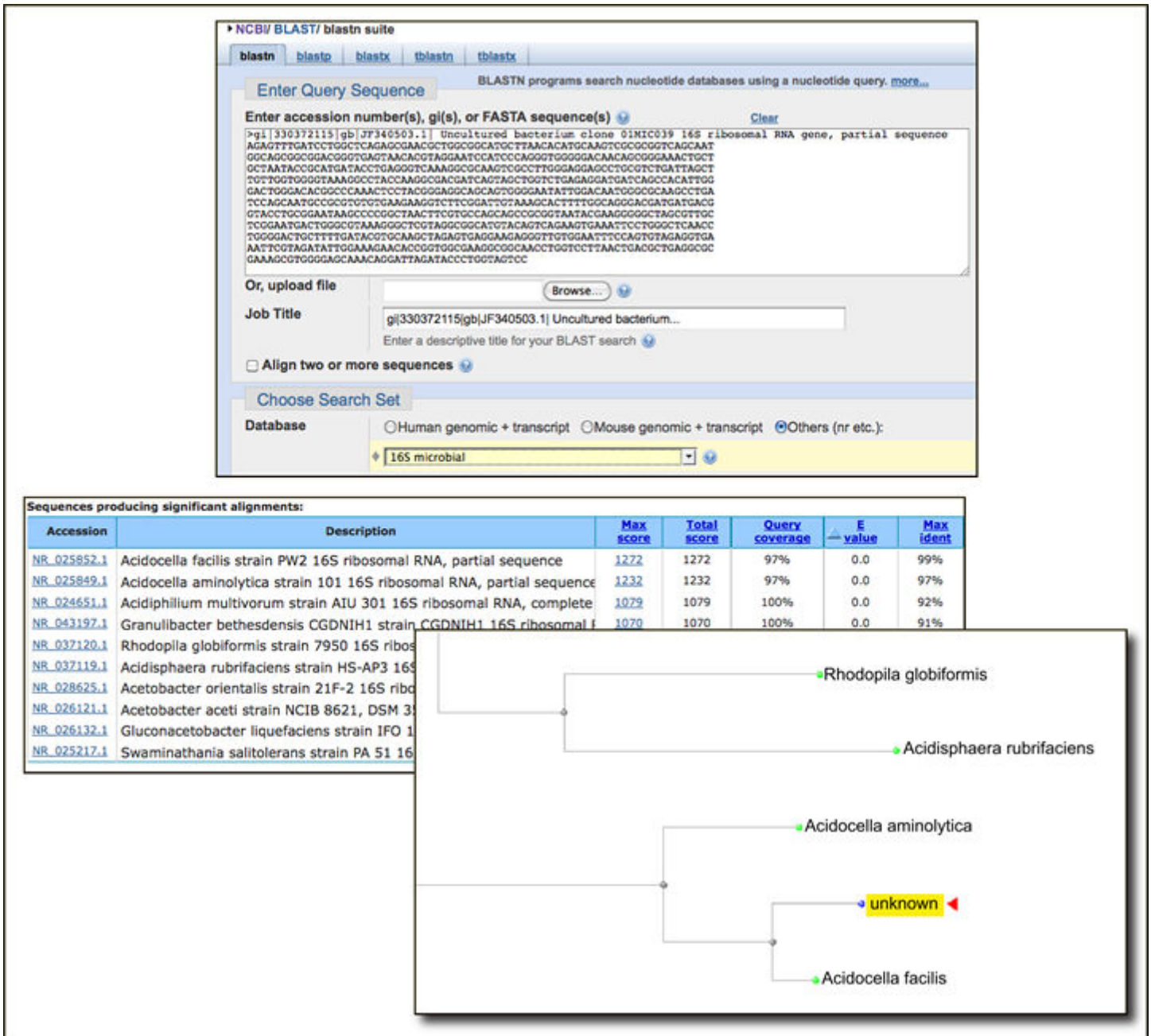


Figure 2. Using the NCBI nucleotide BLAST service with the new 16S microbial rRNA database. *Top panel.* The nucleotide BLAST search form with the 16S microbial database selected. The query sequence (JF340503) is a 16S sequence obtained from an environmental biofilm (PopSet: 330372088). *Center panel.* BLAST results (RID: CUR81/ZY012). The best match is to the 16S ribosomal RNA sequence (NR\_025852) from *Acidocella facilis* strain PW2. The linked BLAST Distance Tree of the results (bottom panel) shows the placement within *Acidocella* at a glance.

## New Builds

Honeybee (*Apis mellifera*), build 5.1 [BioProject, Map Viewer]

Pea aphid (*Acyrtosiphon pisum*), build 2.1 [BioProject, Map Viewer]

Zebrafish (*Danio rerio*), build 5.1 [BioProject, Map Viewer]

Chimpanzee (*Pan troglodytes*), build 3.1 [BioProject, Map Viewer]

## Updated Annotations

Fruit fly (*Drosophila melanogaster*), build 9.4 [[BioProject](#), [Map Viewer](#)]

Horse (*Equus caballus*), EquCab2.0 [[BioProject](#), [Map Viewer](#)]

Dog (*Canis lupus familiaris*), build 2.2 [[BioProject](#), [Map Viewer](#)]

Duck-billed platypus (*Ornithorhynchus anatinus*), build 1.2 [[BioProject](#), [Map Viewer](#)]

Thale cress (*Arabidopsis thaliana*), TAIR 10 [[BioProject](#), [Map Viewer](#)]

Human (*Homo sapiens*), build 37.3 [[BioProject](#), [Map Viewer](#)]

## Human Genome Update

The NCBI human genome annotation has been updated to version 37.3 and is now available in the [Map Viewer](#), the Entrez system and [human genome BLAST](#). The [build statistics](#) have more information on the contents of the release. The update includes the [Genome Reference Consortium](#) sequence patches from [patch 5](#). The patches are currently available as separate sequences from the chromosome assemblies. Patches that correct problems in the current assembly (fix patches) will be incorporated in the next complete genome assembly (build 38).

## Microbial Genomes Update

Fifty-eight finished microbial (archaeal and bacterial) genomes were released during September and October 2011. The original sequence data files submitted to the International Sequence Database Collaboration (INSDC) are available in the [Bacteria directory](#) in the genomes area of the GenBank FTP site. RefSeq provisional versions were released for a selected set of 32 of the complete INSDC microbial genomes during the same period. These are available from the [/genomes/Bacteria](#) directory on the FTP site.

In addition, data from 425 microbial whole genome-shotgun (WGS) sequencing projects were added to the INSDC during this period. The original submitted files are available in the [Bacteria\\_DRAFT](#) directory in the GenBank genomes area. RefSeq provisional versions of 89 WGS microbial projects were released in the [/genomes/Bacteria\\_DRAFT](#) area of the FTP site.

All GenBank and RefSeq microbial genomes are incorporated in the NCBI integrated Entrez search and retrieval system and the BLAST sequence similarity search service.

## GenBank News

GenBank release 186 is available through Entrez, BLAST and from the [GenBank FTP](#) area. The current release incorporates data available as of Oct 13, 2011 and, with the whole-genome shotgun portion, contains 350,733,781,429 bases from 212,788,863 sequence records. [Release notes](#) describe the current state of data and upcoming changes.

## RefSeq News

RefSeq Release 50 is available through Entrez, BLAST, and from the [RefSeq FTP area](#). The current release includes 18.8 million Reference Sequence records from 16,392 different species or strains. The RefSeq [release notes](#) provide more detailed information.

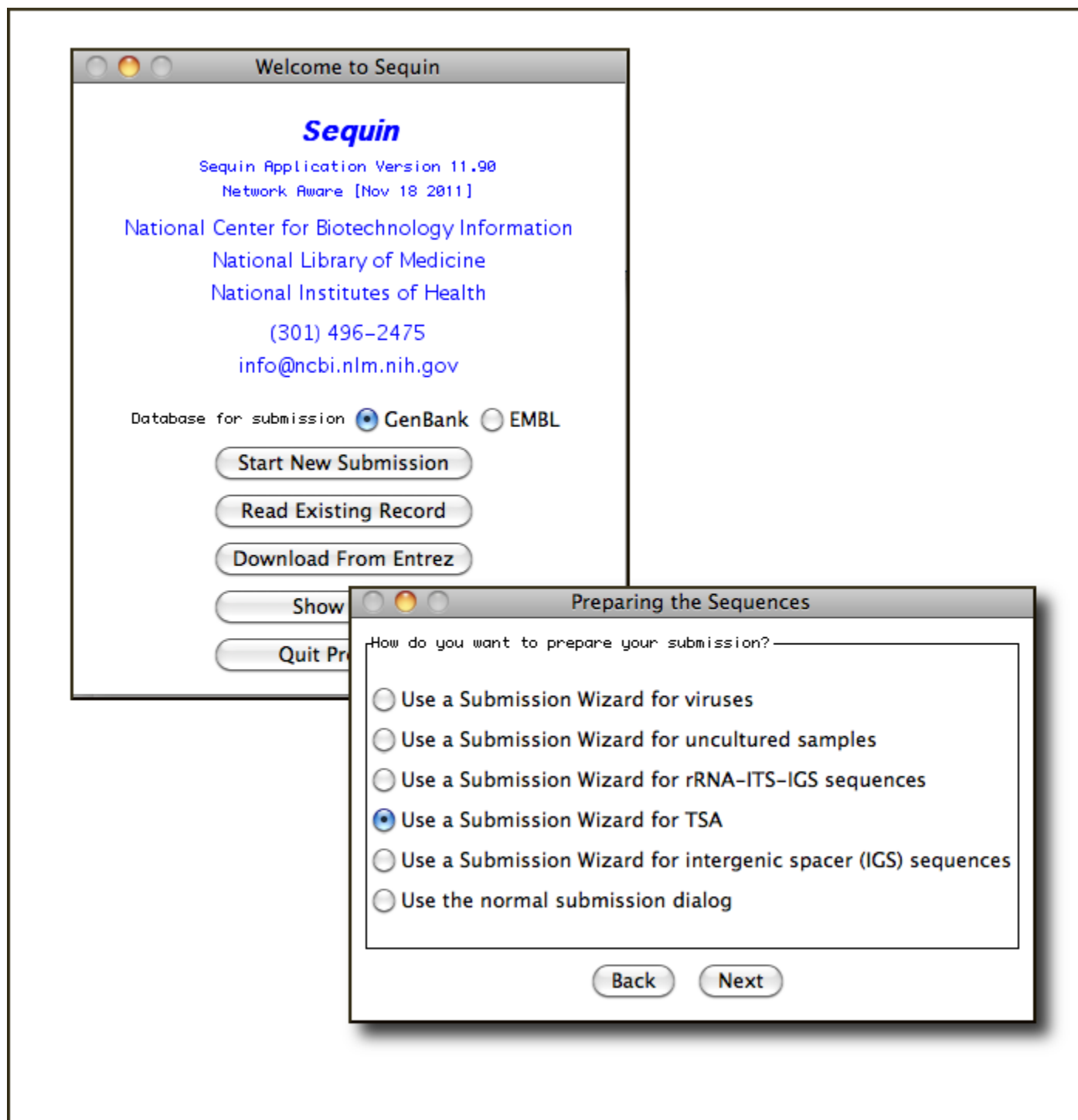


## Conserved Domain Database Update

Version 3.01 of the Conserved Domain Database is now available. The new release contains 298 new or updated NCBI-curated domain models. More detailed statistics are available from the [CDD News page](#). CDD matrices and other information can be downloaded from the [FTP site](#). CDD data are incorporated in the Entrez and BLAST search services at the NCBI Website.

## Sequin Now with Transcriptome Shotgun Assembly and Internal Transcribed Spacer Sequence Submission Wizards

A new version (11.90) of Sequin, the NCBI's standalone submission preparation software, is now available for [download](#). Packages are available for Linux, Unix, Windows, and Mac OSX systems. Improvements include new Submissions Wizards for Transcriptome Shotgun Assemblies (TSA) and ribosomal RNA intergenic spacer sequences (ITS); a Sequencing Method Page for information about the sequencing technology and assembly methods; a Sequence Deletion Tool for removing sequences from the submission; and updated feature and qualifier wizards complying with the latest INSDC Feature Documentation. The [Sequin page](#) has more information, a [Quick guide](#), [FAQs](#), and extensive [help documentation](#) on using Sequin to prepare submissions.



## NCBI C++ Toolkit Major Release

NCBI C++ Toolkit v7.0.0 is now available from the [FTP site](#). The [release notes](#) describe the highlights and contents of this release. The Toolkit contains C++ language sources of NCBI software that can be used to build standalone BLAST, Sequin, Cn3D, and other NCBI tools and utilities. The [NCBI C++ Toolkit Book](#) has in-depth information on working with the toolkit and provides access to source browsers and other useful resources.

## Announce Lists and RSS Feeds

Seventeen topic-specific mailing lists are available that provide email announcements about changes and updates to NCBI resources including dbGaP, BLAST, GenBank, and Sequin. The various lists are described on the [Announcement List summary page](#). Subscribe to the [NCBI Announce list](#) to receive updates on the NCBI News.

Twenty-one [RSS feeds](#) are now available from NCBI including news on PubMed, PubMed Central, NCBI Bookshelf, LinkOut, HomoloGene, UniGene, and NCBI Announce.

NCBI's [Facebook](#) page and [Twitter feed](#) also provide updates on NCBI resources.

Send comments and questions about NCBI resources to [info@ncbi.nlm.nih.gov](mailto:info@ncbi.nlm.nih.gov), or call 301-496-2475 between the hours of 8:30 a.m. and 5:30 p.m. EST, Monday through Friday.