

1. OBJETIVO: Análises prévias para submissão de sequências utilizadas para desenho de Assays TaqMan (Expressão Gênica e Genotipagem).

2. RESPONSÁVEIS: Todos os usuários de Custom TaqMan Assays.

3. DESCRIÇÃO DO PROCEDIMENTO :

O sucesso de customização dos Ensaio TaqMan depende fundamentalmente da qualidade da sequência que será utilizada para o desenho dos oligos. Para isso, durante o processo de escolha das sequências deve-se levar em conta os seguintes pontos:

- ✓ **Relevância biológica;**
- ✓ **Ambiguidades – substituir por N;**
- ✓ **Cruciais análises de bioinformática;**
- ✓ **Escolha dos “target sites”.**

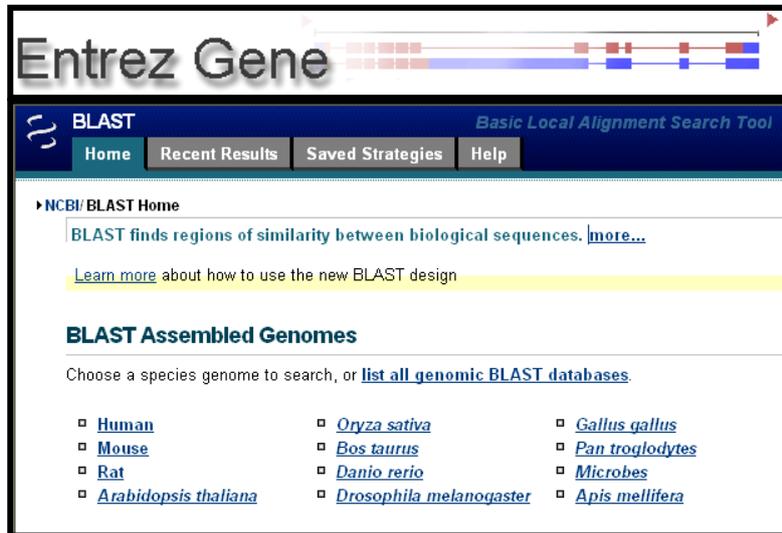
3.1 – Busca da Sequência Alvo

Buscar a sequência de interesse nos bancos de dados relacionados (Ex.: Entrez Gene, VectorBase, Ensembl etc).

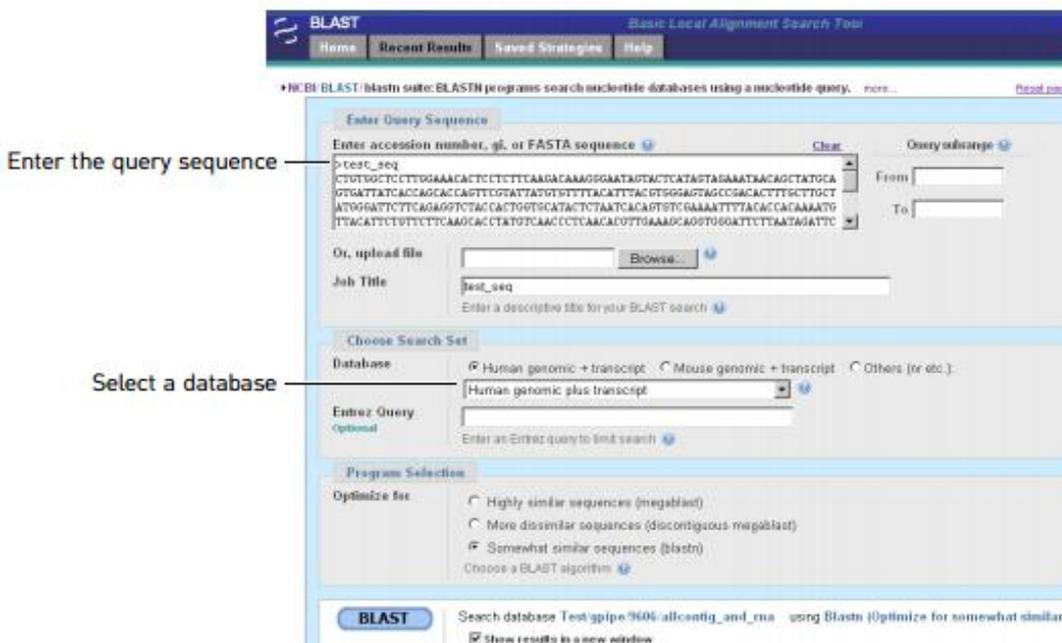
Observação: antes da pesquisa da sequência de estudo, verifique na literatura informações detalhadas sobre suas sequências alvo (por exemplo, se possui ou não isoformas de splicing, variantes, polimorfismos, regiões codificadoras e não codificadoras, e regiões promotoras). Caso seja análise de expressão gênica, verifique sequências de exons e introns e priorize o desenho das sondas e/ou primers (caso seja TaqMan) ou de primers (caso seja intercalante de DNA) em junções exon-exon.

3.2 - Avaliação da Sequência Alvo - Relevância Biológica (BLAST)

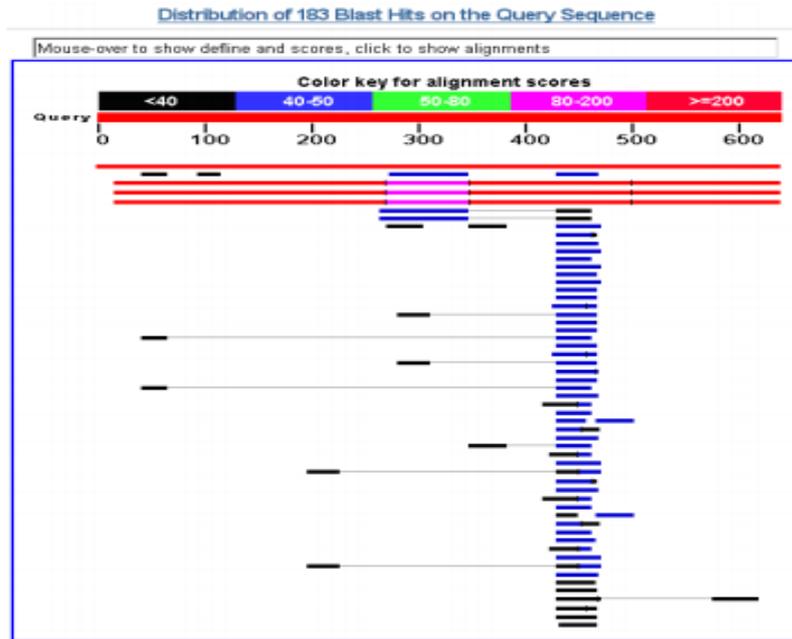
a) Acesse o link <http://blast.ncbi.nlm.nih.gov/> e insira a sequência de interesse (formato fasta):



b) Inserção da sequência alvo (formato fasta)



d) Representação gráfica do resultado da busca (alinhamento) realizado no BLAST



e) Verificar o perfil de *matching* da sequência submetida em relação ao organismo estudado e especificidade da sequência alvo.

Legend for links to other resources: [U](#) UniGene [G](#) GEO [G](#) Gene [S](#) Structure [M](#) Map Viewer

Sequences producing significant alignments:
(Click headers to sort columns)

| Accession | Description | Max score | Total score | Query coverage | E value | Max ident | Links |
|---------------------------------------|--|-----------|-------------|----------------|---------|-----------|---|
| Transcripts | | | | | | | |
| NM_010492.1 | Homo sapiens cystic fibrosis transmembrane conductance regulator (CFTR) | 115.3 | 1153 | 100% | 0.0 | 100% | U G M |
| NM_015045.2 | Homo sapiens ATP-binding cassette, sub-family C (CFTR/MRP), member 10 (ABCC10), mRNA | 39.2 | 46.4 | 11% | 0.040 | 73% | U G M |
| NM_014075.1 | Homo sapiens kinesin family member 14 (KIF14), mRNA | 39.2 | 39.2 | 3% | 5.9 | 95% | U G M |
| Genomic sequences (show first) | | | | | | | |
| NW_027533.1.4 | Homo sapiens chromosome 7 genomic contig, reference assembly | 92.2 | 1262 | 97% | 7e-126 | 100% | |
| NW_023660.1 | Homo sapiens chromosome 7 genomic contig, alternate assembly (Ba) | 92.2 | 1262 | 97% | 7e-126 | 100% | |
| NT_025926.2 | Homo sapiens chromosome 7 genomic contig, alternate assembly (Ba) | 92.2 | 1262 | 97% | 7e-126 | 100% | |
| NW_025925.1.4 | Homo sapiens chromosome 13 genomic contig, reference assembly | 82.2 | 87.3 | 18% | 0.011 | 87% | |
| NW_025912.1 | Homo sapiens chromosome 13 genomic contig, alternate assembly (b) | 82.2 | 87.3 | 18% | 0.011 | 87% | |
| NW_010498.15 | Homo sapiens chromosome 16 genomic contig, reference assembly | 46.4 | 46.4 | 6% | 0.040 | 87% | |
| NT_015926.24 | Homo sapiens chromosome 2 genomic contig, reference assembly | 46.4 | 46.4 | 6% | 0.040 | 85% | |
| NT_015923.18 | Homo sapiens chromosome 1 genomic contig, reference assembly | 46.4 | 124 | 5% | 0.040 | 93% | |
| NW_026462.1 | Homo sapiens chromosome 16 genomic contig, alternate assembly (b) | 46.4 | 46.4 | 6% | 0.040 | 87% | |
| NW_027012.1 | Homo sapiens chromosome 16 genomic contig, alternate assembly (Ba) | 46.4 | 202 | 6% | 0.040 | 90% | |
| NW_027017.1 | Homo sapiens chromosome 1 genomic contig, alternate assembly (Ba) | 46.4 | 46.4 | 5% | 0.040 | 93% | |
| NT_028409.13 | Homo sapiens chromosome X genomic contig, reference assembly | 44.6 | 44.6 | 6% | 0.14 | 85% | |
| NT_034772.5 | Homo sapiens chromosome 5 genomic contig, reference assembly | 44.6 | 44.6 | 6% | 0.14 | 87% | |
| NW_027717.1 | Homo sapiens chromosome X genomic contig, alternate assembly (Ba) | 44.6 | 44.6 | 6% | 0.14 | 85% | |
| NW_027751.1 | Homo sapiens chromosome 5 genomic contig, alternate assembly (Ba) | 44.6 | 44.6 | 6% | 0.14 | 87% | |
| NT_011051.16 | Homo sapiens chromosome X genomic contig, reference assembly | 42.8 | 163 | 5% | 0.48 | 96% | |
| NT_026437.13 | Homo sapiens chromosome 14 genomic contig, reference assembly | 42.8 | 162 | 6% | 0.48 | 90% | |
| NT_026419.13 | Homo sapiens chromosome 12 genomic contig, reference assembly | 42.8 | 81.9 | 10% | 0.48 | 87% | |
| NT_025791.24 | Homo sapiens chromosome 12 genomic contig, reference assembly | 42.8 | 162 | 6% | 0.48 | 96% | |
| NT_014354.18 | Homo sapiens chromosome 8 genomic contig, reference assembly | 42.8 | 93.7 | 6% | 0.48 | 84% | |
| NT_024487.18 | Homo sapiens chromosome 4 genomic contig, reference assembly | 42.8 | 121 | 8% | 0.48 | 95% | |
| NW_027715.1 | Homo sapiens chromosome X genomic contig, alternate assembly (Ba) | 42.8 | 122 | 5% | 0.48 | 90% | |
| NW_025232.1 | Homo sapiens chromosome 14 genomic contig, alternate assembly (b) | 42.8 | 122 | 6% | 0.48 | 90% | |
| NW_025299.1 | Homo sapiens chromosome 12 genomic contig, alternate assembly (b) | 42.8 | 121 | 10% | 0.48 | 87% | |
| NW_023184.1 | Homo sapiens chromosome 6 genomic contig, alternate assembly (Ba) | 42.8 | 242 | 6% | 0.48 | 96% | |
| NW_022162.1 | Homo sapiens chromosome 4 genomic contig, alternate assembly (Ba) | 42.8 | 42.8 | 5% | 0.48 | 84% | |

- f) Checagem da sequência escolhida por meio do perfil de matching acima estabelecido (especificidade da sequência).

```
> ref|NM\_000492.3| UEGM Homo sapiens cystic fibrosis transmembrane conductance regulator
(ATP-binding cassette sub-family C, member 7) (CFTR), mRNA
Length=6132

Score = 1153 bits (1278), Expect = 0.0
Identifier = 639/639 (100%), Gaps = 0/639 (0%)
Strand=Plus/Plus

Query 1 CTGTGGCTCCTTGGAAACACTCCTCTTCAAGACAAAGGGAATAGTACTCATAGTAGAAAT 60
Sbjct 2773 CTGTGGCTCCTTGGAAACACTCCTCTTCAAGACAAAGGGAATAGTACTCATAGTAGAAAT 2832

Query 61 AACAGCTATGCAGTGATTATCACCAGCACCAGTTCGTATTATGTGTTTTACATTTACGTG 120
Sbjct 2833 AACAGCTATGCAGTGATTATCACCAGCACCAGTTCGTATTATGTGTTTTACATTTACGTG 2892

Query 121 GGAGTAGCCGACACTTTGCTTGCTATGGGATTTCCAGAGGTTACCCACTGGTGCATACT 180
Sbjct 2893 GGAGTAGCCGACACTTTGCTTGCTATGGGATTTCCAGAGGTTACCCACTGGTGCATACT 2952

Query 181 CTAATCACAGTGTGAAAATTTTACACCACAAAATGTTACATTCTGTTCTTCAAGCACCT 240
Sbjct 2953 CTAATCACAGTGTGAAAATTTTACACCACAAAATGTTACATTCTGTTCTTCAAGCACCT 3012

Query 241 ATGTCAACCCTCAACACGTTGAAAGCAGGTGGGATTTTAAATAGATTCTCCAAAGATATA 300
Sbjct 3013 ATGTCAACCCTCAACACGTTGAAAGCAGGTGGGATTTTAAATAGATTCTCCAAAGATATA 3072

Query 301 GCAATTTGGATGACCTTCGCCTTTACCATATTTGACTTCATCCAAGTTGTTAATTT 360
Sbjct 3073 GCAATTTGGATGACCTTCGCCTTTACCATATTTGACTTCATCCAAGTTGTTAATTT 3132

Query 361 GTGATTGGAGCTATAGCAGTTGTCGCGATTTTACAACCCATCATCTTTGTTGCAACAGTG 420
Sbjct 3133 GTGATTGGAGCTATAGCAGTTGTCGCGATTTTACAACCCATCATCTTTGTTGCAACAGTG 3192

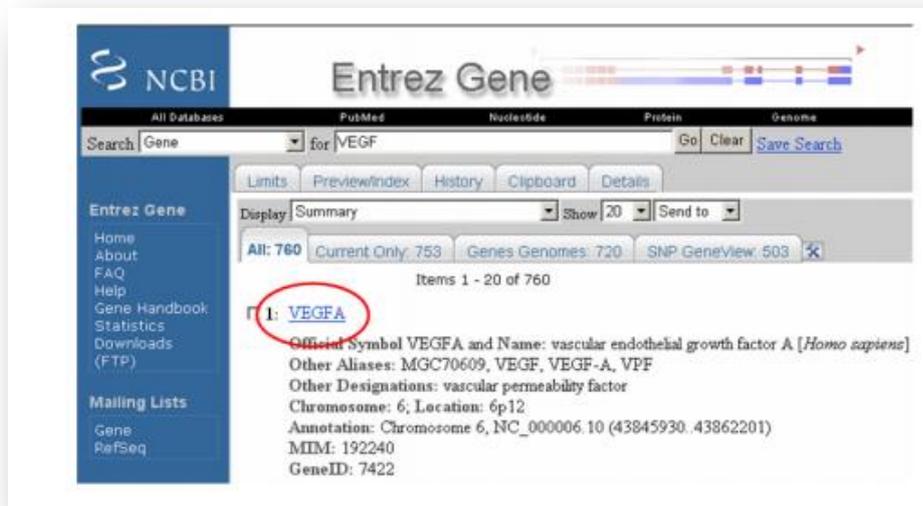
Query 421 CCAGTGATAGTGGCTTTTATTATGTTGAGAGCATATTTCTCCAAACCTCACAGCACTC 480
Sbjct 3193 CCAGTGATAGTGGCTTTTATTATGTTGAGAGCATATTTCTCCAAACCTCACAGCACTC 3252

Query 481 AAACAACCTGGAATCTGAAGCAGGAGTCCAAATTTTCACTCATCTTTGTTACAAGCTTAAAA 540
Sbjct 3253 AAACAACCTGGAATCTGAAGCAGGAGTCCAAATTTTCACTCATCTTTGTTACAAGCTTAAAA 3312

Query 541 GGACTATGGACACTTCGTGCTTCGGACGGCAGCCTTACTTTGAAACTCTGTTCCACAAA 600
Sbjct 3313 GGACTATGGACACTTCGTGCTTCGGACGGCAGCCTTACTTTGAAACTCTGTTCCACAAA 3372

Query 601 GCTCTGAATTTACATACTGCCAACTGGTCTTGTACCTG 639
Sbjct 3373 GCTCTGAATTTACATACTGCCAACTGGTCTTGTACCTG 3411
```

- g) Após avaliar a relevância da sequência e identificar a região de interesse, proceder com o mapeamento físico do gene para encontrar a melhor região para desenho do ensaio. Para isso acesse o site <http://www.ncbi.nlm.nih.gov/pubmed/> e selecione Gene.



NCBI Entrez Gene

All Databases PubMed Nucleotide Protein Genome

Search: Gene for VEGF [Go] [Clear] [Save Search]

Limits Preview/Index History Clipboard Details

Entrez Gene

Home About FAQ Help Gene Handbook Statistics Downloads (FTP) Mailing Lists Gene RefSeq

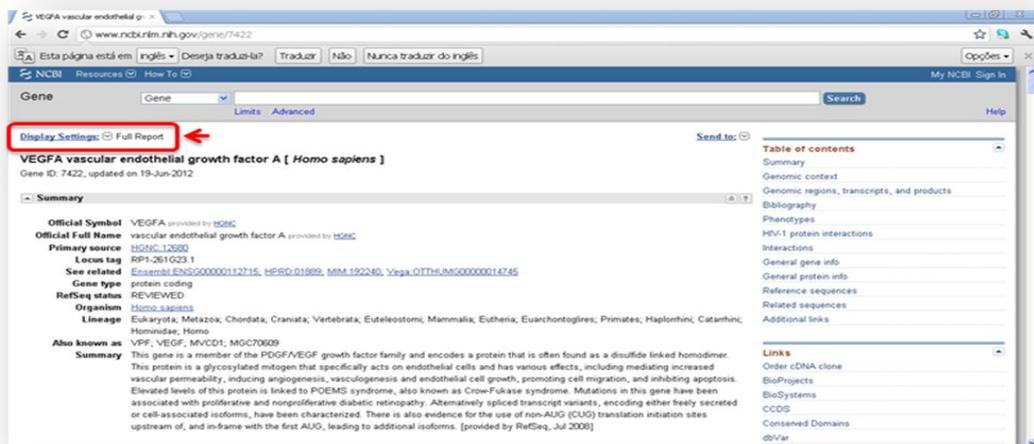
Display: Summary Show 20 Send to

All: 760 Current Only: 753 Genes Genomes: 720 SNP GeneView: 503

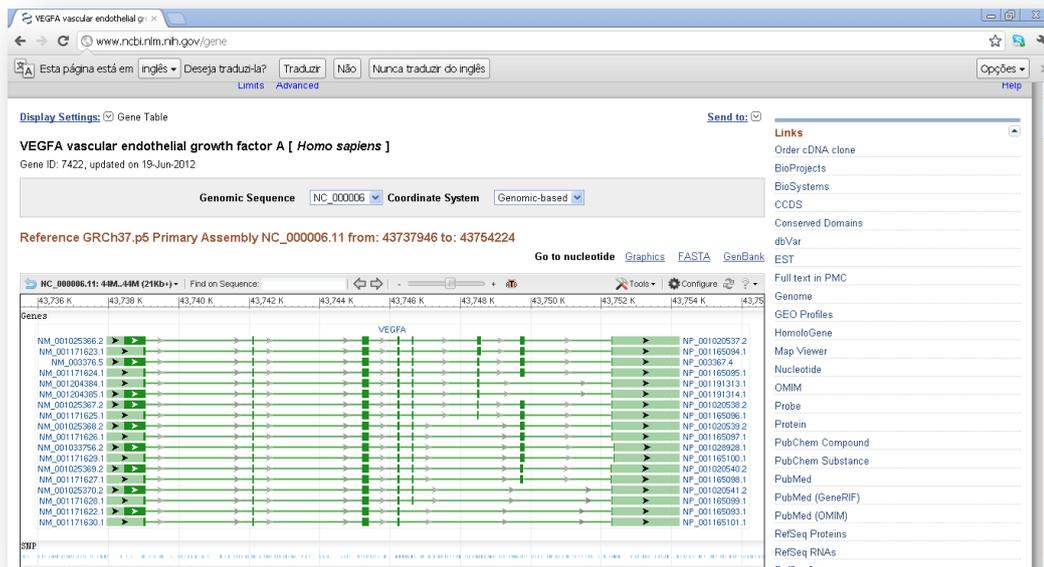
Items 1 - 20 of 760

1: **VEGFA**

Official Symbol VEGFA and Name: vascular endothelial growth factor A [*Homo sapiens*]
 Other Aliases: MGC70609, VEGF, VEGF-A, VPF
 Other Designations: vascular permeability factor
 Chromosome: 6; Location: 6p12
 Annotation: Chromosome 6, NC_000006.10 (43845930..43862201)
 MIM: 192240
 GeneID: 7422



O Mapeamento físico de uma sequência vai possibilitar a escolha/validação da região de interesse que será utilizada no desenho dos oligos. Nesse momento há possibilidade de escolher a sequência inteira ou parte dessa.



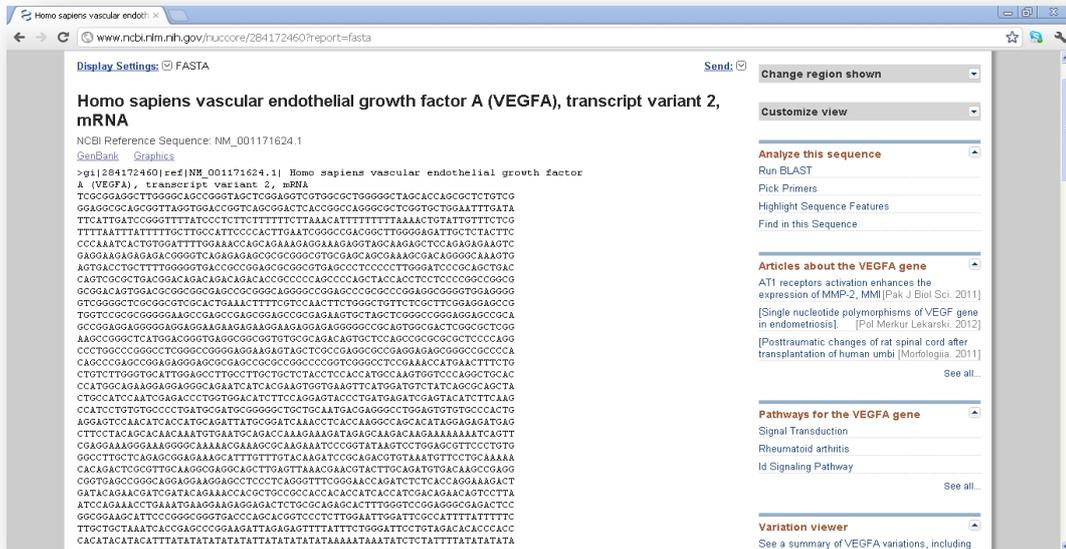
The screenshot shows the NCBI Gene browser for VEGFA (Homo sapiens). The main view displays the genomic sequence with various annotations, including exons and introns. The gene is identified as VEGFA, Gene ID: 7422, updated on 19-Jun-2012. The reference sequence is GRCh37.p5 Primary Assembly NC_000006.11 from 43737946 to 43754224. The interface includes a search bar, navigation tools, and a detailed view of the gene structure with various annotations and links.

Supondo que seja utilizada uma parte da sequência previamente validada:



The screenshot shows the NCBI Nucleotide database for the mRNA sequence of VEGFA (Homo sapiens). The main view displays the mRNA sequence with various annotations, including exons and introns. The gene is identified as Homo sapiens vascular endothelial growth factor A (VEGFA), transcript variant 2, mRNA. The reference sequence is NM_001171624.1. The interface includes a search bar, navigation tools, and a detailed view of the mRNA sequence with various annotations and links.

Pegue a sequência FASTA.



h) Acesse o programa: <http://www.repeatmasker.org/> para fazer o mascaramento de regiões repetitivas e de baixa complexidade.



Services

- [RepeatMasking](#)
- [Protein-based RepeatMasking](#)
- [Pre-Masked Genomes](#)
- [Server Queue Status](#)
- [FEAST - Gene Prediction](#)

Documentation

- [FAQ](#)
- [RepeatMasker](#)
- [Server Configuration](#)

Welcome!

RepeatMasker is a program that screens DNA sequences for interspersed repeats and low complexity DNA sequences. The output of the program is a detailed annotation of the repeats that are present in the query sequence as well as a modified version of the query sequence in which all the annotated repeats have been masked (default: replaced by Ns). On average, almost 50% of a human genomic DNA sequence currently will be masked by the program. Sequence comparisons in RepeatMasker are performed by the program `cross_match`, an efficient implementation of the Smith-Waterman-Gotoh algorithm developed by Phil Green.

Latest News

If you would like to keep up with news and announcements relating to RepeatMasker, you can subscribe to the new [RepeatMasker Announcements List](#).

RepeatModeler Beta: Repeat Discovery Workbench Released
Wednesday, April 16, 2008

| | | | |
|--------------------|-------------------|-------|-----------------|
| Elaborado por: FMU | Revisado por: JCR | Ass.: | Data 23.01.2014 |
|--------------------|-------------------|-------|-----------------|

SEMPRE analisar as sequências antes de submeter (File Builder ou Primer Express).

Sempre inserir nos programas acima:

- 1 – Sequências na orientação 5'-----3';
- 2 – Previamente avaliadas quanto a relevância biológica e especificidade;
- 3 – Previamente analisadas com relação a presença de elementos repetitivos (RepeatMasker);
- 4 – Utilizar somente a sequência no formato **FASTA** nos softwares File Builder e Primer Express;
- 5 – **NÃO retire os N das sequências (tanto no File Builder quanto no Primer Express submeta as sequências com N).**
- 5 – Para SNPs sempre inserir 100 bases 5' e 100 bases 3' do polimorfismo

E.x.:

File Builder:

```
>gnl|dbSNP|rs81505258|allelePos=101|totalLen=201|taxid=9823|snpclass=1|alleles='C/T'|mol=Genomic|build=131  
CCATTTATATCAGCGTTCTTTTTATGAAAAGCATCTGACTAGAAATTCAATAGAGAAGCATGTCATAATCAGTT  
TGAGGCCAAATCTTTAAATATAACA[C/T]GCTAATTTTAGTCCTAAGCATTTCAGCTGTACTTTATTGGTTTACAG  
CTTCATAGGAAGATGTGGCTTCTTAAGGAGGAAAAATCTTTTTTATATGGCTT
```

Primer Express:

```
>gnl|dbSNP|rs81505258|allelePos=101|totalLen=201|taxid=9823|snpclass=1|alleles='C/T'|mol=Genomic|build=131  
CCATTTATATCAGCGTTCTTTTTATGAAAAGCATCTGACTAGAAATTCAATAGAGAAGCATGTCATAATCAGTT  
TGAGGCCAAATCTTTAAATATAACA[Y]GCTAATTTTAGTCCTAAGCATTTCAGCTGTACTTTATTGGTTTACAGCTT  
CATAGGAAGATGTGGCTTCTTAAGGAGGAAAAATCTTTTTTATATGGCTT
```

Abaixo em destaque está um SNP, que NÃO é possível desenhar assays TaqMan para avaliação desse SNP.

Não mascarada

> AY853303

GCCCTCTTAATAAGGCTTATGTGCTTCTGCACAGTTCACCTGACTGTGAAATGGGTATGATTGCCCTTTTGTAC
AGATGGAGAAACTGAGGCTCAGAAATCGCTAAGGGATTGCCACAGGTCCACAGTTCCCAGGCTTCCCAATC
CACTCTCCACTCCCAGGTGTGGAGTGGAGTGGAGAGTCTTGCGCCAGGTCCTCA

Mascarada

>AY853303

GCCCTCTTAATAAGGCTTATGTGCTTCTGCACAGTTCACCTGANNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN
NNCC
CAGGCTTCCAATCCACTCTCCACTCCCAGGTGTGGAGTGGAGTGGAGAGTCTTGCGCCAGGTCCTCA

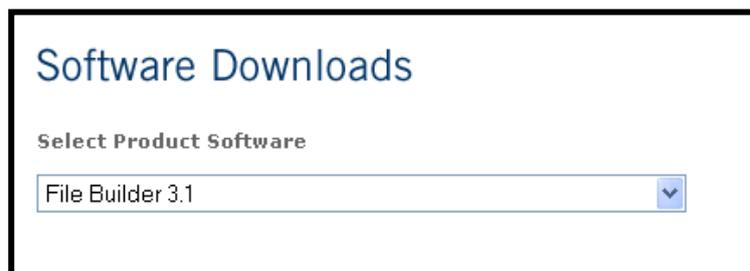
Custom TaqMan Assays

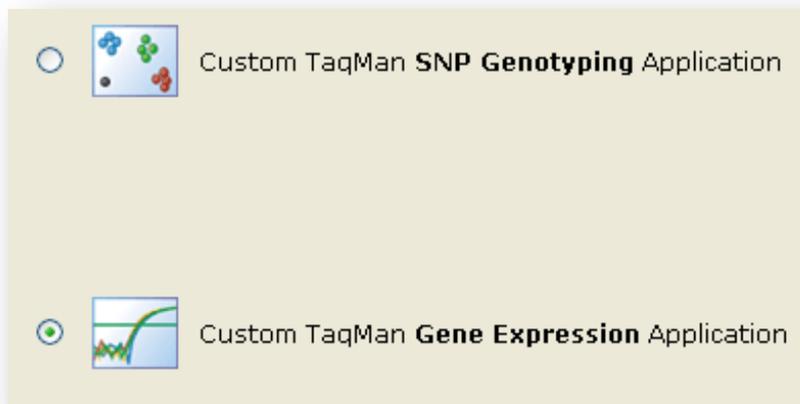
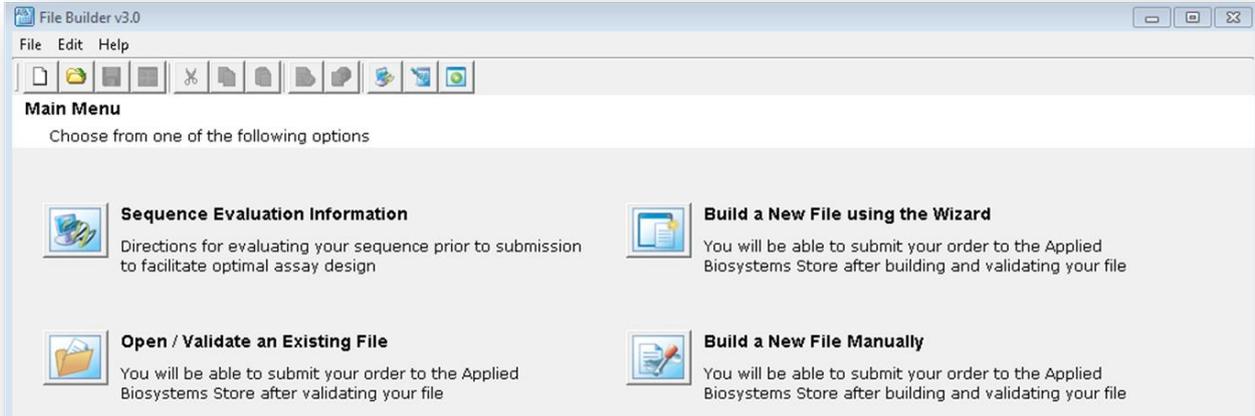
Existem duas possibilidades para o desenho de ensaios customizados:

1. File Builder v3.1 (Gratuito):

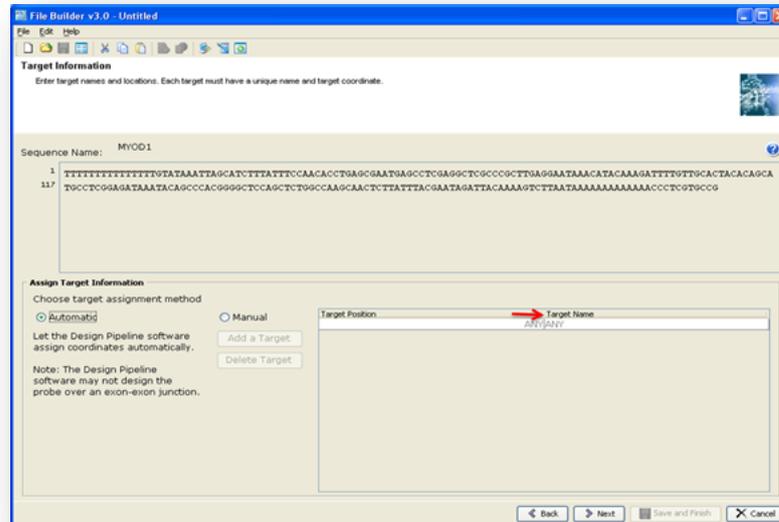
<http://www6.appliedbiosystems.com/support/software/assaysbydesign/installs.cfm>

- a) Usuário envia arquivo e a Life desenha e envia o ensaio em tubo único (a Life não faz as análises de sequências mencionadas anteriormente).
- b) A Life somente realiza desenho de ensaios TaqMan. Não existe a possibilidade de desenho somente das sequências de primers (Ex. Primers para agentes intercalantes).
- c) Software para gerar e validar arquivo de desenho de primers e sondas





- GX Small Scale 20x (140 x 50µl or 360 x 20µl)
Part Number: **4331348**
- GX Medium Scale 20x Concentration 750 (20µl) reactions
Part Number: **4332078**
- GX Large Scale 60x Concentration 2,900 (20µl) reactions
Part Number: **4332079**



File Builder v3.0 - Untitled

File Edit Help

Target Information
Enter target names and locations. Each target must have a unique name and target coordinate.

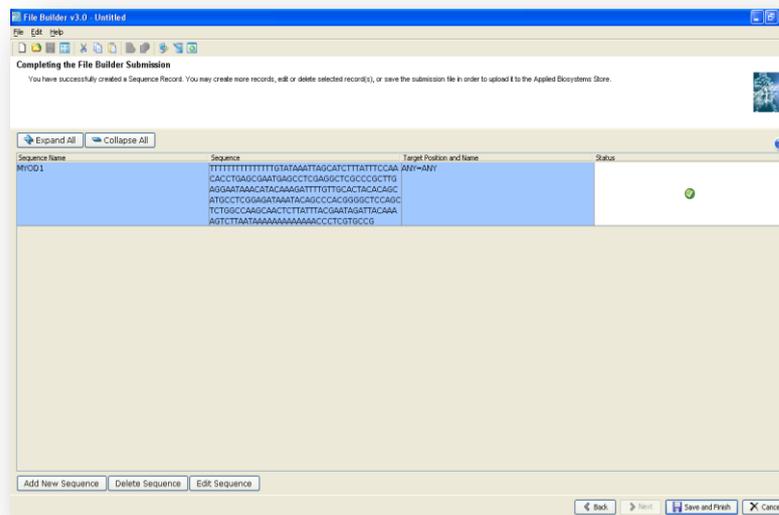
Sequence Name: MYOD1

1
117
TTTTTTTTTTTTTTGTATAAATTAGCATCTTTATTTCCAACACCTGAGGAAATGAGCCTCGAGGCTCGCCCGTTGAGGAATAAACATACAAAGATTTTGTGCCTACACAGCA
TGCCTCGAGATAAATACAGCCACGGGGCTCCAGCTCGGCCAAGCAACTCTTTATTCGAAATAGATTACAAAGTCTTAAATAAAAAAAAAAAACCTCTGTCCG

Assign Target Information
Choose target assignment method
 Automatic Manual
Let the Design Pipeline software assign coordinates automatically.
Add a Target Delete Target
Note: The Design Pipeline software may not design the probe over an exon-exon junction.

Target Position: 217 Target Name: ATGTVV

Back Next Save and Finish Cancel



File Builder v3.0 - Untitled

File Edit Help

Completing the File Builder Submission
You have successfully created a Sequence Record. You may create more records, edit or delete selected records, or save the submission file in order to upload it to the Applied Biosystems Store.

Expand All Collapse All

| Sequence Name | Sequence | Target Position and Name | Status |
|---------------|---|--------------------------|--------|
| MYOD1 | TTTTTTTTTTTTTTGTATAAATTAGCATCTTTATTTCCAA ACCTGAGGAAATGAGCCTCGAGGCTCGCCCGTTG AGGAATAAATACAAAGATTTTGTGCCTACACAGC ATGCCTCGAGATAAATACAGCCACGGGGCTCCAGC TCTGCCAAGCAACTCTTTATTCGAAATAGATTACAA AGTCTTAAATAAAAAAAAAAAACCTCTGTCCG | ANY=ANY | ✓ |

Add New Sequence Delete Sequence Edit Sequence

Back Next Save and Finish Cancel

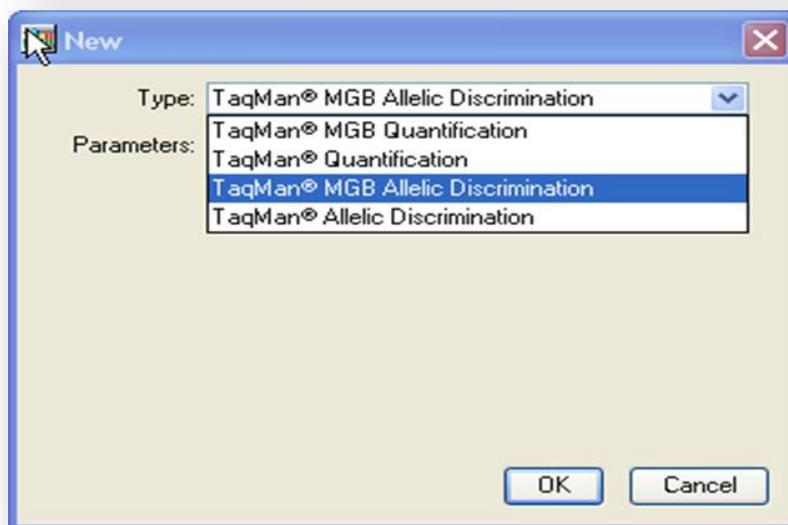
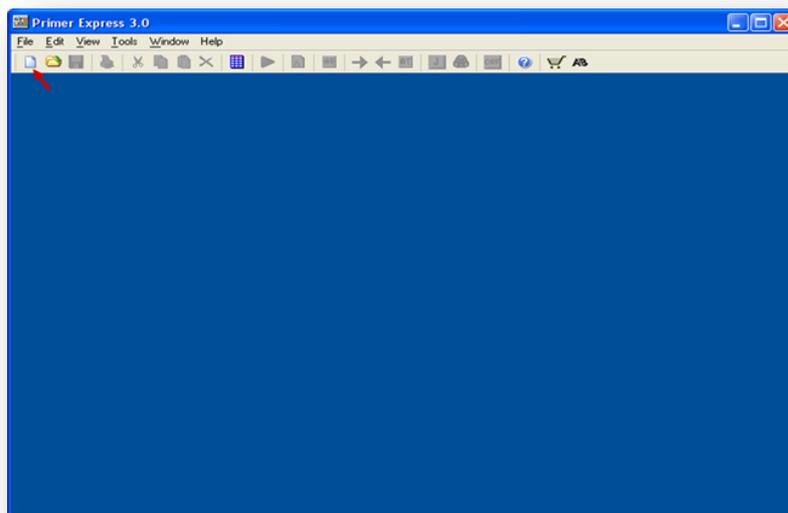
Nota (1): Acima está um exemplo de submissão para desenho de ensaios de expressão gênica. **Caso seja para genotipagem de SNPs o usuário deverá mostrar a localização do SNP da seguinte forma [base1/base2] – por exemplo: [A/G] – sinaliza que trata-se de uma troca de A para G.**

Nota (2): O **Target Name** (em destaque acima) poderá conter no máximo 4 caracteres.

Nota (3): Os arquivos serão salvos em .txt. Enviar esse arquivo anexado para o Customer Care Brasil: orcamento.br@lifetech.com

2. Primer Express (Software de licença única que acompanha todos os instrumentos de qPCR da Life Technologies, também existe a possibilidade para compra separadamente).

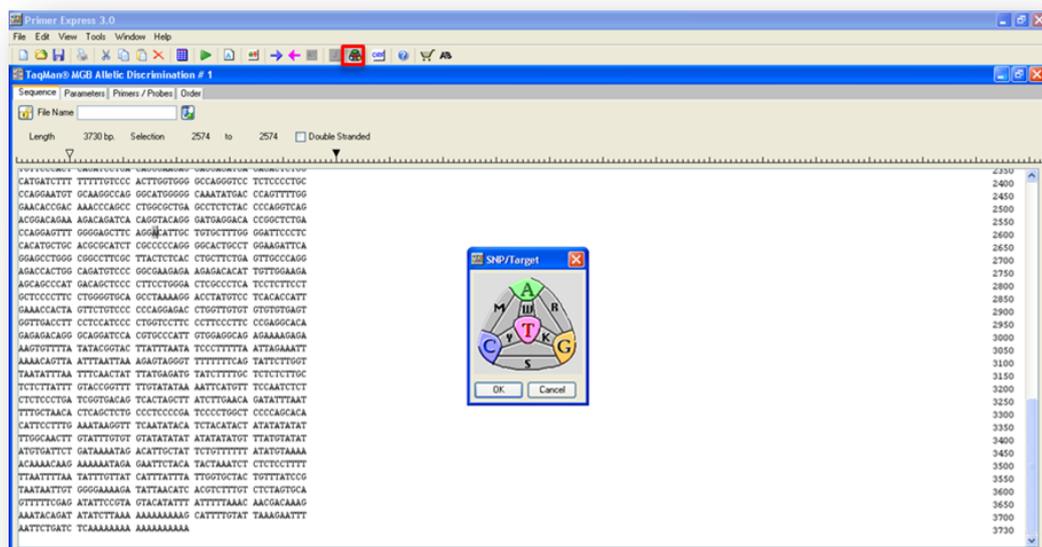
Abra o programa Primer Express, selecione a aplicação (quantificação ou discriminação alélica) e em seguida copie e cole a sequência previamente analisada.



As sondas e os primers desenhados pelo software são utilizados para as seguintes aplicações:

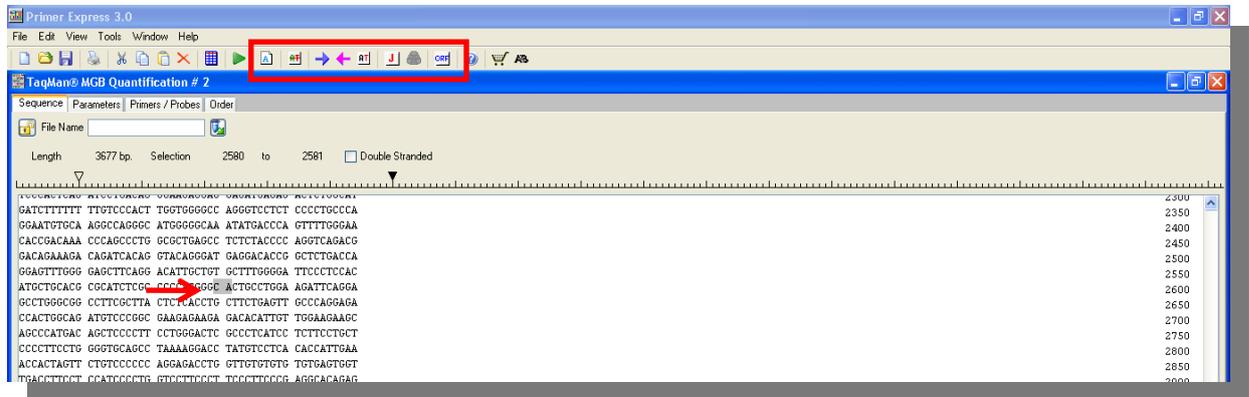
- Quantificação Absoluta/Relativa; Presença/Ausência; e outras aplicações
- **Discriminação Alélica**

No caso de discriminação alélica, selecione o SNP de interesse e vá ao ícone SNP target (em destaque) para que assim possa ser incluído a especificação dos alelos de interesse.

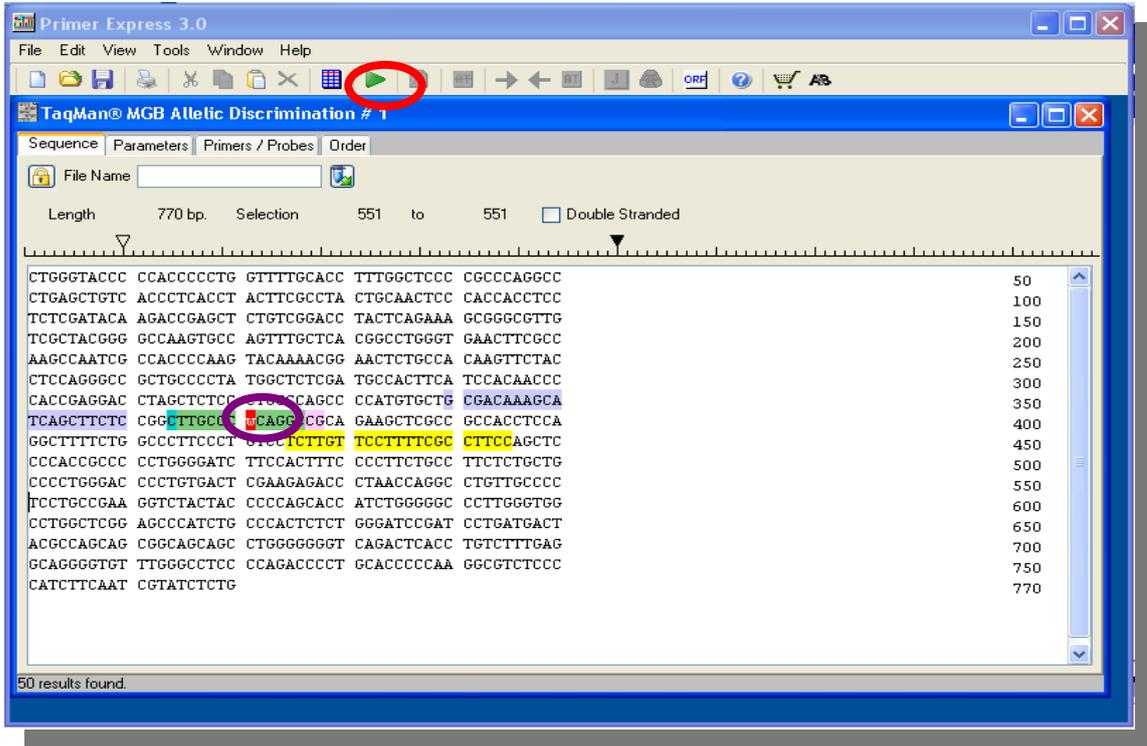


Ícone ativado apenas para discriminação alélica.

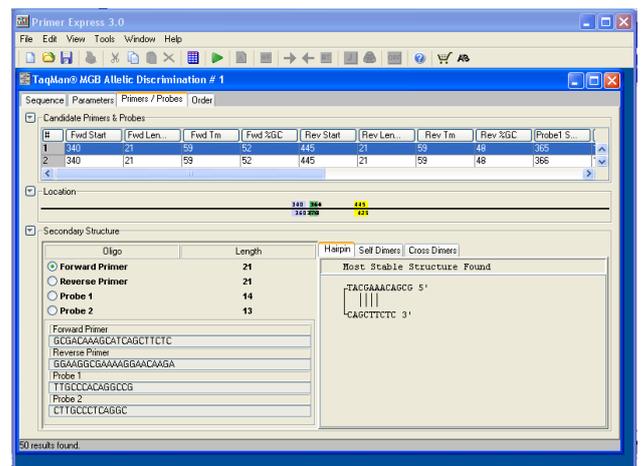
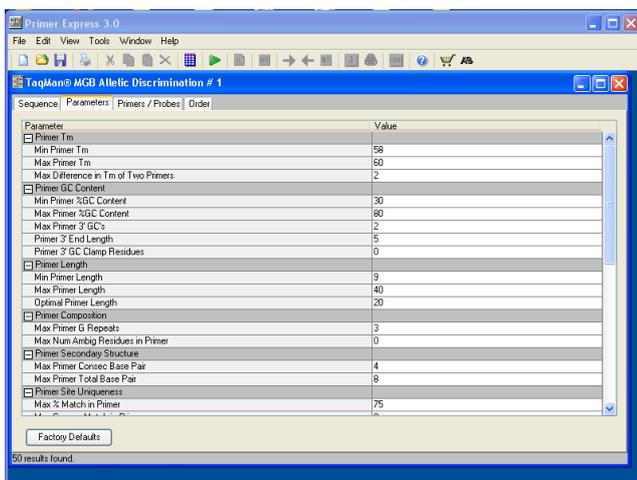
Os ícones para anotações (edições) no Primer Express (abaixo) só são ativados quando destacamos a região a ser editada.



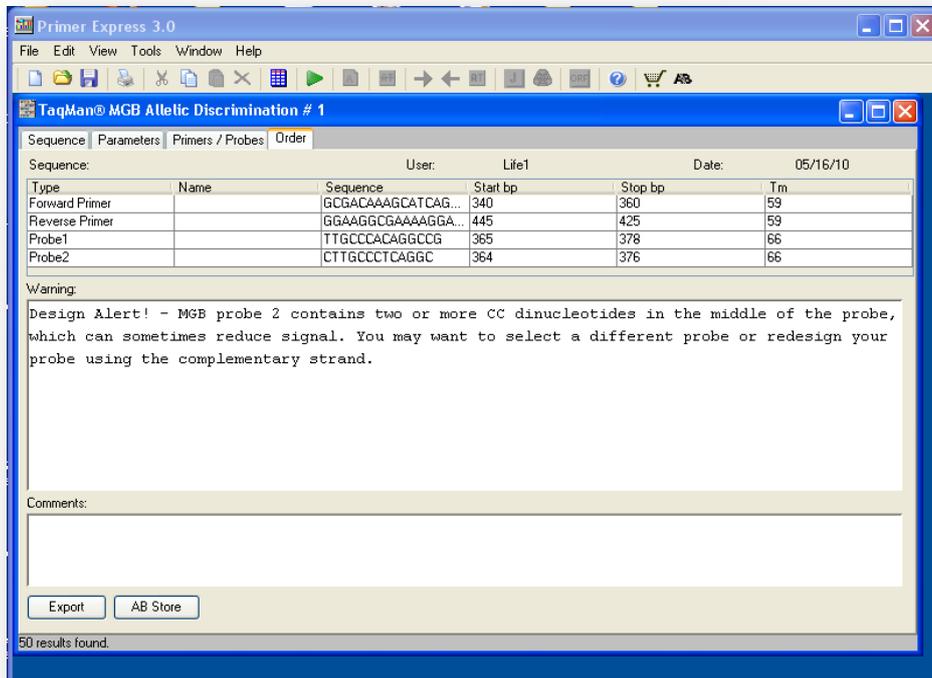
| Annotation | Toolbar Button | Description | Example (shows the default annotation colors) |
|---|---|---|--|
| Clear |  | Removes the annotation from the selected region. Note: You can select Edit > Annotate > Clear All to remove all annotations in the sequence. | CGCAGCAGGT |
| Exclude |  | Excludes the selected region from the primer/probe search. Note: You can select Edit > Find and Exclude to search for a specific sequence, then exclude it. Note: Bases that are excluded (using the Exclude annotation) are not considered in primer and probe design. Masked bases (bases substituted with Ns) are considered. | CGCAGCAGGT |
| Fwd Primer |  | Specifies the selected region as a forward primer. When the software finds primers and probes, it looks only for reverse primers and probes that are compatible with the annotated forward primer. When selecting primer sequences, select sequences close enough to the probe sequence to ensure that amplicon length is within 50 to 150 bases. | CGCAGCAGGT |
| Rev Primer |  | Specifies the selected region as a reverse primer. When the software finds primers and probes, it looks only for forward primers and probes that are compatible with the annotated reverse primer. When selecting primer sequences, select sequences close enough to the probe sequence to ensure that amplicon length is within 50 to 150 bases. | CGCAGCAGGT |
| Probe Quantification documents only |  | Specifies the selected region as a probe. When the software finds primers and probes, it looks only for forward and reverse primers that are compatible with the annotated probe region. | CGCAGCAGGT |
| Junction Quantification documents only |  | Creates a junction annotation across two adjacent bases to mark a junction. When the software finds primers and probes, it attempts to find at least one primer or probe in each primer/probe set that crosses at least one junction. Note: The junction annotation supports only primer and probe design. It does not locate exon-exon junctions. Before applying a junction annotation, reference online bioinformatics databases or existing publications to locate junctions in a sequence. The Junction annotation can be used for a variety of applications, including: <ul style="list-style-type: none"> Exon-exon junction detection, to generate primers that amplify only mRNA (or cDNA made from it), but not genomic DNA. SIRNA and miRNA detection Transgenic DNA detection You can add any number of junction annotations to the sequence. | TGACTTCTCG  ATGGGTACT |
| SNP Target Allelic Discrimination documents only |  | Assigns the selected base as the SNP (single nucleotide polymorphism) target site. When you select this annotation, the SNP Target dialog box is displayed. Click the IUPAC code that represents the possible variations at the SNP target site, then click OK. SNP site location in MGB probes SNP site location in conventional probes |  CGACAAGG |



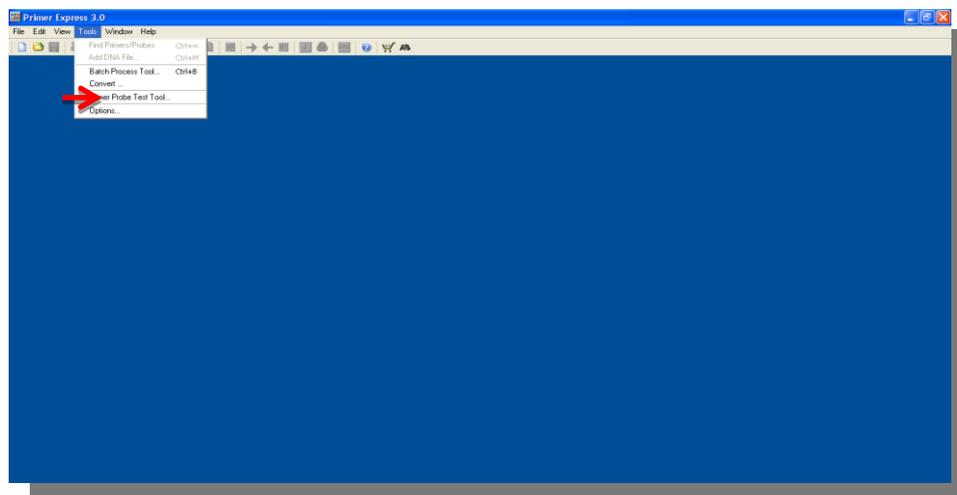
Parâmetros de análise (recomendado deixar *default*): Tela de oligos e estruturas secundárias:

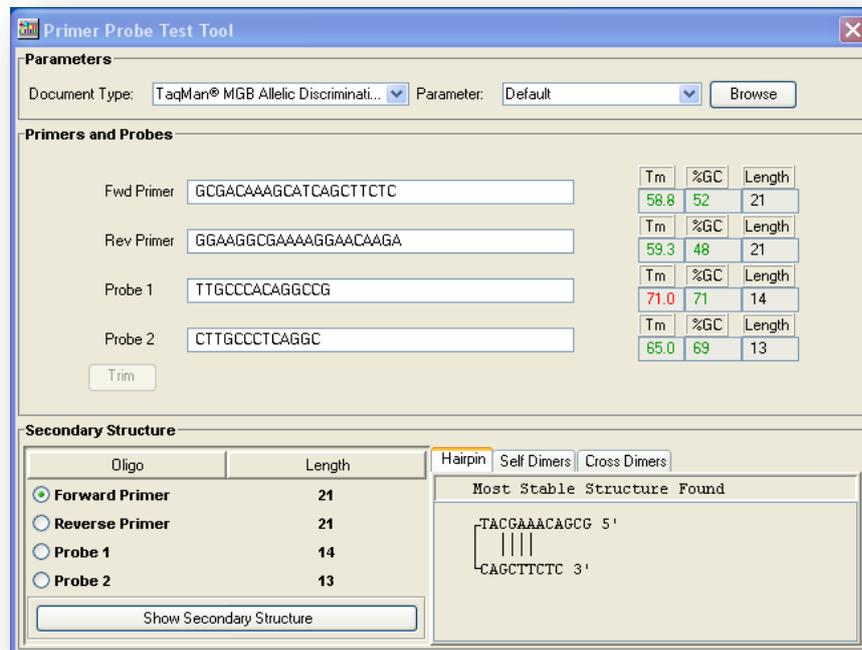


Tela de pedido ("Order"), nesse momento o usuário irá salvar (.txt) o conjunto de oligos escolhidos. Enviar esse arquivo anexado para o Customer Care Brasil: orcamento.br@lifetech.com



Tela para avaliação de oligos previamente desenhados. Vá em *Tools* --- *Primer Probe Test Tool* :





Nota (1): Acima está um exemplo de desenho de oligos para genotipagem (por isso há o desenho de duas probes). Para ensaios de expressão gênica, quantificação, dentre outros, o usuário pode optar pelo uso de sondas ou não (SYBR e MeltDoctor/HRM, por exemplo). Dessa forma, com o software Primer Express é possível tanto escolher para a síntese: primers + sondas TaqMan ou somente primers.

Nota (2): A Life recomenda a utilização de sondas com quencher MGB (para qualquer aplicação).

Nota (3): Os arquivos poderão ser salvos em .txt. Enviar esse arquivo anexado, contendo os primers e sonda(s) TaqMan, ou selecione apenas as sequências dos primers sempre na aba "Order" (no caso de SYBR e HRM) e envie para o Customer Care Brasil: orcamento.br@lifetech.com

Nota Final: Este tutorial tem por objetivo apresentar os requerimentos básicos para submissão de sequências para a Life Technologies realizar o desenho ou para desenho de oligos *in house*. Vale ressaltar que as análises de bioinformática constituem poderosas ferramentas para obtenção de bons oligos, porém não uma garantia de desenho perfeito!