

Tiling Path File (TPF) Specification v1.4: 2/15/2011

What it is: A TPF specifies the order of a set of sequences along a larger molecule(s). Gap types and sizes can be specified. The molecule can be as large as a chromosome and as small as a sequence contig made up of one smaller sequence. A TPF can have one or more sequences/clones.

What it is not: A TPF does not specify the particular version of a sequence, nor does it specify the switch points between adjacent clones in a given scaffold/contig.

Header Information: Header information is provided in a key-value type of structure. Each header field will be prefixed with ##. These fields are available:

- ##Organism (Required)
- ##Assembly Name (Required)
- ##Chromosome (Required)
- ##Strain/Haplotype/Cultivar (Optional)
- ##Type (controlled vocabulary) (Required)
- ##Version (Calculated by Database, Required)
- ##Comment (Optional)

Additional information will be calculated at submission.

- ##Submitter
- ##Create date
- ##Update date

Here is an example header:

```
##ORGANISM: Mus musculus
##ASSEMBLY NAME: Reference
##CHROMOSOME: 1
##STRAIN/HAPLOTYPE/CULTIVAR: C57BL/6J
##TYPE: Complete Chromosome
##Version: 1
##Comment: some useful information here.
##SUBMITTER: Tina Graves, WUGSC
##CREATE DATE: Nov 3 2006 12:16PM
##UPDATE DATE: Nov 5 2006 12:12PM
```

The data defining the actual tiling path should be flanked in header and footer lines to minimize the chance of file truncation. The header and footer lines shall have the following format:

All header "## TAG: value" pairs must occur before this line:

##=== Beginning of TPF Data ===

Required after the TPF data rows:

##=== End of TPF Data ===

Column definitions for sequence lines:

Column 1: accession number. This must be a valid accession assigned by the International DNA Sequence Database (GenBank/EMBL/DDBJ). No version number should be used. It is assumed that the current version of the accession is what is intended. **OPTIONAL**, use "?" if unknown.

Column 2: clone name. If the sequence is defined by a clone, then the clone name should be specified here. Ideally, standard Clone Registry nomenclature will be used. If a sequence is derived from more than one clone, the clone name can be given as "MULTIPLE". **OPTIONAL**, use "?" if unknown.

Column 3: local contig identifier. Name given to the scaffold/contig generated by assembling the listed sequences. **REQUIRED.**

- **NOTE: Use of "." (the "dot" character) is not allowed in contig identifiers.**
- **NOTE: A local contig identifier may only be used once per assembly.**

Column 4: contained status. This line is only used to specify clones that are known to be contained within another clone on the TPF. The only values allowed are 'CONTAINED' and 'CONTAINED_TURNOUT'. **REQUIRED for contained clones, otherwise OPTIONAL.**

Column 5: accession number. This must be a valid accession assigned by the INSDC. No version number should be used. It is assumed that the current version of the accession is what is intended. The accession provided in this column must be in column 1 on another line of the same TPF and must also belong to the same local contig (column 3) as the contained clone. See below for requirements. **REQUIRED if COLUMN 4 is CONTAINED or CONTAINED_TURNOUT, otherwise not valid.**

Column 6: clone name. The clone name used in this column should correspond to the accession listed in column 5, if the sequence is derived from a clone. Ideally, standard Clone Registry nomenclature will be used. See below for requirements. **REQUIRED IF COLUMN 4 is CONTAINED or CONTAINED_TURNOUT, otherwise not valid.**

NOTE1: Columns 5 and 6- these are similar to columns 1 and 2 in that at least one, but not both, requires a value. However, if both are known, please supply both values.

NOTE2: If column 4 is not populated in a sequence line, do not populate columns 4, 5 or 6. If column 4 is populated, then provide values or "?" for columns 5 and 6.

Column definitions for non-sequence lines:

Column 1: GAP. This is the term used and it should always be capitalized. Used to note the gap lines. **REQUIRED**

Column 2: gap type. Specifies the type of gap. **REQUIRED**

Values are:

- ✚ TYPE-1: [Deprecated]- was a place-holder for a picked clone.
- ✚ TYPE-2: clone gap
- ✚ TYPE-3: contig gap- unable to close using available technology
- ✚ Biological Gap: If there is a biological gap such as a centromere, etc. then use the name rather than type-4. This is a controlled vocabulary:
 - ⊖ CENTROMERE
 - ⊖ TELOMERE
 - ⊖ HETEROCHROMATIN
 - ⊖ SHORT-ARM
- ✚ PAR: (Y-chromosome only) The pseudoautosomal regions of the Y chromosome are represented by PAR gaps. Accessions for chr. X-derived PAR sequences are omitted from the chr. Y TPF, but are present in the chr. Y AGP. Switch points for PAR boundary clones will be curated manually.
 - Note: The comments section of the header should be used to provide the bp positions at which the PAR region begins and ends in the relevant accessions. Header comments are entered via the web-based form at the time of submission. This information will be inserted into the ##COMMENT line of the header. It will also be stored in the database and can be retrieved for future use.

Column 3:

- For TYPE-2, TYPE-3 or Biological gap: Gap size. Estimated size x gap. In the absence of submitted gap sizes a default of 50,000 bp will be used for clone and contig gaps. **OPTIONAL**, unless the gap type is 'Biological' then required.
- For PAR gap: accession number of the first sequence to contain the PAR region, as defined on the X chromosome (this accession may contain both non-PAR and PAR sequence)

Column 4:

- For TYPE-2, TYPE-3 or Biological gap: Method used to determine gap size. If a gap size has been estimated experimentally, the method should be noted here. Currently acceptable values:
 - FISH
 - OPTICAL MAP
 - ALIGNMENT
 - PCR
 - FINGERPRINT

- Multiple methods may be entered for a single gap. Methods should be separated by a semi-colon (Example: FISH;OPTICAL MAP;ALIGNMENT).
- **REQUIRED if column 3 is populated and gap type is not biological, otherwise not valid.**
- For PAR gap: accession number of the last sequence to contain the PAR region, as defined on the X chromosome (this accession may contain both non-PAR and PAR sequence)

All columns are tab delimited and lines are terminated by a newline. Lines beginning with a single “#” can occur anywhere within the file, are comments and can be ignored by parsers. Lines beginning with “##” can be ignored by simple parsers but contain structured information about the TPF. Each file should have only one header, so all of the objects described in a file should have common header information.

Validation:

For sequence based lines:

- ✚ If column 1 is populated with a valid accession, column 2 may be unpopulated (using “?” as a placeholder). Ideally, this will only occur in the case where a sequence is not based on a defined clone and will have no other easily identified name.
- ✚ If an accession is supplied, it must be valid according to the IDNSC.
- ✚ A given accession cannot be used more than once per assembly, but may be used in >1 assembly. (The assembly name is listed in row 3 of the TPF header).
- ✚ If column 1 is not populated (using “?” as a placeholder) then column 2 should be populated.
- ✚ Use of the “.” character is not permitted in column 3.
- ✚ If there is not enough information to populate column 1 or 2, then a gap line should be used.
- ✚ If column 4 is populated, then columns 5 and 6 must be populated.
- ✚ If column 4 is not populated, then columns 5 and 6 must not be populated.
- ✚ If column 4 = CONTAINED or CONTAINED_TURNOUT, the accession provided in column 5 must be present in the same local contig (column 3) as the accession in column 1.

For gap lines:

- ✚ Gap type-1 lines are no longer allowed.
- ✚ Gap type-2 lines will default to 50Kb if no other data is provided.
- ✚ Gap type-2 lines may not be adjacent to one another.
- ✚ Gap type-3 lines will default to 50 Kb if no data is provided.
- ✚ Gap type-3 lines may not be adjacent to one another.

- ✚ Gap type-2 and type-3 lines may not be adjacent to one another.
- ✚ TYPE-3 gaps may not be contained within a contig
- ✚ Biological gaps must provide a size estimate in column 3; they do not need to have a method in column 4.
- ✚ Biological gaps may not be contained within a contig
- ✚ There can be consecutive biological gap lines of different types.
- ✚ If a gap size is provided in column 3, the method used to determine this size must be provided in column 4, unless column 2 is a biological gap.
- ✚ PAR gaps are only permitted on the Y chromosome TPF.
- ✚ If the gap type is PAR, the beginning and ending accessions for this region must be listed in columns 3 and 4, respectively.
- ✚ If the gap type is PAR, columns 3 and 4 must both contain valid accessions that are found in column 1 on the chromosome X TPF.

CONTAINED COMPONENTS

Types of Contained Components

A component is considered "contained" when its entire sequence can be aligned to another component on the TPF (excluding any internal alignment gaps). A contained component MUST be marked on a TPF if it is to be included in the corresponding AGP. There are two designators that can be used to mark contained components on the TPF: CONTAINED and CONTAINED_TURNOUT. The designators differ in the switch point selection rules applied to the contained component during AGP production.

A. CONTAINED

TPF components marked with the "CONTAINED" designator will follow the default switching rules during AGP production. Examples for usage are shown.

Example A1: Single contained clone



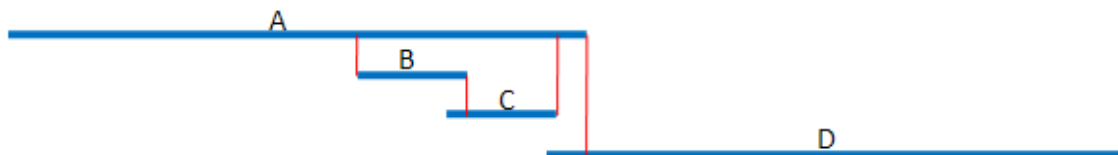
Example A2: Multiple non-overlapping contained clones



TPF shows:

A
B CONTAINED A
C CONTAINED A
D

Example A3: Multiple overlapping contained clones



TPF shows:

A
B CONTAINED A
C CONTAINED A
D

Note that the TPF mark-ups for examples A2 and A3 are identical. The software that produces the AGPs will determine the relationship between clones B and C and produce the appropriate switch points.

IMPORTANT: If the last clone in a chain of contained clones will switch back to the container, all clones in the chain should be marked CONTAINED.

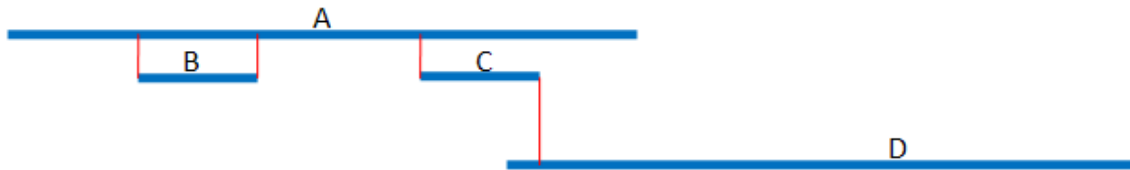
B. CONTAINED_TURNOUT

TPF components marked with the "CONTAINED_TURNOUT" designator will follow alternate switching rules during AGP production. Examples for usage are shown.

Example B1: Single contained clone

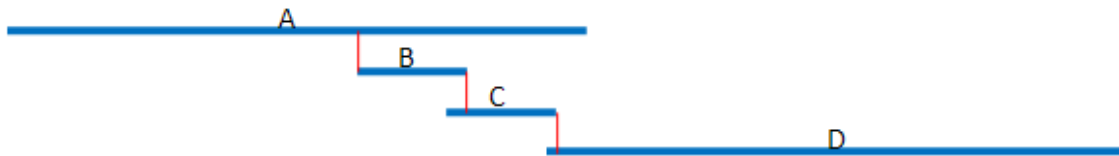


Example B2: Multiple non-overlapping contained clones



TPF shows:
 A
 B CONTAINED A
 C CONTAINED_TURNOUT A
 D

Example B3: Multiple overlapping contained clones



TPF shows:
 A
 B CONTAINED_TURNOUT A
 C CONTAINED_TURNOUT A
 D

IMPORTANT: In Example B3, note the use of CONTAINED_TURNOUT for both components, and contrast to Example A3. **If the last clone in a chain of contained clones will not switch back to the container, the entire chain should be marked as CONTAINED_TURNOUT.**

Additional Info for TPF Representation of Contained Components

- ✚ A component can only be contained by another component present in the same TPF contig
 - If column 4 = CONTAINED or CONTAINED_TURNOUT, the accession provided in column 5 must be present in the same local contig (column 3) as the accession in column 1.
- ✚ In situations of hierarchical containments, the TPF should always display the nearest contained relationship for any contained sequences.
 - Example:
 - A -----
 - B -----
 - C -----
 - TPF should report:
 - B contained in A
 - C contained in B
- ✚ In situations of hierarchical containments the order of the contained sequences on the TPF does not need to reflect the desired sequence order on the AGP.
- ✚ In situations where multiple sequences are contained within the same sequences (but are not hierarchical), the order of the contained sequences on the TPF does not need to reflect the desired sequence order on the AGP.

Specific TPF examples:

1. A TPF may contain one or more partial contigs. For example:

```
##ORGANISM: Mus musculus
##CHROMOSOME: 1
##ASSEMBLY NAME: NOD/MrkTac
##STRAIN/HAPLOTYPE/CULTIVAR: NOD/MrkTac
##TYPE: Contig
##VERSION: 1
##COMMENT: EXAMPLE TPF ONLY
##SUBMITTER: Schneider, NCBI
##CREATE DATE: Nov 3 2006 12:16PM
##UPDATE DATE: Nov 3 2006 12:16PM

##=== Beginning of TPF Data ===

AL645951      DN-29B18      NCBI_Mmchr1_ctg163678
AL596283      DN-257N2      NCBI_Mmchr1_ctg163678
CR936839      DN-120A16     NCBI_Mmchr1_ctg37199
CR936842      DN-189K17     NCBI_Mmchr1_ctg37199
#
AL671997      DN-396O20     NCBI_Mmchr1_ctg163206
AL672024      DN-378G17     NCBI_Mmchr1_ctg163206

##=== End of TPF Data ===
```

The absence of gap lines implies there is no known order for these contigs. Note that contigs may be separated by "#". This is not required and is only done to facilitate reading of the TPF by individuals.

2. A TPF may specify an entire chromosome. For example:

```
##ORGANISM: Mus musculus
##CHROMOSOME: 2
##ASSEMBLY NAME: Reference
##STRAIN/HAPLOTYPE/CULTIVAR: C57BL/6J
##TYPE: Complete Chromosome
##VERSION: 2
##COMMENT: EXAMPLE TPF ONLY
##SUBMITTER: Schneider, NCBI
##CREATE DATE: Oct 29 2007 04:39PM
##UPDATE DATE: Oct 29 2007 04:39PM

##=== Beginning of TPF Data ===

GAP  CENTROMERE      3000000
```

? WI1-1974M19 Mmchr2_ctg1
 ? WI1-923N3 Mmchr2_ctg1
 CU207330 WI1-2764C4 Mmchr2_ctg1
 AL928883 RP23-60E18 Mmchr2_ctg1
 AL732620 RP23-167G19 Mmchr2_ctg1
 AL929080 RP24-129J15 Mmchr2_ctg1
 AL732328 RP23-106P4 Mmchr2_ctg1
 AL844530 RP23-248L2 Mmchr2_ctg1
 CU181741 WI1-2132D7 Mmchr2_ctg1 CONTAINED AL845441 RP23-327I5
 AL845441 RP23-327I5 Mmchr2_ctg1
 AL935139 RP23-74F20 Mmchr2_ctg1
 AL954325 RP23-416H10 Mmchr2_ctg1
 AL928978 RP23-95M7 Mmchr2_ctg1 CONTAINED_TURNOUT AL954325 RP23-416H10
 AL928947 RP23-69N1 Mmchr2_ctg1 CONTAINED_TURNOUT AL954325 RP23-416H10
 AL807832 RP23-198D21 Mmchr2_ctg1
 AL928550 RP23-272I15 Mmchr2_ctg1
 AL807778 RP23-198G1 Mmchr2_ctg1
 BX682541 RP23-349P20 Mmchr2_ctg1
 AL928662 RP23-379M5 Mmchr2_ctg1
 AL732403 RP23-119L20 Mmchr2_ctg1
 BX323054 RP23-329P15 Mmchr2_ctg1
 AL928940 RP23-94J17 Mmchr2_ctg1
 AL929142 RP23-340A13 Mmchr2_ctg1
 AL928958 RP23-344N23 Mmchr2_ctg1
 AL928924 RP23-307E15 Mmchr2_ctg1
 CR388026 RP23-39L16 Mmchr2_ctg1
 AL928735 RP23-413M3 Mmchr2_ctg1
 BX842658 RP23-86F17 Mmchr2_ctg1
 AL845275 RP23-112F5 Mmchr2_ctg1
 AL845515 RP23-256D19 Mmchr2_ctg1
 AL845492 RP23-222P7 Mmchr2_ctg1
 AL929240 RP23-245A10 Mmchr2_ctg1
 AL772190 RP23-124P7 Mmchr2_ctg1
 CR936244 RP24-562B12 Mmchr2_ctg1
 CU207293 WI1-632G16 Mmchr2_ctg1
 AL845485 RP23-237D23 Mmchr2_ctg1
 BX649227 RP23-114C18 Mmchr2_ctg1
 AL840637 RP23-113H3 Mmchr2_ctg1
 AL845488 RP23-215P8 Mmchr2_ctg1
 AL928600 RP23-348O19 Mmchr2_ctg1
 AL929143 RP23-339G18 Mmchr2_ctg1
 AL773590 RP23-181A13 Mmchr2_ctg1
 BX005023 RP24-316N23 Mmchr2_ctg1
 AL928713 RP23-399M5 Mmchr2_ctg1
 AL929149 RP23-93B24 Mmchr2_ctg1

BX293551	RP23-107021	Mmchr2_ctg1	
AL929043	RP23-58H6	Mmchr2_ctg1	CONTAINED BX293551 RP23-107021
AL929187	RP23-8G6	Mmchr2_ctg1	CONTAINED AL929043 RP23-58H6
BX294115	RP24-465I11	Mmchr2_ctg1	
AL929440	RP23-292F11	Mmchr2_ctg1	
AL928665	RP23-294O23	Mmchr2_ctg1	
AL928832	RP23-261N18	Mmchr2_ctg1	
AL929194	RP23-385L14	Mmchr2_ctg1	
AL772377	RP23-141B15	Mmchr2_ctg1	
AL845529	RP23-276D17	Mmchr2_ctg1	
AL928704	RP23-393G10	Mmchr2_ctg1	
AL772367	RP23-119N4	Mmchr2_ctg1	
AL953853	RP23-6M16	Mmchr2_ctg1	
AL772216	RP23-153M22	Mmchr2_ctg1	
AL844485	RP23-436D21	Mmchr2_ctg1	
AL845264	RP23-116F18	Mmchr2_ctg1	
AL772352	RP23-147F6	Mmchr2_ctg1	
AL928715	RP23-373D16	Mmchr2_ctg1	
AL929020	RP23-291L24	Mmchr2_ctg1	
BX679665	RP23-247J17	Mmchr2_ctg1	
AL929179	RP23-428M4	Mmchr2_ctg1	
AL831794	RP23-114B13	Mmchr2_ctg1	
AL845548	RP23-307N14	Mmchr2_ctg1	
AL928795	RP23-38B1	Mmchr2_ctg1	
AL928909	RP23-320I18	Mmchr2_ctg1	
AL845313	RP23-202C2	Mmchr2_ctg1	
AL845533	RP23-232K16	Mmchr2_ctg1	
AL772342	RP23-105D19	Mmchr2_ctg1	
AL928641	RP23-272N6	Mmchr2_ctg1	
AL928560	RP23-353N23	Mmchr2_ctg1	
BX649225	RP23-104N16	Mmchr2_ctg1	
AL929209	RP23-415C3	Mmchr2_ctg1	
AL935271	RP23-336A16	Mmchr2_ctg1	
AL928807	RP23-97D20	Mmchr2_ctg1	
AL773538	RP23-14I24	Mmchr2_ctg1	
AL772303	RP23-185P20	Mmchr2_ctg1	
BX322642	RP24-363O9	Mmchr2_ctg1	
AL928918	RP23-403G13	Mmchr2_ctg1	
AL844560	RP23-334M9	Mmchr2_ctg1	
AL845434	RP23-211O3	Mmchr2_ctg1	
AL845290	RP23-201A19	Mmchr2_ctg1	
AL844558	RP23-378I2	Mmchr2_ctg1	
BX649230	RP23-157J23	Mmchr2_ctg1	
AL935312	RP23-56A7	Mmchr2_ctg1	
AL929165	RP23-446D4	Mmchr2_ctg1	
AL928632	RP23-303M15	Mmchr2_ctg1	
AL929268	RP23-379F6	Mmchr2_ctg1	

AL929158	RP23-92015	Mmchr2_ctg1
AL844166	RP23-107K21	Mmchr2_ctg1
AL845417	RP23-20I9	Mmchr2_ctg1
AL928841	RP23-95M4	Mmchr2_ctg1
AL844839	RP23-191F2	Mmchr2_ctg1
AL845543	RP23-257O6	Mmchr2_ctg1
AL935116	RP23-59I2	Mmchr2_ctg1
AL772224	RP23-131N18	Mmchr2_ctg1
AL845271	RP23-119O19	Mmchr2_ctg1
AL928888	RP23-390M20	Mmchr2_ctg1
AL844888	RP23-193F18	Mmchr2_ctg1
AL845520	RP23-283C20	Mmchr2_ctg1
AL928882	RP23-281I20	Mmchr2_ctg1
BX510346	RP23-188D15	Mmchr2_ctg1
AL929011	RP23-55O12	Mmchr2_ctg1
AL928545	RP23-218A13	Mmchr2_ctg1
AL935297	RP23-419K8	Mmchr2_ctg1
AL772218	RP23-129K21	Mmchr2_ctg1
AL845498	RP23-222D20	Mmchr2_ctg1
AL954131	RP23-32H12	Mmchr2_ctg1
AL928589	RP23-350C1	Mmchr2_ctg1
AL928557	RP23-349H7	Mmchr2_ctg1
AL928620	RP23-319M16	Mmchr2_ctg1
AL845265	RP23-158O8	Mmchr2_ctg1
AL928904	RP23-463M17	Mmchr2_ctg1
AL928680	RP23-396N6	Mmchr2_ctg1
AL928653	RP23-410F9	Mmchr2_ctg1
AL844855	RP23-113K9	Mmchr2_ctg1
AL929034	RP23-90F9	Mmchr2_ctg1
AL928806	RP23-34E4	Mmchr2_ctg1
BX649213	RP24-189E15	Mmchr2_ctg1
AL928877	RP23-52D18	Mmchr2_ctg1
AL928860	RP23-5I15	Mmchr2_ctg1
BX571892	RP23-207B21	Mmchr2_ctg1
AL928572	RP23-333P17	Mmchr2_ctg1
AL845528	RP23-25P15	Mmchr2_ctg1
BX276179	RP23-204B24	Mmchr2_ctg1
AL844538	RP23-289K19	Mmchr2_ctg1
BX649224	RP24-555L22	Mmchr2_ctg1
AL929100	RP23-436G18	Mmchr2_ctg1
BX649226	RP23-105O21	Mmchr2_ctg1
AL773540	RP23-177L19	Mmchr2_ctg1
AL929117	RP23-442K5	Mmchr2_ctg1
AL928939	RP23-294B13	Mmchr2_ctg1
AL929311	RP23-291N2	Mmchr2_ctg1
BX294442	RP23-310O19	Mmchr2_ctg1
AL772387	RP23-133B16	Mmchr2_ctg1

AL805928	RP23-136C12	Mmchr2_ctg1
AL929257	RP23-75A23	Mmchr2_ctg1
AL929064	RP23-442A7	Mmchr2_ctg1
AL929036	RP24-548P4	Mmchr2_ctg1
GAP	TYPE-3 20000	ALIGNMENT
AL928693	RP23-407K8	Mmchr2_ctg2
BX649461	RP23-114M13	Mmchr2_ctg2
CT030013	RP23-282P19	Mmchr2_ctg2
AL845257	RP23-280K4	Mmchr2_ctg2
AL928706	RP23-54E16	Mmchr2_ctg2
CR847856	RP23-447G16	Mmchr2_ctg2
AL773534	RP23-183O12	Mmchr2_ctg2
AL928690	RP23-443L19	Mmchr2_ctg2
AL844844	RP23-103D7	Mmchr2_ctg2
AL844838	RP23-263E15	Mmchr2_ctg2
BX005038	RP24-550O4	Mmchr2_ctg2
AL732430	RP23-176J12	Mmchr2_ctg2
AL732528	RP23-218B17	Mmchr2_ctg2
AL732546	RP23-264D16	Mmchr2_ctg2
BX294112	RP24-316M2	Mmchr2_ctg2
AL732525	RP23-358I10	Mmchr2_ctg2
AL935039	RP24-144B2	Mmchr2_ctg2
AL732585	RP23-226M2	Mmchr2_ctg2
AL732309	RP23-132N23	Mmchr2_ctg2
AL732557	RP23-47P18	Mmchr2_ctg2
AL732590	RP23-464C2	Mmchr2_ctg2
BX649340	RP23-225D24	Mmchr2_ctg2
AL731682	RP23-123F7	Mmchr2_ctg2
AL845455	RP23-325E4	Mmchr2_ctg2
AL773595	RP23-297I16	Mmchr2_ctg2
AL732541	RP23-306D20	Mmchr2_ctg2
BX649422	RP23-70A22	Mmchr2_ctg2
AL732311	RP23-125H15	Mmchr2_ctg2
AL773563	RP23-414L19	Mmchr2_ctg2
AL845266	RP23-449M10	Mmchr2_ctg2
AL954801	RP23-171K6	Mmchr2_ctg2
BX649202	RP24-157P15	Mmchr2_ctg2
AL731552	RP23-113K24	Mmchr2_ctg2
AL772282	RP23-328F3	Mmchr2_ctg2
AL772166	RP23-413I12	Mmchr2_ctg2
AL928953	RP24-83K6	Mmchr2_ctg2
AL732513	RP23-344B24	Mmchr2_ctg2
AL731778	RP23-475B13	Mmchr2_ctg2
AL732616	RP23-466D3	Mmchr2_ctg2
AL772249	RP23-430H1	Mmchr2_ctg2
AL928994	RP24-118E9	Mmchr2_ctg2
AL731851	RP23-362N19	Mmchr2_ctg2

AL732526	RP23-338O4	Mmchr2_ctg2
AL845267	RP23-426D2	Mmchr2_ctg2
AL772379	RP23-170D24	Mmchr2_ctg2
AL772213	RP23-104G16	Mmchr2_ctg2
AL929437	RP23-74P16	Mmchr2_ctg2
AL845323	RP23-304D11	Mmchr2_ctg2
BX294378	RP23-457B12	Mmchr2_ctg2
AL928926	RP23-443G7	Mmchr2_ctg2
BX005298	RP23-261D24	Mmchr2_ctg2
AL845258	RP23-315H12	Mmchr2_ctg2
AL954388	RP23-395P6	Mmchr2_ctg2
AL954299	RP23-399D3	Mmchr2_ctg2
AL928593	RP23-100E9	Mmchr2_ctg2
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