

pubmedKB API

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Li Peng-Hsuan (朋軒)

KB Overview

| | | pubmedKB-BERN | pubmedKB-PTC | pubmedKB-PTC-FT |
|------------|------------------------|---------------|--------------|-------------------|
| Literature | Source | 33M PubMed | 32M PubMed | 2M PubMed Central |
| | Type | Abstract | Abstract | Full text |
| | # PMIDs | 4,348,135 | 10,772,202 | 1,674,383 |
| Entity | # types | 4 | 13 | 14 |
| | # unique (type, name)s | 1,772,821 | 2,718,110 | 2,530,623 |
| | # unique (type, ID)s | 289,629 | 598,074 | 648,891 |
| Relation | # annotators | 4 | 4 | 4 |
| | # annotations | 9,094,498 | 34,393,039 | 41,051,562 |

Entity Types

| pubmedKB-BERN | pubmedKB-PTC | pubmedKB-PTC-FT |
|---------------|-------------------|-------------------|
| gene | Gene | Gene |
| variant | ProteinMutation | ProteinMutation |
| | DNAMutation | DNAMutation |
| | SNP | SNP |
| | CopyNumberVariant | CopyNumberVariant |
| disease | Disease | Disease |
| chemical | Chemical | Chemical |
| - | Species | Species |
| - | CellLine | CellLine |
| - | RefSeq | RefSeq |
| - | GenomicRegion | GenomicRegion |
| - | Chromosome | Chromosome |
| - | Genus | - |
| - | - | DNAAcidChange |
| - | - | DomainMotif |

Entity ID Systems

| pubmedKB-BERN |
|---------------|
| BERN |
| MESH |
| CHEBI |
| OMIM |
| HGNC |
| MIM |
| Ensembl |
| IMGT/GENE-DB |
| miRBase |

| pubmedKB-PTC |
|----------------------|
| MESH |
| NCBITaxID |
| tmVar |
| VariantGroup |
| CorrespondingGene |
| HGVS |
| RS# |
| CA# |
| OMIM |
| CVCL |
| CorrespondingSpecies |
| OriginalGene |
| Chr1~23 |
| ChrX |

| pubmedKB-PTC-FT |
|------------------------|
| MESH |
| CVCL |
| tmVar |
| VariantGroup |
| CorrespondingGene |
| HGVS |
| RS# |
| CA# |
| OMIM |
| CorrespondingSpecies |
| OriginalGene |
| Focus |
| Right |
| Left |
| Chr0~9,01~39,65,88,103 |
| ChrX,Y,XX |

Entity ID Search - Web API

NEN Query

Case sensitive (Y/N): Max length difference: Min similarity: Max #name per query: Max #alias per ID:

Query:

show

Ready

Name: V600E Similarity: 100%

| Type | ID | Frequency of (Type, ID, Name) in all texts | Top 5 aliases by frequency of (Type, ID, Alias) in all texts |
|-----------------|---------------------------|--|--|
| ProteinMutation | tmVar:p SUB V 600 E | 12,046 | V600E (12,046) BRAFV600E (4,433) p.V600E (331) BrafV600E (97) p.Val600Glu (41) |
| ProteinMutation | HGVS:p.V600E | 12,046 | V600E (12,046) BRAFV600E (4,433) p.V600E (331) BrafV600E (97) p.Val600Glu (41) |
| ProteinMutation | CorrespondingGene:673 | 11,801 | V600E (11,801) BRAFV600E (4,426) V600 (339) p.V600E (327) V600K (253) |
| ProteinMutation | RS#:113488022 | 11,617 | V600E (11,617) BRAFV600E (4,433) p.V600E (283) V600 (274) V600E/K (159) |
| ProteinMutation | CA#:123643 | 11,617 | V600E (11,617) BRAFV600E (4,412) p.V600E (283) BrafV600E (97) p.Val600Glu (38) |
| ProteinMutation | VariantGroup:0 | 11,460 | V600E (11,460) V617F (8,378) T790M (5,599) BRAFV600E (4,395) C282Y (3,132) |
| ProteinMutation | VariantGroup:1 | 441 | C282Y (1,209) H63D (1,184) T790M (1,145) Val/Val (812) L858R (796) |
| ProteinMutation | RS#:121913227 | 158 | V600E (158) V600K (128) V600 (59) p.V600E (44) V600R (43) |
| ProteinMutation | CorrespondingSpecies:9606 | 110 | T790M (123) V600E (110) BRAFV600E (93) D614G (86) G12C (60) |
| ProteinMutation | VariantGroup:2 | 103 | Val/Val (366) Met/Met (264) Arg/Arg (243) H63D (227) C282Y (221) |
| ProteinMutation | VariantGroup:3 | 25 | Arg/Arg (125) K103N (77) Y181C (73) Gln/Gln (71) M184V (70) |
| ProteinMutation | CorrespondingGene:109880 | 17 | V600E (17) |
| ProteinMutation | CorrespondingGene:5594 | 16 | K52R (19) E322K (17) serine/threonine (17) S6 (17) V600E (16) |
| ProteinMutation | RS#:121913378 | 14 | V600E (14) V600M (6) V600L (4) V600 (4) V600R (3) |
| ProteinMutation | VariantGroup:4 | 13 | K103N (65) Gln/Gln (51) M184V (45) L180M (41) Arg/Arg (40) |
| ProteinMutation | CorrespondingGene:3845 | 12 | G12D (1,783) G12V (686) G13D (346) p.G12C (106) p.G13D (104) |
| ProteinMutation | CorrespondingGene:7015 | 11 | V600E (11) p-E1A (5) R132H (4) K626A (4) K27M (3) |
| ProteinMutation | CorrespondingGene:22882 | 11 | V600E (11) serine/threonine (9) A85K (5) N71R (2) S621A (2) |

Relation Annotators

| Annotator | Annotation | pubmedKB-BERN | pubmedKB-PTC | pubmedKB-PTC-FT |
|------------|-----------------|---------------|--------------|-----------------|
| odds_ratio | OR, CI, p-value | V | V | V |
| rbert_cre | Relation | V | V | V |
| spacy_ore | Triplet | V | V | V |
| openie_ore | Triplet | V | V | V |

Relation Annotator - Details

| Annotator | Annotation | Explanation | |
|------------|-----------------|------------------------------|---|
| odds_ratio | OR, CI, p-value | OR | Odds ratio extracted from the sentence |
| | | CI | Confidence interval extracted from the sentence |
| | | p-value | p-value statistics extracted from the sentence |
| rbert_cre | Relation | Relation: Cause-associated | VARIANT is a causal variant of DISEASE; or VARIANT is DISEASE-associated |
| | | Relation: Appositive | VARIANT's association with DISEASE is presupposed |
| | | Relation: In-patient | VARIANT occurs in some DISEASE patients |
| spacy_ore | Triplet | (subject, predicate, object) | The combined clause is entailed by the sentence |
| openie_ore | Triplet | (subject, predicate, object) | The combined clause is entailed by the sentence |

Relation Annotation - Web API

REL Query

Entity A

Query by: Type & Name ▾

Type: ProteinMutation ▾

ID: tmVar:p|SUB|V|600|E

Name: V600E

Entity B

Query by: Type & ID ▾

Type: Disease ▾

ID: MESH:D009369

Name: neoplasm

Query

Query by: Un-ordered pair (A, B) ▾

Max #rel per annotator: 20

PMID: 22494505 (ignored if query by entity or entity pair)

show

Ready

| Head | Tail | Annotator | Annotation | PMID | Sentence |
|--|---|------------|--|--------------------------|---|
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | tumors [Disease] MESH:D009369 | odds_ratio | OR: 1.87, CI: 1.01-3.47, p-value: x | 20200438 | ABSTRACT paragraph <i>The prevalence of BRAF V600E was 5.0% in this population, and the mutation was more likely to be found in tumors from cases who were of Ashkenazi Jewish descent [odds ratio (OR), 1.87; 95% confidence interval (95% CI), 1.01-3.47], female (OR, 1.97; P = 1.17-3.31), and older (73.8 years versus 70.3 years; P < 0.001).</i> |
| V600E [ProteinMutation] HGVS:p.V600E tmVar:p SUB V 600 E | cancers [Disease] MESH:D009369 | rbert_cre | Cause-associated: 87.5% | 35028596 | ABSTRACT paragraph <i>The constitutive expression of wild-type BRAF (BRAFWT) and its mutant forms, especially V600E (BRAFV600E), has been linked to multiple cancers.</i> |
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | cancer [Disease] MESH:D009369 | spacy_ore | B-RafV600E, promotes, cancer cell behavior | 22751131 | ABSTRACT paragraph <i>The most prevalent mutation, B-RafV600E, promotes cancer cell behavior through mechanisms that are still incompletely defined.</i> |
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | tumors [Disease] MESH:D009369 | openie_ore | V600E mutation, was detected in, 25 of 27 tumors | 23682579 | ABSTRACT paragraph <i>The BRAF(V600E) mutation was detected in 25 of 27 tumors (92.6%).</i> |

| Head | Tail | Annotator | Annotation | PMID | Sentence |
|--|---|------------|--|--------------------------|---|
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | tumours [Disease] MESH:D009369 | odds_ratio | OR: 12.33, CI: 4.21-41.54, p-value: 1.8x10-7 | 21625944 | RESULTS paragraph We observed a significant association of both the BRAF V600E mutation and MLH1 methylation with CIMP+ tumours (odds ratio (OR) = 12.33, 95% confidence interval (CI) = 4.21-41.54, P = 1.8 x 10-7 and OR = 15.5, 95% CI = 4.54-68.65, P = 2.8 x 10-7, respectively). |
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | tumor [Disease] MESH:D009369 | rbert_cre | In-patient: 75.7% | 23043499 | CASE paragraph Mutation analysis of the patient's tumor material demonstrated the presence of a BRAFV600E mutation and the patient subsequently initiated treatment with a BRAF inhibitor in September 2011. |
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | tumor [Disease] MESH:D009369 | spacy_ore | the BRAFV600E inhibitor, not affect, tumor eradication | 22735805 | DISCUSS paragraph Furthermore, using a tumor model in which tumor eradication is dependent on the activation of tumor antigen specific CD4+ and CD8+ cells and their production of both IFN-gamma and IL-2 demonstrated that the BRAFV600E inhibitor did not affect tumor eradication. |
| V600E [ProteinMutation] CA#:123643 CorrespondingGene:673 HGVS:p.V600E RS#:113488022 VariantGroup:0 tmVar:p SUB V 600 E | cancer [Disease] MESH:D009369 | openie_ore | crVAR V600E, is shared by, 17 cancer types | 27977206 | RESULTS paragraph The crVAR V600E in BRAF protein (ENSP00000288602), which is a therapeutic target of metastatic melanoma, is shared by 17 cancer types. |

API Access

- Source code (programmable API + demo website)
 - https://github.com/jacobvsdaniel/pubmedkb_web
 - Document: https://github.com/jacobvsdaniel/pubmedkb_web/wiki
- Sample dataset (1/10 of pubmedKB-ptc)
 - https://drive.google.com/file/d/10_UmG_ozWSrvFB9vJ0TfY41WHCzswmhm/view?usp=sharing