

Entity	Relative Entropy	Drug Target	Entity Count	Links
PPARGC1B	10.4308		6	 
EP300	10.0487		3	 
PPARGC1A	8.9456		6	 
BRCA1	6.4020		1	 
1 ESR1	4.7803	Yes	3	 

1. Associations of genetic variants in the estrogen receptor coactivators PPARGC1B and EP300 with familial breast cancer.

PubMed 16704985 Authors: Michael Wirtenberger, Sandrine Tchatchou, Kari Hemminki, Julia Schmutzhard, Christian Alfons Meindl, Barbara Wappenschmidt, Marion Kiechle, Norbert Arnold, Bernhard H F Weber, Dieter Niederacher, Claus R Barti 2006-11. Journal: Carcinogenesis Affiliation: Division of Molecular Genetic Epidemiology, Helmholtz-University Group Molecular Cancer Research Center (DKFZ) Heidelberg, Germany. m.wirtenberger@dkfz.de

[Statistics](#)

The mitogen effect of the ovarian steroid estrogen is a strong risk factor for breast cancer development. This effect is mainly mediated by the estrogen receptor alpha, a hormone inducible transcription factor, which activates gene expression through recruiting multiple coactivators including PPARGC1B and EP300. We tested the hypothesis that non-conservative, putative functional amino acid exchanges in PPARGC1B and EP300 act as low-penetrance familial breast cancer risk factors. The analysis of 816 BRCA1/2 mutation-negative familial breast cancer controls revealed an association of the PPARGC1A Thr612Met polymorphism with familial breast cancer (OR = 1.35, 95% CI 1.08-1.68) and high-risk familial breast cancer (OR = 1.51, 95% CI 1.08-2.12, P = 0.017) and bilateral familial breast cancer (OR = 2.30, 95% CI 1.15-4.61, P = 0.020). Logistic regression analyses of the PPARGC1B Ala203Pro variant showed an increased familial breast cancer risk of heterozygote variant allele carriers (OR = 1.48, 95% CI 1.15-1.91, P = 0.002). The genotype-combination analysis of the associated PPARGC1A Thr612Met and PPARGC1B Ala203Pro variant suggests an allele dose-dependent breast cancer risk (P(trend) = 0.0004). Our findings highlight the importance of inherited variants in the estrogen receptor coactivator genes PPARGC1A and PPARGC1B for familial breast cancer. Regarding their impact on estrogen signaling, these polymorphisms might also influence adjuvant anti-estrogen therapy, using a personalized approach, and outcome of breast cancer patients.

- 1 Found entities could be indexed, ranked and linked to other data.
- 2 Highlighting in text corresponding to the entity classes.

ProMiner®: RECOGNITION AND NORMALIZATION OF NAMED ENTITIES IN SCIENTIFIC TEXT

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Challenges

Scientific publications found in abstract databases, full text journals or patents are the main and most up-to-date information source, but the amount of text is overwhelming for most Life Science areas.

Recognition of Life Science terminology is a key prerequisite for performing automatic information retrieval and information extraction. Huge and complex terminologies with high numbers of synonymous expressions, ambiguous terminology and numerous generations of new names and classes present named entity recognition with a real challenge.

ProMiner is a tool for specific terminology recognition and addresses several fundamental issues in name entity recognition in the field of Life Sciences:

- ProMiner can handle voluminous dictionaries, complex thesauri and large controlled vocabularies derived from ontologies
- Regularly updated dictionaries through automatic curation followed by a manual evaluation process
- Mapping of synonyms to reference names and data sources
- Context dependent disambiguation of biomedical termini and resolution of acronyms
- Specific handling of common English word synonyms
- Spelling variants of expressions in the source dictionary can be recognized
- High speed tagging and parallel workflow for multiple dictionaries
- Incorporation of regular expressions (e.g. for the recognition of SNP rs numbers)
- Full text annotation in XML, HTML or PDF format
- Patent annotation

BioCreAtIvE evaluation

Organism (evaluation)	ProMiner®	Best performance
Mouse (BioCreAtIvE04)	0,79	0,79
Fly (BioCreAtIvE04)	0,82	0,82
Yeast (BioCreAtIvE04)	0,90	0,92
Human (BioCreAtIvE07)	0,80	0,81

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Textual information

