



## SFTPC gene

surfactant protein C

### Normal Function

The *SFTPC* gene provides instructions for making a protein called surfactant protein-C (SP-C). This protein is one of four proteins (each produced from a different gene) in surfactant, a mixture of certain fats (called phospholipids) and proteins that lines the lung tissue and makes breathing easy. Without normal surfactant, the tissue surrounding the air sacs in the lungs (the alveoli) sticks together after exhalation (because of a force called surface tension), causing the alveoli to collapse. As a result, filling the lungs with air on each breath becomes very difficult, and the delivery of oxygen to the body is impaired. Surfactant lowers surface tension, easing breathing and avoiding lung collapse. The SP-C protein helps spread the surfactant across the surface of the lung tissue, aiding in the surface tension-lowering property of surfactant.

The phospholipids and proteins that make up surfactant are packaged in cellular structures known as lamellar bodies, which are found in specialized lung cells. The surfactant proteins must go through several processing steps to mature and become functional; some of these steps occur in lamellar bodies.

### Health Conditions Related to Genetic Changes

#### Surfactant dysfunction

More than 35 mutations in the *SFTPC* gene have been identified in people with surfactant dysfunction. When this condition is caused by mutations in the *SFTPC* gene (sometimes called SP-C dysfunction), it can cause severe breathing problems in newborns or gradual onset of milder breathing problems in children or adults.

*SFTPC* gene mutations associated with surfactant dysfunction affect the processing of the SP-C protein. Many of the mutations occur in a particular region of the gene called the BRICHOS domain, which appears to be involved in the processing and cellular placement of the SP-C protein.

Mutations in the *SFTPC* gene result in a reduction or absence of mature SP-C and a buildup of abnormal forms of SP-C. It is unclear which of these outcomes causes the signs and symptoms of SP-C dysfunction. Lack of mature SP-C can lead to abnormal composition of surfactant and decreased surfactant function. The loss of functional surfactant would raise surface tension in the alveoli, causing difficulty breathing and collapse of the lungs. Alternatively, research suggests that abnormally processed SP-C proteins form the wrong three-dimensional shape and accumulate inside lung cells. These misfolded proteins may trigger a cellular response that results in cell damage

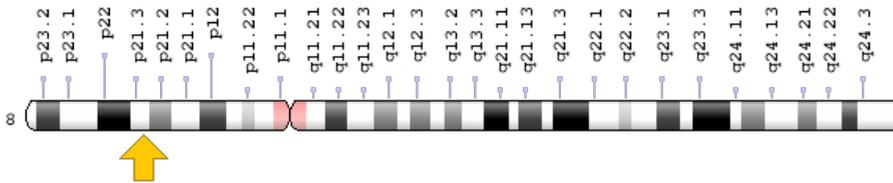
and death. This damage may disrupt surfactant production and release, leading to the breathing problems associated with surfactant dysfunction.

### Idiopathic pulmonary fibrosis

## **Chromosomal Location**

Cytogenetic Location: 8p21.3, which is the short (p) arm of chromosome 8 at position 21.3

Molecular Location: base pairs 22,161,671 to 22,164,479 on chromosome 8 (Homo sapiens Annotation Release 109, GRCh38.p12) (NCBI)



Credit: Genome Decoration Page/NCBI

## **Other Names for This Gene**

- BRICD6
- PSP-C
- PSPC\_HUMAN
- pulmonary surfactant apoprotein-2 SP-C
- pulmonary surfactant-associated protein C
- pulmonary surfactant-associated proteolipid SPL(Val)
- SFTP2
- SMDP2
- SP-C
- SP5

## **Additional Information & Resources**

### Educational Resources

- Molecular Biology of the Cell (fourth edition, 2002): Adjacent Cell Types Collaborate in the Alveoli of the Lungs  
<https://www.ncbi.nlm.nih.gov/books/NBK26875/#A4114>

### Clinical Information from GeneReviews

- Pulmonary Fibrosis, Familial  
<https://www.ncbi.nlm.nih.gov/books/NBK1230>

### Scientific Articles on PubMed

- PubMed  
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28SFTPC%5BTIAB%5D%29+OR+%28surfactant+protein+C%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D>

### Catalog of Genes and Diseases from OMIM

- PULMONARY FIBROSIS, IDIOPATHIC  
<http://omim.org/entry/178500>
- SURFACTANT, PULMONARY-ASSOCIATED PROTEIN C  
<http://omim.org/entry/178620>

### Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology  
[http://atlasgeneticsoncology.org/Genes/GC\\_SFTPC.html](http://atlasgeneticsoncology.org/Genes/GC_SFTPC.html)
- ClinVar  
<https://www.ncbi.nlm.nih.gov/clinvar?term=SFTPC%5Bgene%5D>
- HGNC Gene Symbol Report  
[https://www.genenames.org/data/gene-symbol-report#!/hgnc\\_id/HGNC:10802](https://www.genenames.org/data/gene-symbol-report#!/hgnc_id/HGNC:10802)
- Monarch Initiative  
<https://monarchinitiative.org/gene/NCBIGene:6440>
- NCBI Gene  
<https://www.ncbi.nlm.nih.gov/gene/6440>
- UniProt  
<https://www.uniprot.org/uniprot/P11686>

## Sources for This Summary

- Guillot L, Epaud R, Thouvenin G, Jonard L, Mohsni A, Couderc R, Counil F, de Blic J, Taam RA, Le Bourgeois M, Reix P, Flamein F, Clement A, Feldmann D. New surfactant protein C gene mutations associated with diffuse lung disease. *J Med Genet.* 2009 Jul;46(7):490-4. doi: 10.1136/jmg.2009.066829. Epub 2009 May 13. Erratum in: *J Med Genet.* 2010 Jul;47(7):485.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/19443464>
  - Hamvas A, Nogee LM, White FV, Schuler P, Hackett BP, Huddleston CB, Mendeloff EN, Hsu FF, Wert SE, Gonzales LW, Beers MF, Ballard PL. Progressive lung disease and surfactant dysfunction with a deletion in surfactant protein C gene. *Am J Respir Cell Mol Biol.* 2004 Jun;30(6):771-6. Epub 2003 Dec 4.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/14656744>
  - Mulugeta S, Maguire JA, Newitt JL, Russo SJ, Kotorashvili A, Beers MF. Misfolded BRICHOS SP-C mutant proteins induce apoptosis via caspase-4- and cytochrome c-related mechanisms. *Am J Physiol Lung Cell Mol Physiol.* 2007 Sep;293(3):L720-9. Epub 2007 Jun 22.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/17586700>
  - OMIM: SURFACTANT, PULMONARY-ASSOCIATED PROTEIN C  
<http://omim.org/entry/178620>
  - Weaver TE. Synthesis, processing and secretion of surfactant proteins B and C. *Biochim Biophys Acta.* 1998 Nov 19;1408(2-3):173-9. Review.  
*Citation on PubMed:* <https://www.ncbi.nlm.nih.gov/pubmed/9813310>
- 

Reprinted from Genetics Home Reference:

<https://ghr.nlm.nih.gov/gene/SFTPC>

Reviewed: July 2012

Published: May 14, 2019

Lister Hill National Center for Biomedical Communications  
U.S. National Library of Medicine  
National Institutes of Health  
Department of Health & Human Services