# DISEASE:
# Xeroderma pigmentosum-Cockayne syndrome complex

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| **NAME:** | Xeroderma pigmentosum-Cockayne syndrome complex |
| **DESCRIPTION:** | Xeroderma pigmentosum/Cockayne syndrome complex (XP/CS complex) is characterized by the cutaneous features of xeroderma pigmentosum (XP) (see this term) together with the systemic and neurological features of Cockayne syndrome (CS; see this term). |
| **ORPHACODE:** | 220295 |
| **SYNONYMS:** | XP/CS complex |
| **XREF(S):** | Orphanet<br>OMIM<br>OMIM<br>OMIM<br>OMIM |
| **ANALYTE(S):** | ERCC5<br>ERCC4<br>ERCC2<br>ERCC3 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Ichthyosis (gene panel)

## Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

## Related Analytes

- ERCC excision repair 2, TFIIH core complex helicase subunit
- ERCC excision repair 3, TFIIH core complex helicase subunit
- ERCC excision repair 4, endonuclease catalytic subunit
- ERCC excision repair 5, endonuclease

## Related Gene Panels

- Ichthyosis and erythroderma (98 genes) - KUL