# DISEASE:
# Posterior polymorphous corneal dystrophy

| | |
|---|---|
| **NAME:** | Posterior polymorphous corneal dystrophy |
| **DESCRIPTION:** | A rare mild subtype of posterior corneal dystrophy characterized by small aggregates of apparent vesicles bordered by a gray haze at the level of Descemet membrane, generally with no effect on vision. |
| **ORPHACODE:** | 98973 |
| **SYNONYMS:** | PPCD<br>Posterior polymorphous dystrophy<br>Schlichting dystrophy |
| **XREF(S):** | Orphanet<br>OMIM<br>OMIM<br>ICD-10<br>OMIM |
| **ANALYTE(S):** | VSX1<br>ZEB1<br>COL8A2<br>OVOL2<br>GRHL2 |
| **CREATED:** | 13 May 2019 - 01:02 |

| | |
|---|---|
| **CHANGED:** | 22 Jun 2023 - 16:14 |

---

Source URL: http://gentest.healthdata.be/disease/3619

# RELATED CONTENT

## Related Genetic Tests

- Corneal dystrophy (gene panel)

## Related Laboratories

- Centrum Medische Genetica - UZ Gent

## Related Analytes

- collagen type VIII alpha 2 chain
- grainyhead like transcription factor 2
- ovo like zinc finger 2
- visual system homeobox 1
- zinc finger E-box binding homeobox 1

## Related Gene Panels

- Corneal dystrophy - UGent