# DISEASE:
# PTEN hamartoma tumor syndrome

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| **NAME:** | PTEN hamartoma tumor syndrome |
| **DESCRIPTION:** | PTEN hamartoma tumor syndrome (PHTS) is a term defining a group of clinically heterogeneous disorders united by a germline *PTEN* mutation and the involvement of derivatives of all 3 germ cell layers, manifesting with hamartomas, overgrowth and neoplasia. Currently, subsets carrying clinical diagnoses of Cowden syndrome, Bannayan-Riley-Ruvalcaba syndrome, Proteus and Proteus-like syndromes and SOLAMEN syndrome (see these terms) belong to PHTS. |
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# RELATED CONTENT

## Related Genetic Tests

- Hereditary cancer panel (gene panel)
- PTEN hamartoma tumor syndrome

## Related Laboratories

- Centre de Génétique Médicale UCL

## Related Gene Panels

- Hereditary Cancer Solution (35 genes) - UCL