# DISEASE:
# Acquired partial lipodystrophy

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| **NAME:** | Acquired partial lipodystrophy |
| **DESCRIPTION:** | A rare acquired lipodystrophy characterized by bilateral, symmetrical lipoatrophy of the upper body (face, neck, arms, thorax and sometimes upper abdomen) with sparing of the lower extremities and cephalothoracic progression. The disease may be associated with low serum levels of C3 and presence of C3-nephritic factor. |
| **ORPHACODE:** | 79087 |
| **SYNONYMS:** | Barraquer-Simons syndrome<br>Progressive cephalothoracic lipodystrophy |
| **XREF(S):** | Orphanet<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | LMNB2 |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

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### Related Analytes

- lamin B2

Source URL: http://gentest.healthdata.be/disease/2881