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| <b>POLICY NAME</b> | Lumizyme (alglucosidase) | <b>POLICY #</b> | <b>2477P</b> |
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## Criteria

### Coverage Criteria for the Treatment of Pompe disease

- ☐ **1.1** Diagnosis of Pompe disease, supported by the following:
  - i Enzyme assay showing a deficiency of acid alpha-glucosidase (GAA) activity in the blood, skin, or muscle
  - ii Genetic testing showing a mutation in the GAA gene
- ☐ **1.2** Age 1 year or older
- ☐ **1.3** Prescribed by a geneticist (gene specialist) or specialist in Pompe disease
- ☐ **1.4** Documentation and imaging to rule out presence of an enlarged heart (cardiomyopathy)
- ☐ **1.5** Documentation showing baseline percent-predicted forced vital capacity (FVC) and 6-minute walk test (6MWT)
- ☐ **1.6** Review of chart notes documenting diagnosis and confirming that patient has met all above requirements for treatment with Nexviazyme by both a pharmacist and medical director

### Exclusion Criteria – Any of the following prevents coverage

- ☐ **2.1** Use along with Nexviazyme is considered a duplication and is excluded from coverage.