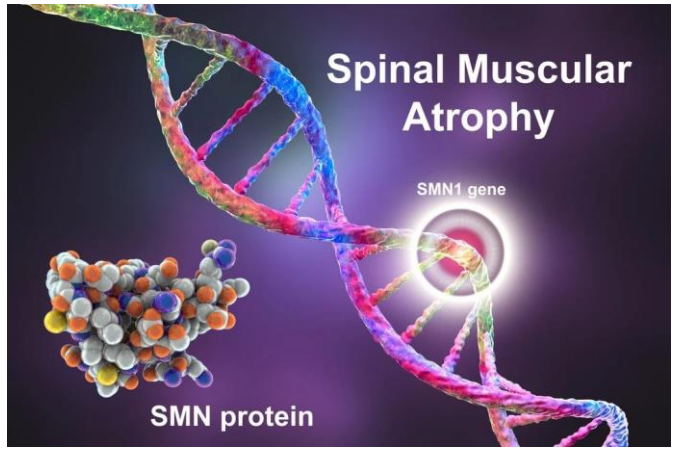
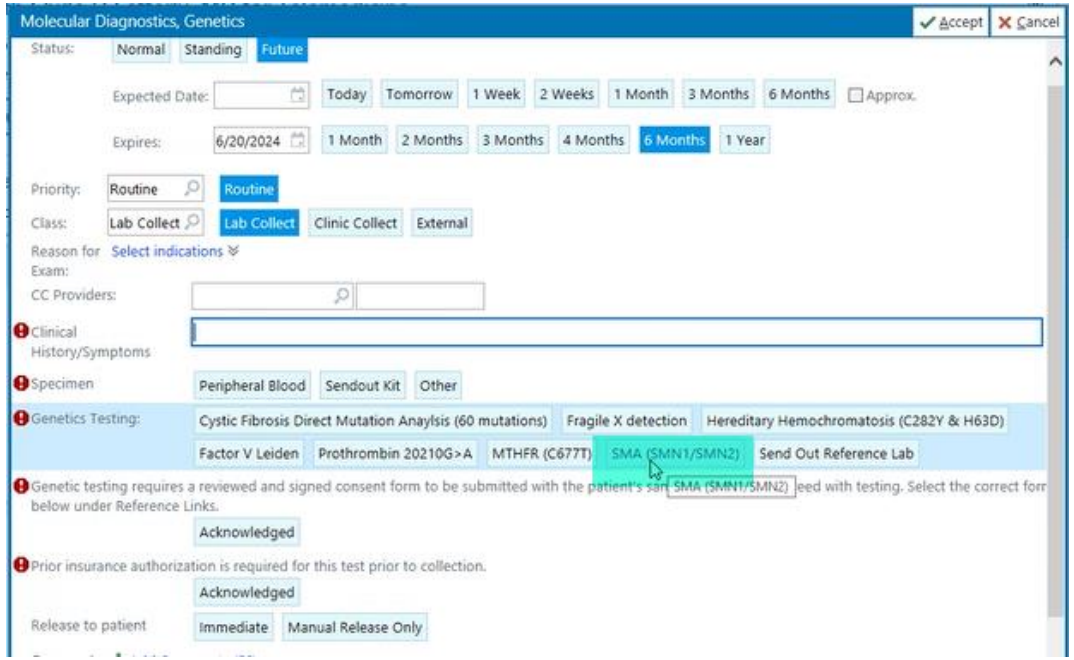
Spinal Muscular Atrophy Genetic Testing In-house

UPSTATE UNIVERSITY HEALTH SYSTEM

- Mutations in SMA (SMN1/SMN2) are one genetic cause of Spinal Muscular Atrophy
- Testing for these mutations can be done for both diagnosis and carrier status
- Currently this testing is a send-out test requiring significant turn-around time





- Starting **February 5, 2024**, this testing will be performed in-house at Upstate's Molecular Laboratory
- Testing for **SMA (SMN1/SMN2)** can be ordered using **LABAP2706** and select the correct test (as shown to the left
- The correct consent for this genetic testing must be completed prior to requesting this assay
- Insurance authorization also must be initiated prior to requesting this assay

For any questions, please contact the Molecular Laboratory (315-464-6806)