## **SUPPLEMENTARY MATERIAL**

RTB Lectin: a novel receptor-independent delivery system for lysosomal enzyme replacement therapies

Walter Acosta<sup>1,4</sup>, Jorge Ayala<sup>1,3</sup>, Maureen C. Dolan<sup>1,2</sup> & Carole L. Cramer <sup>1-3</sup>

<sup>1</sup>Arkansas Biosciences Institute at Arkansas State University-Jonesboro, PO Box 639, State University, AR 72467, USA.

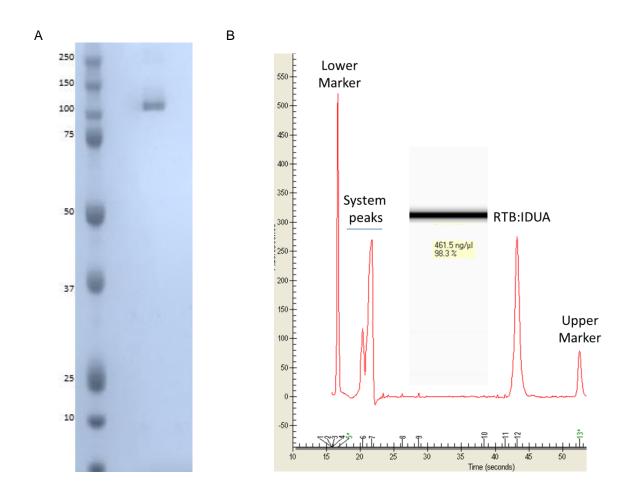
<sup>2</sup>Department of Biological Sciences, Arkansas State University-Jonesboro, State University AR 72467, USA.

<sup>3</sup>BioStrategies LC, PO Box 2428, State University, AR 72467, USA.

<sup>4</sup>Present address: BioStrategies LC, PO Box 2428, State University, AR 72467, USA.

Correspondence should be addressed to C.L.C. (ccramer@astate.edu).

**Supplementary Figure S1. RTB:IDUA protein purity.** After size exclusion, purified protein (1μg, measured by A<sub>280</sub>) was size-separated by SDS-PAGE and Coomassie stained following manufacture instructions (Thermo). A single band of 120 kDa was detected (A). Percentage of purity was further confirmed at 98.3% using an automated electrophoresis station BioRad Experion® Pro260 chip following manufacturer's procedures (B).



**Supplementary Figure S2.** RTB:IDUA protein identification. Purified protein detected by Coomassie staining (e.g., Fig. S1) was analyzed at University of Arkansas for Medical Science Proteomics Core by in-gel trypsin digestion and tandem mass spectrometry (MS/MS). Excluding the patatin signal peptide, there is a 37% coverage (334/895) of the sequence encompassing 32 exclusive unique peptides and 46 exclusive unique spectra.

