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Ulrich Kragh-Hansen, Lorenzo Minchiotti, Andrea Coletta, Konrad Bienk, Monica Galliano, Birgit Schiøtt, Yasunori Iwao, Yu Ishima, Masaki Otagiri

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Mutants and molecular dockings reveal that the primary L-thyroxine binding site in human serum albumin is not the one which can cause familial dysalbuminemic hyperthyroxinemia

**Ulrich Kragh-Hansen <sup>a,\*</sup>, Lorenzo Minchiotti <sup>b</sup>, Andrea Coletta <sup>c</sup>, Konrad Bienk <sup>d</sup>, Monica Galliano <sup>b</sup>, Birgit Schiøtt <sup>c</sup>, Yasunori Iwao <sup>e,1</sup>, Yu Ishima <sup>e</sup>, Masaki Otagiri <sup>e,2</sup>**

<sup>a</sup> *Department of Biomedicine, University of Aarhus, DK-8000 Aarhus C, Denmark*

<sup>b</sup> *Department of Molecular Medicine, University of Pavia, I-27100 Pavia, Italy*

<sup>c</sup> *Department of Chemistry and Interdisciplinary Nanoscience Center, University of Aarhus, DK-8000 Aarhus C, Denmark*

<sup>d</sup> *Department of Molecular Biology & Genetics and Interdisciplinary Nanoscience Center, University of Aarhus, DK-8000 Aarhus C, Denmark*

<sup>e</sup> *Department of Biopharmaceutics, Graduate School of Pharmaceutical Sciences, Kumamoto University, 5-1 Oe-honmachi, Kumamoto 862-0973, Japan*

\* Corresponding author at: Department of Biomedicine, University of Aarhus, Ole Worms Alle 6, Building 1180, DK-8000 Aarhus C, Denmark. Tel.: +45 8716 7798; fax: +45 8613 1160.

*E-mail address:* ukh@biomed.au.dk

<sup>1</sup> Present address: Department of Pharmaceutical Engineering and Drug Delivery Science, School of Pharmaceutical Sciences, University of Shizuoka, 52-1 Yada, Suruga-ku, Shizuoka, 422-8626, Japan

<sup>2</sup> Present address: Faculty of Pharmaceutical Sciences, Sojo University, 4-22-1 Ikeda, Kumamoto 860-0082, Japan

*Abbreviations:* HSA, human serum albumin; rHSA, recombinant HSA; Alb, albumin; Alb A, normal (wild-type) albumin isolated from serum; proAlb, proalbumin; Tr1 – Tr4, L-thyroxine binding sites.

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