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| Katherine A. Flores, MD Immediate-Past Chairwoman Director, UCSF-Fresno, Latino Center for Medical Education \& Research, Fresno, CA | Dear Legislators: |
| David Carlisle, MD, PhD President and CEO Charles R. Drew University of Medicine \& Biomedical Sciences, Los Angeles, CA | On behalf of the Board of Directors of the National Hispanic Medical Association, we urge you to vote yes for HB 7118 , a bill regarding FDA-designated interchangeable biological |
| Nerieda Correa, MD <br> Eastchester Medical Associates, PC, Brorx, NY | drug products. |
| Efrain Fuentes, EdD Los Angeles, CA | This bill authorizes a pharmacist to substitute an alternative biological product when filling |
| Jorge A. Girotti, PhD, MA <br> Associate Dean and Director Special Curricular Programs, Admissions, Hispanic Center of Excellence Assistant Professoc: Medical Education UIC College of Medicine, Chicago IL | a prescription for a prescribed biological product if the alternative biological product is designated as interchangeable with the reference product and communication is provided to the patient and physician that a substitution was made. This bill would also require that the |
| Antonio Linares, MD Medical Director and Regional VP, Anthem Blue Cross Health \& Wellness Solutions, Indianapolis, IN | substitution of a biological product be communicated to the patient. |
| Flavia E. Mercado, MD Medical Director, Inovalon, Inc. Adjunct Professor, Snelville, GA | We recognize the rising use of biologics and biosimilars in our population now aging with |
| Jorge G. Puente, MD Managing Partner <br> Pleasanton Pharma Ventures, Northport, NY | increased chronic disease. Biosimilars go through an extensive review process and manufacturers are required to submit immense studies and data demonstrating a products' |
| Diana Ramos, MD, MPH Directot, Reproductive Health, Los Angeles County Public Health, Laguna Beach CA | efficacy and ensuring it is safe for use by consumers. A pathway for biosimilar regulation in the U.S. was established as a provision of the 2008 Patient Protection and Affordable |
| Vanessa Saucedo, MD, MPH, FAAP Chairwoman, NHMA Council of Young Physicians Union Community Health Center, New York, NY | Care Act (ACA) and in 2012 the FDA issued draft guidelines for biosimilars and a list of biosimilars and interchangeable biological products. |
| Minerva Romero-Arenas, MD, MPH Chairwoman, NHMA Council of Residents General Surgery Resident, Sinai Hospital Baltimore, MD | In summary, the National Hispanic Medical Association recommends your support for HB 7118 before the end of session to clarify the procedures for biosimilar substitution for |
| Eric Molina President, Latino Medical Student Association Baylor College of Medicine, Houston, TX | biologic treatments in a way that increases safety for the patient. We are especially |
| Minerva Campos, MD, MPH Washington, DC | and all persons from Connecticut with chronic diseases. |
| Carlos Corral, MD, FACS New Mexico Cardiovascular Associates Las Cruces, NM |  |
| Claudia H. Zamora, MPA Manager PricewaterhouseCoopers, LLP Washington DC | Sincerely, |
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