(b) The substantial differences in the prices at which drug products are offered to American and foreign purchasers. (P. 49)

(c) Revision of current patent and trademark laws on prescription drugs. (P. 49)

Drug Distributors

5. The Congress should enact legislation requiring that the containers of all dispensed prescription drugs be labeled with the identity, strength and quantity of the product, except where this is waived upon specific orders of the prescribers. (P. 53)

6. Encouragement should be given to the wider use of prepackage dispensing, in which manufacturers prepare and pharmacists dispense tablets and capsules in precounted form, in sealed, pre-labeled containers, and in such numbers as conform to those most frequently prescribed by physicians. (P. 53)

7. The National Center for Health Services Research and Development should develop and support research to improve the efficiency and effective-