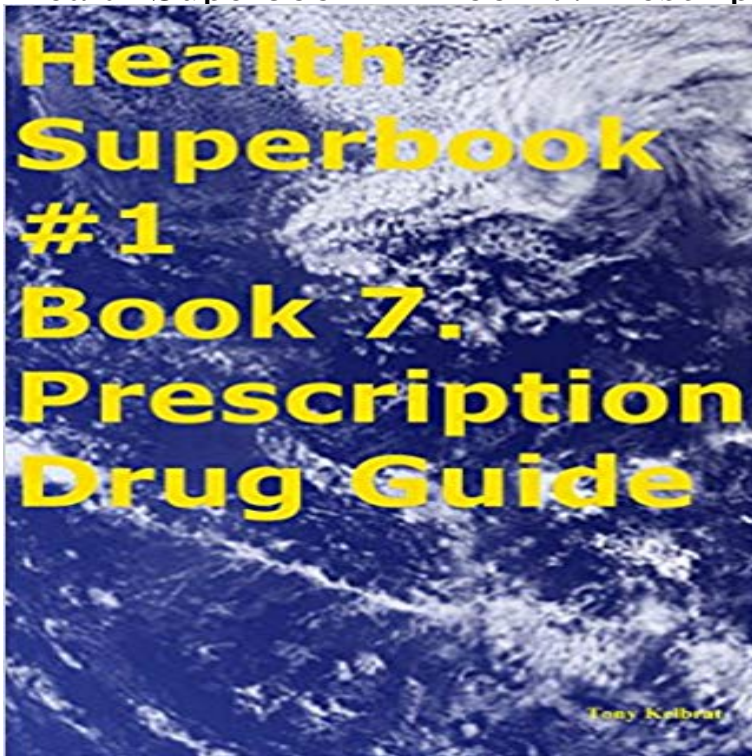


## Health Superbook #1 Book 7. Prescription Drug Guide



Educate yourself about any condition you might have and the treatments from both the Western and holistic points of view. Use both to heal yourself. Like the golden mean, excess in anything is not good for you. Use generic as opposed to brand name drugs to save money. They're drugs whose patent has run out so now anybody can produce them. The original company keeps the right to its brand name so when the doctor writes out your prescription, ask him to write out the generic name of the drug, if applicable as opposed to the brand name which many do, probably because of habit or they get some form of favor from that company. Not all drugs are manufactured in a generic version. The major points of medical drug use are: Know the drug you're using. If it doesn't work, ask for another drug that does the same function. There are many different drugs that treat the same diseases. Follow the dosage instructions. Throw old drugs away. Beware of safety factors in driving and hazardous work when taking certain drugs. If pregnant or breastfeeding, be wary of taking certain drugs. Be wary of taking alcohol with drugs, Keep drugs away from the reach of children. Be wary of drug interactions with other drugs and certain foods. Follow the conditions in which you should take drugs like on an empty stomach, in the morning, before bedtime, etc. Some major drug books are: Essential Guide to Nonprescription Drugs Essential Guide to Prescription Drugs Essential Guide to Psychiatric Drugs Physicians Desk Reference The Complete Drug Reference Books about drugs are at #613.2662, #615.1097, #615.78 and RC483.5 or RM236 to RM301 at the library. Books about naturopathic drugs are at #615.535 or RZ440 at the library. Books about the pharmaceutical industry are at #338.476 or HD9665.5 at the library. Drug Testing Approval Process A new drug is defined in part as: any drug the

composition of which is such that such drug is not generally recognized, among experts qualified by scientific training and experience to evaluate the safety and effectiveness of drugs, as safe and effective for use under the conditions prescribed, recommended, or suggested in the labeling thereof. (21 U.S.C. 321(p)(1)) A key provision of the statute specifies that a new drug cannot be approved and therefore cannot be shipped in interstate commerce, until there is substantial evidence that it is safe and effective for its intended use. The term substantial evidence refers to evidence derived by: adequate and well-controlled investigations, including clinical investigations, by experts qualified by scientific training and experience to evaluate the effectiveness of the drug involved, on the basis of which it could fairly and responsibly be concluded by such experts that the drug will have the effect it purports or is represented to have under the conditions of use prescribed, recommended, or suggested in the labeling or proposed labeling thereof. (21 U.S.C. 355 (d)) Before initiating well-controlled investigations (as specified in the section of the statute quoted above), FDA requires that drugs be studied experimentally in animal systems; if the results of those tests fulfill FDA criteria, FDA allows the drug to proceed to a final stage of testing in specific clinical (human) trials. Any person or company wanting to conduct clinical research on an unapproved drug must submit an investigational new drug (IND) application to FDA. If the application contains sufficient detail to meet FDA's requirements, the IND is allowed to proceed, exempting the sponsor from the FDCA prohibition against shipping unapproved drugs in interstate commerce for the study or studies specified in the IND and ensures that FDA can monitor the clinical research process. Usually, a progression of clinical trials is required, culminating in large, randomized clinical trials.

l'oeil de boeuf

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