# COUNTERFEIT MEDICINES IN LEGITIMATE SUPPLY CHAINS

A <u>study</u> by

B

University of California, San Diego School of Medicine & the Pharmaceutical Security Institute (PSI)

Download <u>here</u> the study Counterfeit Drug Penetration into Global Legitimate Medicine Supply Chains: A Global Health Assessment

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\* Mackey TK, Liang BA, York P, Kubic T. Counterfeit Drug Penetration into Global Legitimate Supply Chains: A Global Assessment. Am J. Trop. Med. Hyg; 2015; doi:10.4269/ajtmh.14-0389 [advanced online publication]





**THERAPEUTIC CATEGORY IS AFFECTED** 52.8% of all counterfeit medicines detected in the legitimate supply chain are lifesaving-related treatments: 21.1% 11.0% 11.6% 9.1% Anti-infective\* Cardiovascular Alimentary Central nervous

**EVERY** 

\*including both branded and generic anti-infectives/antibiotics, such as anti-malarial

system

### **ROUTE OF ADMINISTRATION:**

77.3% 15

were oral dosage formulations

15.4% were injectable biologicals

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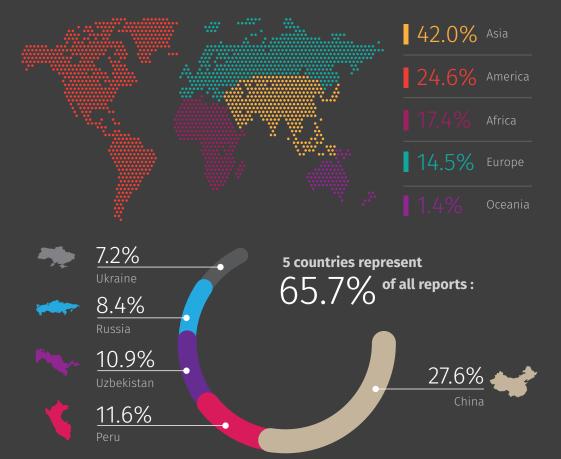
### REPORTS ON COUNTERFEIT MEDICINES ARE LACKING. BASED ON PSI REPORTING SYSTEM ANALYSIS FOUND THAT:

35.2% more than one-third of the nations in the world had their legitimate supply chain compromised

of the reports contained limited information precluding further classification

## SOURCES OF REPORT SHOW A GREAT DIFFERENCE BETWEEN REGIONS:

Middle income markets were most represented



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Improvements in surveillance, including detection of security breaches, data collection, analysis, and dissemination are urgently needed to address the public health need to combat the global counterfeit medicines trade

#### CONCLUSIONS:

Three key needs emerged from this study:

- greater consistency, standardization and robust data collection for analysis,
- enhanced reporting by countries,
- > increased details in the reports.

### **RECOMMENDATIONS:**

- development and coordination of a "single point of contact" system comprised of relevant country authorities to conduct surveillance and reporting counterfeit incidents to a central global system,
- development of a "graded best practices" set of guidelines supporting ongoing training, capacity building, technical assistance, data sharing, and partnership. These guidelines should be developed in cooperation with regulatory authorities, law enforcement, customs, pharmaceutical companies, academic institutions, and other stakeholders.