PROTECT YOUR BUSINESS WITH THE RIGHT SANITIZER

AS THE INVENTORS OF FOAMING HAND SANITIZER

SC JOHNSON PROFESSIONAL® TAKES QUALITY VERY SERIOUSLY.

HAND SANITIZER IS AN FDA REGULATED OVER-THE-COUNTER DRUG WITH REQUIREMENTS FOR HOW PRODUCTS ARE PRODUCED AND PACKAGED[1].

THE FDA HAS RECALLED OVER **200 HAND SANITIZERS** FROM THE MARKET SINCE THE ONSET OF THE PANDEMIC. THESE RECALLS WERE BASED ON VARIOUS FACTORS SUCH AS: **CONTAMINATION WITH POTENTIALLY TOXIC TYPES OF ALCOHOL, NOT ENOUGH ACTIVE INGREDIENT, AND LABELS WITH FALSE, MISLEADING, OR UNPROVEN CLAIMS[2].**

ALL **SC JOHNSON PROFESSIONAL® HAND SANITIZERS** ARE REGISTERED WITH **THE FDA AND ARE SAFE AND EFFECTIVE WHEN USED PROPERLY.**

THERE ARE MANY TYPES AND GRADES OF ALCOHOL. ONLY CERTAIN GRADES OF ETHYL ALCOHOL AND ISOPROPYL ALCOHOL ARE ACCEPTABLE FOR USE AS ACTIVE INGREDIENTS IN ALCOHOL HAND SANITIZER ACCORDING TO THE FDA. OTHER GRADES OF ALCOHOL MAY INCLUDE IMPURITIES, SUCH AS METHANOL, WHICH IS **TOXIC TO HUMANS[2].**

SC JOHNSON PROFESSIONAL® ALCOHOL-BASED HAND SANITIZERS ARE **EXCLUDED FROM USING ALTERNATIVE GRADE ALCOHOL, AS THEY ARE NON-WHO FORMULAS.** THE ETHANOL USED IN SC JOHNSON PROFESSIONAL INSTANT HAND SANITIZERS IS US PHARMACOPEIA (USP) GRADE AS REQUIRED BY THE FDA MONOGRAPH.

THERE ARE CIRCUMSTANCES WHERE HAND HYGIENE IS REQUIRED

BUT ALCOHOL SANITIZERS CANNOT BE USED DUE TO CUSTOMER CONCERNS, INCLUDING RELIGIOUS REASONS OR RISK TO PATIENTS AND/OR USERS SAFETY, E.G. WHERE PATIENTS/USERS MAY DRINK THE PRODUCT (ACCIDENTALLY OR OTHERWISE) DUE TO ALCOHOLISM OR MENTAL HEALTH ISSUES. IN THESE CIRCUMSTANCES, WELL FORMULATED NON-ALCOHOL HAND SANITIZERS THAT HAVE BEEN TESTED FOR ANTIMICROBIAL EFFICACY USING INTERNATIONALLY RECOGNIZED TEST METHODS OR STANDARDS CAN BE AN EFFECTIVE ALTERNATIVE.

YOUR WORLD MEANS THE WORLD TO US.

THAT'S WHY WE GO ABOVE AND BEYOND TO ENSURE YOU KNOW EXACTLY WHAT GOES INTO THE MAKING OF OUR PRODUCTS.

SC JOHNSON PROFESSIONAL® PRODUCTS ARE MANUFACTURED IN FACILITIES WHICH FOLLOW CURRENT GOOD MANUFACTURING PRACTICE (CGMP) REQUIREMENTS. ALL RAW MATERIALS USED FOR PRODUCTION UNDERGO A THOROUGH QUALITY CONTROL PROCESS BEFORE BEING USED FOR MANUFACTURING IN OUR HIGH-QUALITY PRODUCTS. ALL FINISHED GOODS ARE SUBJECT TO EXTENSIVE QUALITY TESTING BEFORE BEING SHIPPED OUT TO OUR CUSTOMERS.

1 - https://www.fda.gov/drugs/coronavirus-covid-19-drugs/hand-sanitizers-covid-19
2 - https://www.fda.gov/consumers/consumer-updates/your-hand-sanitizer-fdas-list-products-you-should-not-use

READ THE LABEL

ALL SANITIZER LABELS MUST CONTAIN THE FOLLOWING INFORMATION:

1. A DRUG FACTS TABLE
2. ACTIVE INGREDIENT STATEMENT - WE RECOMMEND ACTIVE INGREDIENT TO BE ETHANOL WITH AT LEAST 70% CONTENT
3. THE PURPOSE OF THE PRODUCT
4. USE DIRECTIONS

5. A WARNINGS SECTION
6. A LISTING OF INACTIVE INGREDIENTS
7. THE NAME AND ADDRESS OF THE MANUFACTURER
8. AN EXPIRATION DATE AND BATCH (LOT) NUMBER
9. USE A RECOGNIZED MANUFACTURER OR PRODUCTS WITH NDC (NATIONAL DRUG CODE) REGISTERED WITH FDA
10. WE RECOMMEND YOU ENSURE PRODUCT HAS BEEN TESTED AGAINST A BROAD SPECTRUM OF MICRO-ORGANISMS USING RECOGNIZED ASTM TEST METHODS AND DEMONSTRATES A MINIMUM OF 4 LOGS OF EFFICACY IN VITRO AND MEETS FDA REQUIREMENTS FOR IN-VIVO EFFICACY.

There are risks to using bulk hand sanitizer that include: misbranding, unknown product integrity, stability and traceability. We recommend the use of sanitizer in hygienically sealed cartridges or containers that are not refillable to reduce these risks.

ACCORDING TO THE CURRENT FDA MONOGRAPHS FOR TOPICAL ANTISEPTICS, NO SANITIZER CAN CLAIM:

1. EFFICACY AGAINST COVID-19, OR ANY VIRUS
2. EXTENDED PROTECTION SUCH AS “LASTS FOR 24 HOURS”

IF YOUR SANITIZER LABEL IS IN VIOLATION OF ANY OF THESE POINTS, YOU MAY NOT BE PROTECTING YOUR BRAND OR YOUR CUSTOMERS.