



April 22, 2019

Reference: URGENT OTC/DRUG Product Recall

Dear Inopak Customer;

This is to inform you of a product recall involving Over the Counter (OTC) Hand Soap/Sanitizers manufactured, packaged and distributed by Inopak LTD of Ringwood, NJ.

As part of Inopak's on-going commitment to product quality we have determined through our quality processes there may be a potential for a limited number of containers of our hand sanitizers and soaps that are not GMP compliant.

While we have no reported customer issues or adverse events, in an abundance of caution, we are recalling those products manufactured at our Ringwood, NJ facility between January 1, 2016 and June 30, 2018.

Below is a comprehensive list of the Product Name, SKU (part number), product description and batch numbers included in the recall.

OVER THE COUNTER (OTC) PRODUCTS

Product	Description	Batch #
ANTIBACTERIAL FOAM SKU# 5063	Antibacterial Foam Soap	6657, 6666, 6671, 6679 6687, 6695, 6702, 6718, 6730, 6744, 6754, 6760, 6772, 6773, 6784, 6793, 6804, 6812, 6819, 6838, 6846, 6856, 6862, 6877, 6890, 6896 , 6902, 6910, 6919, 6938, 6944, 6953, 6971, 6985, 6990,6998, 7005, 7012, 7018, 7028, 7033, 7039, 7043, 7053, 7060, 7068, 7080, 7092, 7101, 7110, 7116, 7126, 7143, 7153, 7161, 7167, 7175, 7188, 7204, 7208, 7213, 7221, 7232, 7242, 7248, 7267, 7275, 7284, 7287, 7295, 7308, 7314, 7324, 7329, 7336, 7348, 7357, 7363, 7372, 7382, 7389, 7397, 7403, 7411
AQUACIL FHS SKU# BIO-5075	Non-Alcohol Waterless Hand Sanitizer	6736, 6826
DERMAGEL SKU# 5025	Waterless Hand Sanitizer gel	7069, 6648, 6649, 6652, 6654, 6667, 6669, 6675, 6677, 6684, 6686, 6692, 6694, 6703, 6704, 6710, 6717, 6719, 6726, 6727, 6731, 6733, 6739, 6743, 6748, 6751, 6752, 6759, 6763, 6767, 6769, 6771, 6775, 6776, 6780, 6782, 6787, 6788, 6801, 6805, 6808, 6811, 6814, 6815, 6821, 6824, 6825, 6830, 6831, 6836, 6843, 6845, 6852, 6859, 6860, 6874, 6878, 6880, 6881, 6884, 6889, 6897, 6900, 6903, 6905, 6912, 6920, 6922, 6926, 6928, 6931, 6940, 6945, 6948, 6951, 6954, 6962, 6967, 6969, 6972, 6975, 6980, 6986, 6989, 6995, 7002, 7015, 7023, 7029, 7030, 7035, 7044, 7047, 7048, 7051, 7059, 7062, 7066, 7072, 7073, 7079, 7081, 7084, 7085, 7088, 7090, 7095, 7096, 7099, 7102, 7105, 7112, 7113, 7115, 7124, 7125, 7128
E 2 FOAM SKU# 5064F	Antibacterial Foaming Food Handling Soap	7304, 7180, 6707, 7065
E2 SOAP SKU# 5064	Food Handling Soap	7074
HEALTHCARE 2000 SKU# 5050	Antibacterial Hand Soap	6802, 6803, 6909, 6655

MILD HEALTHCARE SKU# 5013	Antibacterial Hand Soap	6656, 6959, 7008
SANIGUARD SKU# 5068	Waterless Foam Hand Sanitizer	6668, 6721
STYLE SKU# 5031	Antibacterial Hand Soap	6660, 6664, 6676, 6682, 6685, 6698, 6701, 6725, 6725, 6732, 6742, 6744, 6746, 6756, 6762, 6770, 6777, 6792, 6798, 6807, 6813, 6816, 6820, 6829, 6839, 6868, 6872, 6875, 6891, 6892, 6904, 6906, 6914, 6916, 6923, 6932, 6939, 6952, 6958, 6963, 6970, 6978, 6982, 6987, 6993, 6997, 7004, 7006, 7024, 7038, 7049, 7054, 7056, 7058, 7078, 7087, 7098, 7104, 7108, 7111, 7129, 7132, 7134, 7138, 7145, 7152, 7157, 7160, 7164, 7170, 7177, 7184, 7190, 7192, 7203, 7207, 7220, 7224, 7231, 7235, 7237, 7243, 7247, 7252, 7258, 7260, 7269, 7281, 7285, 7291, 7296, 7301, 7306, 7311, 7312, 7319, 7322, 7326, 7335, 7338, 7340, 7347, 7351, 7353, 7368, 7377, 7384, 7387, 7394, 7399
WHITE BAJAN SKU# 5012T	Antibacterial Hand Soap	6633, 6827
MEDIWASH SKU# 5050F	Antibacterial Foam Hand Soap	6691

All products were manufactured from January 1, 2016 – June 30, 2018 and all have a 3-year expiration from the date of manufacture.

Expiration Dates would range from January 2019 through June 2021.

All products were packaged in one or more of the following package type and sizes;

- Plastic Bottles – 4 oz., 8 oz., 18 oz., and gallon.
- Pouches – 800 ml., 1000 ml., or 2000 ml.

Immediately examine your inventory and quarantine product subject to recall. In addition, if you may have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this recall notification letter.

This recall should be carried out to the retail level.

Your assistance is appreciated and necessary to prevent any possible harm to users.

Alternatively, this information may also be found, along with return instructions, forms, and process on the Inopak Recall Webpage <http://www.inopak.com/recall>. (Alternatively, you can visit our website home page at [inopak.com](http://www.inopak.com) and then click the “Recall Info” link on the lower-left.)

Here, you will find a login portal to see the list of batches, forms and instructions:

Username: recall

Password: products224

Our recall coordinator, Mr. Nicky DiSarro, we be contacting you in the next few days with additional instructions and follow-up.

If you should have any questions please contact Mr. DiSarro, VP of Sales, at (800) 762-7725. Mr. DiSarro may also be contacted by email at the following address: Nick28@warwick.net

Hours of Operation

Monday – Friday 8:00 am – 5:00 pm. Eastern Standard Time.

Closed Weekends and Holidays

This recall is being made with the knowledge of the Food and Drug Administration.