

Spacers Are Not Interchangeable - Aerosol Evaluation of Online Knock-Off Devices

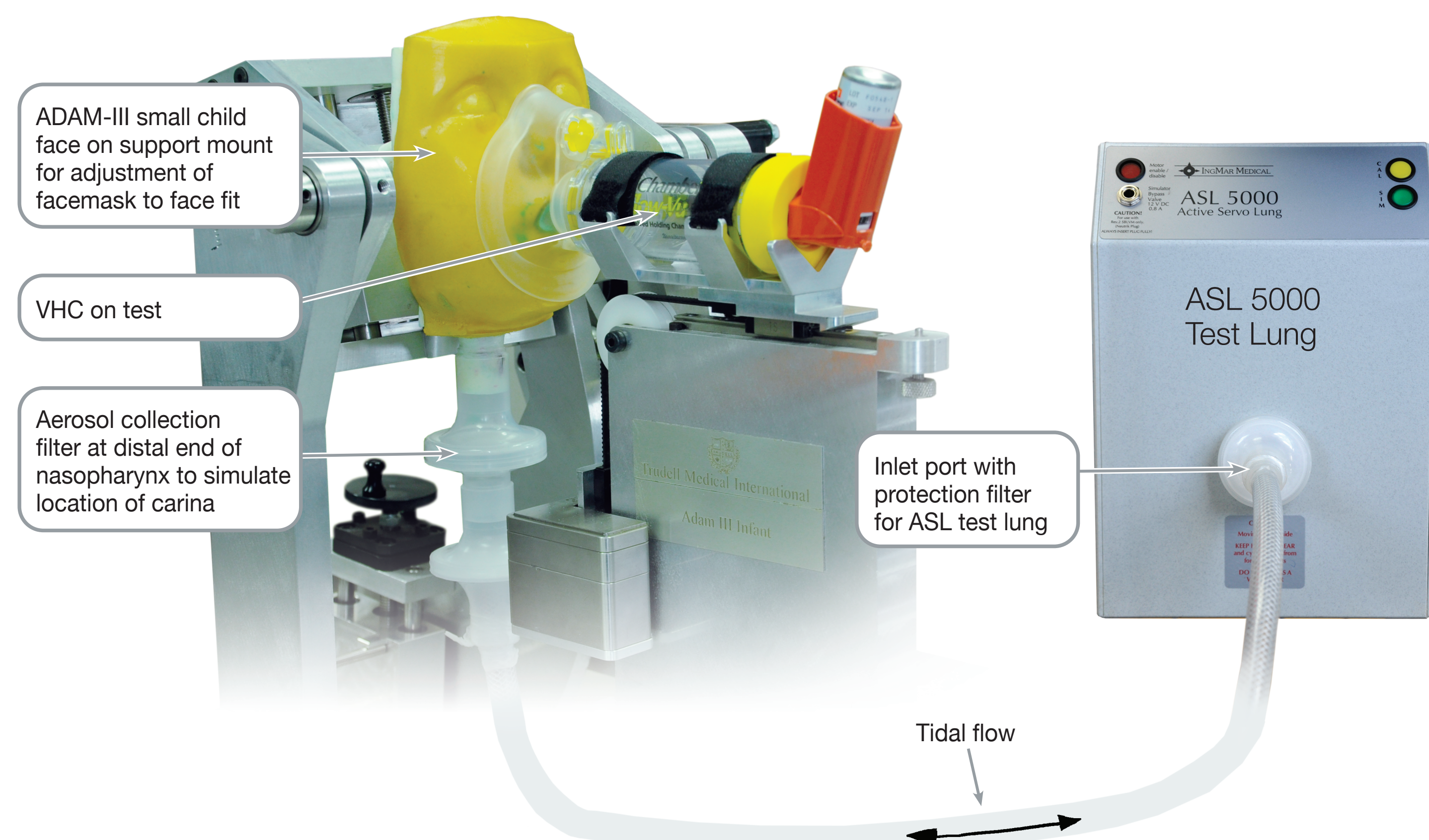
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RATIONALE

Online marketplaces offer convenience and a vast array of options making them an attractive choice for purchasing medical devices. However, spacers are an integral part of the pMDI delivery system, and this convenience comes with inherent dangers and risks that consumers must be aware of. The primary concern lies in the lack of regulation and oversight in online marketplaces, leading to the availability of counterfeit, substandard, and potentially dangerous products.

METHODS

- Chambers were purchased from Amazon.ca and evaluated by breathing simulator mimicking a coordination delay of 2 seconds before inhalation, followed by tidal breathing (tidal volume = 155-mL, I:E ratio = 1:2, rate = 25 cycles/min).
- Each spacer (with mask) was attached to an anatomical model of a 4-year old child face and the airway coupled to the breathing simulator via a filter located at the exit to capture drug particles that penetrated as far as the carina.
- 5-actuations of fluticasone propionate (FP, Flovent⁺ 50) were delivered at 30-s intervals and recovered from specific locations in the aerosol pathway by HPLC-UV spectrophotometry.



RESULTS

Delivery of Flovent⁺ HFA pMDI to the Carina



CONCLUSION

Significantly more medication was delivered to filter/carina with the **AeroChamber Plus[®] Flow-Vu[®] VHC**. (un-paired t-test, $p < 0.001$). While many spacers may visually appear similar, clinicians need to be aware that not all chambers are the same.