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Equipment-free Detection of K⁺ on Paper

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Microfluidic paper-based analytical devices (μ PAD) have been introduced as simpler variants of lab-on-a-chip (LOC) devices, aiming at point-of-care (POC) diagnostics of various analytes. While colorimetric detection is a promising signal output principle, it has difficulty meeting the requirements of the World Health Organization and associated feasibility of commercialization.

This work describes µPADs for the equipment-free detection of K⁺ in a 10 µL serum sample where K⁺ concentration is translated to a distance-based signal. This goal is achieved by separating recognition and detection steps. The recognition part uses an ion-selective film solvent-cast into a glass capillary. This coating contains a charged dye, thioflavin T (ThT+), along with the K+ ionophore valinomycin. Once a 10 µL sample volume is aspirated into the capillary by a commercial pipette, K⁺ in the sample is allowed to be exhaustively exchanged with ThT+, thereby releasing a quantity of ThT+ that reflects the original amount of K+. To allow for a distance-based detection, this ThT⁺ is discharged into a paper channel defined by hydrophobic wax barriers. As the sample flows, ThT+ binds electrostatically to anionic functionalities of the cellulose substrate, which is further enhanced by a polyanionic coating. Higher amounts of K+ translate into a higher quantity of ThT+, in turn resulting in an increased distance of the perceivable color band on the µPAD.

The exhaustive depletion of K⁺ makes it possible to detect K⁺ with high sensitivity in a narrow concentration range, suitable for



Apply a drop of serum through a special capillary into a paper channel and get the result!

Rapid & exhaustive ion exchange in capillary





Schematic illustration of the distance-based analysis of potassium ions with an ion-selective capillary film (top) that exhaustively exchanges potassium ions for the cationic dye ThT⁺. This dye is in turn detected on paper *via* electrostatic interactions, giving a distance-based visual read-out (bottom). Adapted with permission from Y. Soda *et al.*, *ACS Sensors* **2019**, *4*, 670. Copyright (2019) American Chemical Society.



Result of distance-based analysis of K^+ in pooled serum samples. Adapted with permission from Y. Soda *et al.*, *ACS Sensors* **2019**, *4*, 670. Copyright (2019) American Chemical Society.

serum diagnostics (3.5~5 mM). In comparison with traditional ion optodes, this readout principle does not depend on the sample pH and gives a more sensitive response while maintaining a high selectivity. The distance-based readout is more robust than colorimetric detection, which is notoriously difficult to quantify, and does not require any readout equipment. This device principle may pave the way for the practical realization of μ PADs for the detection of ions.

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