

Characterization of Two Inward Shaker K⁺ Channels EgKAT1 and EgKAT2 in Oil Palm

Sandra Espeout¹, Sabine D. Zimmermann², Maelle Delestre¹, Camilo Pesantez¹, Norbert Billotte¹, Rémy Michel¹, Sergi Navarro¹, Reinout Impens³, Florence Jacob⁴, Isabelle Gaillard² and Teresa Cuellar¹

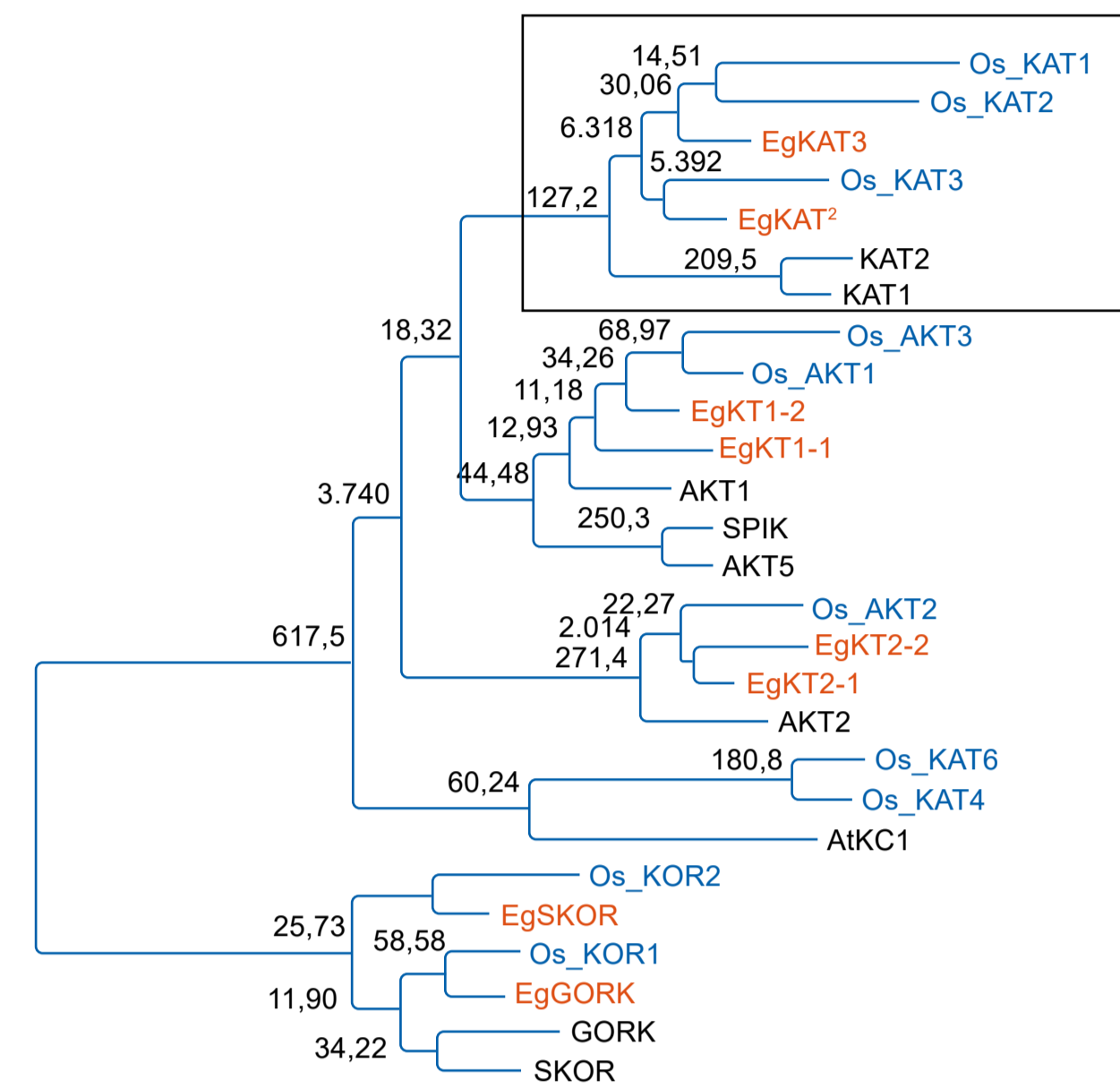
¹UMR AGAP Institut, Univ Montpellier, CIRAD, INRAE, Institut Agro, Montpellier, France, ²IPSIM, Univ Montpellier, CNRS, INRAE, SupAgro, Montpellier, France, ³SIAT, Presco-Plc, Benin City, Nigeria, ⁴PalmElit SAS, F-34980 Montferrier-sur-Lez, France. teresa.cuellar@cirad.fr / sandra.espeout-fois@cirad.fr

Context

- Potassium (K⁺) is a key factor for maintaining high yields in oil palms.
- K⁺ is transported by a large family of channels and transporters.
- Shaker channels are mainly responsible for not only long-distance flows in the plant.
- To maintain high yields while reducing K⁺ fertilization, it is necessary to understand the K⁺ transport mechanisms, in particular the Shaker channels.

>> Molecular and functional characterization of Shaker K⁺ channels in oil palm?

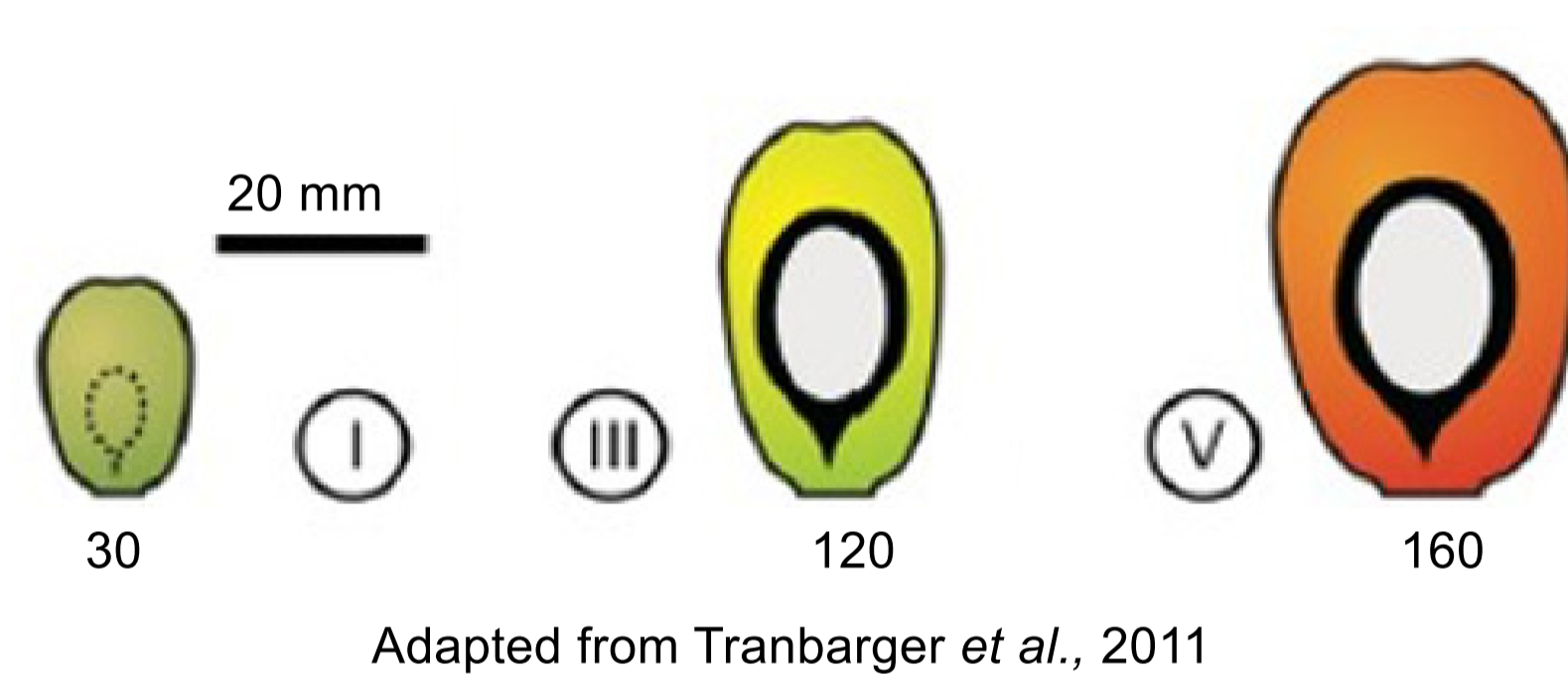
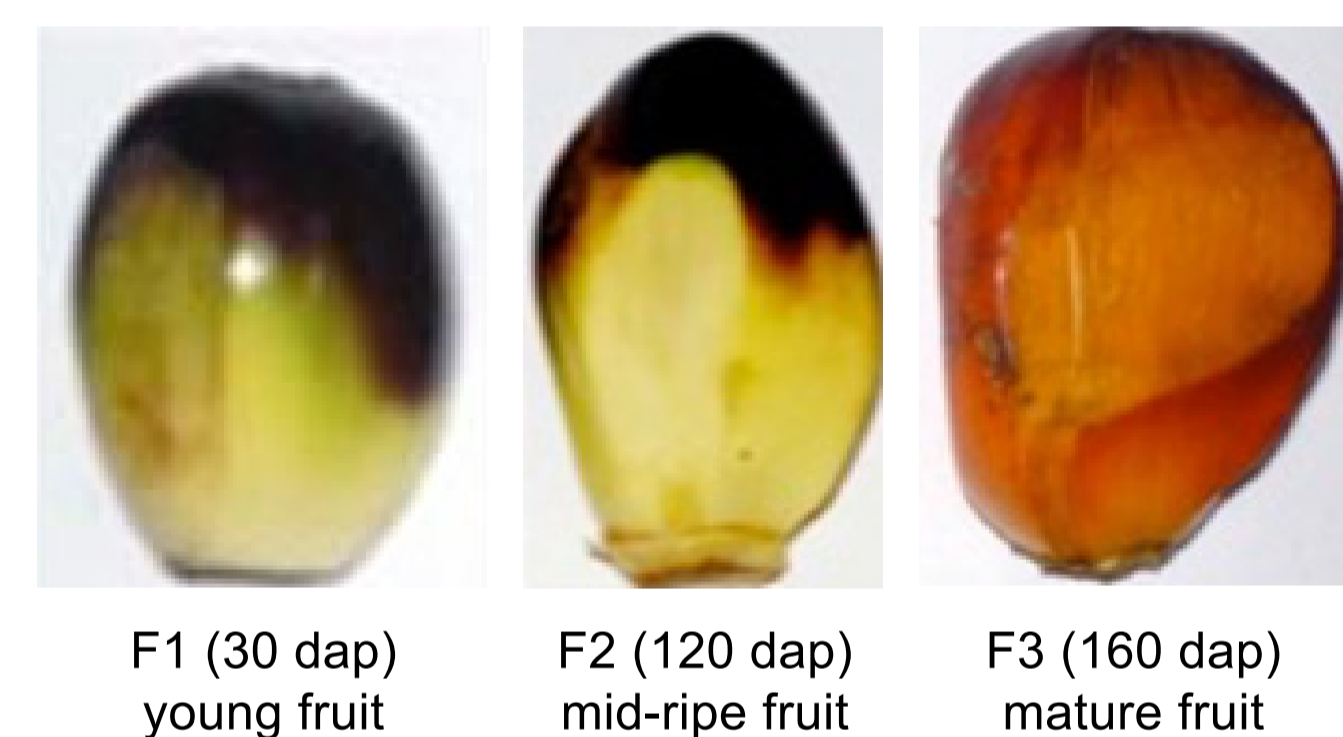
➔ Focus on two Shaker inward channels: EgKAT1 and EgKAT2



Phylogenetic relationships of Shaker Channels in Arabidopsis (black), Rice (blue) and oil palm (red). EgKAT sub family in black box (Monder *et al.*, 2022)

Genetic material

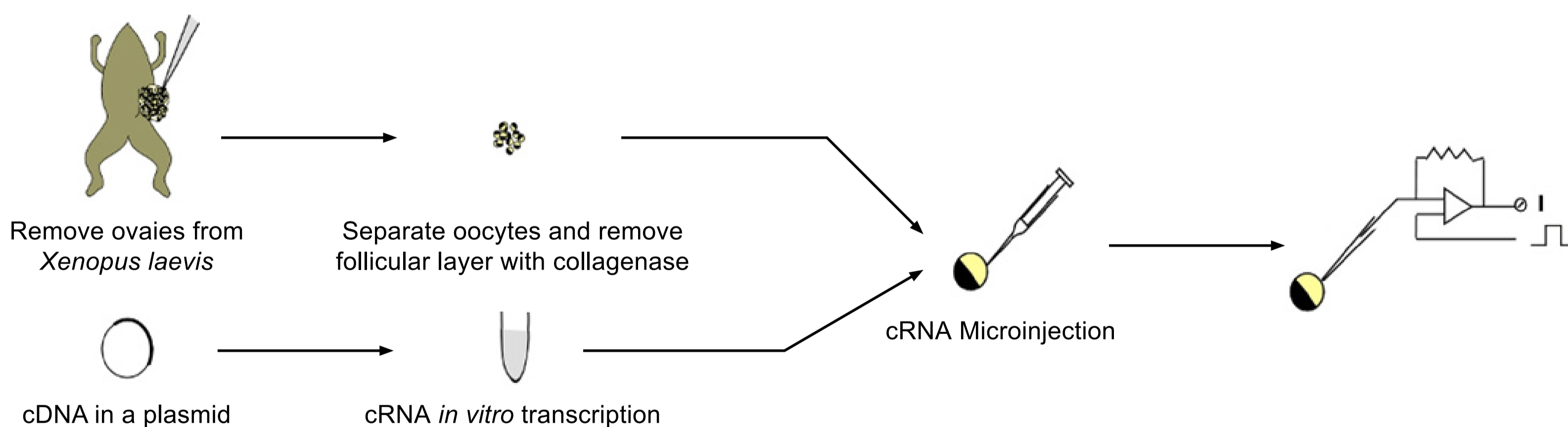
- 27 individuals Deli × La Mé planted in Presco, Nigeria.
- 3 stages of fruits development from adult palm : F1, F2, F3
- 102 individuals Deli × La Mé from greenhouse young plants.



Adapted from Tranbarger *et al.*, 2011

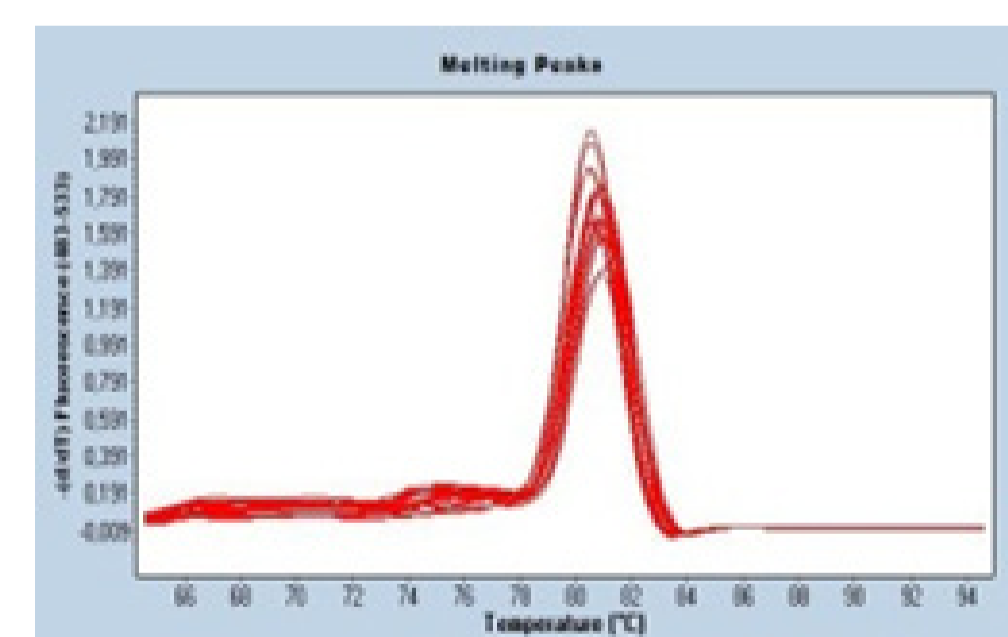
Electrophysiology

- Functional properties of EgKAT1 and EgKAT2 channels in *X. laevis* oocytes by voltage-clamp analyses.



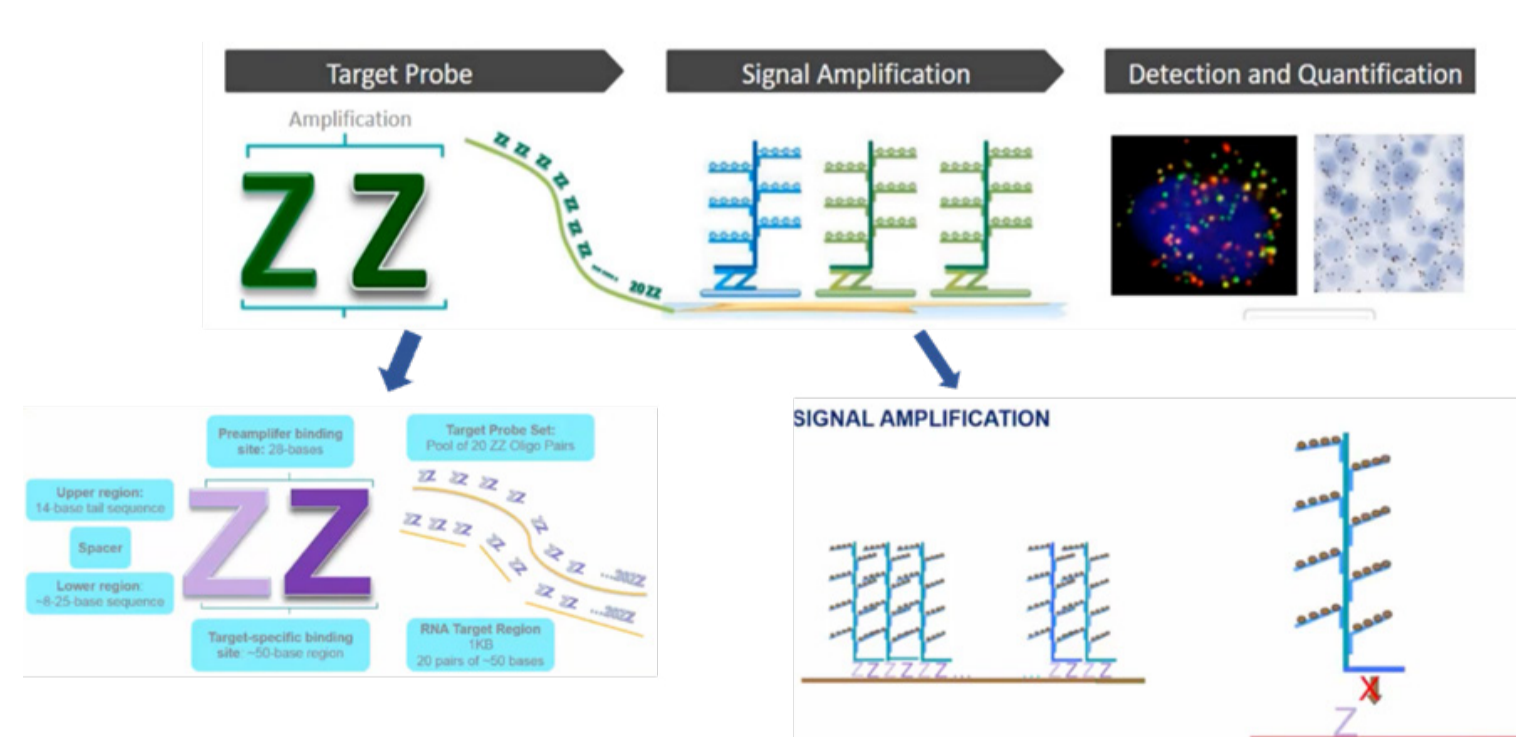
Expression by RT-qPCR

- Tubuline used as housekeeping gene.
- 3 biological replicates.
- 3 young tissues (leaf, meristem and root).
- 3 stages of fruit development.



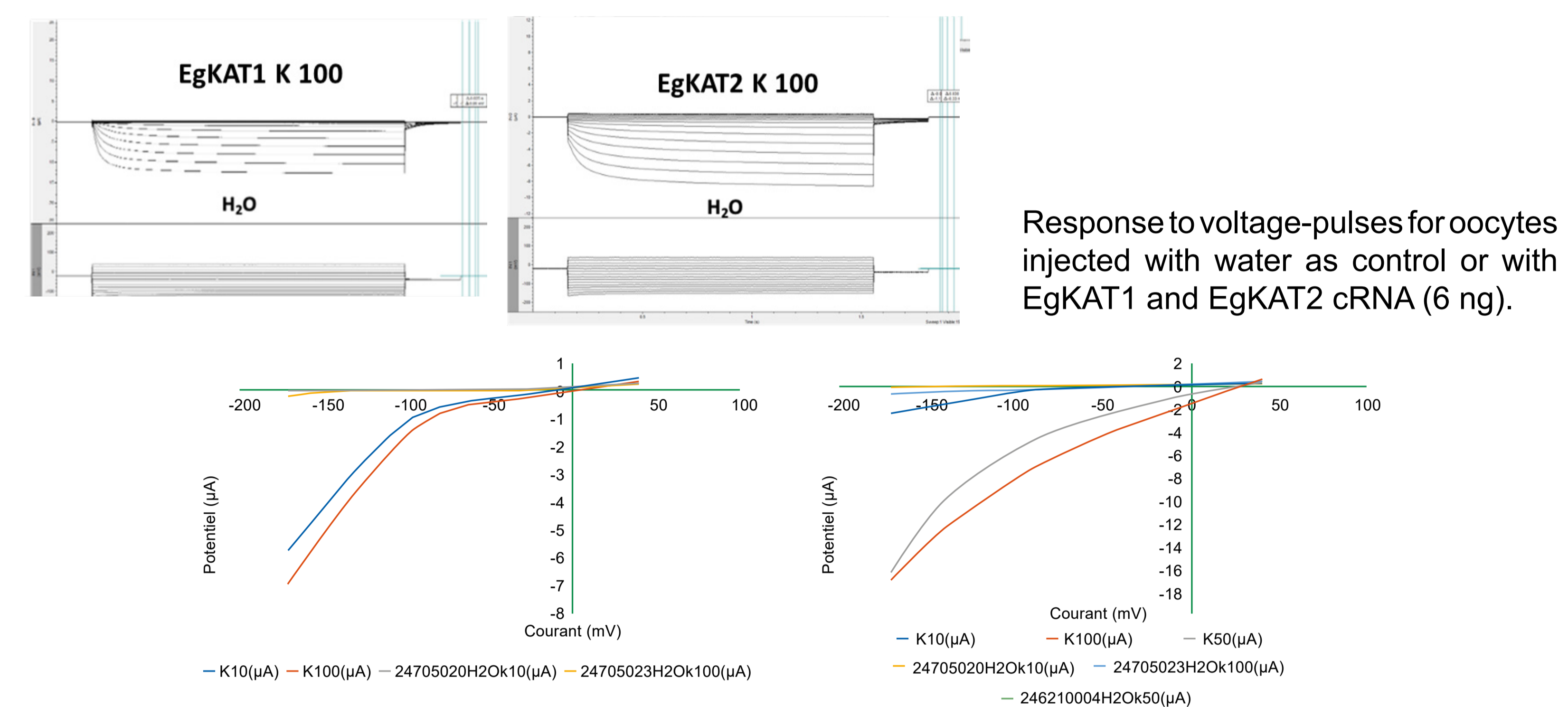
Tissue localization by RNAscope

- New technology *in situ* hybridization for the detection of a single RNA molecule in oil palm tissues.
- (Solanki *et al.*, 2020; Bowling *et al.*, 2014 and Munganyinka *et al.*, 2018).
- Target Probe, signal amplification and chromogenic signal dot detection (RED) <http://www.acdbio.com>



Results

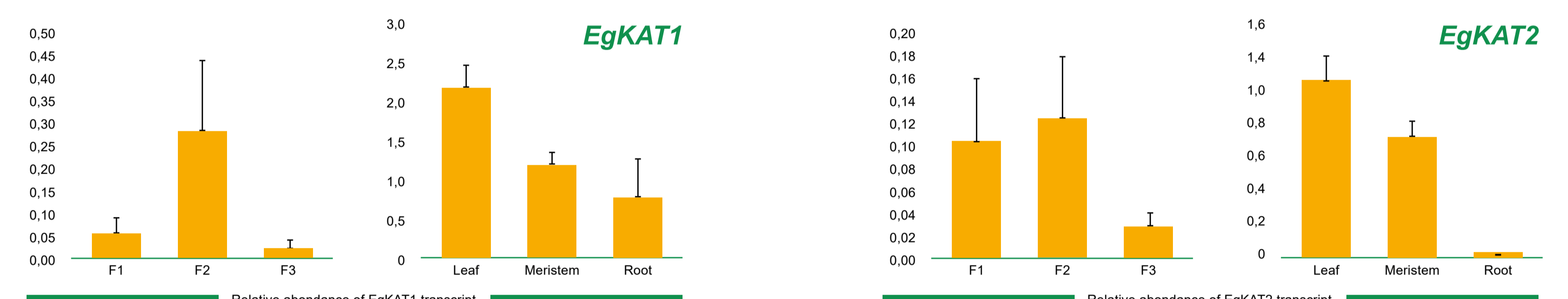
EgKAT1 and EgKAT2 are inwardly rectifying channels with different kinetics. EgKAT1 is activated more quickly.



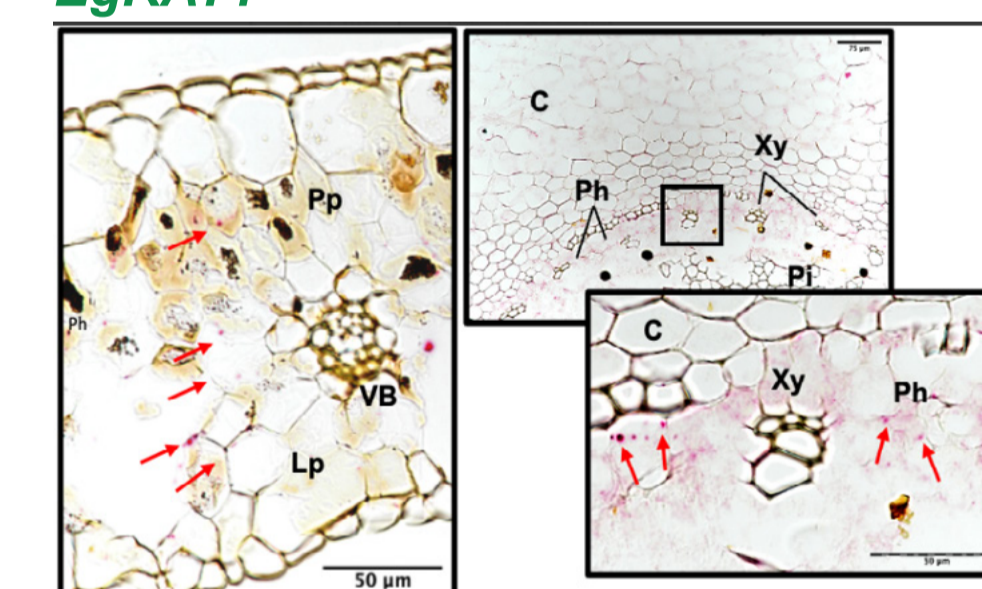
Current-voltage curves for mean currents mediated by EgKAT1 and EgKAT2 in different external K⁺ concentrations (K10, K50 and K100 mM), pH 6,5 (n=7 and n=4, ± SE)

For 2 genes (Kruskal-Wallis test)

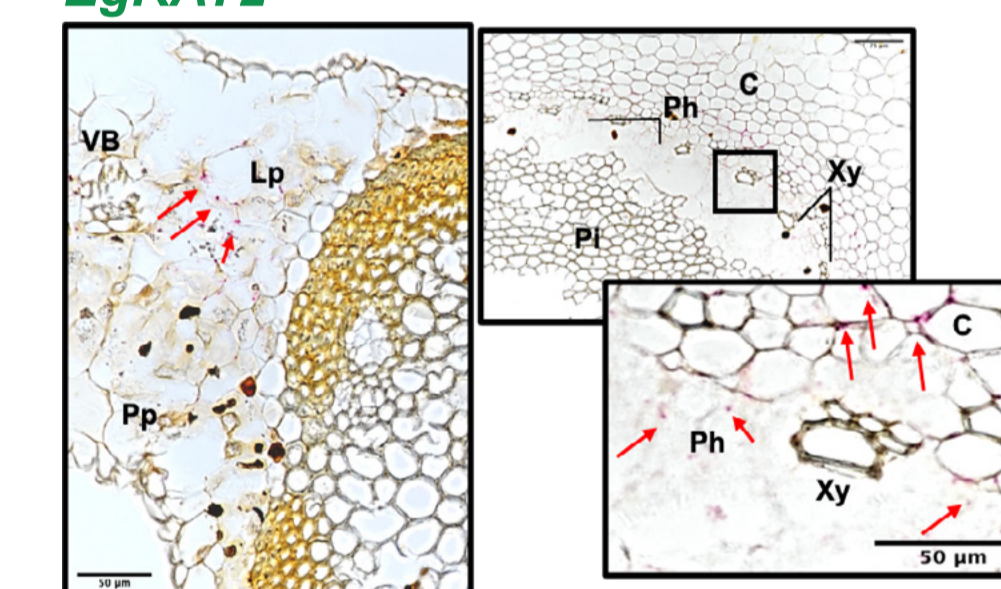
- stronger expression for F2.
- stronger expression for leaves and meristems and weaker for roots in young plants.



EgKAT1



EgKAT2



Transcripts of EgKAT1 and EgKAT2 are localized

- in the phloem and cortex in roots.
- in the parenchyma (palisade and lacunar) cells of the young leaves.
- in the mesocarp cells of F2 fruits and meristem (data not shown).

C, cortex; Lp, Lacunar parenchym; Pp, Palisade parenchyma; Pi, pith; Ph, phloem; VB, vascular bundles; Xy, Xylem

Conclusions

- EgKAT1 and EgKAT2 mediated inward K⁺ uptake currents.
- These genes are expressed in the phloem and cortex in roots, indicating a role in K⁺ uptake from soil solution and parenchyma cells of the leaves suggesting their implication in photosynthesis.

➔ improved yield

>> This work provides a better understanding of the potassium transport mechanism in oil palm and opens up new prospects for research aimed at reducing K⁺ fertilization.

Bibliographic references

Boccaccio, 2001, Thèse pp105; Solanki *et al.*, 2020, Plant Methods 16, 71; Bowling *et al.*, 2014, Appl. in Plant Sci., 2: 1400011; Monder *et al.*, 2022, SFBV. Sept., Munganyinka *et al.*, 2018, Virol J., 14;15(1):128; Tranbarger *et al.*, 2011, Plant Physiol 156, 564-584