Quick start protocol

1.

Add your sample in appropriate buffer to the substrate (remember controls and replicas)

2.

Incubate the reaction at a certain temperature with shaking

3.

Centrifuge (max 12,000 × g)

4.

Transfer the reaction product into the cuvette or other chosen format for the spectrophotometric analysis

5.

Detect the absorbance (**404 nm** for yellow, **517 nm** for red, **595 nm** for blue and **630 nm** forgreen substrate)

More information can be found on our webpage: **www.glycospot.dk**

GlycoSpot

Enzyme screening standard

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