

AgraQuant® Mycotoxin ELISA Test Kit

Romer Labs® has developed a rapid, quantitative enzyme-linked immunosorbent assay (ELISA) for the analysis of mycotoxins in grains, nuts, cereals and other commodities including animal feeds.

Each kit comes complete with 5 to 6 standards, 48 or 96 antibody-coated microwells, 48 or 96 color-coded dilution microwells, conjugate, substrate and stop solution.



AgraQuant® Mycotoxin Kit Performance

- Accurate – Results are comparable with published HPLC method
- Highly Sensitive
- Reproducible – Consistent results obtained in intra- and inter-laboratory settings

	Item No.	Quantitation Range	Limit of Detection
AgraQuant® Total Aflatoxin	COKAQ1100	1 – 20 ppb	1 ppb
AgraQuant® Total Aflatoxin	COKAQ1000	4 – 40 ppb	3 ppb
AgraQuant® Aflatoxin B1	COKAQ8000	2 – 50 ppb	2 ppb
AgraQuant® Ochratoxin	COKAQ2000	2 – 40 ppb	2 ppb
AgraQuant® Total Fumonisin	COKAQ3000	0.25 – 5.0 ppm	0.2 ppm
AgraQuant® DON	COKAQ4000	0.25 – 5.0 ppm	0.2 ppm
AgraQuant® ZON Plus	COKAQ5100	25 – 1000 ppb	20 ppb
AgraQuant® T-2 Toxin	COKAQ6000	25 – 500 ppb	10 ppb
AgraQuant® T-2/HT-2 Toxin	COKAQ6100	37 – 500 ppb	29 ppb
AgraQuant® Aflatoxin M1	COKAQ7100	25 – 500 ppt	18 ppt
AgraQuant® Aflatoxin M1 Plus	COKAQ7200	100 – 2000 ppt	89 ppt

BENEFITS:

- Rapid – 10 to 20 minutes total incubation time (except Aflatoxin M1 kits)
- Stable – up to 12 months shelf life
- Easy – Simple sample extraction and no clean up steps required
- Cost-effective – 48 or 96 breakaway microwell format; minimizes waste and maximizes value
- Convenient – Up to 30 minutes reading time after stopping the reaction



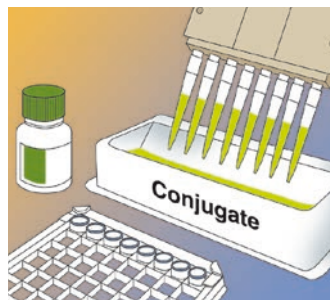
AgraQuant® Mycotoxin ELISA Test Kit

MYCOTOXINS

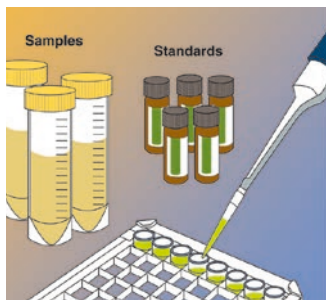
Procedure for AgraQuant® Mycotoxin ELISA

IMPORTANT: Please read kit insert before running the test.

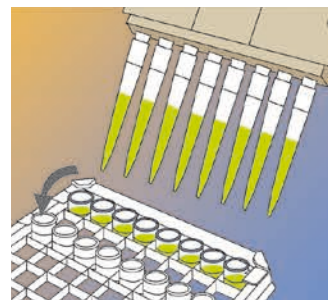
Assay Procedure:



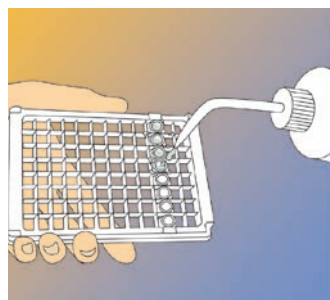
1) Add 200 μ L conjugate into each color-coded dilution well.



2) Add 100 μ L standards or samples to the conjugate.



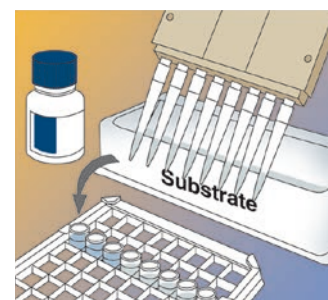
3) Mix well. Transfer 100 μ L content to antibody-coated wells. Incubate for 5 – 60 minutes.



4) Discard contents from the wells and wash wells with deionized water or buffer solution (5x).



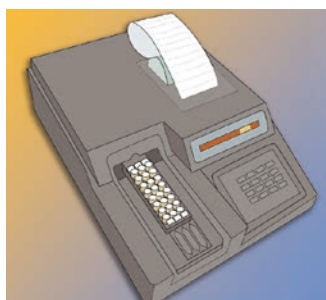
5) Tap dry the wells on absorbent paper towel.



6) Add 100 μ L substrate into each well. Incubate for 5 – 20 minutes.



7) Add 100 μ L stop solution into each well.



8) Analyze results using an ELISA reader with 450 nm filter.

For further information, please contact:

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