# AmplideX<sup>®</sup> PCR/CE CFTR Kit\*

# Providing the broadest coverage of the U.S. population<sup>+</sup> of any commercially available, targeted *CFTR* testing kit

The AmplideX\* PCR/CE *CFTR* Kit\* brings the simplicity and scalability of AmplideX technology to *CFTR* variant detection. Offering the streamlined detection of 67 pathogenic variants, and covering approximately 93%<sup>†</sup> of the U.S. population, the assay provides the broadest coverage of any commercially available, targeted testing kit. The assay is optimized for use on widely established laboratory equipment and delivers genotype results from DNA in under five hours.

# **REDUCED COMPLEXITY**

- Broad coverage<sup>†</sup> provided using only PCR/CE workflow
- Similar testing process to AmplideX PCR/CE *FMR1\** and *SMN1/2* Plus\* kits eases implementation
- Streamlined data analysis via AmplideX Reporter software

#### **OPTIMIZED WORKFLOW**

- Easy-to-use workflow with fewer hands on steps than most competitor assays
- Flexible use on widely installed CE equipment
- <5 hrs from DNA to data

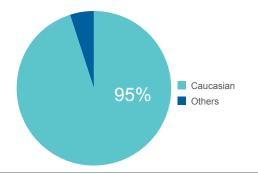
#### **QUALITY RESULTS**

- Built on the latest prevalence data to provide the best coverage<sup>†</sup> for all U.S. ethnicities
- Detects complex yet key *CFTR* mutations (CNVs, STRs, SNPs, INDELs) and resolves zygosity
- Excellent concordance with other methods





Figure 1. AmplideX PCR/CE CFTR Kit\* workflow.



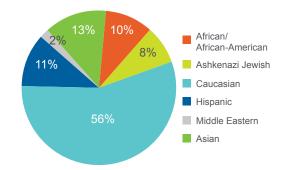


Figure 2A. Ethnic diversity of 89K patients in CFTR2 database.

Figure 2B. Ethnic diversity of 115K subjects in U.S. study.<sup>†</sup>

Commercial Kit Manufacturer/Panel	Full Product Name	# of Variants Detected	% Coverage, CFTR2	% Coverage, U.S.	% Coverage Difference, U.S.
Asuragen	AmplideX PCR/CE CFTR Kit	67	92.1%	93.0%	-
Luminex 97	xTAG Cystic Fibrosis (CFTR) 97 kit (Custom)	97	92.2.%	88.2%	4.8%
Illumina	MiSeqDx Cystic Fibrosis 139 Variant Assay	139	94.3%	87.7%	5.3%
Agena	iPLEXPro CFTR Panel	74	91.5%	86.9%	6.1%
Elucigene	CF-EU2v1	50	90.5%	86.2%	6.8%
Luminex 60	xTAG Cystic Fibrosis ( <i>CFTR</i> ) 60 kit v2	60	91.2%	86.1%	6.9%
Luminex 39	xTAG Cystic Fibrosis ( <i>CFTR</i> ) 39 kit v2	39	89.1%	80.7%	12.3%

Table 1. Percent coverage of variants detected by commercially available kits based on frequencies observed in the CFTR2 database\*\* and the U.S. population.

76 Cell Lines from Coriell Repository					
Sample Agreement with Reference Assay		Sa WT/WT	Overall Sample Agreement		
	WT/WT	22	MUT/MUT 0	MUT/WT	22/22 (100%)
۷/CE As	MUT/MUT	0	24	0	24/24 (100%)
CFTR PCR/CE Assay	MUT/WT	0	0	27	27/27 (100%)

**Table 2A.** Mutations in Cell Lines Detected by Prototype PCR/CECFTR Kit and Sanger Sequencing.

102 Clinical	Samples /	Across 3 Sites

Sample Agreement with Reference Assay		xTAG® Cystic Fibrosis (CFTR) 60 kit v2 or Sanger Sequencing WT/WT MUT/MUT MUT/WT			Overall Sample Agreement
say	WT/WT	34	0	0	34/34 (100%)
۲/CE As	MUT/MUT	0	19	0	19/19 (100%)
CFTR PCR/CE Assay	MUT/WT	0	0	49	49/49 (100%)

**Table 2B.** Mutations in Clinical Samples Detected by Prototype PCR/CE CFTR Kit and Either xTAG® Cystic Fibrosis (CFTR) 60 Kit v2 orSanger Sequencing.

## **ORDERING INFORMATION**

Part Number	Product Description	Number of Reactions
A00076	AmplideX PCR/CE CFTR Kit	50
A00077	AmplideX PCR/CE <i>CFTR</i> Kit	100

## For more information please contact asuragen.com

\*For Research Use Only. Not for use in diagnostic procedures. \*\*The Clinical and Functional Translation of CFTR (CFTR2); available at http://cftr2.org <sup>†</sup>Beauchamp, et al. Genet Med. 2019 Nov;21(11):2569-2576.



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