




# CFTR interactome mapping using the mammalian membrane two-hybrid high-throughput screening system

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## Abstract

Cystic Fibrosis Transmembrane Conductance Regulator (CFTR) is a chloride and bicarbonate channel in secretory epithelia with a critical role in maintaining fluid homeostasis. Mutations in CFTR are associated with Cystic Fibrosis (CF), the most common lethal autosomal recessive disorder in Caucasians. While remarkable treatment advances have been made recently in the form of modulator drugs directly rescuing CFTR dysfunction, there is still considerable scope for improvement of therapeutic effectiveness. Here, we report the application of a high-throughput screening variant of the Mammalian Membrane Two-Hybrid (MaMTH-HTS) to map the protein–protein interactions of wild-type (wt) and mutant CFTR (F508del), in an effort to better understand CF cellular effects and identify new drug targets for patient-specific treatments. Combined with functional validation in multiple disease models, we have uncovered candidate proteins with potential roles in CFTR function/CF pathophysiology, including Fibrinogen Like 2 (FGL2),

which we demonstrate in patient-derived intestinal organoids has a significant effect on CFTR functional expression.

**Keywords** cystic fibrosis; high-throughput screening; integrative computational biology; interactome; mammalian membrane two-hybrid

**Subject Categories** Membranes & Trafficking; Methods & Resources

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## Introduction

Thought to have originated in Europe (Morrall *et al.*, 1994) and spread to other regions with heterozygote advantage (Knudson *et al.*, 1967; Gabriel *et al.*, 1994; Schroeder *et al.*, 1995; Molinski, 2016), Cystic Fibrosis (CF) is the most common autosomal recessive disease in the Caucasian population, occurring in approximately 1/

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3,500 births (Mehta *et al*, 2010) and affecting over 90,000 individuals worldwide. CF affects multiple organs with epithelial tissues including the lungs, pancreas, liver, intestines and the reproductive tract, resulting in significant life-shortening and physical challenges. The most prevalent symptoms develop in the airways of CF patients with accretion of viscous mucus causing obstruction and an increased susceptibility to bacterial infection and neutrophilic inflammation. Development of CF-related diabetes, pancreatic insufficiency, intestinal obstruction and liver disease are also present in individuals affected by CF, albeit with considerable phenotypic variation. Despite the recent approvals of drug therapies that pave the way for personalized medicine treatments of CF with possible disease-modifying effects (Fajac & Wainwright, 2017; Amaral *et al*, 2019; Lopes-Pacheco, 2020), pulmonary function of CF patients progressively decreases due to mucus obstruction/bronchiectasis, recurrent bacterial infection and subsequent inflammation causing high morbidity and mortality (Cutting, 2014). In addition, variation in survival and disease severity is extremely large and this heterogeneity is mostly attributed to differences in environmental factors and patients' genetic background (i.e. modifier genes).

Pathophysiology of CF is caused by mutations in the Cystic Fibrosis Transmembrane Conductance Regulator (*CFTR*) (Riordan *et al*, 1989) gene, whose protein product is localized primarily in the apical membrane of secretory epithelial cells. The *CFTR* protein, regulated by PKA-phosphorylation and nucleotide binding, functions as an anion channel that mediates the flux of chloride ( $\text{Cl}^-$ ) and bicarbonate ( $\text{HCO}_3^-$ ) ions (Riordan, 2008), which provide the driving force for fluid transport and maintenance of ion and fluid homeostasis. As its name suggests, *CFTR* also acts as a regulator influencing the activity of a variety of other channels and transporters and consequently, fluxes of other ions (Li & Naren, 2010). The protein is a member of the human ATP-Binding Cassette (ABC) transporter superfamily and thus has a conserved core structure consisting of two membrane-spanning domains (MSD1 & MSD2) forming the pore of the channel, and two cytosolic nucleotide binding domains (NBD1 & NBD2) for ATP binding/hydrolysis. However, it also contains a unique regulatory domain (RD) that controls the channel's activity based on its phosphorylation state. Notably, *CFTR*'s involvement in promoting the transmembrane flow of anions down their electrochemical gradient (thereby regulating electrolyte and water homeostasis across epithelial cell membranes) is distinct from the activity of other ABC membrane transporters, which direct the movement of organic substrates across cell membranes (Riordan *et al*, 1989; Bear *et al*, 1992; Molinski, 2016).

Being a twelve-pass transmembrane protein, *CFTR* requires a multitude of accessory proteins and cooperative interdomain assembly to progress through a sequence of translational folding events (Riordan, 2005; Wang *et al*, 2006; Du & Lukacs, 2009; Pitzonzo *et al*, 2009). As a result, successful folding and processing of the 1,480 amino acid-long *CFTR* polypeptidic chain into a functional channel is a naturally inefficient process and a significant amount of wild-type (wt) *CFTR* is rapidly degraded before reaching the plasma membrane (PM) (Ward & Kopito, 1994). This intrinsic property of *CFTR* biogenesis leads to the unsurprising observation that mutations which affect this process, or the quantity/quality of *CFTR* protein, are responsible for the majority of CF cases. Such mutations are ultimately associated with the loss of *CFTR* function, most prominently at the surface of the airways, which leads to depletion

of the airway surface fluid and accumulation of dehydrated and persistent mucus, the hallmark feature of CF (Boucher, 2007).

There are currently over 2,100 variants in the *CFTR* gene recorded in the CF Mutation Database (<http://www.genet.sickkids.on.ca/cftr/StatisticsPage.html>), however F508del is by far the predominant mutation, present in ~80% of all individuals with CF worldwide (Cystic Fibrosis Foundation, 2020; Zolin *et al*, 2020). This mutation causes *CFTR* misfolding and subsequent retention in the endoplasmic reticulum (ER) from where it is targeted for degradation (Cheng *et al*, 1990). Interestingly, proper folding and function of this mutant protein can be partially restored by low temperature (Denning *et al*, 1992), which revealed that post-translational processes play important roles in the manifestation of CF (Hutt *et al*, 2010). However, even upon rescue, F508del-*CFTR* displays altered channel activity and reduced protein stability at the cell surface (Okiyoneda *et al*, 2010; Lukacs & Verkman, 2012), suggesting that there are many interacting proteins involved in its cell surface tethering and functionality (Riordan, 2005). Although the genetic cause of CF by *CFTR* mutations has been well studied in the past (Zielenki & Tsui, 1995), considerably less is known about the cellular protein environment of *CFTR* protein in CF.

In an effort to better understand *CFTR* protein regulation and biology, as well as the cellular changes associated with the F508del mutation and CF, we set out to map the shared and differential protein interactions of wt- and the F508del-*CFTR* mutant in mammalian cells. To accomplish this, we applied a newly modified, high-throughput screening (HTS) variant of our previously reported Mammalian Membrane Two-Hybrid (MaMTH) technology (Petschnigg *et al*, 2014, 2017; Saraon *et al*, 2017, 2020, 2021; Yao *et al*, 2017; Aboulizadeh *et al*, 2021) called MaMTH-HTS, to screen the Human Open Reading Frame (ORF)eome V8.1 collection (Yang *et al*, 2011). Subsequent computational and functional analysis of our interactomes identified several interactors with potentially significant roles in *CFTR* regulation and trafficking, furthering our understanding of *CFTR*/CF biology and identifying potential new avenues for future exploration.

## Results

### Design of the mammalian membrane two-hybrid high-throughput screening (MaMTH-HTS) platform

The MaMTH technology is a split ubiquitin-based assay for measurement of the protein-protein interactions (PPIs) of full-length integral membrane proteins directly in the context of living mammalian cells (Johnsson & Varshavsky, 1994; Stagljar *et al*, 1998; Scheper *et al*, 2003; Paumi *et al*, 2008; Petschnigg *et al*, 2014, 2017; Sokolina *et al*, 2017; Yao *et al*, 2017) (Appendix Fig S1). It is highly sensitive and capable of detecting subtle alterations in PPIs in response to environmental factors and mutation state, making it an ideal tool for monitoring dynamic changes between the interactomes of wt- and F508del-*CFTR* proteins in their native plasma membrane environment (Petschnigg *et al*, 2014, 2017; Yao *et al*, 2017).

In its original format, MaMTH functions as an "array-based" assay, whereby a target "bait" protein of interest is tested against candidate interacting "prey" protein partners individually in the wells of microtitre plates (Petschnigg *et al*, 2014, 2017;

Aboulizadeh *et al*, 2021; Grozavu *et al*, 2022) While suitable for medium throughput screens of small cDNA libraries (i.e. hundreds) of candidate interactors, the approach becomes less efficient as library size increases. In order to perform our CFTR MaMTH screens against an ORF library representing a significant portion of human genome (generated from the Human ORFeome V8.1, a large and relatively comprehensive collection consisting of ~12,000 sequence verified ORFs), we therefore needed to modify MaMTH to increase its throughput.

To accomplish this, we assembled a new MaMTH-HTS workflow. In this approach, we employ fluorescence, instead of luminescence, as a readout for PPIs, using an eGFP reporter in place of luciferase (Petschnigg *et al*, 2014) (Appendix Fig S1). This allows for easy enrichment of cells expressing interacting protein pairs using Fluorescence-Activated Cell Sorting (FACS). To help minimize background, tagBFP (a monomeric blue fluorescent protein) and mCherry (a monomeric red fluorescent protein) tags are also attached to MaMTH baits and preys, respectively, via a P2A linker. In this way, the cells can first be FACS-sorted based on the proper expression of “prey” protein of interest (indicated by red fluorescence) and “bait” (indicated by blue fluorescence) prior to selection of the cells containing interacting bait-prey pairs (indicated by green fluorescence). Sorted populations are then expanded, followed by genomic DNA isolation, PCR amplification of prey ORFs and deep sequencing, to determine which “prey” ORFs are enriched in the final population, providing a list of candidate protein interactors. While still new and awaiting substantial benchmarking, in its current format our MaMTH-HTS platform presents a relatively fast and simple means of screening large numbers of interactions involving full-length membrane proteins directly in the environment of living mammalian cells. This potentially allows for detection of interactions that may be missed using systems with more artificial screening conditions or that require the use of only the soluble regions of membrane proteins. A schematic overview of the MaMTH-HTS platform can be found in Fig 1.

### MaMTH-HTS Screening of wt- and F508del-CFTR

In order to map wt- and F508del-CFTR interactomes using the MaMTH-HTS system, we first generated CFTR “bait” constructs and an appropriately tagged “prey” library for high-throughput PPI screening. “Baits” were prepared by cloning wt- and F508del-CFTR ORFs into MaMTH plasmid expressing (under the control of a tetracycline-inducible promoter) the full-length proteins as fusions to a C-terminal bait tag consisting of the C-terminus of ubiquitin (Cub) and a GAL4 transcription factor linked by a P2A sequence to tagBFP (Appendix Fig S2A).

For ORF library generation, we cloned the entire Human ORFeome V8.1 (Yang *et al*, 2011), consisting of ~12,000 ORFs divided into 33 distinct pools, into a CRISPR-compatible MaMTH-HTS “prey” vector. This vector allowed for tetracycline-controlled expression of proteins fused at their N-terminus to the N-terminal fragment of ubiquitin (Nub) and at their C-terminus to a P2A-mCherry tag (Appendix Fig S2B). Following cloning, ORF representation in pools was verified by deep sequencing and individual pools were then stably integrated (using a variant of the *S. pyogenes* CRISPR/Cas9 system (Ran *et al*, 2013) developed in our lab) into the AAVS site of MaMTH HEK293 reporter cells stably expressing

eGFP under the control of a 5XGAL4-UAS. To determine representation of ORFs in these cell lines, integrated preys were amplified by PCR using MaMTH prey-cassette specific primers and deep sequenced, verifying the presence of 10,228 ORFs in the final cell library population (Dataset EV1).

For screens, all 33 ORF “prey” cell pools were combined into a single “master pool” and transfected with either wt- or F508del-CFTR MaMTH bait plasmid. Note that we previously demonstrated that transient transfection of MaMTH CFTR baits does not affect their expected localization (Appendix Fig S3), making this approach ideal for streamlining the screening process. The cells were then grown for 2 days in the presence of 0.5 µg/ml of tetracycline to allow for significant expression of baits, preys and associated fluorophores/fluorophore reporter. Following growth, the cells were harvested and sorted by flow cytometry using a BD FACS Melody (BD Biosciences). Sorting was performed sequentially based on the presence of mCherry fluorescence (indicating cellular expression of prey), tagBFP fluorescence (indicating cellular expression of bait) and finally eGFP fluorescence (indicating a bait/prey interaction). The collected cells were resuspended in appropriate media and grown/expanded for 10 days to increase overall cell number. Genomic DNA was then harvested from the cells and prey ORFs amplified using two distinct sets of MaMTH cassette-specific primers (Table EV1). Purified PCR products were then prepared and deep sequenced separately, providing us with two technical replicates and resulting data analysed using in house bioinformatics software to assemble a preliminary list of putative interactors (Dataset EV2).

### Identification of the wt- and F508del-CFTR Interactomes

Our initial mapping identified a total of 494 unique candidate PPIs across both wt- and F508del-CFTR, from which we were able to flag and filter out 47 as “frequent fliers” based on repeated occurrence across MaMTH-HTS screens of 9 control “baits”, belonging to distinct classes of membrane proteins, as determined using the rank product method (Breitling *et al*, 2004) (Datasets EV2 and EV3).

Of the remaining 447 candidate interactions, a total of 224 interacted with wt-CFTR, with a robust overlap of 209 (~93%) between both technical replicates (Appendix Fig S4A, Dataset EV2). Similarly, a total of 269 unique ORFs interacted with F508del-CFTR, with an overlap of 250 (~93%) between technical replicates (Appendix Fig S4B, Dataset EV2). Collectively, these overlaps demonstrate that our data do not appear to be greatly influenced by technical issues such as primer-specific PCR bias or inconsistencies across sequencing runs. Comparison of the wt- and F508del-CFTR interactomes revealed an overlap of 46 interactors (~10.2%), with wt- and F508del-CFTR having 178 and 223 unique interactions, respectively (Appendix Fig S4C, Dataset EV2). A complete, annotated version of our interactome, sorted by GO Biological process and showing shared/differential interactions and interaction novelty based on functional and predicted PPIs reported in the IID database (Kotlyar *et al*, 2019) is provided in Fig 2 (with additional annotation details provided in Dataset EV4). Notably, our interactors cover a broad range of functions, with some of the largest groups corresponding to signalling, locomotion and, intriguingly, uncharacterized processes. A more focused sub-interactome highlighting previously reported interactions is also provided in Appendix Fig S5.

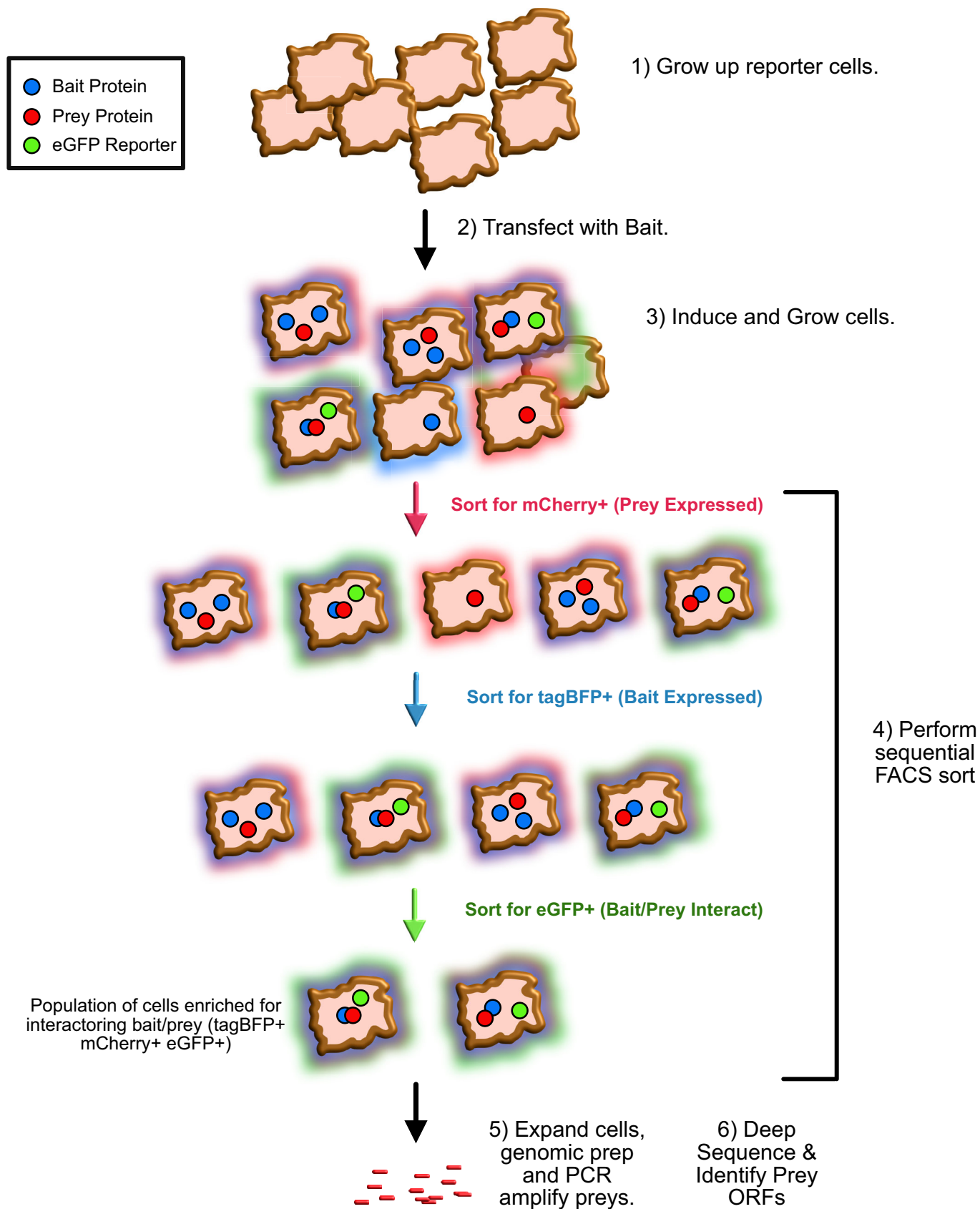
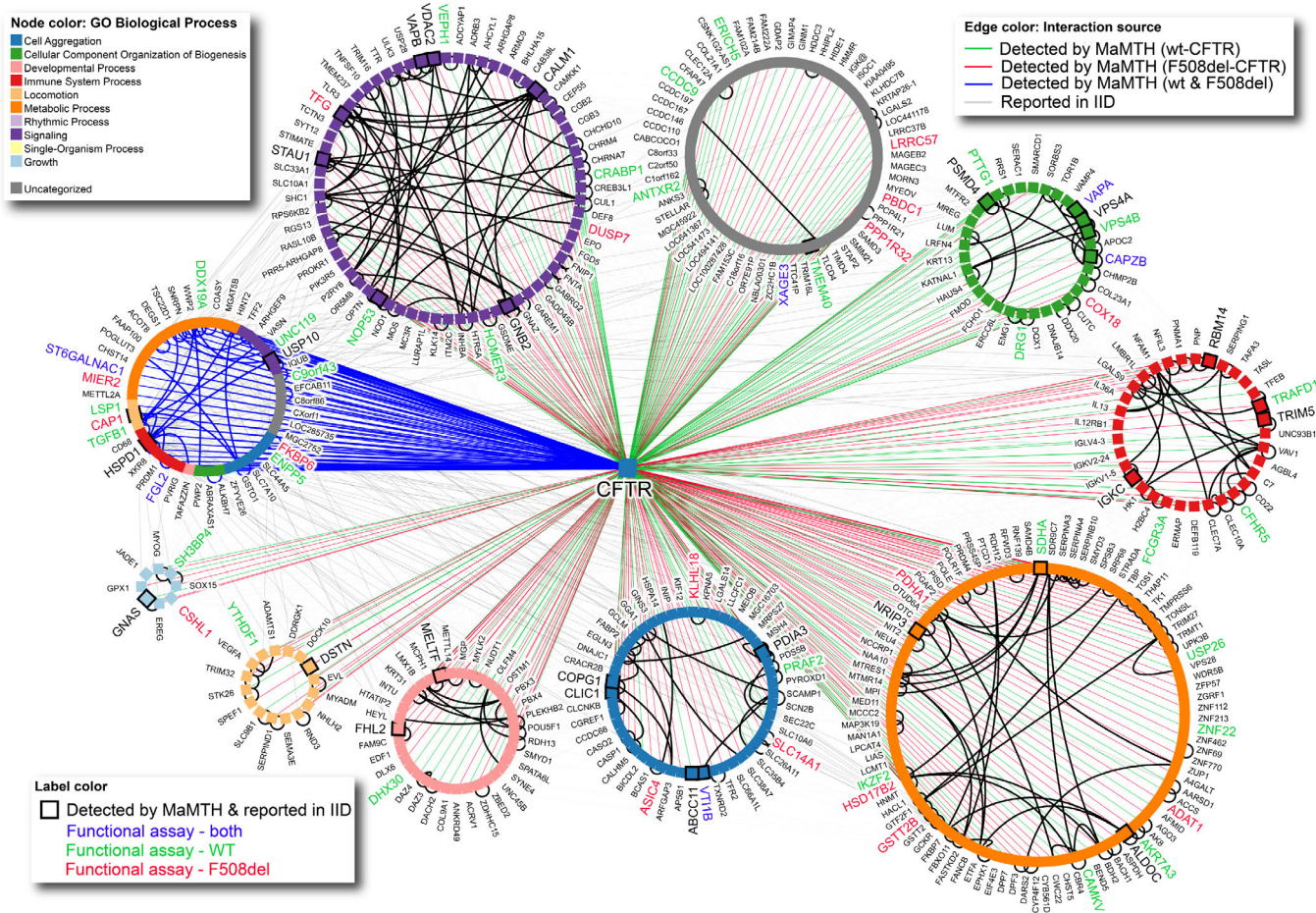


Figure 1.

**Figure 1. The MaMTH-HTS workflow.**

1) Grow up cell lines containing stably integrated prey and reporter. 2) Transfect cells with Bait of Interest. 3) Induce the cells with tetracycline to express Bait and Prey. Continue to grow the cells for 48 h. At this point, we will have a mixed population containing the cells expressing combinations of Bait (along with tagBFP/blue fluorescence) and Prey (along with mCherry/red fluorescence). The cells expressing interacting Bait/Prey pairs will induce eGFP reporter and have additional green fluorescence. 4) Carry out sequential FACS sort, first selecting the cells with red fluorescence (indicating proper Prey expression), then blue fluorescence (indicating proper Bait expression) and finally green fluorescence (indicating Bait/Prey interaction). 5) Expand/grow the cells to increase overall cell number, isolate genomic DNA and PCR amplify using prey cassette specific primers. 6) Deep sequence amplified PCR products to identify prey ORFs enriched in population (indicating potential interactors).



**Figure 2. MaMTH-HTS interactome for wt- and F508del-CFTR.**

Annotated MaMTH-HTS interactome for wt- and F508del-CFTR (green, red and blue edges) expanded with known experimental and predicted interactions from IID v 2021-05 (black/grey edges). Interactions observed in MaMTH-HTS to be shared by both wt and F508del CFTR are indicated by blue edges, while those observed with only wt or F508del are indicated by green and red edges, respectively. Node colour represents Gene Ontology Biological process. Black square outline and larger node label indicate CFTR interactions overlapping with IID. Node label colouring highlights interactors functionally validated in this work.

**Evaluation of CFTR Interactomes using traditional MaMTH**

While MaMTH-HTS is based on the same general principles as our originally published array-based MaMTH assay (Petschnigg et al, 2014, 2017; Aboualizadeh et al, 2021), the important changes introduced into the format (including the use of a multi-step, fluorescence-based selection/reporter system, stably versus transiently expressed

preys, pooled screening that involves testing thousands of potential interactions simultaneously in a mixed population and a deep sequencing readout) suggest that the systems may behave differently in terms of interaction mapping. To further explore this, we re-tested a large subset of 240 identified interactions using the original MaMTH assay with transiently expressed bait-prey pairs in HEK293 cells containing a *Gussia princeps* luciferase reporter system.

Each interaction was tested in three independent trials with three technical replicates. From a total of 138 wt-CFTR interactions, 32 (~23%) produced a statistically significant luminescence signal (with respect to background) in our original MaMTH (Dataset EV5 and Appendix Fig S6A). The proportion of F508del-CFTR interactions was slightly lower, with 26 out of 154 tested (~17%) testing positive (Dataset EV5 and Appendix Fig S6B). These results confirm that while both MaMTH versions can robustly detect certain interactions, MaMTH-HTS was able to detect a considerable number of PPIs not identified by the originally described MaMTH system (Petschnigg *et al*, 2014).

Similarly, our traditional MaMTH format was able to detect interactions missed using our HTS format. Specifically, array-based counter-screening of 86 interactors, identified in MaMTH-HTS as uniquely interacting with wt-CFTR, against F508del-CFTR bait found that 18 (~21%) also interacted with the F508del mutant (Dataset EV5 and Appendix Fig S6C). Conversely, testing of 102 interactors, identified in MaMTH-HTS as uniquely interacting with F508del-CFTR, against wt-CFTR bait revealed that 23 (~23%) also interacted with wt-CFTR (Dataset EV5 and Appendix Fig S6D).

Collectively, while some of these discrepancies are likely attributable to variation/technical issues during screening, our results still suggest that the platforms have highly complementary behaviours suitable for detection of distinct types of interactions.

#### Fluorescence-based assay to monitor CFTR channel activity

To explore the biological relevance of the identified PPIs, we performed a previously reported membrane potential assay (Ahmadi *et al*, 2017) using FLIPR<sup>®</sup> dye to monitor the ability of a subset of 26 interactors, selected based on performance in MaMTH-HTS and traditional MaMTH, to modulate the activity of CFTR baits (Dataset EV6).

For these experiments, each interactor was transiently expressed in HEK 293 cells stably expressing either wt- or F508del-CFTR bait. Activation of CFTR channels in these cells can be detected as membrane depolarization using the FLIPR<sup>®</sup> membrane potential-sensitive dye after stimulation by forskolin (Fsk) and ivacaftor (VX-770), while CFTR Inh<sub>172</sub> inhibitor can impede this fluorescence response (Ahmadi *et al*, 2017). For the wt-CFTR measurements, we compared the maximum Fsk response before and after stimulation. For F508del-CFTR, since it does not produce a detectable signal upon addition of Fsk alone (Fsk plus a potentiator drug such as VX-770 or genistein are required), we compared the fully stimulated response immediately before the addition of CFTR Inh<sub>172</sub> inhibitor to the minimum response reached at the end of each measurement cycle. Using this approach, we detected a total of 15 interactors (of 26 tested in total) whose transient expression appeared to significantly affect wt- and/or F508del-CFTR functional channel activity in response to Fsk and VX-770 potentiator stimulation compared to that of the dimethyl sulfoxide (DMSO) control. Specifically, 12 interactors increased F508del-CFTR channel activity (Fig 3A) and 6 interactors increased wt-CFTR activity (Fig 3B). Of these, 3 interactors (FGL2-Fibrinogen Like 2, VTI1B-Vesicle transport through interaction with t-SNAREs homolog 1B and CAPZB-F-actin-capping protein subunit beta) were “shared” between the wt-CFTR and F508del-CFTR, leading to an increase in the channel activity of both forms of CFTR (Fig 3A and B). Assay

traces for the experiments with these interactors can be found in Appendix Fig S7.

#### Traffic-based siRNA high-content microscopy screen of top MaMTH-HTS interactors

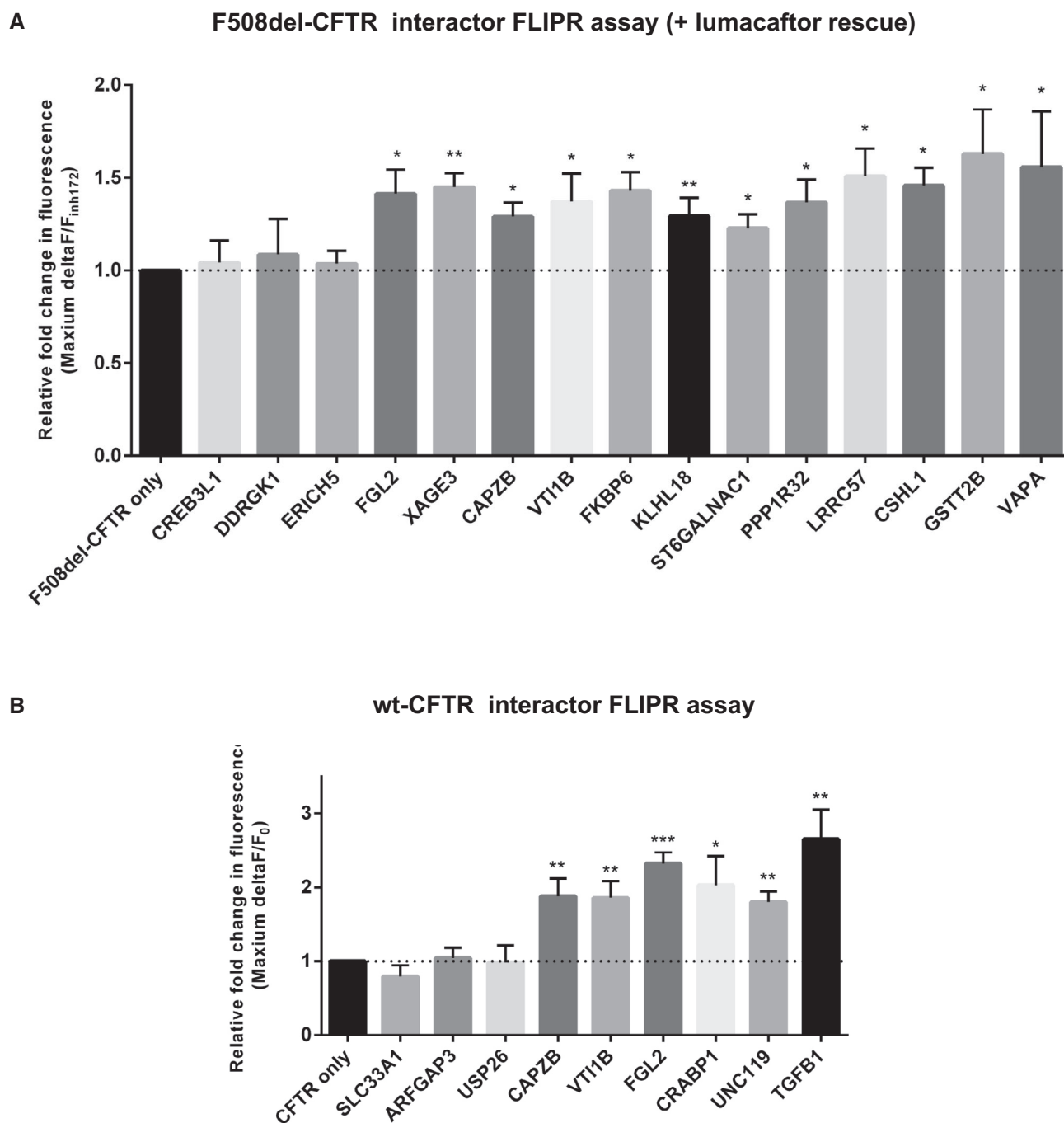
To determine whether interaction partners identified via MaMTH-HTS regulate CFTR (wt- and F508del-) plasma membrane expression, we performed a high-content siRNA-based microscopy assay (Botelho *et al*, 2015) to measure CFTR trafficking upon knocking down the genes encoding for selected interactors. Specifically, a total of 208 genes expressing interaction partners identified with MaMTH-HTS were targeted (using 414 siRNAs, 2 for most gene targets) and the effects of their knock-down on an mCherry-Flag-CFTR traffic reporter expressed in CF bronchial epithelial (CFBE) cells were assessed (Dataset EV7).

By taking the cells treated with a non-targeting siRNA (siNeg1) as a reference and considering the siRNAs generating a significant increase or decrease in PM localization as those with Z-score > +1 or Z-score < -1, respectively, (a strategy successfully followed in several previous related screening assays (Botelho *et al*, 2015; Lérias *et al*, 2018; Santos *et al*, 2020)), we identified 25 genes whose knockdown by at least one siRNA leads to modulation of F508del-CFTR PM expression (Fig 4A, Dataset EV7). Of these, 17 rescued F508del-CFTR PM expression, while 7 worsened its PM trafficking (Fig 4A, Dataset EV7). One gene, MIER2 (Mesoderm induction early response protein 2), produced an ambiguous result, with an increase or decrease in F508del-CFTR PM level observed depending on the siRNA used (Fig 4A, Dataset EV7).

Using the same thresholds, we also identified 36 genes whose siRNA-mediated knockdown by at least one siRNA affected wt-CFTR PM expression. These included 10 genes resulting in an increase in wt-CFTR PM expression and 24 that lead to a decrease (Fig 4B, Dataset EV7). There were also two interactors, SDHA1 (Succinate dehydrogenase [ubiquinone] flavoprotein subunit, mitochondrial) and ZNF22 (Zinc finger protein 22), which produced an ambiguous result similar to that observed with MIER2 siRNA for F508del-CFTR, leading to an increase or decrease in wt-CFTR PM trafficking depending on the siRNA used (Fig 4B, Dataset EV7).

Interestingly, comparison of results revealed that the siRNA targeting one particular gene, CAPZB, led to a significant decrease in trafficking of both wt- and F508del-CFTR (Fig 4A and B and Dataset EV7), highlighting a potentially important role in general localization of CFTR channels to the PM. Disruption of some candidate genes also lead to opposing effects on the CFTR variants. For instance, siRNAs targeting DHX30 (ATP-dependent RNA helicase DHX30), ACCN4 (Acid-sensing ion channel 4), CFHR5 (Complement factor H-related protein 5), YTHDF1 (YTH domain-containing family protein 1) and CSHL1 (Chorionic somatomammotropin hormone-like 1) all lead to a decrease in wt-CFTR trafficking, while that of F508del-CFTR was increased (Fig 4A and B and Dataset EV7). These results suggest a specific role of these genes in intracellular retention of F508del-CFTR, which is alleviated with the knockdown.

Representative images of wt-CFTR localization and F508del-CFTR trafficking rescued in response to siRNA hits (alongside associated controls) are provided in Appendix Fig S8.

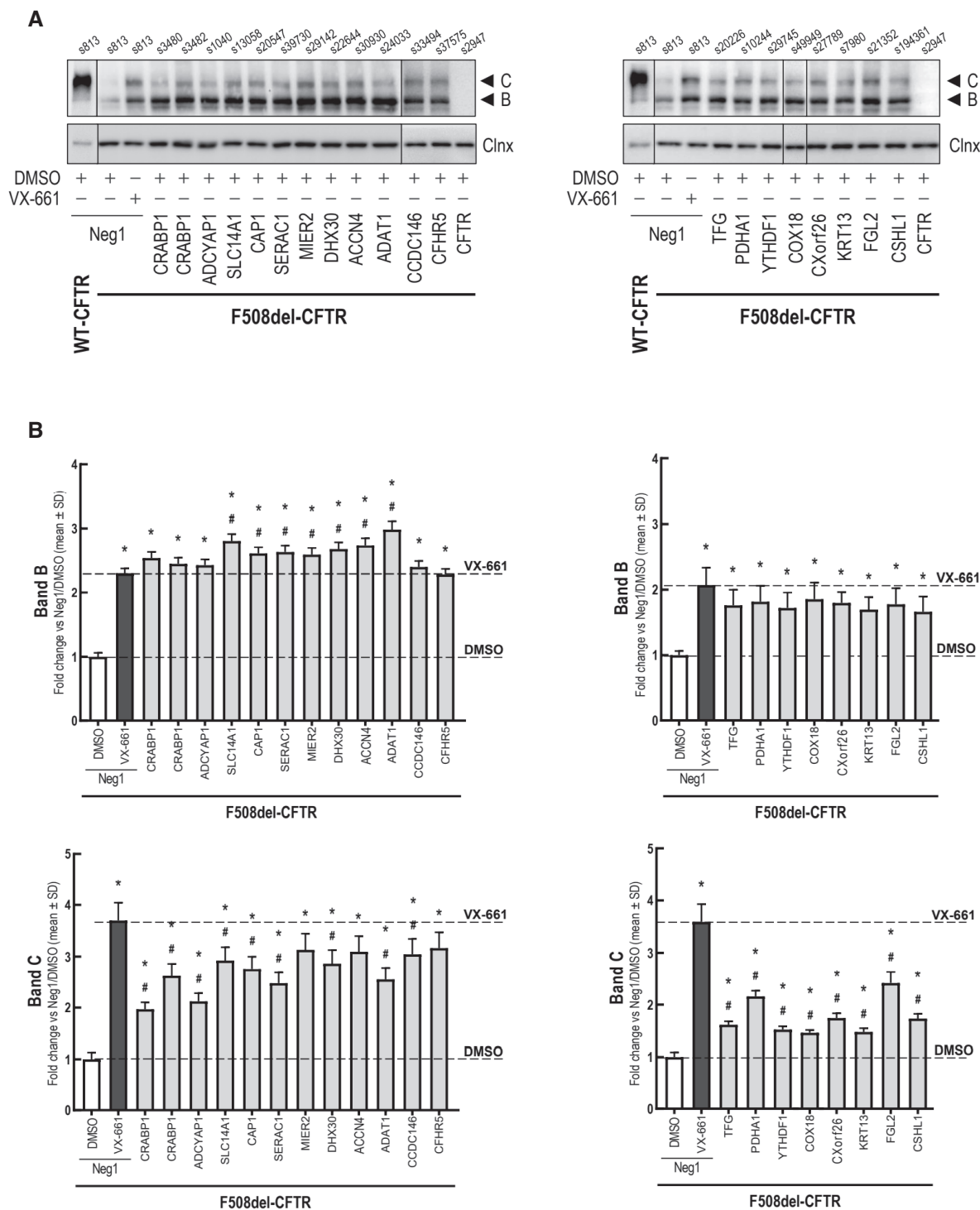


**Figure 3. FLIPR assay results for interactors positively affecting CFTR  $\text{Cl}^-$  channel activity.**

A, B Assays were performed in HEK293 cells for both A) F508del-CFTR and B) wt-CFTR. For the wild-type measurements, we compared the maximum Fsk response (Maximum  $F/F_0$ ; where  $F_0$  represents baseline RFU right before the stimulation). For the F508del-CFTR measurements, since this mutant does not respond well to Fsk alone (Fsk + VX-770 potentiator is required to obtain a detectable signal), we compared the CFTR inh172 (inhibitor) response ( $F_1/F_{\text{CFTRinh172}}$ ; where  $F_1$  represents peak RFU immediately before inhibitor addition and " $F_{\text{CFTRinh172}}$ " is the minimum RFU reached at the end of the CFTR inh172 measurement cycle). For each of wt-CFTR and F508del-CFTR, data for three proteins not detected as interactors (first three bars following CFTR only) are shown as negative controls. Fluorescence signals were normalized to the CFTR-only control and the statistical significance of differences assessed by unpaired t-test (\* $P < 0.05$ , \*\* $P < 0.01$  and \*\*\* $P < 0.001$ ). Samples run in  $n = 3$  replicates, and all values are mean  $\pm$  SD.







involved in retaining F508del-CFTR in the ER or reducing CFTR biogenesis efficacy. Knockdown of FGL2 also led to an increase in the amount of the fully glycosylated form of CFTR, indicating an

improvement in CFTR processing. This result is particularly intriguing, as overexpression of the protein was observed to lead to an increase of CFTR channel activity in our FLIPR assay (Fig 3),

suggesting a potentially complex or cell line-dependent role of the protein in CFTR regulation.

### Orthogonal validation using traditional co-immunoprecipitation (co-IP)

To further validate the quality of our interactome, we performed tag-based co-IP assays on a selected subset of 30 CFTR-interacting proteins identified from the MaMTH-HTS pipeline (Table EV2). For these experiments, we used transiently expressed, V5-tagged wt-CFTR alongside transiently expressed, FLAG-tagged preys in HEK293 cells. Of the 30 interactions tested, we were able to successfully confirm 17 (56%), a substantial proportion providing support for the biological relevance of our identified interactions (Appendix Fig S9A). Based upon our chloride conductance (Fig 3, Dataset EV6) and trafficking validation results (Fig 5), we also specifically examined the interaction of FGL2 with both wt- and F508del-CFTR. Notably, we successfully confirmed the interaction by Co-IP for both protein variants, providing further evidence that FGL2 is an important functional interactor of CFTR (Appendix Fig S9B).

Control blots for the co-IP assays presented in Appendix Fig S9A and B, performed using unrelated GAPDH antibody, are presented in Appendix Fig S9C.

### Knockdown of FGL2 using lentivirus-induced shRNA in patient-derived organoid cultures

In order to further explore the importance of FGL2 in CFTR biogenesis, traffic and function we performed lentivirus-mediated shRNA knockdown of the FGL2 and CFTR genes in patient-derived intestinal organoids and examined their effects using two functional readouts: the 3D-organoid Fsk-induced swelling (FIS) assay (Dekkers *et al*, 2013) and Ussing chamber measurements of organoid-derived 2D-monolayers (Li *et al*, 2004; Molinski *et al*, 2017; Cao *et al*, 2018) (Fig 6A).

Intestinal epithelial organoids were embedded into an extracellular matrix (e.g. Matrigel) that reconstitutes a basement membrane and is indispensable for cell survival, proliferation and maintenance of the 3D structure (Dekkers *et al*, 2013). However, this extracellular matrix prevents lentiviruses from binding to intestinal cells. We therefore applied an optimized version of an efficient gene transduction method reported previously (Van Lidth de Jeude *et al*, 2015) to infect 3D-organoid cultures with lentivirus containing shRNAs

targeting CFTR or FGL2. Using this approach, we were able to successfully knockdown each target gene in intestinal organoids derived from either healthy subjects or F508del homozygous CF patients (Fig 6B).

We next performed a Fsk-induced swelling (FIS) assay on these 3D organoids to measure CFTR function (Dekkers *et al*, 2013). The FIS assay is well established as a relatively simple yet robust approach for the quantification of CFTR function using primary intestinal culture models that recapitulate essential features of the *in vivo* tissue architecture. Exposure of organoids to Fsk leads to a rapid increase in the levels of intracellular cAMP, resulting in opening of the CFTR channel, movement of chloride and fluid into the lumen and a consequent increase in organoid size (due to flow of water following the osmotic gradient) which is tracked in time-course, live-cell microscopy (Dekkers *et al*, 2013).

Using the FIS assay, we noticed a significant drop in the ability of organoids expressing wt-CFTR to swell upon FGL2 knockdown, relative to empty vector control, comparable to the effects observed upon knockdown of CFTR (Fig 6C and D). Interestingly, this effect was not as strong in CF mutant organoids, where FGL2 knockdown only mildly reduced swelling (relative to empty vector control) upon Fsk stimulation in the cells treated with a combination of modulators VX-661+VX-445+VX-770 (Trikafta™) to rescue mutant CFTR activity (Fig 6C and E). This was in strong contrast to CFTR knockdown, which resulted in minimal swelling in CF mutant organoids in response to Fsk in the presence of Trikafta (Fig 6C and E). Additionally, FGL2 knockdown in CF mutant organoids led to increased swelling in response to Fsk in the absence of Trikafta rescue, an effect not observed in either empty vector or CFTR knockdown organoids (Fig 6C and E).

Finally, we examined the transepithelial currents in organoids grown in 2D-monolayer using Ussing chambers, a well-established method for studying epithelial ion transport (Li *et al*, 2004; Molinski *et al*, 2017; Cao *et al*, 2018) (Fig 6F). Consistent with the FIS data, knockdown of both CFTR and FGL2 in wt organoids led to a substantial decrease in the amount of Fsk-stimulated CFTR channel activity (measured as a change in transepithelial current), relative to empty vector control (Fig 6F, left panel). In CF organoids, the effect of FGL2 knockdown was reversed, leading to greater CFTR channel activity in the presence of Trikafta than observed with Trikafta rescue alone (Fig 6F, right panel).

Overall, the results of both of these assays are intriguing, providing further evidence of a role for FGL2 in CFTR function and expression.

### Figure 6. Effects of lentivirus-mediated shRNA knockdown of the FGL2 and CFTR genes in intestinal organoids.

- Schematic of patient-derived intestinal organoid acquisition and expansion followed by two functional assays. Created with BioRender.com.
- qPCR Validation of lentivirus-mediated shRNA knockdown of FGL2 and CFTR in intestinal organoids. Significance of sample differences was calculated using the one-tailed student *t*-test ( $n = 2$  biological replicates). All values are mean  $\pm$  SD. EV = empty vector.
- Representative confocal microscopy images of calcein green labelled FIS organoids of a healthy individual and a CF patient, before and 60 min after stimulation with Fsk (5  $\mu$ M). Scale bar, 200  $\mu$ m.
- Quantification of FIS response in wt-CFTR organoids (ORG-01(HC)) shown in (C) as area under the curve at 60 min. Significance of sample differences was calculated using two-way ANOVA ( $n = 2$  biological replicates). All values are mean  $\pm$  SD.
- Quantification of FIS response in F508del-CFTR organoids (ORG-BX-002 (dF/dF)) shown in (C) as area under the curve at 60 min. Significance of sample differences was calculated using two-way ANOVA ( $n = 2$  biological replicates). All values are mean  $\pm$  SD.
- Tracing of Ussing chamber measurements of the transepithelial current produced by 2D monolayer cultures of specified organoids. The F508del-homozygous monolayers with the knockdowns have been rescued with Trikafta.

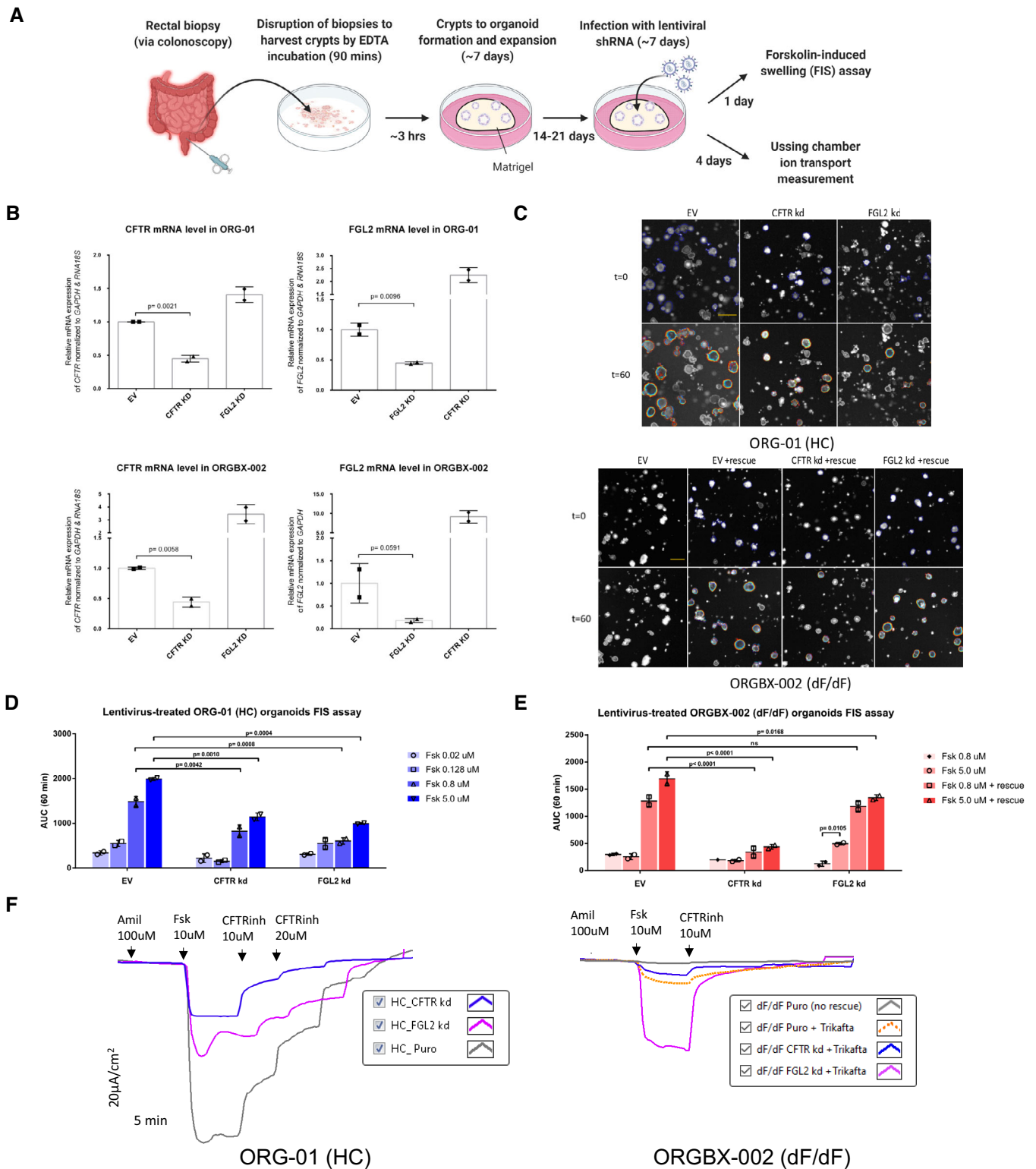


Figure 6.

## Discussion

In this work, we focused on the systematic, high-throughput mapping and characterization of the protein interactomes of wt- and

F508del-CFTR, with the goal of better understanding CFTR biogenesis/function and the cellular changes associated with the CF disease state. To accomplish this, we employed a modified form of our previously reported MaMTH assay (Petschnigg *et al*, 2014, 2017;

Aboulizadeh *et al.*, 2021). While the original MaMTH assay is well suited for mapping the interactions of the full-length forms of complex membrane proteins (such as CFTR) directly in living cells, the arrayed nature of its screening format limits the number of interactions that can be screened in an efficient and cost-effective manner. By introducing a number of changes, including a new fluorophore reporter system, we were able to adapt the system for effective screening of thousands of potential interactors simultaneously, in a pooled manner.

Using this new format, which we term MaMTH-HTS, we were able to screen for PPIs using full-length wt- and F508del-CFTR “baits” against a library of close to 10,000 ORF “preys”, which we constructed from the Human ORFeome V8.1 collection (Yang *et al.*, 2011). Between both baits, we identified a total of 447 candidate interactors, with 224 mapping to wt-CFTR and 269 to F508del-CFTR. Thirty-one of these interactions were previously reported, with the remaining 416 representing new interactions with potential roles in CFTR function. A total of 46 interactors were shared between wt- and F508del-CFTR. This overlap (~10.2%), is lower than that observed in other large-scale CFTR interactome studies, which report a higher percentage of shared proteins (~60% overlap) constituting a “core interactome” containing many ER-, Golgi- and trafficking-associated members involved in CFTR biogenesis (Wang *et al.*, 2006; Pankow *et al.*, 2015). The differences in overlap observed are likely due to the distinct methodologies used, as the CFTR interactome has been mapped heavily by mass spectrometry-based approaches in previous large scale studies (Lim *et al.*, 2018), while our approach looks at interactions of full-length CFTR directly in the membrane of live cells (an environment and context potentially allowing detection of distinct PPIs). Nevertheless, the 31 previously reported interactions confirmed by the MaMTH-HTS help demonstrate the validity of our approach while expanding the scope of the CFTR interactome. Among the detected PPIs, we identified a subgroup of novel interactions for both for wt- and F508del-CFTR (grey circle in Fig 2), which would not likely have been detected by hypothesis-driven approaches. Not surprisingly, these are the proteins with the lowest density of connections, many corresponding to uncharacterized Open Reading Frames (ORFs), that is the so-called dark matter of the genome (Guan & Lazar, 2019). Notwithstanding, these are also conceivably the most interesting PPIs, as they represent still unexplored pathways with the potential to lead to the discovery of novel druggable targets and biomarkers.

Based upon the differences between the traditional MaMTH and our modified MaMTH-HTS format, we suspected that there may be variation in the types of interactions effectively mapped by each method. Comparison of the two approaches using a subset of wt- and F508del-CFTR interactors identified in our MaMTH-HTS screens revealed that this does appear to be the case, with only 20–25% of interactions reproducible in the traditional arrayed format. While some of these differential interactions could be reflective of background inherent in our MaMTH-HTS method, functional testing of several of these revealed that they have a putative role in CFTR trafficking and, in the case of ZNF22, physical interaction was further confirmed by Co-IP, suggesting that these differences are not always due to methodological specificities (summarized in Dataset EV8). Rather, it may indicate that the two methods may better identify different types of interactions, something which could result from steric differences in constructs, the use of stably expressed preys in

MaMTH-HTS versus transiently expressed preys in MaMTH, or even to sensitivity variations resulting from the use of two different reporter systems (i.e. a single *Gaussia princeps* luciferase readout versus the tagBFP/mCherry/eGFP selection), which could lead to detection of stronger versus weaker or stable versus transient interactions. We also observed that in traditional MaMTH testing of a subset of preys identified in MaMTH-HTS as uniquely binding to one CFTR bait (wt- or F508del-CFTR), about 20% also bound to the other CFTR bait. This may provide further support for complementary behaviour between the two assay formats, although technical issues with respect to prey coverage in MaMTH-HTS cannot be ruled out and will be the target of future investigation as we continue to refine and benchmark the format.

To provide preliminary functional characterization of our interactome, we deployed a number of genetic and biochemical assays to address CFTR trafficking and channel activity. These included the FLIPR assay with membrane potential-sensitive dyes (Ahmadi *et al.*, 2017) to measure the effects of interactors on CFTR ion channel activity in cultured cells, targeted siRNA knockdown coupled with high-content microscopy (Botelho *et al.*, 2015) or western blotting to study the role of interactors in CFTR trafficking, co-IP to orthogonally validate selected interactions, and FIS assays (Dekkers *et al.*, 2013) and Ussing chamber (Li *et al.*, 2004) measurements to examine effects of interactor knockdown in patient derived organoids. Although these functional assays only begin to address the potential wealth of information obtainable from our new interactome, we did identify evidence supporting interesting links between some interactors and CFTR function. This dataset also constitutes a valuable resource for multiple future studies to unravel mechanisms of disease in CF and CFTR-related conditions.

The fact that some siRNAs targeting the same gene gave ambiguous results (one with a positive, another with a negative effect) can be due to several reasons. In fact, not all siRNAs target the gene they are aimed at with the same efficiency and some lack specificity. Based on data from multiple high-content screens, it has been estimated that only ~30% of siRNAs effects can be confirmed (Neumann *et al.*, 2010; Simpson *et al.*, 2012; Almaça *et al.*, 2013).

One interesting example of a CFTR PPI is ST6GALNAC1 (ST6 N-Acetylgalactosaminide alpha-2,6-sialyltransferase 1), which was identified as an interactor of both wt- and F508del-CFTR in MaMTH-HTS. Notably, these interactions were also detected in our traditional arrayed MaMTH luciferase assay and the interaction with wt-CFTR was validated by co-IP. Overexpression of ST6GALNAC1 also had a modest effect on improving the functional expression of the F508del-CFTR as seen in the FLIPR assay. ST6GALNAC1 is a sialyltransferase involved in the Golgi transport pathway and plays a role in post-translational modification of lung mucins. A recent study reports an increase in the sialylation of lung mucins in CF pigs, and suggests that this could be linked to an increase in subsequent *Pseudomonas aeruginosa* adhesion and impaired mucociliary transport (Caballero *et al.*, 2021). The authors suggest that lowering the amount of sialylated molecule linked to the lung mucin is worth investigating as a new CF therapeutic strategy for treating bacterial infections. Since we detected a physical interaction between CFTR and ST6GALNAC1, it is interesting to speculate that the altered molecular properties of mucins in the CF pig model could be due to a disruption or modulation of this interaction, something which should be investigated in future studies.

We also identified FKBP6 as a novel interactor of both wt- and F508del-CFTR and found that FKBP6 overexpression led to a modest increase in the function of F508del-CFTR in the FLIPR assay. This finding further supports the growing evidence of a regulatory module formed between ABC transporters and FKBP/HSP90 chaperones responsible for an early step of ER quality control and post-ER trafficking (Wang *et al*, 2006; Geisler & Hegedűs, 2020).

Interestingly, siRNA that targets CAPZB led to a significant decrease in trafficking of both wt- and F508del-CFTR. CAPZB encodes the beta subunit of the barbed-end actin-binding protein, and it functions in regulating actin filament dynamics and stabilization. Although we only found it interacting with mutant CFTR in MaMTH-HTS, it did interact with both wt- and F508del-CFTR in the traditional arrayed MaMTH assay, and overexpression of CAPZB led to an increase in the functional expression of both CFTR variants in FLIPR assays. These results support a role for CAPZB as a key player involved in CFTR and cytoskeletal interaction and acting as a promoter of wt-CFTR trafficking to the PM under EPAC1 activation (Santos *et al*, 2020). Maintaining the CFTR interaction with CAPZB protein thus appears to be critical in promoting CFTR stability at the PM.

Another interesting interactor is VAPA, a type IV integral ER membrane protein in the vesicle-associated membrane protein (VAP) protein family. VAPA is involved in membrane vesicle trafficking by interacting with the SNARE complex. It is also known to be involved in the unfolded protein response (UPR) and to also associate with the cytoskeleton. VAPA interacted only with wt-CFTR in our MaMTH-HTS screen, though it did show an interaction with the mutant in arrayed MaMTH. Interestingly, its siRNA led to a significant decrease in wt-CFTR trafficking, while its overexpression led to increased function of F508del-CFTR in the FLIPR assay, indicating a potentially important role in CFTR trafficking and suggesting that enhancement of VAPA interaction with mutant CFTR could serve as an additional therapeutic avenue to be further explored. It is worth mentioning that one of VAPA's protein family members, VAP, has been reported to play a role in CFTR biogenesis serving as intracellular receptors that couple lipid homeostasis to proteostasis regulation (Ernst *et al*, 2016; Häslér *et al*, 2020). XAGE3 is another wt-CFTR interactor detected in MaMTH-HTS that displays a functional profile in our assays similar to VAPA. Although not much is known about its function aside from its involvement in lung cancer (Zendman *et al*, 2002; Nakagawa *et al*, 2005), it will be interesting to further explore this interaction given the putative role that wt-CFTR plays in protection from epithelial-mesenchymal transition (EMT) and cancer (Amaral *et al*, 2020).

Perhaps the most important functional interactor uncovered from this study is FGL2 (Fibrinogen Like 2, also known as Fibroleukin), whose interaction with both wt- and F508del-CFTR we orthogonally validated using co-IP. This is a 439 amino acid protein belonging to the fibrinogen-related superfamily. It exists as both a type II transmembrane protein found on the surface of macrophages and endothelial cells, and a secreted protein by T cells (Marazzi *et al*, 1998) and it has been associated with liver, interstitium and renal fibrosis by facilitating macrophage polarization (Foerster *et al*, 2010; de Ridder *et al*, 2016; Wu *et al*, 2020). The membrane bound FGL2 has been shown to possess a serine protease activity, cleaving prothrombin to thrombin leading to fibrin deposition and thrombosis and was originally found to be implicated in the pathogenesis of

fulminant hepatitis in humans (Levy *et al*, 2000) and has been used as a biomarker of severe liver diseases, including cancer (Foerster *et al*, 2010; Manns *et al*, 2017; Liu *et al*, 2021). Though little is known about the immunoregulatory properties of FGL2, its expression also seems to play a critical role in the pathogenesis of allograft rejection (Ning *et al*, 2005; Bézie *et al*, 2015) and thus in innate immunity. The soluble FGL2, on the other hand, has been shown to inhibit both T cell proliferation and maturation of bone marrow-derived dendritic cells (Chan *et al*, 2003). Most interestingly, FGL2 transcripts showed a 1.6-fold increase in CFTR-KO mice versus their littermate controls (Guilbault *et al*, 2006).

In our current study, the PPI we observe is most likely between CFTR and the membrane-bound form of FGL2, as the MaMTH system is not suited for detection of secreted proteins (although association with the secreted protein form during trafficking cannot be ruled out). Overexpression of FGL2 in our validation assays led to an enhancement of both wt- and F508del-CFTR activity as measured in the FLIPR membrane potential assays. Additionally, knockdown of the gene led to a decrease in trafficking of wt-CFTR to the PM, though an effect on the mutant was not observed, given it is already absent from the PM. In our intestinal organoid model, knockdown of FGL2 also resulted in a decrease in wt-CFTR channel activity (as measured in both FIS and Ussing chamber assays), consistent with the observed reduction in trafficking observed in our siRNA analysis. However, our results in CF patient organoids were more complex, since FGL2 knockdown apparently did not significantly affect the activity of Trikafta-rescued F508del-CFTR in 3D organoids (as measured in the FIS assay) but led to a substantial increase in activity in 2D monolayers (as measured in using Ussing chamber). Though we cannot rule out the possibility of the difference between 2D monolayer and 3D organoid models, this observed differential effect could be due to the occurrence of FGL2 transcripts in CF versus non-CF tissues (Guilbault *et al*, 2006). It is worth noting that the effects on CFTR transcript levels upon knockdown of FGL2 and vice versa (Fig 6B) suggests some interplay at the transcriptional level is also present.

Altogether, these results suggest a role of the physical interaction of FGL2 with CFTR in its biogenesis and function, and it appears to be clearly important for proper expression of wt-CFTR at the PM. The biology underlying its pronounced effects on mutant CFTR is less clear; however, and it will need to be further investigated. Overall, however, FGL2's reported role in fibrosis and its growing interest as a therapeutic target in treating autoimmune disorders and transplant rejection make it one of the more interesting novel interactors of CFTR uncovered in this study and additional research to determine its potential for use as a therapeutic target in CF seems warranted.

In summary, through the use of MaMTH-HTS, a newly developed high-throughput variant of our MaMTH protein interaction screen designed for integral- and membrane-associated proteins, we were able to successfully map interactomes of wt-CFTR and its primary mutant F508del-CFTR using a library consisting of > 10,000 human ORFs derived from the Human ORFeome V8.1 collection. Alongside the previous study on mapping of the F508del-CFTR interactome using a deep proteomic analysis method (CoPIT) (Pankow *et al*, 2015), the results shown in this work represent one of the largest interactome studies of the wt-CFTR and F508del-CFTR proteins and should be a valuable resource for CF research. The MaMTH-HTS

approach revealed a wide range of new interactions not previously detected by other methods, some of which we demonstrate to have potentially important functional roles in CFTR regulation and biogenesis. Hopefully, the data provided here will prove to be useful for further studies into CF pathophysiology and for the development of innovative biomarkers and better treatment options. It is important to stress, however, that MaMTH-HTS is still new, and that possible issues with background and sensitivity must be acknowledged when evaluating screening results, though such issues will

continue to be explored and addressed as the method undergoes further benchmarking, refinement and application to a greater range of targets. Nevertheless, our results with CFTR clearly demonstrate the ability of the assay to identify novel interactions of biological importance, and we strongly believe that network biology approaches, such as MaMTH-HTS, are increasingly important tools for obtaining unbiased insight into the aetiology of disease. We look forward to the future application of these technologies to further research and drive advancements in human health and disease.

## Materials and Methods

### Reagents and Tools table

Reagent/Resource	Reference or Source	Identifier or catalogue number
<b>Experimental models</b>		
HEK-293 Flp-In cells ( <i>H. sapiens</i> )	ThermoFisher	R75007
HEK-293T cells ( <i>H. sapiens</i> )	ATCC	CRL-3216
Electrocompetent E. coli cells (10G ELITE)	Lucigen	60052-4
CFBE cells ( <i>H. sapiens</i> )	Molinski <i>et al</i> (2015) Botelho <i>et al</i> (2015)	N/A
<b>Recombinant DNA</b>		
MaMTH reporter vector using the pcDNA3.1(–) backbone	Gibson <i>et al</i> (2009) and Saraon <i>et al</i> (2020)	N/A
MaMTH bait destination vector	Gateway cloning technology (Thermo Fisher) and Saraon <i>et al</i> (2020)	N/A
wt- and F508del-CFTR cDNA entry constructs	Dr. Johanna Rommens lab	Ref Seq: NM_000492.3
MaMTH prey destination vector	Saraon <i>et al</i> (2020)	N/A
Plasmid library generation	Human ORFeome V8.1 collection (Yang <i>et al</i> , 2011)	N/A
<b>Antibodies</b>		
Anti-Flag antibody	Sigma	F-1804
Anti-mouse IgG antibody conjugated with Alexa 647	Invitrogen	A-31571
Anti-CFTR 596	Cystic Fibrosis Foundation	N/A
Anti-calnexin	BD Transduction Laboratories	610523
HRP-conjugated goat anti-mouse IgG	Bio-Rad	170-6516
Anti-FLAG antibody	Millipore Sigma	F3165
Anti-V5 antibody	Cell Signaling Technology	13202S
<b>Oligonucleotides and other sequence-based reagents</b>		
PCR primers	This study	Table EV1
Custom library of 421 Ambion Silencer Select siRNAs	This study	Table EV3
Non-targeting Neg1 negative control siRNA	Ambion	4390844
<b>Chemicals, enzymes and other reagents</b>		
DMEM	Wisent	319-005-CL
DMEM high glucose	Corning	10-013-CV
Opti-MEM	Gibco	31985070
IntestiCult™ Human OGM	STEMCELL Technologies	06010
FBS	Gibco	10270
Hygromycin	Bioshop	HYG003.10
Blasticidin	Invivogen	ant-bl

Reagents and Tools table (continued)

Reagent/Resource	Reference or Source	Identifier or catalogue number
Puromycin	Invivogen	ant-pr
Doxycycline	Sigma	9891
Primocin	Invivogen	ant-pm
Vancomycin	Sigma	SBR0001
Gentamycin	Sigma	G1272
Y-2632 (ROCK pathway inhibitor)	Sigma	SCM075
Forskolin	Sigma Aldrich	F3917
VX-661	Selleckchem	S7059
VX-809	Selleckchem	S1565
VX-445	Selleckchem	S8851
CFTRinh-172	Cystic Fibrosis Foundation EMD Millipore Corp	N/A
Amiloride	Spectrum Chemical	TCI-A2599
Phire Tissue Direct Dilution Buffer	Thermo Scientific	N/A
Phire Tissue Direct PCR Master Mix	Thermo Scientific	N/A
QIAquick PCR Purification Kit	Qiagen	N/A
Qiagen Mini Kit	Qiagen	27106
Qiagen Micro Kit	Qiagen	74004
iSCRIPT cDNA synthesis kit	Bio-Rad	1708890
Ssofast™ EvaGreen fluorophore	Bio-Rad	172-5200
CellMask	Invitrogen	H32712
ER-Tracker	Invitrogen	E34250
Hoechst 33342	Sigma	B2261
Calcein green	Invitrogen	C34852
X-tremeGENE 9 DNA Transfection Reagent	Roche	6365787001
Lipofectamine 2000	Invitrogen	11668019
Lipofectamine LTX	Thermo Fisher Scientific	15338100
Laemmli sample buffer	Bio-Rad	161-0747
Protease inhibitor cocktail	Roche	11873580001
Clarity Western ECL Substrate	Bio-Rad	1705061
Blue membrane potential dye	Molecular devices	R8034
384 well plates	Falcon	736-2044
96 well plates	Sarstedt	83.3924
12 well plates	SPL Life Sciences	30012
PureCol®-coated Transwell	Corning	Costar™ 3470
Non-perfused Ussing chamber	Physiologic Instruments	N/A
Drying pearls	Sigma-Aldrich	94098
μMACS™ FLAG magnetic microbeads and columns	Miltenyi Biotec	N/A
Matrigel	BD Biosciences	N/A
<b>Software</b>		
GraphPad Prism	<a href="https://www.graphpad.com">https://www.graphpad.com</a>	N/A
Integrated Interactions Database (IID)	PMID: 34755877 <a href="http://ophid.utoronto.ca/iid">http://ophid.utoronto.ca/iid</a>	Version 2021-05
NaViGaTOR	<a href="http://navigator/ophid.utoronto.ca/navigatorwp">http://navigator/ophid.utoronto.ca/navigatorwp</a>	Version 3.0.15
R programming language/Bowtie 2 Alignment Tool	R Core Team, Bowtie2 alignment tool (Langmead & Salzberg, 2012)	N/A

Reagents and Tools table (continued)

Reagent/Resource	Reference or Source	Identifier or catalogue number
CellProfiler	McQuin <i>et al</i> (2018)	N/A
Image Lab	Bio-Rad	N/A
MATLAB-based organoid swelling capturing program	MATLAB	N/A
<b>Other</b>		
NexteraXT library preparation kit	Illumina	N/A
NextSeq500 System	Illumina	N/A
HiSeq2500 System	Illumina	N/A
BD FACS Melody	BD Biosciences	N/A
Rank product method	Breitling <i>et al</i> (2004)	N/A
Genevac miVac Duo	Thermo Fisher Scientific	N/A
Multidrop™ Combi peristaltic dispenser	Thermo Scientific	5840300
Leica DMI 6000B	Leica	N/A
ChemiDoc XRS	Bio-Rad	N/A
Cellomics ArrayScan, VTI HCS Reader	Thermo Fisher Scientific	N/A
NanoDrop 2000	Thermo Fisher Scientific	N/A

## Methods and Protocols

### Generation of MaMTH-HTS reporter cell line

Reporter vector was generated in a pcDNA3.1(−) backbone using ORFs expressing eGFP under the control of a 5xGAL4 UAS and puromycin resistance marker under the control of a constitutive PGK promoter, via Gibson assembly (Gibson *et al*, 2009). This vector was then stably integrated into FLP-compatible HEK293 cells and the candidate line displaying the best MaMTH-responsive reporter activity (with minimal background) was selected, following our previously described protocol used for generation of *Gaussia princeps* luciferase reporter lines (Saraon *et al*, 2020).

### Generation of MaMTH CFTR bait constructs

CFTR bait constructs were generated using Gateway cloning technology (Thermo Fisher) and destination vector designed in our lab, modified from our previously reported A1160 vector (Saraon *et al*, 2020). These new constructs retained the general structural features of the original vector (including tetracycline induction and Flp-In T-REx compatibility) but also expressed P2A “linked” tagBFP to allow for fluorescence-based detection of bait expression (Appendix Fig S2A). wt- and F508del-CFTR cDNA entry constructs (Ref Seq: NM\_000492.3) used in cloning were generously gifted from Dr. Johanna Rommens.

### Generation of MaMTH Prey construct and library

Gateway- and CRISPR-compatible MaMTH prey destination vector was designed, modified from our previously reported A1245 vector (Saraon *et al*, 2020). This new construct contained the primary features of the earlier plasmid (including P2A “linked” mCherry to allow for fluorescence-based detection of prey expression), alongside a tetracycline-inducible promoter, hygromycin resistance for selection upon stable integration and CRISPR-targeting sites (Appendix Fig S2B).

For plasmid library generation, 33 ORF entry clone pools [generated from the Human ORFeome V8.1 collection (Yang *et al*, 2011), were transferred into this prey vector construct using Gateway cloning (Thermo Fisher). LR reactions were carried out on a per pool basis and transformed into commercial electrocompetent E. coli cells (Lucigen) to maximize efficiency. Colony counts from transformations were used to estimate coverage of individual pools, which were determined to be a minimum of 1,000× in all cases. ORF representation in pools was verified by deep sequencing using NexteraXT library preparation and sequencing on a NextSeq500 system (150 bp single read).

### Generation of MaMTH prey cell line library pools

For generation of MaMTH-HTS cell line library stably expressing preys, MaMTH-ORF-expressing prey plasmids were transfected, on a per pool basis, into HEK293 MaMTH-HTS eGFP reporter cells using X-tremeGene 9 (Roche) alongside plasmids expressing Cas9, AAVS targeting gRNA and prey-plasmid targeting gRNA. The cells were grown for 3 days at 37°C /5% CO<sub>2</sub> in DMEM/10% FBS/1% PenStrep prior to expansion into DMEM/10% FBS/1% PenStrep media containing 100 µg/ml of hygromycin. Plates were grown further until foci appeared (approximately 7 days), at which point all foci were harvested collectively per pool, expanded further and frozen/stored in N<sub>2</sub>(l). Foci counts were used to ensure that pool coverage was at least 30× or greater. To establish ORF representation in cell line pools, genomic DNA was isolated on a per pool basis, amplified by PCR using cassette-specific primers (Table EV1, Primer Set 1) and deep sequenced using NexteraXT library prep and an Illumina HiSeq2500 system (150 bp, single read, Dataset EV1).

### MaMTH-HTS screens

- 1 Plasmid expressing wt-CFTR or F508del-CFTR “Bait” protein fused at its C-terminus to MaMTH-HTS Bait tag (Cub-GAL4TF-P2A-tagBFP) was transfected into a pooled “Prey” library of



HEK293 MaMTH reporter cell lines (containing chromosomally integrated GFP reporter under the control of a GAL4TF promoter) stably expressing members of the Human ORFeome V8.1 collection (~10,000 ORFs) fused at their N-terminus to MaMTH-HTS Prey tag (Nub) and at their C-terminus to P2A-mCherry. Transfections were performed using Xtreme gene 9 transfection reagent (Roche), following manufacturer's protocols, in 6 cm dishes containing 1,000,000 seeded cells per plate (grown overnight to ~80% confluency) with 750 ng of bait DNA.

- 2 Five hours post transfection 0.5 µg/ml Tetracycline was added (to induce Bait and Prey expression) and the cells were grown for 2 days at 37°C/5% CO<sub>2</sub> in DMEM/10% FBS/1% PenStrep media.
- 3 Following growth, the cells were harvested by trypsinization and resuspended in Basic Sorting Buffer (1 × PBS, 5 mM EDTA, 25 mM HEPES pH 7.0, 1% BSA) at a concentration of 1–2 × 10<sup>6</sup> cells/ml.
- 4 The cells were then sorted by Flow Cytometry using a BD FACS Melody (BD Biosciences). Sorts were performed such that the cells were sequentially selected based on the presence of mCherry fluorescence (indicating Prey expression), tagBFP fluorescence (indicating Bait expression) and finally GFP fluorescence (indicating Bait-Prey interaction). The cells were collected in DMEM/25% FBS.
- 5 Collected cells were centrifuged at 200 g for 5 min and pellets resuspended/processed in Phire Tissue Direct Dilution Buffer (Thermo Scientific).
- 6 ORFs were then amplified using Phire Tissue Direct PCR Master Mix (Thermo Scientific) following manufacturer's instructions with the primer sets shown in Table EV1, generating two technical replicates (to use for determining possible primer-specific PCR biases or inconsistencies across sequencing samples).
- 7 PCR products were then purified using a QIAquick PCR Purification Kit (Qiagen).
- 8 Purified PCR products were subjected to Nextera XT library preparation (following manufacturer's instructions) and deep sequenced on an Illumina HiSeq2500 system (150 bp single read).
- 9 Sequencing data was then processed with in-house software scripts developed using the R programming (R Core Team) language and integrating the Bowtie2 alignment tool (Langmead & Salzberg, 2012), removing any ORFs with total counts less than 3 in both replicates, to generate lists of candidate interactors.

### Identifying frequent fliers

Frequent fliers were identified by using the rank product method (Breitling *et al*, 2004) to analyse results of 9 MaMTH-HTS screens of control baits from distinct classes of membrane proteins. The main idea was that frequent fliers would have strong interaction signals from multiple screens; the rank product method (Breitling *et al*, 2004) would calculate the probability of such signals occurring by chance—thus providing a *P*-value for the hypothesis that a prey is a frequent flier. Interaction signal was considered proportional to read counts per million (CPM) from deep sequencing.

The rank product method (Breitling *et al*, 2004) comprised calculation of rank products, *P*-values, and adjusted *P*-values for all preys. Rank products were calculated by ranking CPMs in

each screen (the highest CPM receiving a rank of 1) and then for each prey, calculating the geometric mean of its 9 CPM ranks.

*P*-values were calculated by determining how often rank products were better than from randomized data. This was done through 100,000 iterations of the following steps: CPM rank data were randomized (i.e. CPM ranks in each screen were randomly permuted), rank products were calculated from the randomized data and compared against observed rank products. *P*-values were calculated as the number of times rank products from randomized data were smaller than from observed data, divided by 100,000.

*P*-values were adjusted for multiple testing by the Benjamini-Hochberg method (Benjamini & Hochberg, 1995).

### Annotation of CFTR interactions with known and predicted interactions

Previously detected and predicted CFTR interactions were retrieved from the Integrated Interactions Database (IID) (Kotlyar *et al*, 2021) version 2021-05 (<http://ophid.utoronto.ca/iid>). CFTR interaction partners were annotated with GO Slim cellular components and molecular functions from the GOA database (Huntley *et al*, 2015) (downloaded 2021-06-16), diseases from DisGeNET (Piñero *et al*, 2020) version 7.0, tissues from IID (Kotlyar *et al*, 2021) version 2021-05, and pathways from pathDIP (Rahmati *et al*, 2019) version 4.0 (<http://ophid.utoronto.ca/pathDIP>). CFTR interactions were annotated with mutations from IntAct (Del-Toro *et al*, 2019; Porras *et al*, 2020) version 4.2.17. Annotated interaction data are presented in Dataset EV4.

Network figures were created using the integrated interaction data in NAViGaTOR ver. 3.0.15 (Brown *et al*, 2009); <http://navigator.ophid.utoronto.ca/navigatorwp>). Final networks were exported in SVG file format, finalized in Adobe Illustrator ver. 26.0.1 to include legends, and exported in 300 DPI PNG format for publication.

### Fluorescence -microscopy to monitor CFTR localization

HEK293 cells were transiently transfected with plasmid expressing wt-CFTR or F508-del CFTR fused to MaMTH tag (C-terminus of ubiquitin and LexA artificial transcription factor) and GFP and grown at 37°C/5% CO<sub>2</sub> in DMEM/10% FBS/1% PenStrep media. Following growth, the cells were visualized by fluorescence microscopy using CellMask (Invitrogen) to stain the plasma membrane and ER-Tracker (Invitrogen) to stain the endoplasmic reticulum, following standard protocols.

### Traditional MaMTH assay

Protocols for preparing and performing the MaMTH assay have been previously published (Saraon *et al*, 2017). All the MaMTH data presented are from three independent trials each with at least three technical replicates in a 96-well plate format, unless otherwise stated.

### Membrane potential assay

HEK293 cells were grown on 96 well tissue culture plates until confluent. The blue membrane potential dye (Molecular devices) was dissolved at a concentration of 0.5 mg/ml as reported previously (Molinski *et al*, 2015; Ahmadi *et al*, 2017). After 15–20 min of loading the dye at 37°C, 5% CO<sub>2</sub> and humidified air, the plate was transferred to the microplate reader (Molecular devices). Briefly, the reader was heated to 37°C and the settings for fluorescence whole well scan were turned on, with multiple points being read each well. Upon start of the experiment, baseline

reads of at least 4–5 scans were made, followed by addition of drugs (2.5  $\mu\text{l}$ /well). After addition of each drug again 4–5 scans were done. The experiment ended by addition of inhibitor (CFTRinh-172 10  $\mu\text{M}$ ). Upon completion of experiment, the data were exported and analysed.

#### Cell lines for trafficking assays and western blots

CFTR traffic measurements were performed on Cystic Fibrosis Bronchial Epithelial (CFBE) cells expressing inducible (TET-ON) double-tagged CFTR (Molinski *et al*, 2015) traffic reporters (mCherry-Flag-CFTR) previously described (Botelho *et al*, 2015) and representing either the WT- or F508del-CFTR variants. Western blot validation experiments were performed on CFBE cell lines constitutively expressing wt- or F508del-CFTR.

#### Cell culture for trafficking assays and western blots

CFBE mCherry-Flag-CFTR cells (WT or F508del variants) were cultured in DMEM high glucose (Corning 10-013-CV) supplemented with 10% fetal bovine serum (Gibco 10270), 10  $\mu\text{g}/\text{ml}$  of blasticidin (Invivogen #ant-bl) and 2  $\mu\text{g}/\text{ml}$  of puromycin (Invivogen #ant-pr-1) at 37°C and 5%  $\text{CO}_2$ . CFBE cells constitutively expressing WT-/F508del-CFTR were cultured in equivalent conditions but excluding blasticidin. When seeded to siRNA coated multi-well plates culture media was DMEM supplemented with FBS (0.1% when using VX-661, 10% elsewhere).

#### Preparation of siRNA-coated multi-well plates

Multi-well plates were coated with a custom library of 421 Ambion Silencer Select siRNAs (Table EV3) The library targeted 210 MaMTH-HTS hits (2 siRNAs for most genes) and included the non-targeting Neg1 negative control (Ambion 4390844) and CFTR (s2947) and INCENP (s7424) transfection controls. The library was used to coat multi-well plates for reverse transfection, according to a previously reported protocol (Botelho *et al*, 2015). For the CFTR traffic screen 384 well plates (Falcon 736–2,044) and for western blot assays 12 well plates (SPL Life Sciences 30012) were used. Briefly, siRNAs were diluted to 3  $\mu\text{M}$  with RNase free water. In parallel, a transfection mix comprised of 0.4 M of sucrose in Opti-MEM (Gibco 31985070), RNase-free water and Lipofectamine 2000 (Invitrogen 11668019) in the proportions 1.715:1:1 and a 2 g/l of gelatin solution were prepared. Then, each siRNA solution was mixed with the transfection mix, incubated 20 min at room temperature and mixed with the gelatin solution (proportions 1:1.4:1.4). The resulting solution was then diluted 50-fold in RNase-free water. Finally, the solution was added to each well of the multi-well plates (384 well: 15  $\mu\text{l}$ /well, 0.24 pmol siRNA/well; 12 well: 720  $\mu\text{l}$ /well, 11.4 pmol siRNA/well). Plates were immediately lyophilized on a speedvac (Genevac miVac Duo) at 37°C for about 11 h and then stored in air-tight boxes containing drying pearls (Sigma-Aldrich 94098) at room temperature until needed for bioassays. Traffic assay plates contained 7–18 negative control wells (Neg1). Western blot plates contained 1 negative control sample.

#### CFTR traffic assay

The assay was performed as previously described (Botelho *et al*, 2015) CFBE cells expressing the inducible mCherry-Flag-CFTR reporter (WT- or F508del-CFTR variants) were grown to confluence and split to 50% confluency. Twenty-four hours later, the cells were

trypsinized to antibiotic-free medium and seeded in siRNA coated 384-well plates (1,000 cells/well) using a Multidrop™ Combi peristaltic dispenser (Thermo Scientific 5840300). CFTR expression was induced for 48 h (24 h after seeding) by supplementing the medium with 1  $\mu\text{g}/\text{ml}$  of doxycycline (Sigma 9891). At this point, VX-809 (3  $\mu\text{M}$ ) or VX-661 (5  $\mu\text{M}$ ) (Selleckchem (S7059)) were added to selected wells containing Neg1 siRNA as a positive control for F508del-CFTR traffic rescue. Extracellular Flag tags were immunostained in non-permeabilized cells 72 h after seeding. The primary anti-Flag antibody (Sigma F-1804, 1:500) was incubated 1 h at 4°C, the cells were fixed with PFA 3% for 20 min at 4°C, an anti-mouse IgG antibody conjugated with Alexa 647 (Invitrogen A-31571, 1:500) was incubated 1 h at room temperature and Hoechst 33342 (200 ng/ml, Sigma B2261) was incubated 1 h at room temperature. Four independent biological replicates were performed.

#### High-content image acquisition and analysis

Automated imaging was performed on a Leica DMI 6000B widefield fluorescence microscope using a 10 $\times$ /0.40 objective, EL6000 light source, Hamamatsu Orca-Flash4.0 camera and suitable filter cubes for Hoechst, mCherry and Alexa 647. Each well was imaged at 4 sub-positions. Images were annotated with experimental metadata using the htmrenamer R package (Hagemeyer *et al*, 2020). Cell-based fluorescence quantification was performed with CellProfiler (McQuin *et al*, 2018) in order to quantify CFTR expression (mCherry) or PM localization (Alexa 647), excluding the cells presenting apoptotic or otherwise aberrant nuclei. Numerical data was processed and normalized with a custom R script, which excluded the cells with negligible CFTR expression, out-of-focus images or images with less than 100 cells. For each image, the median Alexa 647 fluorescence intensity across all the cells was computed and converted into a Z-score using the median 5  $\times$  5 well neighbourhood fluorescence as reference. The reported Z-score is the median value across replicate wells and experimental replicates. Screen hits were the siRNAs, which yielded an increase in PM CFTR of Z-score > +1 or decrease in PM CFTR of Z-score < –1.

#### Western blotting (WB)

CFBE cells constitutively expressing WT- or F508del-CFTR were grown to confluence and split to 50% confluency. Twenty-four hours later, the cells were trypsinized to antibiotic-free medium and seeded in siRNA coated 12-well plates (87,500 cells/well). Twenty-four hours later the media was changed to DMEM supplemented with 0.1% FBS containing wither 0.05% DMSO or 5  $\mu\text{M}$  of VX-661. After an additional twenty-four hours, the cells were lysed and subjected to SDS-PAGE. A sample buffer supplemented with reducing Laemmli sample buffer (Bio-Rad 161-0747), protease inhibitor cocktail (Roche 11873580001), benzonase and  $\text{MgCl}_2$  was used. Samples were resolved in a 10% SDS-PAGE gel and transferred into a PVDF membrane. Membrane was blocked with 3% (w/v) non-fat milk (NFM) in tris buffered saline with 0.1% Tween 20 (TBS-T) for 1 h at room temperature and incubated overnight at 4°C with anti-CFTR 596 (Cystic Fibrosis Foundation) or anti-calnexin (BD Transduction Laboratories 610523) diluted 1:3,000 in 3% (w/v) NFM/TBS-T. Membranes were then incubated with HRP-conjugated goat anti-mouse IgG (Bio-Rad 170-6516) diluted 1:3,000 in 3% NFM/TBS-T for 2 h at room temperature. Subsequently, the immunoreactive signals were detected using a Clarity Western ECL Substrate

(Bio-Rad 1705061) and recorded with a ChemiDoc XRS (Bio-Rad). Densitometric analysis was performed on Image Lab (Bio-Rad). The intensity of CFTR bands was always expressed as a fold change versus the calnexin internal control. Statistical analysis consisted of a one-way ANOVA versus the siNeg1 negative control treated with DMSO followed by Dunnett's post hoc test. An equivalent analysis was performed for F508del-CFTR cells treated with siNeg1 and VX-661. The significance level was 0.05.

#### Co-immunoprecipitation (Co-IP)

Co-IP validation was performed on a subset of top CFTR interactors uncovered via MaMTH-HTS ( $n \geq 2$  biological replicates). Briefly, 293T cells were grown in standard high-glucose Dulbecco's modified Eagle medium (DMEM) containing 4 mM of L-glutamine, 1 mM of sodium pyruvate, 10% fetal bovine serum (FBS), penicillin (100 U/ml) and streptomycin (100 µg/ml) under standard conditions (37°C, 5% CO<sub>2</sub>) as previously reported (PMID: 31536960, 29128334). The cells were co-transfected with V5-tagged CFTR and FLAG-tagged preys using Lipofectamine LTX (Thermo Fisher Scientific #15338100), and incubated for 48 h prior to harvesting the cells for Co-IP. Afterwards, the cells were crosslinked for 30 min at room temperature using 0.5 mM of dithiobis succinimidyl propionate (DSP) diluted in PBS. After quenching the crosslinked reaction with the addition of 100 mM of Tris-HCl (pH 7.5) for 10 min at room temperature, the cells were collected by gentle pipetting and centrifuged at 1,300 g for 5 min. The crosslinked pellets were then lysed in 1 ml of RIPA buffer (50 mM of Tris HCl pH7.5, 150 mM of NaCl, 1% Sodium deoxycholate, 0.1% SDS, 1% NP-40, 1 mM of EDTA), containing 1× protease inhibitor cocktail and agitated for 1 h at 4°C. About 100 µl of µMACS DYKDDDDK (also known as FLAG) magnetic microbeads (Miltenyi) were added to the samples and then incubated for additional 4 h at 4°C in agitation. Microbeads suspension was passed through µMACS columns (Miltenyi), washed two times with 1 ml of 0.1% RIPA and 1× protease inhibitor cocktail, followed by another wash with 1 ml detergent free RIPA. Proteins were eluted using 100 µl of 2× Laemmli buffer heated at 95°C. Eluates were then analysed by immunoblotting using anti-FLAG antibody (Millipore Sigma #F3165) directed against overexpressed FLAG-tagged protein corresponding to the interactors, as well as using anti-V5 antibody (Cell Signalling Technology #13202S) directed against the V5-tagged CFTR.

#### Lentivirus-mediated knockdown in the organoids

Lentiviral particles containing shRNA sequences specific for the target proteins were generated in HEK293T cells using the Mission shRNA system with validated shRNA sequences (Sigma-Aldrich) following standard protocols from The RNAi Consortium (TRC) Broad Institute.

The protocol for lentivirus packaging, modified from a previously published method (Hotta *et al*, 2009) was generously provided by Nicolaes Min from Dr. Xi Huang's Lab at The Hospital for Sick Children. Briefly, the virus-containing supernatant media were harvested and filtered through a 0.45-µm syringe filter to remove cell debris and to prevent contamination with virus producer cells. Viruses were then ultracentrifuged for 2 h at 90,000–100,000 g at 4°C. After the centrifugation and discarding of the supernatant, the viral pellets were resuspended in approximately 1 ml of Growth medium (IntestiCult™ Human OGM, STEMCELL Technologies, CAN) and stored in –80°C freezer.

The protocol for lentiviral transduction of the patient-derived intestinal organoids was based on previously published method (Van Lidth de Jeude *et al*, 2015) with minor modifications of scaling down the reagents from the original 48-wells plate to the 24-wells plate format.

Lentivirus production and infection is covered under approval from The Hospital of Sick Children (REB number: 1000058992) and all steps were performed in a biosafety level 2/3-certified laboratory.

#### Quantitative real-time PCR

RNA extraction was performed according to the manufacturer's protocol (Qiagen Micro or Mini Kit) (Hilden, Germany). Briefly, the cells were lysed, RNA extracted and then RNA concentration was measured while using a NanoDrop 2000 instrument (Thermo Fisher Scientific, Waltham, MA, USA). Only samples with a concentration > 100 ng/µl were used, with a 260/280 ratio between 1.8 and 2.1. cDNA synthesis was performed while using reverse transcriptase (iSCRIPT cDNA synthesis kit, Bio-Rad, Hercules, CA, USA) or without reverse transcriptase (negative control). Quantitative real-time PCR was performed while using Eva green (Ssofast Evagreen, Bio-Rad, Hercules, CA, USA) fluorophore in 96-well plates (Bio-Rad, Hercules, CA, USA) and then normalized to GAPDH or RNA18S.

#### Forskolin-induced swelling (FIS) assay

The protocols for FIS assay including the crypt isolation and culturing of human intestinal cells in Matrigel have been described previously (Dekkers *et al*, 2013, 2016; Eckford *et al*, 2014). In short, intestinal crypts were isolated from rectal biopsies and seeded in 50% Matrigel (BD Biosciences) in 24-well plates (~10–30 crypts in three 10-µl Matrigel droplets per well). Growth medium (IntestiCult™ Human OGM, STEMCELL Technologies, CAN) was supplemented with Primocin (1:500; Invivogen). Vancomycin, gentamicin (both from Sigma) and Y-27632 (ROCK pathway inhibitor) were added during the first week of culture. The medium was refreshed every 2–3 days, and resulting organoids were passaged ~1:3–5 every 7–10 days.

Lentivirus-infected rectal organoids were plated in 96-well culture plates in 4 µl of 50% Matrigel containing 20–80 organoids per well and immersed in 100 µl medium. CFTR rescue was achieved by adding modulators (3 µM VX661<sup>+</sup> 3 µM VX445, Selleckchem, Houston, TX, US) in the medium 18–24 h prior to completing the FIS assay. For the FIS assay, organoids were incubated for 30 min with 3 µM of calcein green (Invitrogen) before stimulation with Fsk concentration 0.2–5.0 µM (Sigma Aldrich) in addition to 3 µM VX770 in rescued cultures, and directly analysed by confocal live cell microscopy at 37°C (Cellomics ArrayScan VTI HCS Reader, ThermoFisher Scientific). The increase in swelling over time was captured using a MATLAB-based software program and the area under the curve (AUC,  $t = 60$  min; 2 technical replicates) was calculated using GraphPad Prism.

#### Ussing chamber experiment

The method of measuring the CFTR-dependent intestinal epithelial ion transport properties in rectal organoid-derived monolayers have been described previously (Li *et al*, 2004; Molinski *et al*, 2017; Cao *et al*, 2018; Zomer-van Ommen *et al*, 2018). Briefly, upon dissociation of the 3D intestinal organoid cultures, the cells were grown into

the 2D monolayer on PureCol<sup>®</sup>-coated Transwell (Costar<sup>™</sup> 3470, Corning, Tewksbury, US). They were studied in a non-perfused Ussing chamber (Physiologic Instruments, San Diego, CA). The buffer was maintained at pH 7.4 and 37°C and continuously gassed with 5% CO<sub>2</sub>/95% O<sub>2</sub> mix. The transepithelial potential (V<sub>te</sub>) was recorded and the baseline resistance (R<sub>te</sub>) was measured following repeated, brief short-circuit current pulses (1 μA every 30 s). The results are presented as equivalent transepithelial current (I<sub>eq</sub>), which was calculated using Ohm's law. CFTR function was determined after inhibition of the epithelial sodium channel (ENaC) with amiloride in the apical bath (100 μM, Spectrum Chemical, Gardena, CA) and cAMP activation with forskolin (10 μM, Sigma-Aldrich, US). CFTR activity was quantified as I<sub>eq</sub> difference following forskolin stimulation, CFTRInh-172 (10 μM, EMD Millipore Corp. US) was applied to inhibit CFTR activity.

### Validation strategy

Our validation strategy first involved re-testing a large subset of 240 of the 447 CFTR interactors identified from the MaMTH-HTS screens (Fig 2) using the traditional array-based MaMTH assay. From there, we selected 26 interactors based on their interaction signal strength and consistency in their interaction profile between the two methods. Note that we included both cases where traditional and HTS interactors overlapped and where they did not, focusing largely on interactions confirmed in traditional MaMTH for both WT and mutant (but including some instances where this was not the case for sample diversity). The 26 interactors were then tested in the fluorescence-based FLIPR assay to monitor their effect on the channel activity. Concomitantly, we performed the traffic-based siRNA microscopy screen, targeting 208 genes (based on their availability in the siRNA library) from our original 447 interactors to investigate the trafficking of CFTR to the plasma membrane in the absence of these discovered interactors. This high-content screen was combined with Western blot experiments for validating the post-translational glycosylation pattern of CFTR. From these experiments, we performed final orthogonal validation on some of the most interesting interactors using a co-IP approach and followed up with organoid work. A graphical overview of our validation approach is provided in Appendix Fig S10.

### Statement of informed consent

Informed consent was obtained prior to rectal biopsies using standard endoscopic biopsy forceps.

## Data availability

The data produced in this study are available in the following databases: MaMTH-HTS Interaction Data: IID Integrated Interaction Database (<http://ophid.utoronto.ca/iid>). Part of the IMEx Consortium, (Orchard *et al*, 2012; Porras *et al*, 2020). Deposited as MaMTH-HTS-2021, identifiable with PMID.

**Expanded View** for this article is available online.

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### Author contributions

**Sang Hyun Lim:** Conceptualization; Formal analysis; Validation; Investigation; Writing—original draft; Writing—review & editing. **Jamie Snider:** Conceptualization; Software; Formal analysis; Validation; Investigation; Methodology; Writing—original draft; Writing—review & editing. **Liron Birimberg-Schwartz:** Formal analysis; Investigation. **Wan Ip:** Formal analysis; Investigation. **Joana, C Serralha:** Formal analysis; Investigation. **Hugo, M. Botelho:** Formal analysis; Supervision; Investigation; Writing—original draft; Writing—review & editing. **Miquéias Lopes-Pacheco:** Formal analysis; Investigation. **Madalena, C Pinto:** Investigation. **Mohamed, Taha Moutaoufik:** Formal analysis; Investigation. **Mara Zilocchi:** Investigation. **Onofrio Laselva:** Formal analysis; Investigation. **Mohsen Esmaeili:** Investigation. **Max Kotlyar:** Data curation; Formal analysis; Visualization; Writing—original draft; Writing—review & editing. **Anna Lyakishveva:** Formal analysis; Investigation. **Priscilla Tang:** Formal analysis; Investigation. **Lucía López Vázquez:** Investigation. **Indira Akula:** Investigation. **Farzaneh Aboulzadeh:** Investigation. **Victoria Wong:** Investigation. **Ingrid Grozavu:** Investigation. **Teuta Opacak-Bernardi:** Investigation. **Zhong Yao:** Conceptualization; Methodology. **Meg Mendoza:** Investigation. **Mohan Babu:** Supervision; Writing—original draft. **Igor Jurisica:** Software; Formal analysis; Visualization; Writing—original draft; Writing—review & editing. **Tanja Gonska:** Formal analysis; Supervision; Investigation. **Christine, E Bear:** Conceptualization; Supervision. **Margarida, D Amaral:** Supervision; Writing—original draft; Writing—review & editing. **Igor Stagljär:** Conceptualization; Supervision; Writing—original draft; Project administration.

In addition to the CRediT author contributions listed above, the contributions in detail are:

SHL: performed experiments, analysed data and wrote the manuscript. JS: performed experiments, analysed data and wrote the manuscript. LB-S: performed experiments and analysed data. WI: performed experiments and analysed data. JCS: performed experiments and analysed data. HMB: supervised research, planned, performed and analysed experiments. Contributed to final manuscript. ML-P: performed experiments and analysed. MCP: performed experiments. MTM: performed experiments and analysed data. MZ: performed experiments. OL: performed experiments and analysed data. ME: performed experiments. MK: performed data analysis and interactome assembly. AL: performed and assisted with experiments, assisted with data analysis. PT: performed and assisted with experiments. LLV: performed and assisted with

experiments. IA: performed and assisted with experiments. FA: performed and assisted with experiments. VW: performed and assisted with experiments. IG: performed and assisted with experiments. TO-B: performed and assisted with experiments. ZY: provided assistance with experimental design and implementation. MM: performed experiments. MB: supervised research and contributed to final manuscript. IJ: performed data analysis and interactome assembly. TG: supervised and assisted with experimental design of organoid research. CEB: supervised research and assisted with experimental design. MDA: supervised research and contributed to final manuscript. IS: supervised research and contributed to final manuscript.

### Disclosure and competing interests statement

IS is named as a co-inventor on a patent for the MaMTH technology.

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