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Connectivity Mapping (ssCMap) To Predict A20 Inducing Drugs Anti-inflammatory Action In Cystic Fibrosis

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Cystic Fibrosis (CF) lung disease is characterised by a chronic and exaggerated inflammation in the airways. Despite recent developments in therapeutically overcome the underlying functional defect in CFTR (cystic fibrosis transmembrane conductance regulator), there is still an unmet need to also normalise the inflammatory response. The prolonged and heightened inflammatory response in CF is in part mediated by a lack of intrinsic downregulation of the pro-inflammatory NF-κB pathway. We have previously identified reduced expression of the NF-κB down-regulator A20 in CF as a key target to normalise the inflammatory response.

Here we have used publicly available gene array expression data together with ssCMap (statistically significant connections' map) to successfully predict drugs already licensed for the use in humans to induce A20 mRNA and protein expression and thereby reduce inflammation. The effect of the predicted drugs on A20 and NF-κB(p65) expression (mRNA) as well as pro-inflammatory cytokine release (IL-8) in the presence and absence of bacterial LPS was shown in bronchial epithelial cells lines (16HBE14o-, CBFE410-) and in primary nasal epithelial cells (PNECs) from patients with CF (Phe508del homozygous) and non-CF controls. Additionally, the specificity of the drug action on A20 was confirmed using cell lines with TNFAIP3 (A20) knockdown (siRNA). We also show that the A20 inducing effect of ikarugamycin and quercetin is lower in CF derived airway epithelial cells than in non-CF cells.

A20 (TNFAIP3) | NF-kappaB | Connectivity Mapping | Drug repositioning | CF airway inflammation

Introduction

The response to pathogens, recognised by pattern recognition receptors including Toll-like receptors (TLRs), triggers an acute innate immune response that is mediated by transcription factors such as nuclear factor-kappa-light-chain enhancer of B cells (NF-κB). NF-κB activation promotes the transcription of inflammatory mediators in a tightly regulated process. However, in individuals with underlying chronic inflammatory diseases, this regulation is compromised, leading to constitutive NF-κB activation and persistent inflammation (1-3).

The development of new first-in-class medicines is costly (approximately $1.2 billion for a single FDA-approved drug) and takes between 10 and 15 years (4, 5). Many newly developed drugs perform well in the preclinical testing, but fail when tested in humans (6). Thus alternative approaches using predictive models to identify new drugs are needed. Gene expression connectivity mapping (www.broadinstitute.org/cmap/) is an advanced bioinformatics technique to establish the connections among biological states via gene expression profiles/signatures. One major application of connectivity mapping is to identify potential small molecules able to inhibit a disease state or regulate the expression of a small number of genes (7-9). We used an advanced version of connectivity mapping, ssCMap (statistically significant connections' map) (10), which has been successfully applied to phenotypic targeting and predicting effective drugs in cancer (10). However, this has not yet been applied to chronic inflammatory diseases.

Cystic Fibrosis (CF) is a chronic multi-organ inflammatory disease, caused by mutations in the CFTR gene (Cystic Fibrosis Transmembrane Conductance Regulator) expressed on apical epithelial surfaces. It is the most common lethal genetic disease in Caucasian populations. Lung disease is the primary cause of morbidity and mortality in CF, resulting from dehydration of epithelial surfaces and reduced mucociliary clearance as a consequence of the ionic imbalance created by CFTR mutation. This leads to a cycle of infection and inflammation associated with a progressive reduction in lung function and eventual respiratory failure. A common feature of CF is the heightened, chronic inflammatory response to Pseudomonas aeruginosa (P. aeruginosa), driven by constitutive NF-κB activation in airway and peripheral blood cells (2, 3, 11). Primary nasal epithelial cells (PNECs) from patients with the common F508del/F508del mutation and a milder genotype (R117H/F508del), show a significant increase in NF-κB(p65) which correlates with disease severity (12).

A20 (TNFAIP3) is a central negative regulator of NF-κB activation following stimulation of TLRs and/or TNF-receptor and regulates different signalling pathways such as NF-κB and interferon regulatory factor (IRF) signalling (13). A20 modifies infection. The effect of the predicted drugs on A20 and NF-κB (p65) which correlates with disease severity (12).

Significance

This study reports that publicly available gene array expression data together with statistically significant connections' map (ssCMap) can successfully predict already licensed drugs to modify genes of interest. We applied this bioinformatics approach to the NF-κB regulator A20 (TNFAIP3), which is reduced in Cystic Fibrosis (CF) airway cells. ssCMap predicted drugs that should induce A20 and normalise the inflammatory response in CF. Using airway epithelial cells we show that ikarugamycin and quercetin have anti-inflammatory effects mediated by induction of A20. Using siRNA we confirm that the anti-inflammatory effect of ikarugamycin and quercetin are mainly due to A20 induction. We have identified a process whereby already licensed drugs can be successfully repositioned for chronic inflammatory airway diseases.

Reserved for Publication Footnotes

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Table 1.

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<th>ID of data set</th>
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<th>Platform used</th>
<th>Sample size</th>
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Table 2.

<table>
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<td>A20</td>
<td>Ubiquitination, negative regulator of NF-kB</td>
</tr>
<tr>
<td>ATF3</td>
<td>202672_s_at</td>
<td>Activating Transcription Factor 3</td>
<td>Binds the cAMP response element (CRE), transcriptional repressor (promoters with ATF sites)</td>
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<td>RAB5C small GTPase</td>
<td>201140_s_at</td>
<td>RAB5C</td>
<td>small GTPase, regulates membrane traffic from plasma membrane to early endosomes, ubiquitously expressed</td>
</tr>
<tr>
<td>DENND4A</td>
<td>214787_at</td>
<td>DENN/MADD domain containing 4A</td>
<td>C-myc promoter-binding protein, promotes exchange of GDP to GTP, converting inactive GDP-bound Rab proteins into their active GTP-bound form</td>
</tr>
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</table>

classical immune cells (14, 15) as well as epithelial cells (12), endothelial cells (16), embryonic fibroblasts (17), osteoclasts (18) and pancreatic beta-cells (19) and diverse roles for A20 in innate immunity, apoptosis, autophagy and antigen processing (13, 15, 16, 20). Within the innate immune response, A20 regulates NF-kB signalling at the level of TRAF6 in mouse embryonic fibroblasts (MEFs) and osteoclasts (17, 18). In cultured human airway epithelial cells, A20 is rapidly induced by viral or bacterial compounds (21) and is essential for termination of the TLR4 signal (22). PNECs stimulated with P. aeruginosa LPS show a transient increase in A20, but CF PNECs display lower A20 expression basally and after LPS stimulation (12, 23).

Therefore, A20 induction should have anti-inflammatory effects within the tightly regulated NF-kB signalling pathway as shown by the induction of A20 through giberellin (GA3) in airway epithelial cells. GA3 induced A20, reduced IL-8 secretion, stabilised cytosolic IkBα and reduced NF-kB (p65) activation (24). Here we set out to identify additional compounds able to induce A20. Thus, we performed a compound search using gene expression connectivity mapping to identify existing drugs that could induce A20 expression.

Results

1. Connectivity mapping (sscMap):

The selection of gene array data and creation of the gene signature

Data sets that passed the selection criteria contained human primary nasal epithelial cells (PNECs) and the human bronchial epithelial cell lines CFBE410-13a, Calu-3 and IB3-1 analysed basally and after exposure to P. aeruginosa LPS (Table 1). In total 76 samples from 4 different published gene array data sets were used. Linear expression correlation and GO enrichment analysis for NF-kB pathway genes identified the closest correlates to A20.

Table 2 shows the top 7 genes that subsequently served as the input to the connectivity mapping process.

Prediction of drugs to induce A20 in airway epithelial cells

This study sought small molecular compounds that may enhance A20 expression and as a confirming negative control, those compounds that may inhibit A20 expression. Table 3 summarizes the top candidate drugs identified. The column entitled ‘significance’ shows the significance of drugs based on p values and the column ‘z-score’ shows the correlation of the drugs with the input gene signature. Positive z-scores indicate a positive correlation i.e. the input genes are induced when treated with the particular drug. The significant drugs with the highest positive z-scores along with a negative control were selected for laboratory validation. In addition to p values and z scores, stability of the connections were measured by altering the gene signature and the significance of the connections are given under the column ‘perturbation stability’. Drugs with perturbation stability 1 represent strong connections which remain significant with ‘perturbation’ gene signatures. From these predictions two A20
Fig. 1. Gene expression profile of the gene signature genes associated with A20. 16HBE14o- (grey) and CFBE41o- (green) were stimulated (LPS, 10 μg/ml 0-24 h) and mRNA levels of A20, ATF3, Rab5c, DENND4A, POM121, ICAM-1 and PSNE1 determined as described. * significant difference compared to medium control, * significant differences between genotypes.

inducing drugs (ikarugamycin and quercetin) as well as one non-A20 inducing drug (fluvastatin) were chosen for further investigation.

2. Gene expression of gene signature

Expression of the genes identified as the A20/NF-kB gene signature in CF epithelial cells were analysed by qRT-PCR in 16HBE14o- and CFBE41o- cultured in the presence or absence of LPS for 0-24h (Figure 1).

Expression of the genes identified as the A20/NF-kB gene signature in CF epithelial cells were analysed by qRT-PCR in 16HBE14o- and CFBE41o- cultured in the presence or absence of LPS for 0-24h (Figure 1).

Fig. 2. A20 and p65 mRNA expression of ikarugamycin, quercetin and fluvastatin treated cells. 16HBE14o- and CFBE41o- were pre-incubated with (a) ikarugamycin, (0.01, 1 μM), (b) quercetin (0.1, 100 μM) or (c) fluvastatin (0.1, 1 μM) stimulated (LPS, 10 μg/ml, 0-24 h), A20 mRNA determined (qRT-PCR) and expressed as A20/β-actin relative to the internal control.

LPS induced expression: In 16HBE14o-, A20 mRNA is rapidly induced with expression peaking 1h after LPS exposure (p<0.001 compared to medium, n=5), while CFBE41o- show significantly lower (at 1h p<0.001 vs. 16HBE14o-, n=5) and delayed (maximal induction at 4-8 h, p<0.01 and 0.001 vs. medium, n=5) induction upon LPS stimulation. After LPS, ATF3 and ICAM1 expression was significantly lower in CFBE41o- compared to 16HBE14o- (p<0.01, n=5). Pom121 and PSNE1 expression increased in CFBE41o- compared to medium (8h, p<0.05 and p<0.01) and in CFBE41o- compared to 16HBE14o- (8h, p<0.05 and p<0.001). 16HBE14o- showed a significant reduction in DENND4A and Rab5c expression compared to medium (1h, p<0.05 and 4h, p<0.05, respectively), but there was no significant change in the expression of these genes in CFBE41o-.

3. Effect of A20 inducing drugs on cell lines

LDH release in drug exposed 16HBE14o- and CFBE41o-: LDH release was measured after exposure to the drugs alone (0.01–1000 μM) and with LPS stimulation (supplement Figure 1S).

Quercetin did not cause any LDH release. Overall fluvastatin was almost without effect on LDH; the exceptions were a slight but statistically significant increase at 10 μM alone (CFBE41o-) and in the presence of LPS (in CFBE41o- and 16HBE14o-). Ikarugamycin (1, 100 μM) caused a significantly higher LDH release in both cell types. In LPS stimulated cells, 0.1 μM ikarugamycin and higher concentrations showed a higher LDH release compared to LPS alone, but this did not reach statistical significance (supplement Figure 1S).

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Malcolmson et al. 2015 Connectivity mapping to predict A20 inducing drugs - Figures

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Fig. 3. Effect of ikarugamycin, quercetin and fluvastatin on A20 and p65 protein expression. 16HBE14o- and CFBE41o- were pre-incubated with (a) ikarugamycin (0.01, 1 μM), (b) quercetin (0.1, 100 μM) or (c) fluvastatin (0.1, 1 μM) and stimulated (LPS, 10 μg/ml, 0-24 h). Cytosolic A20 and p65 protein was determined by Western Blotting: 1 = Ctr, 2 = LPS, 3 = drug at lower conc., 4 = drug at lower conc. + LPS, 5 = drug at higher conc., 6 = drug at higher conc. + LPS.

LPS stimulated IL-8 release in drug pre-treated 16HBE14o- and CFBE41o-: To assess the anti-inflammatory potential of the selected drugs, cells were pre-treated with the drug for 1h, stimulated with LPS and the IL-8 release measured and the relative IC_{50} calculated (supplement S3). In 16HBE14o-, all drugs reduced IL-8 release by at least 50% with an IC_{50} of 15.6 μM for ikarugamycin, 0.09 μM for quercetin and 0.11 μM for fluvastatin. In CFBE41o-, only quercetin (IC_{50} 0.03 μM) and ikarugamycin (IC_{50} 0.001 μM) pre-treatment were able to reduce release by 50%. In contrast, pre-treatment of CFBE41o- with 1 μM ikarugamycin caused a significant increase in IL-8 release compared to LPS alone (LPS 269.9 ± 47.9 pg/ml vs. 590.7 ± 82.6 pg/ml, p<0.05, n=5). Therefore, a meaningful calculation of the relative IC_{50} for IL-8 release in ikarugamycin treated CFBE41o- cells was not possible.

A20 mRNA induction in drug treated 16HBE14o- and CFBE41o-: To elucidate if ikarugamycin and quercetin facilitate their anti-inflammatory action though the induction of A20 as predicted, A20 mRNA was determined by qRT-PCR. Fluvastatin was included as a negative control. Using the LDH and IL-8 release data two drug concentrations were selected for further investigations (supplement S3). In 16HBE14o-, LPS stimulation caused a significant induction of A20 1h after stimulation (Figure 2). Ikarugamycin (0.01 μM) alone did not cause a significant induction of A20, but additional LPS stimulation caused a significant A20 induction at 24h (p<0.05 vs LPS 24h). 1 μM ikarugamycin significantly induced A20 at 4h on its own, but the higher A20 expression in the presence of LPS (1-24h) did not reach statistical significance. Quercetin pre-treatment did not induce significant levels of A20 mRNA at 0.1 μM, alone or in the presence of LPS. However, at 100 μM quercetin A20 mRNA was significantly induced alone at 1h and 4h (p<0.05 vs. medium 1h and 4h) and in the presence of LPS at 4h (p<0.05 vs. LPS 4h). Fluvastatin alone did not induce A20 mRNA at any time or concentration (Figure 2), in the presence of LPS fluvastatin pre-treatment caused a significant reduction in A20 mRNA at both concentrations tested (p<0.001 for 0.1 μM+LPS 1h vs. LPS 1h; p<0.05 for 1 μM+LPS

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Fig. 5. Effect of ikarugamycin, quercetin and fluvastatin on IL-8 release from PNECs from (a) healthy controls and (b) patients with CF. The release of IL-8 (pg/ml) was determined using a commercially available IL-8 ELISA kit. Statistical analysis was performed using Wilcoxon paired ranked t-test.

Effect of selected components on NF-κB (p65) mRNA in 16HBE14o- and CFBE41o-: Next we investigated if A20 induction altered NF-κB(p65) mRNA levels (Figure 2). LPS stimulation caused a significant induction of p65 1h after stimulation in 16HBE14o- (p<0.01 – p<0.001 vs. medium). Ikarugamycin (0.01 μM) was without significant effect on p65 mRNA. At 1 μM, ikarugamycin alone induced p65 at 24h (p<0.01 vs. medium 24h) and in the presence of LPS at 1h and 4h (both p<0.05 vs. LPS). Quercetin alone showed no effect on p65 mRNA levels, but when stimulated with LPS both concentrations of quercetin (0.1, 100 μM) significantly induced p65 at 24h (p<0.05 vs. medium 24h; p<0.01 for 1 μM+LPS 24h vs. LPS 24h).

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Fig. 6. Effect of ikarugamycin, quercetin and fluvastatin on A20 and p65 mRNA expression in PNECs from (a) healthy controls and (b) patients with CF. Cells were pre-incubated with ikarugamycin, quercetin or fluvastatin at the indicated concentrations (0.01–100 μM) and then stimulated (LPS, 10 μg/ml, 0–24 h). A20 mRNA was determined by qRT-PCR and expressed as A20/β-actin relative to internal control.

Table 3.

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<th>Drug</th>
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<th>Perturbation stability</th>
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<td>HMG-CoA reductase inhibitor</td>
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μM) significantly reduced p65 mRNA levels at 1h (p<0.01 vs LPS 1h). Similar to quercetin, fluvastatin alone showed no effect.
on p65 mRNA levels, but after LPS stimulation fluvastatin (0.1, 100 µM) significantly reduced p65 mRNA levels at 1h (p<0.001 and p<0.05 vs LPS 1h). In CFBE41o-LPS significantly induced p65 at 1h, 4h and 24h (p<0.05-0.001 vs medium) and overall CFBE41o-exhibited higher expression levels of p65 at 4h and 24h than 16HBE14o- (Figure 2). Ikarugamycin was without significant effect on p65 mRNA expression at any concentration or time point, although overall expression levels appear higher at 1 µM, when stimulated with LPS (Figure 2). In CFBE41o-, quercetin (0.1 µM) did not affect p65 mRNA levels. At 100 µM, quercetin significantly induced p65 (p<0.05; quercetin alone 100 µM 24h vs. medium 24h and quercetin 100 µM + LPS 24h vs. LPS 24h). Fluvastatin caused a significant reduction in p65 mRNA at both concentrations and all time points after LPS stimulation (p<0.05 and p<0.01 for 0.1 µM+LPS vs. LPS; p<0.05 and p<0.01 for 1 µM+LPS vs. LPS) (Figure 2).

Effect of predicted drugs on A20 and p65 protein expression:

We then determined the effect of the drugs on cytosolic A20 and p65 protein by Western Blotting using the same selected concentrations than before. Ikarugamycin (0.01 µM) induced A20 protein in both 16HBE14o- and CFBE41o-, with less A20 protein induction showed similar results with a significant IL-8 reduction at both cell types (Figure 3a). Quercetin treatment caused a strong induction of A20 protein at both concentrations (0.1, 100 µM) in 16HBE14o- and to a lower degree in CFBE41o-. Quercetin (100 µM) reduced cytosolic p65 in 16HBE14o- and in CFBE41o- (Figure 3b). Fluvastatin did not induce A20 protein at either concentration (0.1, 1 µM) in both 16HBE14o- and CFBE41o-cells. Fluvastatin pretreatment reduced cytosolic p65 protein in 16HBE14o- cells though this was only apparent at the higher concentration in CFBE41o- (Figure 3c).

Specificity of the drug effect on A20 mRNA expression using A20 siRNA: To confirm that the effect of the selected drugs is facilitated through A20 induction, we used siRNA to knock down A20 expression in 16HBE14o- cells as previously described (30). Cells were pre-treated with quercetin or ikarugamycin prior to LPS and IL-8 determined. Results (Figure 4) showed that in 16HBE14o-LPS significantly induced IL-8 (p<0.05 compared to untreated control), but when A20 was knocked down IL-8 increased further (although not significantly different from LPS alone). When cells are pre-treated with quercetin (100 µM) or ikarugamycin (1 µM), the LPS induced IL-8 release was significantly reduced (p<0.05). However, when A20 is knocked down IL-8 levels were different from LPS control (Figure 4).

Effect of A20 inducing drugs on PNECs Effect on IL-8 release:

LPS significantly induced IL-8 release from PNECs from non-CF and CF patients (non-CF; p<0.01, CF; p<0.05, Wilcoxon paired test, Figure 5). IL-8 release from CF PNECs was significantly higher than from non-CF PNECs (600.6±36.1 pg/ml vs. 315.8±62.8 pg/ml; p<0.01, Mann-Whitney test). In non-CF PNECs (Figure 5a), pre-treatment with ikarugamycin at 0.01 µM increased A20 mRNA which reached significance at 1h, 4h and 24h (both p<0.05 vs medium) and additionally above LPS induction (1h and 24h, both p<0.05 vs LPS). Similar to non-CF PNECs, fluvastatin treatment of CF PNECs had no significant effect on A20 mRNA expression levels.

NF-κB (p65) induction in PNECs: PNECs from non-CF control subjects respond to LPS exposure with a significant increase in NF-κB (p65) at 1h and 4h (p<0.01 vs medium control). Thereafter, p65 mRNA expression returns to its corresponding medium control value (Figure 6). PNECs from patients with CF however show the expected high levels of p65 throughout the 24h studied (p<0.01 at 1h, p<0.05 at 24h and medium control) (Figure 6).

In non-CF PNECs, ikarugamycin treatment alone did not change p65 mRNA levels (vs medium control), but in the presence of LPS p65 mRNA was significantly reduced (p<0.05 at 1h, p<0.01 at 4h vs LPS). Overall the higher concentration of ikarugamycin induced p65 mRNA levels with a significant increase at 4h (p<0.05 vs medium control). However, in the presence of LPS, p65 levels remained not significantly different from those after LPS exposure at 1h and 24h, but were significantly lower compared to LPS alone at 4h (p<0.05) (Figure 6). The lower concentration of 0.01 µM quercetin alone did not modify basal p65 mRNA. After subsequent LPS challenge p65 mRNA significantly decreased at 1h (p<0.01 vs LPS), and p65 levels increased in a similar manner to LPS alone. However, 100 µM quercetin alone significantly reduced p65 induction at 1h and 4h (p<0.05 and p<0.01 vs. medium control). In the presence of LPS this reduction of p65 mRNA reached statistical significance at 4h and 24h (both p<0.05 vs LPS) (Figure 6). Fluvastatin (0.1 µM) induced p65 at 4h (p<0.05 vs. medium 4h), but p65 levels remain significantly lower when LPS is added (p<0.05, vs LPS p<0.01 vs LPS). Similarly, quercetin treatment with the lower concentration (0.01 µM) alone and in the presence of LPS had no effect on A20 mRNA levels (Figure 6), while 100 µM quercetin significantly induced A20 mRNA alone (4h, p<0.05 vs medium control) and additionally above LPS induction (1h and 24h, both p<0.05 vs LPS). Similar to non-CF PNECs, fluvastatin treatment of CF PNECs had no significant effect on A20 mRNA expression levels.

NF-κB (p65) induction in PNECs: PNECs from non-CF control subjects respond to LPS exposure with a significant increase in NF-κB (p65) at 1h and 4h (p<0.01 vs medium control). Thereafter, p65 mRNA expression returns to its corresponding medium control value (Figure 6). PNECs from patients with CF however show the expected high levels of p65 throughout the 24h studied (p<0.01 at 1h, p<0.05 at 24h and medium control) (Figure 6).
0.1 μM quercetin in the presence of LPS significantly reduced p65 at 24h (p<0.05 vs LPS). Treatment with 100 μM quercetin significantly reduced p65 mRNA in the presence of LPS at all time points (all p<0.05 vs LPS) (Figure 6). Fluvasatin alone had no significant effect on p65 mRNA levels, but significantly reduced LPS induced p65 at 1h, 4h and 24h (p<0.05 vs. medium 1h, 4h or 24h), while the higher concentration of fluvasatin (1 μM) showed no significant effect on basal or LPS induced p65 mRNA (Figure 6).

Discussion
Airways infection and the subsequent inflammation are deleterious for patients suffering from CF. Current drugs targeting the mutated CFTR (potentiators/correctors) improve expression and function of CFTR on epithelial surfaces and patients showed improved lung function and reduced frequency of pulmonary exacerbations, hospitalization and use of intravenous antibiotics, but augmented CFTR function failed to reduce inflammatory markers in sputum (e.g. IL-1,-6,-8) (25) and heterogeneous responses to the treatment have been reported (26), suggesting that CFTR correction/potentiation may not directly improve the underlying compromised immune responses. The negative NF-κB regulator A20 (TNFAIP3) is reduced in CF airway epithelial cells, basally and after LPS stimulation (23) and is associated with markers of inflammation and decreased lung function (12). A20 silencing increased TRAF6 and NF-κB activity (18), and A20 over-expression had protective effects in airway inflammation in ‘asthmatic mice’ (27), suggesting that A20 augmentation normalises the inflammatory response in the airways.

In order to find agents to induce A20 in CF we employed sscMAP, which has been widely used in drug development uncovering potential new indications for existing drugs as well as predicting side effects (28). Using disease specific publicly available gene array data ( GEO data sets), we used connectivity mapping to firstly identify the target gene (A20) related gene signature and to secondly predict already licensed drugs to induce A20 expression. We included a total of 76 gene array data from primary nasal epithelial cells and cell lines commonly used in CF research (Table 1). Gene array databases were first selected in August 2013, but a recent (Jan 2016) search revealed no further significant published gene array data on CF (primary nasal) epithelial cells.

The applied linear regression model (Pearson’s correlation coefficient) is an established robust method to identify the correlates of a known gene expression estimating the strength of a linear relationship between two random normally distributed variables (29). The application of the linear regression model with GO (gene ontology) selection revealed a gene signature of 6 genes additionally to the seed gene A20 (Table 2) and we confirmed that the expression of these genes is similarly reduced in CF epithelial cells, basally and after LPS stimulation (Figure 1). The identified A20 correlates were AFT3, a transcriptional repressor that binds to CAMP response elements (CRE); Rab5c, a small ubiquitously expressed GTPase; DENND4A, which encodes the C-Myc Promoter Binding Protein (MBP-1); POM121, a nuclear transmembrane protein and essential component of the nuclear pore complex; ICAM1, a cell surface glycoprotein typically expressed on endothelial and immune cells, especially during inflammation and PSEN 1 (Presenilin 1), a catalytic component of γ-secretase and a DREAM binding protein. Further descriptions of these genes and their involvements in inflammation can be found in the online appendix. These genes, as a combined gene signature, were then input into the sscMAP process comparing the gene expression of the gene signature with the gene expression in the reference database (www.broadinstitute.org), which was obtained from systematic microarray gene expression profiling.

ScsMAP predicted a short list of drugs that should modify the expression profile of the gene signature genes, including A20. Those drugs included azacyclonel, ikarugamycin, quercetin and karakoline (Table 3). Azacyclonel is a drug used in psychotic individuals (30). We excluded azacyclonel, as its use requires special permission through relevant governmental authorities. Interestingly, the anti-histamine terfenadine is metabolised to azacyclonel and terfenadine (31). Karakoline is a highly toxic plant diterpenoid (32) and the pharmacological effects of preparations of Aconitum roots are attributed to diterpenoid alkaloids (33). The anti-inflammatory activity of gibberelin (GA3), also a plant-derived diterpenoid, is mediated through A20 induction (24). We therefore selected ikarugamycin and quercetin for further studies.

Ikarugamycin is a macrolide antibiotic with cytostatic effects against Gram-positive bacteria. We show that ikarugamycin exhibits anti-inflammatory properties in LPS stimulated airway cells. In 16HBE14o- ikarugamycin showed a dose-dependent reduction of LPS-induced IL-8 release (supplement S4), through induction of A20 and reduction of p65 (Figure 2). 16HBE14o- and CFBE41o- did not show reduced cell viability at concentrations lower than 1 μM, higher concentrations increased LDH release suggesting a cytotoxic effect (supplement S2). CFBE41o- appear more sensitive to ikarugamycin treatment (supplement S4), which made it not possible to calculate a meaningful relative IC50 value (supplement S3), although p65 protein expression was not increased (Figure 2). In HL-60 cells, ikarugamycin reduced cell viability and increased DNA fragmentation starting at 0.1 μM (IC50 of 0.22 μM), while MCF-7 cells and peripheral blood mononuclear cells showed higher resistance. Furthermore, ikarugamycin treatment of HL-60 cells caused a significant caspase activation, increase in intracellular calcium and p38 MAP kinase activation (34). However, investigating the pro-apoptotic mechanisms in bronchial epithelial cells was beyond the scope of this study. Nonetheless, our ikarugamycin data at near cytotoxic levels add valuable information: Firstly, sscMap correctly predicted that ikarugamycin would induce A20 mRNA but sscMap does not predict the physiological effect of the gene induction. CF cells overall show a limited ability to induce A20, however, our results show that – given the right stimulus – CF cells are indeed able to induce A20 mRNA and the high induction of A20 at near cytotoxic levels may be able to counteract the pro-apoptotic stimulation of ikarugamycin.

Quercetin, a flavonoid, is known for its anti-inflammatory effects. In vivo studies have shown antioxidant, anti-inflammatory, anti-tumour and even anti-infectious properties of quercetin, which are predicted through its anti-inflammation signaling pathways such as NF-κB (35). In lung epithelial cells, quercetin inhibited IL-1 and TNF-α induced iκBα degradation and NF-κB activity through modification of the MAPK pathway (AP-1) (36). ScsMap correctly predicted that quercetin could induce A20 mRNA, adding a new mechanism for the anti-inflammatory effects of quercetin. It also significantly reduced LPS-induced IL-8 release in both cell types with a relative IC50 of 0.15 and 0.04 μM in 16HBE14o- and CFBE41o-, respectively. Quercetin at concentrations up to 1000 μM did not show any cytotoxicity, although in neuronal cell cultures quercetin higher than 100 μM was cytotoxic (37). Within the in vivo antioxidant network, quercetin has been described to be oxidised and to yield an ortho-quinone, which, in absence of reducing glutathione, can oxidise protein thiols impairing enzyme activities (38). We have not investigated the antioxidant status of our cell culture, but we took precautions to minimise oxidation when preparing our quercetin dilutions.

To further investigate the A20-dependent mechanism of the anti-inflammatory action of quercetin and ikarugamycin, we used A20 knock down in 16HBE14o-. As previously described for the A20 inducing anti-inflammatory compound gibberelin (24), we were able to confirm that the anti-inflammatory effects of the
predicted drugs was indeed mainly mediated by the induction of A20.

We also tested fluvastatin, which was predicted not to affect or reduce A20 gene expression (negative z-score). Although fluvastatin exerted anti-inflammatory effects (IL-8) in both cell lines, our data show that this was not mediated by the induction of A20 (mRNA), clearly confirming the sscMap prediction. In asthma, fluvastatin inhibits eosinophil adhesion to ICAM-1 (39) and fibroblast proliferation (40). Using similar concentrations we did not observe any reduced proliferation. Fluvastatin at a concentration range similar to those we used, reduced basal and LPS-induced IL-8 release from LPS stimulated whole blood cells, with CF cells appearing more sensitive to fluvastatin than control cells (IC50: 19.1 μM in non-CF cells, 4.6 μM in CF blood cells) (41). In isolated LPS-stimulated peripheral blood monocytes from patients with chronic kidney disease fluvastatin had a significant anti-inflammatory effect (IL-8, IL-6) at a concentration range of 0.0001-1 μM (42). Patients with heart transplants receiving 40 mg fluvastatin/day for 4 weeks showed a significant reduction in total cholesterol levels and a maximum blood fluvastatin concentration of 2.11 and 3.77 μM. These studies suggest that we have covered a physiologically relevant range of fluvastatin. However, fluvastatin metabolism may be affected by concomitant therapies, especially substances competing with cytochrome enzymes and in such cases fluvastatin levels may need to be monitored (43). Any reactions with other therapies (as they would appear in patients with CF) were not investigated in our manuscript as they would have been beyond the scope of the study.

Similar to ikrugamycin, fluvastatin has been described to have pro-apoptotic effects e.g. in human lymphoma cells, human smooth muscle cells and in rat neonatal cardiac myocytes or rat vascular smooth muscle cells (44, 45), mediated through activation of caspase-3, reactive oxygen species and activation of p38 MAPK (44, 55). However, statins, through their inhibition of 3-hydroxy-3-methylglutaryl-CoA (HMG-CoA), but not through induction of A20, may still have therapeutic potential in airway and systemic inflammation in CF (41).

Overall, our study shows that connectivity mapping (sscMap) can predict A20 inducing drugs. Pre-treatment of cells with both ikrugamycin and quercetin reduces LPS-induced IL-8 secretion by induction of A20. In non-CF PNECs both drugs upregulated A20 and reduced IL-8 and p65 mRNA at lower concentrations than in cell lines. CF PNECs, however, have a reduced and delayed A20 inducing response to LPS and to the tested drugs and a significant A20 induction appears at the higher concentration of the drugs tested, which might be near to the cytotoxic effect. A20 reduces apoptosis (46, 47) and mutational loss of A20 resulted in rapid apoptosis and inflammation in hematopoietic cells (48). We did not determine markers of apoptosis in our study, but the huge increase in A20 mRNA may indicate a possible counteraction to pro-apoptotic changes in response to ikrugamycin treatment. Of particular interest, this may indicate a higher susceptibility/sensitivity of CF cells to pro-apoptotic stimuli.

Our study has several limitations. Firstly, for the sscMAP process a huge number of gene array samples are required and although the database search gave a high number of initial results, upon detailed inspection several gene array studies could not be included. Connectivity mapping uses gene array data run on Affimetrix platforms and we selected those performed on these platforms. Several published gene array studies were performed in cell lines. However, the majority of samples selected were PNECs (n=40), but we also included data from cell lines. Furthermore, every published dataset has been performed using a specific experimental design with respect to treatments and time points. We selected experiments that used either no stimulation or exposure to *P. aeruginosa* LPS or to *P. aeruginosa* itself. A sample size of 50-100 individual samples is a statistically acceptable sample size to produce an unbiased result and we used 76 individual gene array samples.

Secondly, the reference database was generated using different cell lines: MCF7, IL60, PC3 and SKMEL5. Although all of human origin, none of these cell lines are airway derived. We therefore confirmed the effect of the predicted drugs in airway relevant and disease specific cell lines, determined an effective drug concentration in our disease model and confirmed their effect in primary cells. Additionally, factors such as the interaction between various signalling pathways and the interplay between genes can affect the functions of the predicted and validated drugs when used in humans.

The aim of our study was to investigate the potential of sscMap to predict A20-inducing drugs from a list of drugs already licenced for the use in humans to make them available for drug repositioning. As a proof of concept we have focused on the LPS induced expression of A20, p65 and cytokines IL-6 and IL-8. However, in addition to its direct regulation of TLR-induced NF-κB activation, A20 is also involved in the negative regulation of the of the NFκB3 inflammasome via TRIF (TRIF)-RIPK3 (49) may also inhibit inflammation induced regulated necrosis (Ripk3 via RIPK3 (50), adding further levels of action and complexity to the anti-inflammatory action of A20. Therefore, future work analysing further NFκB driven cytokines such as TNFα and IL-1β would also indicate if the predicted drugs are able to modify A20 action on the inflammasome.

**Summary**

To date there is still a need for alternative anti-inflammatory drugs for patients with CF as restoring CFTR function with potentiators and correctors does not directly affect the inherent innate immune defect. The exaggerated inflammatory response is in part due to the lack of the NF-κB regulator A20 and pharmacological induction of A20 is anti-inflammatory. We have shown here that sscMap is a potent tool to predict effective drugs that can modify A20 without totally inhibiting NF-κB. This is particularly important in the clinical setting as pharmacological suppression of inflammation may increase the incidence of infective exacerbations (51). In addition, A20 inducing drugs have to be carefully adjusted as in addition to the A20 induction e.g. to date there is still a need for alternative anti-inflammatory drugs for patients with CF as restoring CFTR function with potentiators and correctors does not directly affect the inherent innate immune defect. The exaggerated inflammatory response is in part due to the lack of the NF-κB regulator A20 and pharmacological induction of A20 is anti-inflammatory. We have shown here that sscMap is a potent tool to predict effective drugs that can modify A20 without totally inhibiting NF-κB. This is particularly important in the clinical setting as pharmacological suppression of inflammation may increase the incidence of infective exacerbations (51). In addition, A20 inducing drugs have to be carefully adjusted as in addition to the A20 induction e.g. to date there is still a need for alternative anti-inflammatory drugs for patients with CF as restoring CFTR function with potentiators and correctors does not directly affect the inherent innate immune defect. The exaggerated inflammatory response is in part due to the lack of the NF-κB regulator A20 and pharmacological induction of A20 is anti-inflammatory. We have shown here that sscMap is a potent tool to predict effective drugs that can modify A20 without totally inhibiting NF-κB. This is particularly important in the clinical setting as pharmacological suppression of inflammation may increase the incidence of infective exacerbations (51). In addition, A20 inducing drugs have to be carefully adjusted as in addition to the A20 induction e.g. to date there is still a need for alternative anti-inflammatory drugs for patients with CF as restoring CFTR function with potentiators and correctors does not directly affect the inherent innate immune defect. The exaggerated inflammatory response is in part due to the lack of the NF-κB regulator A20 and pharmacological induction of A20 is anti-inflammatory. We have shown here that sscMap is a potent tool to predict effective drugs that can modify A20 without totally inhibiting NF-κB. This is particularly important in the clinical setting as pharmacological suppression of inflammation may increase the incidence of infective exacerbations (51). In addition, A20 inducing drugs have to be carefully adjusted as in addition to the A20 induction e.g. to
gene expression reference profile was assessed via a connection score. The subsequent gene signature (including A20) was used to further filter the A20 correlates identifying those that meet the criteria of significance. The significant r values were selected as correlates of A20. Gene ontology (GO) enrichment analysis (http://geneontology.org/) was applied to the significant correlates of A20. Gene silencing was assessed by transfections (49, 50). Transfection efficiency was quantified. Equal amounts of RNA (250 ng) were reverse transcribed into cDNA. cDNA was used for qPCR. All data are presented as the means ± SEM. Differences between groups were analysed using the Kruskal-Wallis non-parametric ANOVA with Dunn’s post-test (*p < 0.05, **p < 0.01 and ***p < 0.001), s and Tukey’s multiple comparisons test. Rat lung fibroblasts (RLF, ATCC) were cultured as previously described (32) and given in supplement table S1. Relative expression of β-actin was assessed using primer sequences previously described (23) and in a previous study (55). Control participants did not have any acute airways disease at the time of recruitment. The authors would like to express their thanks to all CF patients and volunteers that took part in this study. BM and HW were in part supported by a BBSRC/MRC/EPSCR co-funded grant (BB/I00905/1). We thank Declan McGuan and Fiona Manzendorf Kovulia (University of Ulster) for technical assistance with the transfection experiments.


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