The Role of Isocitrate Lyase (ICL1) in the Metabolic Adaptation of Candida albicans Biofilms

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Abstract

Background: A major characteristic of Candida biofilm cells that differentiates them from free-floating cells is their high tolerance to antifungal drugs. This high resistance is attributed to particular biofilm properties, including the accumulation of extrapolymeric substances, morphogenetic switching, and metabolic flexibility.

Objectives: This study evaluated the roles of metabolic processes (in particular the glyoxylate cycle) on biofilm formation, antifungal drug resistance, morphology, and cell wall components.

Methods: Growth, adhesion, biofilm formation, and cell wall carbohydrate composition were quantified for isogenic Candida albicans ICL1/ICL1, ICL1/icl1, and icl1/icl1 strains. The morphology and topography of these strains were compared by light microscopy and scanning electron microscopy. FKS1 (glucan synthase), ERG11 (14-α-demethylase), and CDR2 (efflux pump) mRNA levels were quantified using qRT-PCR.

Results: The ICL1/icl1 and icl1/icl1 strains formed similar biofilms and exhibited analogous drug-tolerance levels to the control ICL1/ICL1 strains. Furthermore, the drug sequestration ability of β-1, 3-glucan, a major carbohydrate component of the extracellular matrix, was not impaired. However, the inactivation of ICL did impair morphogenesis. ICL1 deletion also had a considerable effect on the expression of the FKS1, ERG11, and CDR2 genes. FKS1 and ERG11 were upregulated in ICL1/icl1 and icl1/icl1 cells throughout the biofilm developmental stages, and CDR2 was upregulated at the early phase. However, their expression was downregulated compared to the control ICL1/ICL1 strain.

Conclusions: We conclude that the glyoxylate cycle is not a specific determinant of biofilm drug resistance.

Keywords: Biofilms, Isocitrate Lyase (ICL1), Resistance, Candida albicans

1. Background

Fungal pathogens are gaining increasing attention as infection rates increase due to advances in healthcare and rising numbers of susceptible immunocompromised patients (1,2), and their major impact on human health is becoming better appreciated. Despite advancements in the management of hospital-acquired infections, Candida albicans, a commensal organism that helps maintain microbiota normalcy, is the fourth most common isolated nosocomial bloodstream pathogen, after Staphylococcus aureus, accounting for approximately 80% of candidemia cases with a 40% fatality rate (3-9). These microbes exist predominantly as attached aggregated communities in nature, forming an irreversible assemblage that is strongly influenced by the growth medium, the attachment surfaces, and the environment. Microcolonies of cells are sheltered in self-secreted polymeric substances and communicate via quorum sensing (10-12).

Biofilm growth is a unique virulence trait and defense mechanism of C. albicans that is associated with infectious diseases and the clinical use of implanted medical devices, such as central catheters and intracardiac prosthetic valves. Candida albicans causes life-threatening conditions, such as urinary tract infection and endocarditis, with a mortality rate as high as 30% (10, 13, 14). Despite treatment, such an infection may re-establish itself. Sessile cells are phenotypically and genetically different from free-floating cells. A characteristic attribute of biofilms, unlike planktonics, is resistance to antifungal agents, particularly azole, polyenes, and recently, echinocandins. Candida albicans sessile cell resistivity is always expressed, irrespective
of definite phenotypic variation and the expression of specific drug-resistance genes, entailing complex multicellular processes (11, 15, 16). Due to this property, in most clinical cases, replacement of the infected devices is the only treatment option (13, 17, 18). Thus, it is imperative to develop effective strategies for control. Recent research has begun to shed light on how attached microbial communities develop resistance to antifungal agents, proposing various involved mechanisms (13, 14). This peculiarity has been reported by previous studies to be a multifactorial phenomenon not regulated by a singular mechanism, which was confirmed by this study.

2. Objectives

In this study, we examined whether *C. albicans* biofilm development depends on the glyoxylate pathway since this is essential for the assimilation of many alternative carbon sources in the absence of glucose, and we investigated whether this pathway contributes to the high tolerance of biofilm cells to antifungal drug therapy. In addition, we examined whether a correlation exists between cell carbohydrates and matrix glucan concentrations during biofilm development.

3. Methods

3.1. Strains

*Candida albicans* wildtype strain ICL1/ICL1, heterozygous mutant ICL1/icl1, and homozygous mutant icl1/icl1 were obtained (Table 1). Strains were stored in 15% (vol/vol) glycerol at -80°C and routinely maintained on YPD (1% yeast extract, 2% peptone, and 2% dextrose; Sigma-Aldrich, USA) medium inoculated with single colony cells from overnight-grown stock culture, in an orbital shaker at 200 rpm and 35°C. The overnight grown wildtype, single-knockout, and double-knockout *C. albicans* cells were harvested by centrifuging at 3,000 rpm for 5 minutes at 4°C (3×). The cells were standardized using the spectrophotometer to OD600 of 0.1. Next, 100 µL of the standardized inoculum was pipetted into the selected wells and incubated for various time intervals, depicting their developmental growth phases (6, 24, and 48 hours) at 37°C. This experiment was performed in 2-4 replicates.

Biofilm cultivation on cellulose filter paper (15 mm in diameter, 0.2 µm pore size) was carried out using flat-bottomed 6-well plates. Strains were inoculated into 100 mL of YPL, YPD, and RPMI 1640 supplemented broths and incubated overnight in an orbital shaker at 35°C and 200 rpm. After the centrifugation of the previously grown primary culture, the cell pellets were resuspended in sterile 10 mL 0.15 M PBS and cell density was standardized to 1 × 10^7 CFU/mL. Next, 2 mL of each inoculum was added to 6-well plates containing sterilized 15 mm filter membranes, and incubated for 90 minutes at 37°C for adhesion of cells to the membranes, without shaking. After the adhesion phase, the medium was carefully aspirated and the membranes were gently washed in 2 mL sterile 0.15 MPBS twice to remove non-adherent cells. The membranes were carefully transferred with a sterile forceps into new 6-well plates containing 2 mL of YPD and YPL plates containing 2 mL of YPD and YPL and supplemented with RPMI-1640. The plates were incubated for different time intervals (6, 24, and 48 hours) at 37°C, representing the early, intermediate, and mature phases, respectively, of biofilm formation.

### 3.3. MTT Reduction Assay

A colorimetric MTT reduction assay was performed as described by Hawser and Douglas in 1995 (22). Stock solution (0.5%, 5 mg/mL) was prepared by dissolving 5 mg of MTT salt into 1 mL of sterile PBS. Working solution was prepared by diluting the stock 1:10 (V/V).

### 3.4. Antifungal Susceptibility

For antifungal susceptibility testing of sessile cells, biofilms that formed on the microtiter plates at the various time intervals (6, 24, and 48 hours) were challenged with antifungal drugs. The media were aspirated off and gently washed twice with sterile PBS, then 100 µL of YPD and YPL...
broth and RPMI 1640 medium containing serially diluted concentrations of antifungals (fluconazole 256 µg/mL and amphotericin B 16 µg/mL) was added and the plates were incubated for another 24 hours at 37°C. The MTT reduction assay at 405 nm was performed as described (22). The experiment was performed in 2-4 replicates.

3.5. Biofilm Compound Light Microscopy

The developmental stages were examined using a compound light. At different time intervals, the biofilms grown in lactate medium were repeatedly washed with PBS and viewed at 20× magnification.

3.6. Biofilm Scanning Electron Microscopy (SEM)

The biofilms were fixed using McDowell-Trump fixative on cellulose filter paper. The membranes were serially dehydrated by ethanol washes and immersed in hexamethyldisilazane (HMDS) (Sigma-Aldrich, USA). After drying, the dehydrated substrates were mounted on the stub. The studs were sputter-coated with gold and palladium (Au/Pd 60/40).

3.7. Cell Wall Polysaccharide Isolation and Composition

Sessile cell soluble extracellular polysaccharides, insoluble extracellular polysaccharides, and intracellular polysaccharides were extracted using 1 M NaOH. Into all of the tubes containing sycamore extracellular polysaccharides (SEPS), insoluble extracellular polysaccharides (IEPS), and intracellular polysaccharides (IPS), 5 mL of absolute ethanol was added, then the tubes were incubated at -20°C for 30 - 60 minutes. All tubes were centrifuged and washed twice with pre-chilled 75% ethanol. The carbohydrate content of each fraction was measured using the phenol-sulfuric acid method, and the results were normalized to a standard curve.

3.8. Beta-1, 3-Glucan Measurement

The matrix glucan was measured using a limulus lysate-based assay as previously described by Nett et al. (14). Biofilms were dislodged and samples were sonicated for 5 minutes at 7 W in PBS, then centrifuged 3× at 10,000 rpm for 5 minutes at 4°C to separate the cells from the soluble extracellular polysaccharide with the matrix material. Beta-1, 3-glucan was measured using the Glucatell (1, 3)-β-D-glucan detection reagent kit.

3.9. RNA Isolation and RT-PCR Expression Analysis

At each developmental stage, the biofilms were washed 3× with PBS. TRI reagent was used for RNA extraction. Pre-chilled chloroform was added for separation, and 500 µL of pre-chilled isopropanol was added to the clear aqueous upper part for precipitation. The supernatant was removed and the resulting pellets were washed with 75% ethanol and resuspended in 40 µL of dH2O. Complementary DNA was generated from the RNA using a Tetro cDNA synthesis kit according to the manufacturer’s instructions. Expression levels of the desired genes were quantified using light-cycle 480 SYBR green kits according to the manufacturer’s instructions, with a small amount of optimization. Reactions were run at denaturation for 15 seconds and at melting temperature for 15 seconds, with an extension time of 30 seconds for approximately 40 cycles (Table 2) (23).

| Primer | Nucleotide Sequence (5’ - 3’) | Target mRNA | Product Size |
|--------|-----------------------------|-------------|--------------|
| CaFKS1 | F CACGATTGAGCCGTTGGTG | CaFKS1 | 169 |
| R TGCTTGCAATGAGAATG | |
| CaERG11 | F ATGGGGTCCGAATGTAATG | CaERG11 | 208 |
| R TTCTTTGAGCAGCATCACG | |
| CaCDR2 | F CCCAGGTCATGTGGGTAATG | CaCDR2 | 200 |
| R CAGCGACTCCAACATAGCA | |
| CaACT1 | F GTGTAAGCAGCCGATCCA | CaACT1 | 174 |
| R GTGTAAGCAGCCGATCCA | |

3.10. Statistical Analyses

Analyses were performed using SPSS 17.0 for Windows. One-way ANOVA was performed to analyze the differences among multiple means. According to the tests of homogeneity of variances, Bonferroni-Dunn was carried out for post-hoc multiple comparisons. P values of < 0.05 were considered statistically significant.

4. Results

4.1. Biofilm Formation is Independent of Isocitrate Lyase

The icl1/icl1 mutants formed biofilms, as revealed by MTT quantification, at each developmental stage. Interestingly, ICL1/icl1 and icl1/icl1 cells were metabolically active at
a similar level to the wildtype cells in the presence and absence of glucose. Figure 2A - 2C depict a significantly (P < 0.05) higher degree of formation in glucose and lactate media in the wildtype and icl1/icl1 strains than in ICL1/ICL1. However, metabolic activity increased with maturation in all strains. There was no significant difference among the three strains on RPMI 1640. At maturity, the strains’ growth displayed no obvious differences on YPL and YPD. This result indicates that ICL1 is not essential for growth under these conditions.

4.2. Resistance Phenotype is Independent of Isocitrate Lyase

We investigated ICL’s impact on C. albicans biofilm resistivity. Biofilms at each developmental phase on a non-carbohydrate medium (lactate) were treated with 256 µg/mL of fluconazole and 16 µg/mL of amphotericin B based on SMICs. Figure 3A - 3C show that the reference, ICL1/ICL1, and icl1/icl1 strains exhibited high resistance to fluconazole at the 256 µg/mL concentration at all stages (as well as in the YPD and RPMI media; data not shown). Growth was characterized by high metabolic activity that did not deviate significantly from untreated biofilms. As seen in Figure 3B - 3C, a 20% inhibitory effect was conserved with 16 µg/mL amphotericin B on ICL1/ICL1 and icl1/icl1 biofilms at 6 hours. However, resistance built up at the intermediate and mature phases, as seen in Figure 3B - 3C. In addition, resistance increased with metabolic activity regardless of the supporting medium and the in vitro materials used. This result indicates that the glyoxylate cycle is not essential for resistance to these antifungal drugs.

4.3. Biofilms Grow in Three Developmental Stages

Our results confirmed that biofilm development progresses in three distinct stages. The early phase (Figure 4A - 4C) showed free-floating cells attached to the substratum and to each other in microcolonies. The intermediate phase (Figure 4D - 4F) shows active replication, evident through increased numbers of microcolonies, with heterogeneous communities consisting of hyphae and pseudohyphae in the wildtype cells (Figure 4D). Filamentous morphogenesis was absent in the mutant biofilms (Figure 4E - 4I).

4.4. Isocitrate Lyase (ICL) Alters Morphogenesis

In an attempt to visualize the role of ICL on biofilm structural organization and morphology during challenges with antifungal drugs, mature biofilms of the reference and reconstructed mutants on cellulose membranes were treated with 256 µg/mL of fluconazole, then examined under SEM and compared to untreated biofilms. Selected concentrations were based on generated susceptibility testing data. The reference strain-treated biofilm (Figure 5A) conformed to previous descriptions [33]. This community presented a highly heterogeneous assemblage, with visible pseudohyphae, hyphae, and blastospores encased within the ECM and divided into two distinct layers. Blastospores were more concentrated and tightly packed at the basal region, while the upper region consisted of pseudohyphae and true hyphae networks. Untreated wildtype biofilms (Figure 5B) exhibited substantially less hyphae growth, consisting mostly of budding, densely packed, clustered yeast cells. Hyphae growth was incapacitated and defective in the reconstructed mutants, with no visible bilayer in either the treated or the untreated biofilms. The treated ICL1/ICL1 strain seen in Figure 5C showed resistance, as indicated by the steadiness of metabolic activity and the detectable cluster of yeast cells in the basal region. The untreated biofilm seen in Figure 5D was characterized by a mass cluster of viable cells with defective hyphae growth. The icl1/icl1 strain (Figure 5E) exhibited the same basal composition as ICL1/ICL1, but with smaller and denser adhered, mature, and budding blastospores. The generated images suggest that ICL influences morphogenesis under these conditions.

4.5. ECM β-1, 3-Glucan is Dependent on Cell Wall Polysaccharides

We designed a hypothesis to answer whether sessile cells generate glucan using alternative generated hexoses in the absence of the glyoxylate pathway in relative proportion to when glyoxylate is present. Also, we further explored the extent of glucan defenses against treatment at each developmental stage. Table 3 reveals that the cell wall carbohydrates of treated and untreated strains increased with time. Compared to the wildtype cells, hexose accumulation in untreated and treated icl1/icl1 and ICL1/ICL1 was not impaired despite the absence of ICL. These data indicate that hexoses were synthesized through other active pathways. Synthesis was more amplified in treated than in untreated biofilms, and the increment was proportional to time.

Sequestration activity through matrix β-1, 3-glucan encoded by the FKS1 gene mediates tolerance to azole drugs [40, 41]. To determine the β-1, 3-glucan concentration in the matrix, C. albicans wildtype, ICL1/ICL1, and icl1/icl1 biofilms on YPL medium were measured using a Glucatell (1, 3)-β-D-glucan detection reagent kit based on limulus lysate. Figure 6A - 6B show that the increase was significantly higher in wildtype and icl1/icl1 compared to ICL1/ICL1 at all stages. At the mature stages, ICL1/ICL1 showed a greater rate than the mutant strain, and the icl1/icl1 glucan level was moderately higher than that of ICL1/ICL1. After treatment, as de-
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Figure 1. Inverted Microscopy Images of C. albicans Biofilms at Different Growth Stages on RPMI 1640 Medium

These images depict biofilm formation on a polystyrene surface under static conditions through the three developmental phases. A, Growing predominantly as yeast cells, attachment of the cells to the potential growth material and to each other was observed; B, metabolic activity increased with cellular density in correlation with the duration of cultivation. This image shows cells at the maturation phase; C, Microcommunities became heterogeneous and clustered. Apparent production and visible accumulation of slime extracellular matrix (ECM) enclosing the sessile yeast and filamentous cells, which increased with maturity, was observed.

4.5. Transcript Profile of Associated Resistance Genes Regulated by Isocitrate Lyase

We investigated the extent of ICL’s regulatory role on FKS1, ERG11, and CDR2 at the mRNA expression level using real-time qPCR. ACT1 was used as a housekeeping gene. The expression profiles of ICL1/icl1 and icl1/icl1 were compared to that of the wildtype strain. Figure 7A shows that the expression level of FKS1 in the mutants was downregulated compared to the wildtype. FKS1 transcript levels were relatively high at the intermediate phase in the mutants, although no fold to the wildtype was suggested, and expression decreased at maturity. Expression of ERG11 in icl1/icl1 regressed over time, unlike that of ICL1/icl1, which increased at the intermediate stage and regressed at maturity (Figure 7B). The drug efflux-pump level was also reduced with maturation. The ICL1/icl1CDR2 level was higher than that of the wildtype at the early phase and decreased to reference levels at maturity. Overall, the transcript levels of the studied genes were downregulated to the wildtype, as shown in Figure 7C. Therefore, our data show that ICL exerts a substantial regulatory effect on FKS1, ERG11, and CDR2 expression.

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| Table 3. Comparison of Carbohydrate Concentrations of Biofilm Sessile Cell Walls Before and After Treatment |
| --- |
| Candida albicans strain | Untreated, µg/mL | Treated, µg/mL | Correlation With ECM Glucan |
| CaICL1/#CL1 | Increase with time | Lower concentration | Higher concentration |
| hours | 49.4 ± 0.0 | 49.3 ± 0.6 | + |
| hours | 97.0 ± 0.0 | 98.0 ± 0.1 | + |
| hours | 126.0 ± 0.5 | 125.2 ± 0.1 | + |
| CaICL1/#icl1 | Increase with time | Lower concentration | Higher concentration |
| hours | 26.0 ± 0.3 | 31.03 ± 1.0 | + |
| hours | 22.5 ± 0.6 | 65.1 ± 0.2 | + |
| hours | 32.2 ± 0.3 | 141.0 ± 0.2 | + |
| Caicl1/#icl1 | Increase with time | Lower concentration | Higher concentration |
| hours | 19.6 ± 0.0 | 23.3 ± 0.6 | + |
| hours | 54.0 ± 0.0 | 151.0 ± 0.0 | + |
| hours | 46.0 ± 0.0 | 149 ± 0.6 | + |
Figure 2. A, The chart represents the average metabolic activity OD of C. albicans wildtype biofilms on YPL, YPD, and RPMI 1640 media, measured with the MTT assay at each developmental stage (early, intermediate, and maturation). The results are representative of three independent experiments. Absorbance is plotted against growth intervals (time). B, Heterozygous CaICL1/icl1 mutant biofilm on YPL, YPD, and RPMI 1640 media. Formation at early, intermediate, and maturation stages. The results are representative of three independent experiments. Absorbance is plotted against growth intervals (time). Independent experiments were performed in triplicate; C, Homozygous CaICL1/icl1 mutant biofilm on YPL, YPD, and RPMI 1640 media. Formation is shown at the early, intermediate, and maturation stages. The results are representative of three independent experiments. Absorbance is plotted against growth intervals (time). Independent experiments were performed in triplicate.
Figure 3. A, C. albicans wildtype susceptibility testing with fluconazole 128 µg/mL and amphotericin B 16 µg/mL on lactate-containing medium. ANOVA was carried out for the statistical analysis to calculate the significant difference at each developmental level; B, Heterozygous C(IC1)/icl mutant susceptibility testing with fluconazole 128 µg/mL and amphotericin B 16 µg/mL on lactate-containing medium. ANOVA was carried out for the statistical analysis to calculate the significant difference at each developmental level. Independent experiments were performed in triplicate.
Figure 4. Developmental Stages of Biofilms
Figure 5. SEM images of reference and knockout strain biofilms at maturity, before and after treatment. Biofilms were cultured for 48 hours at 37°C under static conditions in YPL medium and visualized at 500× magnification. A, Treated wildtype ICL1/ICL1; B, untreated wildtype ICL1/ICL1; C, treated ICL1/icl1; D, untreated ICL1/icl1; E, treated icl1/icl1, and F, untreated icl1/icl1. Images have been reconstructed to permit clear viewing.
Figure 6. A. Quantification of β-1,3-glucan present in the ECM of wildtype and mutant biofilms not treated with fluconazole at the early, intermediate, and mature phases, respectively; B. Quantification of β-1,3-glucan present in the ECM of the wildtype and mutant biofilms treated with 128 µg/ml fluconazole at the early, intermediate, and mature phases, respectively.
Figure 7. A, Comparative Relative Transcriptional Expression of FKS1; Results were normalized to the internal control CaACT1; B, Comparative relative transcriptional expression of ERG11. Results were normalized to the housekeeping gene CaACT1; C, Comparative relative transcriptional expression of CDR2. Results were normalized to the internal control CaACT1.
5. Discussion

Metabolic flexibility aids *C. albicans* in establishing infections (19, 24, 25). We examined the possible roles of ICL1, a major glyoxylate-synthesizing enzyme, in the distinctive traits of biofilms. Biofilm formation by ICL1/*icl1* and *icl1/icl1* showed a partial impact of ICL, as metabolic activity remained high. The continual growth of ICL1 mutants on non-fatty-acid compounds showed that glyoxylate is not the sole pathway capable of synthesizing C4 molecules for gluconeogenesis, suggesting the presence of similar derivatives and other catabolic pathways. The glyoxylate cycle is preferred because it consumes less energy; however, it can be bypassed in the presence of glucose, farnesol, lactate, malic acid, and succinic acid. Moreover, glucose and fatty acids are more energetically favorable and thus their presence represses lactate assimilation (19, 26).

Lactate, malic acid, and succinic acid derived from pyruvate can be oxidized to oxaloacetate through the pyruvate dehydrogenase pathway. This supports the observation that ICL1/*icl1* and *icl1/icl1* are able to form extensive biofilms in lactate-containing media. The low assimilation observed in ICL1/*icl1* (Figure 2B), later compensated by a rapid growth at the intermediate phase until the matured stage, could be due to incomplete dependence on the pyruvate decarboxylase pathway because of ICL1 allele partial activity. The continual growth of ICL1 mutants (28). Previous works have shown that ERG11 transcripts are higher in young biofilms, which was supported by the findings of this study (31, 32). Upregulation of ERG11 at the early and intermediate stages suggests that its role is streamlined to certain stages, and its expression indicates a response to fluconazole treatment through sterol accumulation and mutation. High ERG11 expression in ICL1/*icl1* suggests that ICL is needed for ERG11 functionality. However, resistance was maintained, suggestive of influences by other mechanisms. The role of efflux-pump genes has been addressed (31, 33, 34). These genes are responsible for the expulsion of drugs from cells. The results of the present study revealed the function of CDR2 to be early-phase-specific. Nevertheless, resistance was exhibited at all stages, suggestive of other genetic influences that are yet to be identified. Thus, ICL may be a part of the synergistic interactions and mechanisms that mediate biofilm resistance phenotypes.

Biofilm formation by *C. albicans* ICL heterozygous and homozygous mutants was found to not be regulated by ICL, as metabolic activity remained high, suggesting that ICL is a negative regulator for other synthesizing enzymes.
and pathways, thereby advancing the cell’s growth and homeostasis. The resistance phenotype was expressed at each formation stage, increasing with maturation. True hyphae morphogenesis impairment indicates filamentous growth as a prominent feature of biofilms, regulated by the preferred non-fermentable carbohydrate-absorption pathway.

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Footnote

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