Chapter

The Application of Solid State Fermentation for Obtaining Substances Useful in Healthcare

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Abstract

In the current review we summarised the research involving solid state fermentation (SSF) for the production of compounds that could be used in healthcare (terpenoids, polyphenols, fibrinolytic enzymes, mycophenolic acid and others). We described several groups of obtained agents which hold various activity: antimicrobial, anti-inflammatory, immunosuppressive, anticoagulant and others (e.g. anticancer or anti-diabetic). It seems that especially terpenoids and polyphenols could be useful in that field, however, other substances such as enzymes and fatty acids play important role as well. We described main groups of microorganisms that are applied in SSF of those compounds, particularly Bacillus genus and fungi, and where possible provided information regarding genes involved in those processes. We also compared various approaches toward optimisation of SSF.

Keywords: solid state fermentation, healthcare, agricultural waste, Bacillus, fungi

1. Introduction

Solid state fermentation (SSF) is a process during which microorganisms (in the presence of small amounts of water) transform agro-industrial waste into valuable compounds [1]. Based on the literature research, wheat bran was commonly used for those processes (Table 1). It is composed of about 53% of dietary fibre (xylans, lignin, cellulose, and galactan, fructans) and contains variety of phenolic acids e.g. ferulic acid, vanillic acid, coumaric acid, caffeic acid, and chlorogenic acid [71]. Researchers also applied other materials rich in polysaccharides (e.g. rice, whole grain wheat, millet, barley) or simple sugars (e.g. fruit pomace) (Table 1). Other substrates which were utilised for SSF are not only sources of carbohydrates, but also protein e.g. soybeans, lentil flour, silkworm larvae, fish meal, cuttle fish waste and king oyster mushroom (Table 1). The selection of waste products used in SSF should ensure the proper balance of nutrients to allow microbial growth and production of terpenoids, polyphenols, enzymes, biosurfactants, short chain fatty acids or others. Therefore, industrial waste with a high content of carbohydrates, protein, pectin or lipids is a suitable substrate.

There were various review papers regarding SSF but in the current chapter we focused only on selected substances which could be used in healthcare and demonstrate antimicrobial, anti-inflammatory properties or/and are immunosuppressants, anticoagulants and anticancer agents, e.g. enzymes, surfactants, terpenoids,
| Name of the substance | Microorganism | Agricultural waste | Reference |
|-----------------------|---------------|-------------------|-----------|
| Antimicrobial properties | Streptomyces cavourensii TN638 | Immobilised bacterial spores (XAD-16) on potato dextrose agar | [2] |
| Surfactin homologues | Bacillus natto NT-6 | Potato dextrose medium | [3] |
| Biosurfactant | Bacillus subtilis SPBI | Millet | [4] |
| Surfactin | Bacillus pumilus UFPEDA 448 | Okara and sugarcane bagasse | [5] |
| Biosurfactant | Tremetes versicolor TV-6 | Two-phase olive mill waste, wheat bran and olive stone | [6] |
| Biosurfactant | Aspergillus niger | Wheat bran and corn cob | [7] |
| Not specified | Pediococcus acidilactici KTU-05-7 | Milk thistle seeds | [8] |
| Not specified | Bacillus licheniformis | Wheat bran, soybean meal, yeast, fish meal | [9] |
| Sambacide | Fusarium sambucinum B10.2 | Potato | [10] |
| γ-Decalactone | Yarrowia lipolytica W29 (ATCC 20460) | Luffa sponge, cellulose sponge, corn cob, castor seed | [11] |
| Phenolic acids | Pleurotus sapidus | Sunflower seed hulls, golden rice straw and husks | [12] |
| Curcumin | Trichoderma strains | Turmeric | [13] |
| Coumarins and oxylipins | Aspergillus oryzae KCCM 12698 | Malt extract agar | [14] |
| Phenolic compounds | Lentinus edodes | Cranberry pomace | [15] |
| Phenolic compounds | Trichoderma strains | Commercial turmeric | [16] |
| Phenolic compounds | Trichoderma strains | Ginger powder | [16] |
| Phenolic compounds | Trichoderma reesi | Garden cress seeds | [17] |
| Phenolic compounds | Aspergillus oryzae NCH 42 | Chinese cucumber, Chinese sage, houpu magnolia, liquorice root | [18] |
| Phenolic compounds | Bacillus clausii | Spent coffee grounds (Arabica) | [19] |
| Anti-inflammatory agents | Trametes versicolor TV-6 | Grape pomace | [20] |
| Not specified | Taiwanefungus campophoratus obtained by SSF | | [21] |
| Rutin | Rhizopus oligosporus NRRL 2710 | Buckwheat groats | [22] |
| Betulinic acid | Inonotus obliquus | The spent substrate of king oyster mushroom, grain including corn, rice grain, white birch and mulberry powder | [23] |
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| Name of the substance                        | Microorganism                        | Agricultural waste                        | Reference |
|----------------------------------------------|---------------------------------------|-------------------------------------------|-----------|
| Phenolic compounds                          | Xylaria nigripes                      | Wheat bran                                | [24]      |
| limonene-1,2-diol, α-terpineol, (−)-carvone, | Diasporthe sp. (Phomopsis sp.)         | Orange peel and bagasse                   | [25]      |
| α-tocopherol, dihydrocarnarveol and valencene|                                        |                                           |           |
| Phenolic compounds, lignans                 | Pediococcus acidilactici LUHS29       | Grounded barley by-products               | [26]      |
| α-Pinene                                     | Saccharomyces cerevisiae AXAZ-1        | Mixed solid and liquid food industry      | [27]      |
| δ-Octalactone                                |                                       | Dried and grounded sugarcane bagasse     | [28]      |
| γ-Unedecalactone                             |                                       |                                           |           |
| γ-Dodecalactone                              |                                       |                                           |           |
| cis-Linaloloxide, Phenanthrene               | Trichoderma viride Pers, ex Fr.; Aspergillus niger van Tieghem | Pu-erh tea                                | [29]      |
| Various terpenes                             | Antrodia camphorata                   | Millet                                    | [30]      |
| Neochlorogenic acid), chlorogenic acid, rutin, | Aspergillus niger ATCC-6275 Rhizopus oligosporus ATCC-22959 | Stones and pomace from fully ripened apricot | [31]      |
| 6’acetyl-glucoside                           |                                       |                                           |           |
| Quercetin and phenolic acids: gallic, vanillic, | Lactobacillus plantarum CECT 748 ATCC 14917 | Cowpeas                                   | [32]      |
| p-hydroxybenzoic, ferulic                    |                                       |                                           |           |
| Phenolic compounds                          | B. subtilis BCRC 14715                | Black soybeans                            | [33]      |
| Daidzin, daidzein, genistin and genistein    | Eurotium cristatum YL-1               | Soybeans seeds                            | [34]      |
| Gallic acid                                  | Rhizopus oryzae (RO IIT RB-13, NRRL 21498) Aspergillus foetidus (GMRB013 MTCC 3557) | Powdered fruits of Myrobalan and Teri pod | [35]      |
| 4-hydroxybenzoic acid, 4-hydroxy-3-methoxybenzoic compound | Rhizopus oryzae RCK2012 | Whole grain wheat                         | [36]      |
| 3,4-di-hydroxybenzoic acid, ferulic acid, vanillic acid, quercetin | Aspergillus oryzae LBA01, A. niger LBA02 | Lentil flour                              | [37]      |

**Immunosupresants**

| Mycophenolic acid                           | Penicillium brevicompactum DSM 2215 | Rice bran-potato peel mixture            | [38]      |
| Mycophenolic acid                           | Penicillium brevicompactum ATCC 16024 (AFI 668) | Pearl barley                            | [39]      |
| Mycophenolic acid                           | Penicillium brevicompactum ATCC 16024 | Wheat bran                               | [40]      |
| Mycophenolic acid                           | Penicillium brevicompactum MTCC 8010 | Rice bran                                | [41]      |
| Mycophenolic acid                           | Penicillium brevicompactum (various strains) | Various agricultural waste               | [42]      |
| Mycophenolic acid                           | Penicillium roqueforti (AG101 and LG109) | Sugarcane bagasse                       | [43]      |
| Name of the substance | Microorganism | Agricultural waste | Reference |
|-----------------------|---------------|--------------------|-----------|
| Cyclosporin A         | Tolypocladium inflatum MTCC 557 | Hydrolysed wheat bran flour and coconut oil cake | [44] |
| Cyclosporin A         | Tolypocladium inflatum MTCC 557 | Wheat bran flour and coconut oil cake | [45] |
| Cyclosporin A         | Tolypocladium inflatum ATCC 34921 | Wheat bran | [46] |
| Cyclosporin A         | Tolypocladium inflatum DRCC 106 (mutated strain) | Wheat bran | [47] |
| Cyclosporin A         | Tolypocladium sp. | Wheat bran | [48] |
| Tacrolimus            | Streptomyces hygroscopicus | Various agricultural waste | [49] |

**Anticoagulant agents**

| Name of the substance         | Microorganism                  | Agricultural waste                  | Reference |
|-------------------------------|--------------------------------|-------------------------------------|-----------|
| Halotolerant chitinase        | Citrobacter freundii str. nov. haritD11 | Wheat bran with fish scale | [50] |
| Halotolerant Chitinase        | Citrobacter freundii str. nov. haritD11 | Wheat bran with shrimp shellfish | [51] |
| Fibrinolytic enzyme           | Bacillus amyloiquefaciens LSSE-62 | Chickpeas | [52] |
| Fibrinolytic enzyme           | Bacillus sp. IND6               | Wheat bran | [53] |
| Fibrinolytic enzyme           | Bacillus sp. IND12              | Cow dung | [54] |
| Nattokinase                   | Bacillus subtilis natto         | Soybean | [55] |
| Fibrinolytic enzyme           | Bacillus subtilis XZ1125        | Soybean meal | [56] |
| Fibrinolytic enzyme           | Bacillus subtilis WR350         | Corn steep | [57] |
| Fibrinolytic enzyme           | Bacillus halodurans IND18       | Wheat bran | [58] |
| Fibrinolytic enzyme           | Bacillus cereus GD55            | Apple pomace | [59] |
| Fibrinolytic enzyme           | Bacillus cereus IND5            | Cuttle fish waste and cow dung | [60] |
| Fibrinolytic enzyme           | Paenibacillus sp. IND8          | Wheat bran | [61] |
| Fibrinolytic enzyme           | Pseudalteromonas sp. IND11      | Sun-dried cow dung | [62] |
| Fibrinolytic enzyme           | Xanthomonas oryzae IND3         | Cow dung | [63] |
| Fibrinolytic enzyme           | Bacillus firmae NA-1            | Soybean grits | [64] |
| Fibrinolytic enzyme           | Mucor subtilissimus UCP 1262    | Wheat bran | [65] |
| Fibrinolytic enzyme           | Fusarium oxysporum              | Rice chaff | [66] |
| Fibrinolytic enzyme           | Fusarium oxysporum              | Rice chaff | [67] |

**Anticancer agents**

| Name of the substance       | Microorganism | Agricultural waste | Reference |
|----------------------------|---------------|--------------------|-----------|
| Short chain fatty acids     | Aspergillus kawachii KCCM 32819 | Silkworm larvae powder | [68] |
| Putative phytoestrogen      | Aspergillus fumigatus F-993 or A. awamori FB-133 | Defatted soybean | [69] |
| Andrastin A and C           | Penicillium expansum KACC 40815 | Malt extract agar | [70] |

**Table 1.**
Examples of substances produced by solid state fermentation that could be used in healthcare.
polyphenols and short chain fatty acids. We also described main groups of microorganisms that were involved in cited studies and compared various approaches for optimising SSF.

2. Main properties of substances obtained from SSF

2.1 Antimicrobial properties

In the majority of cited studies (Table 1) authors did not verify which particular compound contributed to antimicrobial properties. In most of cases they concluded that polyphenols contributed to that phenomenon [13, 15–19] because in comparison to control groups, extracts obtained after SSF demonstrated stronger antimicrobial effects containing more phenolic compounds (PC) at the same time. In the paper written by Mohamed et al. [13, 72] authors did not carry out detailed qualitative and quantitative analysis of fungal metabolites – they assumed that only curcumin would be the substance demonstrating antibacterial properties.

Some studies involved detailed analysis of polyphenol profiles and authors assigned antibacterial and antifungal properties to phenolic acids which concentration was increased by Pleurotus sapidus [12]. Others indicated that antimicrobial activity was achieved due to the occurrence of coumarins and oxylipins detected in post-fermentation extracts when Aspergillus oryzae KCCM 12698 was used for SSF [14]. Kaaniche et al. [2] additionally analysed structures of obtained bioactive compounds and they proved that four most potent antimicrobials produced by Streptomyces cavourensis TN638 were macrotetrolides. Similar approach was applied to identify antimicrobial compounds produced by Fusarium sambucinum B10.2 and proved it was sambacide [10]. When surfactants produced by various Bacillus strains were tested for antimicrobial properties, researchers additionally tested their properties like emulsification activities [4] or tensioactive activity [5]. Except for latest reports regarding surfactin, we did not include antibiotics in our chapter because currently there are various resistant strains so some alternatives are required.

The majority of identified antimicrobial compounds demonstrated activity equal to [2, 13, 17] or greater [10, 13, 15, 16] than well-known antibiotics. In some cases authors did not provide results for control samples so it was not possible to assess how those substances were effective, however, inhibition zones in diffusion disk method were very prominent [4, 9, 14, 15]. In other studies MIC (Minimum Inhibitory Concentration) of extracted substances were not higher than for antibiotics, however, since those substances were obtained from agricultural waste which is a cost effective substrate, they still could be considered as potential antimicrobials [2, 12, 18, 19]. Only metabolites produced by Pediococcus acidilactici KT-05-7 demonstrated very weak antimicrobial properties [8].

2.2 Anti-inflammatory agents

Anti-inflammatory properties of terpenoids were already described in various reviews [73, 74] but they were not investigated in cited papers so we did not discuss obtained results. It is worth mentioning that each extract obtained after SSF contained at least one compound that could demonstrate such activity: lactones which were produced by Trichoderma viride EMCC-107 [28]; limonene-1,2-diol, α-terpineol, (−)-carvone, α-tocopherol produced by Diaporthe sp. KY113119 [25]; 1-terpineol, L-linalool produced by Antrodia camphorata [30]; betulinic acid – Inonotus obliquus and [23]; α-pinene produced by Saccharomyces cerevisiae AXAZ-1 and Kluyveromyces marxianus IMB3 [27].
In the case of polyphenols we took the same approach – we only summarised the research that investigated anti-inflammatory properties of extracts obtained after SSF. When grape pomace was treated with *Trametes versicolor* TV-6 the concentration of phenolic acids, flavan-3-ols and rutin increased while the concentration of anthocyanins decreased. Those changes resulted in enhanced anti-inflammatory activity of obtained extracts which was measured by the inhibition of 5-lipoxygenase and hyaluronidase [20]. Also polyphenols produced by *Xylaria nigripes* demonstrated enhanced anti-inflammatory properties which were verified by the inhibition of cyclooxygenase-2 [24]. Additionally, in both cases, obtained extracts demonstrated neuroprotective properties. Studies carried out by Yin et al. [75] demonstrated that *A. niger* was able to release ferulic acid bound to various polysaccharides in the wheat bran. Obtained substances exhibited stronger anti-inflammatory activity than that of free ferulic acid which probably took place due to the presence of accompanying compounds in obtained extracts. Moreover, those released compounds could significantly inhibit intracellular malondialdehyde formation and the LPS-induced inflammation. It is difficult to assess which extracts demonstrated greater activity because authors of all abovementioned studies used different way to express results – the provided IC₅₀ values or % of inhibition.

In the majority of cited papers anti-inflammatory properties of extracts containing polyphenols were not verified despite the fact that there were some compounds among them which demonstrate such activity [76, 77]. Research mainly focused on antioxidant properties [20, 22, 23, 31]. This applies to various studies involving organisms providing increase of particular phenolic compounds: *Rhizopus oligo-sporus* NRRL 2710 – rutin [22], *Prunus armeniaca* L – cinnamic acids and selected flavonols [31]; *Lactobacillus plantarum* CECT 748 – hydroxybenzoic acids and flavonols [78]; *Eurotium cristatum* YL-1 – isoflavones (daidzin, daidzein, genistin and genistein) [34]; *Rhizopus oryzae* and *Aspergillus foetidus* – tannins [35]; *Rhizopus oryzae* RCK2012 – phenolic acids [36]; *Aspergillus oryzae* LBA01 – 3,4-di- hydroxy-benzoic acid, ferulic acid, vanillic acid and quercetin [37].

**2.3 Immunosuppressants**

Immunosuppressants could be obtained by SSF as well. It seems that one of the most common substance which was detected in studies involving SSF is myco-phenolic acid (MA, Table 1). It works as a blocker in producing precursors for the synthesis of RNA and DNA, so as the result it blocks proliferative response of T and B lymphocytes [79]. In the research that involved SSF, authors did not verify properties of the obtained MA.

Another substance belonging to that group, classified as calcineurin inhibitor [80], is cyclosporin A (CA). This substance is used not only in transplant patients but also in treatment of glomerular disease. CA prevents calcineurin-dependent transcription in activated T cells. Based on studies that were aiming to produce CA we concluded that authors did not verify properties of obtained substances but they focused on various methods for its extraction and purification. They mostly used butyl acetate [44, 48] or ethyl acetate [47]. Tacrolimus could be also produced by SSF by *Streptococcus hygroscopicus*. The whole process of bacterial cultivation and the extraction of that compounds was covered by the patent [49].

**2.4 Anticoagulants**

Another group of substances that could be produced by SSF is anticoagulants. Blood coagulation is a physiological process, which consists of a series of coagulation factors and proteolytic activation steps, which lead to the production of
thrombin – the main coagulation enzyme. The majority of research was focused on fibrinolytic enzymes (FE, Table 1) – subtilisin, however, in one case it was nattokinase [55]. In the majority of cited papers authors verified anticoagulant properties of obtained enzymes and they used different methods – some of them applied spectrophotometric method which measured the increase of turbidity at 275 nm caused by added enzyme [53, 54, 58, 60, 62, 65] and others measured zones of clearance in solid media containing fibrinogen [64, 66, 67].

Another compound which could be considered as a putative anticoagulant is halotolerant chitinase. Its properties were confirmed [50, 51] by testing how this enzyme could dissolve fibrin in time. Moreover, Meruvu et al. [51] showed that it held antifungal activity.

2.5 Anticancer agents

Terpenoids are well-known for their cytotoxic activity and there were various studies investigating such activity against cancer cells. Anticancer mechanism of terpene or essential oils that contain them was described in various review papers [81–83]. Several compounds belonging to that class could be obtained by SSF (Table 1): limonene from orange waste [25]; linalool, geraniol and β-caryophyllene from millet [28, 30]; andrastin A and C on malt extract agar [70]. Properties of extracts that contained those and other terpenoids were not investigated in cited papers so we decided not to discuss that aspect in the current chapter.

Another group of bioactive compounds which was shown to hold anticancer activity is polyphenols [84, 85]. The following substances occurred in cited reviews and studies that we summarised in Table 1: cinnamic acids [22], daidzein and genistein [86], quercetin [35, 87], and tannins [34]. Since authors of cited research papers (Table 1) did not test obtained extracts against those properties, we did not discuss that phenomenon.

In the study of Cho et al. (2019) authors demonstrated that fatty acids detected in extracts obtained from silkworm larvae powder fermented by Aspergillus kawachii demonstrated such activity against human hepatocellular carcinoma [68]. It was shown that fermentation increased concentration of those compounds, especially oleic and linoleic acids. This phenomenon took place due to the enhancement of cell apoptosis and suppression of protein responsible for preventing the apoptosis. The value of that research is particularly significant because so far, polyunsaturated fatty acids (PUFAs) attracted most attention in the context of colorectal cancer [87].

3. Groups of microorganisms demonstrating greatest potential for the production of health-promoting properties in SSF

3.1 Bacillus genus

It was demonstrated that the representatives of the Bacillus genus were able to produce fibrinolytic enzymes by SSF [52–58, 60]. That ability is mostly assigned to the expression of fibE gene which encodes enzyme called subtilisin [88] which solubilises blood clots. Gene expression was not investigated in cited papers so the ability to produce those enzymes by tested strains could be the result of other genes expression.

Bacillus genus is successfully used in solid-state fermentation to improve antimicrobial activity of fermented food. Rochín-Medina et al. [19] who tried to determine optimal bioprocessing conditions for SSF of spent coffee grounds by Bacillus clausii achieved increase of flavonoid and total phenolic contents by 13 and 36%,
respectively. SSF also enhanced antibacterial activity of obtained extracts. That phenomenon could be explained by the fact that *Bacillus* sp. strains could metabolise fibre which releases phenolic compounds as a result of lignocellulolytic activity, and demonstrate strong correlation between the increase of phenolic compounds and the synthesis of cellulases and pectinases [19]. Those enzymes can break down plant cell wall components which leads to the hydrolysis of ester bonds that bind phenolic compounds [19].

The representatives of the *Bacillus* genus are able to produce surfactants as well (Table 1). There are various enzymes involved in the production of surfactin which form multienzyme peptide synthetase complex. One of those proteins is Srf (which consists of three units A, B and C). There are also two other genes involved in that process – *sfp* and *comX*. However, the role and interactions of protein and genes was described in more details in other review papers [89], therefore, we did not discuss that phenomenon in details.

### 3.2 Other bacteria

There are also other bacteria that were tested against the production of substances holding the potential for application in healthcare. For instance, there are several species that are able to produce fibrinolytic enzymes which could serve as potential anticoagulants: *Paenibacillus* sp. IND8 [61], *Pseudoalteromonas* sp. IND11 [62], *Xanthomonas oryzae* IND3 [63] or *Citrobacter freundii* nov. *haritD11* [50, 51]. In the case of *Paenibacillus* its ability for producing such enzymes could be assigned to the expression of *PPFE-I* gene [90]. It seems that the synthesis of protein encoded by that gene could be stimulated by Zn$^{2+}$, Mg$^{2+}$ and Fe$^+$ so the concentration of these ions could be considered in future studies focusing on optimisation of enzyme production. In the case of *Citrobacter freundii*, molecular mechanism seems much simpler because so far, only *chiX* gene was assigned to its ability for chitinase production [91]. It is still unclear which genes are involved in the production of anticoagulant agent by *Pseudoalteromonas* or *Xanthomonas oryzae* so it is an aspect that could be investigated in the future, since the quantity of the enzyme was very prominent – up to 1,388 U/ml.

It was also demonstrated that *Streptococcus hygroscopicus* is able to produce immunosuppressant in SSF based on agricultural waste with added supplements [49] and it seems that *aroA, fkbN*, and *luxR* genes are mostly responsible for that phenomenon [92]. On the other hand, molecular mechanisms standing behind the ability of *Yarrowia lipolytica* W29 (ATCC 20460) to produce γ-decalactone [11] which analogues demonstrate antiviral and antifungal properties [93] is simpler because it involves only *POX2* overexpression [94]. Similarly, in the case of *Lactobacillus planta-rum* CECT 748 [95] which increased the concentration of particular phenolic compounds, probably only *est_1092* gene was involved – it encodes phenolic esterase [96].

None of the cited paper evaluated molecular mechanisms involved in processes carried out by tested strains.

### 3.3 Fungi

The majority of research involving SSF is carried out with various fungi (Table 1). Many of analysed examples focused on the increase of phenolic compounds. One of the genera which were involved in that process was *Trichoderma* [13, 16, 17]. Those fungi are known to produce cellulolytic, ligninolytic and xylanolytic enzymes [97] and it has been demonstrated in other studies that cellulase could significantly increase concentrations of various polyphenols e.g. caffeic acid, vanillin, p-coumaric acid, and ferulic acid [98]. In fact, similar strategies
might apply to Aspergillus spp. [22, 37, 99] because representatives of that genus could produce enzymes demonstrating such activities [98] as well. Rhizopus oryzae [36, 100] synthesizes cellulase, xylanase and pectinase [101]. Other fungi applied in other cited studies could produce the following enzymes that participate in polyphenol increase: Lentinus edodes [15] – xylanase and cellulase [102], Pleurotus sapidus – ligninolytic enzymes [12], Trametes versicolor TV-6 – β-glucosidase [20].

Genetic background of the synthesis of abovementioned enzymes by Trichoderma spp. and Aspergillus spp. was revised by Amore et al. [103] while in the case of R. oryzae and P. chrysosporium it was described by Battaglia et al. [104] so we did not provide details of those processes, especially that many genes are involved.

There are also some fungi which could be natural sources of polyphenols: Taiwanofungus camphoratus [21], Inonotus obliquus [23], and Xylaria nigripes [24]. In all cases SSF increased their anti-inflammatory properties by increasing concentration of particular phenolic compounds which could also originate from substrates that were used for their cultivation and those were: spent substrate of Pleurotus eryngii, sunflower seed hulls, corn and rice grain, white birch and mulberry in the case of I. obliquus; wheat bran in the case of X. nigripes.

Another significant group of bioactive compounds which concentration was increased by SSF was terpenes (Table 1) and those processes were carried out by various fungi: Fusarium sambucinum B10 [10], Penicillium expansum KACC 40815 [70], Diaporthe sp. KY113119 [25], Antrodia camphorata [30], Saccharomyces cerevisiae AXAZ1-1 and Kluyveromyces marxianus IMB3 [27], Trichoderma viride EMCC-107 [28], and Aspergillus niger van Tieghem [29]. A. camphorate is the natural source of terpenes and their concentration was increased by selecting millet as the main substrate for SSF. K. marxianus and S. cerevisiae could probably increase the concentration of tested compounds by releasing terpenes from their glycosidic forms by β-glucosidases. In the case of Fusarium spp., Aspergillus spp. and Penicillium spp. it was demonstrated that those genera could produce several sesquiterpene synthases [105]. As in the case of polyphenols, molecular background of terpene transformations is very complex so we decided not to describe it in the current chapter, but refer to the review of Quin et al. [105] instead. It must be highlighted that except for the study regarding P. expansum KACC 40815 demonstrating that terpenoid cyclase was mostly involved in described processes [70], none of the authors investigated enzymatic activity during terpene transformation nor determined gene expression.

Based on cited papers it might be stated that Penicillium brevicopactum is most common in the research regarding production of mycophenolic acid in SSF [38–43]. Those fungi produce polyketide synthase encoded by mpaC gene along with other enzymes such as: protein transacylase, β-ketoacylsynthase, acyltransferase, acyl carrier protein, and methyltransferase (MT) domains [106]. There is another taxon which is able to produce immunosuppressants, namely cyclosporin A, and it is Tolypocladium inflatum [44–48] which has got nonribosomal peptide synthetase that encodes for cyclosporin synthetase (simA gene) [107].

Mucor subtilissimus UCP 1262 [65] and Fusarium oxysporum [66, 67] were shown to produce fibrinolytic enzymes. It has been already demonstrated that FP gene is responsible for encoding fibrinolytic protease in Fusarium sp. [108] but in the case of Mucor sp. it remains unknown. Molecular background of A. kawachii KCCM 32819 production of short chain fatty acids could be the same as for A. nidulans and other filamentous ascomycetes – farB gene is mostly responsible for that ability, however, farA participates as well [109]. Genetic background of biosurfactant production in moulds is still unknown.
3.4 Modified strains

It has been demonstrated that in some cases the yield of microbial metabolites significantly increased when the strain was subjected to gamma ray – it increased the quantity of obtained mycophenolic acid produced by two strains of *Penicillium requeforti* [43] in comparison of other cited studies which involved unmodified *Penicillium brevicompactum* [39, 41]. On the other hand, UV radiation was used for the modification of *Tolypocladium inflatum* which was used for the production of Cyclosporin A [47], however, in the case of that substance there were other studies which resulted in much higher yields [44, 45].

4. Optimisation of SSF conditions

One of the approaches that were applied for the optimisation of the concentration of bioactive substances by SSF is to provide pre-treatment to the main substrate. Heat-treatment was one of the main methods e.g. autoclaving, moistening in boiling water, cooking with deionised water, drying, freezing, freeze-drying, vacuum-drying and roasting. It inactivates native microorganisms and enzymes.

Once the main substrate is prepared, there are other experimental conditions that need to be optimised. In cited papers the major factor contributing to obtained results was moisture which was at least 50% [69, 110]. The next crucial factor was medium composition. In few cases authors used solid media commonly used for the cultivation of microorganisms [2, 70]. However, in the majority of cases authors added some nutrients to agricultural waste to provide better results of releasing bioactive compounds. In the case of mycophenolic acid produced *P. brevibactum* MTCC 8010 from rice bran those supplements were: peptone, KH₂PO₄, glycine and methionine [41]. On the other hand, in the case of *P. brevicompactum* ATCC 16024, the addition of mannitol or (NH₄)₂HPO₄ to pearl barley did not enhance the MA synthesis [39]. Surprisingly, the quantity of MA was higher in the latter case – 5.47 g/kg of the substrate in comparison to 4.5 g/kg under optimised conditions. It seems that pearl barley has got chemical composition which is more preferable for obtaining MA.

On the other hand, Plackett-Burman design was applied for the optimisation of cyclosporin A production by *Tolypocladium inflatum* MTCC 557 which resulted in 8,166 mg/kg [45] which is 1.26-fold higher than in studies carried out by Survase et al. [44, 45] who applied the same fungal strain or even 45.62-fold higher than reported by Nisha and Ramasamy [46] who used *T. inflatum* ATTC 34921 strain. In the last study authors firstly used Plackett-Burman design for the selection of nutrients and later on, they used half-factorial central composite rotatable design (CCRD) of response surface methodology (RSM) to select optimum concentrations of those substances which resulted in more than 10-fold increase of tested compound [47].

When the production of fibrinolytic enzyme was optimised with RSM, its activity increased 3 times in the case of *Pseudoalteromonas* sp. IND11 [62]; 4 times in the case of *Xanthomonas oryzae* sp. IND3 [63]; while when central composite design (CCD) was used, 4.5-fold increase was noted when *Paenibacillus* sp. IND8 was used [61]. In the case of *Xanthomonas oryzae* sp. IND3, CCD was additionally used for estimating optimal values of the following variables: sucrose, yeast extract, and pH of the medium [63]. Among all mentioned microorganisms *Paenibacillus* sp. IND8 produced greatest quantities of the enzyme – 4,418 U/ml.

Various *Bacillus* strains were used for the production of fibrinolytic enzyme (Table 1) and those proved to be more efficient enzyme producer
that abovementioned bacteria. When authors used two-level factorial design in the case of \textit{Bacillus} sp. IND12 examining the impact of moisture, sucrose, and MgSO$_4$ levels added to the cow dung, maximum enzyme activity reached 4,143 U/g [54]. Further, orthogonal design (corn steep powder, sucrose and MgSO$_4$ $\cdot$ 7H$_2$O) provided even greater enzyme activity (5,865 U/ml) in 100 l fermenter when applying \textit{Bacillus subtilis} WR350 [57]. However, the greatest activity was achieved when \textit{Bacillus halodurans} IND18 was used. It produced 6,851 U/g when two-level full factorial design was applied and the optimum conditions were as follows: 1% peptone, 80% moisture and pH 8.32, using wheat bran as the main substrate [58].

Ghribi et al. [4] took a different approach for the optimisation of surfactin production – firstly, authors applied Plackett-Burman design to assess which of the five variables were the most important and then they optimised the process with CCD involving three selected variables. They found parameters (temperature – 37°C, inoculum age – 14 h, and moisture – 88%) that were the most favourable for the production of surfactin and increased its yield by 2-fold (up to 2 g/l). Sun et al. [3, 111] carried out step-by-step optimisation and found out that the addition of attapulgite by 1.96-fold (4.3782 g/kg). This would suggest that \textit{B. natto} NT-6 was the most suitable for that application among all tested strains.

Optimization of halotolerant chitinase was carried out using RSM – Box Behnken method which involved \textit{Citrobacter freundii} and that process slightly improved enzyme production in comparison to initial optimisation experiments from 112.43 U/g dry substance to 124.73 U/g dry substance [51]. That optimisation was mainly focused on the ratio of main substrates (wheat bran and shrimp shellfish), temperature and moisture content. Similarly, minor changes were observed when wheat bran and powdered fish scales were used for the statistical optimisation of chitinase production [50].

5. Conclusions

Solid-state fermentation could provide various substances useful in healthcare: antimicrobials, immunosuppressants, anticoagulants, substances holding anti-inflammatory properties and anticancer agents. It seems that polyphenols and terpenes are especially versatile in their applications. The majority of studies involved various fungi mainly due to their enzymatic activity which supports the release of bioactive compounds from agricultural waste. Molecular mechanisms of those processes are usually very complex; however, they remain unknown for some fungi. Further studies are necessary to assess which genes could be expressed during those processes because those could be used for modification of microorganisms to increase their yield. It is also important to bear in mind that SSF requires the presence of various supplements and fermentation could be optimised by statistical tools, especially Response Surface Methodology and Central Composite Design. In some cases step-by-step optimisation could be sufficient.

Conflict of interest

The authors declare no conflict of interest.
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References

[1] Verduzco-oliva R, Gutierrez-uribe JA. Beyond Enzyme Production: Solid State Fermentation (SSF) as an Alternative Approach to Produce Antioxidant Polysaccharides. Sustainability. 2020;12: 1-14. doi:10.3390/su2020495

[2] Kaaniche F, Hamed A, Elleuch L, Chakchouk-mtibaa A, Mellouli L. Purification and characterization of seven bioactive compounds from the newly isolated Streptomyces cavourensis TN638 strain via solid-state fermentation. Microb Pathog. 2020; 104106. doi:10.1016/j.micpath.2020.104106

[3] Sun D, Liao J, Sun L, Wang Y, Liu Y, Deng Q, et al. Effect of media and fermentation conditions on surfactin and iturin homologues produced by Bacillus natto NT - 6: LC – MS analysis. AMB Express. 2019;9: 1-9. doi:10.1186/s13568-019-0845-y

[4] Ghribi D, Abdelkefi-mesrati L, Mnif I, Kammoun R, Ayadi I, Saadaoui I, et al. Investigation of Antimicrobial Activity and Statistical Optimization of Bacillus subtilis SPB1 Biosurfactant Production in Solid-State Fermentation. J Biomed Biotechnol. 2012;2012: 1-12. doi:10.1155/2012/373682

[5] Slivinski CT, Mallmann E, De Araújo JM, Mitchell DA, Krieger N. Production of surfactin by Bacillus pumilus UFPEDA 448 in solid-state fermentation using a medium based on okara with sugarcane bagasse as a bulking agent. Process Biochem. 2012;47: 1848-1855. doi:10.1016/j.procbio.2012.06.014

[6] Lourenço LA, Alberton Magina MD, Tavares LBB, Guelli Ulson de Souza SMA, Garcia Román M, Altmajer Vaz D. Biosurfactant production by Trametes versicolor grown on two-phase olive mill waste in solid-state fermentation. Environ Technol (United Kingdom). 2018;39: 3066-3076. doi:10.1080/0959330.2017.1374471

[7] Kreling NE, Simon V, Fagundes VD, Thomé A, Colla LM. Simultaneous Production of Lipases and Biosurfactants in Solid-State Fermentation and Use in Bioremediation. J Environ Eng. 2020;146: 04020105. doi:10.1061/(ASCE)EE.1943-7870.0001785

[8] Juodeikiene G, Cizeikiene D, Ceskeviciute V, Vidmantiene D, Basinskiene L, Akuneca I, et al. Solid-State Fermentation of Silybum marianum L. Seeds Used as Additive to Increase the Nutritional Value of Wheat Bread. Food Technol Biotechnol. 2013;51: 528-538.

[9] Lin E-R, Cheng Y-H, Hsiao FS-H, Proskura, Witold S, Dybus A, Yu Y-H. Optimization of solid-state fermentation conditions of Bacillus licheniformis and its effects on Clostridium perfringens -induced necrotic enteritis in broilers. Rev Basiliera Zootec. 2019;48: 1-14. doi:10.1590/rbz4820170298

[10] Dong J, Cai L, Li X, Duan R, Shu Y, Chen F, et al. Production of a new tetracyclic triterpene sulfate metabolite sambacide by solid-state cultivated Fusarium sambucinum B10.2 using potato as substrate. Bioresour Technol. 2016. doi:10.1016/j.biortech.2016.07.014

[11] Try S, De-Coninck J, Voilley A, Chunhieng T, Waché Y. Solid state fermentation for the production of γ-decalactones by Yarrowia lipolytica. Process Biochem. 2018;64: 9-15. doi:10.1016/j.procbio.2017.10.004

[12] Pinela J, Omarini AB, Stojković D, Barros L, Postemsky PD, Calhelha RC,
et al. Biotransformation of rice and sunflower side-streams by dikaryotic and monokaryotic strains of Pleurotus sapidus: Impact on phenolic profiles and bioactive properties. Food Res Int. 2020;132. doi:10.1016/j.foodres.2020.109094

[13] Mohamed SA, Saleh RM, Kabli SA, Al-garni SM. Influence of solid state fermentation by Trichoderma spp. on solubility, phenolic content, antioxidant, and antimicrobial activities of commercial turmeric. Biosci Biotechnol Biochem. 2016;8451: 1-9. doi: 10.1080/09168451.2015.1136879

[14] Son SY, Lee S, Singh D, Lee N, Lee D, Lee CH. Comprehensive Secondary Metabolite Profiling Toward Delineating the Solid and Submerged-State Fermentation of Aspergillus oryzae KCCM 12698. Front Microbiol. 2018;9: 1-12. doi:10.3389/fmicb.2018.01076

[15] Vattem DA, Lin YT, Labbe RG, Shetty K. Phenolic antioxidant mobilization in cranberry pomace by solid-state bioprocessing using food grade fungus Lentinus edodes and effect on antimicrobial activity against select food borne pathogens. Innov Food Sci Emerg Technol. 2004;5: 81-91. doi:10.1016/j.ifset.2003.09.002

[16] Saleh RM, Kabli SA, Al-Garni SM, Al-Ghamdi MA, Abdel-Aty AM, Mohamed SA. Solid-state fermentation by Trichoderma viride for enhancing phenolic content, antioxidant and antimicrobial activities in ginger. Food Res Int. 2020;132. doi:10.1016/j.foodres.2020.109094

[17] Abdel-Aty AM, Bassuiny RI, Barakat AZ, Mohamed SA. Upgrading the phenolic content, antioxidant and antimicrobial activities of garden cress seeds using solid-state fermentation by Trichoderma reesei. J Appl Microbiol. 2019;127: 1454-1467. doi:10.1111/jam.14394

[18] Wen Y, Yan L, Chen C. Effects of fermentation treatment on antioxidant and antimicrobial activities of four common Chinese herbal medicinal residues by Aspergillus oryzae. J Food Drug Anal. 2013;21: 219-226. doi:10.1016/j.jfda.2013.05.013

[19] Rochín-Medina JJ, Ramírez K, Rangel-Peraza JG, Bustos-Terrones YA. Increase of content and bioactivity of total phenolic compounds from spent coffee grounds through solid state fermentation by Bacillus clausii. J Food Sci Technol. 2018;55: 915-923. doi:10.1007/s13197-017-2998-5

[20] Bucić-Kojić A, Fernandes F, Silva T, Planinić M, Tišma M, Šelo G, et al. Enhancement of the anti-inflammatory properties of grape pomace treated by: Trametes versicolor. Food Funct. 2020;11: 680-688. doi:10.1039/c9fo02296a

[21] Liu DZ, Liang HJ, Chen CH, Su CH, Lee TH, Huang CT, et al. Comparative anti-inflammatory characterization of wild fruiting body, liquid-state fermentation, and solid-state culture of Taiwanofungus camphoratus in microglia and the mechanism of its action. J Ethnopharmacol. 2007;113: 45-53. doi:10.1016/j.jep.2007.03.037

[22] Wronkowska M, Honke J, Piskula Mariusz K. Effect of solid-state fermentation with Rhizopus oligosporus on bioactive compounds and antioxidant capacity of raw and roasted buckwheat groats. Ital J Sci Food. 2015;27: 424-431. doi:10.14674/1120-1770/ijfs.v373

[23] Chen HJ, Chen YS, Liu SL, Liou BK, Chen CS. The Increase of Bioactive Ingredients by Solid State Fermentation of Inonotus obliquus with Spent Substrate. Waste and Biomass Valorization. 2019. doi:10.1007/s12649-019-00911-7

[24] Divate RD, Wang CC, Chou ST, Chang CT, Wang PM, Chung YC.
Using wheat bran and soybean meal as solid state fermentation substances for the production of Xylaria nigripes with bioactivities. J Taiwan Inst Chem Eng. 2017;70:127-133. doi:10.1016/j.jtice.2016.11.003

[25] Jucoski Bier MC, Medeiros ABP, De Kimpe N, Soccol CR. Evaluation of antioxidant activity of the fermented product from the biotransformation of R-(+)-limonene in solid-state fermentation of orange waste by Diaporthe sp. Biotechnol Res Innov. 2019;3:168-176. doi:10.1016/j.biori.2019.01.002

[26] Bédard F, Hammami R, Zirah S, Rebuffat S, Fliss I, Biron E. Synthesis, antimicrobial activity and conformational analysis of the class IIa bacteriocin pediocin PA-1 and analogs thereof /631/45/611 /631/326/22 /639/638/54/978 /639/638/30/92419 /639/638/40/3/977 /140/131 /82/6 /82/16 article. Sci Rep. 2018;8:1-13. doi:10.1038/s41598-018-27225-3

[27] Aggelopoulos T, Katsieris K, Bekatorou A, Pandey A, Banat IM, Koutinas AA. Solid state fermentation of food waste mixtures for single cell protein, aroma volatiles and fat production. Food Chem. 2014;145:710-716. doi:10.1016/j.foodchem.2013.07.105

[28] Fadel HHM, Mahmoud MG, Asker MMS, Lotfy SN. Characterization and evaluation of coconut aroma produced by Trichoderma viride EMCC-107 in solid state fermentation on sugarcane bagasse. Electron J Biotechnol. 2015;18:5-9. doi:10.1016/j.ejbt.2014.10.006

[29] Chun-hua Z, Zhi-guo S, Wen-xia Y, Ya-li Li, Ting-ting SUN. Effects on Aroma Composition of Pu-erh Tea under Solid State Fermentation with Different Beneficial Microorganisms. J Tea Sci. 2010;30:251-258.

[30] Xia Y, Zhang B, Li W, Xu G. Changes in volatile compound composition of Antrodia camphorata during solid state fermentation. J Sci Food Agric. 2011;91:2463-2470. doi:10.1002/jsfa.4488

[31] Dulf FV, Vodnar DC, Dulf EH, Pintea A. Phenolic compounds, flavonoids, lipids and antioxidant potential of apricot (Prunus armeniaca L.) pomace fermented by two filamentous fungal strains in solid state system. Chem Cent J. 2017;11:1-10. doi:10.1186/s13065-017-0323-z

[32] Leite P, Silva C, Salgado JM, Belo I. Simultaneous production of lignocellulolytic enzymes and extraction of antioxidant compounds by solid-state fermentation of agro-industrial wastes. Ind Crops Prod. 2019;137:315-322. doi:10.1016/j.indcrop.2019.04.044

[33] Juan MY, Chou CC. Enhancement of antioxidant activity, total phenolic and flavonoid content of black soybeans by solid state fermentation with Bacillus subtilis BCRC 14715. Food Microbiol. 2010;27:586-591. doi:10.1016/j.fm.2009.11.002

[34] Chen Y, Wang Y, Chen J, Tang H, Wang C, Li Z, et al. Bioprocessing of soybeans (Glycine max L.) by solid-state fermentation with Eurotium cristatum YL-1 improves total phenolic content, isoflavone glycones, and antioxidant activity. RSC Adv. 2020;10:16928-16941. doi:10.1039/c9ra10344a

[35] Mukherjee G, Banerjee R. Biosynthesis of tannase and gallic acid from tannin rich substrates by Rhizopus oryzae and Aspergillus foetidus. J Basic Microbiol. 2004;44:42-48. doi:10.1002/jobm.200310317

[36] Dey TB, Kuhad RC. Enhanced production and extraction of phenolic compounds from wheat by solid-state fermentation with Rhizopus oryzae RCK2012. Biotechnol Reports. 2014;4:120-127. doi:10.1016/j.btre.2014.09.006
[37] Magro AEA, Silva LC, Rasera GB, de Castro RJS. Solid-state fermentation as an efficient strategy for the biotransformation of lentils: enhancing their antioxidant and antidiabetic potentials. Bioresour Bioprocess. 2019;6. doi:10.1186/s40643-019-0273-5

[38] Alpak I, Askin Uzel R, Sargin S, Yesil-Celiktas O. Supercritical CO2 extraction of an immunosuppressant produced by solid-state fermentation. J CO2 Util. 2018;27: 398-404. doi:10.1016/j.jcou.2018.08.014

[39] Alani F, Grove JA, Anderson WA, Moo-Young M. Mycophenolic acid production in solid-state fermentation using a packed-bed bioreactor. Biochem Eng J. 2009;44: 106-110. doi:10.1016/j.bej.2008.11.005

[40] Sadhukhan AK, Ramana Murthy M V, Ajaya Kumar R, Mohan EVS, Vandana G, Bhar C, et al. Optimization of mycophenolic acid production in solid state fermentation using response surface methodology. J Ind Microbiol Biotechnol. 1999;22: 33-38. doi:10.1038/sj.jim.2900597

[41] Patel G, Patil MD, Soni S, Chisti Y, Banerjee UC. Production of Mycophenolic Acid by Penicillium brevicompactum Using Solid State Fermentation. Appl Biochem Biotechnol. 2017;182: 97-109. doi:10.1007/s12010-016-2313-3

[42] Tiwari S, Santhyanathan D, Thomas E, Melarkode R, Ramavana G, Suryanarayan S. Fed batch solid state fermentation for the production of mycophenolic acid. India: World Intellectual Property Organization International, International Bureau; WO2003106690A1, 2003.

[43] El-Sayed ESR, Ahmed AS, Ismaiel AA. Agro-industrial byproducts for production of the immunosuppressant mycophenolic acid by Penicillium roqueforti under solid-state fermentation: Enhanced production by ultraviolet and gamma irradiation. Biocatal Agric Biotechnol. 2019;18: 101015. doi:10.1016/j.bcab.2019.01.053

[44] Survase SA, Shaligram NS, Pansuriya RC, Annapure US, Singhal RS. A novel medium for the enhanced production of Cyclosporin A by Tolypocladium inflatum MTCC 557 using solid state fermentation. J Microbiol Biotechnol. 2009;19: 462-467. doi:10.4014/jmb.0805.324

[45] Survase AS, Annapure US, Singhal RS. Statistical optimization for improved production of cyclosporin A in solid-state fermentation. J Microbiol Biotechnol. 2009;19: 1385-1392. doi:10.4014/jmb.0901.0035

[46] Nisha AK, Ramasamy K. Cyclosporin production in various solid substrate media by Tolypocladium inflatum (ATCC 34921). Biotechnology. 2008. pp. 357-359. doi:10.3923/biotech.2008.357.359

[47] Ramana Murthy M V, Mohan EVS, SadhukhanAK.Cyclosporin-Aproduction by Tolypocladium inflatum using solid state fermentation. Process Biochem. 1999;34: 269-280. doi:10.1016/S0032-9592(98)00095-8

[48] Sekar C, Rajasekar VW, Balaraman K. Production of cyclosporin A by solid state fermentation. Bioprocess Eng. 1997;17: 257-259. doi:10.1007/s12010-016-2313-3

[49] Ul NA, Khandige S, Patil NS, Khedkar AP, Melarkode R, Gururaja R, et al. Solid state fermentation and fed batch for the production of an immunosuppressant. India: World Intellectual Property Organization International, International Bureau; WO2004069845A1, 2004.

[50] Meruvu H, Meruvu HBT. Anticoagulant activity of partially
purified chitinase produced by Citrobacter freundii str. nov. haritD11 by fermentation of wheat bran coupled with fish scales. SN Appl Sci. 2019;1: 1-9. doi:10.1007/s42452-018-0097-5

[51] Meruvu H, Meruvu HBT. Biosynthesis and Characterization of an Anticoagulant Chitinase from Fermented Wheat Bran & Shrimp Shells’ Substratum. Trends Pept Protein Sci. 2018;3: 1-9. doi:https://doi.org/10.22037/tppps.v3i0.23165

[52] Wei X, Luo M, Xu L, Zhang Y, Lin X, Kong P, et al. Production of fibrinolytic enzyme from Bacillus amyloliquefaciens by fermentation of chickpeas, with the evaluation of the anticoagulant and antioxidant properties of chickpeas. J Agric Food Chem. 2011;59: 3957-3963. doi:10.1021/jf1049535

[53] Almalki MA, Dhas DSD, Vijayaraghavan P. Optimization of Fibrinolytic Enzyme Production from Bacillus sp. IND6 in Solid State Fermentation. J Sci Res Reports. 2017;16: 1-8. doi:10.9734/JSRR/2017/36748

[54] Vijayaraghavan P, Rajendran P, Gnana S, Vincent P, Arun A, Al-dhabi NA, et al. Novel Sequential Screening and Enhanced Production of Fibrinolytic Enzyme by Bacillus sp. IND12 Using Response Surface Methodology in Solid-State Fermentation. Biomed Res Int. 2017;2017: 1-13. doi:doi.org/10.1155/2017/3909657 Research

[55] Xin X, Ambati RR, Cai Z, Lei B. Development of universal purification protocols for fibrinolytic enzyme-producing bacilli. CyTA - J Food. 2019;17: 112-120. doi:10.1080/19476337.2018.1561521

[56] Bi H, Zhao H, Lu F, Zhang C, Bie X, Lu Z. Improvement of the nutritional quality and fibrinolytic enzyme activity of soybean meal by fermentation of Bacillus subtilis. J Food Process Preserv. 2014;39: 1-8. doi:10.1111/jfpp.12340

[57] Wu R, Chen G, Pan S, Ze J, Liang Z. Cost-effective fibrinolytic enzyme production by Bacillus subtilis WR350 using medium supplemented with corn steep powder and sucrose. Sci Rep. 2019;9: 1-10. doi:10.1038/s41598-019-43371-8

[58] Venkatanagaraju E, Divakar G. Production and Purification of Alkaline Fibrinolytic Enzyme from Bacillus cereus GD55 under Solid State. Curr Trends Biotechnol Pharm. 2015;9: 348-356.

[59] Devadhasan G, Arumugaperumal B. Bio-prospecting of cuttle fish waste and cow dung for the production of fibrinolytic enzyme from Bacillus cereus IND5 in solid state fermentation. 3 Biotech. 2016;6: 1-13. doi:10.1007/s13205-016-0553-0

[60] Vijayaraghavan P, Gnana S, Vincent P, Valan M, Al-dhabi NA. Statistical optimization of fibrinolytic enzyme production by Pseudoalteromonas sp. IND11 using cow dung substrate by response surface methodology. Sci World J. 2014;2014: 1-9.

[61] Vijayaraghavan P, Gnana S, Vincent P. medium Optimization for the Production of Fibrinolytic Enzyme by Paenibacillus sp. IND8 Using Response Surface Methodology. Sci World J. 2014;2014: 1-9.

[62] Vijayaraghavan P, Gnana S, Vincent P. Statistical optimization of fibrinolytic enzyme production by Pseudoalteromonas sp. IND11 using cow dung substrate by response surface methodology. Springerplus. 2014;3: 1-10.
Biotechnological Applications of Biomass

low-cost culture medium by response surface methodology. Saudi J Biol Sci. 2019;26: 217-224. doi:10.1016/j.sjbs.2018.08.029

[64] Seo J-H, Lee S-P. Production of Fibrinolytic Enzyme from Soybean Grits Fermented by Bacillus firmus NA-1. J Med. Food. 2004;7: 442-449.

[65] Pajeú T, Emmanuelle A, Souza C, Marcos R, Brandão P, Campos-takaki GM De, et al. Purification of a fibrinolytic protease from Mucor subtilissimus UCP 1262 by aqueous two-phase systems (PEG/sulfate). J Chromatogr B. 2016;1025: 16-24. doi:10.1016/j.jchromb.2016.04.046

[66] Tao S, Beihui L, Peng L, Deming L, Zuohu L. New solid-state fermentation process for repeated batch production of fibrinolytic enzyme by Fusarium oxysporum. Process Biochem. 1998;33: 419-422. doi:10.1016/S0032-9592(97)00096-4

[67] Tao S, Peng L, Beihui L, Deming L, Zuohu L. Solid state fermentation of rice chaff for fibrinolytic enzyme production by Fusarium oxysporum. Biotechnol Lett. 1997;19: 465-467.

[68] Cho HD, Min HJ, Won YS, Ahn HY, Cho YS, Seo K Il. Solid state fermentation process with Aspergillus kawachii enhances the cancer-suppressive potential of silkworm larva in hepatocellular carcinoma cells. BMC Complement Altern Med. 2019;19: 241. doi:10.1186/s12906-019-2649-7

[69] Ghanem KZ, MahranMZ, Ramadan MM, Ghanem HZ, Fadel M, Mahmoud MH. A comparative study on flavour components and therapeutic properties of unfermented and fermented defatted soybean meal extract. Sci Rep. 2020;10: 1-7. doi:10.1038/s41598-020-62907-x

[70] Kim HY, Heo DY, Park HM, Singh D, Lee CH. Metabolomic and transcriptomic comparison of solid-state and submerged fermentation of Penicillium expansum KACC 40815. PLoS One. 2016;11: 1-14. doi:10.1371/journal.pone.0149012

[71] Onipe OO, Jideani AIO, Beswa D. Composition and functionality of wheat bran and its application in some cereal food products. Int J Food Sci Technol. 2015;50: 2509-2518. doi:10.1111/ijfs.12935

[72] Mohamed SA, Saleh RM, Kabli SA, Al-garni SM. Influence of solid state fermentation by Trichoderma spp. on solubility, phenolic content, antioxidant, and antimicrobial activities of commercial turmeric. Biosci Biotechnol Biochem. 2016;8451: 1-9. doi:10.1080/09168451.2015.1136879

[73] José Serrano Vega R, Campos Xolalpa N, Josabad Alonso Castro A, Pérez González C, Pérez Ramos J, Pérez Gutiérrez S. Terpenes from Natural Products with Potential Anti-Inflammatory Activity. Terpenes and Terpenoids. 2018. pp. 59-85. doi:10.5772/intechopen.73215

[74] Kim T, Song B, Cho KS, Lee IS. Therapeutic potential of volatile terpenes and terpenoids from forests for inflammatory diseases. Int J Mol Sci. 2020;21. doi:10.3390/ijms21062187

[75] Yin ZN, Wu WJ, Sun CZ, Liu HF, Chen WB, Zhan QP, et al. Antioxidant and Anti-inflammatory Capacity of Ferulic Acid Released from Wheat Bran by Solid-state Fermentation of Aspergillus niger. Biomed Environ Sci. 2019;32: 11-21. doi:10.3967/bes2019.002

[76] Joseph S V, Edirisinghe I, Burton-Freeman BM. Fruit Polyphenols: A Review of Anti-inflammatory Effects in Humans. Crit Rev Food Sci Nutr. 2016;56: 419-444. doi:10.1080/10408398.2013.767221

[77] Yahfoufi N, Alsadi N, Jambi M, Matar C. The immunomodulatory and
The Application of Solid State Fermentation for Obtaining Substances Useful in Healthcare

DOI: http://dx.doi.org/10.5772/intechopen.94296

- Anti-inflammatory role of polyphenols. Nutrients. 2018;10: 1-23. doi:10.3390/nu10111618
- Dueñas M, Fernández D, Hernández T, Estrella I, Muñoz R. Bioactive phenolic compounds of cowpeas (Vigna sinensis L). Modifications by fermentation with natural microflora and with Lactobacillus plantarum ATCC 14917. J Sci Food Agric. 2005;85: 297-304. doi:10.1002/jsfa.1924
- Kitchin JES, Pomeranz MK, Pak G, Washenik K, Shupack JL. Rediscovering mycophenolic acid: A review of its mechanism, side effects, and potential uses. J Am Acad Dermatol. 1997;37: 445-449. doi:10.1016/S0190-9622(97)70147-6
- Wiseman AC. Immunosuppressive medications. Clin J Am Soc Nephrol. 2016;11: 332-343. doi:10.2215/CJN.08570814
- Huang M, Lu J-J, Huang M-Q, Bao J-L, Chen X-P, Wang Y-T. Terpenoids: natural products for cancer therapy. Expert Opin Investig Drugs. 2012;21: 1801-1818. doi:10.1517/13543784.2012.727395
- Cox-georgian D, Ramadoss N, Dona C, Basu C. Therapeutic and Medicinal Uses of Terpenes. In: Joshee N, Dhekney S, Parajuli P, editors. Medicinal Plants. Springer Nature Switzerland; pp. 333-359. doi:https://doi.org/10.1007/978-3-030-31629-5_15
- Blowman K, Magalhães M, Lemos MFL. Anticancer Properties of Essential Oils and Other Natural Products. Evidence-Based Complement Altern Med. 2018;2018: 1-12. doi:10.1155/2018/3149362 Review
- Fresco P, Borges F, Diniz C, Marques MPM. New insights on the anticancer properties of dietary polyphenols. Med Res Rev. 2006;26: 747-766. doi:10.1002/med.20060
- Mileo AM, Miccadi E. Polyphenols as Modulator of Oxidative Stress in Cancer Disease: New Therapeutic Strategies. Oxid Med Cell Longev. 2016;2016. doi:10.1155/2016/6475624
- Cao L, Guo X, Liu G, Song Y, Ho CT, Hou R, et al. A comparative analysis for the volatile compounds of various Chinese dark teas using combinatory metabolomics and fungal solid-state fermentation. J Food Drug Anal. 2018;26: 112-123. doi:10.1016/j.jfda.2016.11.020
- Ohara T, Mori T. Antiproliferative Effects of Short-chain Fatty Acids on Human Colorectal Cancer Cells via Gene Expression Inhibition. Anticancer Res. 2019;39: 4659-4666. doi:10.21873/anticancerres.13647
- Ghasemi Y, Dabbagh F, Ghasemian A. Cloning of a fibrinolytic enzyme (subtilisin) gene from bacillus subtilis in Escherichia coli. Mol Biotechnol. 2012;52: 1-7. doi:10.1007/s12033-011-9467-6
- Krieger N, Neto DC, Mitchell DA. Production of microbial biosurfactants by solid-state cultivation. Adv Exp Med Biol. 2010;672: 203-210. doi:10.1007/978-1-4419-5979-9_15
- Lv F, Zhang C, Guo F, Lu Y, Bie X, Qian H, et al. Expression, purification, and characterization of a recombined fibrinolytic enzyme from endophytic Paenibacillus polymyxa EJS-3 in Escherichia coli. Food Sci Biotechnol. 2015;24: 125-131. doi:10.1007/s10068-015-0018-y
- Xu J, Yang Y, Liu Y, Ran C, Li J, He S, et al. Characterization of an exo-chitinase from a Citrobacter strain isolated from the intestine content of large yellow croakers. Acta Microbiol Sin. 2016;56: 1089-1104. Available: https://pubmed.ncbi.nlm.nih.gov/29733170/
[92] Wang C, Yang B, Wang J. Generation of Streptomyces hygroscopicus cell factories with enhanced ascomycin production by combined elicitation and pathway-engineering strategies. Biotechnol Bioeng. 2019; 1-14. doi:10.1002/bit.27158

[93] Kishimoto N, Sugihara S, Mochida K, Fujita T. In Vitro Antifungal Analogs and Antiviral Utilized as Activities of γ- and δ-Lactone Analogs Utilized as Food Flavoring. Biocontrol Sci. 2005;10: 31-36. doi:10.4265/bio.10.31

[94] Guo Y, Song H, Wang Z, Ding Y. Expression of POX2 gene and disruption of POX3 genes in the industrial Yarrowia lipolytica on the γ-decalactone production. Microbiol Res. 2012;167: 246-252. doi:10.1016/j.micres.2011.10.003

[95] Park MJ, General T, Lee SP. Physicochemical properties of roasted soybean flour bioconverted by solid-state fermentation using Bacillus subtilis and Lactobacillus plantarum. Prev Nutr Food Sci. 2012;17: 36-45. doi:10.3746/pnf.2012.17.1.036

[96] Esteban-Torres M, Landete JM, Reverón I, Santamaría L, de las Rivas B, Muñoz R. A Lactobacillus plantarum Esterase Active on a Broad Range of. Appl Environ Microbiol. 2015;81: 3235-3242. doi:10.1128/AEM.00323-15

[97] Arnau J, Yaver D, Hjort CM. Strategies and Challenges for the Development of Industrial Enzymes Using Fungal Cell Factories. 1st ed. In: Nevalainen H, editor. Grand Challenges in Fungal Biotechnology. 1st ed. Springer International Publishing; 2020. pp. 179-210. doi:10.1007/978-3-030-29541-7_7

[98] Chen D, Shi J, Hu X. Enhancement of polyphenol content and antioxidant capacity of oat (Avena nuda L.) bran by cellulase treatment. Appl Biol Chem. 2016;59: 397-403. doi:10.1007/s13765-016-0171-x

[99] Salgado JM, Abrunhosa L, Venâncio A, Domínguez JM, Belo I. Integrated use of residues from olive mill and winery for lipase production by solid state fermentation with Aspergillus sp. Appl Biochem Biotechnol. 2014;172: 1832-1845. doi:10.1007/s12010-013-0613-4

[100] Das P, Mukherjee S, Sen R. Genetic regulations of the biosynthesis of microbial surfactants: An overview. Biotechnol Genet Eng Rev. 2008;25: 165-186. doi:10.5661/bger-25-165

[101] Ghosh B, Ray RR. Current Commercial Perspective of Rhizopus oryzae: A Review. J Appl Sci. 2011;11: 2470-2486. doi:10.3923/jas.2011.2470.2486

[102] Mata G, Salbones D, Pérez-merlo R. Hydrolytic enzyme activities in shiitake mushroom (Lentinula edodes) strains cultivated on coffee pulp. Rev Argent Microbiol. 2016;48: 191-195. doi:10.1016/j.ram.2016.05.008

[103] Amore A, Giacobbe S, Faraco V. Regulation of Cellulase and Hemicellulase Gene Expression in Fungi. Curr Genomics. 2013;14: 230-249. doi:10.2174/138920291340400002

[104] Battaglia E, Benoit I, Brink J Van Den, Wiebenga A, Coutinho PM, Henrissat B, et al. Carbohydrate-active enzymes from the zygomycete fungus Rhizopus oryzae: a highly specialized approach to carbohydrate degradation depicted at genome level. BMC Genomics. 2011;12: 38. doi:10.1186/1471-2164-12-38

[105] Quin MB, Flynn CM, Schmidt-Dannert C. Traversing the fungal terpenome. Nat Prod Rep. 2015;31: 1449-1473. doi:10.1039/c4np00075g:Traversing
[106] Regueira TB, Kildegaard KR, Hansen BG, Mortensen UH, Hertweck C, Nielsen J. Molecular basis for mycophenolic acid biosynthesis in Penicillium brevicompactum. Appl Environ Microbiol. 2011;77: 3035-3043. doi:10.1128/AEM.03015-10

[107] Bushley KE, Raja R, Jaiswal P, Cumbie JS, Nonogaki M, Boyd AE, et al. The Genome of Tolypocladium inflatum: Evolution, Organization, and Expression of the Cyclosporin Biosynthetic Gene Cluster. PLOS Genet. 2013;9: 20. doi:10.1371/journal.pgen.1003496

[108] Lu C, Chen S. Fibrinolytic Enzymes from Medicinal Mushrooms. In: Faraggi E, editor. Protein Structure. InTech; 2012. pp. 338-362. doi:10.5772/38221

[109] Hynes MJ, Murray SL, Duncan A, Khew GS, Davis MA. Regulatory genes controlling fatty acid catabolism and peroxisomal functions in the filamentous fungus Aspergillus nidulans. Eukaryot Cell. 2006;5: 794-805. doi:10.1128/EC.5.5.794-805.2006

[110] Nascimento TP, Sales AE, Porto CS, Marcos R, Brandão P, Maria G, et al. Production and Characterization of New Fibrinolytic Protease from Mucor subtilissimus UCP 1262 in Solid-State Fermentation. Adv Enzym Res. 2015;3: 81-91.

[111] Sun D, Liao J, Sun L, Wang Y, Liu Y, Deng Q, et al. Effect of media and fermentation conditions on surfactin and iturin homologues produced by Bacillus natto NT-6: LC – MS analysis. AMB Express. 2019;9: 1-9. doi:10.1186/s13568-019-0845-y