An Update on Oligosaccharides and Their Esters from Traditional Chinese Medicines: Chemical Structures and Biological Activities

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A great number of naturally occurring oligosaccharides and oligosaccharide esters have been isolated from traditional Chinese medicinal plants, which are used widely in Asia and show prominent curative effects in the prevention and treatment of kinds of diseases. Numerous in vitro and in vivo experiments have revealed that oligosaccharides and their esters exhibited various activities, including antioxidant, antidepressant, cytotoxic, antineoplastic, anti-inflammatory, neuroprotective, cerebral protective, antidiabetic, plant growth-regulatory, and immunopotentiating activities. This review summarizes the investigations on the distribution, chemical structures, and bioactivities of natural oligosaccharides and their esters from traditional Chinese medicines between 2003 and 2013.

1. Introduction

Oligosaccharides and their esters, a significant group of phytochemical compounds, are widely distributed in the roots, rhizomes, stems, barks, leaves, aerial, and whole parts of medicinal plants. They not only serve as the energy storage components, but also play a vital role in the treatment of diseases. Before 2003, there have been a number of reviews and reports in respect to the isolation and structure elucidation of oligosaccharides and their esters from Chinese medicinal plants [1–3], but few biological activities such as cancer chemopreventive, and protein kinase C inhibitory activities had been reported [4–6]. With the development of isolation and identification techniques [7–11], a larger number of oligosaccharides and their esters have been endlessly identified from traditional Chinese medicines in the past decades. These compounds have a wide variety of structure types because of the assembly of different monosaccharide units, the combination of various linking styles and the existence of kinds of substituents. And more promising biological activities associated with some of the oligosaccharides and their esters have been discovered. In vitro and in vivo investigations have demonstrated that they displayed antioxidant, antidepressant, anti-inflammatory, neuroprotective, cerebral protective, antidiabetic, cytotoxic, antineoplastic, plant growth-regulatory, and immunopotentiating activities, and so forth. This review aims to provide a systemic summary of the studies on the distribution, chemical structures and biological activities of naturally occurring oligosaccharides and their esters from traditional Chinese medicines in the past decades. Among these compounds, the number of oligosaccharide esters is much greater than that of oligosaccharides, and the disaccharide esters are a very valuable source of active compounds. This information may help readers understand the structure characteristics and therapeutic indications of oligosaccharides and their esters from traditional Chinese medicines and offer clues to the development of new drugs.

2. Chemical Structures

Phytochemical investigations of traditional Chinese medicines have shown that many botanical families, including Polygalaceae, Liliaceae, Asteraceae, Polygonaceae, Smilacaceae, Scrophulariaceae, Asclepiadaceae, Arecaceae,
Orobanchaceae, Acanthaceae, Rosaceae, Musaceae, Spargan-
iaceae, Leguminosae, Equisetaceae, Boraginaceae, Iridaceae,
Alismataceae, Lamiaceae, Araliaceae, Rubiaceae, Oleaceae,
Apocynaceae, Caryophyllaceae, Aspleniaceae and Trilliaceae,
are rich in oligosaccharides and their esters. Oligosaccharides
show diversified structures because of the type and the
number of monosaccharides, as well as the position of
glycosidic bonds. And oligosaccharide esters also
display distinctive structural diversity largely owing to the
number, type, and position of O-substituent units, including
phenylpropanoid groups (e.g., coumaroyl, feruloyl, caffeoyl,
sinapoyl, 3,4,5-trimethoxycinnamoyl, and cinnamoyl),
benzoyl, p-methoxybenzoyl, and p-hydroxybenzoyl groups
(Figure 2). Moreover, the double bonds of phenylpropanoid
groups possess trans and cis isomeric forms, of which the
trans forms widely exist in nature. Hence, according to
the number of monosaccharides and the characteristics of
chemical structures, these oligosaccharide esters could be
categorized into 7 large groups.

2.1. Oligosaccharides. All compounds of this group (Table 1
and Figure 1) merely consist of various monosaccharides
without O-substituents. In addition to the well-known
sucrose, β-D-glucopyranosyl(1 → 2)-β-D-glucopyranoside
(1) was isolated from Camptosorus sibiricus [12]. The
oligosaccharides of raffinose (3), stachyose (19), and
verbascose (21), all of which belong to the Raffinose family,
possess one, two or three galactopyranosyl units linked to
sucrose and have been found in the rhizomes and roots of
Alisma orientalis [13], Lycopus lucidus [14], Rehmannia
glutinosa [15, 16], Salvia miltiorrhiza [17], and Scrophularia
ningpoensis [18]. Manninotriose (4) and verbascotetraose (5)
consisting of galactopyranosyl units and a glucopyranosyl
unit have been isolated from Alisma orientalis [13].

Five oligosaccharides comprising 1-kestose (6), nystose
(7), 1-β-Fructofuranosynystose (8), hexascarabide (9), and
heptasaccharide (10) consisting of fructofuranose and glu-
copyranose have been isolated from the aerial parts and roots
of Gynura divaricata subsp. formosana [19], Morinda offici-
nalis [20–22], Saussurea lappa [23], and Aralia cordata [24].
Two water-soluble oligosaccharides (11, 12) composed of two
or three types of monosaccharides including glucopyranose,
fructopyranose, and fructofuranose have been obtained from
the whole plants of Blumea riparia [25, 26].

Besides, malt-o-oligosaccharides (17, n = 0–8) consist-
ing of α-D-glucopyranosyl residues assembled by (1 → 4)-
linkages and inulo-oligosaccharides (18, n = 1–3) consisting of
only fructosyl residues formed by (2 → 1)-linkages have been
found in the roots of Panax ginseng [27, 28] and Morinda
officinalis [20], respectively. Three noteworthy oligosaccha-
rides (2, 13, 14) formed by α-D-glucopyranosyl units with
(1 → 6)-linkages and a (1 → 4)-linkage have been found in the
roots of Panax ginseng [28]. And two linear oligosaccharides
termed heptasaccharide (15) and octasaccharide (16) consist-
ing of glucose and mannose monomers were identified from
the rhizomes of Paris polyphylla var. yunnanensis [29, 30].
A pentasaccharide, stellarioside (20) consisting of a raffinose
backbone with two galactosyl residues bound to the fructosyl
and glucosyl moieties was identified from the stems of
Stellaria media [31].

Oligosaccharides (Table 1) are composed of seven kinds of
deoxyhexoses including cymaropyranose, canaropyra-
nose, digitoxopyranose, oleandropyranose, digitalopyranose,
cymaropyranurolactone, and oleandronic acid-δ-lactone.
Oleandronic acid-δ-lactone exhibits the boat and chair con-
formations. The hydroxyl, methyl, and acetyl groups are
located at the equatorial (e) and axial (a) bonds in the chair
conformation of deoxyhexoses. These oligosaccharides were
isolated from the traditional Chinese medicines including the
roots of Periploca forrestii, the root barks of P. sepium, the
stems of P. calophylla, and the barks of Parabarrium huatingii.

2.2. Oligosaccharide Esters

2.2.1. Phenylpropanoid-Derived Disaccharide Esters. Phenyl-
propanoid-derived disaccharide esters (Table 2 and Figure 3)
account for a considerable proportion of oligosaccharide
esters and mainly possess a core of sucrose carrying a
varying number of O-substituents, includingphenyl-
propanoid groups, acetyl, benzoyl, p-methoxybenzoyl, and
p-hydroxybenzoyl groups. Phenylpropanoid substituents are
just present at 1′, 3′, 4′, 6′ positions of β-D-fructofuranosyl
unit in compounds 35–97, whereas they appear at 2, 3, 4,
6 positions of α-D-glucopyranosyl moiety in compounds
98–101. Moreover, compounds 76–97 are mainly esterified
with acetyl groups along with a phenylpropanoyl substi-
tuent, coumaroyl, feruloyl, or 3,4,5-trimethoxycinnamoyl
group. Interestingly, the phenylpropanoid substituents are
only attached to the 3′ position of sucrose. The two
sugar rings of compounds 102–128 both possess phenyl-
propanoid substituents. These oligosaccharide esters have
been found in the roots and rhizomes of Polygala tricornis,
P. tenuifolia, Fagopyrum tataricum, Scrophularia ningpoensis,
Cynanchum amplexicaule, Smilax riparia, Paris polyphylla
var. yunnanensis, Smilacis glabrae, Fagopyrum dibotrys,
and Sparganium stoloniferum, the underground parts of Tri-
lium kamtschaticum, the stems of Polygonum sachalinensis,
P. cuspidatum, P. hydropiper, Smilax china, and Calamus
quisquetinervius, the aerial parts of Polygala sibirica, Smilax
bracteata, Heterosmilax erythrantha, and Musella lasiocarpa,
the leaves of Persicaria hydropiper and Polygonum hydropiper,
the whole plants of Bidens parviflora and Polygala hongkon-
gensis, and the flower buds of Prunus mume.

Cistanoside F (129) has been found in the stems of
Cistanthe tubulosa [80] and C. sinensis [81], the barks of Pau-
lownia tomentosa var. tomentosa [82], and the aerial parts
of Acanthus ilicifolius [83]. Cistanoside I (130) has also
been isolated from the stems of the Cistanthe plants [84].
Both of them are composed of glucosyl and rhamnosyl
groups connected by a 1 → 3 glycosidic bond. In addition,
6,6′-sucrose ester of (1α,2x3,β,4β)-3,4-bis(4-hydroxy-
phenyl)-1,2-cyclobutanedicarboxylic acid (131) with a bis(4-
hydroxyphenyl) cyclobutanedicarboxyl group as the acyl unit
in the molecule structure was isolated from the whole plants
of Bidens parviflora [61].
| No. | Name                  | R₁    | R₂      | R₃    | R₄      | R₅    | R₆    | Source          | Parts       | Reference |
|-----|-----------------------|-------|---------|-------|---------|-------|-------|----------------|-------------|-----------|
| 22  | Perifosaccharide A    | OH(e) | OCH₃(e) | H     | OH(e)   | OH    | A     | Periploca forrestii | Roots       | [32]      |
| 23  | Perifosaccharide B    | OH(e) | OCH₃(a) | H     | OH(e)   | OH    | A     | Periploca forrestii | Roots       | [32]      |
| 24  | Perifosaccharide C    | OH(e) | OCH₃(e) | H     | OH(e)   | OCH₃  | A     | Periploca forrestii | Roots       | [32]      |
| 25  | Perifosaccharide D    | OAc(e) | OCH₃(e) | H     | OH(e)   | OH    | A     | Periploca forrestii | Roots       | [32]      |
| 26  | Perisaccharide A      | OH(a) | OCH₃(e) | OAc   | OH(e)   | OCH₃  | A     | Periploca sepium   | Root barks  | [33]      |
| 27  | Perisaccharide B      | OAc(e) | OCH₃(a) | H     | OH(e)   | OH    | A     | Periploca sepium   | Periploca calophylla | Root barks | [33, 34] |
| 28  | Perisaccharide C      | OH(a) | OCH₃(e) | OAc   | OCH₃(a) | OH    | A     | Periploca sepium   | Root barks  | [33]      |
| 29  | Perisaccharide D      | OAc(e) | OCH₃(a) | H     | OH(e)   | OCH₃  | A     | Periploca calophylla | stems       | [34]      |
| 30  | Perisesaccharide B    | OH(a) | OCH₃(e) | OAc   | OH(e)   | OCH₃  | C     | Periploca sepium   | Root barks  | [35]      |
| 31  | Perisesaccharide C    | OH(a) | OCH₃(e) | OH    | OCH₃(a) | OCH₃  | C     | Periploca sepium   | Root barks  | [35]      |
| 32  | Perisesaccharide D    | OH(a) | OCH₃(e) | OH    | OCH₃(a) | OH    | C     | Periploca sepium   | Root barks  | [35]      |
| 33  | Perisesaccharide E    | OH(a) | OCH₃(e) | OAc   | OH(e)   | OH    | C     | Periploca sepium   | Root barks  | [35]      |
| 34  | Cymaropyranurolactone | OH(a) | OCH₃(e) | OH    | OCH₃(e) | OCH₃  | B     | Parabarium huatingii | Barks       | [36]      |

See Scheme 1.
2.2.4. Phenylpropanoid-Derived Trisaccharide Esters. In this group, all oligosaccharides including glucopyranose, fructofuranose, and rhapnopyranose with different types of fatty acid residues which attach to the 6 or 6′ position of sucrose. These fatty acids include oleandric acid, palmitic acid, linolenic acid, myristic acid, hexadeca-7,10,13-trienoic acid, and hexadeca-7,10-dienoic acid. The above sucrose fatty acid esters have been found in the rhizomes of Astragalus membranaceus and the roots of Equisetum hiemale.

2.2.5. Phenylpropanoid-Derived Tetrasaccharide Esters. Phenylpropanoid-derived tetrasaccharide esters (Table 5) consisting of three glucopyranosyl units and a fructofuranosyl unit have been isolated from the roots of Polygala tenuifolia. The nonsugar moieties of these oligosaccharides include coumaroyl, feruloyl, sinapoyl, and benzoyl groups, respectively.

2.2.6. Phenylpropanoid-Derived Pentasaccharide Esters. As shown in Table 6, oligosaccharide esters (160–179) possessing a skeleton of five sugar residues have been isolated from the roots of Polygala tenuifolia. The sugar residues are composed of two types of monosaccharides including fructofuranose and glucopyranose, which are esterified with acetyl, benzoyl, rhamnose-substituted/nonsubstituted coumaroyl, and rhamnose-substituted/nonsubstituted feruloyl groups. Other than that, a structure-complex oligosaccharide polymer shown in Figure 5, polygalajaponicose I (180), consisting of a pentasaccharide backbone esterified with feruloyl, coumaroyl, rhamnosyl-coumaroyl, acetyl, and benzoyl groups has been obtained from the roots of P. japonica [90].

2.2.7. Others. Polygalatenosides A–C (181–183) (Figure 6) containing a galactosyl unit and a polygalolinosyl unit esterified with benzoyl groups at 3, 4 and 6 positions have been found in the roots of Polygala tenuifolia [91]. Three sucrose esters, including polygalatenoside D (190), telephiose F (191), and 6-O-benzylsucrose (192), possess one benzoyl group, two benzoyl groups, and a p-methoxybenzoyl group, respectively. They were isolated from the roots of P. tenuifolia [91], the whole plants of P. telephioide (92), and the roots of P. tricorin (37). Six trisaccharide esters, named telephiose A–E and G (184–189) with substituents of acetyl and benzoyl groups, were isolated from the whole plants of P. telephioide [92, 93]. Moreover, a trisaccharide ester (193), pubescenside A from the flowers of Syringa pubescens, possesses a fatty acid residue [94].

3. Biological Activities of Oligosaccharides and Their Esters

The oligosaccharides and oligosaccharide esters from Chinese medicinal plants are important products with diversified structures, which have triggered an increasing number of studies carried out on the isolated compounds. And thus diverse pharmacological activities have been proved. Among the isolated compounds, oligosaccharides, phenylpropanoid-derived disaccharide esters and trisaccharide esters, fatty acid-derived disaccharide esters, and others from the families Polygonaceae, Asclepiadaceae, Rubiaceae, Polyalaceae, Liliaceae, Smilacaceae, Arecaceae, Orobancheaceae, Scrophulariaceae, Acanthaceae, Rosaceae, Sparganiaceae, Leguminosae, and Equisetaceae have shown significant pharmacological activities including antioxidant,
Figure 1: Continued.
Figure 1

D: benzoyl  E: \( p \)-methoxybenzoyl  F: \( p \)-hydroxybenzoyl  G: coumaroyl

I: feruloyl  J: caffeoyl  K: sinapoyl

L: 3,4,5-trimethoxycinnamoyl  M: cinnamoyl

Figure 2
| Number | Name                  | R_1 | R_2 | R_3 | R_4 | R_5 | R_6 | R_7 | R_8 | Source                  | Parts            | Reference         |
|--------|-----------------------|-----|-----|-----|-----|-----|-----|-----|-----|-------------------------|------------------|------------------|
| 35     | Sibiricose A_6        | H   | H   | H   | H   | K   | H   | H   | H   | Polygala tricornis      | Roots            | [37–40]          |
|        | 3'-O-Feruloylsucrose/Sibiricose A_5 | H   | H   | H   | H   | I   | H   | H   | H   | Polygala tenuifolia    | Root barks       |                  |
| 36     | Glomeratose A         | H   | H   | H   | H   | L   | H   | H   | H   | Polygala tricornis      | Underground parts | [38–42]          |
| 37     | Lapathoside D         | H   | H   | H   | H   | G   | G   | H   | H   | Polygonum sachalinense  | Roots            | [37]             |
| 38     | Glomeratose A         | H   | H   | H   | H   | I   | I   | H   | H   | Trillium kamtschaticum  | Stems            | [43]             |
|        | Tenuifoliside A       | F   | H   | H   | H   | L   | H   | H   | H   | Trillium kamtschaticum  | Underground parts |                  |
| 39     | Helonioside A         | H   | H   | H   | H   | I   | I   | H   | H   | Smilax bracteata       | Aerial parts     | [41, 44, 45]     |
| 40     | Helonioside B         | Ac  | H   | H   | H   | I   | I   | H   | H   | Smilax riparia          | Stems            | [44, 46–48]      |
| 41     | Parispolyside F       | H   | H   | H   | H   | G   | I   | H   | H   | Paris Polyphylla var. yunnanensis | Rhizomes      | [49, 50]          |
| 42     | Hydropiperoside       | H   | H   | H   | H   | G   | G   | H   | G   | Polygonum sachalinense  | Stems            | [43, 51, 52]     |
| 43     | Tricornose B          | D   | Ac  | H   | H   | L   | H   | H   | H   | Persicaria hydropiper   | Leaves           |                  |
| 44     | Tenuifoliside A       | F   | H   | H   | H   | L   | H   | H   | H   | Polygala tricornis      | Roots            | [37]             |
| 45     | Tatariside A          | Ac  | H   | H   | H   | Ac  | G   | G   | H   | Polygala tenuifolia     | Whole plants     | [38, 40, 53–56]  |
| 46     | Tatariside E          | H   | H   | H   | H   | Ac  | G   | H   | Ac  | Fagopyrum tataricum     | Stems            | [57]             |
| 47     | Tatariside G          | H   | H   | H   | H   | I   | G   | H   | G   | Fagopyrum tataricum     | Roots            | [57]             |
| 48     | Smiglaside A          | Ac  | Ac  | H   | Ac  | I   | I   | H   | I   | Smilax riparia          | Roots and rhizomes |                  |
| 49     | Smiglaside B          | Ac  | H   | H   | Ac  | I   | I   | H   | I   | Smilax riparia          | Roots and rhizomes |                  |
| 50     | Smiglaside E          | Ac  | H   | H   | Ac  | I   | I   | H   | G   | Smilax china            | Stems            | [46]             |
| 51     | Smilaside A           | Ac  | Ac  | H   | Ac  | I   | I   | H   | G   | Smilax china            | Stems            | [46]             |
| 52     | Smilaside B           | H   | H   | H   | Ac  | I   | I   | H   | H   | Smilax china            | Stems            | [46]             |
| 53     | Smilaside C           | H   | H   | H   | H   | I   | I   | H   | G   | Smilax china            | Stems            | [46]             |
| 54     | Smilaside D           | H   | H   | H   | H   | I   | I   | Ac  | G   | Smilax china            | Stems            | [46]             |
Table 2: Continued.

| Number | Name          | R₁ | R₂ | R₃ | R₄ | R₅ | R₆ | R₇ | R₈ | Source                  | Parts        | Reference |
|--------|---------------|----|----|----|----|----|----|----|----|-------------------------|--------------|-----------|
| 55     | Smilaside E   | Ac | H  | H  | H  | I  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44, 46]  |
| 56     | Smilaside F   | Ac | H  | H  | Ac | G  | I  | H  | G  | Smilax china            | Stems        | [46]      |
| 57     | Smilaside G   | H  | H  | H  | H  | G  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44, 58]  |
| 58     | Smilaside H   | H  | H  | H  | Ac | G  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44]      |
| 59     | Smilaside I   | Ac | H  | H  | H  | G  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44]      |
| 60     | Smilaside J   | H  | H  | H  | H  | G  | I  | H  | I  | Smilax bracteata        | Aerial parts | [44, 58]  |
| 61     | Smilaside K   | H  | H  | H  | Ac | I  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44]      |
| 62     | Smilaside L   | H  | H  | H  | H  | I  | I  | H  | I  | Smilax bracteata        | Aerial parts | [44, 58]  |
| 63     | Smilaside M   | Ac | H  | H  | Ac | Cis-feruloyl | I | H  | H  | Smilax riparia          | Roots and rhizomes | [59]      |
| 64     | Smilaside N   | Ac | H  | H  | Ac | I | Cis-feruloyl | H  | H  | Smilax riparia          | Roots and rhizomes | [59]      |
| 65     | Smilaside P   | H  | H  | H  | Ac | I  | I  | H  | I  | Smilax riparia          | Roots and rhizomes | [47]      |
| 66     | 5',4',6'--Tri-feruloylsucrose | H  | H  | H  | H  | I  | I  | I  | H  | Smilax riparia          | Rhizomes and roots | [60]      |
| 67     | 6'-O-Coumaroylsucrose | H  | H  | H  | H  | H  | G  | H  | H  | Bidens parviflora       | Whole plants | [61]      |
| 68     | 1'-O-Coumaroyl-6'-O-feruloylsucrose | H  | H  | H  | H  | H  | I  | H  | G  | Smilax bracteata        | Aerial parts | [44]      |
| 69     | 4-O-Benzoyl-3',3,4,5-trimethoxycinnamoylsucrose | H  | D  | H  | H  | L  | H  | H  | H  | Polygala tricornis       | Roots        | [37]      |
| 70     | 6-O-Benzoyl-3',3,4,5-trimethoxycinnamoylsucrose | D  | H  | H  | L  | H  | H  | H  | H  | Polygala tricornis       | Roots        | [37]      |
| 71     | 6-O-Benzoyl-3',3,4,5-sinapoylsucrose | D  | H  | H  | K  | H  | H  | H  | H  | Polygala tricornis       | Roots        | [37]      |
| 72     | 6-O-p-Methoxybenzoyl-3',3,4,5-trimethoxycinnamoylsucrose | E  | H  | H  | L  | H  | H  | H  | H  | Polygala tenuifolia     | Roots        | [56]      |
| 73     | 2,6-Di-acetyl-3',6'-di-feruloylsucrose | Ac | H  | H  | Ac | I  | I  | H  | H  | Smilax china            | Stems        | [46-48, 59]|
| 74     | 2,6-Di-acetyl-3',6'-cis-feruloyl-6'-trans-feruloylsucrose | Ac | H  | H  | Ac | Cis-feruloy | I | H  | H  | Smilax riparia          | Roots and rhizomes | [59]      |
| Number | Name | R₁ | R₂ | R₃ | R₄ | R₅ | R₆ | R₇ | R₈ | Source | Parts | Reference |
|--------|------|----|----|----|----|----|----|----|----|--------|-------|------------|
| 75     | 2,6-Di-acetyl-3'-O-trans-feruloyl-6'-cis-feruloylsucrose | Ac | H | H | Ac | I | Cis-feruloyl | H | H | *Smilax riparia* | Roots and rhizomes | [59] |
| 76     | Regaloside A | Ac | H | H | H | I | H | H | H | *Trillium kamtschaticum* | Underground parts | [41] |
| 77     | Tricosorne A | Ac | H | H | H | L | H | H | H | *Polygala tricornis* | Flower buds | [37] |
| 78     | Mumeose A | H | H | Ac | G | H | H | H | H | *Prunus mume* | Flower buds | [62, 63] |
| 79     | Mumeose B | Ac | H | Ac | H | G | H | H | H | *Prunus mume* | Flower buds | [62, 63] |
| 80     | Mumeose C | Ac | H | Ac | Ac | G | H | H | H | *Prunus mume* | Flower buds | [62, 63] |
| 81     | Mumeose D | Ac | Ac | Ac | Ac | G | H | H | Ac | *Prunus mume* | Flower buds | [62, 63] |
| 82     | Mumeose E | Ac | Ac | Ac | Ac | Cis-coumaroyl | H | H | Ac | *Prunus mume* | Flower buds | [62, 63] |
| 83     | Mumeose F | Ac | Ac | Ac | H | G | H | H | H | *Prunus mume* | Flower buds | [63] |
| 84     | Mumeose G | Ac | H | Ac | H | G | H | Ac | H | *Prunus mume* | Flower buds | [63] |
| 85     | Mumeose H | H | H | Ac | Ac | G | H | Ac | H | *Prunus mume* | Flower buds | [63] |
| 86     | Mumeose I | Ac | Ac | Ac | Ac | G | H | Ac | H | *Prunus mume* | Flower buds | [63] |
| 87     | Mumeose J | Ac | Ac | Ac | Ac | G | H | Ac | Ac | *Prunus mume* | Flower buds | [63] |
| 88     | Mumeose K | H | H | Ac | Ac | G | H | Ac | H | *Prunus mume* | Flower buds | [64] |
| 89     | Mumeose L | Ac | H | Ac | Ac | G | H | Ac | H | *Prunus mume* | Flower buds | [64] |
| 90     | Mumeose M | Ac | Ac | Ac | H | G | H | Ac | Ac | *Prunus mume* | Flower buds | [64] |
| 91     | Mumeose N | Ac | Ac | Ac | H | G | Ac | Ac | H | *Prunus mume* | Flower buds | [64] |
| 92     | Mumeose O | Ac | Ac | Ac | H | G | Ac | Ac | Ac | *Prunus mume* | Flower buds | [64] |
| 93     | Mumeose P | Ac | Ac | Ac | Ac | G | H | H | Ac | *Musella lasiocarpa* | Aerial parts | [65] |
| 94     | Sibirioside A | H | H | H | H | H | H | H | H | *Musella lasiocarpa* | Aerial parts | [65] |
| 95     | 1',2,3,4,6-O-Penta-acetyl-3'-O-trans-coumaroylsucrose | Ac | Ac | Ac | Ac | Cis-coumaroyl | H | H | Ac | *Musella lasiocarpa* | Aerial parts | [65] |
| 96     | 1',2,3,4,6-O-Penta-acetyl-3'-O-cis-coumaroylsucrose | Ac | Ac | Ac | Ac | Cis-coumaroyl | H | H | Ac | *Musella lasiocarpa* | Aerial parts | [65] |
| 97     | 1',2,3,6-O-Tetra-acetyl-3'-O-cis-feruloylsucrose | Ac | H | Ac | Ac | Cis-feruloyl | H | H | Ac | *Sparganium stoloniferum* | Rhizomes | [66] |
| 98     | 1',2,4,6-O-Tetra-acetyl-3'-O-trans-feruloylsucrose | Ac | Ac | Ac | Ac | H | H | H | Ac | *Sparganium stoloniferum* | Rhizomes | [66] |
| 99     | Sibricose A₁ | M | H | H | H | H | H | H | H | *Scrophularia ningpoensis* | Roots | [67] |
| 100    | 6-O-Caffeoylsucrose | K | H | H | H | H | H | H | H | *Scrophularia ningpoensis* | Roots | [68] |
| 101    | Acreteside | J | H | H | H | H | H | H | H | *Scrophularia ningpoensis* | Roots | [68] |
| Number | Name                  | $R_1$ | $R_2$ | $R_3$ | $R_4$ | $R_5$ | $R_6$ | $R_7$ | $R_8$ | Source                  | Parts     | Reference     |
|--------|-----------------------|-------|-------|-------|-------|-------|-------|-------|-------|--------------------------|-----------|--------------|
| 102    | Tenuifoliside B       | F     | H     | H     | H     | K     | H     | H     | H     | *Polygala tenuifolia*   | Roots     | [42, 56]     |
| 103    | Tenuifoliside C       | K     | H     | H     | H     | L     | H     | H     | H     | *Polygala tenuifolia*   | Roots     | [37, 42, 56] |
| 104    | Heterosmilasides     | H     | H     | I     | H     | H     | I     | H     | H     | *Heterosmilax*          | Aerial parts | [48]         |
| 105    | Quiquesetinerviuside A| H     | I     | H     | H     | I     | I     | H     | H     | *Calamus quiquesetinervius* | Stems    | [70]         |
| 106    | Quiquesetinerviuside B| Ac    | I     | H     | H     | I     | I     | H     | H     | *Calamus quiquesetinervius* | Stems    | [70]         |
| 107    | Quiquesetinerviuside C| H     | I     | H     | Ac    | I     | I     | H     | H     | *Calamus quiquesetinervius* | Stems    | [70]         |
| 108    | Quiquesetinerviuside D| Ac    | G     | H     | H     | I     | I     | H     | H     | *Calamus quiquesetinervius* | Stems    | [70]         |
| 109    | Quiquesetinerviuside E| H     | G     | H     | Ac    | I     | I     | H     | H     | *Calamus quiquesetinervius* | Stems    | [70]         |
| 110    | Vanicoside A          | I     | H     | H     | Ac    | G     | G     | H     | G     | *Polygonum cuspidatum*   | Stems    | [51, 52, 71, 72] |
|        |                       |       |       |       |       |       |       |       |       | *Polygonum hydropiper*   | Leaves    | [51, 52, 71, 72] |
|        |                       |       |       |       |       |       |       |       |       | *Polygonum sachalinensis*| Rhizomes  | [51, 52, 71, 72] |
| 111    | Vanicoside B          | I     | H     | H     | H     | G     | G     | H     | G     | *Polygonum cuspidatum*   | Stems    | [43, 51, 52, 71, 72] |
|        |                       |       |       |       |       |       |       |       |       | *Polygonum hydropiper*   | Leaves    | [51]         |
|        |                       |       |       |       |       |       |       |       |       | *Polygonum sachalinensis*| Stems and leaves | [72]     |
| 114    | Lapathoside A         | I     | H     | H     | H     | G     | G     | H     | I     | *Polygonum cuspidatum*   | Stems    | [51, 73]     |
| 115    | Lapathoside C         | I     | H     | H     | H     | G     | G     | H     | H     | *Polygonum sachalinensis*| Stems    | [43, 52]     |
| 116    | Diboside A            | G     | H     | H     | H     | G     | I     | H     | G     | *Fagopyrum dibotrys*     | Roots     | [43, 57]     |
| 117    | Hidropiperoside A     | I     | H     | H     | H     | G     | H     | G     | G     | *Polygonum hydropiper*   | Stems and leaves | [72]     |
| 118    | Hidropiperoside B     | I     | H     | H     | Ac    | G     | G     | H     | I     | *Polygonum hydropiper*   | Stems and leaves | [72]     |
| 119    | Tatariside B          | I     | H     | H     | Ac    | G     | G     | H     | Ac    | *Fagopyrum tataricum*    | Roots     | [57]         |
Table 2: Continued.

| Number | Name                        | R₁  | R₂  | R₃  | R₄  | R₅  | R₆  | R₇  | Source                                  | Parts          | Reference                  |
|--------|-----------------------------|-----|-----|-----|-----|-----|-----|-----|-----------------------------------------|----------------|-------------------------------|
| 120    | Tatariside C                | I   | Ac  | H   | Ac  | G   | G   | H   | Ac                                      | Roots          | [57]                         |
| 121    | Tatariside D                | G   | H   | H   | Ac  | G   | I   | H   | H                                       | Roots          | [57]                         |
| 122    | Tatariside F                | G   | H   | H   | H   | I   | I   | H   | G                                       | Roots          | [57]                         |
| 123    | 3',6'-O-Di-sinapoylsucrose  | K   | H   | H   | H   | K   | H   | H   | H                                       | Root barks     | [37–39, 41, 53–56, 69, 74] |
| 124    | 6,6'-O-Di-coumaroylsucrose  | G   | H   | H   | H   | G   | G   | H   | H                                       | Whole plants   | [61]                         |
| 125    | 1',3',6'-O-Tri-coumaroyl-6-feruloylsucrose | I   | H   | H   | H   | G   | G   | H   | G                                       | Roots          | [75, 76]                    |
| 126    | 3',6'-O-Di-coumaroyl-1',6'-O-di-feruloylsucrose | I   | H   | H   | H   | G   | G   | H   | I                                       | Roots          | [75, 76]                    |
| 127    | 3'-O-Coumaroyl-1',6'-O-tri-feruloylsucrose | I   | H   | H   | H   | I   | G   | H   | I                                       | Roots          | [75, 76]                    |
| 128    | 1',3',6'-6-O-Tetra-feruloylsucrose | I   | H   | H   | H   | I   | I   | H   | I                                       | Roots          | [75, 76]                    |

See Scheme 2.
antidepressant, cytotoxic, antineoplastic, anti-inflammatory, antidiabetic, plant growth-regulatory, neuroprotective, and cerebral protective activities. Lignan-derived disaccharide esters, phenylpropanoid-derived tetrasaccharide esters, and pentasaccharide esters with biological activities have not been reported. Aside from the isolated constituents, oligosaccharide esters with biological activities have not been identified from medicinal plants by using different kinds of assay methods, which include DPPH radical scavenging assay, hydroxyl radical scavenging assay, superoxide anion scavenging assay, and ABTS radical scavenging method [96].

3.1. Antioxidant Activity. The adverse effects of oxidative stress proposed to play significant roles in the pathogenesis of cardiovascular diseases, atherosclerosis, hypertension, cancer, diabetes mellitus, neurodegenerative diseases, rheumatoid arthritis, ischemia/reperfusion injury, and ageing have become an inevitable and serious issue [95, 96]. Scientists have thus made great efforts to explore antioxidants from medicinal plants by using different kinds of assay methods, which include DPPH radical scavenging assay, hydroxyl radical scavenging assay, superoxide anion scavenging assay, and ABTS radical scavenging method [96].

Lapathosides C and D, hydropiperoside, vanicoside B, hidropiperosides A and B, lapathoside A, and diboside A were isolated from the Polygonum, Persicaria, and Fagopyrum genera belonging to the Polygonaceae family. The DPPH test revealed that free radical-scavenging activity of the isolated compounds termed lapathoside C (115), hydropiperoside (42), vanicoside B (111), and lapathoside D (38) increased in turn, and lapathoside D exhibited strongest scavenging ability with an IC₅₀ of 0.088 μM [43]. Hidropiperosides A and B (117, 118) were reported to show obvious antioxidant response to DPPH radicals with the SC₅₀ values of 23.4 and 26.7 μg/mL, respectively, while vanicoside E moderately exhibited the same activity with a SC₅₀ value of 49.0 μg/mL [72]. Lapathoside A (114) and diboside A (116) just showed lower antioxidant activities with the SC₅₀ values of 199.48 and 165.52 μM, respectively [73].

Smiglasides A and B, smilaside P, 2,6-di-acetyl-3’,6’-di-feruloylsucrose, helonioside B, smilasides G–L, and heterosmilaside were isolated from the Heterosmilax and Smilax genera. Compared with ascorbic acid (IC₅₀ 143.52 μM) used as positive control, smiglasides A and B, and smilaside P (48, 49, 65) (IC₅₀ 339.58, 330.66 and 314.49 μM, resp.) showed higher antioxidant activities than 2,6-di-acetyl-3’,6’-di-feruloylsucrose (73) and helonioside B (40) (IC₅₀ 631.66 and 518.27 μM, resp.) [47]. Additionally, Nhiem et al. reported that helonioside B, heterosmilaside (104), and 2,6-di-acetyl-3’,6’-di-feruloylsucrose exhibited important DPPH radical scavenging activities with the SC₅₀ values of 9.1, 12.7 and 8.7 μg/mL, respectively [48]. Compared with smilasides G–I (57–59) (ED₅₀ 68.5–79.4 μM), smilasides J–L (60–62) showed higher radical scavenging activities with an ED₅₀ value of 26.7–32.7 μM [44].

Five quikesetinerviuisides A–E (105–109) isolated from the Calamus genus showed weak DPPH scavenging activities (IC₅₀ 60.4–101.8 μM) but exhibited better hydroxyl radical scavenging activities (IC₅₀ 3.6–8.4 μM). Moreover, quikesetinerviuiside C showed superoxide anion scavenging activity with an IC₅₀ value of about 184.3 μM [70]. Liu et al. investigated the antioxidant capacity of 3’,6-O-di-sinapoylsucrose (DISS) (123) by using the accelerated senescence-prone, short-lived mice (SAMP) in vivo. The analyses indicated that the activities of antioxidant enzymes of SOD and glutathione peroxidase ascended obviously in SAMP mice when amended with DISS 50 mg/kg. Moreover, DISS could downregulate and even restore the level of malondialdehyde in SAMP model group [97].

From the above studies, it can be concluded that oligosaccharide esters with antioxidant activities have been identified in the Polygonaceae, Lilaceae, Smilacaceae, and Aracaceae families. The results of the antioxidant assays show that the increased number of phenolic hydroxyl groups and acetyl groups could produce higher antioxidant activity. Fan et al. indicated that the increased number of phenylpropenoid groups was not beneficial to free radical scavenging activity [43]. Zhang et al. pointed out that oligosaccharide esters with feruloyl groups exhibited better antioxidant activities than those with coumaroyl groups [44].
Table 3: Fatty acid-derived disaccharide esters.

| Number | Name                                    | R₁  | R₂  | Source              | Parts       | Reference |
|--------|-----------------------------------------|-----|-----|---------------------|-------------|-----------|
| 132    | 6'-O-Linoleylsucrose                     | H   | O   | Astragalus membranaceus | Roots       | [77]      |
| 133    | 6'-O-Palmitoylsucrose                    | H   | P   | Astragalus membranaceus | Roots       | [77]      |
| 134    | 6-O-Palmitoylsucrose                     | P   | H   | Astragalus membranaceus | Roots       | [77]      |
| 135    | 6'-O-Linolenoylsucrose                   | H   | N   | Astragalus membranaceus | Aerial parts | [77, 78] |
| 136    | 6-O-Linoleylsucrose                      | O   | H   | Astragalus membranaceus | Roots       | [77]      |
| 137    | 6-O-Myristoylsucrose                     | Q   | H   | Astragalus membranaceus | Roots       | [77]      |
| 138    | 6-O-[(7Z,10Z,13Z)-Hexadeca-7,10,13-trienoyl]sucrose | H | R | Equisetum hiemale | Aerial parts | [78]      |
| 139    | 6-O-[(7Z,10Z)-Hexadeca-7,10-dienoyl]sucrose | H  | S  | Equisetum hiemale | Aerial parts | [78]      |

See Scheme 3.

3.2. Antidepressant Activity. The oligosaccharides obtained from the Morinda genus not only show specific antidepressant and antistress activities but also have no suppression or excitatory effects on central nervous system as well. What is more, they can be taken orally with little toxicity [21]. The inulin-type hexasaccharide (IHS) (9) from Morinda officinalis obviously exhibited cytoprotective activity, which contributed to the antidepressant effect, not only by providing the PC12 with protection against Cort-induced lesion with IHS 0.625 and 1.25 \( \mu \)M, but also by reducing the Cort-induced \( [Ca^{2+}]_o \) overloading with IHS 2.5 and 10 \( \mu \)M. IHS 5 and 10 \( \mu \)M upregulated the nerve growth factor mRNA expression in Cort-induced PC12 cells [22]. Polygalatenosides A (181) and B (182) were isolated from the Polygala genus. They significantly inhibited the isotope-labeled RTI-55 binding to norepinephrine transporter protein with the IC\(_{50}\) values of 30.0 and 6.04 \( \mu \)M, respectively [91].

DISS and tenuifoliside A were isolated from the Polygala and Cynanchum genera. Liu et al. investigated the antidepressant effect of YZ ethanol extract based on the tail suspension test (TST) and forced swimming test (FST), which are the ease-of-use and widely-accepted models for estimating antidepressant activities in mice. The results indicated that YZ-50 fraction at a dose of 200 mg/kg was able to significantly decrease the immobility time in TST. Furthermore, YZ-50 possessed ability to inhibit corticosterone-induced injury of human neuroblastoma SH-SY5Y cells. What is more, DISS (123) and tenuifoliside A (44), two major compounds of YZ-50 fraction, showed effective protective response to the lesion in SY5Y cells [53]. The antidepressant-like effect of DISS at
Figure 4

Table 4: Phenylpropanoid-derived trisaccharide esters.

| Number | Name        | R₁ | R₂ | R₃ | R₄ | R₅ | R₆ | R₇ | Source            | Parts  | Reference |
|--------|-------------|----|----|----|----|----|----|----|-------------------|--------|-----------|
| 146    | Tricornose C| K  | H  | H  | H  | H  | L  | H  | Polygala tricornis| Roots  | [37]      |
| 147    | Tricornose D| K  | H  | H  | H  | L  | H  | H  | Polygala tricornis| Roots  | [37]      |
| 148    | Tricornose E| K  | H  | H  | H  | K  | L  | H  | Polygala tricornis| Roots  | [37]      |
| 149    | Tricornose F| K  | H  | H  | I  | L  | H  | H  | Polygala tricornis| Roots  | [37]      |

See Scheme 4.

Table 4: Phenylpropanoid-derived trisaccharide esters.

3.3. Cytotoxic and Antineoplastic Activities. Smilasides A–F and P, smiglasides A and B, smilaside P, and helonioside A were isolated from the *Smilax*, *Trillium*, and *Paris* genera. Kuo et al. obtained smilasides A–F ([51–56]) and evaluated their cytotoxicity against human tumor cell lines comprising human oral epithelium carcinoma (KB), human cervical carcinoma (Hela), human colon tumor (DLD-1), human breast adenocarcinoma (MCF-7), human lung carcinoma (A-549), and human medulloblastoma (Med) cells by MTT assay. Experimental data indicated that all but smilaside C showed cytotoxicity against three to six human tumor cell lines (ED₅₀ = 5.1–13.0 μg/mL), and smilasides D–F (ED₅₀ = 2.7–5.0 μg/mL) displayed strong cytotoxic activities against DLD-1 cells [46]. Wang et al. reported the antitumor constituents of

the doses of 5, 10, and 20 mg/kg was also tested in chronically mild stressed rats. DISS was able to exhibit antidepressant activity by upregulating the expression of noradrenergic-regulated plasticity genes including cell adhesion molecule L1, brain-derived neurotrophic factor, laminin, and cAMP response element binding protein factor in hippocampus [98]. DISS improved the reward reaction by increasing sucrose intake and obviously decreased the levels of serum cortisol, adrenocorticotropic hormone, and corticotropin-releasing factor. Further, DISS played an enhanced role in the expression of mineralocorticoid receptor, together with glucocorticoid receptor mRNA [99].
Table 5: Phenylpropanoid-derived tetrasaccharide esters.

| Number | Name        | R₁ | R₂ | R₃ | Source             | Parts   | Reference |
|--------|-------------|----|----|----|--------------------|---------|-----------|
| 154    | Tricornose G | K  | H  | K  | Polygala tricornis | Roots   | [37]      |
| 155    | Tricornose H | K  | K  | K  | Polygala tricornis | Roots   | [37]      |
| 156    | Tricornose I | K  | K  | L  | Polygala tricornis | Roots   | [37]      |
| 157    | Tricornose J | K  | I  | L  | Polygala tricornis | Roots   | [37]      |
| 158    | Tricornose K | K  | I  | K  | Polygala tricornis | Roots   | [37]      |
| 159    | Tricornose L | K  | G  | K  | Polygala tricornis | Roots   | [37]      |

See Scheme 5.

**Smilax riparia**, including smiglasides A (48) and B (49), 2,6-di-acetyl-3',6'-di-feruloylsucrose (73), helonioside B (40), and smilaside P (65). Only smiglasides A and B, and smilaside P exhibited cytotoxicity against human tumor cell lines with different inhibitory concentrations comparing with cisplatin and paclitaxel as positive controls [47]. Helonioside A (39) exhibited higher cytotoxicity with the increase of concentration (0.1-100 μg/mL) [45]. Tatarisides A–G (45, 119–121, 46, 122, 47) and diboside A (116) from the *Fagopyrum* genus exerted cytotoxic activities against different human cell lines, and the cytotoxicity of tatariside C was the most remarkable with the IC₅₀ values ranging from 6.44 to 7.49 μg/mL [57].

1',2,3,6-O-Tetra-acetyl-3'-O-cis-feruloylsucrose (95) from the *Sparganium* plants exhibited extremely weak cytotoxicity against the growth of mice Lung Adenocarcinoma 795 cell lines with an IC₅₀ value of 116 μg/mL [66]. SnS-2, oligosaccharides mixture, including raffinose (3), stachyose (19), and verbascose (21) from the roots of *Scrophularia ningpoensis*, had antitumor activity against the growth of Lewis pulmonary carcinoma cells transplanted into mice [18].

Disaccharide esters and oligosaccharides mixture from the Liliaceae, Polygonaceae, Sparganiaceae, and Scrophulariaceae families showed effective cytotoxic and antineoplastic activities. The study results indicated that feruloyl and acetyl groups play an important role in mediating cytotoxicity, which seems to be related to the substitution position of feruloyl groups. The feruloyl groups at C-6 or C-1' are vital for cytotoxicity. In addition, the increased number of acetyl groups could induce higher tumoricidal activity.
### Table 6: Phenylpropanoid-derived pentasaccharide esters.

| Number | Name         | \(R_1\) | \(R_2\) | \(R_3\) | \(R_4\) | \(R_5\) | \(R_6\) | Source          | Parts   | Reference |
|--------|--------------|---------|---------|---------|---------|---------|---------|-----------------|---------|-----------|
| 160    | Tenuifoliose A | G       | D       | I       | Ac      | Ac      | Ac      | Polygala tenuifolia | Roots   | [38, 42]  |
| 161    | Tenuifoliose B | G       | D       | I       | H       | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 162    | Tenuifoliose C | G       | D       | I       | H       | H       | H       | Polygala tenuifolia | Roots   | [42]      |
| 163    | Tenuifoliose D | G       | D       | U       | Ac      | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 164    | Tenuifoliose E | G       | D       | U       | Ac      | H       | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 165    | Tenuifoliose F | G       | D       | G       | Ac      | Ac      | Ac      | Polygala tenuifolia | Roots   | [38, 42, 79]|
| 166    | Tenuifoliose G | G       | D       | G       | Ac      | H       | Ac      | Polygala tenuifolia | Roots   | [42, 79]  |
| 167    | Tenuifoliose H | G       | D       | G       | H       | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 168    | Tenuifoliose I | G       | D       | G       | H       | H       | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 169    | Tenuifoliose J | G       | D       | T       | Ac      | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 170    | Tenuifoliose K | G       | D       | T       | Ac      | H       | Ac      | Polygala tenuifolia | Roots   | [42, 79]  |
| 171    | Tenuifoliose L | I       | D       | I       | Ac      | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 172    | Tenuifoliose M | I       | D       | I       | H       | Ac      | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 173    | Tenuifoliose N | I       | D       | I       | H       | H       | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 174    | Tenuifoliose O | I       | D       | T       | Ac      | H       | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 175    | Tenuifoliose P | I       | D       | T       | H       | Ac      | Ac      | Polygala tenuifolia | Roots   | [79]      |
| 176    | Tenuifoliose Q | G       | D       | G       | H       | H       | H       | Polygala tenuifolia | Roots   | [42]      |
| 177    | Tenuifoliose R | G       | D       | I       | H       | H       | H       | Polygala tenuifolia | Roots   | [42]      |
| 178    | Tenuifoliose S | G       | D       | U       | Ac      | H       | Ac      | Polygala tenuifolia | Roots   | [42]      |
| 179    | Tenuifoliose T | I       | D       | I       | H       | H       | H       | Polygala tenuifolia | Roots   | [42]      |

See Scheme 6.

3.4. Anti-Inflammatory Activity. Inflammation, an important basic pathological process, is a defense response of biopsy with vascular system to damage stimuli such as pathogens, impaired cells and tissues, and physical and chemical factors. However, if the process of inflammatory response cannot end normally when cell debris and pathogens were cleared, the biological defence response will become causative factor and bring about many diseases, such as diabetes, cardiovascular diseases, metabolic syndrome, and cancer [100, 101].

Tenuifoliside A (44) from the *Polygala* genus exhibited strong anti-inflammatory effect not only by suppressing the production of NO, but also by reducing the production of iNOS, prostaglandin E2, cyclooxygenase-2, and proinflammatory cytokines through the inhibition of the mitogen-activated protein kinases pathway and NF-\( \kappa \)-B pathway [102].

The anti-inflammatory activities of quiquesetinerviusides D (108) and E (109) from the *Calamus* genus were evaluated in RAW 264.7 cells. Both of them showed significant inhibitory effects against the production of LPS-stimulated NO with the IC\(_{50}\) values of 9.0–29.5 \(\mu\)M [70].

Six disaccharide fatty acid esters (132–137) were isolated from the *Astragalus* and *Equisetum* genera. The anti-inflammatory effects of these isolated compounds have also been documented. The activation of NF-\( \kappa \)-B could upregulate the expression of proinflammatory cytokines inducible nitric oxide synthase (iNOS) and tumor necrosis factor alpha (TNF-\( \alpha \)). The NF-\( \kappa \)-B inhibitory activities of compounds 132–137 were tested in HepG2 cells stimulated with TNF-\( \alpha \). All of these compounds could significantly restrain TNF-\( \alpha \)-induced NF-\( \kappa \)-B transcriptional activities with the IC\(_{50}\) values of 4.4–24.7 \(\mu\)M. Li et al. pointed out that olefinic bonds and the length of the fatty acid moiety contributed to the NF-\( \kappa \)-B inhibitory activity. Furthermore, the inhibition increased significantly with the increase of the number of olefinic bonds on the aliphatic moiety [77]. These results may provide a scientific basis for the development of new anti-inflammatory agents.

3.5. Neuroprotective and Cerebral Protective Activities. As we all know, glutamate works as a major excitatory amino acid neurotransmitter in the mammalian central nervous system.
3.6. Antidiabetic Activity. Diabetes mellitus, a chronic debilitating metabolic disease, is characterized by high blood glucose content and comprises three types termed type I, type II, and gestational diabetes [106]. Stachyose (19) extract (a part) from *Rehmannia glutinosa* obviously exhibited the activity of downregulating fasting plasma glucose level and partially keeping from hyperglycemia induced by adrenaline and glucose without obvious dose-dependent effect. Other than that, *in vivo* tests in rats induced by alloxan revealed that stachyose extract at the dose of 200 mg/kg significantly decreased blood-sugar level [15].

Diboside A, lapathosides C and D, vanicosides A and B, and hydropiperoside were isolated from the *Fagopyrum, Polygonum*, and *Persicaria* genera belonging to the Polygonaceae family. Diboside A (116) could potentially inhibit α-amylase activity with an IC$_{50}$ of 26.9 μM and thus retard the starch digestion rate, which is helpful for diabetic individuals in controlling blood sugar level [107]. Lapathoside D (38) exerted stronger activity of α-glucosidase inhibition with an IC$_{50}$ value of 0.113 mM than acarbose which was chosen as a positive drug for the treatment of type II diabetes [43]. Vanicoside B (111) was reported to have higher β-glucosidase inhibitory activity with an IC$_{50}$ of 59.9 μM because of the acetyl moiety of the latter possibly decreasing inhibitory activity of vanicoside A [71].

Fujimoto et al. investigated the inhibitory effects of mumeoses F–O (83–92) from the *Prunus* genus on aldose reductase and discovered that caffeoyl groups are crucial for the inhibitory effect on aldose reductase. And thus, mumeoses F, G, H, J, K, L, M and N (IC$_{50}$ = 22–77 μM), with a coumaroyl group and acetyl groups, inhibited moderately aldose reductase from reducing glucose to sorbitol, which is associated with the chronic complications of diabetes [63, 64].

3.7. Elicitors and Regulators. Oligosaccharides are quite propitious for encoding biological information because of diverse monosaccharide units and complex molecular structures and they are therefore first described as biological system and plays a crucial role in several physiological processes [103]. However, the accumulation of glutamate induces diverse acute and chronic neurodegenerative diseases, such as epilepsy, ischemic stroke, and Parkinson’s disease, as well as Alzheimer’s disease [104]. DISS (123) isolated from the *Polygala* genus exhibited neuroprotective effect against glutamate-induced SH-SY5Y neuronal cell damage. The in vitro test demonstrated that DISS (0.6, 6 and 60 μmol/L) played a critical role in increasing cell viability, controlling lactate dehydrogenase and attenuated apoptosis ranging from 1.95% to 2.58% [105].

Tenuifoliside B (102) from the *Polygala* genus was able to significantly shorten the coma time of KCN-induced anoxia mice at the doses of 3 and 10 mg/kg, and it played an important role in ameliorating the scopolamine-induced impairment of performance in passive avoidance task in rats and enhancing the tremors induced by oxotremorine in mice. These results together demonstrated that tenuifoliside B possessed cognitive improving and cerebral protective effects [56].
signals in plants [108]. Oligosaccharides from the cell wall fragments of plants and fungi are powerful signal molecules, such as the elicitors of plant defence response and the regulators of plant growth, and they are capable of exerting biological activities at exceedingly low concentrations [109]. Heptasaccharide (HS) (15) and octasaccharide (OS) (16) isolated from the Paris genus possessed plant growth-regulatory activities [29, 30]. The two oligosaccharides significantly promoted the proliferation of Paris polyphylla var. yunnanensis roots at the doses of 2.5–20 mg/L. The octasaccharide had the most obvious effect on the growth of Panax japonicus var. major hairy roots at a dose of 30 mg/L, while the other had the most positive effect on saponin accumulation of Panax japonicus var. major hairy roots at a dose of 10 mg/L.
Similarly, Zhou et al. evaluated the stimulating effects of HS and OS on the root growth and saponin production of Panax ginseng hairy roots, which were induced from the plant roots infected with Agrobacterium rhizogenes strain A4. The results showed that there was a maximum effect on the hairy roots growth and saponin accumulation on day 10. Compared with control group, the root biomass dry weight was increased by more than 1.7-fold while the total saponin content of roots increased by more than 1-fold when these two oligosaccharides were added to the hairy root at a dose of 30 mg/L [30]. The above data illustrate that HS and OS could serve as the plant growth-regulators not only in their original species but also in others.

3.8. Immunopotentiating Activity. Macrophages are important targets of investigations on cytaphagy, cellular immunity, and molecular immunology. Therefore, they are deemed to play a vital role in host defense comprising phagocytosis, proteolytic processing, pathogenic agent, apoptosis, cytokines production, and foreign antigens presentation [110]. The water-extracted oligosaccharides from Panax ginseng (WGOS) exhibited better immunopotentiating activity by increasing phagocytic function of macrophages and promoting NO, TNF-α and reactive oxygen species production [110]. In addition, Wan et al. have obtained maltoligosaccharides (17, n = 3–8) and three oligosaccharides (2, 13, 14) from the Panax ginseng roots. The in vitro bioassay pointed out that WGOS could serve as efficacious stimulators of B and T lymphocytes [28]. These studies provided enlightenment that the mixture of oligosaccharides from Chinese herbal medicine exhibits significant effect on immune system.

3.9. Others. Acetylcholinesterase (AChE) inhibitors show good therapeutic effects on myasthenia gravis, glaucoma, and Alzheimer's disease through reversible enzyme inhibition so as to increase the accumulation of acetylcholine in the synapse and then promote and prolong the function of acetylcholine. Vanicoside B (III) showed AChE inhibitory activity with an IC₅₀ of 0.062 mM, while hydropiperoside (42), and lapathosides C (115) and D (38) just exhibited weak enzyme inhibitory activity [43].

Wang et al. has explored low molecular mass carbohydrate polymer from Panax ginseng roots and obtained 30% ethanol elution (PGO) which included peptides and oligosaccharides (17, n = 0–5) identified as maltose, maltotriose, maltotetraose, maltopentaose, maltohexaose, and maltoheptaose. Pharmacological experiments revealed that PGO could significantly enhance the memory in scopolamine-induced memory deficit rats [27].

Cistanoside F (129) and kankanose (150) were isolated from the Cistanche, Paulownia, and Acanthus genera. Pharmacological experiments showed that cistanoside F and kankanose significantly exhibited vasorelaxant effects on the noradrenaline-induced contraction of thoracic aorta from rats [80].

4. Conclusion

Traditional Chinese medicine from natural kingdom plays an indelible role in the treatment of human diseases, and it has aroused the attention of those who have engaged in medicinal pharmaceutical chemistry. Therefore, scientists have made great contributions day after day to investigate the valid chemicals from traditional Chinese medicines. In the past decades, about 193 oligosaccharides and their esters have been identified from traditional Chinese medicinal plants. On the one hand, only a few oligosaccharides and their mixtures were investigated and just exhibited antidepressant, antineoplastic, antidiabetic, plant growth-regulatory, immunopotentiating, and enhanced memory activities. More exploratory work is still needed to excavate biological and pharmacological activities of oligosaccharides. On the other hand, oligosaccharide esters exhibited multi-advantageous activities. Bioassays have revealed that antioxidant, cytotoxic, antineoplastic, and anti-inflammatory activities are the most notable bioactivities. Of course, to search for the natural products with these activities is a hotpot in the contemporary drug research. Oligosaccharide esters provide a vast treasure trove for medical researchers. After considering the current studies, it should be taken as future directions to make more mechanism of action studies and clinical trials to further evaluate its potential as new drugs. Moreover, the structure-activity relationships discussed in this review will provide reference information for further exploring their relationships and continually discovering the new bioactive oligosaccharide esters.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

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