From metagenomic gene discovery to enzymatic breakdown of crosslinks in plant cell wall biopolymers

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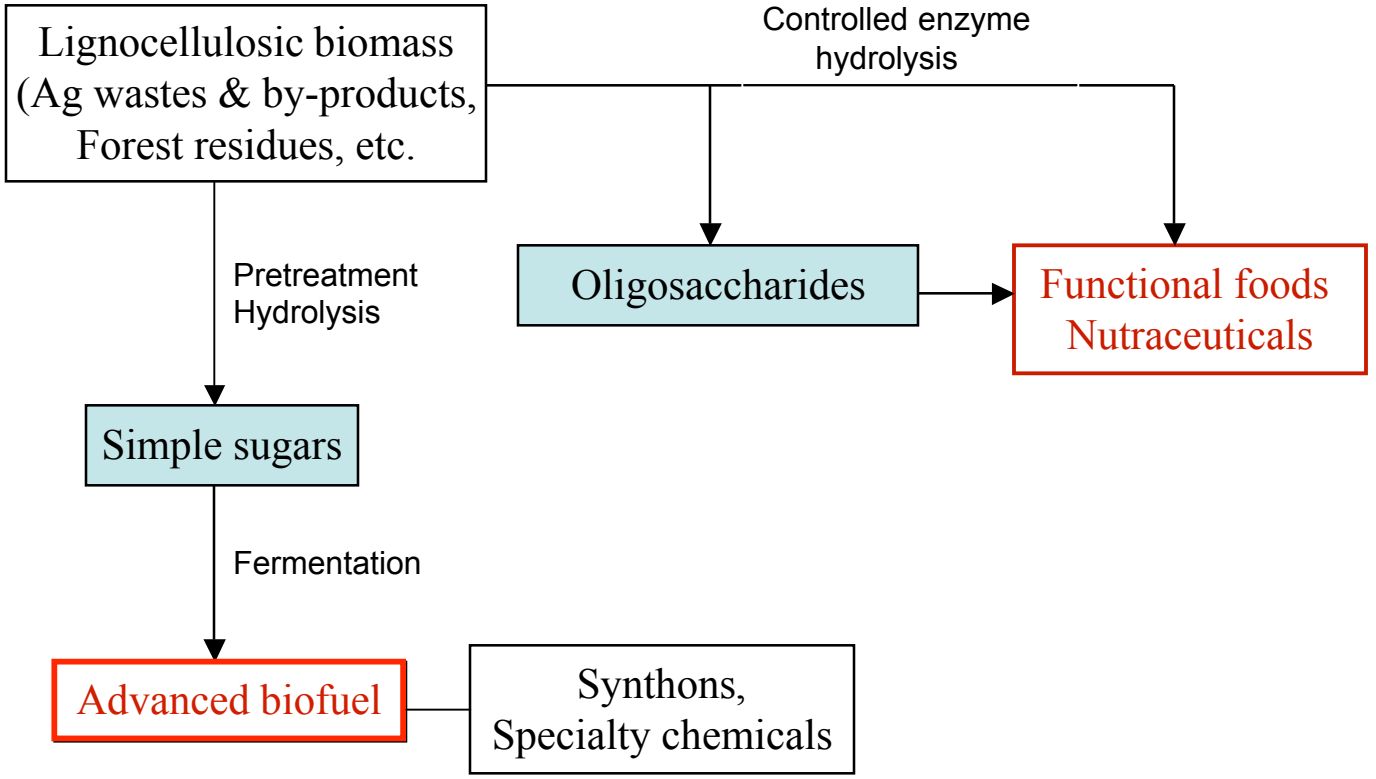
Research Objectives:

- *Research key enzymes and their cooperative mode of action on breaking down plant cell wall polymers.*

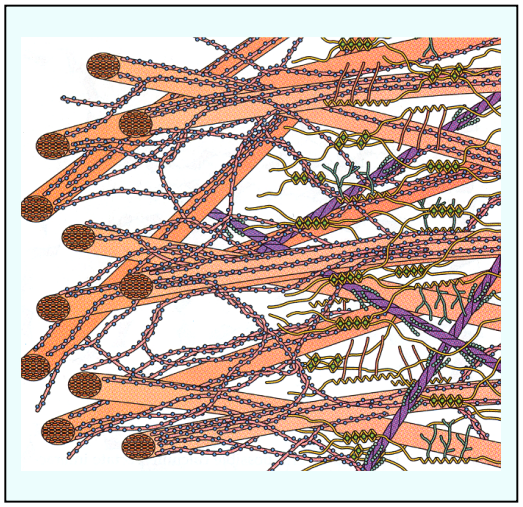
- *Research the efficacy of using hemicellulases and other non-cellulase enzymes prior to, during, or after pretreatment.*

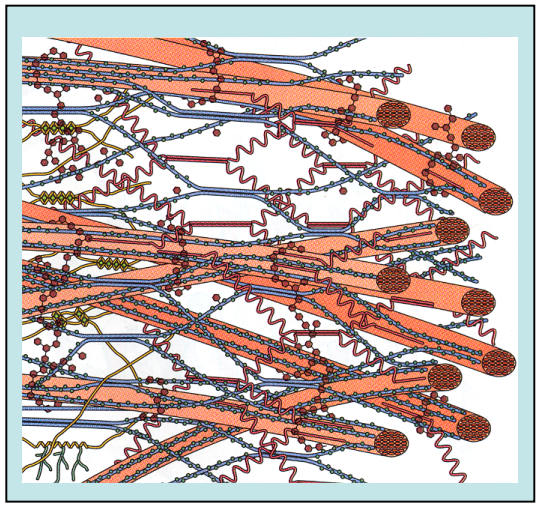
For conversion of agricultural wastes to biofuel and high-value products

Biomass conversion to high value products



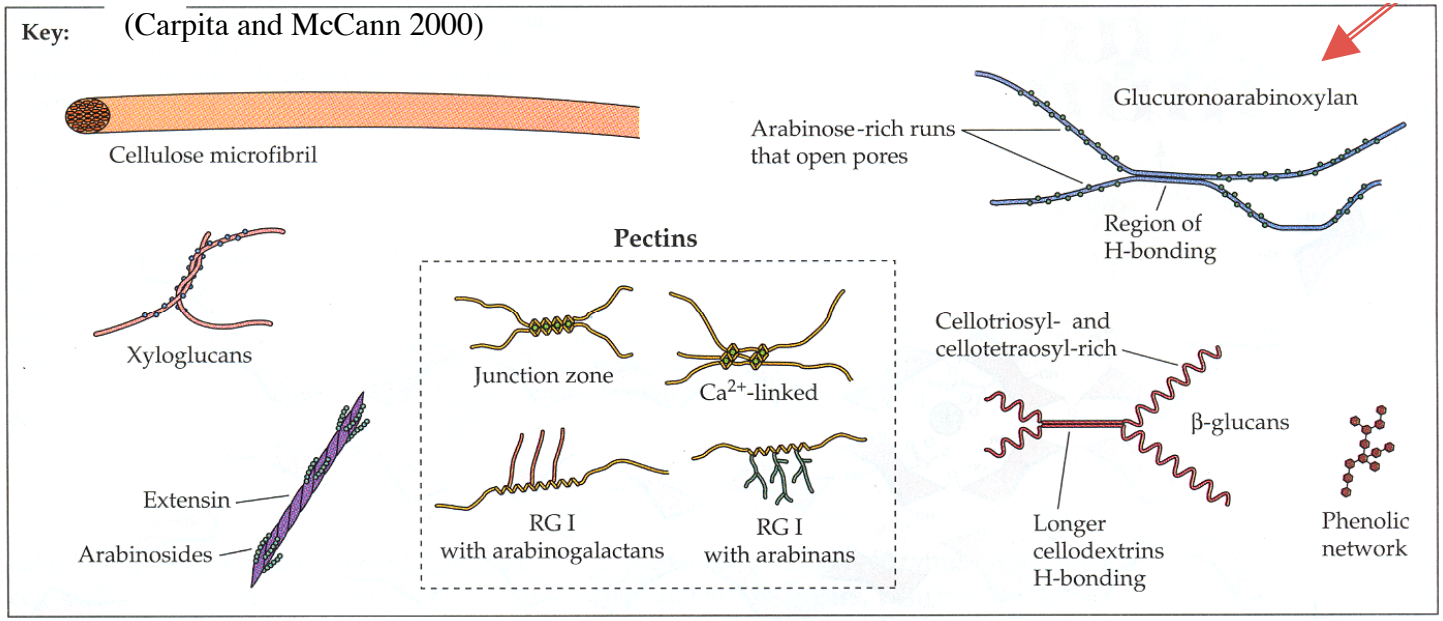
Cell wall polymers





Biomass source	Cellulose	Hemi-	Lignin
	(% dry weight)		
Agricultural residues	38	32	17
Corn fiber	15	35	8
Corn cob	45	35	15
Corn stover	40	25	17
Rice straw	35	25	12
Wheat straw	30	50	20
Sugarcane bagasse	40	24	25
Herbaceous energy crops	45	30	15
Switch grass	45	30	12
Coastal Bermuda grasss	25	35	6
Sorted municipal solid waste	45	9	10
Underutilized & short rotation hardwoods	50	23	22

(Saha 2003; Wyman 1994)



Type I -
All dicots & one-half of monocots (roses, sunflowers, maples, oaks beans, spinach, sugar beet)

Type II -
The "commelinoid" of monocots (palms, gingers, cypresses, and grasses- including grain crops)

Biomass Degradation - *Known enzyme groups found in nature*

Cellulose: (3)

endoglucanase, exoglucanase, cellobiosidase

Hemicellulose: (15):

Glucuronoarabinoxylan: (6) *endoxylanase, β-xylosidase, α-L-arabinoxylanosidase, α-glucuronisdase, acetylxylan esterase, feruloyl esterase*

Xyloglucan: (6) *endo-xyloglucanase, xyloglucosidase, β-glucosidase, α-xylosidase, β-galactosidase, fucosidase*

β-Glucan: (3) *β-glucan endohydrolases, β-glucan exohydrolases, β-glucosidase*

Pectins: (17)

Hydrolases: (5) *endopolygalacturonase, exopolygalacturonase, rhamnogalacturonan hydrolase, rhamnogalacturonan rhamnohydrolase, xylogalacturonanhydrolase*

Lyases: (4) *endopectate lyase, exopectate lyase, pectin lyase (endo), rhamnogalacturonan lyase*

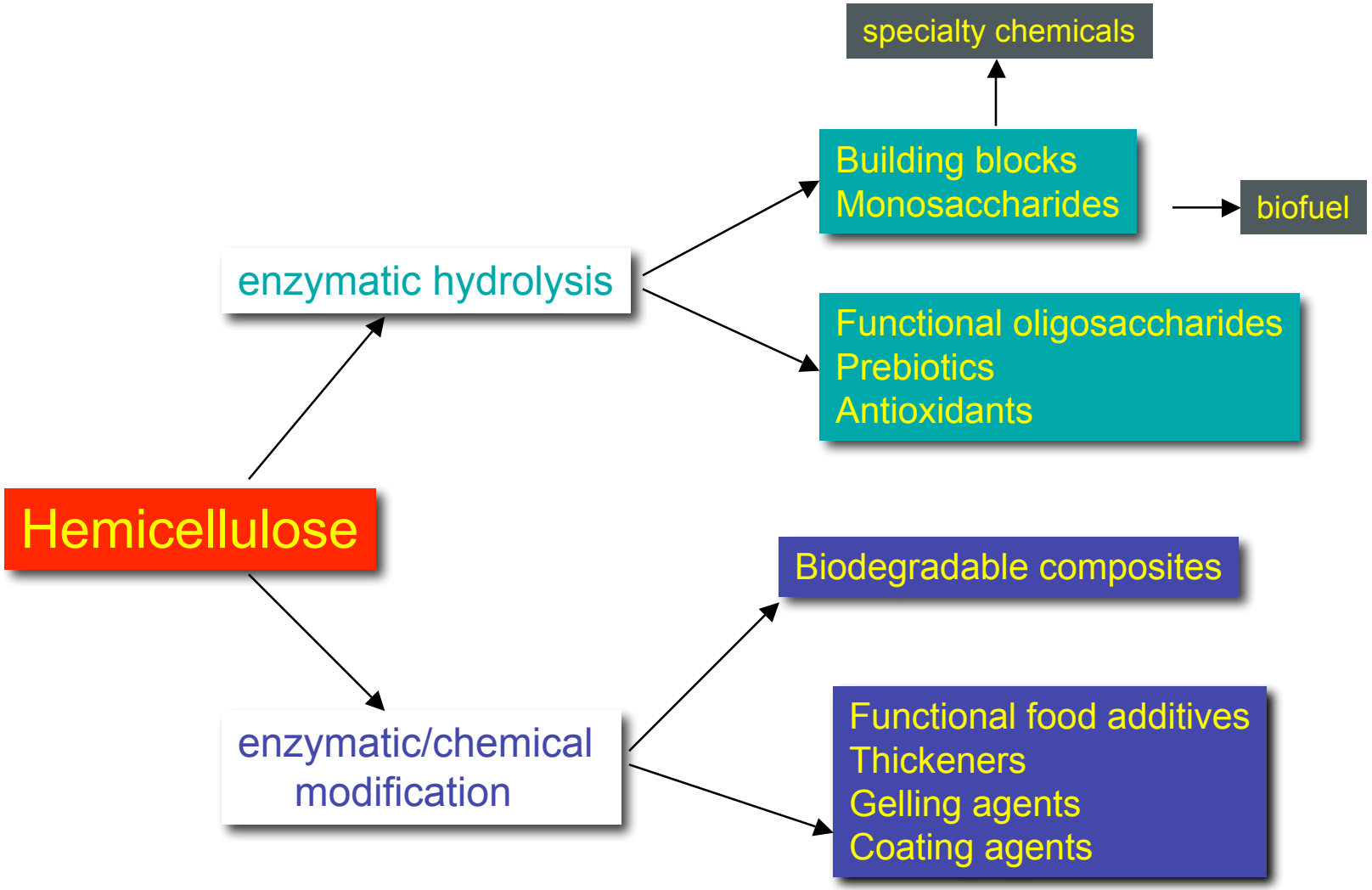
Esterases: (3) *pectin methyl esterase, pectin acetyl esterase, rhamnogalacturonan acetyl esterase*

Auxiliary enzymes: (5) *galactanase, arabinanase, β-galactosidase, α-L-furanosidase, feruloyl esterases*

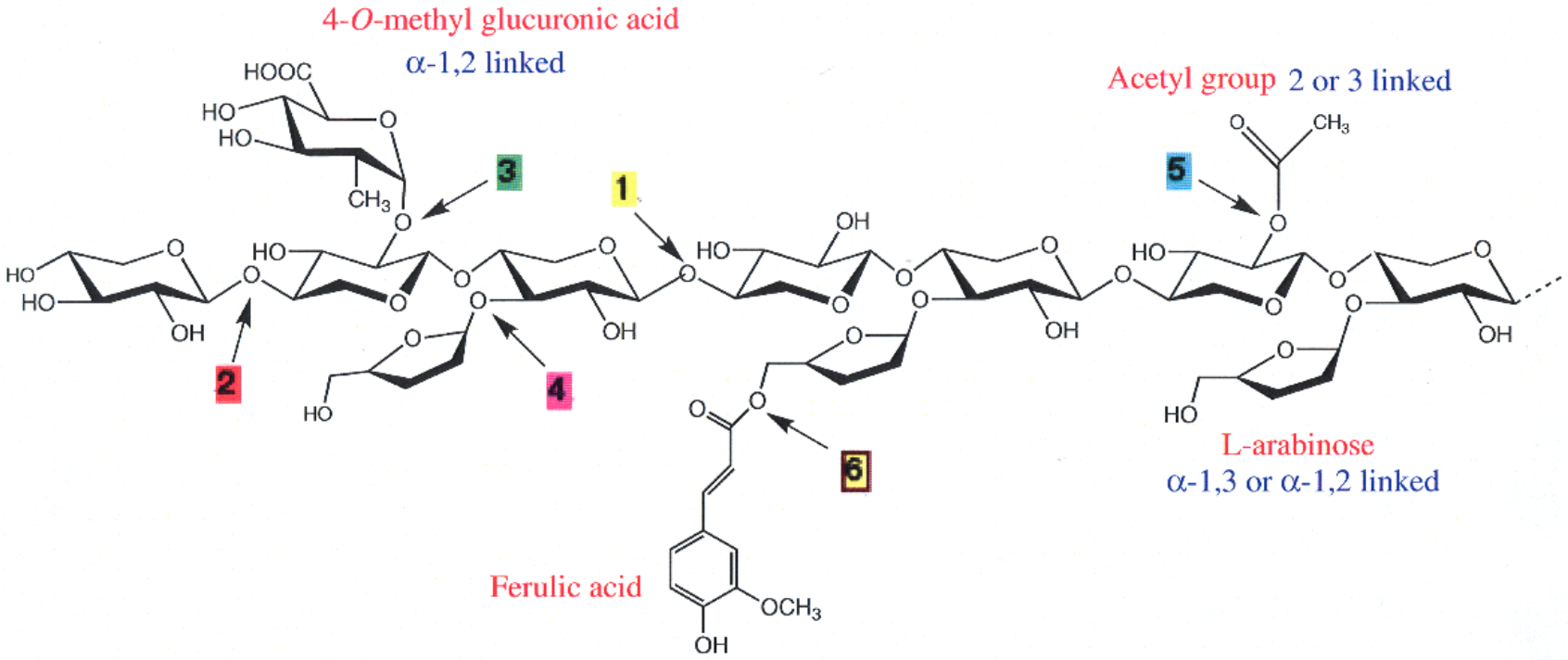
Lignin: (3) *lignin peroxidase, manganese-dependent peroxidase, laccase*

USDA ARS WRRC

Bioconversion of hemicellulose



Glucuronoarabinoxylan



1 endo-β-1,4-xylanase (EC3.2.1.8)

2 β-xylosidase (EC 3.2.1.37)

3 α-glucuronidase (EC 3.2.1.139)

4 α-L-arabinofuranosidase (EC 3.2.1.55)

5 acetylxylan esterase (EC 3.1.1.72)

6 feruloyl esterase (EC 3.1.1.79)

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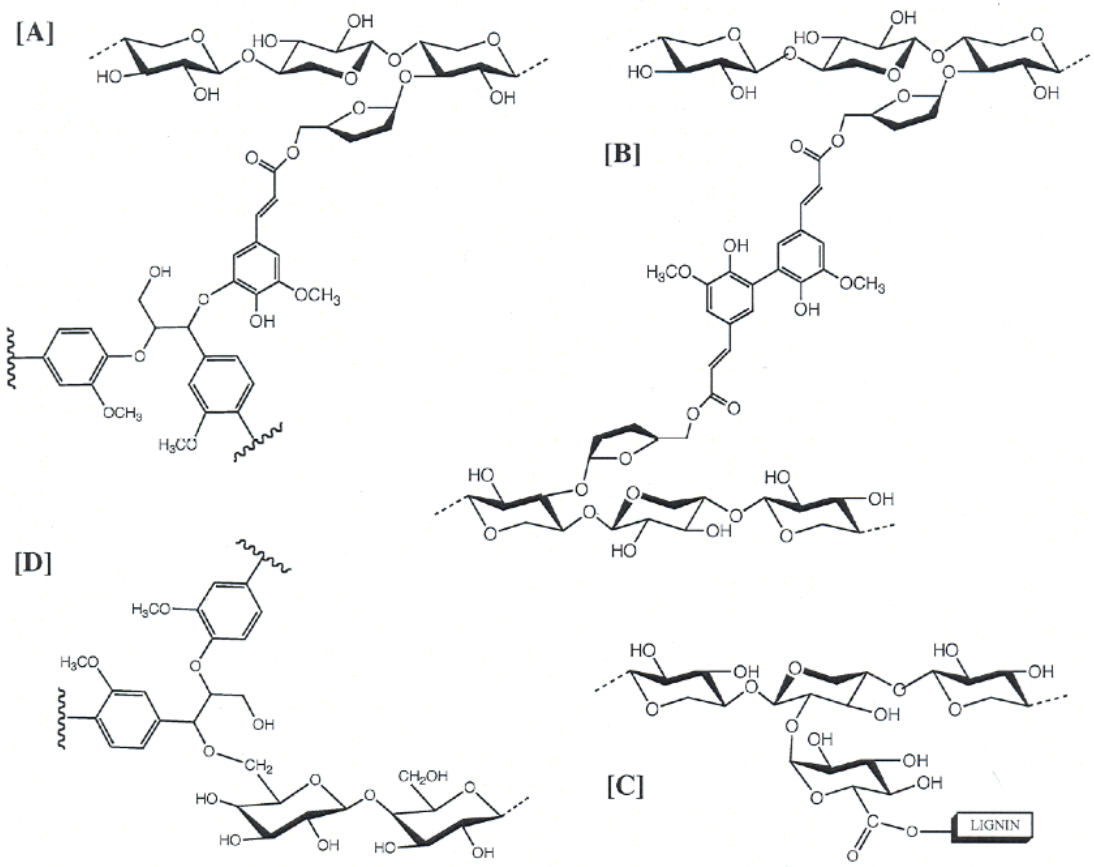
Chemical structures of crosslinks



(A) xylan-ferulic acid-lignin; (B) arabinoxylan-diferulic acid-arabinoxylan
(C) xylan-glucuronic acid-lignin; (D) pectin-galactan-lignin

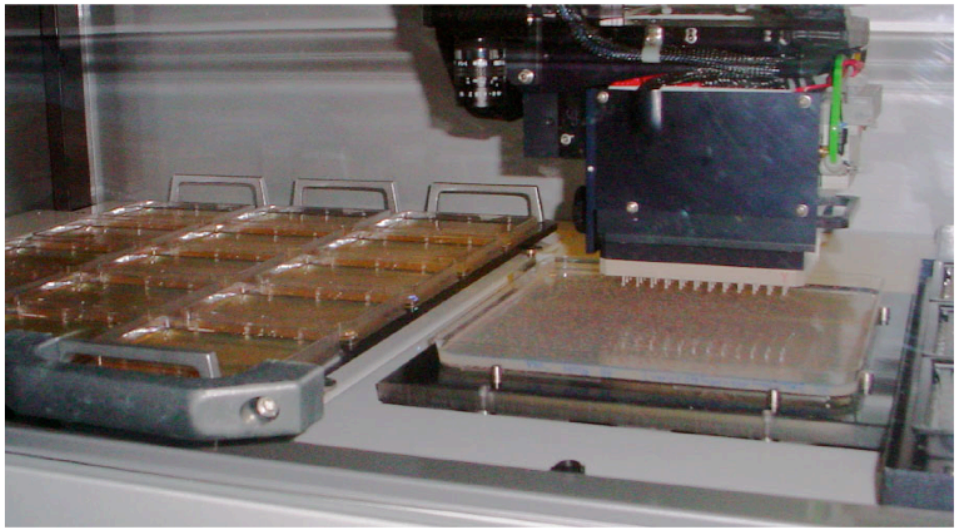
D. Wong (2006) (2009) Applied Biochem. Biotechnol D. Wong (2008) Protein Journal

Novel Genes from Metagenomes

"Metagenome" = the collective genomes of all microorganisms in a given habitat (Handerisman et al. 1998; Schmidt et al. 1991)

D. Wog (2010) *In*: Metagenomics: Theory, Methods, and Applications

Q-Pix



Q-Bot

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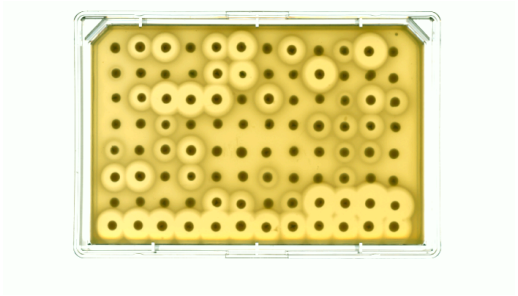
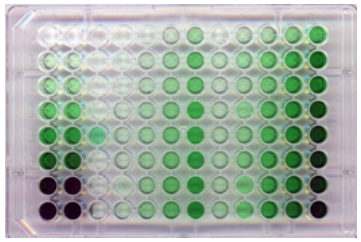
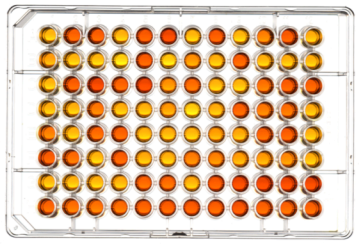
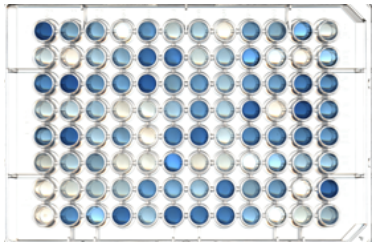
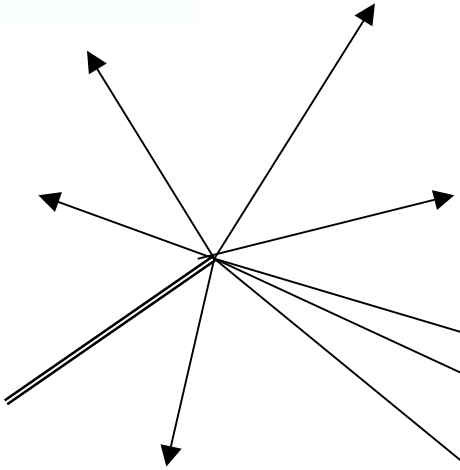
High-throughput screening active clones

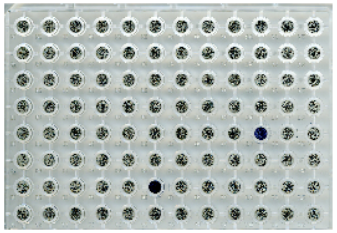


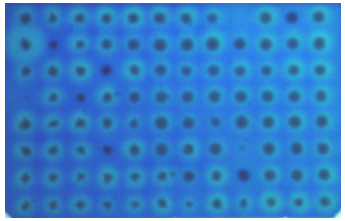


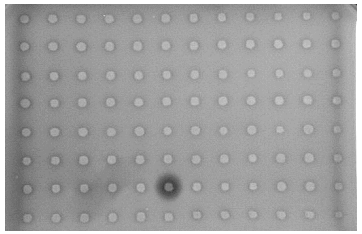


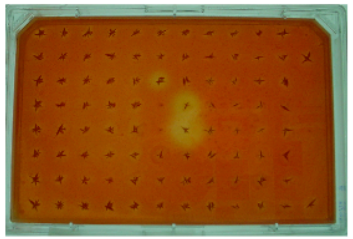








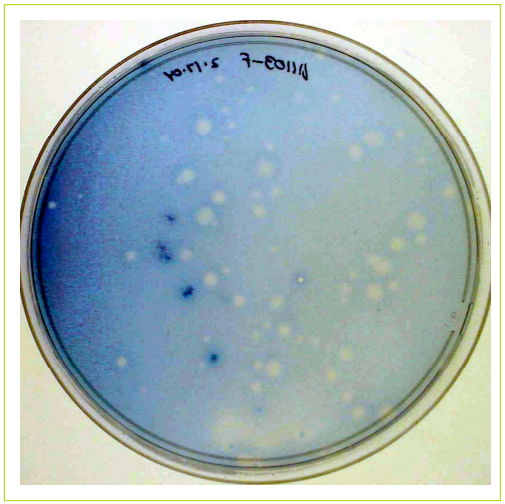




Q-tray

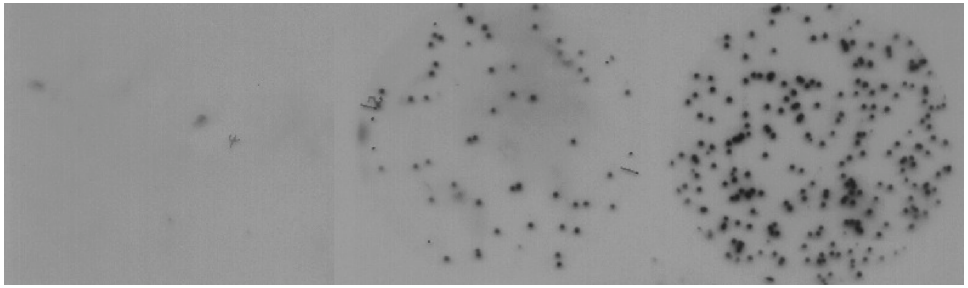
Wong et al. US Patent 8,361,764 B1

High-throughput screening active clones

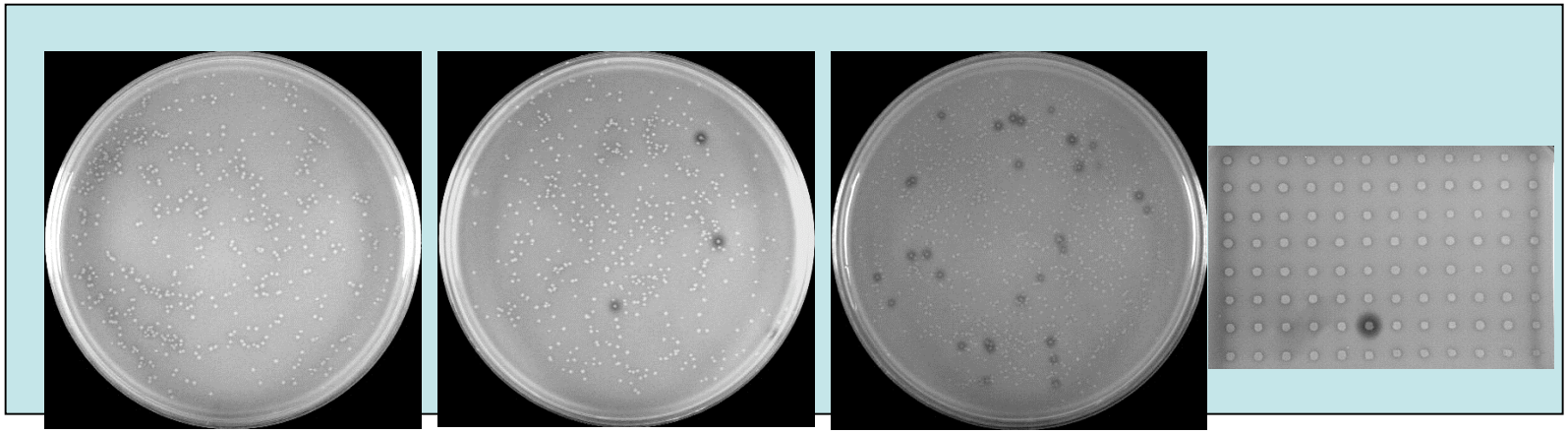


Screening for xylanase genes from metagenomic libraries

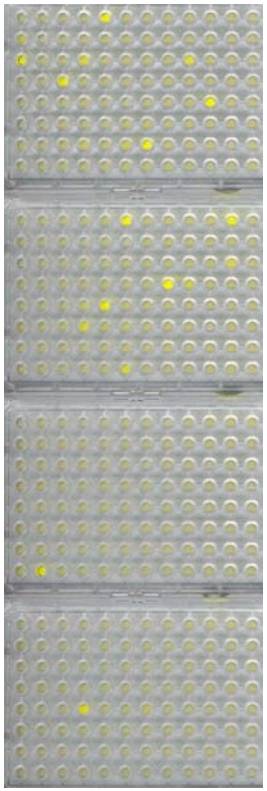
Screening for amylase genes from cDNA (λZapII) libraries



Screening for feruloyl esterase genes from metagenomic libraries

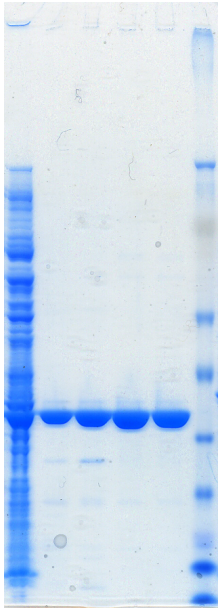


FAE gene isolation, cloning and expression



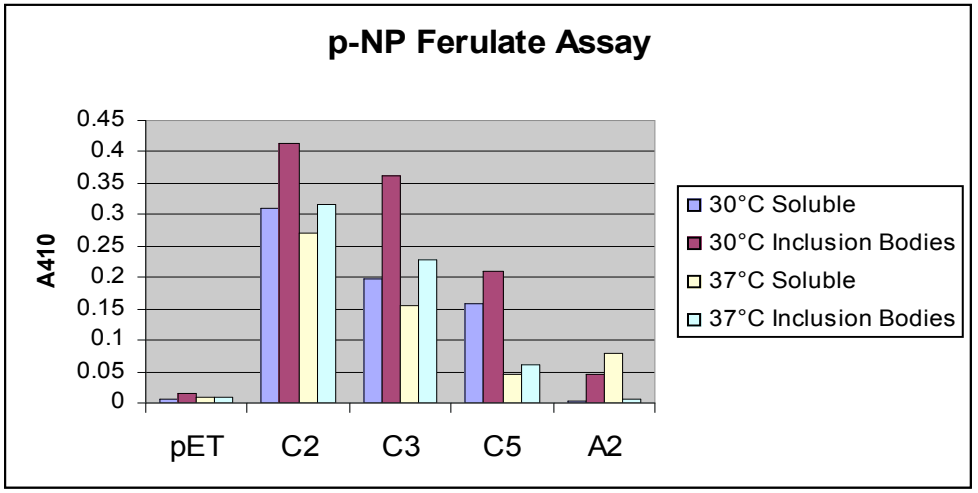
High-throughput screening

Rescreen positives on ethyl ferulate



pET vector/ *E. coli* BL21

Rescreen positives by liquid assay



Wong et al. (2013) US Patent 8,361,764 B1

FAE genes and proteins characterization

Clone Name	Gene Location (Insert Size)	Reference Sequence	% Identity to Ref Seq	Number of Residues (Includes tag)	SEQ ID NO	Recombinant Size (kDa)
Rum-C1	1811-2497 (687 bp)	ZP_01061020.1 Hydrolase of ¯ Family	45	229	1	25.2
Rum-C2	587-1396 (810 bp)	ZP_01061020.1 Hydrolase of ¯ Family	45	269	2	29.7
Rum-C3	663-1343 (681 bp)	ZP_01061020.1 Hydrolase of ¯ Family	46	381	3	41.9
Rum6-C4	112-831 (720 bp)	AAF70241 Feruloyl Esterase A	41	246	4	27.1
Rum-C5	60-1001 (942 bp)	ZP_01122099.1 probable lipase/esterase	32	416	5	45
Rum-C6	2871-3827 (957 bp)	YP_169760.1 hypothetical protein lipase/esterase	34	422	6	45.6
Rum-C7	447-1277 (831 bp)	ZP_01777732 putative esterase	48	277	7	32.3
Rum-C8	1849-3207 (1359 bp)	ZP_01061020.1 Hydrolase of ¯ Family	47	382	8	41.9
Rum-C9	66-1403 (1338 bp)	ZP_01061020.1 Hydrolase of ¯ Family	47	383	9	41.9
Mul-C10	181-1281 (1101 bp)	ZP_01421329 Beta-lactamase	44	367	10	39.7
Mul-C11	116-1057 (942 bp)	YP_583166 alpha/beta hydrolase	54	322	11	34.7
Mul-C12	1495-2412 (918 bp)	YP_578455 metallophosphoesterase	43	319	12	34.8

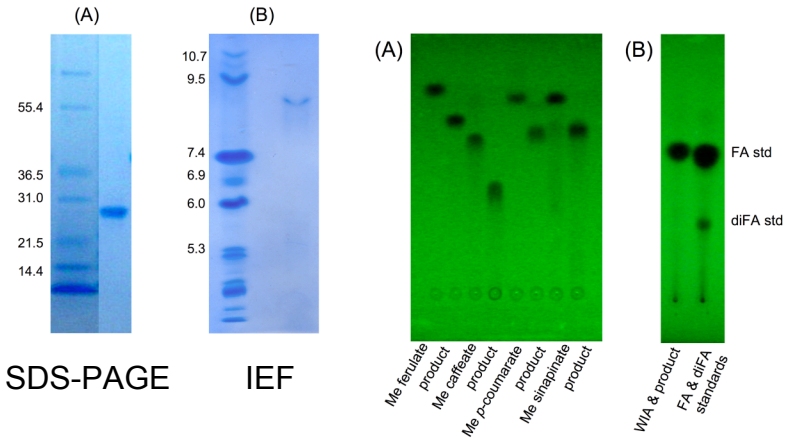
Genbank 15, WRRC 12

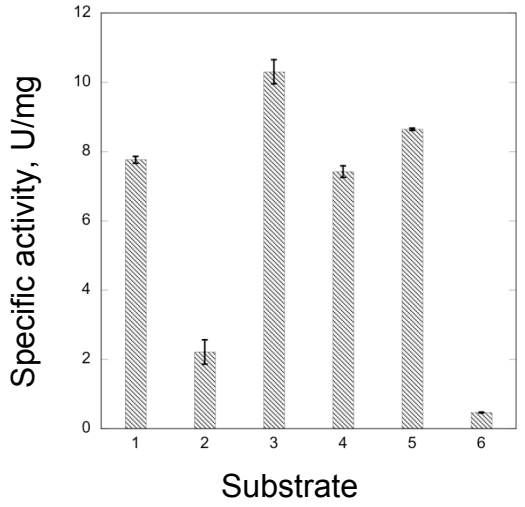
Wong et al. (2013) US Patent 8,361,764 B1

Novel Feruloyl Esterases - Phylogenetic Tree

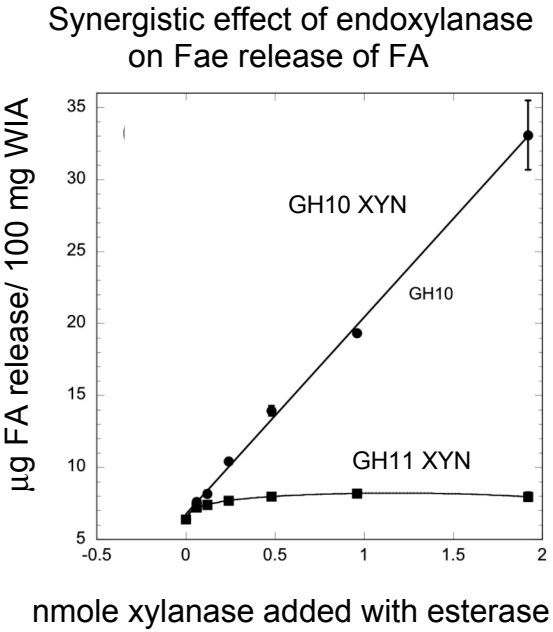
USDA ARS WRRC

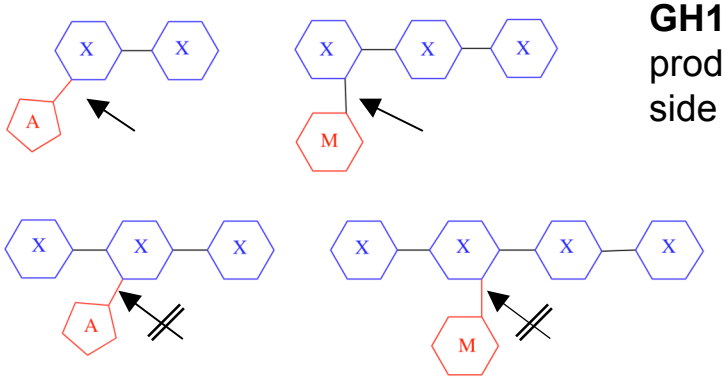
Fae clone 2 enzyme characterization





1. Corn fiber
2. Corn bran
3. Wheat bran
4. Wheat arabinoxylan
5. Rice bran
6. switchgrass





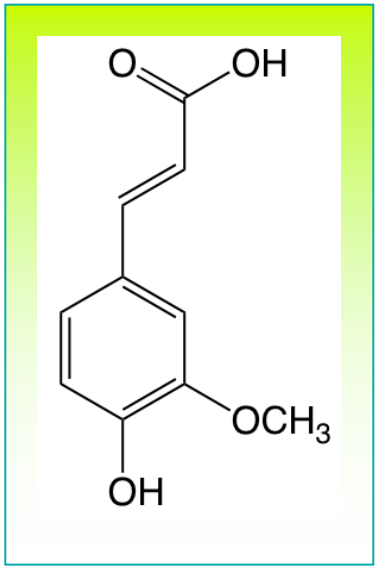
GH10 hydrolysis products with end-linked side groups

GH11 products : internal side groups

The Fae 2 enzyme can hydrolyze only or at a much higher rate araf-ferulic acid side groups linked to the non-reducing end.

Wong et al. (2013) J. Ind. Microbiol. Biotechnol.

Ferulic acid and derivatives



Properties:
- improve anti-oxidative activity
- increase lipid solubility
- enhance gelling effect

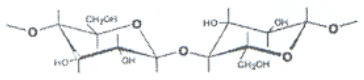
Applications:
- protect against photo-oxidative damage
- enhance wound management aid
- synthesis of vanillin
- produce thermal stable biogel

Release of ferulic acid:
- prevent filter blockage in brewing
- increase loaf volume in baking
- increase biomass degradation

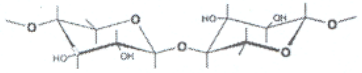
Barley: 0.14% dry w/w
Wheat bran: 0.66%
Sugar beet pulp: 0.8%
Corn bran: 3.1%
Corn fiber: 3-6%
Switchgrass: 0.3-0.5%

USDA ARS WRRC

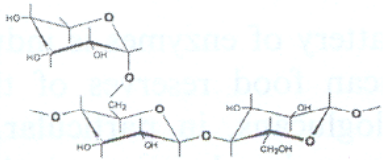
Xyloglucan



Cellulose

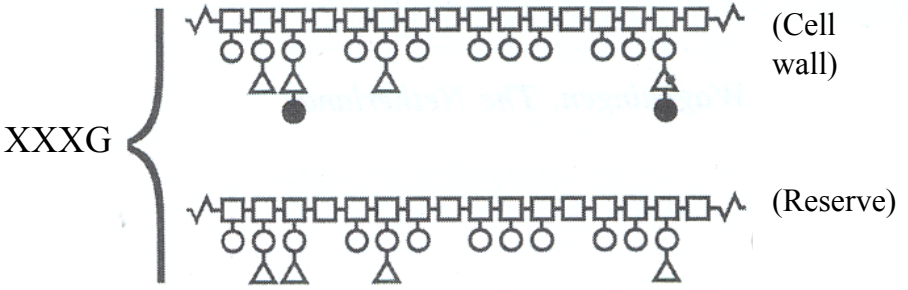


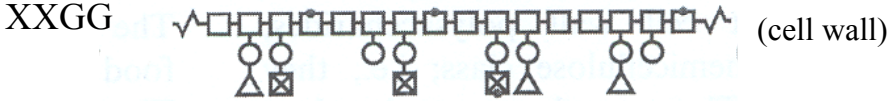
Xylan

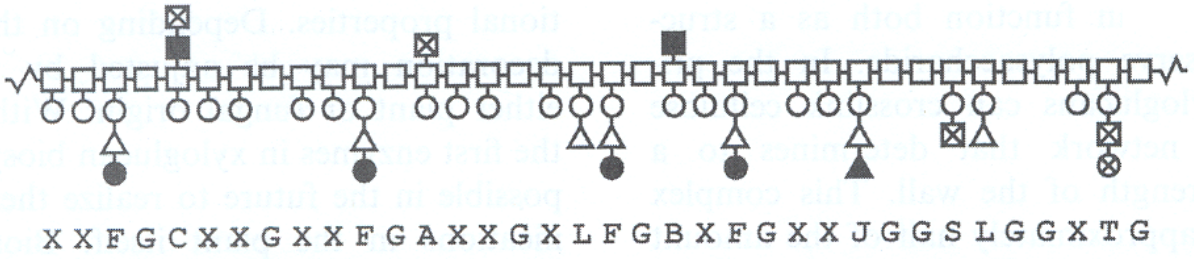


Xyloglucan

General branching pattern



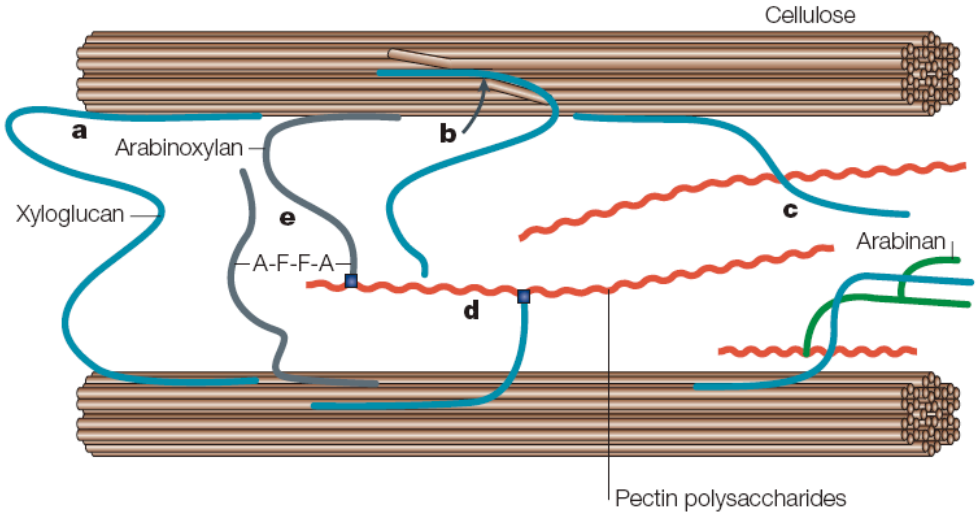




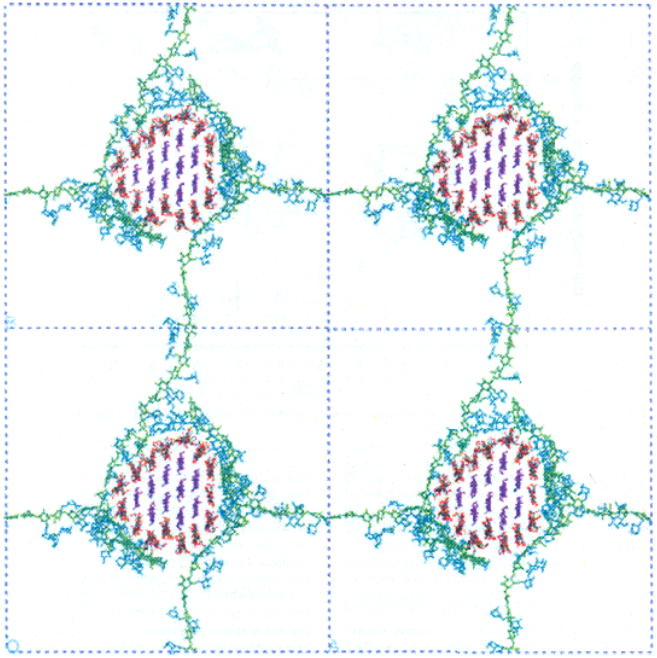
Hypothetical xyloglucan

Vincken (2003) In: *Handbook of Food Enzymology*

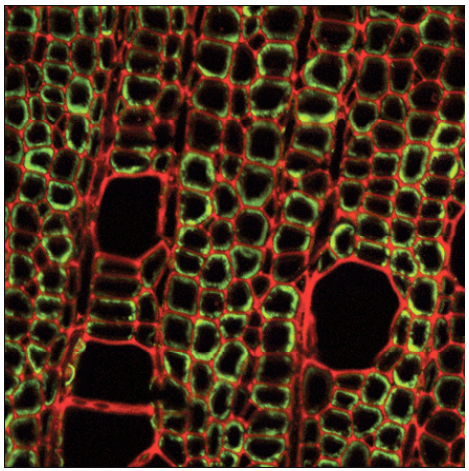
Xyloglucan in cell wall



Cosgrove, D. Nature Rev. Mol. Cell Biol. 2005



Hanus, J & Mazeau, K. Biopolymers 2006

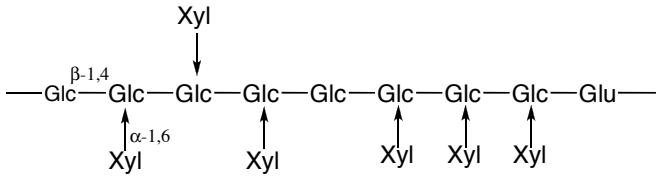


Mellerowicz et al. Ann. Bot. 2008

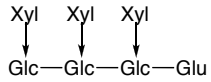
XEG5A: xyloglucan-specific endo-β-1,4-glucanase

Substrate	Relative activity
Tamarind xylogluca n	5.6
HDP xyloglucan oligosaccharide	2.6
CMC	1.0
Cellulose, xylan, arabinoxylan, pectin	n.d.

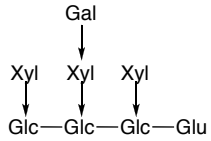
Tamarind xyloglucan



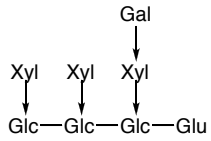
Blocks

(1) 

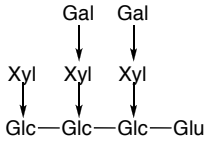
XXXG (heptasaccharide), XG7, DP7

(2) 

XLXG Ooctasaccharide), XG8, DP8



XXLG Ooctasaccharide), XG8, DP8

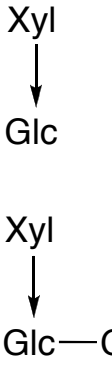
(3) 

XXLG (nonasaccharide). XG9. DP9

Tamarind xyloglucan *or*
HDP xyloglucan oligosaccharide

XEG5A

XXXG, XLXG/XXLG, XLLG

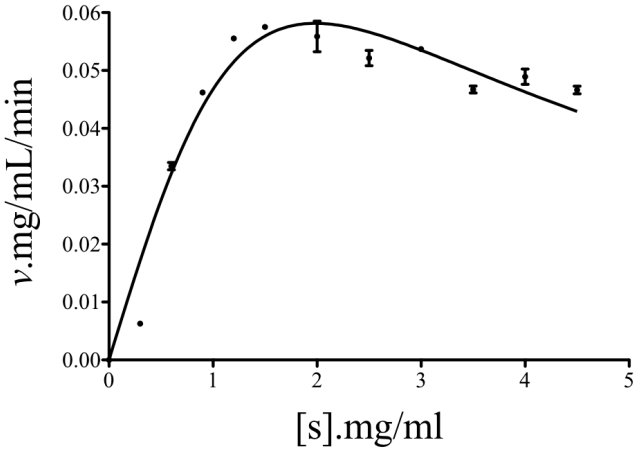


XG (isoprimeverose); XG2, DP2

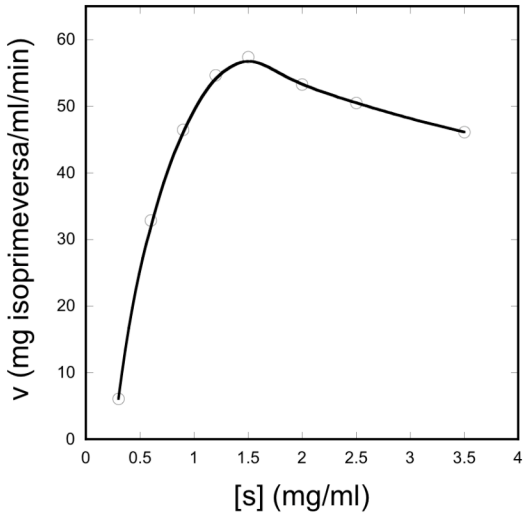
XGG (xylosyl-cellobiose); XG3, DP3

Wong et al. (2010) *Applied Microbiol. Biotechnol.*

XEG5A: xyloglucan-specific endo-β-1,4-glucanase

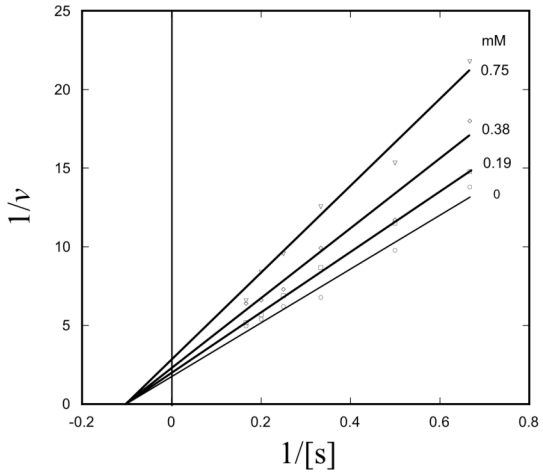


Xyloglucan oligosaccharide HDP-XGP

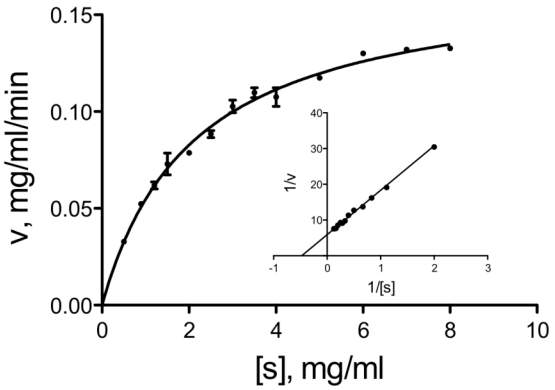


Xyloglucan oligosaccharide DP14

Xyloglucan oligosaccharid
DP7 (XXXG):
K_i = 1.46 ± 0.13 mM



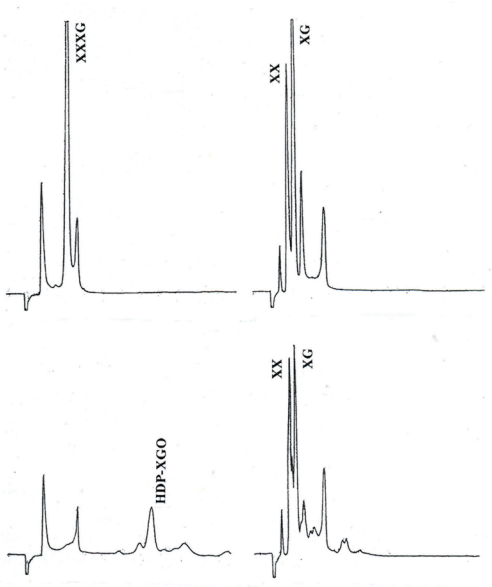
XEG5B: xyloglucan-specific exo-β-1,4-glucanase [EC 3.2.1.155]



K_m = 2.12 ± 0.13
K_{cat} = 47.6 min^{-1}
(Tamarind xyloglucan, 37°C, pH 6.0)

Active only on xyloglucan & HDP-XGO
Activity not detected -CMC, Avicel, acid cellulose, lichenan, laminarin, xylan, arabinoxylan

HPLC analysis of end-products



Tamarind xyloglucan *or*
HDP xyloglucan oligosaccharide

↓ XEG5B

?

XXXG, XLXG/XXLG, XLLG

↓

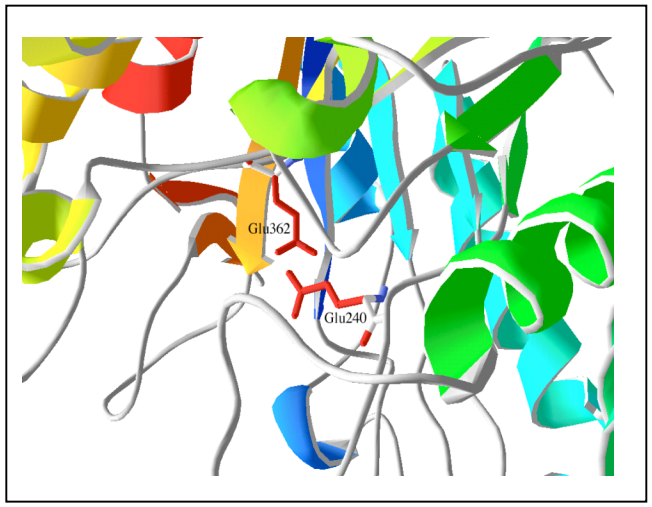
Xyl—Glc — isoprimeverose; XG2, DP2

Xyl
Glc—Glc — xylosyl-cellobiose; XG3, DP3

Wong et al. (2010) *Protein & Peptide Lett.*

XEG 5A & 5B Structures

Gene: 1293 bp
Protein: 431 aa (20 aa signal peptide, 45 kD)
Structure: $(\alpha/\beta)_8$ fold, GH family 5
Catalytic: Glu240, Glu362, retaining



XEG5A - endo

Gene: 1548 bp
Protein: 516 aa (18 aa signal peptide, 45.5 kD)
Structure: $(\alpha/\beta)_8$ fold GH family 5
Catalytic: Glu431, Glu293, retaining

XEG5B - exo

Uses
* Surfactants derived from oligoxyloglucans
* Biocomposites by grafting cellulose surfaces with xyloglucans
* Thermally reversible xyloglucan gels for slow-release delivery

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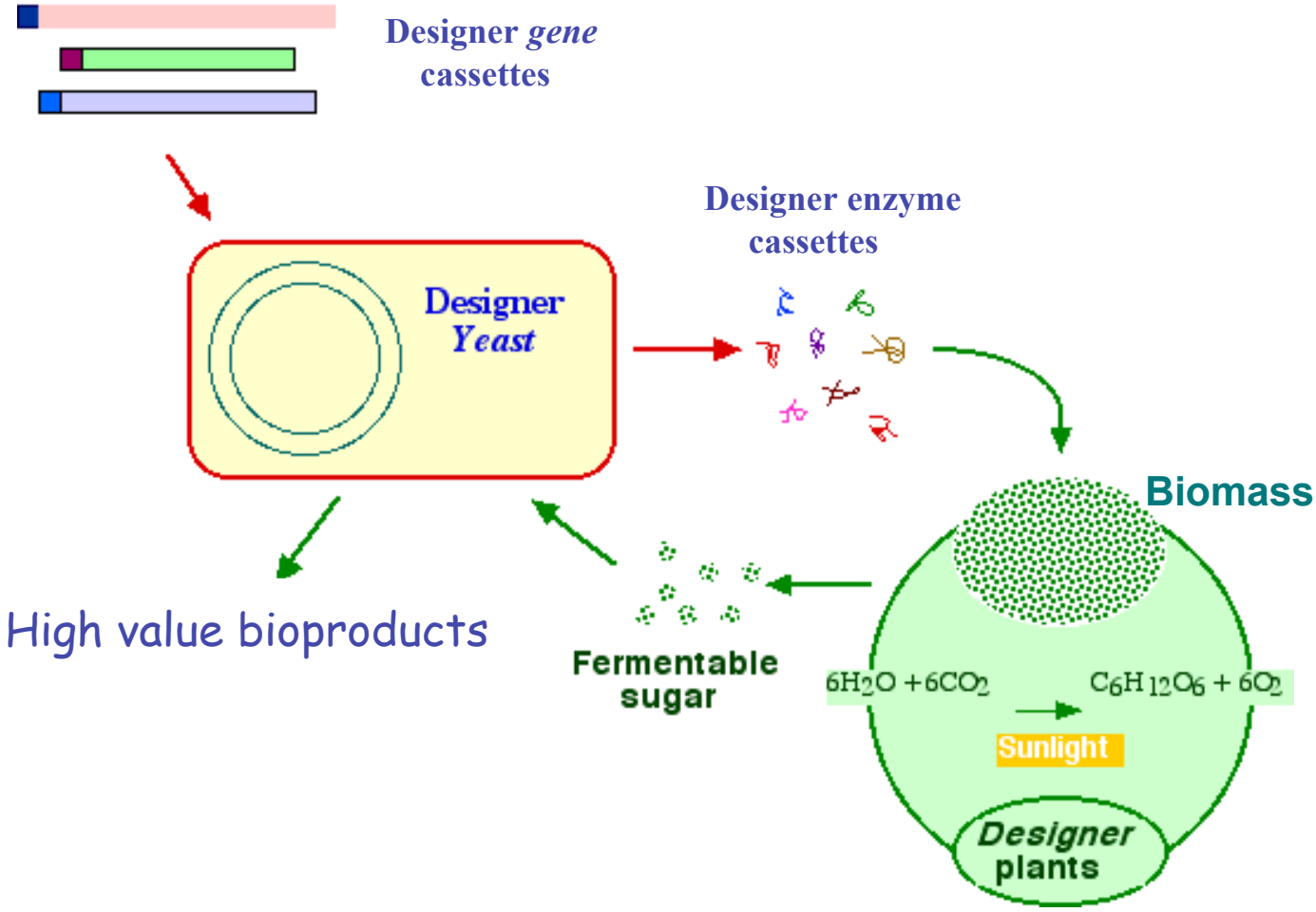
Engineering yeast



Chromosomal integration of genes in yeast

- Integration of multiple genes
- at precise locations in the yeast chromosome
- resulting in stable clones
- rapid testing of enzyme combinations in a cell-based system

Feruloyl esterase *endo-Xylanase* *endo-Cellulase*

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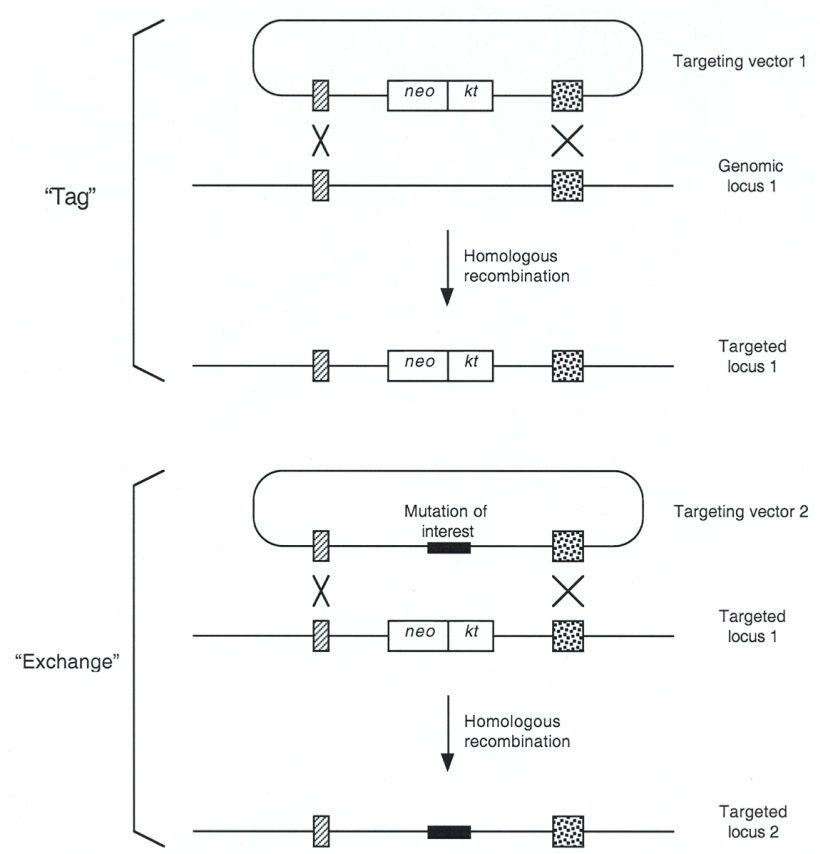
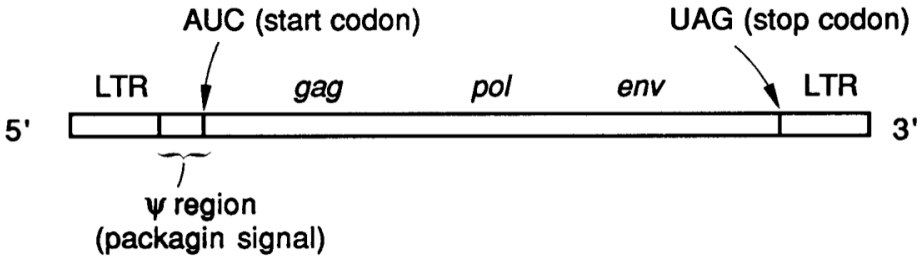
Homologous recombination



Double-hit replacement



Retrovirus RNA genome

LTR = long terminal repeats
gag = viral core protein
pol = reverse transcriptase
env = envelop
psi region = packaging signal

D. Wong *(2006) The ABCs of Gene Cloning*

Ty elements in *Saccharomyces cerevisiae*

Table 1. Distribution of Ty Elements by Chromosome

Chr. no.	Number of insertions[a] Ty1	Ty2	Ty3	Ty4	Ty5	Total Ty insertions on chromosome
I	6 (1)	1	1	0	0	8
II	13 (2)	2 (1)	1	2	0	18
III	11	2 (1)	0	1	2 (1)	16
IV	20 (5)	3 (3)	6	2	0	31
V	18 (2)	5	4	3	2	32
VI	8	1 (1)	0	1	0	10
VII	25 (3)	2 (2)	6 (1)	4	1	38
VIII	14 (1)	1	3	3 (1)	1	22
IX	5	1	2 (1)	1	0	9
X	14 (2)	4	1	3 (1)	0	22
XI	11	1	1	0	1	14
XII	18 (4)	3 (2)	4	1	0	26
XIII	15 (4)	2	0	3	0	20
XIV	5 (2)	1 (1)	4	3	0	13
XV	18 (2)	3 (2)	4	2	0	27
XVI	16 (4)	2	4	3 (1)	0	25
Genome total	217 (32)	34 (13)	41 (2)	32 (3)	7 (1)	331

[a]Insertions include full-length elements, solo LTRs, and LTR fragments. Numbers of full-length elements are shown in parentheses.

From *Genome Res*. 8:464-478 (1998)

Single gene clones:

endoglucanase, exoglucanase, b-glucosidase, feruloyl esterase, endoxylaase, α-amylase, glucoamylase,

Double clones (co-integration):

(1) endoxylanase + feruloyl esterase

(2) α-amylase/glucoamylase

Triple clones:

(1) endoglucanase + exoglucanase + β-glucosidase

(2) endoglucanase + endoxylanase + feruloyl esterase

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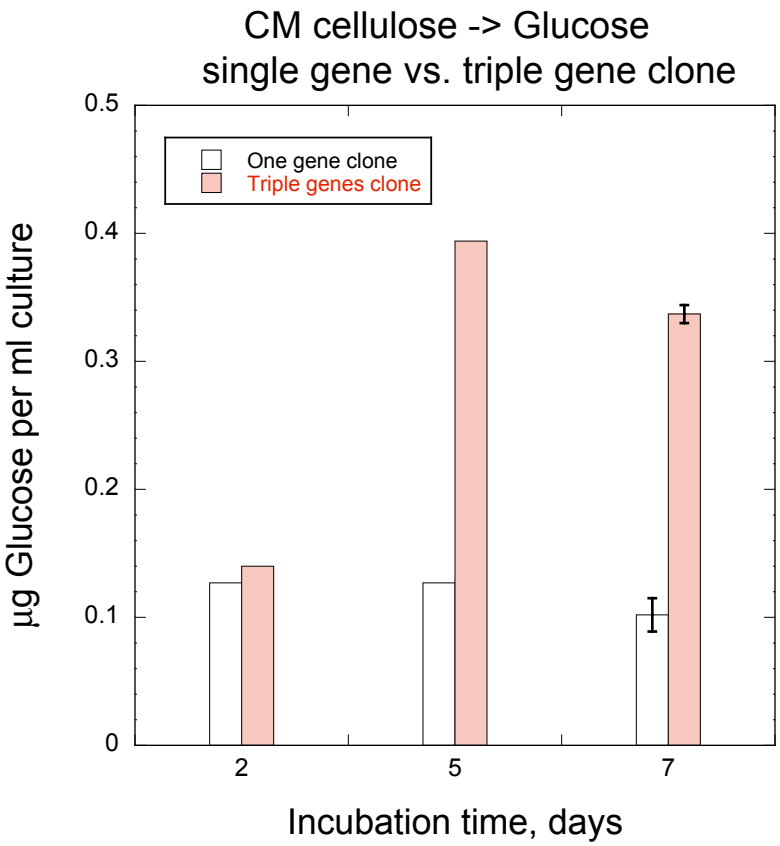
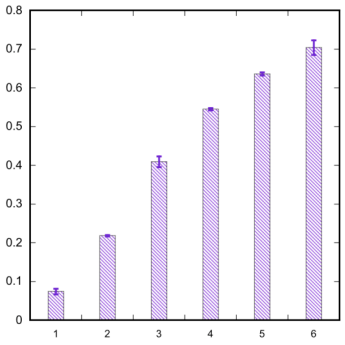
Triple Clones

endoglucanase + endoxylanase + feruloyl esterase

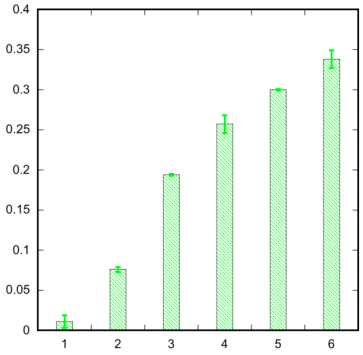
endoglucanase + exoglucanase + β-glucosidase



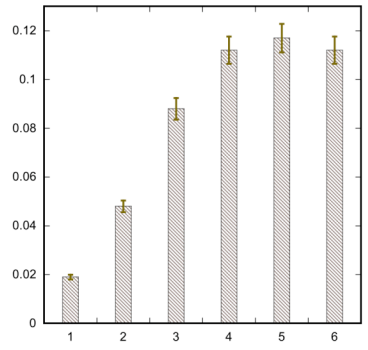


CM celllulose activity

Arabinoxylan activity



NP ferulate activity



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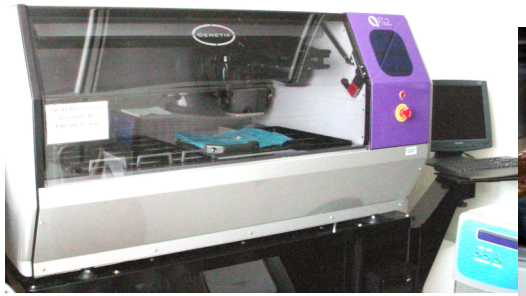
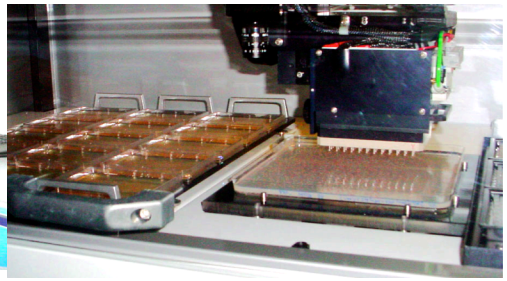
High-Throughput Functional Screening for active clones in Libraries

Metagenomic libraries

•Gene discovery
•Gene variants





Q-Pix

Liquid handling

Capabilities

Incubator-shaker
for microplates

Microplate
reader

•Assay development
•Assay under bioprocess
condition

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Genes and Enzymes

Enzymes	*Genes*	*Expressed*	*Purified*	*Characterized*
ferulic acid esterase*	12	10	10	10
α-glucuronidase*	(1)	1	1	1
glucuronoyl esterase*	(1)	1	1	1
arabinanase*	9	4	3	2
xyloglucanase*	6	2	1	1
mannanase-glucanase (bifunctional)*	1	1	1	1
rhamnogalacturonanase	1	1	0	0
galactanase	4			
arabinofuranosidase	2			
endoxylanase	2			
Galactomannanase	1			
acetylxylan esterase	2			

Yeast vectors episomal, integrative

Yeast clones single gene, double genes, triple genes

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ADDITIONAL ENZYME/PROTEIN RESEARCH AREAS:

- Mutangenesis and directed evolution of enzymes
- Engineering *yeast* for consolidated bioprocessing
- Enzymatic fragmentation of crystalline and complex biopolymers
- Enzymatic modification of food proteins

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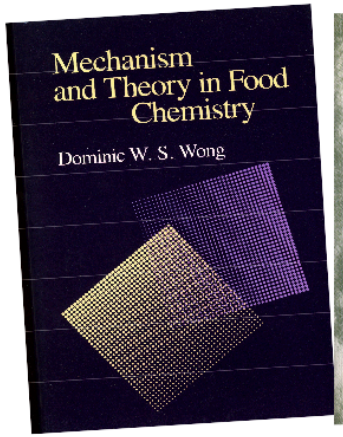
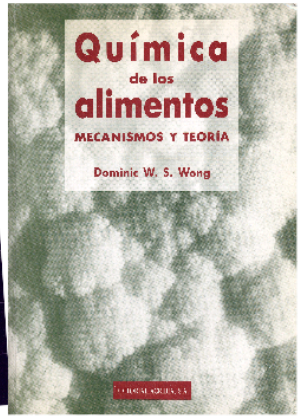
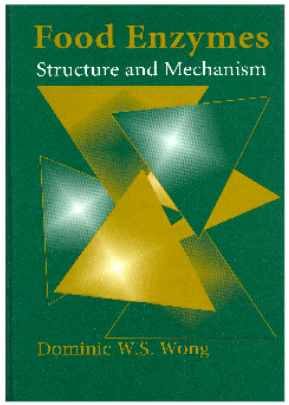
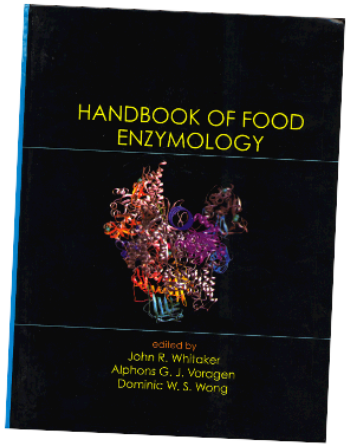
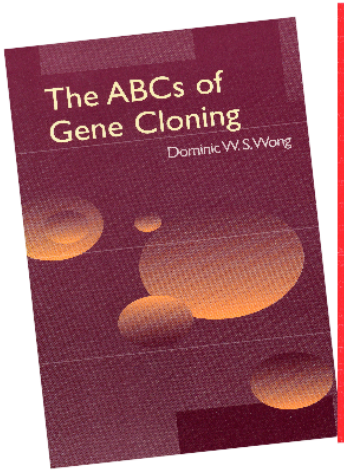
Backgrounds

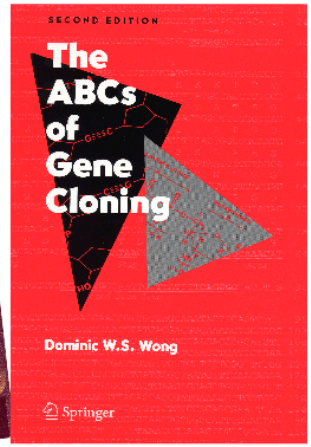






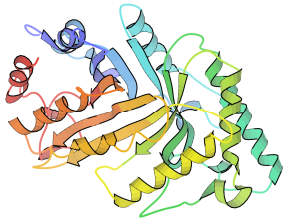






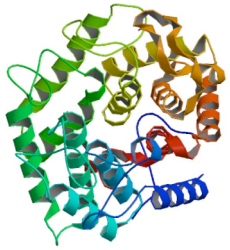
Enzymes.....

Enzymes.....



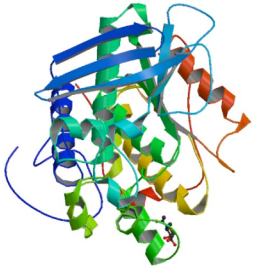
α–amylase

Feruloyl esterase

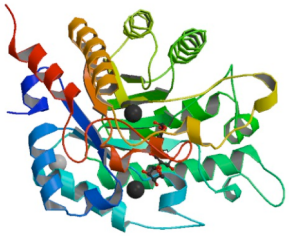


Endoglucanase

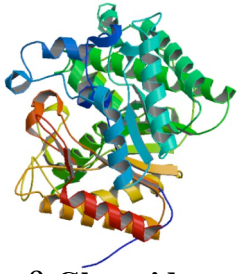
Cellobiohydrolase



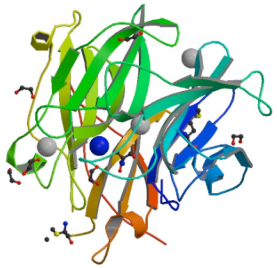
Glucuronoyl esterase



Endoxylanase



β-Glucosidase



β-Xylosidase

Glucoamylase
(catalytic domain)

- Sarah Batt
- Victor chan
- Kurt Wagschal
- Rena Kibblewhite
- Charles Lee
- Amanda McCormack
- Meiling Shang
- David Wan
- Dominic Wong

- Dr. Peter Biely, Slovak Academy of Science
- Dr. Mario Murakami, LNBio, Brazil
- Dr. Mary Jo Zidwick, Cargill Biotech Center
- Professor Roy Doi, UC Davis
- Dr. Hans Liao, OPX Biotechnologies

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Thank You.....