

# Ascospore Formation in the Yeast *Saccharomyces cerevisiae*

Aaron M. Neiman\*

Department of Biochemistry and Cell Biology, SUNY Stony Brook,  
 Stony Brook, New York 11794-5215

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## INTRODUCTION

The generation of ascospores is a defining feature of the fungal phylum *Ascomycota*. Ascospores are generally found in clusters of four or eight spores within a single mother cell, the ascus. These spores are formed as a means of packaging post-meiotic nuclei. As such, they represent the “gametic” stage of the life cycle in these fungi. The creation of these specialized cells requires a cell division mechanism distinct from that used during vegetative growth of fungal cells. This review will focus on this process in the best-studied model, the baker’s yeast *Saccharomyces cerevisiae*.

## OVERVIEW OF ASCOSPORE FORMATION

An overview of the cytological events of sporulation in *S. cerevisiae* is shown in Fig. 1. Diploid cells of *S. cerevisiae* modify their growth in response to nutrient availability. In the presence of nutrients they grow in budding form. The presence of a poor nitrogen source such as proline will trigger the onset

of mitotic growth in a pseudohyphal form (46). The complete absence of nitrogen, and the presence of a nonfermentable carbon source such as acetate, causes the cells to exit the mitotic cycle, undergo meiosis, and sporulate (40).

When triggered to enter the sporulation program, cells exit the mitotic cycle from the G<sub>1</sub> phase (40, 80). This is followed by premeiotic DNA synthesis and entry into the meiotic divisions. As with mitosis in yeast, meiosis is “closed”; that is, it takes place without breakdown of the nuclear envelope. The spindle microtubules are nucleated from the nuclear face of spindle pole bodies (SPBs) that span the nuclear envelope (21). In G<sub>1</sub> cells, there is a single SPB present in the cell. This SPB duplicates prior to meiosis I to generate the two poles of the meiosis I spindle. SPB duplication proceeds by a conservative process in which the daughter SPB is assembled adjacent to the mother and then migrates to the opposite side of the nucleus to form a spindle (2). As the cell exits from meiosis I, the SPBs duplicate again and the two meiosis II spindles are generated (Fig. 1B). Separation of the sister chromatids during anaphase II pulls a haploid set of chromosomes into each of four lobes of the nucleus. These lobes then pinch off at the end of meiosis II to generate four haploid nuclei.

The process of spore formation begins during meiosis II. Vesicles are recruited to the cytoplasmic side of each of the

\* Mailing address: Department of Biochemistry and Cell Biology, SUNY Stony Brook, Stony Brook, NY 11794-5215. Phone: (631) 632-1543. Fax: (631) 632-8575. E-mail: Aaron.Neiman@sunysb.edu.

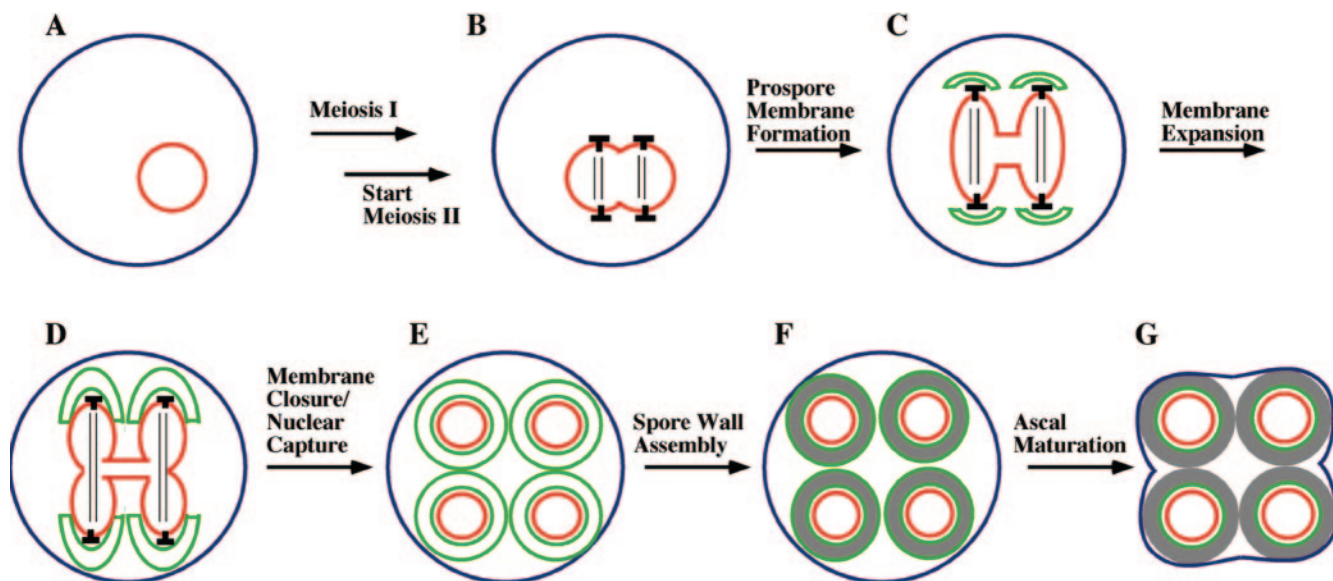


FIG. 1. (A to G) Overview of the stages of spore and ascus formation. In the presence of a nonfermentable carbon source, diploid cells starved for nitrogen will undergo meiosis. During the second meiotic division, the SPBs (indicated as  $\top$ ), which are embedded in the nuclear envelope (shown in red), become sites for formation of prospore membranes (shown in green). As meiosis II proceeds, the prospore membranes expand and engulf the forming haploid nuclei. After nuclear division, each prospore membrane closes on itself to capture a haploid nucleus within two distinct membranes. Spore wall synthesis then begins in the lumen between the two prospore membrane-derived membranes. After spore wall synthesis is complete, the mother cell collapses to form the ascus.

four SPBs, where they coalesce to form flattened double-membrane sheets termed prospore membranes (often referred to as the prospore wall in the older literature). (Fig. 1C) (21, 99). These four prospore membranes expand during the course of meiosis II so that at the time of nuclear division each prospore membrane completely engulfs the nuclear lobe to which it is anchored via the SPB (Fig. 1D). After nuclear division, the open end of each prospore membrane closes off so that each daughter nucleus and its associated cytoplasm is completely encapsulated within a double membrane (Fig. 1E). The closure of the prospore membranes is a cytokinesis event, generating four autonomous prospores. Of the two distinct membranes derived from the prospore membrane, the one closest to the nucleus now serves as the plasma membrane of the prospore.

The prospores mature into spores through the synthesis of a spore wall. Assembly of the spore wall occurs initially in the lumen between the two prospore membrane-derived membranes (Fig. 1F) (91). As described below, the outer of these two membranes breaks down during the process of spore wall formation. Once complete, the spore wall confers on the spore resistance to a variety of environmental stresses (156). After the completion of the spore wall, the now anucleate mother cell, which remains intact throughout sporulation, is remodeled to serve as an ascus that encapsulates the four spores of the tetrad (Fig. 1G). A mutation that delays or blocks the shrinking of the ascus around the spore has been described, suggesting that ascus maturation may be an active process, rather than a simple lysis of the anucleate cell (105). The mechanisms controlling this process remain unknown, although vacuolar proteases have been suggested to play a role in the remodeling (187).

#### AN UNDERLYING TRANSCRIPTIONAL CASCADE

Starvation and entry into meiosis are prerequisites for spore formation to begin. Both the control of entry into meiosis and the details of meiotic chromosome segregation have been the subject of earlier reviews (57, 86, 98, 138, 160) and will only be touched on here. The coordination of spore formation with these other events is achieved through an underlying transcriptional cascade (175). Studies using Northern analysis identified a number of transcripts whose expression was induced by sporulation conditions (56, 126). Time course studies grouped these transcripts broadly into three categories: early, middle, and late genes. As more genes have been characterized, sub-categories such as early-middle and mid-late have been added to the classifications (13, 120).

The use of transcriptional microarrays to examine the pattern of transcription during sporulation has confirmed and expanded on these original studies (25, 130). Several waves of induction are seen. Immediately upon transfer to sporulation medium, many genes involved in metabolic adaptation are induced. Subsequent to that immediate response, a set of transcripts termed the early genes are upregulated. Many of these genes are involved in meiotic chromosome metabolism. After the early genes, a second large set of genes are coordinately induced. These middle genes include those involved in progression through the meiotic divisions, such as genes encoding cyclins and components of the anaphase-promoting complex, as well as many of the genes encoding proteins directly involved in the construction of the spore (described below) (25, 130). The transcription factor *NDT80* is required for the induction of these middle genes (26). The common regulation of

meiotic functions and spore assembly functions by Ndt80p may help maintain coordination between the two processes.

After the middle genes, at a time roughly correlated with the completion of meiosis II, a smaller set of mid-late genes are induced, and subsequent to that a set of "late" genes are induced. While some of the late genes are required for the assembly of the spore wall, others appear to be more general stress response genes, and still others are haploid-specific genes whose induction may correlate with restoration of the haploid state in the spores (130).

The characterization of sporulation-induced genes has been a fruitful source for study of meiosis- and sporulation-defective mutants. In particular, Nasmyth and colleagues (133) constructed a set of ~300 strains, each deleted for a different sporulation-induced gene. Analyses of this collection have yielded more than 40 mutants with a variety of sporulation defects (27, 133). It is noteworthy, however, that, despite their regulated expression, deletion of most of these genes does not cause a detectable sporulation phenotype. Thus, sporulation-induced expression is not strongly correlated with an obvious sporulation function.

#### A GENETIC DESCRIPTION OF SPORE FORMATION

Beginning with the pioneering studies by Esposito and Esposito (38), more than 30 years of work has identified a large number of sporulation-defective mutants, some by directed screens and others by happenstance (40, 80). The identification and analysis of these mutants provide the basis for much of what is known about sporulation.

More recently, the construction of the yeast knockout collection has provided an invaluable new tool that allows the development of an even more comprehensive picture of the genes involved in sporulation. In addition to systematic screens of a collection of sporulation-induced genes (133) or a subset of the knockout collection (12), three different groups have used this resource to screen the homozygous diploid knockouts of all the nonessential yeast genes for sporulation defects. Importantly, each group has done the screening in a slightly different way. One such study screened the knockout collection directly in the light microscope for the presence of spores (36). A second screened the collection by using the microarray-based technique of phenotypic profiling (34, 152). In this analysis, sporulation was operationally defined as the acquisition of resistance to cell wall-degrading enzymes. The third study first crossed the knockout collection to a different genetic background, SK-1, which displays enhanced sporulation (92). Hybrid strains homozygous for each knockout were generated, and these strains were then screened visually for sporulation (92).

The result of these efforts is that we now have a much more complete picture of the genetic requirements for sporulation. Each of these studies identified >200 mutants defective in sporulation. Perhaps not surprisingly given the different techniques and strain backgrounds involved, these mutants fall into three distinct but overlapping sets. Table 1 presents a list of the 289 genes identified in at least two of the three screens. Based on annotations available for each of these genes at the *Saccharomyces* Genome Database ([www.yeastgenome.org](http://www.yeastgenome.org)), they have been divided into five broad categories: (i) genes in-

involved in mitochondrial/metabolic functions, (ii) vacuolar and autophagy genes, (iii) genes previously characterized as having premeiotic or meiotic defects, (iv) genes with known defects in prospore membrane or spore wall assembly, and (v) genes whose role in sporulation has not yet been defined.

The majority of sporulation-defective mutants arrest prior to the meiotic divisions (36). Mutants in the first three categories above all fall into this class. Because it occurs in the presence of a nonfermentable carbon source, sporulation is an obligatorily aerobic process, and respiratory-incompetent yeast cells arrest upon transfer to sporulation medium (110, 169). This is the reason for the large number of genes with known mitochondrial functions listed in the first group in Table 1. Similarly, because sporulation takes place in the absence of a nitrogen source, during this process all the amino acids used for new protein synthesis must be scavenged by recycling of pre-existing proteins. The repeated identification of many genes involved in delivery of proteins to the vacuole or in autophagy (second group in Table 1), which are necessary for protein recycling, is consistent with insufficient amino acid availability blocking entry into the meiotic divisions. Included in the third group of genes are many required for proper chromosome segregation during meiosis as well as transcription factors required for the induction of early genes.

The fourth, and smallest, group of genes are those known to be required for assembly of the spore per se. The majority of these genes are induced by Ndt80p at the onset of meiosis I (25, 130). The functions of many of them will be described in detail below. Finally, the fifth group consists of another 80 genes whose roles in sporulation have yet to be determined. While this group includes genes with various functions, it is noteworthy that a large fraction (~30%) appear to be involved in gene expression as subunits of either chromatin remodeling or mRNA processing complexes. Many of these genes may, therefore, be involved in regulation of meiotic gene expression.

Even this large list does not present a comprehensive picture of the genes required for sporulation. Essential genes were not assayed in any of the genomic screens, and many are likely to be required for sporulation. Moreover, because of the nature of the assays involved, mutations that make visible but inviable spores, such as mutants defective in meiotic chromosome segregation, are absent from the list in Table 1. Similarly, mutants that form defective but visible spore walls, such as those with outer spore wall defects (see below), are underrepresented in the two visual screens. Nonetheless, these studies have provided us with a broad overview of the genetic requirements for spore formation. The remainder of this review will focus on the cellular events involved in formation of spores and the genes known to participate in these events.

#### TO MAKE A SPORE

The process of spore construction requires two events in which cellular structures are assembled de novo. First, prospore membranes must be generated around the daughter nuclei to create prospores. Second, the resulting prospores must be surrounded by a protective spore wall. These events will be considered in turn.

TABLE 1. Genes required for efficient sporulation in *Saccharomyces cerevisiae*

		Genes involved in <sup>a</sup> :							
Mitochondria/metabolism		Vacuole/autophagy		Meiosis/early sporulation		Spore formation		Unknown sporulation role	
Gene	ORF	Gene	ORF	Gene	ORF	Gene	ORF	Gene	ORF
<i>ACS1</i>	<b>YAL054c</b>	<i>AUT7</i>	<b>YBL078c</b>	<i>SPO7</i>	<b>YAL009w</b>	<i>GIP1</i>	<b>YBR045c</b>	<i>ATS1</i>	YAL020c
<i>PET112</i>	YBL080c	<i>CCZ1</i>	YBR131w	<i>MUM2</i>	YBR057c	<i>SPO71</i>	<b>YDR104c</b>	<i>CCR4</i>	<b>YAL021c</b>
<i>COQ1</i>	YBR003w	<i>APG12</i>	<b>YBR217w</b>	<i>MMS4</i>	YBR098w	<i>SWM1</i>	<b>YDR260c</b>	<i>FUN12</i>	YAL035w
<i>SCO1</i>	YBR037c	<i>BPH1</i>	YCR032w	<i>KAR4</i>	<b>YCL055w</b>	<i>DIT1</i>	YDR403w	<i>GPB2</i>	YAL056w
<i>TPS1</i>	YBR126c	<i>CVT17</i>	<b>YCR068w</b>	<i>CSM1</i>	YCR086w	<i>SPO73</i>	YER046w		YAL068c
<i>ADY2</i>	YCR010c	<i>BRE1</i>	YDL074c	<i>RAD57</i>	YDL059c	<i>ISC10</i>	<b>YER180c</b>	<i>NUP170</i>	YBL079w
<i>SLM3</i>	YDL033c	<i>VAM6</i>	YDL077c	<i>RAD55</i>	YDR076w	<i>SPO74</i>	<b>YGL170c</b>	<i>HMT1</i>	YBR034c
<i>MRPL1</i>	YDR116c	<i>ATG9</i>	<b>YDL149w</b>	<i>GSG1</i>	YDR108w	<i>SPR3</i>	YGR059w	<i>FAT1</i>	YBR041w
<i>KGD2</i>	YDR148c	<i>VPS41</i>	<b>YDR080w</b>	<i>HPR1</i>	YDR138w	<i>CRH1</i>	YGR189c	<i>UBP14</i>	YBR058c
<i>CBS2</i>	YDR197w	<i>BRS52</i>	YDR484w	<i>UME6</i>	<b>YDR207c</b>	<i>AMA1</i>	<b>YGR225w</b>	<i>RPL19A</i>	YBR084c-a
<i>TCM10</i>	YDR350c	<i>VPS3</i>	YDR495c	<i>RMD5</i>	<b>YDR255c</b>	<i>NDT80</i>	YHR124w		YBR090c
<i>ATP17</i>	YDR377w	<i>VAC8</i>	YEL013w	<i>ZIP1</i>	YDR285w	<i>CHS7</i>	YHR142w	<i>SGF29</i>	YCL010c
<i>SHE9</i>	<b>YDR393W</b>	<i>PRB1</i>	<b>YEL060c</b>	<i>EM11</i>	<b>YDR512c</b>	<i>SSP1</i>	<b>YHR184w</b>	<i>THR4</i>	<b>YCR053w</b>
<i>ACN9</i>	YDR511w	<i>ATG18</i>	YFR021w	<i>RAD51</i>	YER095w	<i>ADY1</i>	<b>YHR185c</b>	<i>SED4</i>	YCR067c
<i>QCR7</i>	YDR529c	<i>MON1</i>	<b>YGL124c</b>	<i>DMC1</i>	YER179w	<i>SPO14</i>	YKR031c	<i>STP4</i>	YDL048c
<i>RIP1</i>	<b>YEL024w</b>	<i>ATG1</i>	<b>YGL180w</b>	<i>RIM15</i>	YFL033c	<i>SPO75</i>	<b>YLL005c</b>		YDL072c
<i>AFG3</i>	YER017c	<i>VD30</i>	YGL227w	<i>RMD8</i>	YFR048w	<i>OSW2</i>	<b>YLR054c</b>	<i>UFD2</i>	YDL190c
<i>PET117</i>	YER058w	<i>CLC1</i>	YGR167w	<i>HOP2</i>	YGL033c	<i>SEC22</i>	YLR268w	<i>SNF3</i>	YDL194w
<i>ICL1</i>	<b>YER065c</b>	<i>PPA1</i>	YHR026w	<i>SAE2</i>	<b>YGL175c</b>	<i>CDA2</i>	YLR308w	<i>GCS1</i>	YDL226c
<i>PDA1</i>	YER178w	<i>ATG7</i>	<b>YHR171w</b>	<i>DOC1</i>	YGL240w	<i>CHS5</i>	YLR330w	<i>MAF1</i>	YDR005c
<i>RMD9</i>	YGL107c	<i>VPS53</i>	YJL029c	<i>ZIP2</i>	YGL249w	<i>SPO77</i>	YLR341w	<i>CIS1</i>	YDR022c
<i>MET13</i>	YGL125w	<i>SNX4</i>	YJL036w	<i>RMD11</i>	<b>YHL023c</b>	<i>SMA2</i>	<b>YML066c</b>		YDR117c
<i>HAP2</i>	<b>YGL237c</b>	<i>VPS25</i>	YJR102c	<i>RIM4</i>	YHL024w	<i>SPO20</i>	<b>YMR017w</b>	<i>STB3</i>	YDR169c
<i>SHY1</i>	YGR112w	<i>VPS24</i>	<b>YKL041w</b>	<i>RIM101</i>	YHL027w	<i>MSO1</i>	<b>YNR049c</b>	<i>CDC40</i>	YDR364c
	YGR150c	<i>VPS13</i>	<b>YLL040c</b>	<i>NEM1</i>	YHR004c	<i>SPO21</i>	<b>YOL091w</b>	<i>RVS167</i>	YDR388w
<i>CBP4</i>	YGR174c	<i>ATG10</i>	<b>YLL042c</b>	<i>SPO13</i>	YHR014w	<i>MPC54</i>	<b>YOR177c</b>	<i>SAC7</i>	YDR389w
<i>QCR9</i>	YGR183c	<i>SNF7</i>	YLR025w	<i>SAE3</i>	YHR079c-a	<i>SSP2</i>	<b>YOR242c</b>	<i>SPT3</i>	YDR392w
<i>LSC2</i>	YGR244c	<i>PEP3</i>	YLR148w	<i>NAM8</i>	YHR086w	<i>OSW1</i>	<b>YOR255w</b>	<i>SSN2</i>	YDR443c
<i>CBP2</i>	<b>YHL038c</b>	<i>VPS36</i>	YLR417w	<i>SPO12</i>	YHR152w	<i>MUM3</i>	<b>YOR298w</b>	<i>SNF1</i>	YDR477w
<i>RRF1</i>	YHR038w	<i>ATG17</i>	YLR423c	<i>SPO16</i>	YHR153c	<i>SMA1</i>	<b>YPL027w</b>	<i>NPR2</i>	YEL062w
<i>HTD2</i>	YHR067w	<i>YPT7</i>	<b>YML001w</b>	<i>MND2</i>	YIR025w	<i>SPO19</i>	<b>YPL130w</b>	<i>CKB1</i>	YGL019w
<i>MSR1</i>	YHR091c	<i>VPS20</i>	YMR077c	<i>IME2</i>	<b>YJL106w</b>	<i>SSO1</i>	<b>YPL232w</b>	<i>SGF73</i>	YGL066w
	<b>YHR116w</b>	<i>ATG16</i>	<b>YMR159c</b>	<i>IDS2</i>	YJL146w	<i>SMK1</i>	<b>YPR054w</b>	<i>ARC1</i>	YGL105w
<i>MRPL6</i>	YHR147c	<i>ATG2</i>	YNL242w	<i>IME1</i>	<b>YJR094c</b>			<i>RTF1</i>	<b>YGL244w</b>
<i>MDM31</i>	YHR194w	<i>VPS27</i>	YNR006w	<i>CST9</i>	YLR394w			<i>PEX31</i>	YGR004w
<i>PFK26</i>	YIL107c	<i>AUT1</i>	YNR007c	<i>RIM11</i>	<b>YMR139w</b>			<i>BTN2</i>	YGR142w
<i>KGD2</i>	YIL125w	<i>PEP12</i>	YOR036w	<i>RIM13</i>	YMR154c			<i>GCN5</i>	YGR252w
	YIL157c	<i>VPS5</i>	YOR069w	<i>SPO1</i>	YNL012w			<i>ZUO1</i>	YGR285c
<i>PET191</i>	YJR034w	<i>SNF8</i>	YPL002c	<i>MCK1</i>	YNL307c			<i>PRS3</i>	<b>YHL011c</b>
<i>NFU1</i>	YKL040c	<i>ATG5</i>	<b>YPL149w</b>	<i>EM15</i>	<b>YOL071w</b>			<i>ARD1</i>	YHR013c
<i>MDH1</i>	<b>YKL085w</b>	<i>ATG11</i>	YPR049c	<i>ME15</i>	YPL121c				YHR132w-a
<i>CYT2</i>	YKL087c	<i>VPS4</i>	YPR173c	<i>REC8</i>	<b>YPR007c</b>			<i>RPN10</i>	<b>YHR200w</b>
<i>HAP4</i>	YKL109w	<i>ATG13</i>	YPR185w	<i>CLB5</i>	<b>YPR120c</b>			<i>RPL34B</i>	<b>YIL052c</b>
<i>SDH1</i>	YKL148c								YIL060w
<i>PCK1</i>	<b>YKR097w</b>							<i>LAS21</i>	YJL062w
<i>MMM1</i>	YLL006w								YJL160c
<i>COX17</i>	<b>YLL009c</b>								YJR008w
<i>SDH2</i>	<b>YLL041c</b>								YKL033w-a
<i>MEF1</i>	YLR069c								YKL054c
<i>CSF1</i>	YLR087c							<i>NUP120</i>	YKL057c
<i>MSS51</i>	<b>YLR203c</b>							<i>MUD2</i>	YKL074c
	YLR368w							<i>CTK1</i>	YKL139w
<i>FBP1</i>	<b>YLR377c</b>							<i>CBT1</i>	YKL208w
<i>NAM2</i>	YLR382c							<i>DOA1</i>	YKL213c
<i>PIF1</i>	YML061c								YKR089c
<i>ATP18</i>	YML081c-a								YLL033w
<i>NDI1</i>	YML120c							<i>UBI4</i>	<b>YLL039c</b>
<i>RSF1</i>	YMR030w							<i>AAT2</i>	YLR027c
<i>PET111</i>	YMR257c							<i>IES3</i>	YLR052w
<i>PPA2</i>	YMR267w								YLR149c
<i>CAT8</i>	YMR280c							<i>DCS1</i>	YLR270w
<i>AEP2</i>	YMR282c							<i>HMG2</i>	<b>YLR450w</b>
<i>MRPL33</i>	<b>YMR286w</b>							<i>GSF2</i>	YML048w
<i>MSU1</i>	YMR287c								YMR010w
<i>COX5A</i>	YNL052w							<i>SRT1</i>	<b>YMR101c</b>

Continued on following page

TABLE 1—Continued

Genes involved in <sup>a</sup> :									
Mitochondria/metabolism		Vacuole/autophagy		Meiosis/early sporulation		Spore formation		Unknown sporulation role	
Gene	ORF	Gene	ORF	Gene	ORF	Gene	ORF	Gene	ORF
<i>MLS1</i>	<b>YNL117w</b>							<i>STO1</i>	YMR125w
<i>ATP11</i>	YNL315c								YMR196w
<i>COQ2</i>	YNR041c							<i>SSN8</i>	YNL025c
<i>PET494</i>	YNR045w							<i>WHI3</i>	YNL197c
<i>IFM1</i>	YOL023w							<i>CAF40</i>	YNL288w
<i>MDH2</i>	YOL126c							<i>POP2</i>	YNR052c
<b><i>CYT1</i></b>	<b>YOR065w</b>							<i>SIN3</i>	YOL004w
<i>CAT5</i>	YOR125c								YOR008c-a
<b><i>IDH2</i></b>	<b>YOR136w</b>							<i>WHI5</i>	YOR083w
<i>LSC1</i>	YOR142w							<i>LEO1</i>	YOR123w
<i>MRPL23</i>	YOR150w							<i>RIS1</i>	YOR191w
<i>PET123</i>	YOR158w								YOR291w
<i>LIP5</i>	YOR196c							<i>SSN3</i>	YPL042w
<i>HAP5</i>	YOR358w							<i>LGE1</i>	YPL055c
<i>MRPS16</i>	YPL013c							<i>TGS1</i>	YPL157w
<i>LPE10</i>	YPL060w								YPL166w
<i>MSD1</i>	YPL104w							<i>CBC2</i>	YPL178w
<b><i>MRP51</i></b>	<b>YPL118w</b>								YPL208w
<b><i>COX10</i></b>	<b>YPL172c</b>							<b><i>UBA3</i></b>	<b>YPR066w</b>
<i>MRPL40</i>	YPL173w								
<i>CBP3</i>	YPL215w								

<sup>a</sup> Genes listed were identified as causing sporulation-defective phenotypes in two of the three genomic screens (34, 91, 132). Overlapping or dubious open reading frames (ORFs) have been removed from the list. Genes in boldface were found in all three screens.

### Formation and Growth of the Prospore Membrane

The formation and growth of the prospore membrane occur in several steps (Fig. 2). Initially, the SPB is modified to allow its cytoplasmic surface to act as a site for the homotypic fusion of secretory vesicles into a membrane sheet (4, 31, 76) (Fig. 2A to C). A specialized SNARE complex is then required for the fusion event itself (104). After the initial formation of a membrane cap on the SPB, the membrane continues to grow by vesicle addition. During membrane expansion, membrane-associated complexes are required properly shape the membrane so that its growth is properly constrained and ultimately leads to nuclear capture (Fig. 2D and E). The details of these different steps in membrane formation are described below.

**Modification of the SPB.** The SPB is the sole microtubule-organizing center in an *S. cerevisiae* cell and spans the nuclear envelope so that it has distinct nuclear and cytoplasmic faces (67). The SPB is a multilaminar structure with a nuclear face, or inner plaque, from which spindle microtubules are nucleated; a central plaque, which contains the Spc42 protein, spanning the nuclear envelope; and a cytoplasmic face, or outer plaque, which is the organizing center for cytoplasmic microtubules (Fig. 3). In vegetative cells, the outer plaque consists primarily of three proteins, Cnm67p, Nud1p, and Spc72p (183). Spc72p acts a receptor for the gamma-tubulin complex, anchoring cytoplasmic microtubules to the SPB (75).

At the onset of meiosis II, Spc72p is lost from the cytoplasmic face of the SPB and replaced by a several meiosis-specific proteins, which together make up the meiosis II outer plaque (76, 113) (Fig. 3). This change in composition converts the function of the cytoplasmic face of the SPB from microtubule nucleation to membrane nucleation. The meiosis II outer plaque is the site of formation of the prospore membrane (52, 100). Early work established a correlation between the number

of SPBs displaying an outer plaque and the number of prospore membranes formed, suggesting that the meiosis II outer plaque is essential for prospore membrane formation (31). More recently, several gene products that are components of this structure have been identified. The first of these was Cnm67p, a coiled-coil protein that is present on the vegetative SPB outer plaque (10, 183). The original study noted that *cnm67* homozygous mutants fail to sporulate, suggesting that Cnm67p is also required for assembly of the meiosis II outer plaque, a prediction that was subsequently confirmed (4, 10). Of the other identified components of the vegetative outer plaque, only Nud1p is also present in outer plaques in meiosis II (76). The remaining genes known to encode components of this structure are meiosis specific in their expression. These are *MPC54*, *MPC70/SPO21*, *SPO74*, and *ADY4* (4, 76, 113).

Figure 3 shows the proposed arrangement of these proteins within the meiosis II outer plaque (76, 113). The large electron density that is in contact with the prospore membrane is composed of the Mpc54, Spo21, and Spo74 proteins. Two of these proteins, Spo21p and Mpc54p, contain predicted coiled-coil domains, while Spo74p is globular (4, 76, 113). By analogy to areas of lower electron density in other parts of the SPB (149), the lower density in the middle of the Mpc54p/Spo21p/Spo74p region is thought to be the coiled-coil regions of the Mpc54 and Spo21 proteins (Fig. 3). Individual deletion of any of these three genes produces a common phenotype: the outer plaque structure is ablated, and no prospore membrane forms (4, 76, 113). Thus, assembly of the outermost layers of the plaque requires all three components, and, as inferred in the earlier studies, this structure is essential for the coalescence of vesicles into a prospore membrane.

The Nud1 and Ady4 proteins are thought to reside on the inner side of the Mpc54p/Spo21p/Spo74p complex (Fig. 3)

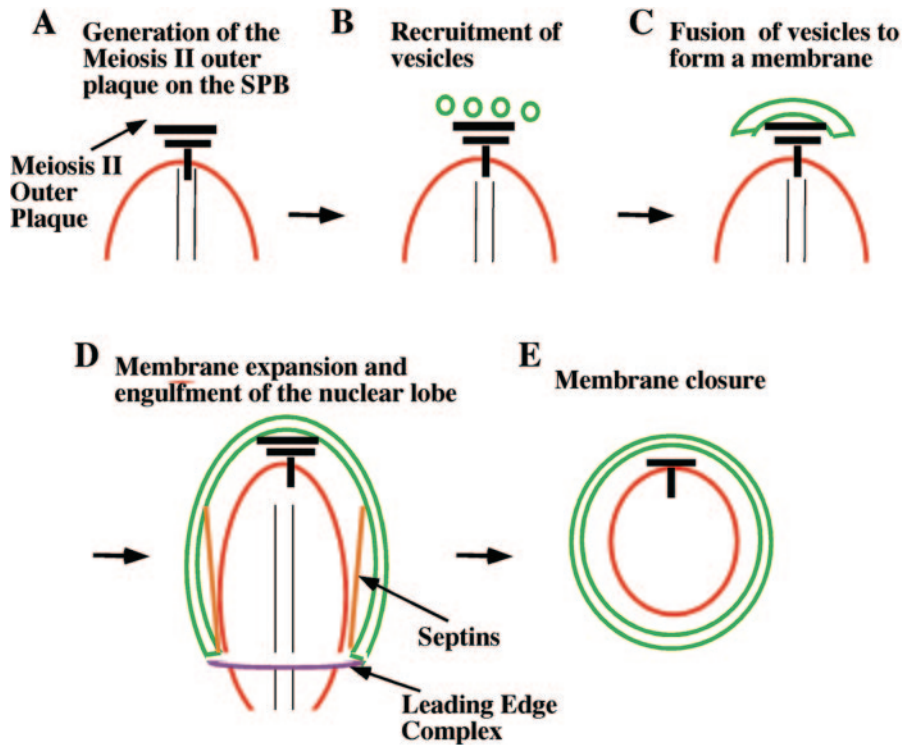


FIG. 2. Stages of prospore membrane growth. (A) As cells enter meiosis II, the meiosis II outer plaque is formed on the cytoplasmic face of the SPB (black bar). (B and C) The meiosis II outer plaque becomes a site for the recruitment and subsequent fusion of secretory vesicles to form a prospore membrane (green). (D) As the prospore membrane expands to engulf a daughter nucleus, its growth is controlled by two membrane-associated complexes: the septins (orange), which form sheets thought to run down the nuclear-proximal side of the prospore membrane, and the leading-edge complex (purple), which forms a ring structure at the membrane lip. (E) Closure of the prospore membrane completes cytokinesis. All three of the prospore membrane-associated complexes, i.e., the septins, leading-edge complex, and meiosis II outer plaque, disassemble at about the time of prospore membrane closure.

(113). Nud1p is also present in vegetative outer plaques, where it serves as the SPB anchor for Spc72p and for components of the mitotic exit network (51). The function of Nud1p at the SPB during sporulation has not been described. Ady4p appears

to be a minor component of the meiotic SPB. Though not essential for formation of the meiosis II outer plaque, it is important for stability of the structure. In *ady4* mutants a fraction of the outer plaques appear to disassemble during meiosis II, leading

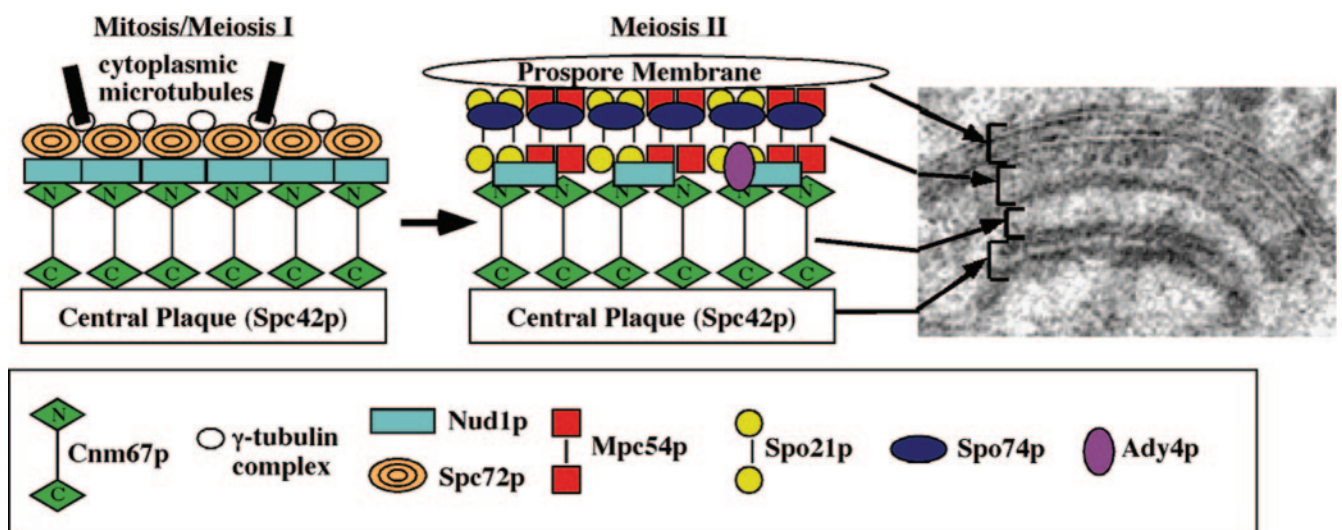


FIG. 3. Organization of proteins within the meiosis II outer plaque. The changes in organization and composition between a mitotic/meiosis I outer plaque and a meiosis II outer plaque are shown in the cartoon. The Spc72p and  $\gamma$ -tubulin complex proteins are removed and replaced with Ady4p, Mpc54p, Spo21p, and Spo74p, leading to a conversion from microtubule to membrane nucleating activity. In the upper right is shown an electron micrograph of a meiosis II outer plaque with associated prospore membrane. The proposed correspondence between the arrangement of proteins in the model and the structure as seen in the electron micrograph is indicated.

to the failure of some prospore membranes to package nuclei and the production of asci with fewer than four spores (113).

The large area of low density between the outer plaque and the central plaque has been demonstrated in vegetative cells to correspond to the coiled-coil domain of Cnm67p (149) (Fig. 3). It is thought that Cnm67p also links the central plaque to the outer plaque in the meiotic SPB (4). The particular importance of the meiosis II outer plaque in allowing membrane formation is shown by the distinct phenotype of *cnm67* mutants during sporulation (4). In *cnm67* cells, no outer plaque structures are seen at the SPB; in fact, even less residual electron density is present than in *mpc54*, *spo21*, or *spo74* mutants. Unlike these other mutants, a fraction of *cnm67* cells do form prospore membranes; however, these membranes are not anchored to the SPB and fail to capture nuclei. This ectopic formation of prospore membranes is proposed to result from the formation of *MPC54/SPO21/SPO74* complexes that are not anchored to the rest of the SPB, an interpretation supported by the observation that ectopic membranes are absent in a *cnm67 spo21* double mutant (4).

The identification of multiple components of the meiosis II outer plaque has led to the demonstration that this structure is essential both for the coalescence of vesicles into a prospore membrane and to anchor that membrane to the nucleus and facilitate nuclear capture. The mechanisms by which this structure binds to the membrane and promotes fusion remain unknown. However, as described in the next section, the critical role of the meiosis II outer plaque in membrane formation allows the cell to regulate spore formation by regulating this modification of the SPB.

**(i) Regulation of SPB assembly by carbon availability.** Sporulation of diploid cells occurs in response to starvation conditions, specifically the absence of nitrogen and the presence of a nonfermentable carbon source such as acetate. In a seminal study, Davidow et al. (31) defined a secondary response to the depletion of the carbon source during the process of sporulation, the formation of a particular form of two-spored ascus or dyad. Using a strain background that was temperature sensitive for progression out of meiotic prophase, the authors found that the strain could be held in prophase by incubation at 37°C and that when released by transfer to lower temperature, it proceeded through meiosis and sporulated. If the strain was held at 37°C for up to 24 h, it formed tetrads upon a shift to lower temperature. However, when the temperature was lowered after arrest for greater than 24 h, the cells predominantly formed dyads. Dissection and analysis of marker genes segregating in the dyads revealed that the spores were viable and haploid. Moreover, for heterozygous markers that were tightly centromere linked, there was always one spore of each genotype. That is, the chromosomes present in the two spores were homologues, not sister chromatids. This segregation pattern distinguished these dyads from other mutants that form dyads of two haploid spores in which centromeric markers segregate randomly or dyads which contain two diploid spores (73, 173). Because of the chromosome segregation pattern, these dyads were termed nonsister dyads (NSDs). An outline of NSD formation is shown in Fig. 4.

Two critical features of NSD formation were elucidated in this study. First, it was shown that the switch from tetrads to NSDs is triggered by the depletion of acetate from the me-

dium. This suggests that NSD formation is a regulated response to carbon depletion. In fact, a subsequent study found that transfer of cells from sporulation medium to water prior to meiosis I could trigger the same phenomenon (158). Second, in cells forming NSDs, the basis for the formation of two spores was shown to be that only two of the four SPBs, one on each of the two meiosis II spindles, are modified on their outer plaque. As a consequence, only two prospore membranes form and, therefore, only two spores. Because one pole from each spindle is modified, the resulting spores will contain centromeres from homologous chromosomes, which separate from each other at the first meiotic division (Fig. 4). Thus, NSD formation is a conservation mechanism that allows a cell that is committed to sporulation to save resources by reducing the number of daughter cells it will form. Moreover, this reduction results from the regulation of SPB modification in response to carbon availability. This regulated change in SPB modification will be referred to as the NSD response.

Subsequent studies identified two mutants that formed NSDs at high frequency even under conditions where acetate was plentiful (33, 115). The first of these mutant genes, *hfd1-1*, has been lost (T. Iino, personal communication). The second, *ady1*, encodes a nucleus-localized protein that triggers NSD formation by causing only two of the four SPBs to be modified, similar to the metabolic response (33). Although the molecular function of *ADY1* remains unknown, it may play a role in SPB modification in response to carbon limitation.

A third mutant condition that causes NSD formation is haplo-insufficiency of outer plaque components. Strains heterozygous for a deletion of *SPO21*, *SPO74*, or *MPC54* form a high percentage of NSDs under conditions where wild-type cells would form tetrads (4, 113, 181; A. M. Neiman, unpublished observations). Additionally, a particular allele of the central plaque component gene *SPC42* has been reported to increase NSD formation (62). Not all mutations affecting the SPB lead to this phenotype, however, as *cnm67* heterozygotes form tetrads and *ady4* homozygotes form random, rather than nonsister, dyads (A. M. Neiman and M. Nickas, unpublished observations) (113). The consistent pattern that emerges from these studies is that decreasing the abundance of any one of the outermost components of the meiosis II outer plaque leads to formation of NSDs.

The studies described above suggested that *SPO21*, *SPO74*, and/or *MPC54* might be a regulatory target of the NSD response. Direct observation of the localization of these three proteins in cells undergoing the NSD response revealed that Spo74p and Spo21p are absent from two of the four spindle poles during meiosis II but that Mpc54p and the other outer plaque components are still found predominantly on all four SPBs (111). Thus, localization of Spo74p and Spo21p to the SPB seems to be regulated in response to carbon availability. Under normal conditions, these two proteins are interdependent for localization to the SPB: Spo74p does not localize to spindle poles in a *spo21* mutant and vice versa, so regulation of either one of the proteins would account for the effects seen (113).

**(ii) Asymmetric control of mother and daughter SPBs during sporulation.** How the localization of Spo21p and Spo74p to the pole is regulated during NSD response remains an open

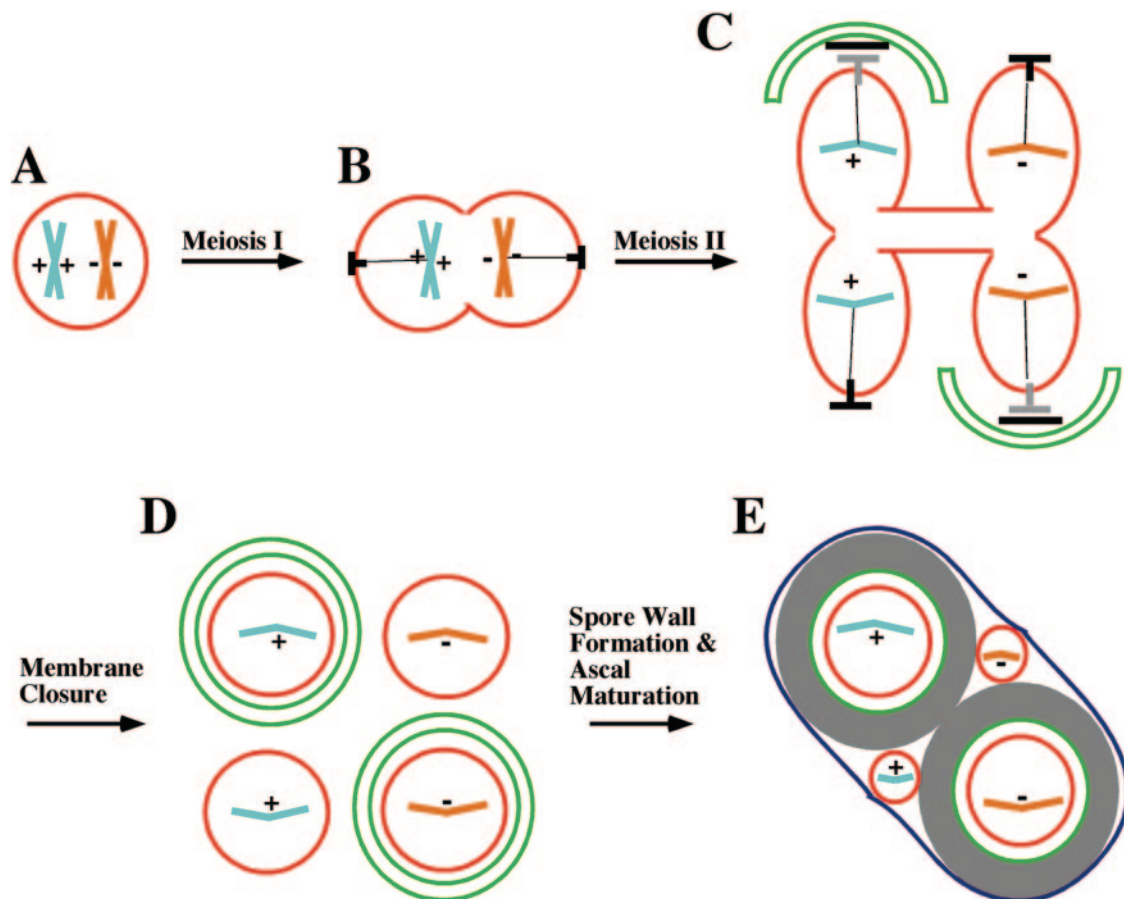


FIG. 4. Nonsister dyad formation. (A to C) During the first meiotic division, homologous chromosomes segregate to opposite poles, and at the second meiotic division, sister chromatids are separated. (C) When sporulated under acetate-depleted conditions, only two of the four SPBs, one from each spindle, form meiosis II outer plaques. (D) As a result, only two prospore membranes are formed and only two nuclei are packaged into spores. (E) The unpackaged nuclei degenerate during ascus maturation, resulting in a two-spored ascus. If a heterozygous centromere-linked marker (indicated by + and -) is followed, an NSD will contain one spore with each allele. The gray  $\mp$  indicates the daughter SPB formed at meiosis II (see text).

question, but some insight into this issue was afforded by the identification of which SPBs these proteins are recruited to (or excluded from) during the NSD response. At the meiosis I/meiosis II transition, the two SPBs present during meiosis I duplicate to form the two poles of each meiosis II spindle so that each meiosis II spindle contains a mother pole and daughter pole (Fig. 4). During mitotic growth, the asymmetric localization of the Kar9 protein to the mother, but not daughter, poles is important for proper orientation of the mitotic spindle and for nuclear segregation into the bud (88, 127). *KAR9* plays no obvious role in sporulation (36), but, by analogy, formation of NSDs could be explained if the mother and daughter poles are distinct in their capacity to form meiosis II outer plaques under NSD conditions.

By using an Spc42 fusion protein that specifically marks the older, mother SPBs (127), it was shown that during the NSD response the Spo21 and Spo74 proteins are found almost exclusively on the daughter SPBs in meiosis II, indicating that NSDs are formed by modification of only the daughter SPBs (Fig. 4) (111). Remarkably, when NSDs are triggered by haploinsufficiency of *spo21*, it is again only the daughter SPBs that get modified (M. Nickas and A. M. Neiman, unpublished ob-

servations). While the molecular mechanisms at work remain to be elucidated, these results allow a more precise description of the phenomenon: the depletion of the carbon source during the process of sporulation leads to a failure to recruit the Spo74 and Spo21 proteins to the mother SPBs. Any model describing these events must account for the fact that simply lowering the amount of these proteins is sufficient to generate the asymmetric use of the mother and daughter SPBs. One proposed explanation is that at the meiosis I/meiosis II transition the daughter SPBs are modified—that is, assemble membrane-organizing outer plaques—before the mother SPBs (111). This may be because the mother SPBs have a preexisting microtubule-nucleating outer plaque that must be rearranged. If the outer plaque components are limiting, the daughter SPBs use up the available pool and so the mother SPBs are not modified. If assembly of the outer plaque structure was highly cooperative, then limitation of any one of the components would cause this effect. This model can account for the haploinsufficient phenotype of *mpc54*, *spo21*, or *spo74* mutants and suggests that limiting expression of *SPO21* and or *SPO74* could be the basis for regulation by carbon depletion. Though appealingly simple, this proposal has yet to be rigorously tested.



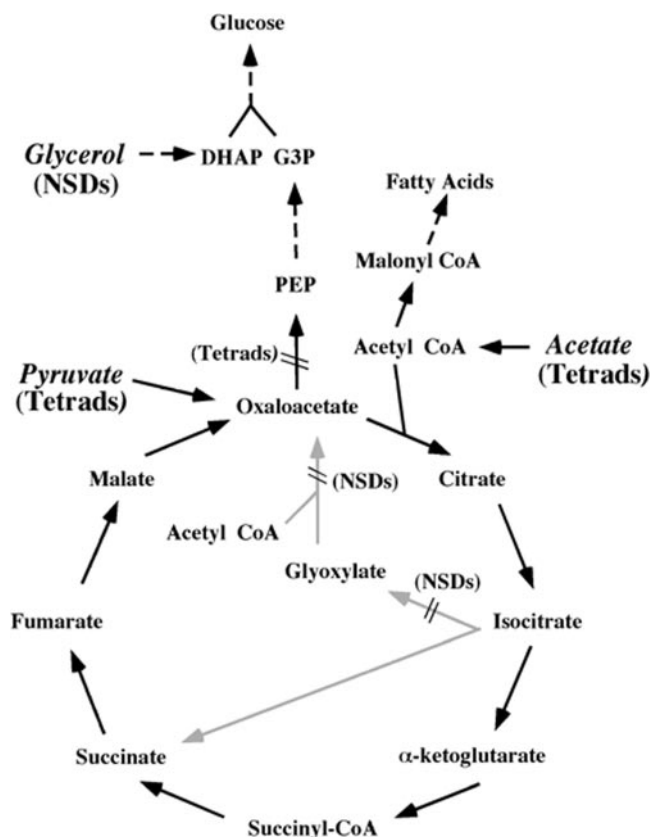


FIG. 5. Pathways of acetate metabolism in sporulating cells. Carbon sources mentioned in the text are shown in italic, and whether cells form tetrads or NSDs when sporulated on those carbon sources is indicated. Positions at which mutations in the glyoxylate cycle or gluconeogenesis lead to NSD or tetrad formation in the presence of acetate are also indicated. Gray arrows denote metabolic reactions unique to the glyoxylate cycle. Dashed lines indicate multiple reactions. DHAP, dihydroxyacetone phosphate. G3P, Glyceraldehyde-3-phosphate. PEP, phosphoenolpyruvate.

Much remains to be learned about the regulatory mechanisms governing assembly of this membrane-organizing center.

**(iii) An intermediary metabolite regulates SPB modification.**

The original description of the NSD response demonstrated that depletion of acetate from the medium triggered this response (31). What remained unclear was whether the cell was responding directly to the disappearance of acetate or perhaps to the absence of some downstream product of acetate metabolism. In sporulation medium, acetate is used by the cell as both the energy source and the source of structural carbon for biosynthesis of macromolecules (39). The metabolic pathways of acetate usage are summarized in Fig. 5. For ATP generation, acetate is converted to acetyl coenzyme A (acetyl-CoA) and oxidized in the tricarboxylic acid (TCA) cycle to generate reducing equivalents for oxidative phosphorylation (157). For biosynthetic purposes, the acetyl-CoA must feed into synthesis of lipids, nucleotides, and polysaccharides (there is no net synthesis of amino acids during sporulation because of the absence of a nitrogen source) (129, 137, 157). Carboxylation of acetyl-CoA to malonyl-CoA provides the building blocks for lipid biosynthesis. However, nucleotide synthesis for DNA rep-

lication and synthesis of the polysaccharide layers of the spore wall require that the acetate first be converted into glucose (129). Gluconeogenesis consumes two molecules of the TCA cycle intermediate oxaloacetate for each glucose molecule produced. Oxaloacetate is also required as a catalytic intermediate in the TCA cycle. To prevent gluconeogenic consumption of oxaloacetate from interfering with oxidation of acetyl-CoA, the yeast cell generates oxaloacetate via the glyoxylate cycle (77, 157) (Fig. 5). The glyoxylate cycle consumes two acetyl-CoA molecules and produces a molecule of succinate without donating electrons to the electron transport chain. Succinate is then oxidized to oxaloacetate and provides the starting point for gluconeogenesis. It should be noted that many of the glyoxylate cycle enzymes are located in different cellular compartments from the analogous TCA cycle enzymes, and it is a cytoplasmic pool of oxaloacetate that is used for gluconeogenesis as opposed to the oxaloacetate that functions in the TCA cycle, which is located in the mitochondrial matrix (59, 87, 90, 96, 159).

If a product of acetate metabolism is used by the cell to detect the availability of the environmental carbon source, then blocks in acetate metabolism might lead to the formation of NSDs even in the presence of acetate. This “sensor metabolite” could be an end product of acetate metabolism, such as glucose or ATP, or an intermediate in acetate usage. Strikingly, when mutants blocked in the glyoxylate cycle or the subsequent conversion of succinate to fumarate were sporulated in acetate, about 50% of the cells formed NSDs, indicating that these cells were undergoing the NSD response (111). These mutations do not effect the TCA cycle or ATP generation, and therefore these results implicate a downstream product of the glyoxylate cycle as the sensor metabolite. By contrast, mutations that block the committed steps of gluconeogenesis did not cause NSD production (111). These results suggest that the sensor metabolite is produced even when gluconeogenesis is blocked. Together these genetic tests map the sensor to one of three metabolites: fumarate, malate, or oxaloacetate (Fig. 5).

Further support for the idea that one of these three molecules is used by the cell as a carbon sensor was provided by changing the carbon source for sporulation. Sporulation on pyruvate as the principal carbon source restored tetrad formation to the glyoxylate cycle mutants. This suggests that the sensor intermediate is produced from pyruvate. To feed gluconeogenesis, pyruvate is converted directly to oxaloacetate, bypassing the need for the glyoxylate cycle (Fig. 5). Thus, these experiments suggest that a cytoplasmic pool of oxaloacetate (or some derivative) is the sensor intermediate. Consistent with this interpretation, when wild-type cells are sporulated on glycerol as the sole carbon source, they form almost exclusively NSDs (111). The conversion of glycerol to glucose, or its conversion to acetyl-CoA for oxidation in the TCA cycle, never requires the generation of cytoplasmic oxaloacetate (Fig. 5). Thus, under these conditions, even though the cell has ample carbon available to meet both its energy and biosynthetic needs, it behaves as though carbon were limiting.

In sum, the NSD response allows the cell to conserve resources when it has committed to meiosis and sporulation but environmental conditions may not support the formation of four daughter cells. As cells sporulate in acetate, increased flux through the glyoxylate pathway generates a increase in the

cytoplasmic pool of oxaloacetate. By a mechanism yet to be explained, the presence of oxaloacetate is measured by the cell as an indicator of the available biosynthetic carbon pool. If sufficient oxaloacetate is present, the cell modifies all four spindle pole bodies and a tetrad is formed. If oxaloacetate is not present, then Spo21p and Spo74p are not recruited to the mother SPBs and NSDs are formed. It is interesting that the NSD/tetrad decision appears to be tuned to an intermediate in acetate metabolism. Studies of sporulation on different carbon sources demonstrated that acetate is the preferred carbon source for sporulation (95). Acetic acid-producing bacteria and yeasts are major components of the microbial flora that grow on broken grapes (43). It is tempting to speculate that in nature *S. cerevisiae* grows in competition with acetate-producing bacteria and, therefore, that acetate is the carbon source most likely to be present at times of nitrogen depletion. Perhaps for this reason *S. cerevisiae* has tuned its sporulation program to respond to acetate levels.

**Initial formation of the prospore membrane.** Once the meiosis II outer plaque has assembled, it becomes a site for the docking of precursor vesicles for the prospore membrane (104). These vesicles are post-Golgi vesicles that carry proteins destined for an extracellular compartment and are analogous to secretory vesicles in vegetative cells (106). Furthermore, the fusion of these vesicles requires many of the same proteins necessary for fusion of secretory vesicles at the plasma membrane (106). In contrast to secretory vesicles in vegetative cells, which are delivered to the periphery along actin cables (132), in sporulating cells vesicles are delivered initially to the spindle poles and then to the growing prospore membrane. Several lines of evidence suggest that actin does not play a critical role in vesicle delivery during sporulation. Mutations of many actin-organizing proteins that disrupt vesicle delivery in vegetative cells, as well as mutations of the *ACT1* gene itself, do not significantly affect sporulation efficiency (34, 36, 92, 182). Moreover, while treatment of sporulating cells with the actin-depolymerizing drug Latrunculin A blocks sporulation, it does not inhibit prospore membrane growth (A. Coluccio and A. M. Neiman, unpublished observations). Taken together, these observations suggest that actin-based transport does not play a critical role in this process. Delivery apparently does not require cytoplasmic microtubules, which are absent by this stage of sporulation due to the changes in the SPB. The mechanism of vesicle delivery to the prospore membrane is unknown and remains an important challenge in our understanding of the process.

In addition to this change in vesicle delivery, the fusion of vesicles with the prospore membrane has different genetic requirements than fusion of vesicles with the plasma membrane. After secretory vesicles are delivered to the plasma membrane, the fusion of the vesicle membrane and the plasma membrane is then mediated by a heterotrimeric SNARE complex (141). Analogous SNARE complexes function for each fusion event in the secretory pathway, and much of the specificity of vesicle/target membrane interactions that is required for proper trafficking through the secretory pathway is thought to be based on cognate interactions between SNARE proteins (123, 124, 142). SNARE complexes are based on the formation of a four-helix bundle between different subunits of the hetero-oligomer (162). Three of the helices are provided by proteins associated with

the target membrane (the t-SNAREs) and one is provided by a protein on the surface of the vesicle (the v-SNARE). The v-SNARE and at least one of the t-SNARE proteins are transmembrane proteins, and assembly of the bundle drives these transmembrane domains together, possibly leading directly to the fusion of the two lipid bilayers (180).

For fusion of vesicles with the plasma membrane, the t-SNARE consists of the Sso1 or Sso2 protein, which are redundant for this purpose, in partnership with the Sec9 protein (1, 11). Sso1p or Sso2p provides the transmembrane domain and one helix of the oligomer, whereas Sec9p is a peripheral membrane protein that provides two helices. The v-SNARE is encoded by a second redundant pair of genes, *SNC1* and *SNC2* (131). The first genetic distinction between vesicle fusion at the plasma membrane or prospore membrane was the finding that the t-SNARE *SEC9* was dispensable for sporulation (106). During sporulation, Sec9p is replaced by the related, sporulation-specific Spo20 protein. Although Spo20p and Sec9p are homologous, they are largely distinct in their sites of action. Ectopic expression of *SPO20* in vegetative cells cannot rescue the growth defect of a *sec9-ts* mutant, nor can expression of *SEC9* under the *SPO20* promoter rescue the sporulation defect in a *spo20* cell (106). However, there is some overlap of function in sporulation, as a *spo20* single mutant produces abnormal prospore membranes whereas a *spo20 sec9-ts* mutant displays no prospore membranes at all, suggesting that *SEC9* is capable of supporting at least abortive prospore membrane assembly (106). In chimera studies, the major determinant of sporulation function of Spo20p mapped to an amino-terminal region outside of the domain in which it is homologous to Sec9p (107).

Also indicative of the close relationship between Sec9p and Spo20p, both bind in vitro to the Sso1 protein and can form ternary complexes with Sso1p and Snc2p with similar efficiency (107). Indirect immunofluorescence using polyclonal antibodies to the Sso and Snc proteins demonstrated that these SNAREs localize to the prospore membrane. Based on these observations, the change in fusion functions between the plasma membrane and the prospore membrane was proposed to be a switch in one subunit of the SNARE complex, an exchange between Spo20p and Sec9p (107).

However, the requirements for vesicle fusion at the prospore membrane are more complicated than this initial picture, as there is further specialization of the SNARE complex (Fig. 6). A study of *sso1* mutants revealed that an *sso1* single mutant, which has no strong vegetative phenotype, is completely defective in sporulation (66). The *sso1* phenotype could not be rescued by overexpression of *SSO2*. A subsequent chimera study suggested that the specificity of *SSO1* for sporulation lay partly in its amino-terminal region (the only region of the proteins with significant sequence divergence) and, surprisingly, in the 3' untranslated region of the *SSO1* gene (119). The 3' untranslated region did not seem to have a significant effect on the levels of the Sso1 protein, so the basis for this effect remains mysterious. An examination of the sporulation defect in *sso1* cells revealed that prospore membranes are absent in the mutant, indicating that *SSO1* is required for the fusion of vesicles to form a prospore membrane (H. Nakanishi and A. M. Neiman, unpublished data). Thus, Sso1p and Sso2p are redundant for

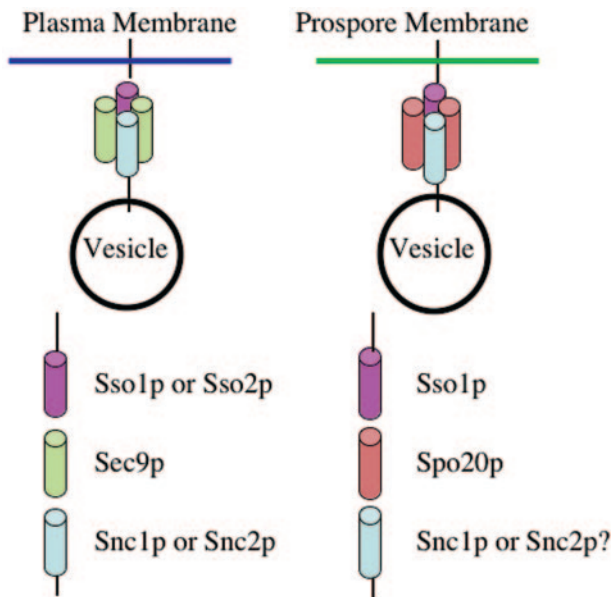


FIG. 6. Specific SNARE complexes mediate fusion with the plasma membrane and prospore membrane. Secretory vesicle fusion with the plasma membrane is mediated by a heterotrimer consisting of Sso1p or Sso2p, Sec9p, and Snc1p or Snc2p (left panel). At the prospore membrane the fusion of secretory vesicles requires Sso1p, Spo20p (although Sec9p can substitute to a limited extent), and probably Snc1p or Snc2p. The roles of Snc1p and Snc2p in this process are inferred from the protein localization but have not been directly demonstrated (107).

fusion at the plasma membrane, but only Sso1p can function during fusion at the prospore membrane.

Studies of the lipid-modifying enzyme phospholipase D suggest that changes in lipid composition are also important for assembly of the prospore membrane (143). This enzyme, which hydrolyzes the choline head group of phosphatidylcholine to produce phosphatidic acid, is encoded by the *SPO14* gene, and the Spo14 protein is localized to the prospore membrane during sporulation (140, 143). Originally identified on the basis of its sporulation defect, this gene is transcribed in both vegetative and sporulating cells, but deletion of the gene has significant phenotypes only during sporulation. Deletion of *SPO14* leads to a failure to form prospore membranes (143). Prospore membranes are also absent in cells expressing a point mutation in *SPO14* that encodes a catalytically inactive protein, indicating that enzyme activity is critical for function (143). In the presence of 1-butanol, phospholipase D will convert phosphatidylcholine into choline and phosphatidylbutanol rather than phosphatidic acid. Addition of 1-butanol, but not 2-butanol, to sporulation medium blocks sporulation (144). Together these results indicate that it is the production of phosphatidic acid rather than the turnover of phosphatidylcholine which is in some way critical for membrane assembly (144).

One role for Spo14-generated phosphatidic acid in prospore membrane formation may be in enhancing the activity of Spo20p. As noted earlier, the amino-terminal domain of Spo20p contains a region crucial for the function of this protein at the prospore membrane (103). This crucial "activating" function is a lipid binding domain that can bind to phosphatidic acid and is essential for localization of the Spo20p *in vivo* (103). More-

over, translocation of Spo14p to the plasma membrane in vegetative cells, an arrangement analogous to that on the prospore membrane, allows Spo20p to replace Sec9p for fusion of vesicles at the plasma membrane (28). This is true even for derivatives of Spo20p that lack the phosphatidic acid binding domain. However, the phenotype of a *spo14* mutant is more extreme than that of a *spo20* mutant (106, 143). Therefore, although *SPO14* may be important for the function of Spo20p-containing SNAREs, phosphatidic acid may play additional roles in prospore membrane assembly as well. Deciphering the interaction between SNARE specificity and lipid composition during fusion at the prospore membrane is an important challenge that will shed light more generally on how the specificity of fusion is regulated in the secretory pathway.

**Membrane expansion.** Once an initial membrane cap has been established by fusion of vesicles on the SPB, the membrane grows to engulf the nuclear lobe by continued vesicle fusion. The growth of the membrane must be controlled so that it obtains the proper shape and dimensions to enclose a nucleus. These events require several additional genes that are not necessary for the membrane fusion per se (101, 133, 163). While the functions of most of these genes are not known, two membrane-associated complexes that are implicated in the control of membrane expansion have been identified: the septins and the leading-edge complex.

**(i) Septins.** Septins are a conserved family of filament-forming proteins (72). In vegetative cells, five distinct septin proteins, Cdc3, Cdc10, Cdc11, Cdc12, and Sep7, form a ring structure at the bud neck (48, 89). The septin ring plays a variety of important roles as a scaffold for the organization of complexes involved in cytokinesis, cell wall deposition, and cell cycle regulation (48, 89). In addition, the septin ring appears to form a diffusion barrier that aids in differentiating the bud from the mother cell (5). In sporulating cells, septin rings on the cell plasma membrane are disassembled and the septins relocalized to the forming prospore membrane (41).

At least three and possibly all five of the septins found in septin rings are localized to the prospore membrane, and two of the genes, *CDC3* and *CDC10*, are strongly upregulated by Ndt80p in mid-sporulation (25, 68, 89). In addition, two sporulation-specific septin proteins, Spr3p and Spr28p, are also found in the prospore membrane-associated septin structures. Several lines of evidence suggest that the organization of the septins on the prospore membrane is distinct from that in the septin ring. First, while the septin ring in vegetative cells is a relatively static structure, during sporulation the septins move as the prospore membrane expands (35, 41). Second, their arrangement on the prospore membrane is different than that at the bud neck. By immunofluorescence, the septins appear first to be organized as rings near the SPB when prospore membranes are small and then to expand into a pair of bars that run parallel to the long axis of the prospore membrane as the membrane expands (Fig. 2) (41). Three-dimensional reconstructions reveal these bars to be a pair of sheets running on opposite sides of the membrane (163). Moreover, these sheets contain the Spr3 and Spr28 proteins, which cannot be incorporated into the septin ring when expressed in vegetative cells (41; H. Tachikawa, personal communication). Finally, in vegetative cells mutation of any one septin (all except for *SEP7*) leads to the disappearance of bud neck rings. By con-

trast, mutation of individual septins, or even an *spr3 spr28 cdc10* triple mutation, has no strong effect on the ability of the other septins to organize on the prospore membrane (35, 41; A. M. Neiman, unpublished observations). Thus, the organization of these filaments may be distinct in sporulating cells.

What is the role of the septins in spore formation? Not only does mutation of one or multiple septins during sporulation not disrupt the localization of the other septins, the mutants do not have a strong sporulation phenotype (35, 41; A. M. Neiman, unpublished observations). This suggests that if, as implied by their localization pattern, the septins play an important role in spore formation, then they must function redundantly in this process. Analogous to their role in vegetative cells, the septins may help to localize other proteins to specific areas of the prospore membrane, for example, the Gip1 protein. *GIP1* encodes a sporulation-specific regulatory subunit of the Glc7 protein phosphatase (170). During sporulation, Gip1p and Glc7p colocalize with the septins throughout prospore membrane formation (163). Mutation of *gip1* or alleles of *glc7* that cannot interact with Gip1p leads to the failure of septins to localize to the prospore membrane, indicating that these proteins both organize and may be part of the septin complex (163).

Because *gip1* mutants lack organized septins, the phenotype of the mutant may provide some insight into the role of septins in spore formation. In fact, prospore membrane formation is largely normal in *gip1* mutants. Four membranes are formed per cell, and these membranes, though they appear slightly smaller than in wild-type cells, capture nuclei efficiently (163). The most striking defect in *gip1* mutants appears to be in triggering the assembly of the spore wall in response to closure of the prospore membrane. Whether this phenotype is due to the absence of organized septins or the absence of an independent function of *GIP1* remains to be determined, but the phenotype of *gip1* raises the possibility that septins may be involved in monitoring membrane growth and closure rather than in controlling membrane growth.

(ii) **The leading-edge complex.** Electron micrographs reveal an electron-dense coat, termed the leading-edge complex, located at the lip of each growing prospore membrane (21, 25, 76, 101). Three components of this coat have been identified: Ssp1p, Ady3p, and Don1p (101, 112). Localization of these proteins by immunofluorescence and green fluorescent protein tagging confirms that they form a ring structure at the mouth of the growing prospore membrane (101, 112). *SSP1* and *DON1* were originally identified on the basis of their sporulation-specific expression (101, 102). *ADY3*, by contrast, was identified by its ability to bind to meiotic SPB components in both two-hybrid and copurification studies (63, 101, 171). At early times of prospore membrane assembly, Ady3p is found on the SPB, whereas the other two subunits are localized in a punctate pattern in the cytosol, suggesting an association with transport vesicles (101). As the vesicles coalesce into a membrane, all three proteins can be found at the SPB. It has been suggested that this pattern of localization indicates a role for these proteins in the recruitment of vesicles to the SPB (101). As the prospore membrane expands, the localization patterns of the three proteins resolve into a ring structure that remains associated with the leading edge of the prospore membrane (Fig. 2). This structure is distinct from the septin sheets de-

scribed above. Double-labeling studies indicate that the septins run backwards from the leading edge towards the SPB (101). However, the assemblies of the two structures appear to be independent: septin localization is retained in leading-edge complex mutants, and the leading-edge complex is unaffected in a *gip1* mutant where septin localization is lost (163).

The organization of Ssp1p, Don1p, and Ady3p within the leading-edge complex is not known, but localization dependence studies suggest a stratified arrangement (101, 112). Deletion of *DON1* does not affect the localization of the other two proteins. Deletion of *ADY3* causes the delocalization of Don1p, but Ssp1p remains in a ring at the prospore membrane lip. Deletion of *SSP1* causes the disappearance of all three proteins. Therefore, Ssp1p may be the most membrane-proximal component, serving to connect Ady3p to the leading edge, and Ady3p in turn anchors Don1p.

Mirroring this stratified arrangement, the sporulation phenotypes of the three mutants are progressively severe. Mutation of *DON1* produces no obvious sporulation defect (76). As suggested by its name, loss of *ADY3* (*Accumulates DYads*) leads to an increase in asci with fewer than four spores (133). The spores present in these dyads are haploid and are packaged randomly with respect to centromere-linked markers, unlike the nonsister dyads described above (101, 112). In fact, in addition to dyads, the culture contains significant numbers of asci with no spores, as well as monads and triads. Surprisingly, the defect in spore formation is caused by a failure of individual spores in the *ady3* mutant to form mature spore walls. Cells lacking *ADY3* show no apparent defects in prospore membrane formation or nuclear capture by the prospore membrane, but a significant fraction of these prospores fail to elaborate a spore wall (112).

Mutation of *SSP1* causes the most severe phenotype: a complete block to sporulation. In these cells, prospore membranes are still formed, but they are grossly abnormal (101). The membranes are tubular rather than round and appear to be adherent to the nuclear envelope. The membranes also occasionally grow in the wrong direction, resulting in a failure to capture nuclei. As in wild-type cells, four membranes are formed; however, in *ssp1* cells these membranes often fragment. The *ssp1* phenotype demonstrates the necessity for the leading-edge complex (or at least Ssp1p) for proper membrane growth.

While the phenotype of *ssp1* mutants is dramatic, it is unclear how loss of *SSP1* results in these abnormalities. One proposed explanation is that in the absence of the leading-edge complex, secretory vesicles may fuse only with the outer membrane of the prospore membrane, creating a disproportionate force that pushes the prospore membrane onto the nuclear envelope (101). An alternative possibility is that the leading-edge complex performs functions analogous to the septin ring in budding cells. For example, it could function as a diffusion barrier that distinguishes the inner prospore membrane, which will ultimately become the plasma membrane of the spore, from the outer prospore membrane, which will ultimately be degraded (see below). Although no protein whose localization is asymmetric between these two regions of the prospore membrane has been described, it is tempting to speculate that such proteins exist and that the loss of this asymmetry in *ssp1* mu-

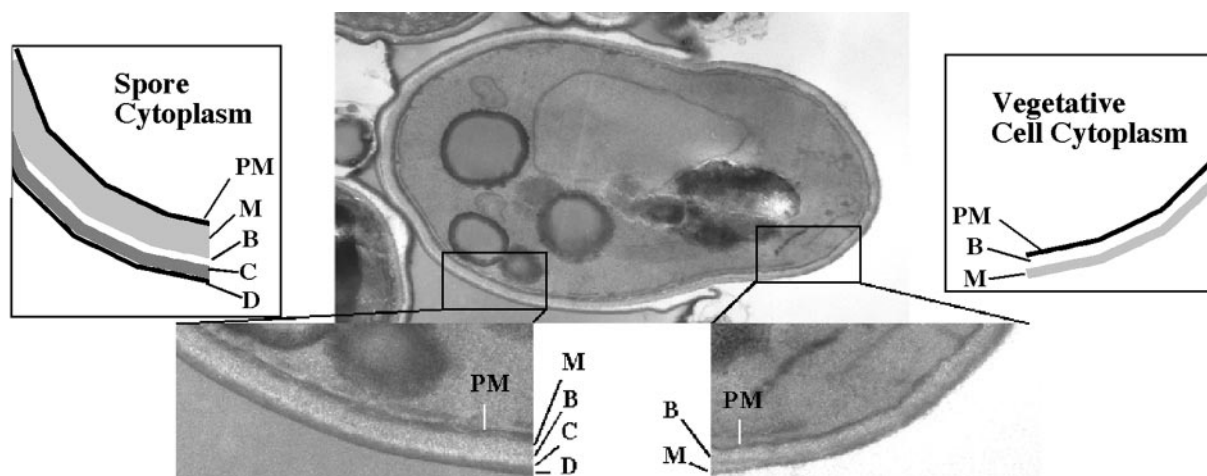


FIG. 7. Comparison of the spore wall and the vegetative cell wall. An electron micrograph of a germinating ascospore is shown. On the left side, the cell is surrounded by spore wall with its four layers, mannan, beta-glucan, chitosan, and dityrosine (indicated by M, B, C, and D, respectively, in the close-up and in the cartoon). On the right side, the tip of the germinating cell is bounded by vegetative cell wall with its predominant beta-glucan and mannan layers (indicated by B and M). The mannan layer of the cell wall appears to be continuous with that of the spore wall. PM, plasma membrane.

tants might lead to some of the membrane defects seen in the mutant.

The position of the leading-edge complex at the mouth of the prospore membrane also raises the possibility that it could play a role in the segregation of organelles into the spore or even in the closure of the membrane. A failure in one of these processes could account for the subsequent spore wall defects seen in *ady3* mutants (112). With respect to closure, the leading-edge complex might function in a fashion similar to that of the septins at the bud neck in defining the site of closure. Alternatively, the leading-edge complex might take a more active role in causing the membrane to constrict and close. Whether the leading-edge complex is dynamic or static is an important area for future study.

**Membrane closure.** The closure of the prospore membrane is a cytokinetic event, the molecular mechanism of which is unknown. After closure, the cytoplasms of the prospores are distinct from each other and from the mother cell (now ascus) cytoplasm. In electron micrographs of wild-type cells, deposition of spore wall material (identifiable as an expansion of the prospore membrane lumen) is seen only in cells in which the prospore membrane is closed, suggesting that closure necessarily precedes the onset of spore wall formation. This inference is supported by analysis of the *gip1* mutant (163). In these cells, prospore membrane formation proceeds normally up to and perhaps including closure, but subsequent spore wall deposition does not occur. This suggests that *gip1* mutants are defective either in the process of closure or in a signaling pathway which monitors closure and triggers subsequent events. Although roles for *GIP1* (and possibly septins) and the leading-edge complex in membrane closure have been postulated, this remains a largely unexplored area. Several recent studies have described a role for vesicle fusion in the terminal stages of cytokinesis following mitosis in both yeast and animal cells (20, 65, 153, 179). Understanding prospore membrane closure may, therefore, provide more general insights into cell division.

### Spore Wall Assembly

The closure of the prospore membrane completes the act of cell division. However, proper differentiation into stress-resistant ascospores requires assembly of the spore wall. As with prospore membrane formation, this is a *de novo* assembly process. That is, spore wall assembly begins in the lumen between the two membranes derived from the prospore membrane, where there is no preexisting wall structure to serve as a template. Thus, this process is an excellent one with which to explore the assembly of a complex extracellular structure.

**Description of the spore wall.** The spore wall is a more extensive structure than the vegetative cell wall. For comparison, Fig. 7 shows an electron micrograph of a germinating spore, allowing both the spore wall and vegetative cell wall to be seen on one cell. The vegetative wall consists of two major layers, an inner layer (closer to the plasma membrane) consisting primarily of beta-glucan (chains of beta-1,3-linked glucose) with some chitin (beta-1,4-linked *N*-acetylglucosamine) and an outer "mannan" layer consisting of proteins that have been heavily N and O glycosylated with primarily mannose side chains (74). By contrast, the spore wall consists of four layers (156). The inner two layers consist predominantly of mannan and beta-1,3-glucans, although their order is reversed with respect to the plasma membrane; the mannan is interior to the beta-glucan (78). The outer two layers are specific to the spore. Immediately outside of the beta-glucan is a layer of chitosan (beta-1,4-linked glucosamine) (16). Finally, coating the chitosan is a thin layer that consists in large part of dityrosine molecules (18). Unlike the dityrosine found in the extracellular matrices of metazoans (6, 71, 81, 108), the dityrosine in the spore wall is not created by cross-linking between peptides but rather is synthesized directly from the amino acid tyrosine (14, 17). It is these outer spore wall layers that confer upon the spore much of its resistance to environmental stress (13).

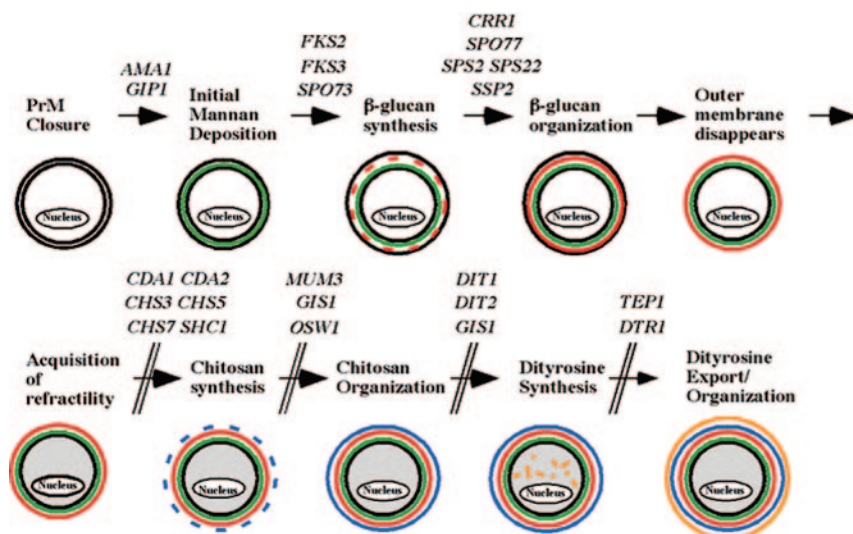


FIG. 8. A pathway of spore wall assembly. The steps in assembly of the spore wall are shown. Genes shown to be required for specific steps are indicated (13, 24, 28, 34, 42, 55, 94, 121, 145–147, 163, 168). Adapted from reference 27.

In wild-type cells, formation of the spore wall begins after prospore membrane closure. The layers of the spore wall appear to be laid down in a temporal order, beginning with the innermost mannan layer and moving outward to beta-glucan, then chitosan, and finally dityrosine (163). These observations suggest that feedback mechanisms must exist to coordinate the completion of one layer with the beginning of synthesis of the next layer.

**A pathway of spore wall assembly.** Through the analysis of mutants defective in construction of the spore wall, the outlines of the assembly process are beginning to emerge. Two general types of mutations blocking spore wall assembly have been described. Mutation of some genes results in heterotypic spore wall defects, in which different spores within the same ascus display different defects, whereas other mutations cause a more uniform arrest in assembly (27, 45, 79, 172). An analysis of the arrest points in a collection of such mutants led to the definition of the following stages of assembly (Fig. 8) (29).

(i) **Initiation.** As described above, the *GPI1* gene appears to be required for the initiation of spore wall formation in response to membrane closure. Mutation of a second gene, *AMA1*, displays a block to spore wall synthesis similar to that of *gip1*. However, *ama1* lacks the septin defects seen in *gip1* mutants. *AMA1* encodes a meiosis-specific regulatory subunit of the anaphase-promoting complex, an E3 ubiquitin ligase (30). Recent studies suggest that Ama1p is regulated so that it activates the anaphase-promoting complex only late in meiosis II (114, 125). These data suggest that *AMA1* may be involved in coordinating the onset of spore wall formation with the completion of meiosis II.

(ii) **Formation of the mannan and beta-glucan layers.** Following initiation, the first stage of spore wall formation is a deposition of material into the lumen between the two membranes formed from the prospore membrane. This expansion apparently begins with the deposition of mannoproteins, as suggested by the phenotype of a *spo73* mutant, which exhibits expansion of the lumen without production of beta-glucans (27).

Assembly of the beta-1,3-glucan layer requires two elements: a beta-glucan synthase that is located in the spore plasma membrane to synthesize and extrude the initial beta-glucan chains into the lumen between the spore plasma membrane and outer membrane (the two membranes derived from the prospore membrane) and wall proteins in this luminal space to “stitch and weave” these initial chains into the properly assembled wall. There are three genes in *S. cerevisiae* that encode catalytic subunits of the synthase. *FKS1* encodes the primary synthase during vegetative growth (61, 94). *GSC2/FKS2* seems to encode the predominant synthase for assembly of the spore wall, although *FKS3* may contribute some function as well (34, 94). Several genes that appear to be important for proper assembly of the beta-glucan layer have been identified, including *SPO77*, *SPS2*, *SPS22*, *CRR1*, and *SSP2* (27, 49, 147). It is of note in this regard that some of these genes are sporulation-specific members of sequence families that are involved in cell wall synthesis in vegetative cells. For example, *SPS2* and *SPS22* are related to *ECM33* and *PST1*, while *CRR1* is similar to the glucohydrolase genes *CRH1* and *CRH2* (27, 49, 122). Additionally, there are sporulation-specific members of the *GAS* and *PIR* families of cell wall proteins (25). These sporulation-specific proteins may play roles in the assembly of mannan or beta-glucan layers of the spore wall that are analogous to those played by their counterparts in construction of the vegetative cell wall.

Two other events occur during this stage of spore wall formation that are linked with the completion of beta-glucan layer assembly. First, during this phase the outer membrane derived from the prospore membrane disappears. Second, the spores acquire refractility in the light microscope. Mutants that are blocked in the early stages of spore wall synthesis appear by light microscopy not to form spores, whereas those affected in later steps do make visible spores. However, these spores are not as resistant to environmental challenge as wild-type spores.

(iii) **Formation of the chitosan layer.** Once the beta-glucan layer is completed, the spore initiates synthesis of the chitosan layer. The chitosan polymer itself is assembled similarly to

beta-1,3-glucan in that a spore plasma membrane-associated synthase, in this case Chs3p, binds nucleotide sugars in the cytosol, couples them, and extrudes the polymer through the spore plasma membrane (53). In vegetative cells, Chs3p is the chitin synthase responsible for the chitin ring around the bud scar. During sporulation, Chs3p again synthesizes chitin (beta-1,4-*N*-acetylglucosamine), but two sporulation-specific chitin deacetylases are present in the spore wall, Cda1p and Cda2p, which deacetylate the chitin, converting it to chitosan (i.e., beta-1,4-glucosamine) (23, 97, 121). The conversion of chitin to chitosan is essential for organization of the outer spore wall layers (24).

In vegetative cells, the activity of Chs3p is controlled both by allosteric regulation of Chs3p and by regulated delivery of Chs3p to the surface (32, 145, 167, 168, 186). Several of the proteins involved in regulated delivery are also important for formation of the chitosan layer during sporulation (34, 168). However, at least one of the allosteric regulators, Chs4p, is replaced during sporulation by a related protein, Shc1p, and other sporulation-specific functions may regulate Chs3p localization during sporulation (see below) (64, 146).

As with beta-glucan synthesis, once the polymer is synthesized by Chs3p it must be properly assembled into the wall. At least two genes, *MUM3* and *OSW1*, that have mutant phenotypes consistent with a role in chitosan layer assembly have been identified (27). While the molecular function of these genes is not known, *MUM3* encodes a protein with homology to acyl transferases, consistent with a possible catalytic role in wall assembly (109).

During assembly of the chitosan layer, interspore bridges are formed (29). Chitosan not only surrounds each spore but also branches, creating bridges that connect adjacent spores of a tetrad. These bridges serve to physically connect spores of a tetrad together even when the overlying ascus is removed.

**(iv) Formation of the dityrosine layer.** Unlike the underlying layers, the dityrosine layer is composed primarily neither of polysaccharide nor of protein. Approximately 50% of the mass of this layer is composed of nonpeptide *N,N*-bisformyl-dityrosine (15). The remaining constituents are not well characterized, nor is it known how the dityrosine monomers are coupled to the wall.

The dityrosine molecules are synthesized in the cytoplasm of the spore by the action of the *DIT1* and *DIT2* gene products (14). These two mid-late genes are divergently transcribed from a common promoter and appear to be induced only after closure of the prospore membrane, as shown by their absence of expression in a *gip1* mutant and by the spore-autonomous phenotype of heterozygous *dit1* or *dit2* mutants (13, 163). The two enzymes act sequentially. Dit1p acts as a formyltransferase that modifies the amino group on L-tyrosine, and Dit2p is a member of the cytochrome P450 family that couples the benzyl rings of two *N*-formyl-tyrosine molecules (14). Both enzymes are cytoplasmic, so the newly formed dityrosine must be exported to the spore wall for assembly. This is achieved through the action of a sporulation-specific ATP-dependent transporter, Dtr1p, localized in the prospore membrane (42). The expression of these three genes, *DIT1*, *DIT2*, and *DTR1*, in vegetative cells is sufficient to allow the production and export of dityrosine (42). The dityrosine is not incorporated into the cell wall under these conditions, presumably due to the ab-

sence of the appropriate enzymes and/or substrates to allow its incorporation.

As with the interior layers, assembly of the dityrosine must involve extracellular enzymes that assemble it into a polymerized structure. The identities and activities of these enzymes remain to be determined. Other mutants, such as *gis1* and *tep1* mutants, that have defects specifically in the dityrosine layer of the spore wall have been identified (27, 55). However, these gene products are nuclear and cytoplasmic, respectively, and so their effects on coupling of dityrosine into the spore wall must be indirect. Homologues of the *DIT1*, *DIT2*, and *DTR1* genes are found in a variety of other fungi, including pathogens such as *Candida albicans* and *Coccidioides immitis*. In *C. albicans*, dityrosine has been identified in the cell wall (154). Given the importance of dityrosine in conferring stress resistance to *S. cerevisiae* spores, it will be of interest to learn if these genes play a role in allowing pathogenic fungi to survive in the host.

**Regulating spore wall assembly.** The events of assembly described above are inferred largely from the phenotypes of mutants that are blocked at specific points in the process. As noted, several mutations that produce heterogeneous defects in the spore wall have been described (27, 45, 79, 161, 172, 177). These include the protein kinases Cak1p, Mps1p, Smk1p, and Sps1p as well as an integral membrane protein, Spo75p. The heterogeneous phenotypes of these mutants raise the possibility that these gene products are involved in the coordination of the different stages of assembly. Certainly, the fact that many of these genes encode protein kinases would be consistent with roles for these gene products in signaling pathways.

The two best-characterized of these are Sps1p, which is a member of the Ste20/Pak family of kinases, and Smk1p, a mitogen-activated protein (MAP) kinase (45, 79). Most characterized MAP kinases are regulated by a cascade involving upstream kinases from the MEK and MEKK kinase families. In the yeast mating response, the Sps1-related kinase Ste20p triggers a cascade that ultimately leads to the activation of the Smk1-related kinase Fus3p (54, 84). Therefore, it was originally suggested that Sps1p and Smk1p might be part of a similar cascade (45, 79). In the interim, however, it has become clear that no sporulation-specific MEK or MEKK homologues are present in *S. cerevisiae* to link Sps1p and Smk1p. Moreover, *SPS1* is not required for the activation of Smk1p (148). Thus, Sps1p does not seem to trigger a MAP kinase cascade.

A recent report indicates that Sps1p plays a role in the trafficking of both the Gsc2p beta-glucan synthase and the Chs3p chitin synthase to the spore plasma membrane. In wild-type spores, both proteins are found uniformly around the spore periphery, but in *sps1* mutants these proteins accumulate in a punctate pattern within the spore (64). During sporulation, the activity of these two synthases may be regulated, at least in part, by localization, and Sps1p may be involved in coordinating the delivery of these enzymes to the spore surface. Importantly, *smk1* mutants had no effect on delivery of Chs3p, providing further evidence that Sps1p and Smk1p have distinct functions (64).

Deletion of *smk1* produces a heterogeneous spore wall phenotype (79). However, an elegant genetic study demonstrated that hypomorphic forms of *smk1* arrest more uniformly, with the precise stage of arrest varying with the level of activity of the protein (176). This is consistent with a model in which Smk1p

activity is required at different stages to move spore wall assembly in an ordered fashion (176). More recently, Smk1p has been found to bind to the beta-glucan synthase Gsc2p (58). Surprisingly, Gsc2p activity is increased in an *smk1* mutant, suggesting that Smk1p inhibits Gsc2p. Reduction of beta-glucan synthesis in the *smk1* mutant by deletion of *GSC2* restores chitosan deposition, suggesting that active beta-glucan synthesis might inhibit Chs3p activity or delivery. Thus, Smk1p may regulate later stages of spore wall assembly indirectly through Gsc2p (58).

Like other MAP kinases, Smk1p is subject to two regulatory phosphorylations, one on a threonine and one on a tyrosine, both of which are located in the same loop in the enzyme structure, and these phosphorylations are required for Smk1p activity (148). Thus, there must be an activating kinase for Smk1p. Hypomorphic mutations of the essential *CAK1* or *MPS1* genes produce spore wall defects similar to those seen in *smk1* mutants, although unlike Smk1p, both of these kinases have distinct functions earlier in sporulation (19, 148, 150, 161, 177). In vegetative cells, *CAK1* is required for phosphorylation of Cdc28p and other cyclin-dependent kinases on a threonine in an analogous position to the activating phosphorylation sites in Smk1p (37, 69, 117, 184). Smk1p is not phosphorylated in a *cak1* hypomorphic mutant (148). Additionally, Mps1p is a mixed-specificity kinase capable of phosphorylating both threonine and tyrosine residues (83). It is tempting to speculate that one or both of these kinases are responsible for activating Smk1p, although direct tests for Cak1p phosphorylation of Smk1p in vitro were not successful (148). The kinases activating the Smk1p MAP kinase remain to be determined.

### S. *CEREVISIAE* AS A MODEL FOR SPORULATION IN OTHER ASCOMYCETES

Many fungi form asexual (i.e., mitotic) spores of various types (e.g., conidiospores in *Aspergillus* or chlamydospores in *Candida*). The formation of these asexual spores proceeds by different mechanisms than those described here. However, the process of meiotic or ascospore formation appears to be cytologically similar in all ascomycetes (47, 135, 164, 165). Thus, the molecular mechanisms defined for *S. cerevisiae* are likely to be conserved in other species.

The best-studied example of sporulation in another ascomycete is in the fission yeast *Schizosaccharomyces pombe* (reviewed in reference 151). In this yeast, the sporulation process is very similar to that described for *S. cerevisiae*. Spore formation begins with a modification of the SPBs and the subsequent creation of novel membranes, termed forespore membranes. Each forespore membrane engulfs a haploid nucleus, and, after closure, spore walls are assembled. Again the mother cell becomes a protective ascus around the spores.

Despite these similarities, however, conservation of sporulation functions at the level of protein sequence is surprisingly low. For example, although *S. pombe* analogues of the meiotic SPB and leading-edge complex proteins have been identified (60, 116), there is no sequence identity between the proteins in the two yeasts. Comparison of transcriptional microarray studies with *S. cerevisiae* and *S. pombe* indicates that relatively few of the sporulation-induced genes in *S. cerevisiae* have homologues whose expression is similarly induced during sporula-

tion in fission yeast (93). Those genes for which their homologues display conserved regulation are associated predominantly with spore wall assembly and include *SPO71*, *SPO73*, *AMA1*, the septin genes, and the *SPS2* and *SPS22* genes. Of these, only the *S. pombe* knockout of the *SPS2/SPS22* homologue, *meu10*, has been described. Encouragingly, it produces a phenotype very similar to that of the corresponding *S. cerevisiae* mutant, suggesting that the function and regulation of this gene have been conserved (27, 166). Given the overall similarity in the events of sporulation, it is likely that *S. cerevisiae* will serve as a good model for understanding the process in other fungi even if identification of analogous functions is not readily done by BLAST search.

### S. *CEREVISIAE* SPORULATION AS A MODEL FOR MEMBRANE BIOGENESIS EVENTS

In addition to ascospore formation in other fungi, an understanding of the various aspects of spore formation is relevant to more general events in eukaryotic cells. Of particular interest are other instances in which new membrane compartments are formed de novo. Examples of this include cellularization of the syncytial blastoderm during *Drosophila* embryogenesis, cell division during mitosis in plant cells and in apicomplexan parasites such as *Toxoplasma*, and formation of transverse tubules (t-tubules) during myotube differentiation in mammals (9, 44, 85, 134, 155, 174).

The studies of yeast sporulation described above indicate that prospore membrane formation requires both a specialized substrate, the meiosis II outer plaque, on which the membrane initially forms and a specific SNARE complex to mediate the vesicle fusion. These two requirements may be generally true of the process of membrane formation. In the mitotic division of plant cells, for example, secretory vesicles congress in the middle of the cell postmitotically and subsequently fuse to form a flattened membrane sheet termed a phragmoplast. The phragmoplast then expands to connect to the preexisting plasma membrane to complete cytokinesis (9, 155, 174). In *Arabidopsis*, the fusion of vesicles to form a phragmoplast has been shown to require both a mitosis-specific SNARE, *KNOLLE*, and a specific *SECI* homologue, *KEULE* (3, 82, 178). Thus, fusion of these vesicles appears to be mediated by a specialized SNARE complex. Moreover, in the electron microscope these vesicles are seen to be connected by electron-dense linkers that are proposed to be docking complexes (118). The protein constituents of these linkers have not been identified, but they may be a surface necessary for this fusion event, analogous to the meiosis II outer plaque.

In the case of t-tubules, the cellular event is not a cell division but the formation of deep invaginations of the plasma membrane of myotubes that allow an action potential generated at the neuromuscular synapse to be brought to the sarcomere (70). t-tubules are not formed by pulling down of the plasma membrane but rather by the generation of a tubular network within the cytosol that secondarily connects to the plasma membrane (44). A similar structure would result if, instead of closing on itself, a prospore membrane were to fuse with the plasma membrane. The genes specifying t-tubule formation have not been described. It is interesting to note, however, that the muscle-specific SNARE VAMP5 has been found



to localize to t-tubules (185). Given the similarity to prospore membrane formation, it seems possible that VAMP5 participates in a specialized SNARE complex involved in assembly of this compartment. Thus, though the specific proteins may not be conserved, the principles gleaned from the study of spore formation provide insight into analogous events in other organisms.

### PERSPECTIVES AND FUTURE DIRECTIONS

The broad outline of the cytological steps of spore construction in *S. cerevisiae* and the overall similarity of sporulation in a variety of other ascomycetes were established by a plethora of early electron microscopy studies (7, 8, 22, 50, 52, 91, 99, 128, 136, 139). Those studies firmly established the enclosure of the nucleus within a double membrane and the subsequent formation of the spore wall within the lumens of those membranes as the central features of this cell division event. More recently, the identification and characterization of mutants defective in this process have begun to reveal the molecular mechanisms and regulation of some of the detailed steps required for spore assembly. These studies both illuminate our understanding of the process and highlight critical gaps in our knowledge.

Although there is a great deal still to be learned, several questions stand out as particularly important. (i) How does the meiosis II outer plaque promote vesicle fusion? Although many of the components of this complex are now known, how the structure promotes the fusion of vesicles and acts to anchor the prospore membrane remains unknown. (ii) How is the specificity of SNAREs at the prospore membrane defined? The fusion of secretory vesicles with the prospore membrane or plasma membrane requires specific combinations from an overlapping set of SNARE proteins. How the specificity of SNARE combinations for a specific membrane is achieved remains to be determined. (iii) How does membrane closure occur? What provides the motive force to bring the ends of the prospore membrane together is not known. The leading-edge complex marks the ends of the prospore membrane and disassembles after the membrane closes. Whether the leading-edge complex aids in membrane closure or acts more passively is an important issue to be resolved. (iv) How is the assembly of the different spore wall layers coordinated? Assembly of the spore wall proceeds in a temporal order. Feedback pathways must exist to monitor the assembly of one layer and trigger the deposition of the subsequent layer. Much remains to be learned here, about both the signals being monitored and the signaling pathways that respond to those signals. The answers to these questions will provide a rich source of insight both into the process of spore formation and, more generally, into many areas of cell biology, including vesicular transport, membrane fusion, cytokinesis, and cell wall assembly.

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### REFERENCES

1. Aalto, M. K., H. Ronne, and S. Keranen. 1993. Yeast syntaxins Sso1p and Sso2p belong to a family of related membrane proteins that function in vesicular transport. *EMBO J.* **12**:4095–4104.
2. Adams, I. R., and J. V. Kilmartin. 2000. Spindle pole body duplication: a model for centrosome duplication? *Trends Cell Biol.* **10**:329–335.
3. Assaad, F. F., Y. Huet, U. Mayer, and G. Jurgens. 2001. The cytokinesis gene *KEULE* encodes a Sec1 protein that binds the syntaxin *KNOLLE*. *J. Cell Biol.* **152**:531–543.
4. Bajgier, B. K., M. Malzone, M. Nickas, and A. M. Neiman. 2001. SPO21 is required for meiosis-specific modification of the spindle pole body in yeast. *Mol. Biol. Cell* **12**:1611–1621.
5. Barral, Y., V. Mermall, M. S. Mooseker, and M. Snyder. 2000. Compartmentalization of the cell cortex by septins is required for maintenance of cell polarity in yeast. *Mol. Cell* **5**:841–851.
6. Baudry, N., P. J. Lejeune, P. Niccoli, L. Vinet, P. Carayon, and B. Mallet. 1996. Dityrosine bridge formation and thyroid hormone synthesis are tightly linked and are both dependent on N-glycans. *FEBS Lett.* **396**:223–226.
7. Beckett, A., R. Barton, and I. M. Wilson. 1968. Fine structure of the wall and appendage formation in ascospores of *Podospora anserina*. *J. Gen. Microbiol.* **53**:89–94.
8. Beckett, A., R. F. Illingworth, and A. H. Rose. 1973. Ascospore wall development in *Saccharomyces cerevisiae*. *J. Bacteriol.* **113**:1054–1057.
9. Bednarek, S. Y., and T. G. Falbel. 2002. Membrane trafficking during plant cytokinesis. *Traffic* **3**:621–629.
10. Brachat, A., J. V. Kilmartin, A. Wach, and P. Philippsen. 1998. *Saccharomyces cerevisiae* cells with defective spindle pole body outer plaques accomplish nuclear migration via half-bridge-organized microtubules. *Mol. Biol. Cell* **9**:977–991.
11. Brennwald, P., B. Kearns, K. Champion, S. Keranen, V. Bankaitis, and P. Novick. 1994. Sec9 is a SNAP-25-like component of a yeast SNARE complex that may be the effector of Sec4 function in exocytosis. *Cell* **79**:245–258.
12. Briza, P., E. Bogengruber, A. Thur, M. Rutzler, M. Munsterkotter, I. W. Dawes, and M. Breitenbach. 2002. Systematic analysis of sporulation phenotypes in 624 non-lethal homozygous deletion strains of *Saccharomyces cerevisiae*. *Yeast* **19**:403–422.
13. Briza, P., M. Breitenbach, A. Ellinger, and J. Segall. 1990. Isolation of two developmentally regulated genes involved in spore wall maturation in *Saccharomyces cerevisiae*. *Genes Dev.* **4**:1775–1789.
14. Briza, P., M. Eckerstorfer, and M. Breitenbach. 1994. The sporulation-specific enzymes encoded by the *DIT1* and *DIT2* genes catalyze a two-step reaction leading to a soluble DL-dityrosine-containing precursor of the yeast spore wall. *Proc. Natl. Acad. Sci. USA* **91**:4524–4528.
15. Briza, P., A. Ellinger, G. Winkler, and M. Breitenbach. 1990. Characterization of a DL-dityrosine-containing macromolecule from yeast ascospore walls. *J. Biol. Chem.* **265**:15118–15123.
16. Briza, P., A. Ellinger, G. Winkler, and M. Breitenbach. 1988. Chemical composition of the yeast ascospore wall. The second outer layer consists of chitosan. *J. Biol. Chem.* **263**:11569–11574.
17. Briza, P., H. Kalchhauser, E. Pittenauer, G. Allmaier, and M. Breitenbach. 1996. N,N'-Bisformyl dityrosine is an in vivo precursor of the yeast ascospore wall. *Eur. J. Biochem.* **239**:124–131.
18. Briza, P., G. Winkler, H. Kalchhauser, and M. Breitenbach. 1986. Dityrosine is a prominent component of the yeast ascospore wall. A proof of its structure. *J. Biol. Chem.* **261**:4288–4294.
19. Bungard, D., M. Reed, and E. Winter. 2004. *RSC1* and *RSC2* are required for expression of mid-late sporulation-specific genes in *Saccharomyces cerevisiae*. *Eukaryot. Cell* **3**:910–918.
20. Burgess, R. W., D. L. Deitcher, and T. L. Schwarz. 1997. The synaptic protein syntaxin1 is required for cellularization of *Drosophila* embryos. *J. Cell Biol.* **138**:861–875.
21. Byers, B. 1981. Cytology of the yeast life cycle, p. 59–96. *In* J. N. Strathern, E. W. Jones, and J. R. Broach (ed.), *The molecular biology of the yeast Saccharomyces: life cycle and inheritance*. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.
22. Carroll, G. C. 1967. The ultrastructure of ascospore delimitation in *Saccharomyces cerevisiae*. *J. Cell Biol.* **33**:218–224.
23. Christodoulidou, A., V. Bouriotis, and G. Thireos. 1996. Two sporulation-specific chitin deacetylase-encoding genes are required for the ascospore wall rigidity of *Saccharomyces cerevisiae*. *J. Biol. Chem.* **271**:31420–31425.
24. Christodoulidou, A., P. Briza, A. Ellinger, and V. Bouriotis. 1999. Yeast ascospore wall assembly requires two chitin deacetylase isozymes. *FEBS Lett.* **460**:275–279.
25. Chu, S., J. DeRisi, M. Eisen, J. Mulholland, D. Botstein, P. O. Brown, and I. Herskowitz. 1998. The transcriptional program of sporulation in budding yeast. *Science* **282**:699–705.
26. Chu, S., and I. Herskowitz. 1998. Gametogenesis in yeast is regulated by a transcriptional cascade dependent on Ndt80. *Mol. Cell* **1**:685–696.
27. Coluccio, A., E. Bogengruber, M. N. Conrad, M. E. Dresser, P. Briza, and

- A. M. Neiman. 2004. Morphogenetic pathway of spore wall assembly in *Saccharomyces cerevisiae*. *Eukaryot. Cell* **3**:1464–1475.
28. Coluccio, A., M. Malzone, and A. M. Neiman. 2004. Genetic evidence of a role for membrane lipid composition in the regulation of soluble NEM-sensitive factor receptor function in *Saccharomyces cerevisiae*. *Genetics* **166**:89–97.
  29. Coluccio, A., and A. M. Neiman. 2004. Interspore bridges: a new feature of the *Saccharomyces cerevisiae* spore wall. *Microbiology* **150**:3189–3196.
  30. Cooper, K. F., M. J. Mallory, D. B. Egeland, M. Jarnik, and R. Strich. 2000. Ama1p is a meiosis-specific regulator of the anaphase promoting complex/cyclosome in yeast. *Proc. Natl. Acad. Sci. USA* **97**:14548–14553.
  31. Davidow, L. S., L. Goetsch, and B. Byers. 1980. Preferential occurrence of nonsister spores in two-spored asci of *Saccharomyces cerevisiae*: evidence for regulation of spore-wall formation by the spindle pole body. *Genetics* **94**:581–595.
  32. DeMarini, D. J., A. E. Adams, H. Fares, C. De Virgilio, G. Valle, J. S. Chuang, and J. R. Pringle. 1997. A septin-based hierarchy of proteins required for localized deposition of chitin in the *Saccharomyces cerevisiae* cell wall. *J. Cell Biol.* **139**:75–93.
  33. Deng, C., and W. S. Saunders. 2001. ADY1, a novel gene required for prospore membrane formation at selected spindle poles in *Saccharomyces cerevisiae*. *Mol. Biol. Cell* **12**:2646–2659.
  34. Deutschbauer, A. M., R. M. Williams, A. M. Chu, and R. W. Davis. 2002. Parallel phenotypic analysis of sporulation and postgermination growth in *Saccharomyces cerevisiae*. *Proc. Natl. Acad. Sci. USA* **99**:15530–15535.
  35. De Virgilio, C., D. J. DeMarini, and J. R. Pringle. 1996. SPR28, a sixth member of the septin gene family in *Saccharomyces cerevisiae* that is expressed specifically in sporulating cells. *Microbiology* **142**:2897–2905.
  36. Enyenihi, A. H., and W. S. Saunders. 2003. Large-scale functional genomic analysis of sporulation and meiosis in *Saccharomyces cerevisiae*. *Genetics* **163**:47–54.
  37. Espinoza, F. H., A. Farrell, H. Erdjument-Bromage, P. Tempst, and D. O. Morgan. 1996. A cyclin-dependent kinase-activating kinase (CAK) in budding yeast unrelated to vertebrate CAK. *Science* **273**:1714–1717.
  38. Esposito, M. S., and R. E. Esposito. 1969. The genetic control of sporulation in *Saccharomyces*. I. The isolation of temperature-sensitive sporulation-deficient mutants. *Genetics* **61**:79–89.
  39. Esposito, M. S., R. E. Esposito, M. Arnaud, and H. O. Halvorson. 1969. Acetate utilization and macromolecular synthesis during sporulation of yeast. *J. Bacteriol.* **100**:180–186.
  40. Esposito, R. E., and S. Klapholz. 1981. Meiosis and ascospore development, p. 211–287. *In* J. N. Strathern, E. W. Jones, and J. R. Broach (ed.), *The molecular biology of the yeast Saccharomyces: life cycle and inheritance*. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.
  41. Fares, H., L. Goetsch, and J. R. Pringle. 1996. Identification of a developmentally regulated septin and involvement of the septins in spore formation in *Saccharomyces cerevisiae*. *J. Cell Biol.* **132**:399–411.
  42. Felder, T., E. Bogengruber, S. Tenreiro, A. Ellinger, I. Sa-Correia, and P. Briza. 2002. Dtr1p, a multidrug resistance transporter of the major facilitator superfamily, plays an essential role in spore wall maturation in *Saccharomyces cerevisiae*. *Eukaryot. Cell* **1**:799–810.
  43. Fleet, G. H. 2003. Yeast interactions and wine flavour. *Int. J. Food Microbiol.* **86**:11–22.
  44. Flucher, B. E., M. Terasaki, H. M. Chin, T. J. Beeler, and M. P. Daniels. 1991. Biogenesis of transverse tubules in skeletal muscle in vitro. *Dev. Biol.* **145**:77–90.
  45. Friesen, H., R. Lutz, S. Doyle, and J. Segall. 1994. Mutation of the SPS1-encoded protein kinase of *Saccharomyces cerevisiae* leads to defects in transcription and morphology during spore formation. *Genes Dev.* **8**:2162–2175.
  46. Gancedo, J. M. 2001. Control of pseudohyphae formation in *Saccharomyces cerevisiae*. *FEMS Microbiol. Rev.* **25**:107–123.
  47. Garrison, R. G., F. Mariat, K. S. Boyd, and H. Fromentin. 1979. Perithecial ultrastructure and formation of ascospores of *Ceratocystis stenoceras* (Robak) C. Moreau Ann. Microbiol. (Paris) **130A**:3–21.
  48. Gladfelter, A. S., J. R. Pringle, and D. J. Lew. 2001. The septin cortex at the yeast mother-bud neck. *Curr. Opin. Microbiol.* **4**:681–689.
  49. Gomez-Esquer, F., J. M. Rodriguez-Pena, G. Diaz, E. Rodriguez, P. Briza, C. Nombela, and J. Arroyo. 2004. CRR1, a gene encoding a putative transglycosidase, is required for proper spore wall assembly in *Saccharomyces cerevisiae*. *Microbiology* **150**:3269–3280.
  50. Greenhalgh, G. N., and L. V. Evans. 1968. The developing ascospore wall of *Hypoxylon fragiforme*. *J. R. Microsc. Soc.* **88**:545–556.
  51. Gruneberg, U., K. Campbell, C. Simpson, J. Grindlay, and E. Schiebel. 2000. Nud1p links astral microtubule organization and the control of exit from mitosis. *EMBO J.* **19**:6475–6488.
  52. Guth, E., T. Hashimoto, and S. F. Conti. 1972. Morphogenesis of ascospores in *Saccharomyces cerevisiae*. *J. Bacteriol.* **109**:869–880.
  53. Henar Valdivieso, M., A. Duran, and C. Roncero. 1999. Chitin synthases in yeast and fungi. *EXS* **87**:55–69.
  54. Herskowitz, I. 1995. MAP kinase pathways in yeast: for mating and more. *Cell* **80**:187–197.
  55. Heymont, J., L. Berenfeld, J. Collins, A. Kaganovich, B. Maynes, A. Moulin, I. Ratskovskaya, P. P. Poon, G. C. Johnston, M. Kamenetsky, J. DeSilva, H. Sun, G. A. Petsko, and J. Engebrecht. 2000. TEPI, the yeast homolog of the human tumor suppressor gene PTEN/MMAC1/TEP1, is linked to the phosphatidylinositol pathway and plays a role in the developmental process of sporulation. *Proc. Natl. Acad. Sci. USA* **97**:12672–12677.
  56. Holaway, B. L., D. J. Lehman, D. A. Primerano, P. T. Magee, and M. J. Clancy. 1985. Sporulation-regulated genes of *Saccharomyces cerevisiae*. *Curr. Genet.* **10**:163–169.
  57. Honigberg, S. M. 2004. Ime2p and Cdc28p: co-pilots driving meiotic development. *J. Cell Biochem.* **92**:1025–1033.
  58. Huang, L. S., H. K. Doherty, and I. Herskowitz. 2005. The Smk1p MAP kinase negatively regulates Gsc2p, a 1,3- $\beta$ -glucan synthase, during spore wall morphogenesis in *Saccharomyces cerevisiae*. *Proc. Natl. Acad. Sci. USA* **102**:12431–12436.
  59. Huh, W. K., J. V. Falvo, L. C. Gerke, A. S. Carroll, R. W. Howson, J. S. Weissman, and E. K. O'Shea. 2003. Global analysis of protein localization in budding yeast. *Nature* **425**:686–691.
  60. Ikemoto, S., T. Nakamura, M. Kubo, and C. Shimoda. 2000. *S. pombe* sporulation-specific coiled-coil protein Spo15p is localized to the spindle pole body and essential for its modification. *J. Cell Sci.* **113**:545–554.
  61. Inoue, S. B., N. Takewaki, T. Takasuka, T. Mio, M. Adachi, Y. Fujii, C. Miyamoto, M. Arisawa, Y. Furuichi, and T. Watanabe. 1995. Characterization and gene cloning of 1,3- $\beta$ -D-glucan synthase from *Saccharomyces cerevisiae*. *Eur. J. Biochem.* **231**:845–854.
  62. Ishihara, S., A. Hirata, M. Minemura, S. Nogami, and Y. Ohya. 2001. A mutation in SPC42, which encodes a component of the spindle pole body, results in production of two-spored asci in *Saccharomyces cerevisiae*. *Mol. Genet. Genomics* **265**:585–595.
  63. Ito, T., T. Chiba, R. Ozawa, M. Yoshida, M. Hattori, and Y. Sakaki. 2001. A comprehensive two-hybrid analysis to explore the yeast protein interactome. *Proc. Natl. Acad. Sci. USA* **98**:4569–4574.
  64. Iwamoto, M. A., S. R. Fairclough, S. A. Rudge, and J. Engebrecht. 2005. *Saccharomyces cerevisiae* Sps1p regulates trafficking of enzymes required for spore wall synthesis. *Eukaryot. Cell* **4**:536–544.
  65. Jantsch-Plunger, V., and M. Glotzer. 1999. Depletion of syntaxins in the early *Caenorhabditis elegans* embryo reveals a role for membrane fusion events in cytokinesis. *Curr. Biol.* **9**:738–745.
  66. Jantti, J., M. K. Aalto, M. Oyen, L. Sundqvist, S. Keranen, and H. Ronne. 2002. Characterization of temperature-sensitive mutations in the yeast syntaxin 1 homologues Sso1p and Sso2p, and evidence of a distinct function for Sso1p in sporulation. *J. Cell Sci.* **115**:409–420.
  67. Jaspersen, S. L., and M. Winey. 2004. The budding yeast spindle pole body: structure, duplication, and function. *Annu. Rev. Cell Dev. Biol.* **20**:1–28.
  68. Kaback, D. B., and L. R. Feldberg. 1985. *Saccharomyces cerevisiae* exhibits a sporulation-specific temporal pattern of transcript accumulation. *Mol. Cell Biol.* **5**:751–761.
  69. Kaldis, P., A. Sutton, and M. J. Solomon. 1996. The Cdk-activating kinase (CAK) from budding yeast. *Cell* **86**:553–564.
  70. Katz, B. 1966. Nerve, muscle, and synapse. McGraw-Hill, New York, N.Y.
  71. Kay, E. S., and B. M. Shapiro. 1987. Ovoperoxidase assembly into the sea urchin fertilization envelope and dityrosine crosslinking. *Dev. Biol.* **121**:325–334.
  72. Kinoshita, M. 2003. The septins. *Genome Biol.* **4**:236.
  73. Klapholz, S., and R. E. Esposito. 1980. Recombination and chromosome segregation during the single division meiosis in SPO12-1 and SPO13-1 diploids. *Genetics* **96**:589–611.
  74. Klis, F. M., P. Mol, K. Hellingerwerf, and S. Brul. 2002. Dynamics of cell wall structure in *Saccharomyces cerevisiae*. *FEMS Microbiol. Rev.* **26**:239–256.
  75. Knop, M., and E. Schiebel. 1998. Receptors determine the cellular localization of a gamma-tubulin complex and thereby the site of microtubule formation. *EMBO J.* **17**:3952–3967.
  76. Knop, M., and K. Strasser. 2000. Role of the spindle pole body of yeast in mediating assembly of the prospore membrane during meiosis. *EMBO J.* **19**:3657–3667.
  77. Kornberg, H. L. 1966. Anaplerotic sequences and their role in metabolism, p. 1–31. *In* P. N. Campbell and G. D. Greville (ed.), *Essays in biochemistry*, vol. 2. Academic Press, London, United Kingdom.
  78. Kreger-Van Rij, N. J. 1978. Electron microscopy of germinating ascospores of *Saccharomyces cerevisiae*. *Arch. Microbiol.* **117**:73–77.
  79. Krisak, L., R. Strich, R. S. Winters, J. P. Hall, M. J. Mallory, D. Kreitzer, R. S. Tuan, and E. Winter. 1994. SMK1, a developmentally regulated MAP kinase, is required for spore wall assembly in *Saccharomyces cerevisiae*. *Genes Dev.* **8**:2151–2161.
  80. Kupiec, M., B. Byers, and R. E. Esposito. 1997. Meiosis and sporulation in *Saccharomyces cerevisiae*, p. 889–1036. *In* J. R. Pringle, J. R. Broach, and E. W. Jones (ed.), *The molecular and cell biology of the yeast Saccharomyces*. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y.
  81. Lassandro, F., M. Sebastiano, F. Zei, and P. Bazzicalupo. 1994. The role of dityrosine formation in the crosslinking of CUT-2, the product of a second cuticlin gene of *Caenorhabditis elegans*. *Mol. Biochem. Parasitol.* **65**:147–159.
  82. Lauber, M. H., I. Waizenegger, T. Steinmann, H. Schwarz, U. Mayer, I.

- Hwang, W. Lukowitz, and G. Jurgens. 1997. The Arabidopsis KNOLLE protein is a cytokinesis-specific syntaxin. *J. Cell Biol.* **139**:1485–1493.
83. Lauze, E., B. Stoelcker, F. C. Luca, E. Weiss, A. R. Schutz, and M. Winey. 1995. Yeast spindle pole body duplication gene MPS1 encodes an essential dual specificity protein kinase. *EMBO J.* **14**:1655–16563.
84. Leberer, E., D. Dignard, D. Harcus, D. Y. Thomas, and M. Whiteway. 1992. The protein kinase homologue Ste20p is required to link the yeast pheromone response G-protein beta gamma subunits to downstream signalling components. *EMBO J.* **11**:4815–4824.
85. Lecuit, T. 2004. Junctions and vesicular trafficking during *Drosophila* cellularization. *J. Cell Sci.* **117**:3427–3433.
86. Lee, B., and A. Amon. 2001. Meiosis: how to create a specialized cell cycle. *Curr. Opin. Cell Biol.* **13**:770–777.
87. Lewin, A. S., V. Hines, and G. M. Small. 1990. Citrate synthase encoded by the *CIT2* gene of *Saccharomyces cerevisiae* is peroxisomal. *Mol. Cell. Biol.* **10**:1399–1405.
88. Liakopoulos, D., J. Kusch, S. Grava, J. Vogel, and Y. Barral. 2003. Asymmetric loading of Kar9 onto spindle poles and microtubules ensures proper spindle alignment. *Cell* **112**:561–574.
89. Longtine, M. S., and E. Bi. 2003. Regulation of septin organization and function in yeast. *Trends Cell Biol.* **13**:403–409.
90. Lowenstein, J. M. 1969. Citric acid cycle: control and compartmentation. Marcel Dekker, New York, N.Y.
91. Lynn, R. R., and P. T. Magee. 1970. Development of the spore wall during ascospore formation in *Saccharomyces cerevisiae*. *J. Cell Biol.* **44**:688–692.
92. Marston, A. L., W. H. Tham, H. Shah, and A. Amon. 2004. A genome-wide screen identifies genes required for centromeric cohesion. *Science* **303**:1367–1370.
93. Mata, J., R. Lyne, G. Burns, and J. Bahler. 2002. The transcriptional program of meiosis and sporulation in fission yeast. *Nat. Genet.* **32**:143–147.
94. Mazur, P., N. Morin, W. Baginsky, M. el-Sherbeini, J. A. Cemas, J. B. Nielsen, and F. Foor. 1995. Differential expression and function of two homologous subunits of yeast 1,3-beta-D-glucan synthase. *Mol. Cell. Biol.* **15**:5671–5681.
95. Miller, J. J. 1957. The metabolism of yeast sporulation. II. Stimulation and inhibition by monosaccharides. *Can. J. Microbiol.* **3**:81–90.
96. Minard, K. L., and L. McAlister-Henn. 1991. Isolation, nucleotide sequence analysis, and disruption of the *MDH2* gene from *Saccharomyces cerevisiae*: evidence for three isozymes of yeast malate dehydrogenase. *Mol. Cell. Biol.* **11**:370–380.
97. Mishra, C., C. E. Semino, K. J. McCreath, H. de la Vega, B. J. Jones, C. A. Specht, and P. W. Robbins. 1997. Cloning and expression of two chitin deacetylase genes of *Saccharomyces cerevisiae*. *Yeast* **13**:327–336.
98. Mitchell, A. P. 1994. Control of meiotic gene expression in *Saccharomyces cerevisiae*. *Microbiol. Rev.* **58**:56–70.
99. Moens, P. B. 1971. Fine structure of ascospore development in the yeast *Saccharomyces cerevisiae*. *Can. J. Microbiol.* **17**:507–510.
100. Moens, P. B., and E. Rapport. 1971. Spindles, spindle plaques, and meiosis in the yeast *Saccharomyces cerevisiae* (Hansen). *J. Cell Biol.* **50**:344–361.
101. Moreno-Borchart, A. C., K. Strasser, M. G. Finkbeiner, A. Shevchenko, and M. Knop. 2001. Prospore membrane formation linked to the leading edge protein (LEP) coat assembly. *EMBO J.* **20**:6946–6957.
102. Nag, D. K., M. P. Koonce, and J. Axelrod. 1997. *SSP1*, a gene necessary for proper completion of meiotic divisions and spore formation in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **17**:7029–7039.
103. Nakanishi, H., P. de los Santos, and A. M. Neiman. 2004. Positive and negative regulation of a SNARE protein by control of intracellular localization. *Mol. Biol. Cell* **15**:1802–1815.
104. Nakanishi, H., M. Morishita, C. L. Schwartz, A. Coluccio, J. Engebrecht, and A. M. Neiman. Phospholipase D is required to promote vesicle fusion during sporulation in yeast. Submitted for publication.
105. Neugeborn, L., and A. P. Mitchell. 1991. The yeast MCK1 gene encodes a protein kinase homolog that activates early meiotic gene expression. *Genes Dev.* **5**:533–548.
106. Neiman, A. M. 1998. Prospore membrane formation defines a developmentally regulated branch of the secretory pathway in yeast. *J. Cell Biol.* **140**:29–37.
107. Neiman, A. M., L. Katz, and P. J. Brenwald. 2000. Identification of domains required for developmentally regulated SNARE function in *Saccharomyces cerevisiae*. *Genetics* **155**:1643–1655.
108. Nelson, R. E., L. I. Fessler, Y. Takagi, B. Blumberg, D. R. Keene, P. F. Olson, C. G. Parker, and J. H. Fessler. 1994. Peroxidase: a novel enzyme-matrix protein of *Drosophila* development. *EMBO J.* **13**:3438–3447.
109. Neuwald, A. F. 1997. Barth syndrome may be due to an acyltransferase deficiency. *Curr. Biol.* **7**:R465–466.
110. Newlon, M. C., and B. D. Hall. 1978. Inhibition of yeast sporulation by ethidium bromide. *Mol. Gen. Genet.* **165**:113–114.
111. Nickas, M. E., A. E. Diamond, M. J. Yang, and A. M. Neiman. 2004. Regulation of spindle pole function by an intermediary metabolite. *Mol. Biol. Cell* **15**:2606–2616.
112. Nickas, M. E., and A. M. Neiman. 2002. Ady3p links spindle pole body function to spore wall synthesis in *Saccharomyces cerevisiae*. *Genetics* **160**:1439–1450.
113. Nickas, M. E., C. Schwartz, and A. M. Neiman. 2003. Ady4p and Spo74p are components of the meiotic spindle pole body that promote growth of the prospore membrane in *Saccharomyces cerevisiae*. *Eukaryot. Cell* **2**:431–445.
114. Oelschlaegel, T., M. Schwickart, J. Matos, A. Bogdanova, A. Camasses, J. Havlis, A. Shevchenko, and W. Zachariae. 2005. The yeast APC/C subunit Mnd2 prevents premature sister chromatid separation triggered by the meiosis-specific APC/C-Ama1. *Cell* **120**:773–788.
115. Okamoto, S., and T. Iino. 1981. Selective abortion of two nonsister nuclei in a developing ascus of the hfd-1 mutant in *Saccharomyces cerevisiae*. *Genetics* **99**:197–209.
116. Okuzaki, D., W. Satake, A. Hirata, and H. Nojima. 2003. Fission yeast *meu14+* is required for proper nuclear division and accurate forespore membrane formation during meiosis II. *J. Cell Sci.* **116**:2721–2735.
117. Ostapenko, D., and M. J. Solomon. 2005. Phosphorylation by Cak1 regulates the C-terminal domain kinase Ctk1 in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **25**:3906–3913.
118. Otegui, M. S., and L. A. Staehelin. 2004. Electron tomographic analysis of post-meiotic cytokinesis during pollen development in *Arabidopsis thaliana*. *Planta* **218**:501–515.
119. Oyen, M., J. Jantti, S. Keranen, and H. Ronne. 2004. Mapping of sporulation-specific functions in the yeast syntaxin gene SSO1. *Curr. Genet.* **45**:76–82.
120. Pak, J., and J. Segall. 2002. Regulation of the premiddle and middle phases of expression of the *NDT80* gene during sporulation of *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **22**:6417–6429.
121. Pammer, M., P. Briza, A. Ellinger, T. Schuster, R. Stucka, H. Feldmann, and M. Breitenbach. 1992. DIT101 (CSD2, CAL1), a cell cycle-regulated yeast gene required for synthesis of chitin in cell walls and chitosan in spore walls. *Yeast* **8**:1089–1099.
122. Pardo, M., L. Monteoliva, P. Vazquez, R. Martinez, G. Molero, C. Nombela, and C. Gil. 2004. PST1 and ECM33 encode two yeast cell surface GPI proteins important for cell wall integrity. *Microbiology* **150**:4157–4170.
123. Parlati, F., O. Varlamov, K. Paz, J. A. McNew, D. Hurtado, T. H. Sollner, and J. E. Rothman. 2002. Distinct SNARE complexes mediating membrane fusion in Golgi transport based on combinatorial specificity. *Proc. Natl. Acad. Sci. USA* **99**:5424–5429.
124. Pelham, H. R. 1999. SNAREs and the secretory pathway—lessons from yeast. *Exp. Cell Res.* **247**:1–8.
125. Penkner, A. M., S. Prinz, S. Ferscha, and F. Klein. 2005. Mnd2, an essential antagonist of the anaphase-promoting complex during meiotic prophase. *Cell* **120**:789–801.
126. Percival-Smith, A., and J. Segall. 1984. Isolation of DNA sequences preferentially expressed during sporulation in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **4**:142–150.
127. Pereira, G., T. U. Tanaka, K. Nasmyth, and E. Schiebel. 2001. Modes of spindle pole body inheritance and segregation of the Bfa1p-Bub2p checkpoint protein complex. *EMBO J.* **20**:6359–6370.
128. Plurad, S. B. 1972. Fine structure of ascosporeogenesis in *Nematospora coryli* Peglion, a pathogenic yeast. *J. Bacteriol.* **109**:927–929.
129. Pontremoli, S., and E. Grazi. 1968. Gluconeogenesis, p. 259–295. In F. Dickens, P. J. Randle, and W. J. Whelan (ed.), *Carbohydrate metabolism and its disorders*, vol. 1. Academic Press, New York, N.Y.
130. Primig, M., R. M. Williams, E. A. Winzeler, G. G. Tevzadze, A. R. Conway, S. Y. Hwang, R. W. Davis, and R. E. Esposito. 2000. The core meiotic transcriptome in budding yeasts. *Nat. Genet.* **26**:415–423.
131. Protopopov, V., B. Govindan, P. Novick, and J. E. Gerst. 1993. Homologs of the synaptobrevin/VAMP family of synaptic vesicle proteins function on the late secretory pathway in *S. cerevisiae*. *Cell* **74**:855–861.
132. Pruyne, D. W., D. H. Schott, and A. Bretscher. 1998. Tropomyosin-containing actin cables direct the Myo2p-dependent polarized delivery of secretory vesicles in budding yeast. *J. Cell Biol.* **143**:1931–1945.
133. Rabitsch, K. P., A. Toth, M. Galova, A. Schleiffer, G. Schaffner, E. Aigner, R. Rupp, A. M. Penkner, A. C. Moreno-Borchart, M. Primig, R. E. Esposito, F. Klein, M. Knop, and K. Nasmyth. 2001. A screen for genes required for meiosis and spore formation based on whole-genome expression. *Curr. Biol.* **11**:1001–1009.
134. Radke, J. R., B. Striepen, M. N. Guerini, M. E. Jerome, D. S. Roos, and M. W. White. 2001. Defining the cell cycle for the tachyzoite stage of *Toxoplasma gondii*. *Mol. Biochem. Parasitol.* **115**:165–175.
135. Raju, N. B. 1980. Meiosis and ascospore genesis in *Neurospora*. *Eur. J. Cell Biol.* **23**:208–223.
136. Reeves, F., Jr. 1967. The fine structure of ascospore formation in *Pyrenopeziza domesticum*. *Mycologia* **59**:1018–1033.
137. Reeves, H. C., R. Rabin, W. S. Wegener, and S. J. Ajl. 1967. Fatty acid synthesis and metabolism in microorganisms. *Annu. Rev. Microbiol.* **21**:225–256.
138. Roeder, G. S. 1995. Sex and the single cell: meiosis in yeast. *Proc. Natl. Acad. Sci. USA* **92**:10450–10456.

139. Rooney, L., and P. B. Moens. 1973. The fine structure of ascospore delimitation in the yeast *Wickerhamia fluorescens*. *Can. J. Microbiol.* **19**:1389–1392.
140. Rose, K., S. A. Rudge, M. A. Frohman, A. J. Morris, and J. Engebrecht. 1995. Phospholipase D signaling is essential for meiosis. *Proc. Natl. Acad. Sci. USA* **92**:12151–12155.
141. Rossi, G., A. Salminen, L. M. Rice, A. T. Brunger, and P. Brennwald. 1997. Analysis of a yeast SNARE complex reveals remarkable similarity to the neuronal SNARE complex and a novel function for the C terminus of the SNAP-25 homolog, Sec9. *J. Biol. Chem.* **272**:16610–16617.
142. Rothman, J. E. 1994. Intracellular membrane fusion. *Adv. Second Messenger Phosphoprotein Res.* **29**:81–96.
143. Rudge, S. A., A. J. Morris, and J. Engebrecht. 1998. Relocalization of phospholipase D activity mediates membrane formation during meiosis. *J. Cell Biol.* **140**:81–90.
144. Rudge, S. A., V. A. Sciorra, M. Iwamoto, C. Zhou, T. Strahl, A. J. Morris, J. Thorer, and J. Engebrecht. 2004. Roles of phosphoinositides and of Spo14p (phospholipase D)-generated phosphatidic acid during yeast sporulation. *Mol. Biol. Cell* **15**:207–218.
145. Santos, B., A. Duran, and M. H. Valdivieso. 1997. *CHS5*, a gene involved in chitin synthesis and mating in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **17**:2485–2496.
146. Sanz, M., J. A. Trilla, A. Duran, and C. Roncero. 2002. Control of chitin synthesis through Shc1p, a functional homologue of Chs4p specifically induced during sporulation. *Mol. Microbiol.* **43**:1183–1195.
147. Sarkar, P. K., M. A. Florczyk, K. A. McDonough, and D. K. Nag. 2002. SSP2, a sporulation-specific gene necessary for outer spore wall assembly in the yeast *Saccharomyces cerevisiae*. *Mol. Genet. Genomics.* **267**:348–358.
148. Schaber, M., A. Lindgren, K. Schindler, D. Bungard, P. Kaldis, and E. Winter. 2002. *CAK1* promotes meiosis and spore formation in *Saccharomyces cerevisiae* in a *CDC28*-independent fashion. *Mol. Cell. Biol.* **22**:57–68.
149. Schaerer, F., G. Morgan, M. Winey, and P. Philippsen. 2001. Cnm67p is a spacer protein of the *Saccharomyces cerevisiae* spindle pole body outer plaque. *Mol. Biol. Cell* **12**:2519–2533.
150. Schindler, K., K. R. Benjamin, A. Martin, A. Boglioli, I. Herskowitz, and E. Winter. 2003. The Cdk-activating kinase Cak1p promotes meiotic S phase through Ime2p. *Mol. Cell. Biol.* **23**:8718–8728.
151. Shimoda, C. 2004. Forespore membrane assembly in yeast: coordinating SPBs and membrane trafficking. *J. Cell Sci.* **117**:389–396.
152. Shoemaker, D. D., D. A. Lashkari, D. Morris, M. Mittmann, and R. W. Davis. 1996. Quantitative phenotypic analysis of yeast deletion mutants using a highly parallel molecular bar-coding strategy. *Nat. Genet.* **14**:450–456.
153. Shuster, C. B., and D. R. Burgess. 2002. Targeted new membrane addition in the cleavage furrow is a late, separate event in cytokinesis. *Proc. Natl. Acad. Sci. USA* **99**:3633–3638.
154. Smail, E. H., P. Briza, A. Panagos, and L. Berenfeld. 1995. *Candida albicans* cell walls contain the fluorescent cross-linking amino acid dityrosine. *Infect. Immun.* **63**:4078–4083.
155. Smith, L. G. 1999. Divide and conquer: cytokinesis in plant cells. *Curr. Opin. Plant Biol.* **2**:447–453.
156. Smits, G. J., H. van den Ende, and F. M. Klis. 2001. Differential regulation of cell wall biogenesis during growth and development in yeast. *Microbiology* **147**:781–794.
157. Sols, A., C. Gancedo, and G. DelaFuente. 1971. Energy-yielding metabolism in yeasts, p. 271–307. In A. H. Rose and J. S. Harrison (ed.), *The yeasts*, vol. 2. Academic Press, New York, N.Y.
158. Srivastava, P. K., S. Harashima, and Y. Oshima. 1981. Formation of two-spored asci by interrupted sporulation in *Saccharomyces cerevisiae*. *J. Gen. Microbiol.* **123**:39–48.
159. Steffan, J. S., and L. McAlister-Henn. 1992. Isolation and characterization of the yeast gene encoding the MDH3 isozyme of malate dehydrogenase. *J. Biol. Chem.* **267**:24708–24715.
160. Stern, B. M. 2003. FEARless in meiosis. *Mol. Cell* **11**:1123–1125.
161. Straight, P. D., T. H. Giddings, Jr., and M. Winey. 2000. Mps1p regulates meiotic spindle pole body duplication in addition to having novel roles during sporulation. *Mol. Biol. Cell* **11**:3525–3537.
162. Sutton, R. B., D. Fasshauer, R. Jahn, and A. T. Brunger. 1998. Crystal structure of a SNARE complex involved in synaptic exocytosis at 2.4 Å resolution. *Nature* **395**:347–353.
163. Tachikawa, H., A. Bloecher, K. Tatchell, and A. M. Neiman. 2001. A Gip1p-Glc7p phosphatase complex regulates septin organization and spore wall formation. *J. Cell Biol.* **155**:797–808.
164. Tanaka, K., and A. Hirata. 1982. Ascospore development in the fission yeasts *Schizosaccharomyces pombe* and *S. japonicus*. *J. Cell Sci.* **56**:263–279.
165. Thompson-Coffe, C., and D. Zickler. 1993. Cytoskeleton interactions in the ascus development and sporulation of *Sordaria macrospora*. *J. Cell Sci.* **104**:883–898.
166. Tougan, T., Y. Chiba, Y. Kakihara, A. Hirata, and H. Nojima. 2002. Meu10 is required for spore wall maturation in *Schizosaccharomyces pombe*. *Genes Cells* **7**:217–231.
167. Trilla, J. A., T. Cos, A. Duran, and C. Roncero. 1997. Characterization of *CHS4* (*CAL2*), a gene of *Saccharomyces cerevisiae* involved in chitin biosynthesis and allelic to *SKT5* and *CSD4*. *Yeast* **13**:795–807.
168. Trilla, J. A., A. Duran, and C. Roncero. 1999. Chs7p, a new protein involved in the control of protein export from the endoplasmic reticulum that is specifically engaged in the regulation of chitin synthesis in *Saccharomyces cerevisiae*. *J. Cell Biol.* **145**:1153–1163.
169. Tsuboi, M., K. Kondo, and N. Yanagishima. 1974. Inhibition of sporulation by ethidium bromide and its reversal by fermentable sugars in *Saccharomyces cerevisiae*. *Arch. Microbiol.* **99**:295–305.
170. Tu, J., W. Song, and M. Carlson. 1996. Protein phosphatase type 1 interacts with proteins required for meiosis and other cellular processes in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **16**:4199–4206.
171. Uetz, P., L. Giot, G. Cagney, T. A. Mansfield, R. S. Judson, J. R. Knight, D. Lockshon, V. Narayan, M. Srinivasan, P. Pochart, A. Qureshi-Emili, Y. Li, B. Godwin, D. Conover, T. Kalbfleisch, G. Vijayadomodar, M. Yang, M. Johnston, S. Fields, and J. M. Rothberg. 2000. A comprehensive analysis of protein-protein interactions in *Saccharomyces cerevisiae*. *Nature* **403**:623–627.
172. Ufano, S., P. San-Segundo, F. del Rey, and C. R. Vazquez de Aldana. 1999. *SWMI*, a developmentally regulated gene, is required for spore wall assembly in *Saccharomyces cerevisiae*. *Mol. Cell. Biol.* **19**:2118–2129.
173. Uno, I., K. Matsumoto, A. Hirata, and T. Ishikawa. 1985. Outer plaque assembly and spore encapsulation are defective during sporulation of adenylate cyclase-deficient mutants of *Saccharomyces cerevisiae*. *J. Cell Biol.* **100**:1854–1862.
174. Verma, D. P. 2001. Cytokinesis and building of the cell plate in plants. *Annu. Rev. Plant Physiol. Plant Mol. Biol.* **52**:751–784.
175. Vershon, A. K., and M. Pierce. 2000. Transcriptional regulation of meiosis in yeast. *Curr. Opin. Cell Biol.* **12**:334–339.
176. Wagner, M., P. Briza, M. Pierce, and E. Winter. 1999. Distinct steps in yeast spore morphogenesis require distinct SMK1 MAP kinase thresholds. *Genetics* **151**:1327–1340.
177. Wagner, M., M. Pierce, and E. Winter. 1997. The CDK-activating kinase *CAK1* can dosage suppress sporulation defects of *smk1* MAP kinase mutants and is required for spore wall morphogenesis in *Saccharomyces cerevisiae*. *EMBO J.* **16**:1305–1317.
178. Waizenegger, I., W. Lukowitz, F. Assaad, H. Schwarz, G. Jurgens, and U. Mayer. 2000. The Arabidopsis *KNOLLE* and *KEULE* genes interact to promote vesicle fusion during cytokinesis. *Curr. Biol.* **10**:1371–1374.
179. Wang, H., X. Tang, J. Liu, S. Trautmann, D. Balasundaram, D. McCollum, and M. K. Balasubramanian. 2002. The multiprotein exocyst complex is essential for cell separation in *Schizosaccharomyces pombe*. *Mol. Biol. Cell* **13**:515–529.
180. Weber, T., B. V. Zemelman, J. A. McNew, B. Westermann, M. Gmachl, F. Parlati, T. H. Sollner, and J. E. Rothman. 1998. SNAREpins: minimal machinery for membrane fusion. *Cell* **92**:759–772.
181. Wesp, A., S. Prinz, and G. R. Fink. 2001. Conservative duplication of spindle poles during meiosis in *Saccharomyces cerevisiae*. *J. Bacteriol.* **183**:2372–2375.
182. Whitacre, J., D. Davis, K. Toenjes, S. Brower, and A. Adams. 2001. Generation of an isogenic collection of yeast actin mutants and identification of three interrelated phenotypes. *Genetics* **157**:533–543.
183. Wigge, P. A., O. N. Jensen, S. Holmes, S. Soues, M. Mann, and J. V. Kilmartin. 1998. Analysis of the *Saccharomyces cerevisiae* spindle pole by matrix-assisted laser desorption/ionization (MALDI) mass spectrometry. *J. Cell Biol.* **141**:967–977.
184. Yao, S., and G. Prelich. 2002. Activation of the Bur1-Bur2 cyclin-dependent kinase complex by Cak1. *Mol. Cell. Biol.* **22**:6750–6758.
185. Zeng, Q., V. N. Subramaniam, S. H. Wong, B. L. Tang, R. G. Parton, S. Rea, D. E. James, and W. Hong. 1998. A novel synaptobrevin/VAMP homologous protein (*VAMP5*) is increased during in vitro myogenesis and present in the plasma membrane. *Mol. Biol. Cell* **9**:2423–2437.
186. Ziman, M., J. S. Chuang, M. Tsung, S. Hamamoto, and R. Schekman. 1998. *Chs6p*-dependent anterograde transport of *Chs3p* from the chitosome to the plasma membrane in *Saccharomyces cerevisiae*. *Mol. Biol. Cell* **9**:1565–1576.
187. Zubenko, G. S., and E. W. Jones. 1981. Protein degradation, meiosis and sporulation in proteinase-deficient mutants of *Saccharomyces cerevisiae*. *Genetics* **97**:45–64.