Search for inhibitors of endocytosis
Intended specificity and unintended consequences

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We discuss here the variety of approaches that have been taken to inhibit different forms of endocytosis. Typically, both non-specific and specific chemical inhibitors of endocytosis are tried in order to "classify" entry of a new plasma membrane protein into one of the various types of endocytosis. This classification can be confirmed through genetic approaches of protein depletion or overexpression of mutants of known endocytosis machinery components. Although some new compounds have been designed to be selective in biochemical assays, we caution investigators to be alert to the unintended consequences that sometimes arise when these compounds are applied to intact cells.

Endocytosis is a process that cells use to bring extracellular material and plasma membrane into the cell interior. Once internalized, the fluid, membrane proteins and membrane lipids meet different fates by trafficking to different compartments: to late endosomes and lysosomes for degradation, to recycling endosomes for recycling back to the PM, to the trans Golgi network or to other destinations in the cell. Endocytosis is important for the proper signaling and regulation of cell surface receptors, delivery of nutrients into the cell, establishment and maintenance of cell polarity, and the turnover of PM proteins and lipids. Additionally, bacterial toxins and pathogens use endocytosis as a mode of entry to the cell interior. Understanding how this process occurs and how it is regulated is a goal for cell biologists. Complicating this is the fact that endocytosis takes many forms.

Classically, endocytosis can be divided into pinocytosis and phagocytosis (Fig. 1). Pinocytosis involves internalization of fluid while phagocytosis, an actin-dependent process, involves the internalization of large particles such as bacteria. Pinocytosis can be further divided into those that are dependent on the clathrin coat (clathrin-mediated endocytosis, CME) or those that are independent of clathrin (clathrin-independent endocytosis, CIE). CME has been extensively studied for the past 30 years and the machinery involved in selecting the cargo and initiating and completing the process is well understood. CME requires adaptor proteins that select and concentrate cargo into clathrin-coated pits and depends on the dynamin GTPase to facilitate vesicle scission. By contrast, the variety of forms of CIE observed in different cells has presented a complicated picture, making descriptions of these pathways less clear.

CIE is involved in the internalization of glycolipid-binding toxins, glycosylphosphatidyl inositol-anchored proteins (GPI-AP), and many cell surface proteins (channels, transporters, proteins involved in cell-cell and cell-matrix interactions and in cellular immune function). CIE occurs independently of adaptor proteins and clathrin coats, and mostly does not require dynamin for vesicle scission. CIE pathways are an active area of study. Thus far, it has been shown that the small GTPase Arf6 is associated with the uptake and sorting of many plasma membrane proteins while some lipid-raft associated pathways (called CLIC/GEEC) are involved in endocytosis of GPI-AP. Rho proteins have been implicated in yet

Keywords: endocytosis, pinocytosis, phagocytosis, clathrin-independent endocytosis, clathrin-mediated endocytosis, inhibitor, chemical inhibitor

Submitted: 01/21/13
Revised: 02/11/13
Accepted: 02/11/13
http://dx.doi.org/10.4161/cl.23967
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another CIE pathway. It is likely that the Arf6 and CLIC-GEEC pathways are closely related since they both are clathrin- and dynamin-independent, cholesterol-dependent and carry GPI-AP into the cell. Finally, macropinocytosis is a stimulated form of CIE where large pinosomes are brought into the cell interior as a consequence of cellular protrusions in an actin-dependent process.

To better understand the different types of endocytosis, cell biologists have sought ways to block this process through chemical and genetic means. The use of such inhibitors can reveal molecular components required for, and the physiological consequences of blocking, specific forms of endocytosis. These studies contribute to an understanding of the basic mechanism(s) of endocytosis and help define modes of cellular entry for medically relevant components such as signaling receptors, and bacterial and viral pathogens. In this commentary we discuss the approaches that have been taken to block particular forms of endocytosis (see Table 1). These include the use of non-specific chemical inhibitors, the new generation of selective pharmacologic agents, and genetic approaches designed to target a particular form of endocytosis. We will discuss advantages and limitations of each approach.

**Classical Chemical Inhibitors of Endocytosis**

An attractive feature of chemical inhibitors is that they may be administered acutely to reveal direct inhibition of a particular process. Reversibility of the block can also help support the notion that the effect is specific on the particular mode of endocytosis being examined. Nevertheless, there is always a possibility of indirect effects of the agent influencing another process that will affect that form of endocytosis. For example, inhibitors of actin polymerization, cytochalasin D or latrunculin block both actin-dependent phagocytosis and macropinocytosis, which is not surprising since macropinosomes and phagosomes are coated with actin. Although actin polymerization appears to be required for endocytosis in yeast, this is less clear in mammalian cells. Inhibitors of actin polymerization have variable effects on transferrin endocytosis by CME depending on the cell line examined and experimental conditions used. Some forms of CIE are not dependent on actin polymerization however recycling of endosomal membrane back to the cell surface is dependent on actin, which could complicate interpretations of experiments. Thus, care should be taken to ensure that a requirement for a process is a direct and specific one and not a downstream consequence of inhibiting another process.

Over the past 25 years a number of chemical inhibitors have been used with the intent to block CME. In many instances and cell types this is the case; however, these treatments are general cellular perturbants and thus suffer from unknown global effects on the cell (see Table 1). For example, potassium depletion was found to block CME by causing aggregation of clathrin in the cytoplasm thus removing it from functioning at the cell surface. However, significant reduction of potassium in the cell leads to a reduction in protein and DNA synthesis. Hypertonic sucrose is another treatment used to inhibit CME yet it also causes cell shrinkage and may lead to changes in cortical actin cytoskeleton. Cytosol acidification blocks CME by freezing clathrin-coated pits at the cell surface; however, other effects on the actin cytoskeleton and macropinocytosis have been reported. Chlorpromazine is a cationic amphipathic drug that causes the assembly of adaptor proteins and...
clathrin on endosomal membranes thus depleting it from the PM, leading to a block in CME. However this drug also causes an inhibition of receptor recycling and inhibits CIE in some cells. Indeed, a study testing a number of these inhibitors on four different cell lines revealed that these compounds are not very specific and that the efficacy, even for inhibiting CME, varies in different cell lines. GPI-APs and toxins that bind to membrane lipids enter cells via CIE and various manipulations of PM cholesterol have been used to suggest a mechanism requiring cholesterol and glycosphingolipid-enriched membrane or “lipid raft domains.”

Cyclodextrins such as methyl-β-cyclodextrin have been used to deplete cells of PM cholesterol, leading to a block in endocytosis of various toxins and GPI-APs. Although this treatment is effective and can be reversed by cholesterol repletion, it results in profound changes in cell surface structure and can affect other endocytic pathways including macroendocytosis and CME. As an alternative to extracting cholesterol from cells, the cholesterol-binding agent filipin can be used to bind to cell surface cholesterol and rapidly inhibit endocytosis of GPI-APs. Although free PM cholesterol is required for endocytosis of GPI-APs and lipid-binding toxins, it is also necessary for entry of a number of PM proteins (MHC, Glut1, CD44, CD98 and CD147) that are not lipid-raft partitioning proteins but enter cells by CIE. At higher concentrations of filipin CME can also be inhibited, thus demonstrating that most, if not all, forms of endocytosis are dependent upon PM cholesterol. One compound that consistently blocks macroendocytosis is amiloride, which acts to inhibit Na+/H+ exchange at the cell surface.

### Table 1. Chemical and genetic endocytosis inhibitors

| Endocytosis inhibitors | Pathways targeted | Mode of action | Comments |
|------------------------|------------------|----------------|----------|
| **Chemical inhibitors** |                  |                |          |
| Hypertonic sucrose     | CME              | Traps clathrin in microcages | Nonspecific; interferes with fluid phase macropinocytosis |
| Potassium depletion    | CME              | Aggregates clathrin | Nonspecific; affects actin cytoskeleton |
| Cytosol acidification  | CME              | Inhibits the scission of the clathrin pits from the membrane | Interferes with macroendocytosis and actin cytoskeleton |
| Chlorpromazine         | CME              | Translocates clathrin and AP2 from the cell surface to intracellular endosomes | Inhibits CIE in some cells |
| Monodansylcadaverine   | CME              | Stabilizes CCVs | Global changes in actin dynamics |
| Phynylarsine oxide     | CME              | Not clearly known. | Inhibits macroendocytosis and phagocytosis |
| Chloroquine            | CME              | Affects the function of CCVs |          |
| Monensin               | CME              | Affects the proton gradient |          |
| Phentothiazines        | CME; phagocytosis | Affects the formation of CCVs |          |
| Methyl-β-cyclodextrin   | Lipid raft       | Removes cholesterol out of the plasma membrane | Interferes fluid phase endocytosis and CME |
| Filipin                | CIE              | Binds to cholesterol in the membrane | Toxic at higher concentration; inhibits CME |
| Cytochalasin D, latrunculin | Phagocytosis; macroendocytosis | Depolymerizes F-actin | Affects most endocytic pathways |
| Amiloride              | Macropinocytosis | Inhibits Na+/H+ exchange | May affect actin |
| **Pharmacological inhibitors** |                  |                |          |
| Dynasore               | CME              | Blocks GTPase activity of dynamin | Interferes with actin |
| Dynol, dynoves         | CME              | Blocks GTPase activity of dynamin I | Interferes with actin |
| Pitstop 2              | CME              | Interferes with binding of proteins to the N-terminal domain of clathrin | Most forms of CIE affected; causes decrease in PM mobility |
| **Genetic approaches** |                  |                |          |
| Dynamin mutant, Dyn K44A | CME            | Defective in GTP hydrolysis | Enhances fluid phase uptake |
| AP180C67               | CME              | Sequesters clathrin |          |
| Eps15 mutant           | CME              | Inhibits clathrin pits assembly | Other secondary effects (e.g., changes in gene expression) might occur due to overexpression and knocking down for several days |
| Clathrin Hub mutant    | CME              | Dominant negative mutant of clathrin |          |
| siRNA of clathrin      | CME              | Blocks formation of clathrin pits |          |
| siRNA of AP2           | CME              | Blocks formation of AP2-dependent clathrin pits |          |

CME, clathrin-mediated endocytosis; CIE, clathrin-independent endocytosis.
this compound appears to be fairly specific for blocking this process, macropinocytosis is generally easy to identify, even without the use of inhibitors.

Taken together, many, if not all, of the aforementioned inhibitors are agents that act in a non-specific way and thus caution is in order when interpreting the findings. Nevertheless, the effects of these widely available inhibitors on endocytosis of a particular cargo protein in comparison to other cargo proteins can help characterize or place that cargo as entering cells via a particular class of endocytosis.

Targeted Chemical Inhibitors of Endocytosis

In an attempt to develop potent and specific pharmacological inhibitors of endocytosis, chemical libraries have been screened for their ability to inhibit the GTPase activity of dynamin. Dynasore was the first compound identified to block dynamin’s GTPase activity in a biochemical assay and it has been shown to rapidly and reversibly block CME in cells.39 Dynasore has been widely used and has spawned the development of related compounds, the dynols and dyngoes, with different characteristics and affinities for dynamins 1 and 2.30 The development of these compounds was assisted by knowledge of the biochemistry and structure of dynamin. Overall, the advantage of these dynamin-targeted drugs is their rapid time of action and reversibility. Interestingly, dynasore treatment was shown to affect cortical actin, reinforcing earlier studies reporting roles for dynamin in the actin cytoskeleton.32 Hence the idea that dynasore acts solely as a CME inhibitor is misleading. Rather, the effects of dynasore may reveal other roles of dynamin.

Success with the chemical screen for dynamin inhibitors stimulated a search for compounds that would specifically block the formation of clathrin-coated vesicles at the cell surface. The N-terminal domain of clathrin heavy chain interacts with many cellular proteins, including amphiphysin33 and these interactions are critical for clathrin function. A chemical screen was performed to identify compounds that would block the binding of amphiphysin to the N-terminal domain of clathrin. The screen identified “pitstops” 1 and 2 as compounds that blocked this interaction in biochemical assays and also blocked CME in cells.34 The authors showed that endocytosis of shiga toxin was not affected by pitstop treatment, implying that the CIE mode of entry was not affected.34 Although this compound was a potent inhibitor of CME and presumably was working in part through this specific effect on blocking the binding of proteins to the N-terminal domain of clathrin, pitstop 2 has other unexpected targets in cells.

We found that pitstop 2 also inhibits CIE, and its ability to block CIE is still observed in cells where clathrin had been depleted by siRNA35 suggesting that this compound was affecting other cellular targets. Indeed, we found that treatment of cells with pitstop 2 results in a severe decrease in lateral mobility of cell surface proteins. Both integral membrane proteins and peripheral cytosolic proteins were essentially made immobile after addition of pitstop 2 to the cells.35 Ironically, cellular entry of shiga toxin, which von Kleist et al. used as a representative CIE cargo, could still occur in the presence of pitstop, probably due to the ability of shiga toxin to cross-link lipids and force entry into cells.36 Thus, despite the initial characterization of pitstop that indicated target specificity, in cells these new compounds appear to have additional targets that lead to severe changes in the cell surface.

Genetic Approaches to Endocytosis Inhibitors

To avoid the problem of non-specific effects of chemical inhibitors, genetic approaches have been used to inhibit endocytosis, in particular CME, by altering the expression of specific proteins. These have included the expression of mutant forms of critical proteins involved in endocytosis and siRNA-mediated depletion of these proteins. One of the first genetic approaches used was the expression of a mutant form of dynamin, K44A, patterned after a temperature-sensitive mutant in Drosophila. Expression of dynamin 2 K44A inhibits CME and it has been widely used to demonstrate that an endocytic event requires functional dynamin for fission.37 However, Damke et al. also noted that as a result of the block in CME invoked by Dyn2K44A the rate of clathrin-independent fluid endocytosis increased.38 Interestingly, recent reports have shown a more general role for dynamin in regulating cortical actin structure.31,32,39 Additional studies have targeted other regulatory proteins of CME such as the clathrin-associated proteins AP180 and Eps15. Expression of the carboxyl-terminal clathrin-binding domain of AP180 (AP180C)30 or a truncated form of Eps15 lacking the epsin homology (EH) domain40 can effectively block the formation of clathrin-coated pits. Expression of the C-terminal (Hub) region of clathrin also effectively blocks CME.32 Concerns have been raised that the overexpression of wild type and dominant negative forms of proteins might lead to many indirect effects and consequently investigators turned to methods to block CME by silencing the expression of players of the pathway such as clathrin heavy chain and the μ2 subunit of the AP2 adaptor complex. Knocking down these proteins by siRNA or shRNA can clearly indicate whether endocytosis of a particular type of PM protein requires clathrin and which adaptor protein.33,44 A drawback of this approach is that the time it takes to deplete a cell of these proteins can be considerable (3–7 d) and during this time the cell may adapt and even alter gene expression such that one cannot be assured that only CME is impacted. Also, loss of a protein like clathrin impairs trafficking at the TGN and to and from the lysosome,45,46 which might lead to defects in trafficking and lysosomal function. One clever approach to circumvent this drawback has been the use of the “knock-sideways” clathrin depletion scheme developed by Robinson and colleagues. Stable cell lines depleted of endogenous clathrin heavy chain and expressing a tagged form can be treated with rapamycin, causing the clathrin to be translocated to the mitochondria thus acutely depleting the cell of clathrin.47 The knock-sideways approach is acute and comes closest to a specific genetic inhibition similar to that imposed by the original temperature-sensitive mutant of dynamin in Drosophila. Drawbacks of this approach.
include the time required to prepare the cell lines expressing the two tagged proteins, the concern that during the translocation of clathrin other proteins may also be diverted and the possible cellular effects of inhibiting the mTOR pathway in cells.

**Perspective**

In conclusion, as our understanding of the different forms of endocytoses increases, it becomes more difficult to embrace one particular inhibitor as being diagnostic for a particular mode of endocytosis. Thus, results with endocytosis inhibitors, whether identified from a screen of a general endocytic process or from a screen of inhibitors for a specific biochemical activity, need to be interpreted carefully. The use of targeted pharmacological inhibitors (dynasore) appear to offer better choices than the non-selective chemical inhibitors (potassium depletion and cytokol acidification) but once again these experiments should be supported with corroborating evidence from the expression of mutant proteins and depletion of cellular proteins. The problems encountered with the “old” inhibitors, should not deter us from discovering “new” inhibitors. The identification and use of inhibitors that act quickly, specifically and reversibly will be invaluable as we set out to understand the physiological functions and mechanisms of different endocytic modes of cellular entry.

**Acknowledgments**

We thank L. Maldonado-Baez, D. Karabasheva, and J. Caviston for comments. This work was supported by the Intramural Research Program of the National Heart, Lung and Blood Institute, NIH.

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