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1 Endocytosis in context-dependent regulation of individual and collective cell 2 properties

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16 17 18 **Abstract**

19
20 Endocytosis allows cells to transport particles and molecules across the plasma
21 membrane. In addition, it is involved in the termination of signalling through receptor
22 downmodulation and degradation. This “traditional” outlook has been substantially
23 modified, in recent years, by discoveries that endocytosis and subsequent trafficking
24 routes have a profound impact on the positive regulation and propagation of signals,
25 being key for spatio-temporal regulation of signal transmission in cells. Accordingly,
26 endocytosis and membrane traffic regulate virtually every aspect of cell physiology,
27 and are frequently subverted in pathological conditions. Two key aspects of endocytic
28 control over signalling are coming into focus: context-dependency and long-range
29 effects. First, endocytic-regulated outputs are not stereotyped, but heavily dependent
30 on cell-specific regulation of endocytic networks. Second, endocytic regulation has
31 impact not only on individual cells, but also on the behaviour of cellular collectives.
32 Herein, we will discuss recent advancements in these areas, highlighting how
33 endocytic trafficking impacts complex cell properties, including cell polarity and
34 collective cell migration, and the relevance of these mechanisms to disease, in
35 particular cancer.

36 37 **[H1] Introduction**

38
39 Endocytosis is used by cells to internalize various types of molecules, including
40 nutrients, and fluids, which could not otherwise pass through the plasma
41 membrane^{1,2}. While this has probably represented the initial driving force behind its
42 emergence in evolution, the system has been exploited to actively regulate various
43 forms of communication within the cell and between the cell and its environment.
44 Signalling receptors, for instance, are internalized upon engagement by cognate
45 ligands and frequently targeted for degradation in the lysosome, resulting in long-term
46 signalling attenuation^{3,4}. In addition, regardless of their interaction with extracellular
47 moieties, many surface-resident molecules (mostly, but not exclusively, proteins) are

48 internalized and either degraded or recycled back to the cell surface, thereby providing
49 a mechanism through which the cell controls and adjusts its repertoire of plasma
50 membrane-resident molecules for various functional purposes. Yet, these are only
51 particular facets of endocytosis, whose impact on cellular homeostasis appears much
52 wider (see Supplementary Table 1).

53 A modern view of endocytic trafficking is that of a “vast program, deeply engrained
54 in the cellular masterplan and inextricably intertwined with signaling, which
55 constitutes the major communication infrastructure in the cell”¹. At the individual cell
56 level, for instance, endosomes represent major signalling stations. This is embodied in
57 the concept of the “signalling endosome”: a platform capable of sustaining signalling
58 by numerous mechanisms, including assembly of endosome-specific signalling
59 complexes, crosstalk, regulation of signal persistence in intracellular compartments,
60 and signal computing and resolution in time and space⁵. Endocytosis also controls the
61 execution of polarized cell functions through the redistribution of surface molecules
62 towards sites of polarized activities. In this case, fast and site-directed redistribution of
63 membrane proteins is not achieved by planar diffusion on the plasma membrane, but
64 rather by cycles of endocytosis and directed recycling⁶⁻⁹.

65 Although these activities are largely pertinent to the workings of individual cells, it
66 is becoming increasingly clear that also cellular collectives are controlled by
67 endocytosis. This is remarkable, as it entails that endocytic events occurring on the
68 level of a single cell must be synchronized, frequently spanning a distance of hundreds
69 of cells, to contribute to a coordinated behaviour¹⁰⁻¹³.

70 Mechanistically, endocytosis has long been considered a rather stereotyped
71 process, irrespective of the cell type and (to some extent) of the transported cargo.
72 This vision started to change with the realization that clathrin-coated pits, responsible
73 for clathrin-mediated endocytosis (CME), have varying compositions and contain a
74 plethora of non-obligatory components that confer specificity to the uptake process. In
75 addition, a wealth of non-clathrin endocytosis (NCE) routes exist, which also exhibit
76 context specificity, being present only in certain cell types^{2,15,16}. This heterogeneity in
77 endocytic mechanisms is reflected in differences in effector functions in cells, both at
78 the individual and the collective cellular level, which increasingly appear to depend on
79 the varying “endocytic landscapes” of different cell types. With this expression, we
80 mean the varying cellular composition in endocytic and trafficking proteins that can
81 determine the presence or absence of a certain endocytic route and/or confer a
82 different cargo specificity of the same endocytic route in different cells. In turn, this
83 may determine diverse biological outcomes in response to the same signalling input,
84 as a function of the cellular endocytic context.

85 Given the pervasive nature of endocytosis and its emerging roles in the control of
86 virtually every cellular phenotype (Supplementary Table 1), it is not surprising that its
87 subversion is relevant to human pathologies^{1,17,18}. As one example, in cancer, the
88 context-dependency of endocytic control over signalling might have a major role in the
89 migratory/invasive phenotype. Indeed, endocytic pathways seem to be preferentially
90 involved in the acquisition of pro-metastatic traits vs. other tumor properties^{6,18} (see
91 Supplementary Tables 2 and 3).

92 In this Review, we discuss the pleiotropic roles of endocytosis in cell regulation and
93 the importance of these mechanisms in physiology and pathology, especially cancer.
94 Many of these facets of endocytosis have been excellently reviewed elsewhere^{5-7,10-13}.

95 Herein, we will concentrate on an emerging trait: that of context-dependency of
96 endocytic regulation, with respect to different cell types and different stimuli they are
97 exposed to. We will also highlight the emerging view that endocytic trafficking,
98 although occurring at the single-cell level, has an impact on tissue-wide properties and
99 behaviours, such as tissue polarity and collective cell migration.

101 **[H1] Endocytic regulation in context**

102
103 In this section, we will use paradigmatic examples to illustrate how the greater context
104 – encompassing type of cargo as well as cell type and expression of proteins
105 comprising endocytic machinery – in which endocytosis operates influences its impact
106 on cell function in physiology and cancer. We will concentrate predominantly on the
107 regulation of signalling by receptor tyrosine kinases (RTKs) and G protein-coupled
108 receptors (GPCRs). For discussions on the impact of endocytic routes on intracellular
109 communication mediated by membrane contact sites¹⁹⁻²¹, on endocytosis in neuronal
110 communication and synaptic function²²⁻²⁴, and on the trafficking of adhesion
111 molecules, such as integrins and cadherins²⁵⁻²⁷, we refer the reader to several
112 exhaustive reviews.

114 **[H2] Differential regulation of signalling by distinct clathrin-coated structures.**

115 Different types of clathrin-coated structures exist at the surface of mammalian cells.
116 These include dynamic curved clathrin-coated pits that are heterogeneous in
117 dimension, dynamics and composition and are responsible for constitutive and ligand-
118 induced CME of different cargoes (Figure 1) as well as long-lived large flat clathrin
119 lattices, also known as coated plaques²⁸⁻³³, which act as signalling and adhesion
120 platforms³⁴, in addition to other functions (Box 1).

121 CME is characterized by a core of molecular components, prominently including:
122 clathrin; its main adaptor, adaptor protein 2 (AP2), which bridges cargoes to
123 clathrin^{35,36}; and the GTPase, dynamin, responsible for endocytic vesicle fission^{37,38} (see
124 example in Figure 1a). Depending on the cellular context and the type of cargo, other
125 adaptors — such as DAB2, ARH, epsins, EPS15 and EPS15L1 as well as arrestins — can
126 link cargoes to the clathrin machinery, producing clathrin-coated pits with varying
127 characteristics^{14,36,39} (see example in Figure 1b). In addition, AP2 levels are regulated in
128 a cell context-dependent manner, impinging on CME dynamics⁴⁰ and on fate(s) and
129 function(s) of signalling receptors (see below). Thus, the view that clathrin-coated pits
130 represent a uniform population is now outdated; it is now evident that distinct
131 clathrin-coated pits exist, with differential roles in the control of receptor fate and
132 signalling²⁸.

133 This is exemplified by the regulated endocytosis of GPCRs, which occurs
134 preferentially through subsets of distinct clathrin-coated pits characterized by: the
135 engagement of specific adaptors (such as β -arrestins)⁴¹; specific regulation, (such as
136 receptor ubiquitylation)⁴¹; increased surface residence time/slow internalization rates
137 (achieved by receptor–actin cytoskeleton interactions) (Figure 1c)⁴²⁻⁴⁴. GPCR
138 recruitment to specific subsets of clathrin-coated pits and cargo (ligand-bound GPCRs)-
139 dependent control of clathrin-coated pit dynamics and composition, are thought to
140 reduce competition with other CME cargoes, allowing the generation of clathrin-
141 coated pits with specialized functions⁴². Interestingly, β -arrestin-mediated CME varies

142 depending on the type of GPCR that is activated, as exemplified by the β -subfamily of
143 adrenergic receptors (AR). At variance with the other family members, the β 1-AR is
144 endocytosis-incompetent, meaning it is not internalized, yet, is still able to induce β -
145 arrestin accumulation and signalling in clathrin-coated pits⁴⁵⁻⁴⁷. Although the molecular
146 mechanisms responsible for this accumulation and activation of β -arrestin are not
147 completely elucidated, computational simulations suggest that the active
148 conformation of β -arrestin is stabilized by binding to the receptor tail and persists even
149 after detachment from the receptor; this conformational change renders β -arrestin
150 competent for binding to clathrin and AP2⁴⁷.

151 Similar to GPCRs, the epidermal growth factor receptor (EGFR) is also internalized
152 through distinct classes of clathrin-coated pits, distinguished by the presence or
153 absence of AP2 (Figure 1d), which differently regulate EGFR recycling/degradation and
154 thereby signalling, in a context-specific fashion (*e.g.*, fibroblasts *vs.* epithelial cells),
155 depending on the type of internalizing clathrin-coated pit (AP2-proficient or AP2-
156 less)⁴⁸⁻⁵¹. Specialized and context-specific functions of clathrin-coated pits also appear
157 to be conferred by the different members of the epsin family of endocytic adaptor
158 proteins⁵²⁻⁵⁴. Epsin1 (EPN1) and EPN2 are ubiquitously expressed and are implicated in
159 the CME of several plasma membrane receptors, such as the aforementioned EGFR, as
160 well as vascular endothelial growth factor receptors (VEGFR), NOTCH and WNT
161 receptors^{48,55-60}. By contrast, EPN3 shows a more restricted pattern of expression,
162 suggesting cell-specific functions^{52,61}. Indeed, EPN3 has a selective role in the
163 regulation of E-cadherin (CDH1) endocytosis and turnover and, when
164 amplified/overexpressed, contributes to breast cancer development through the
165 induction of a partial epithelial- mesenchymal transition (EMT) state (see also
166 subsection Endocytosis and EMT below)⁵³.

167 Different dynamin isoforms also mark distinct types of clathrin-coated pits.
168 Dynamin1 (DYN1) has traditionally been viewed as neuronal-specific and involved in
169 synaptic vesicle recycling, where its activity is regulated through phosphorylation–
170 dephosphorylation cycles⁶². By contrast, dynamin2 (DYN2) is constitutively active and
171 responsible for ubiquitous and constitutive endocytosis in all cell types⁶²⁻⁶⁶. Increasing
172 evidence, however, indicates that DYN1 can regulate CME also in non-neuronal cells,
173 and is selectively recruited to subsets of clathrin-coated pits, distinct to those
174 containing DYN2 (refs. ^{66,67}). In these cells, similarly to neurons, DYN1 is kept inactive
175 through GSK3 β -dependent phosphorylation^{36,65,68}. This inhibition is released upon EGF
176 stimulation, leading to DYN1 activation. Thus, at variance with DYN2, DYN1 appears to
177 be selectively activated by specific ligands. Interestingly, dynamins are deregulated in
178 cancer⁶⁹. DYN2 overexpression promotes invasiveness in different cancers, including
179 hepatocellular carcinoma, prostate and pancreatic cancer⁷⁰⁻⁷². DYN1 is aberrantly
180 activated in non-small cell lung cancer where GSK3 β function is inhibited, leading to
181 dysregulated CME of the EGFR and to the acquisition of migratory/invasive properties
182 (Figure 1e)^{66,73}. In addition, DYN1 activation, via calcineurin-dependent
183 dephosphorylation, upregulates endocytosis of death receptors, inhibiting apoptosis
184 and contributing to cancer cell survival⁷⁴.

185 Thus, the specialization of the machinery employed in clathrin-coated pits — driven
186 by the engagement of different cargoes, adaptor repertoire and cellular context (such
187 as expression of particular components of the machinery or their differential
188 regulation) — can impinge on cell behaviour in physiology and in cancer.

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[H2] Context-dependent regulation of signalling by NCE and integration of outputs.

NCE comprises several internalization mechanisms differing in morphology of the internalizing structures at the plasma membrane, machinery, cargo and regulation^{2,15,16}. Herein, we will not cover endocytosis through caveolae and refer the reader to extensive reviews on the subject^{75,76}.

The existence of so many NCE mechanisms might be, in part, due to inaccurate classification, as historically they were defined solely on the basis of clathrin-independence. Indeed – as we gain a deeper understanding of the specific features of NCE mechanisms, their overlapping characteristics and dependency on cellular context – it is becoming clear that some of them might represent “variations on a theme”, rather than truly distinct processes (Figure 2).

CLIC/GEEC (clathrin-independent carrier/GPI-anchored protein-enriched early endosomal compartment)⁷⁷⁻⁷⁹ is an NCE mechanism of central relevance: it relies on pleomorphic tubular endocytic intermediates and displays high endocytic capacity (Figure 2a)², rendering it suitable for mediating large plasma membrane rearrangements. CLIC/GEEC is thus implicated in plasma membrane turnover during cell spreading and migration⁸⁰ and in the regulation of membrane tension in adherent cells⁸¹. Indeed, upon a sudden reduction in plasma membrane tension, CLIC/GEEC is transiently activated to remove excess “floppy” membrane invaginations, thereby restoring the initial plasma membrane tensile state^{79,81}. The opposite is also true: perturbation of CLIC/GEEC directly decreases plasma membrane tension. In this circuitry, vinculin acts as the CLIC/GEEC-sensing mechanotransducer at focal adhesions^{81,82}. The mechanical buffering role of CLIC/GEEC differs from that of caveolae, which passively buffer the increase in tension also in non-adherent cells and seem to be critical for anchorage-independent growth⁸³. An interesting view of CLIC/GEEC is that of a prototype coatless NCE mechanism mediated by a conserved core of molecular components, while other NCE routes might represent specialized variants. For instance, fast endophilin-mediated endocytosis (FEME)⁸⁴ has many commonalities with CLIC/GEEC⁸⁵, but differs in that it is strictly regulated by receptor activation and displays rapid kinetics (Figure 2b)⁸⁴.

Another NCE mechanism, crucial to the fate and signalling of its cargo, is EGFR-NCE^{56,86}. This route internalizes the EGFR at high EGF doses in cellular contexts distinct from those where FEME and CLIC/GEEC are active (Figure 2c)^{86,87}. EGFR-NCE relies on contacts between the plasma membrane and the endoplasmic reticulum (ER), which might generate the pulling force for the formation of plasma membrane tubular invaginations. These contacts are also necessary for localized Ca²⁺ release from the ER, required for NCE vesicle fission⁸⁷. Since this route targets the EGFR to degradation and ultimately restricts its downstream signalling^{86,87}, it could represent a tumour suppressor pathway, as EGFR is commonly upregulated in cancer^{88,89}. Of note, a preferential degradative fate for internalized receptors might characterize other NCE mechanisms, as shown – for instance – in the case of the β and γ subunits of the IL2 receptor (Figure 2d)^{90,91}. However, it is worth pointing out that the degradative route does not necessarily preclude signalling from the internalized receptor, as shown for EGFR-NCE (see subsection Endomembrane dynamics controls collective motion).

Similarly to the EGFR, other RTKs are internalized via different endocytic routes. This is the case of MET (the receptor for hepatocyte growth factor), PDGFR (platelet-

236 derived growth factor receptor), VEGFR2, IGF1R (insulin-like growth factor 1 receptor)
237 and NGFR (nerve growth factor receptor)^{2,85}. For instance, IGF1R enters the cell via
238 CME or through caveolae depending on the dose of IGF1 (refs. ^{92,93}). Similarly,
239 depending on ligand concentrations, PDGFRB switches between CME and the
240 CLIC/GEEC pathway⁹⁴⁻⁹⁶. These RTKs are also internalized via FEME⁸⁴. Also in these
241 cases, the choice of entry route might specify the signalling outcome, as exemplified by
242 the VEGFR2 system^{84,97-99}. Postnatal retinal angiogenesis is controlled by spatially
243 regulated VEGFR2 endocytosis: mature vessels in the central retina display slow
244 VEGFR2 endocytosis, while, at the edge of the growing vasculature, tip cells show high
245 internalization rate, promoting the extension of the vessel network. In these cells,
246 VEGFR2 is internalized both via ephrin-B2/DAB2-mediated CME and endophilin-A2-
247 mediated NCE (resembling the FEME pathway). Consequently, CME of VEGFR2
248 promotes ERK signalling and vessel sprouting, whereas NCE regulates the signaling
249 effector PAK, front–rear polarization, and migration of tip cells⁹⁸⁻¹⁰⁰.

250 Macropinocytosis is a form of bulk NCE, which internalizes extracellular fluids and
251 macromolecules in large heterogeneous vesicles (Figure 2e)¹⁰¹. In mammalian cells, it
252 displays remarkable cell-specific variations in terms of molecular mechanism, fate of
253 the macropinosomes (recycled or degraded), and regulation¹⁰². In macrophages and
254 immature dendritic cells, macropinocytosis is constitutively active and critical for
255 antigen presentation¹⁰². Conversely, epithelial cells show barely detectable levels of
256 macropinocytosis in normal growing conditions. However, micropinocytosis is induced
257 in epithelia by growth factors and pathogens, and is upregulated in cancer¹⁰² by
258 activation of oncogenes, such as RAS and SRC¹⁰³⁻¹⁰⁵. Under nutrient-deprived
259 conditions, when the supply from the vasculature is insufficient, cancer cells can
260 scavenge nutrients (mainly albumin) from the extracellular environment by
261 macropinocytosis¹⁰⁶. This provides a sufficient intracellular pool of amino acids (in
262 particular glutamine) to meet the demands for energy production and nucleotide
263 biosynthesis.

264 Through macropinocytosis, cancer cells can also internalize necrotic debris (which
265 probably provides a higher quality source of nutrients), in a process called
266 necrocytosis. This debris is degraded, providing amino acids, fatty acids, sugars and
267 nucleotides used in anabolic pathways^{107,108}. In prostate cancer cells, in nutrient
268 deprivation conditions, necrocytosis is stimulated by PTEN loss, which cooperates with
269 active AMPK to stimulate RAC1 -dependent macropinocytosis. AMPK is also required
270 for RAS-driven macropinocytosis under nutrient restriction, indicating a general role of
271 this kinase in macropinosome formation¹⁰⁷. Context is again at play, since cancer cell
272 lines displaying similar oncogenic alterations vary in the extent and modality
273 (constitutive or inducible) of necrocytosis¹⁰⁸. Importantly, necrocytosis promotes
274 therapy resistance as it relieves stress induced by drugs targeting anabolic
275 pathways¹⁰⁸; thus, its inhibition could be exploited to target therapy resistant
276 tumours¹⁰⁹.

277

278 **[H2] Context-dependent role of endosomes in signalling and cellular responses.**

279 Although the activity of signalling receptors starts at the plasma membrane, it is now
280 abundantly clear that it persists in the various endocytic routes. In particular,
281 endosomes are critical signalling 'stations' that: sustain signalling originating from the
282 plasma membrane; are able to direct signalling through the recruitment of specific

283 effector molecules; create membrane microdomains where receptors are sequestered,
284 upon specific post-translational modifications, and sorted to their final fate, thereby
285 regulating signalling outcomes^{5,110}. The centrality of endosomes as signalling platforms
286 has been established for several signalling pathways in physiology and in cancer^{1,5}.
287 Here, we will focus on examples relevant to the idea of context-dependency.

288 GPCRs induce signalling from multiple cellular locations^{111,112}. At the plasma
289 membrane, GPCRs signal through the so-called “canonical pathway”, leading to
290 adenylyl cyclase activation and cAMP production. Upon prolonged ligand exposure, β -
291 arrestins bind to phosphorylated GPCRs, extinguishing the signals and inducing
292 receptor internalization (Figure 1c). β -arrestins also promote “non-canonical” GPCR
293 signalling by their scaffolding function and activation of signalling pathways centred on
294 ERK, p38 and JNK¹¹¹. Canonical and non-canonical GPCR signalling can occur at the
295 plasma membrane and from the endosomal station^{111,112}, in a context-dependent
296 manner (see subsection Differential regulation of signalling by distinct clathrin-coated
297 structures) and further demonstrated by in vivo studies of *Drosophila melanogaster*
298 gastrulation, where tissue level regulation of GPCR endocytosis by specific factors and
299 the dynamic partitioning of active receptors in different plasma membrane
300 compartments, *i.e.*, flat plasma membrane or invaginations, determine different
301 patterns of myosin II activation in mesodermal vs. ectodermal cells leading to
302 differential tissue specification^{113,114}. In addition, for some GPCRs, signalling was
303 shown to be transmitted also from the Golgi compartment^{115,116} and from the
304 nucleus¹¹⁷, regulating pathways distinct from those activated at the plasma
305 membrane. Finally, adenylyl cyclases themselves are regulated by trafficking and
306 localize to different cellular compartments, thus representing an additional variable for
307 the diversification of GPCR signalling¹¹⁸⁻¹²¹. The emerging picture is that of a complex
308 pattern of regulation, orchestrated by “localization biases” (*i.e.*, the cellular
309 compartment from which the signal originates) and dependent on cellular context,
310 where different membrane compositions and specific sets of adaptors/interactors
311 determine biological output¹²²⁻¹²⁴. This has enormous therapeutic implications. GPCRs
312 are easily druggable: approximately 35% of all approved drugs are directed against
313 them¹²⁵. Further advances in this field will largely depend on knowledge of spatial
314 resolution of GPCR signalling, to design more effective drugs with localization bias that
315 would target a specific signalling output¹²⁶. For instance, endosomal signalling of
316 GPCRs has been involved in chronic inflammation and pain, and revealed to be an
317 effective therapeutic target through the use of endosomally-directed GPCR
318 antagonists¹²⁷⁻¹²⁹. These types of drugs may also benefit cancer patients; this is a
319 particularly urgent need since, despite extensive knowledge of altered GPCR
320 signalling¹³⁰⁻¹³² in tumours, only a few anti-GPCR drugs are approved for therapy in this
321 disease¹³³.

322 Similarly, EGFR signalling is transmitted and regulated both at the plasma
323 membrane and from endosomes, where specific signalling platforms are assembled to
324 sustain and/or specify the signalling output (Figure 2c)^{1,5}. A clear example of this
325 regulation is represented by ERK1/2 signalling, which is involved in multiple cellular
326 outputs, including proliferation, migration and survival^{134,135}. Waves of ERK activation
327 were observed in epithelial cell sheets during wound healing¹³⁶ and their propagation
328 serves as a directional cue for collective cell migration¹³⁷. This might be due to
329 endosome-originated signals, as sustained ERK1/2 activation from endosomes is

330 necessary to exert a productive collective migratory response in breast epithelial
331 cells^{7,138}. Of note, in this instance, the EGFR is internalized through NCE-EGFR, a cell-
332 specific, and hence context-dependent, process (see subsection Context-dependent
333 regulation of signalling by NCE and integration of outputs). We will further discuss
334 these issues later (subsection Endomembrane dynamics controls collective motion).

335 In addition to the recruitment of endosomal signalling effectors, the modulation of
336 early endosome homotypic fusion represents another mechanism that regulates EGFR
337 signal amplitude and duration, and thereby cellular response. This is achieved by the
338 direct regulation of the fusion–fission machinery exerted by active EGFRs, leading to
339 modulation of the number of endosomes, which may act as quanta signalling
340 platforms that contain a pre-set number of activated receptors, thereby ultimately
341 determining context-dependent programmes driving cell proliferation versus cell
342 differentiation¹³⁹.

343 Finally, it is well established that endosomal membrane compartmentalization, that
344 is, the specification of specialized membrane microdomains on the limiting membrane
345 of endosomes, has a crucial role in receptor sorting and fate at endosomal signalling
346 stations^{110,140} with impact on the duration and specificity of signalling outputs. RAB
347 GTPases are pivotal in this process, as they contribute to the definition of endosomal
348 microdomains^{141,142}. An example of how RABs define endosomal microdomains was
349 elucidated studying RAB5 (refs. ^{141,143}): active RAB5, initially recruited to endosomes,
350 can further recruit its own GEF, creating a local, positive feedback loop of activation. In
351 addition, RAB5 directly modifies the local lipid environment stabilizing its own
352 membrane recruitment, thus providing another level of positive feedback regulation
353 that determines its patterning¹⁴⁴⁻¹⁴⁹. Notably, three highly conserved RAB5-encoding
354 genes exist, which share biochemical and biological activity. However, they can also
355 display specific functions depending on the cell context (as a function of their relative
356 levels of expression¹⁵⁰), signalling pathways that they impinge upon, and possibly
357 subcellular localization¹⁵¹⁻¹⁵³. Additionally, only the expression of RAB5A, but not
358 RAB5B or RAB5C, is elevated in breast carcinomas and is capable of reawakening cell
359 locomotion of jammed epithelial monolayers of breast tissue cells by differentially
360 impacting on endosomal signalling (see section Endomembrane dynamics controls
361 collective motion below).

362 A similar mechanism has been described for specification of endosomal
363 microdomains by the small GTPase CDC42 (refs. ^{154,155}) and for organization of plasma
364 membrane microdomains by K-RAS¹⁵⁶. Such non-linear dynamics of recruitment and
365 activation are typical of self-organizing systems that form spatial patterns on
366 membranes¹⁵⁷, possibly representing a general mechanism for spatial organization of
367 GTPases inside cells and for their role in generating membrane microdomains critical
368 for receptor fate and sorting along the endocytic pathway¹⁵⁸.

369 **[H1] Endocytosis in tissue morphogenesis**

370
371
372 In this and in the following section, we will discuss the role of endocytosis and
373 endocytic trafficking in the control of epithelial cellular collectives. In this section, we
374 will highlight emerging principles relative to the role of endocytosis in tissue
375 organization by focusing on the establishment of apical-basal cell polarity during

376 lumenogenesis of epithelial sheets, and tissue extension and cell shape morphogenesis
377 in embryo development and adult epithelia.

378

379 **[H2] Polarized transport controls apical-basal cell polarity and lumenogenesis**

380 A defining feature of epithelial and glandular tissues is a pronounced apical–basal
381 asymmetry, which is important for establishing barrier function, polarized
382 transport^{159,160}, and successful lumenogenesis/tubulogenesis that gives these tissues
383 their shape^{161,162}. Failure of these mechanisms results in altered epithelial function,
384 dysmorphologies and malignant transformation¹⁶³.

385 Directed membrane trafficking is critical for the polarized distribution of molecules
386 in the cell and for the generation of specialized membranous domains in individual
387 epithelial cells, but also for the coordination of epithelial polarization during lumen
388 formation and the morphogenesis of glandular tissues¹⁶⁰⁻¹⁶². There is extensive
389 interplay between membrane trafficking and cell polarity (illustrated in Figure 3),
390 whereby epithelial polarization involves not only the formation of apical and
391 basolateral domains, but a whole topological re-organization of intracellular trafficking
392 pathways along the apical-basal axis.

393 Two major membrane routes are used to establish apical-basal polarity in individual
394 cells¹⁶⁴. The first one involves recycling of polarity molecules from the plasma
395 membrane followed by their sorting and targeted delivery to a polarized surface. The
396 second one involves regulated trafficking of newly synthesized polarity determinants.
397 Both of these routes are accomplished by polarized membrane trafficking regulated by
398 RAB GTPases¹⁶⁵, which are key regulators of membrane trafficking also in non-
399 polarized cells; however, of the ~70 mammalian RABs, only a dozen has been
400 specifically implicated in regulating apical-basal cell polarity¹⁶⁶. Furthermore, while
401 some RABs operate regardless of environmental conditions, others control apical-basal
402 cell polarity solely during 3D morphogenetic events, but not in 2D, indicating context-
403 dependency¹⁶⁶.

404 Central to the establishment and maintenance of apical-basal polarity is the correct
405 topological distribution of membrane-associated apical-basal cell polarity complexes
406 (Supplementary Box 1) and lipid domains, organelles, and cytoskeletal structures¹⁶⁷
407 (Figure 3). A detailed description of apical-basal cell polarity complexes, their
408 interaction with trafficking components, and their routes, is outside the scope of this
409 review, but several excellent reviews on this topic are available¹⁶⁸⁻¹⁷⁰.

410 How these events, occurring on a cellular level, orchestrate the morphogenesis of
411 tube-shaped and sac-shaped structures is the object of intense scrutiny. A model
412 system that reproduces the early steps of apical lumen formation and expansion is
413 MDCK cells in 3D culture^{160,171}. Following cell division, individual MDCK cells remain
414 connected by a midbody, which serves as the spatial instructive cue for the creation of
415 an apical membrane initiation site, thus establishing the location of the nascent lumen
416 (Figure 4a). This process involves multiple trafficking steps. The first establishes the
417 structural identity of the apical membrane initiation site through the coordination and
418 re-organization of microtubules and branched actin filaments, via RAC1 GTPase-WAVE
419 pathway and the adhesion protein, cingulin. This latter protein binds the central
420 spindle microtubules and is recruited to the apical membrane initiation site via a direct
421 interaction with the midbody and the tight junction protein, ZO1. Cingulin then can
422 activate Arp2/3 complex to promote branched polymerization, which in turn recruits

423 more ZO1 — and consequently more cingulin — installing a positive feedforward loop
424 critical for reinforcing the apical localization of these proteins¹⁷². Cingulin also serves
425 as a tethering platform for the anchoring of apical endosomal vesicles by directly
426 interacting with FIP5, a RAB11-interacting protein^{172,173} (Figure 4b). Consequently,
427 transmembrane proteins, such as podocalyxin (a classical apical marker) and Crumbs
428 polarity complex— which undergo transcytosis from the extracellular matrix abutting
429 surface to the apical membrane initiation site via RAB11-marked endo/exocytic
430 structures (which are most likely also positive for RAB8; see discussion in the following
431 paragraph) — can be hooked to the apical membrane initiation site through
432 interaction with cingulin and FIP5. Alternatively, association of apical polarity proteins
433 with the apical membrane initiation site could occur through binding to RAB35, which
434 in addition to cingulin has been proposed to act as an anchoring platform for polarity
435 determinants^{171,174}. Although the exact molecular role of RAB35 is still debated, there
436 is a consensus that its function is important for the correct transport of apical and
437 basolateral determinants^{166,175}.

438 Additionally, RAB11 can also activate RAB8 through the GEF RAB3IP (aka Rabin8) on
439 Rab11 and RAB8 double-positive endosomes. Both RAB8 and RAB11 can then recruit
440 myosin motor, MYO5B and the EXOC6 (aka Sec15A) exocyst subunit in order to
441 transport recycling vesicles that carry critical polarity determinants to the apical
442 membrane initiation site¹⁷⁶, and they also interact with EXOC5 (aka Sec10) at the apical
443 membrane initiation site to promote vesicle docking (Fig. 4B, inset)¹⁷¹.

444 Finally, apical membrane initiation site is characterized by a unique phospholipid
445 composition, being enriched in di-phosphorylated phospholipids, including PI(3,4)P2
446 (ref. ¹⁷⁷) and PI(4,5)P2 (ref. ¹⁷⁸). This lipid environment is critical for the asymmetric
447 recruitment of annexin-2 and of the CDC42 apical polarity complex, which is in turn
448 capable of binding key apical polarity determinants, PAR3 and PAR6¹⁷⁸. These lipids
449 also ensure the correct localization and function of membrane-curvature sensor
450 proteins. One such protein, IRSP53 (aka BAIAP2), binds to PI(4,5)P2-rich membranes. In
451 the early stages of cyst formation, IRSP53 re-localizes from the plasma membrane to
452 endosomal structures and to the apical membrane initiation site, where, through its I-
453 BAR domain (which binds to negatively curved membranes), it ensures the integrity
454 and shape of the opposing membranes of the two neighbouring cells for correct lumen
455 generation. Loss of IRSP53 results in the formation of aberrant cytoplasmic bridges
456 that interconnect adjacent cells and interrupt the continuity of the nascent lumen,
457 yielding aberrant multiple lumens¹⁷⁹.

458 It is still unclear how, after the establishment of the apical membrane initiation site,
459 a lumen develops and expands. It is likely that a balance of forces between tensile
460 junctional elements and cell–substrate adhesion, contributes to the position and shape
461 of the de novo lumen^{180,181}. Additionally, lumen opening might be driven by: reduced
462 actomyosin contractility at adherens junctions, which allows fluctuation of the
463 membrane at the interface between two adjacent cells; and/or by increased
464 intraluminal hydrostatic pressure, which follows the establishment of tight junctions
465 and the polarized flux of water through ion channels (Figure 4c). Notably, as the lumen
466 inflates the cell volume decreases, suggesting that the system is isochoric and
467 proceeds through volume conservation between the cells and the luminal space¹⁸².

468 An invariable feature of carcinomas is the loss of epithelial stereotypical
469 architecture. Observations in *D. melanogaster* identified polarity regulators as tumour

470 suppressors. By contrast, studies in mammals suggest oncogenic dominant roles of
471 these proteins¹⁸³. A number of polarity determinants – including PARD6B, SCRIB, PRKCI
472 and DLGs – are amplified or display aberrant cellular distribution in cancer¹⁸³. This
473 suggests that carcinomas might exploit polarity proteins to promote their progression.
474 For example, polarity proteins, when overexpressed, might switch from controlling
475 apical-basal cell polarity to regulate subcellular polarity (intracellular asymmetry),
476 which can improve cellular fitness to execute functions such as proliferation,
477 apoptosis, stress adaptation, stemness and organelle biology¹⁶³. Membrane trafficking
478 processes are expected to be directly involved in the regulation of this polarity switch
479 and may thus contribute to the onset and progression of carcinomas.

480

481 **[H2] Tissue organization by mechanosensitive membrane trafficking.**

482 At variance with apical-basal cell polarity, planar cell polarity is a system of information
483 that provides cues along the axis parallel to the epithelial plane, and requires the
484 establishment of asymmetry within individual cells and the alignment of these
485 asymmetries across dozens or hundreds of cells. This is achieved through the proximal-
486 to-distal distribution, along the plane, of six planar cell polarity proteins
487 (Supplementary Box 1)¹⁸⁴⁻¹⁸⁷. The planar distribution of these proteins is driven by a
488 series of feedback loops, typically operative at the single cell level and dependent on
489 endocytosis and polarized membrane trafficking, as described in recent reviews^{188,189}.
490 Here, we will focus on additional, more recently described, endocytic mechanisms that
491 respond to tissue mechanics and control individual cell shape in the context of
492 supracellular, tissue-level organization.

493 One such mechanism involves the control of cell geometry and of the actomyosin
494 force-generation machinery, which regulate tissue rearrangements and size, as
495 elucidated in several developmental systems¹⁹⁰, including germband extension in *D.*
496 *melanogaster*^{191,192}. Here, tissues undergo elongation to attain their final shape
497 through a process of cell intercalation that drives the so-called convergent extension
498 — a cell rearrangement whereby cells intercalate in an oriented fashion causing tissues
499 to shrink along the dorso-ventral axis and to elongate along the antero-posterior axis
500 (Figure 5a). At the level of neighbouring cells, dorso-ventral intercalation is the result
501 of the contraction of the interface between two dorso-ventral neighbours, followed by
502 the extension of the interface between the antero-posterior neighbours¹⁹³. This occurs
503 through the combination of a tension-producing actomyosin network and adherens
504 junctions, which are governed by spatially restricted membrane trafficking, specifically
505 occurring along the surface of neighbouring cells¹⁹⁴. Indeed, during germband
506 extension in *D. melanogaster*, clathrin and AP2 become enriched at the contractile
507 interface of neighbouring cells in the antero-posterior direction¹⁹⁵; here, polarized
508 actomyosin contractility drives clustering and internalization of E-Cadherin, which
509 leads to junctional shrinkage by reducing local cell–cell adhesion¹⁹⁵.

510 The interplay between actomyosin contractility and endocytosis goes beyond
511 regulation of planar cell polarity, impinging also on the homeostatic control of tension
512 state and cell size in tissues, as occurs during the contraction of amnioserosa during
513 dorsal closure in *D. melanogaster*¹⁹⁶, and in adherens junction remodelling during
514 morphogenetic cell shape changes in mammalian epithelial tissues¹⁹⁷. In the former
515 example (Figure 5b), adherens junctions tune their length and maintain junctional
516 straightness and tension while the whole tissue undergoes a global contraction¹⁹⁸.

517 When the tissue contracts, junctions shorten resulting in excess plasma membrane
518 that folds inwards, causing a reduction in membrane tension. Junctional and
519 membrane tension is restored by promoting removal of excess or folded plasma
520 membrane at junctions, through endocytosis¹⁹⁸. The opposite occurs when junctions
521 are stretched, which results in the redistribution of actomyosin to adherens junction
522 for reinforcement, which increases tension that is then restored through inhibition of
523 endocytosis. Similarly, in various mammalian epithelial monolayers, endocytic
524 pathways act as mechanosensitive networks to stabilize junctional tension and length.
525 In these systems, junctions are dynamically remodelled by short pulses of RHOA-
526 mediated actomyosin contractility¹⁹⁷. For pulses of RHOA activation that induce a
527 contraction capable of overcoming a defined strain threshold (*i.e.*, the maximal
528 deformation that epithelial cells within an epithelial tissue can undergo without
529 rupturing) associated with morphogenetic tissue shaping, junctions are permanently
530 relaxed and subsequently shortened in a process that requires endocytosis¹⁹⁷.

531 These findings highlight that changes in cell and tissue size are frequently driven by
532 pulsatile processes, where a directed dynamic process (*e.g.*, shortening of a junction)
533 alternates with periods in which force-generating networks are restored. This cycling
534 requires consolidation or “ratcheting” to lead to permanent changes, by preventing
535 the reversal of cell shape once the contractile network has dissipated. Membrane
536 trafficking operates as such a ratchet to ensure the removal of slackened membrane,
537 as mediated by RAB35 in germband extension^{199,200}. At the antero-posterior interface,
538 RAB35 accumulates in compartments that dynamically grow and shrink on the minute
539 timescale. Genetic removal of RAB35 does not prevent the contractile cycles of
540 junctional length but impedes the permanent shortening of the cell interface (Figure
541 5b).

542 It is noteworthy that trafficking regulators, such as RAB35, appear to play multiple
543 roles in controlling the proper abscission of dividing cells²⁰¹ and integrin trafficking
544 during glandular morphogenesis¹⁶⁶, in addition to their role in tissue adaptation to
545 mechanical stresses. Indeed, the ability of tissues to adapt and respond to mechanical
546 perturbations has emerged as a key mechanism to ensure proper organ and
547 organismal development; whereas, failure to mount this mechanoresponse can lead to
548 patterning and structural abnormalities in the developing embryo. RAB35 also controls
549 the ability of cells to internalize nutrients through macropinocytosis²⁰² and was
550 identified as a pro-oncogenic mutated RAB capable of interacting with the regulatory
551 subunit of PI3K (p85) — a key oncogene — enhancing the activity of this kinase²⁰³.
552 How this pro-oncogenic function of RAB35 impinges on glandular morphogenesis and
553 tissue mechanoadaptation is unknown, but it is tempting to speculate that RAB35
554 might be at the centre of a trafficking network that coordinates tissue shape, division
555 and growth, thus directly influencing tumorigenesis when its activity is perturbed²⁰⁴.

556 557 **[H1] Endocytosis in collective cell migration**

558
559 One of the best characterized functions of endocytosis, beyond simply uptake of
560 extracellular cargo, is in the regulation of cell migration in physiology and in invasive
561 cell migration in cancer, as discussed in recent reviews^{18, 205-207}. In this section, we will
562 focus on the regulation of cell migration by endocytosis, in the context of collective
563 modalities of migration with emphasis to their subversion in cancer (see also

564 Supplementary Table 2). In particular, we will discuss three emerging aspects of
565 endocytic regulation of collective migration: promoting persistent migration of cell
566 monolayers, modulation of tissue mechanical and material-like properties and
567 induction of EMT.

568

569 **[H2] Endomembrane dynamics controls collective motion.**

570 A function of endocytosis is the coordination of soluble and mechanical cues during
571 individual cell migration^{7,10,208-210}. Frequently, however, cells are embedded in
572 multicellular ensembles, where the motion of the whole cell collective is overarching
573 with respect to individual cell locomotion. This occurs during the development of
574 epithelial layers (skin, digestive or respiratory tracts), in glandular organs, or during
575 wound healing. Collective motion may also take the form of cell clusters that maintain
576 cohesive interactions while migrating, as in the case of border cell migration in *D.*
577 *melanogaster*²¹¹ or during the dissemination of epithelial cancers^{212,213}. In these
578 situations, individual cells must coordinate their movement with that of their
579 neighbours, while keeping tight cell–cell interactions.

580 Multicellular migration is ruled by the biochemical and physical interactions that
581 cells establish with one another and their environment²¹⁴. Indeed, physical forces
582 exerted by individual cells on their substrate or propagated, long-range, in
583 multicellular cohorts through cell–cell adhesions (via cadherins at adherens junctions),
584 are principal determinants of multicellular dynamics²¹⁵. Establishing and maintaining
585 adherens junction strength, dynamics and polarity during cell migration is achieved by
586 trafficking membrane cadherins. One example here is collective migration of
587 astrocytes — major glial cells of the central nervous system — during development
588 and wounding²¹⁶. Astrocytes migrate collectively through maintenance of connections
589 via N-Cadherin (aka CDH2). Developmental or wounding cues induce polarization of
590 astrocyte monolayer with the establishment of “leader cells” that determine the
591 migratory front. These leader cells extend lamellipodia, where actin treadmilling and
592 actomyosin contractility generate a retrograde flow of actin. N-Cadherin travels
593 backwards along this flow (retrograde transport along the plasma membrane). At the
594 back, N-Cadherin is endocytosed and through polarized trafficking delivered to the
595 front, to form new junctions²¹⁶, virtually installing a membrane trafficking treadmilling
596 process for N-Cadherin (Figure 6a), which can support persistent and cohesive
597 migration of the entire cell sheet.

598 Another example is the collective motion of the neural crest in *Xenopus laevis*²¹⁷,
599 where cells become fully migratory before they complete cell–cell dissociation,
600 thereby moving as collective clusters. A signalling mechanism, affecting junctional
601 adhesion strength, is triggered by lysophosphatidic acid (LPA)²¹⁸. In the neural crest,
602 the interaction of LPA with its receptor, LPAR, specifically affects collective motion by
603 modulating the extent of cell–cell cohesion through internalization of N-Cadherin²¹⁹.
604 Neural crest collectives undergo a transition from a solid-like state, where local cell
605 rearrangements (the motion of a cell with respect to its neighbours) are not permitted,
606 to a fluid-like state, where reduced Cadherin adhesion allows local cell
607 rearrangement²¹⁹. This results in fluidization of neural crest clusters, which can more
608 easily migrate into confined spaces, while retaining cell–cell cohesion.

609 LPA is a signalling lipid, abundant in blood, skin and ascitic fluids, which acts both as
610 mitogen and chemotactic agent. Melanoma and pancreatic cancer cells can

611 breakdown LPA, generating outward-facing local gradients — low in the tumour and
612 high in the surrounding environment — that guide metastatic cell dispersal through a
613 self-generated LPA chemotactic gradient^{220,221}. Engagement of the LPAR stimulates its
614 endocytosis in early endosomes. N-WASP, a promoter of branched actin
615 polymerization, together with the endocytic protein SNX18, coordinate recycling of
616 internalized LPAR from RAB11–RAB8-double positive endocytic tubules back to the
617 plasma membrane preventing its degradation. The N-WASP-mediated LPAR recycling
618 then ensures constant replenishment of the receptor at the plasma membrane
619 allowing persistent RHOA activation, resulting in increased cell contractility, generation
620 of traction forces and matrix remodelling for efficient cell migration. Loss of N-WASP
621 affects invasiveness of pancreatic cancer cells pointing to N-WASP-dependent
622 chemotaxis to LPA gradients as a motivator of egress from primary tumours and
623 tropism towards metastatic sites²²¹. In HeLa cells, the hyperactivation of the LPAR
624 might also occur through trapping the receptor at the plasma membrane in coated
625 plaques²²² (Box 1): reduced LPAR internalization prolongs its downstream signalling,
626 representing an alternative mechanism to recycling for regulating actomyosin-based
627 contractility. Whether coated plaques are operative in pancreatic cancer cells remains
628 an issue for future investigations; notably, these structures are increased in cells plated
629 on stiff substrates³³, suggesting that the hyper-stiff, stromal-rich, microenvironment of
630 pancreatic cancer cells might favour their formation.

631 These findings highlight how an integrated biochemical and physical perspective is
632 necessary for a holistic comprehension of the dynamics of multicellular entities and
633 tissues. Such dynamics can be better understood by considering cell collectives as
634 ensembles of “active particulate matter” that are, nevertheless, governed by structural
635 and dynamic physical properties typical of amorphous viscoelastic materials^{223,224}.
636 During tissue growth, cells are free to move, as in a fluid, but their motion becomes
637 constrained as cell density increases. At a critical cell density, motility ceases and
638 collectives rigidify undergoing a liquid-like to solid-like transition referred to as tissue
639 jamming²²⁵⁻²²⁹. This transition is thought to promote the development of elasticity and
640 of barrier properties, but also to act as a suppressive mechanism for the growth of
641 oncogenic clones.

642 Recently, it was shown that global perturbations of endocytosis — achieved by
643 modulating the master regulator of early endosome biogenesis, RAB5A — impinges on
644 biomechanical properties of cellular collectives. Increased RAB5A levels were sufficient
645 to re-awaken the motility of “jammed” epithelial monolayers by triggering millimetre-
646 scale, coherent and ballistic locomotion of multicellular streams that flow like
647 “flocking” fluids²³⁰, through the modulation of EGFR signalling (Figure 6b and 6c). In
648 mammary epithelial cells, RAB5A induced EGFR-NCE, leading to accumulation of
649 activated receptors in endosomes, whose size and number were remarkably
650 elevated²³¹, and to elevation of endosomal ERK1/2 signalling, in keeping with the
651 concept that endosomes act as quanta-like platforms where phosphorylated EGFR can
652 be packaged at constant mean amounts¹³⁹. In turn, this caused the hyper-
653 phosphorylation of the Arp2/3 nucleating promoting factor, WAVE2, fuelling the
654 extension of lamellipodia and directed cell motility²³¹ (Figure 6b).

655 Tumour cells can exploit this endocytic-mediated regulation of cell mechanics to
656 facilitate their collective dissemination. Consistently, mammary tumour spheroids
657 overexpressing RAB5 display persistent angular rotation and elevated local cell

658 rearrangements²³¹. This leads to the generation of large traction forces on the
659 surrounding extracellular matrix, resulting in its remodelling and the generation of
660 tracks and channels into which tumour cells invade collectively. In another study,
661 lowering of cell–cell adhesion with concomitant increase in 3D matrix confinement
662 have been shown to affect unjamming transition and to foster collective invasion in
663 breast cancer models²³². These dynamics are reminiscent of those during body axis
664 elongation in the zebrafish embryo, where a gradual solid-like to fluid-like transition is
665 critical for promoting the elongation of the entire body axis²³³. Within this context,
666 membrane trafficking might be a general mechanism to control the transition from a
667 solid-like/jammed to a liquid-like/unjammed state, leading to progressive fluidization
668 and collective motion.

669

670 [H2] Endocytosis and EMT

671 In addition to solid-to-liquid transition, another process contributing to collective
672 motion of tissues is EMT. Unlike solid–liquid transitions, however, which rely on
673 mechanical remodelling, EMT primarily involves changes of individual cell identity and
674 state. Accordingly, recent evidence established that unjamming and EMT are very
675 distinct processes differing not only in molecular machinery, but also in physical,
676 dynamic, geometrical and structural properties²³⁴. Indeed, unjamming and EMT might
677 be viewed as complementary gateways to cell migration, driving the escape of
678 epithelial cells from a static, rigid and sessile state.

679 EMT is a process in which cells shift from a sessile epithelial to a migratory/invasive
680 mesenchymal-like state²³⁵. Physiologically, it is involved in morphogenetic events,
681 including gastrulation and neural crest migration²³⁶. Pathologically, EMT is exploited by
682 cancer cells to acquire invasive/metastatic ability²³⁷⁻²³⁹ — a process also connected
683 with the acquisition of a cancer stem cell-like phenotype and drug resistance²⁴⁰⁻²⁴³.

684 Various mechanisms act as initiators of EMT in cancer, including: growth factors and
685 cytokines, components of the extracellular matrix and mechanical cues. Given the
686 involvement of endocytosis in cell response to all these factors, it is not surprising that
687 it controls EMT, at multiple steps in the process^{244,245}. The common distal effectors, on
688 which all these signalling pathways converge, are a set of transcription factors, which
689 include Snail, Slug and ZEB1/2²⁴⁶⁻²⁴⁸. These transcription factors orchestrate the
690 transcriptional repression of epithelial-specific genes that are critical in cell adhesion
691 (*e.g.*, E-Cadherin), and the transcriptional activation of mesenchymal genes that
692 reshape the cytoskeleton and the plasma membrane to permit migration (*e.g.*, N-
693 Cadherin and vimentin). Thus, the actual enactment of the EMT program is generally
694 viewed as a transcriptionally-driven process. This view is changing.

695 The key to understanding this renewed vision is the finding that, at least in some
696 cases, EMT is not a binary process involving cell state conversion, but rather involves a
697 plastic state in which cells can exhibit intermediate phenotypes, retaining some
698 epithelial characteristics while acquiring some mesenchymal ones^{236,237}. This state,
699 called partial or plastic EMT (P-EMT), appears to be advantageous for cancer cells to
700 oscillate between a mesenchymal phenotype, necessary to migrate out of the primary
701 tumor, and an epithelial identity, necessary for survival and proliferation once they
702 reach the final metastatic niche²⁴⁹. Cell context is emerging as a determining factor for
703 the decision to adopt a full EMT vs. a P-EMT state: in this latter instance, endocytic
704 mechanisms appear paramount over transcriptional ones. Cancer cells adopting the

705 “endocytic” modality to achieve P-EMT, lose their epithelial phenotype through
706 surface protein internalization rather than through the action of EMT-promoting
707 transcription factors²⁵⁰. In particular, E-Cadherin is actively internalized and
708 sequestered in RAB11-positive recycling endosomes, consistent with the notion that
709 the molecular alteration behind the cell state shift resides in a blockade of the
710 physiological recycling of E-Cadherin to the plasma membrane.

711 In this contention, we note that the “endocytic mechanism” of P-EMT has the
712 characteristics of being more rapidly reversible (through re-induction of recycling) and
713 perhaps less extensive than the transcriptional reprogramming, thus being more adapt
714 to the plastic state of P-EMT. It is also interesting that cells adopting the endocytic-
715 based P-EMT modality, exhibit a collective, rather than individual, mode of migration,
716 characteristic of cells displaying full EMT²⁵⁰. Furthermore, it has been shown that
717 cancer cells adopting different degrees of EMT, tend to metastasize to different sites²⁵¹
718 raising the interesting question of whether a primary alteration of endocytic
719 mechanisms, in some tumours, might underlie metastatic organotropism.

720

721 **[H1] Conclusions and perspectives**

722

723 Since the first hints, more than a quarter of century ago, that endocytosis could sustain
724 and/or diversify cell signalling²⁵²⁻²⁵⁴, rather than merely extinguishing it, a wealth of
725 literature has accumulated, which not only supports this concept but extends it to
726 virtually every area of cell regulation^{1,255}. The field is now mature to address the next
727 question: that of the significance of cellular context of endocytic regulation.

728 In this vision, the same signalling circuitry could elicit substantially different
729 biological outputs based on the “endocytic landscape” in which it is embedded and
730 regulated by. Although the concept of the existence of specific endocytic landscapes in
731 cells, herein proposed, is at present molecularly vague, we would like to draw one
732 example that may support it. The EGFR is internalized by two types of CME, by FEME
733 and by EGFR-NCE, which all impact differently on receptor fate, signalling properties,
734 and timing of signals (see section Endocytic regulation in context). They also display
735 cell context specificity. Thus, cell-specific “endocytic landscapes” are projected to
736 differentially affect the EGFR signalling output — an issue of great pathophysiological
737 relevance, given the importance of this RTK to cancer^{88,89}.

738 There are clever approaches being pursued to investigate this hypothesis. One is
739 that of exploiting isogenic models of cells differentiated in vitro from multipotent cells
740 (embryonic stem cells or induced pluripotent cells). Using this approach, it was shown
741 that isogenic, diversely differentiated cells display remarkable variations in the
742 dynamics and structure of clathrin coats⁴⁰. These differences hinged on context-
743 dependent variations in AP2 levels and in the requirement for PI3K activity⁴⁰. Another
744 approach is to employ organoids that recapitulate the complex differentiation patterns
745 that occur in vivo²⁵⁶. This approach, coupled to advanced imaging to detect live
746 endocytic dynamics and to single-cell sequencing, to decipher the endocytic landscape
747 of individual cells, should substantially improve our understanding of cell context in
748 individual and collective cellular behaviour.

749 The same view could be extended to the domain of endocytic alterations in cancer.
750 In this case, perhaps, the simple idea of endocytic proteins acting as oncogenes or
751 tumour suppressors, albeit valid in some cases^{1,17,18}, might be too narrow. The

752 evidence that the endocytic context might skew the output of a signalling pathway
753 towards different biological outcomes, opens up the possibility that differences in the
754 expression of groups of endocytic proteins (the “endocytic landscape”) might render
755 cells more or less prone to respond to oncogenic insults and, therefore affect the
756 propensity of cells for tumorigenic transformation. In this contention, we note that
757 genetic alterations of endocytic proteins are infrequent in cancer^{1,17,18}; rather, the
758 more common alteration is their over- or under-expression, which often correlates
759 with metastasis and disease outcome^{6,18} (see also Supplementary Tables 2 and 3) To
760 extend our knowledge, we should go beyond the “one protein at the time” approach
761 and embrace comprehensive profiling of tumours, to identify endocytic landscapes
762 that might define the propensity of certain tumors (or subgroups of tumours) to adopt
763 different individual or collective cell behaviours (even in response to the same
764 signalling input), especially those relevant to the invasive/metastatic phenotype.

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777 included in Supplementary Information.

778 **Competing interests**

779 The authors declare no competing interests.

780 **Figure Legends**

781 **Figure 1. Heterogeneity of clathrin-coated pits.**

782 Clathrin-mediated endocytosis (CME) is a heterogeneous process, with a variety of
783 clathrin-coated pits (CCPs) that differ in their composition, leading to different fates of
784 the endocytic cargo. **a.** In physiological conditions, transferrin (Tf) receptor (TfR)
785 internalization depends on clathrin adaptor AP2 only⁴⁸⁻⁵¹. After endocytosis, the iron
786 bound to Tf is released in an endosomal compartment and the Tf–TfR complex is
787 recycled back to the surface²⁵⁷. **b.** Low-density lipoprotein (LDL) receptor (LDLR)
788 endocytosis relies on the cargo-specific adaptors, DAB2 and ARH, in addition to AP2
789 (refs. ²⁵⁸⁻²⁶³). In the endosomal compartment, the LDL moiety is released and
790 committed to lysosomal degradation (and the LDL-bound cholesterol is released),

799 while the “empty” receptor is recycled to the cell surface²⁶⁴. **c.** Agonist-activated G
800 protein-coupled receptors (GPCRs) bind heterotrimeric G-proteins (α , β , γ) which
801 triggers “canonical” G-protein-mediated signalling at the plasma membrane¹¹¹.
802 Receptor desensitization occurs by phosphorylation (P) of the active receptor by GPCR
803 kinases (GRKs) and subsequent binding of β -arrestin, resulting in dissociation of the G-
804 protein complex and of its interaction with the receptor. Further desensitization occurs
805 when the β -arrestin bound GPCR is sequestered into distinct CCPs and internalized, to
806 be either recycled back to the plasma membrane or degraded in the lysosome (not
807 shown)⁴¹. Receptor ubiquitylation (Ub) and PDZ-containing scaffold proteins, linking
808 the GPCR to the actin cytoskeleton, determine increased surface retention of GPCR-
809 containing CCPs and their slower kinetics, promoting cargo clustering and β -arrestin-
810 dependent “non-canonical” signalling⁴¹⁻⁴⁴. **d.** Epidermal growth factor receptor (EGFR)
811 can be internalized through different CCPs⁵¹: AP2-dependent CCPs mostly commit the
812 receptor to recycling, sustaining signalling; AP2-independent CCPs, which rely on the
813 EPS15 and/or EPS15L1 and EPN1 adaptors, target the EGFR to lysosomal
814 degradation⁵¹. **e.** Dynamin2 (DYN2) is ubiquitously and constitutively involved in CCP
815 maturation and fission (parts a-d)⁶²⁻⁶⁶. By contrast, dynamin1 (DYN1) is employed in
816 clathrin-mediated endocytosis only under specific conditions; *e.g.*, in EGF stimulated
817 non-small cell lung cancer cells where the inhibitory constraints of GSK3 β -dependent
818 DYN1 phosphorylation are removed^{36,65,68}. In cancer, the GSK3 β kinase can be
819 inactivated (not shown) or targeted for degradation due to aberrant activation of AKT
820 kinase, leading to DYN1 dephosphorylation and aberrant activation, and eventually to
821 deregulated EGFR endocytosis, signalling and migration^{66,73}.

822

823 **Figure 2. NCE mechanisms and cellular contexts.**

824 **a.** CLIC/GEEC (clathrin-independent carrier/GPI-anchored protein-enriched early
825 endosomal compartment) requires extracellular clustering of glycosylated cargoes
826 (through galectins) to induce membrane bending, and small GTPases (ARF1, CDC42)
827 that coordinate the activation of actin nucleators (Arp2/3) and regulators (NWASP)^{2,15}.
828 Bar-domain proteins (GRAF1, IRSP53, PICK1) control the initial phase of vesicle
829 formation, while endophilinA2 (EndoA2) is required for fission, together with the
830 pulling force of the actin cytoskeleton and the microtubule-associated motor protein,
831 dynein: a mechanism known as “friction-mediated scission”^{265,266}. The internalized
832 vesicle fuses with an endosomal GEEC compartment from which cargoes are
833 recycled^{2,15}. **b.** Fast endophilin-mediated endocytosis (FEME), which is active in the
834 same cells as CLIC/GEEC⁸⁵, also requires the action of actin, actin nucleators, and BAR-
835 domain proteins. EndoA2 is required for vesicle formation and fission (together with
836 dynein, microtubules and DYN2)⁸⁵. FEME internalizes epidermal growth factor receptor
837 (EGFR) stimulated with high EGF, and other cargoes, including other receptor tyrosine
838 kinases (RTKs) and G protein-coupled receptors (GPCRs)⁸⁴. **c.** High EGF also induces
839 EGFR-NCE, which requires EGFR ubiquitylation (Ub) and Ub receptors (EPS15, EPS15L1
840 and EPN1)⁵⁶, plasma membrane–ER contacts mediated by the ER transmembrane
841 protein RTN3, and Ca²⁺ release at plasma membrane–ER contacts (of note, some
842 mechanistic aspects of EGFR-NCE are still missing, *e.g.*: Which are the factors involved
843 in plasma membrane–ER tethering? Is RTN3 directly recruited to the EGFR? Which is
844 the exact role of EGFR ubiquitylation and Ub receptors?)⁸⁷. EGFR-dependent signalling
845 occurs at the plasma membrane and in endosomes through ERK1/2 (refs. ^{139,231}).

846 Notably, activated ERK1/2 and EGFR might not accumulate on the same endosome
847 after CME²⁶⁷. They might, instead, do so in cells expressing high EGFR and treated with
848 high EGF concentrations, or expressing elevated RAB5A (ref. ²³¹) (see also Fig. 6),
849 where EGFR-NCE and massive activation of pinocytotic processes occur²⁶⁸. Eventually,
850 EGFR-NCE targets the EGFR to lysosomal degradation, restricting signalling in a cell
851 context-dependent fashion^{86,269}. **d.** The interleukin-2 (IL-2) receptor (IL-2R) complex
852 (comprising α , β , subunits) is expressed in lymphocytes and signals through JAK–
853 STAT²⁷⁰. IL-2R is internalized via an NCE mechanism occurring in proximity of actin-
854 enriched membrane protrusions that facilitate actin polymerization and plasma
855 membrane invagination^{90,271-273}. Once internalized, the α chain is recycled, whereas
856 the β and γ chains are degraded⁹¹. **e.** Macropinocytosis internalizes pathogens,
857 apoptotic cells/debris and proteins through the formation of actin-dependent plasma
858 membrane protrusions¹⁰¹. Internalized materials are delivered to the lysosome. In the
859 lysosome, pathogen-derived antigens are generated and bind to MHC receptors (in
860 immune cells), which are targeted to the plasma membrane for antigen
861 presentation¹⁰². Apoptotic cells/debris and proteins are degraded in lysosomes and
862 used by epithelial cells (particularly, cancer cells) for nutrient scavenging^{103,107-109}.

863

864 **Figure 3. Membrane trafficking in apical-basal cell polarity.**

865 The apical-basal polarity complexes (see Supplementary Box 1) are shown as green
866 ovals, including their main protein components. Epithelial polarity is maintained
867 through coordinated and polarized trafficking of the various components of these
868 complexes, achieved through distinct endosomal intermediates, regulated by different
869 RAB GTPases, as shown²⁷⁴. There is bidirectional connection between polarity
870 complexes and membrane traffic. For instance, RAB11-dependent apical transport
871 pathways reinforce the localization of the apical PAR complex, while apical PAR3,
872 controls the subapical positioning of RAB11-positive endosomes^{171,275}.

873 A number of other apical and basolateral cargoes enter spatially distinct,
874 peripherally localized apical or basolateral early endosomes (AEE or BEE), respectively,
875 and undergo alternative fates²⁷⁶. They can be either recycled to the plasma membrane
876 (not shown), or routed to RAB8—RAB10-double positive common recycling
877 endosomes (CRE)^{277,278}. CREs are so named because they are the target of both apical
878 and basal cargoes, which are then either shuttled to the basolateral surface, or
879 delivered to the apical recycling endosomal (ARE) compartments, before being
880 directed to the apical surface²⁷⁶. For example, basolateral proteins – such as E-
881 Cadherin or transferrin receptor – are targeted from CRE to the basolateral plasma
882 membrane via a pathway that involves the AP1B coat protein as well as the exocyst
883 protein complex (not shown)²⁷⁹. An example of an apical cargo that traffics via AREs to
884 the apical plasma membrane is H⁺/K⁺-ATPase of gastric parietal cells²⁸⁰.
885 Neosynthesized proteins can also be delivered, via Golgi, to their destinations (apical
886 or basolateral) either directly (not shown) or indirectly through AREs or CREs. The
887 green arrow points to the aPKC/LGL circuitry described in Supplementary Box 1.

888

889 **Figure 4. Membrane trafficking in apical lumen formation.**

890 Lumen formation in epithelial monolayers requires coordination between cell division,
891 actin polymerization and membrane trafficking^{160,171,281} (events are shown across two
892 cells for clarity but they occur in both cells). **a.** Following cell division, at the midbody,

893 the apical membrane initiation site (dashed box) is established, through the
894 coordination and re-organization of microtubules (not shown) and branched actin
895 filaments, via RAC1—WAVE to promote recruitment of cingulin. A positive feedback
896 loop (asterisk) involving branched actin polymerization, ZO1 and cingulin further
897 sustains the establishment of the apical membrane initiation site (whereby cingulin
898 promotes branched actin polymerization, which supports recruitment of ZO1 and
899 consequently recruitment of cingulin via its interaction with ZO1)¹⁷². At this stage,
900 early polarity determinants such as podocalyxin or IRSP53, are present on the plasma
901 membrane facing the outer surface. **b.** Multiple endocytic and transport events (1)
902 direct apical determinants (podocalyxin and IRSP53 are shown, but there are several
903 others) to the apical membrane initiation site, where the exocytosis of membrane and
904 apical proteins results in formation of a nascent lumen (2). IRSP53 senses and stabilizes
905 the negative curvature of the membrane via its I-BAR domain, which ensures the
906 integrity and shape of the opposing membranes of the two neighbouring cells for
907 correct lumen generation. RAB11—RAB8-double positive endosomes are critical in this
908 phase (3 and inset)¹⁷¹, as they become restricted towards the luminal side, ideally
909 positioned to dock and fuse at the apical membrane through multiple interactions.
910 These endosomes can be tethered to the apical membrane initiation site by binding to
911 cingulin (via FIP5—RAB11 (4)), by binding to RAB35 directly (5) or through tethering
912 with the exocyst complex (including MYO5B, Sec15a and SEC10)²⁸² (6, inset). Among
913 the cargoes transported by the RAB8—RAB11 vesicles, are annexin2 and CDC42, which
914 are directed apically to interact with apical polarity complex component aPKC (7, inset;
915 see also Supplementary Box 1 for details on epithelial polarity)¹⁷¹. Thus, a self-
916 sustained polarized transport system of apical determinants is essential and precedes
917 the formation of a tight junction-delimited lumen. **c.** Subsequent cell divisions expand
918 the luminal space and increase the size of the luminal cyst through a combination of
919 water influx through solute carriers that increases hydrostatic luminal pressure (blue
920 arrows) and reduced actomyosin contractility (red arrows; the exact molecular
921 mechanisms are unclear) along the forming tight and adherens junctions.

922

923 **Figure 5. Endocytosis and actomyosin in the control of cell shape and tissue**
924 **elongation.**

925 **A.** Diagrammatic representation of cell intercalation in *Drosophila melanogaster*
926 germband extension¹⁹³ (the arrows show the developmental axes: A, anterior; P,
927 posterior; D, dorsal; V, ventral.). T1, T2, T3 indicate the different phases of the process.
928 During the T1-to-T2 transition, the antero-posterior interface between two adjacent
929 cells shrinks to a four-cell vertex (T2, which defines a state where the sides of four
930 adjacent cells converged into a single focal point indicated with a red dot). This is
931 thought to occur because of increased actomyosin medial contraction that generates
932 forces directed along the antero-posterior plane (1). This event is accompanied by
933 elevated and localized clathrin-mediated endocytosis (CME) of junctional molecules,
934 such as E-Cadherin^{193,195} (2). In the T2-to-T3 transition, the four-cell vertex expands to
935 generate a new cell-junctional surface oriented along the dorso-ventral interface that
936 leads to effective tissue elongation along the dorso-ventral axis. **b.** Contraction of
937 amnioserosa in dorsal closure in *D. melanogaster*¹⁹⁶, and in morphogenetic junction
938 remodelling in mammalian epithelial tissues¹⁹⁷. In the initial step, changes in RHOA-
939 mediated activity result in increased medial-apical actomyosin contractility (1) that

940 causes cell areas to oscillate leading to increased junctional tension (2). During area
941 oscillations, RAB35 endocytic activity is enhanced resulting in increased endocytosis
942 (3). This aims at reduction of plasma membrane tension. Concomitantly,
943 internalization of the plasma membrane and various plasma membrane cargo proteins
944 (*e.g.*, E-Cadherin) leads to shortening of the junctional interfaces¹⁹⁶ (4). Myosin II and
945 CME are required to terminate the RAB35 intermediate compartment (not shown) and
946 to direct endocytosed materials to endosomes. This process of internalization of the
947 plasma membrane and junctional molecules functions as an effective ratcheting device
948 to remove excess membrane and to maintain the reduced length of cell–cell junctions.
949 However, through not yet fully understood mechanisms, cargoes – such as CDH1 – can
950 be routed back from endosomes to the interface between the two cells (5), thereby
951 replenishing the membrane and interfacial components and ensuring the reversibility
952 of the interface contraction (6).

953

954 **Figure 6: Endomembranes in the dynamics of collective motion.**

955 **a.** The treadmilling cycle of N-Cadherin in developmental collective motion of
956 astrocytes. Cells at the front show protrusive edges. F-actin attachment to adherens
957 junction components (catenins and N-Cadherin) drives their retrograde flow (1) along
958 lateral contacts (2). At the rear, phosphorylation (P) of p120-catenin by GSK3 untangles
959 the complex (3) allowing for N-Cadherin endocytosis (4) and polarized recycling to the
960 leading edge where GSK3 is inactive (5)²¹⁶. **b.** In epithelial 2D and 3D collectives that
961 depend on epidermal growth factor receptor (EGFR) activation for proliferation and
962 motility, the elevation of RAB5A triggers non-clathrin endocytosis (NCE) of the
963 activated EGFR leading to the concomitant expansion in the size and number of
964 endosomes. This results in sustained ERK1/2 signalling from endosomes. ERK1/2
965 signalling promotes the phosphorylation of WAVE2 (WAVE2-P)²³¹, which, in the
966 presence of activated RAC1, leads to the activation of Arp2/3 complex and branched
967 actin nucleation^{283,284} leading to the formation of “cryptic lamellipodia”²⁸⁵. These
968 structures extend beneath neighbouring cells and drive persistent, coordinated cell
969 motion and efficient cell re-orientation, which, together with elevated junctional
970 tension (not shown), results in unjamming of the entire monolayer^{138,231,286,287}. **c.**
971 Snapshots representative of two distinct states of densely packed epithelial collectives
972 in the presence or absence of RAB5A expression¹³⁸. The velocity vectors (black arrows)
973 of each individual centroid are shown. Jammed immobile, control monolayers display
974 randomly oriented poorly motile cells. RAB5A expression increases cell velocity and
975 promotes the alignment of the velocity vectors. This results in monolayers displaying a
976 flocking mode of motion where long-range collective motion and short-range, local cell
977 arrangements are permitted (not shown). The colour code indicates regions within the
978 monolayer with velocity vectors that are either parallel (yellow) or antiparallel (blue)
979 to the mean direction of migration.

980

981 **Box 1. Clathrin-coated plaques: role in mechanosensing, adhesion and mitosis.**

982 Clathrin-coated plaques (CPLs) are long-lived plasma membrane structures enriched in
983 integrins and various signalling receptors but not actin^{32,33}, that are regulated by the
984 physical properties of the extracellular matrix (ECM). As the substrate rigidity
985 increases, CPLs assemble and expand. By linking CPLs to the substrate, integrin $\alpha\upsilon\beta 5$
986 delays their budding, in a process called “frustrated endocytosis”, ultimately stabilizing

987 them³³. Thus, CPLs might represent mechanosensitive adhesion platforms, generated
988 as a consequence of “frustrated endocytosis”³³ (see figure). CPLs are present in
989 interphase cells (left in figure), together with “canonical” focal adhesion complexes,
990 which are enriched in integrins, focal adhesion kinase (FAK), actin, vinculin, talin and
991 paxillin. At mitosis (right in figure), the actin cytoskeleton is reorganized to form a
992 contractile cortical network of actin filaments, which promotes a cellular
993 morphological change, known as “rounding up”. During cell rounding for mitosis, a
994 decrease in cell surface area is achieved and endocytosis continues, albeit at a lower
995 rate, while recycling is impaired²⁸⁸⁻²⁹⁰. At the same time, “canonical” focal adhesion
996 complexes are disassembled, while “mitotic” adhesion complexes are established to
997 preserve the interaction with the substrate needed to achieve daughter cell re-
998 spreading and mitotic spindle orientation²⁹¹⁻²⁹⁴. These mitotic adhesion complexes and
999 CPLs are likely the same type of structure³⁴ since they share several properties. Both
1000 types of structures are very stable with slow plasma membrane turnover; they are
1001 composed mainly of integrins, while actin is not enriched and does not play any role in
1002 their dynamics; they are enriched in endocytic proteins. In addition, their number and
1003 size increase with the increase of the rigidity of the substrate²⁹³. Another function of
1004 CPLs (not shown in the figure) is to act as adhesive structures in non-mitotic cells
1005 migrating in 3D environments. In this case, they are assembled on linear collagen
1006 fibres and provide multiple anchoring sites to the ECM along cell protrusions in order
1007 to support 3D cell migration, in cooperation with classical focal adhesions²⁹⁵.

1008

1009

1010 **Glossary**

1011

1012 **Receptor tyrosine kinases**

1013 A family of plasma membrane proteins (~ 60 genes in humans) that function as high
1014 affinity binding sites for growth factors and cytokines and transduce signals
1015 intracellularly through their intrinsic tyrosine kinase activity.

1016

1017 **G protein-coupled receptors**

1018 A vast family of plasma membrane receptors (more than 800 genes in humans)
1019 characterized by seven transmembrane regions. They transduce signals through a
1020 variety of modes, among which the best characterized one is the coupling with
1021 heterotrimeric G proteins.

1022

1023 **Arrestins**

1024 A family of proteins that act as multifunctional scaffolding proteins, regulating
1025 trafficking and signaling of transmembrane receptors, particularly of GPCRs. They are
1026 involved in receptor desensitization, endocytosis and ubiquitylation. They can also
1027 function as positive effectors of GPCRs through their scaffolding abilities. The arrestin
1028 family comprises visual arrestins, β -arrestins (non-visual arrestins) and α -arrestins

1029

1030 **Epsin family of endocytic adaptor proteins**

1031 A family of endocytic proteins composed of 3 paralogs: EPN1, EPN2 and EPN3,
1032 characterized by the presence of an epsin N-terminal homology (ENTH) domain
1033 involved in phosphoinositide binding at the plasma membrane, ubiquitin binding

1034 motifs (UIMs), as well as motifs that bind to clathrin, AP2 and other endocytic proteins.
1035 They are involved in both CME, where they play a role in clathrin-coat assembly and
1036 cargo recruitment, and in EGFR-NCE.

1037

1038 **Epithelial-mesenchymal transition (EMT)**

1039 This is a process, of great relevance in embryogenesis, through which epithelial cells
1040 lose polarity and cell–cell adhesion contacts (sessile state) to acquire characteristics of
1041 migratory mesenchymal-like cells. In physiology, typically, after migrating, these cells
1042 re-acquire an epithelial phenotype through the opposite process of mesenchymal–
1043 epithelial transition (MET).

1044

1045 **Death receptors**

1046 Death receptors are type I transmembrane proteins belonging to the tumor necrosis
1047 factor/nerve growth factor superfamily. They are activated upon binding to various
1048 agonists (such as FASLG, TNFA or TRAIL). They typically trigger the so-called apoptotic
1049 extrinsic pathway, yet they can also activate multiple alternative signaling pathways
1050 with opposing outcomes (survival/proliferation vs. cell death) depending on the cell
1051 context.

1052

1053 **Caveolae**

1054 Small flask-shaped invaginations of the plasma membrane (50–80 nm) that can be
1055 morphologically identified by the presence of coat-like proteins, caveolins, and that
1056 are particularly abundant in tissues involved in lipid homeostasis or subjected to
1057 mechanical challenges like adipocytes, muscle and endothelial cells.

1058

1059 **Vinculin**

1060 A protein involved in the formation of focal adhesions that links surface structures
1061 (integrins) to the actin cytoskeleton (through binding to F-actin).

1062

1063 **Focal adhesions**

1064 Cell-to-matrix adhesion structures involved in the transmission and regulation of
1065 signals between the extracellular matrix and the intracellular environment. They are
1066 large and dynamic protein complexes established through integrins (which bind to the
1067 extracellular matrix), vinculin, F-actin and several regulatory components (up to
1068 hundred different proteins, according to the state of the cell). Focal adhesions have
1069 roles in signal transduction, cell motility, cell cycle regulation and several other cellular
1070 phenotypes. They represent one of the main sensors/effectors in cellular
1071 mechanosensing.

1072

1073 **Tip cells**

1074 During angiogenesis, new vessels that sprout from existing ones are guided by a leader
1075 cell that drives the extension of the sprout and senses the environment for guidance
1076 cues.

1077

1078 **ERK signalling**

1079 Is the signalling mediated by the activation of the extracellular signal regulated kinases
1080 (ERKs, also called mitogen-activated protein kinases or MAPKs). This signalling is

1081 mediated by the sequential activation of the small GTPases RAS, and a cascade of
1082 kinases (RAF, MEK, and ERK1,2) that transduce a signal from a receptor, located on the
1083 cell surface or on endosomes, to regulate a number of fundamental biological
1084 function, including cell proliferation, differentiation and migration.

1085

1086 **PAK**

1087 A family of serine/threonine protein kinases that includes six members in mammals.
1088 They serve as targets for the small GTPases CDC42 and RAC and have been implicated
1089 in a wide range of biological activities.

1090

1091 **PTEN**

1092 A lipid phosphatase (phosphatidylinositol 3,4,5 triphosphate 3-phosphatase). It
1093 catalyses the conversion of PI(3,4,5)P3 to PI(4,5)P2, thereby antagonizing the action of
1094 PI3K and the activation of AKT. It represents one of the most frequently lost tumor
1095 suppressors in human cancers

1096

1097 **AMPK**

1098 AMP-activated protein kinase or 5' adenosine monophosphate-activated protein
1099 kinase. It is a heterotrimeric protein complex endowed with serine/threonine kinase
1100 activity which regulates the energy metabolism, mostly acting on glucose and fatty
1101 acid metabolism.

1102

1103 **RAC1**

1104 A member of RHO subfamily of small GTPases that plays a central role in controlling
1105 the activity of protein complexes that are necessary to remodel the actin cytoskeleton
1106 during migration.

1107

1108 **P38**

1109 A member of a class of mitogen-activated protein kinases (MAPKs) that are responsive
1110 to stress stimuli, such as cytokines, ultraviolet irradiation, heat shock, and osmotic
1111 shock, and are involved in cell differentiation, apoptosis and autophagy.

1112

1113 **JNK**

1114 JNK (or c-Jun N-terminal Kinase), is a member of a family of protein kinases, which play
1115 a central role in stress signalling pathways implicated in gene expression, neuronal
1116 plasticity, regeneration, cell death, and regulation of cellular senescence.

1117

1118 **RAB GTPases**

1119 A subfamily of small GTPases that includes more than 70 members in mammals and
1120 regulates several key steps of membrane trafficking, including vesicle formation,
1121 vesicle movement along actin and tubulin networks, and membrane fusion.

1122

1123 **GEF**

1124 This term defines broadly a vast group of proteins (frequently unrelated) which all
1125 possess Guanine Nucleotide Exchange Factor activity, *i.e.*, the ability to convert G
1126 proteins from an inactive GDP-bound to an active GTP-bound form.

1127

1128 **“Jammed” epithelial monolayers**

1129 The dynamics of epithelia has been described in terms of jamming transitions. During
1130 this transition, collective motion ceases, cells can no longer exchange neighbours, and
1131 monolayers become static and rigid, displaying a behaviour similar to that of
1132 ensembles of dense and packed inactive particles, such as coffee in a chute or sand in
1133 a pile.

1134

1135 **WAVE**

1136 A key component of a pentameric actin nucleation promoting complex which acts
1137 downstream of the GTPase RAC, and is necessary for activating the ARP2/3 complex
1138 for the generation of branched actin networks.

1139

1140 **Midbody**

1141 Central region of the thin cytoplasmic bridge that connects cells at the end of
1142 cytokinesis. It consists mostly of microtubules, together with various other types of
1143 proteins (400-500). It functions as a platform to mediate abscission, the process of
1144 severing the intercellular bridge. It is also endowed with numerous other functions,
1145 including determination of cell fate and asymmetric post-abscission signal
1146 transduction.

1147

1148 **Tight junction**

1149 A cell-to-cell junction formed by a multiprotein complex. This type of junction is
1150 established through homotypic interactions between adhesion molecules (occludins,
1151 claudins, JAMs) present on the surface of abutting cells. Tight junctions mark the
1152 border between the apical and the basolateral surfaces in epithelial cells and control
1153 the formation of functionally distinct apical domains. They are also present in
1154 endothelial cells and astrocytes and establish the blood-brain barrier. One of their
1155 major function is to seal the epithelia by preventing leakage of water and small
1156 molecular weight solutes.

1157

1158 **Arp2/3 complex**

1159 A seven-subunit complex that upon activation promotes the branched elongation of
1160 the actin network by binding to the side of mother filaments.

1161

1162 **Crumbs polarity complex**

1163 A multiprotein complex composed of three members originally identified in
1164 *Drosophila*, Crumbs, Pals1 and PatJ. This complex plays a key role in specifying the
1165 apical plasma membrane domain of epithelial cells and in controlling cell shape in both
1166 invertebrates and vertebrates.

1167

1168 **Transcytosis**

1169 A process in which molecules are transported across cellular barriers. It involves the
1170 endocytosis of molecules (typically plasma membrane proteins or extracellular
1171 molecules captured through interaction with surface receptors) at one side of the cell
1172 and their vesicle-mediated transport to another side, where they are released through
1173 exocytosis. It contributes to the establishment of apical-basal cell polarity by
1174 transferring transmembrane proteins between distinct plasma membrane domains. It

1175 is involved in many other processes, for instance the crossing of the blood-brain
1176 barrier.

1177

1178 **Exocyst**

1179 An octameric protein complex involved in vesicle trafficking, specifically in the
1180 tethering and spatial targeting of vesicles to the plasma membrane prior to vesicle
1181 fusion.

1182

1183 **Annexin-2**

1184 A 36-kDa calcium-dependent, phospholipid-binding protein that functions in
1185 promoting the exocytosis of intracellular proteins to the extracellular space.

1186

1187 **CDC42 apical polarity complex**

1188 CDC42 is a highly conserved RHO-family GTPase that regulates cell polarity in many
1189 eukaryotes. It directly interacts with PAR6, and regulates, through this protein, the
1190 activity of the atypical protein kinase C, aPKC.

1191

1192 **BAR domain**

1193 This is a banana-shaped protein domain capable of sensing membrane curvature by
1194 binding preferentially to curved membranes. BAR domains are named after three
1195 proteins in which they were originally identified: BIN1, bridging interactor 1; AMPH,
1196 amphiphysin; and Rvs167, the yeast homolog of amphiphysin.

1197

1198 **Adherens junctions**

1199 Cadherin-based cell-to-cell junctions present in epithelial and endothelial cells,
1200 frequently in a more basal position with respect to tight junctions.

1201

1202 **Isochoric**

1203 A process in which the volume of a closed system does not change. It is synonym of
1204 isovolumetric.

1205

1206 **Germband**

1207 In *Drosophila melanogaster*, the ventral part of the embryo that forms during
1208 gastrulation and gives rise to the segmented trunk of the animal (gnathal, thoracic,
1209 abdominal segments). It includes the mesoderm, ventral ectoderm and dorsal
1210 epidermis but excludes the dorsal-most tissue of the embryo, the amnioserosa.

1211

1212 **Amnioserosa**

1213 In *Drosophila melanogaster*, a short-lived extraembryonic tissue with a critical role in
1214 dorsal closure and other early developmental morphogenetic events.

1215

1216 **Border cells**

1217 A cluster of cells that migrate from the anterior tip of the *Drosophila* egg chamber to
1218 the border of the oocyte at stage 9 of *Drosophila* oogenesis. These cells perform a
1219 stereotypical collective migration on the intervening nurse cells, and reach the oocyte.

1220 They are required for the formation of the micropyle, the eggshell structure through
1221 which sperm enters the egg.

1222

1223 **Lamellipodia**

1224 Thin membrane protrusion present at the leading edge of migrating cells, mostly
1225 constituted by a flat network of actin.

1226

1227 **Treadmilling**

1228 A process characterizing filamentous multimeric protein structures within the cell and
1229 mostly used in reference to filamentous actin (F-actin). When actin subunits (G-actin)
1230 are constantly added at one end of the filament and removed from the opposite one,
1231 the net effect is the treadmilling of the filament which is used, for instance, to
1232 generate motion. The term is also used, more generally, for other biological processes
1233 in which “treadmilling” of molecules or organelles occurs.

1234

1235 **Neural crest**

1236 A temporary group of cells established during vertebrate development, which forms
1237 after gastrulation at the border between the neural plate and the surrounding
1238 ectoderm. After closure of the neural tube (due to the folding of the neural plate into
1239 itself), the neural crest runs along the roof plate of the neural tube. At this stage,
1240 neural crest cells undergo epithelial-mesenchymal transition and migrate to the
1241 periphery, where they give origin to various cell lineages.

1242

1243 **Ascitic fluid**

1244 An abnormal accumulation of fluid in the abdominal cavity frequently caused by liver
1245 disease or cirrhosis, cancers, specifically ovarian and colon cancer, and heart failure.

1246

1247 **AKT kinase**

1248 The three members of the human AKT serine-threonine protein kinase family are often
1249 referred to as protein kinase B alpha, beta, and gamma. These proteins are
1250 phosphorylated by phosphoinositide 3-kinase (PI3K). AKT/PI3K forms a key component
1251 of many signalling pathways that involve the binding of membrane-bound ligands such
1252 as receptor tyrosine kinases.

1253

1254 **Galectins**

1255 A class of proteins that bind specifically to β -galactoside sugars, such as N-acetyl-
1256 lactosamine. Galectins are secreted in the extracellular space, where they encounter
1257 galactose-containing glycoproteins and glycolipids. Binding of galectins to glycosylated
1258 proteins, such as CD44 and $\alpha 5 \beta 1$ integrin, triggers galectin oligomerisation, which
1259 allows their interaction with glycosphingolipids and the generation of plasma
1260 membrane curvature, leading to the formation of clathrin-independent endocytic
1261 carriers (CLICs).

1262

1263

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