# PDinsight user manual

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#### 1 General information

PDinsight is a python program for predicting tissue (cell-cell) level effective permeabilities for symplasmic transport from ultrastructural (EM) data. A typical use would be to compare these values to tissue level assessments of effective permeabilities based on photobleaching / photoactivation of carboxyfluorecein (CF) (Rutschow et al., 2011) or suitable GFP derivates (e.g., DRONPA-s (Gerlitz et al., 2018)). The first version of PDinsight was released along with "From plasmodesma geometry to effective symplasmic permeability through biophysical modelling", Deinum et al. eLife 2019. Mathematical notation in this manual follows (Deinum et al., 2019). If using PDinsight in a publication, please cite this manuscript.

PDinsight is written in python 3. If available, it uses the numpy module, but does not strictly depend upon that. The program has different modes for computing the parameter requirements for a given effective symplasmic wall permeability  $P(\alpha)$  for particles of radius  $\alpha$  and related quantities. The different modes and the relevant parameters are controlled from a parameter file (default: parameters.txt). A graphical user interface (GUI) is provided to help the user create parameter files and run PDinsight. The GUI is written using TKinter, which is included in standard installation of python.

For electron microscopists, who typically have access to many ultrastructural parameters, but often do not know  $P(\alpha)$ , the mode computeVals will be useful. This computes the expected  $P(\alpha)$  values when taking all parameters at face value. Comparison with the sub-nano channel model is possible. In principle, all model parameters must be defined, but missing parameters may be explored using lists of possible values or left at a default (e.g., cell length L and a triangular distribution of PDs), as these have little influence on  $P(\alpha)$ . If estimates of PD density and distribution are missing, the mode computeUnitVals, which computes  $\Pi(\alpha)$  (a "unit permeability", i.e.,

assuming a density of 1 PD/ $\mu$ m<sup>2</sup> and  $f_{ih} = 1$ ), could be useful. In this mode, comparison with the sub-nano channel is impossible.

In tissue level experiments,  $P(\alpha)$  is typically measured, but not all ultrastructural parameters will be known. It is likely that PD density  $\rho$  and radius  $(R_n)$ , assuming straight channels) are poorly known. In this case, mode computeRnDensityGraph will be useful, or a combination of modes computeDens and computeAperture and a number of guesses for  $R_n/\bar{\alpha}$  (maximum  $\alpha$  that would fit through the channel) or  $\rho$ , respectively. Additional poorly known parameters can be explored as suggested above. If uncertain, it is strongly advised to explore PD length l ( $\approx$ cell wall thickness). For thick cell walls  $(l \geq 200 \text{ nm})$ , it may be worth exploring the effects of increased central radius  $(R_c > R_n)$ . This is currently only possible in modes computeVals and computeUnitVals.

PDinsight is published under the GNU general public licence v3.0.

#### 2 Modes

**Major modes** The core of PDinsight is computing effective permeabilities  $(P(\alpha))$  for symplasmic transport based on all model parameters mentioned in the manuscript. This is in mode computeVals. The same computations are used in other modes, which compute the requirements for obtaining a given target value (or set of values) of  $P(\alpha)$ . In mode computeDens, required densities  $(\rho)$  are computed for given values of maximum particle size  $\bar{\alpha}$  (and other parameters). In mode computeAperture, required apertures, given as  $\bar{\alpha}$  as well as neck radius  $R_n$ , are computed for given values of  $\rho$  (and other parameters). In mode computeRnDensityGraph,  $R_n$ ,  $\rho$  curves are computed that together yield a target  $P(\alpha)$ . The corresponding values of  $\bar{\alpha}$  are also reported. These curves can be visualized using any plotting program. Mode sensitivityAnalysis computes so-called elasticities (normalized partial derivatives) around a given set (or sets) of parameters. These elasticities tell how sensitive calculated values of  $P(\alpha)$ ,  $\Pi(\alpha)$  and constituents like  $f_{ih}$ (a correction factor for the fact that the cell wall is only symplasmically permeable where the PDs are) are on the parameters involved.

Auxillary modes In computing  $P(\alpha)$ , correction factor  $f_{ih}$  is automatically included. For specific cases such as modelling studies, however, it may be useful to calculate  $f_{ih}$  separately. For this purpose, several modes exist for exploring inhomogeneity factor  $f_{ih}$ : computeFih\_subNano (function of  $\bar{\alpha}$ ; also output values for sub-nano model), computeFih\_pitField\_dens (function of  $\rho$ ), computeFih\_pitField\_xMax (function of  $\bar{\alpha}$ ) and computeTwinning

(function of  $\rho_{pits}$ , cluster density).

By default, computations are performed for the unobstructed sleeve model (Deinum et al., 2019). Most computations can also be performed for the subnano channel model Liesche and Schulz (2013); Comtet et al. (2017). Using switch compSubNano, values for the sub-nano model are also computed.

# 3 Graphical user interface

The GUI to PDinsight is written to facilitate the creation of parameter files and also has a button to run PDinsight directly based on the parameters displayed. In contrast to the generic parameter file, the GUI only shows fields for parameters that are actually used in a specific run mode. The first step of using the gui is to select the run mode (choose from: computeVals, computeUnitVals, computeDens, computeAperture, computeRnDensityGraph and sensitivityAnalysis), followed by "load default parameters". This overwrites any user input from previous modes. All required parameters are shown as text entry fields, with radio buttons for the relevant options. Basic validation occurs on the fly (valid input type, etc). When done, click "Run PDinsight" to write a parameter file and run the program or "Export parameter file" to write a parameter file only. Note that switching modes requires clicking "load default parameters", i.e., a fresh start.

Additional information can be obtained by clicking the "Info" button and, for certain parameters, clicking on the parameter name. If additional information is available, the mouse cursor changes into a question mark.

### 4 Command line usage

**Command line usage** (linux): python PDinsight\_vXX.py PARAMETER-FILE.

(windows): first open a command prompt window (e.g., by searching for "cmd" in the search bar) and go to the directory where PDinsight is located. Then type: py PDinsight\_vXX.py PARAMETERFILE

Using the GUI from the command line (linux): python PDinsightGUI\_vYY.py or (windows): py PDinsightGUI\_vYY.py The correct version of PDinsight\_vXX.py should also be available in the current directory.

In the above, XX and YY should be replaced by the respective version numbers.

## 5 Output files

By default, all outputs are tab separated files (.tsv). This can be switched to csv using the option "outputType" (options "tsv" and "csv").

### 6 Program maintenance

The latest version of PDinsight + documentation can be downloaded from Github: .... It is recommended to update both the core (PDinsight\_vXX.py) and the GUI (PDinsightGUI\_vYY.py) simultaneously. If, for whatever reason you would like to update only the core, please note that you have to adapt the following line in the GUI by hand:

import PDinsight\_v1XX as pd

### 7 Model parameters

See Figure 1 for an overview of single channel PD parameters and PD geometry. See Table 1 for a list of model parameters (PDinsight names) and the corresponding mathematical symbols used in Deinum et al. (2019). See Table 2 for available modes and Table 3 for available options.

#### References

- Comtet J, Turgeon R, Stroock A. Phloem loading through plasmodesmata: a biophysical analysis. Plant physiology. 2017; p. pp-01041.
- **Deinum EE**, Benitez-Alfonso Y, Mulder BM. From plasmodesma geometry to effective symplastic permeability through biophysical modelling. BioRxiv. 2019; p. 668715.
- **Gerlitz N**, Gerum R, Sauer N, Stadler R. Photoinducible DRONPA-s: a new tool for investigating cell–cell connectivity. The Plant Journal. 2018; 94(5):751–766.
- **Liesche J**, Schulz A. Modeling the parameters for plasmodesmal sugar filtering in active symplasmic phloem loaders. Front Plant Sci. 2013; 4:207. http://dx.doi.org/10.3389/fpls.2013.00207, doi: 10.3389/fpls.2013.00207.

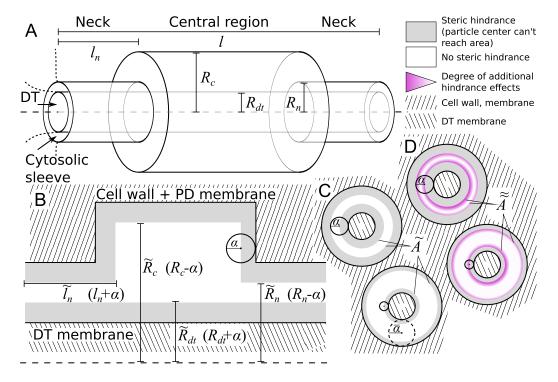


Figure 1: Overview of single channel PD parameters. Figure from Deinum et al. (2019). A: structural parameters as used for input (PD dimensions). B-D: Cross sections showing how internally, available volumes are adjusted for steric hindrance (B,C) and additional (hydrodynamic) hindrance effects (D) depending on particle size  $\alpha$ . A particle with radius  $\bar{\alpha}$  would touch both channel walls in the narrowest (neck) region.

Table 1: List of parameters and mathematical symbols (Deinum et al., 2019). [List] indicates that (depending on mode), a list of values may be entered.

PDinsight	0 -	User unit	Description
X	$\alpha$	nm	Particle radius
xMax	$\bar{\alpha}$	nm	Maximum particle radius that fits
			through a (model) PD
diff	D	$\mu\mathrm{m}^2/\mathrm{s}$	Particle size dependent diffusion con-
			stant
Fih	$f_{ih}$	-	Correction factor for inhomogeneous
			wall permeability $(0 \le f_{ih} \le 1)$
Lneck[List], Lneck2[List]	$l_n$	nm	Neck length
Lpd	l	nm	Total PD length
Lcell	L	$\mu\mathrm{m}$	Cell length
Rc[List]	$R_c$	nm	Central region radius
Rdt[List]	$R_{dt}$	nm	DT radius
Rn[List], Rn2[List]	$R_n$	nm	Neck radius
dens[List]	ho	$PDs / \mu m^2$	PD density
pit[List]	p	-	number of PDs per cluster ("pit
			field"). Options: 1, 2, 3, 4, 5, 6, 7,
			12, 19.
dPit[List]	p	nm	nearest neighbour distance between
			PDs inside pit field.
Peff	$P(\alpha)$	$\mu\mathrm{m/s}$	symplasmic permeability (for particles
			of size $\alpha$ ) of the entire cell wall
Punit	$\Pi(\alpha)$	$\mu\mathrm{m/s}$	symplasmic permeability (for particles
			of size $\alpha$ ) of a single PD per unit of cell
			wall surface, without correction factor
			$f_{ih} (\Pi(\alpha) = \frac{P(\alpha)}{f_{ih}\rho}).$
$\operatorname{grid}[\operatorname{List}]$			PDs (or pit fields) are assumed to
			spaced on a grid. Options: "triangu-
			lar", "square", "hexagonal" or "hex"
			and "random".
file Tag			Prefix for all output files.

Table 2: List of modes. Modes indicated with \* are only available through the command line interface.

Name	Description
computeRnDensityGraph	Compute a graph of $R_n$ , $\rho$ pairs that matches a given
	$P(\alpha)$ given structural parameters etc.
computeDens	Compute required density $\rho$ given a (list of) $\bar{\alpha}$ values.
compute Aperture	Compute required PD aperture (and related $\bar{\alpha}$ ) given
	a (list of) $\rho$ values.
compute Vals	Compute $P(\alpha)$ given structural parameters etc.
computeUnitVals	Compute $\Pi(\alpha)$ given single PD structural parameters
	only.
sensitivityAnalysis	Compute elasticities (normalized partial derivatives)
	around a given set (or sets) of parameters.
$computeFih\_subNano$	Compute $f_{ih}$ , including comparison with sub-nano
	channel model, for $\bar{\alpha} \in [2\alpha, 50]$ nm.
computeFih_pitField_xMax	Compute $f_{ih}$ , possibly for multiple values of $p$ , for
	$\bar{\alpha} \in [2\alpha, 50] \text{ nm}.$
$computeFih_pitField\_dens$	Compute $f_{ih}$ , possibly for multiple values of $p$ , for
	$ \rho \in [0.1, 30] \text{ PDs}/\mu\text{m}^2. $
compute Twinning	Compute $f_{ih}$ , possibly for multiple values of $p$ , where
	the total density is $\rho p$ and $\rho \in [0.1, 30] \text{ PDs}/\mu\text{m}^2$ .

	Table 3: List of options	
Name	Description	
compSubNano	Compare with sub-nano channel model.	
compute Cluster Increase	If true, dens indicates cluster density rather than total	
	density. If false, dens indicates total density.	
$\operatorname{printRn}$	(Additionally) include $R_n$ in output. For some modes,	
	$R_n$ is always printed.	
doNotCombine	If true, "list" parameters must contain either 1 or $n$	
	entries, where $n$ is the same for all. PDinsight uses the	
	$i^{th}$ entry (or single value) of each list as a set. Useful	
	for evaluating raw data. If false, PDinsight evaluates	
	all possible combinations of entries in all lists.	
$\operatorname{asymmetricPDs}$	Allow for different neck dimensions on either side of	
	the PD. If true, Rn2[List] and Lneck2[list] are re-	
	quired.	

Rutschow HL, Baskin TI, Kramer EM. Regulation of solute flux through plasmodesmata in the root meristem. Plant Physiol. 2011 Apr; 155(4):1817–1826. http://dx.doi.org/10.1104/pp.110.168187, doi: 10.1104/pp.110.168187.