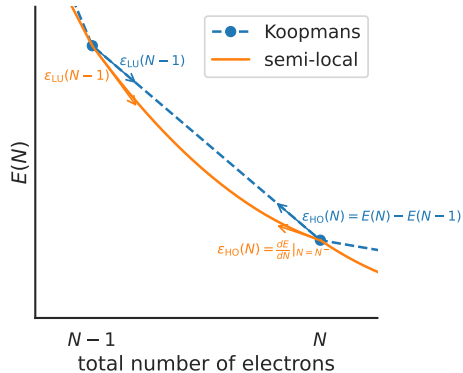


Accurate charged and neutral excitations with Koopmans functionals

Goal: spectral properties (charged excitations) with a functional theory



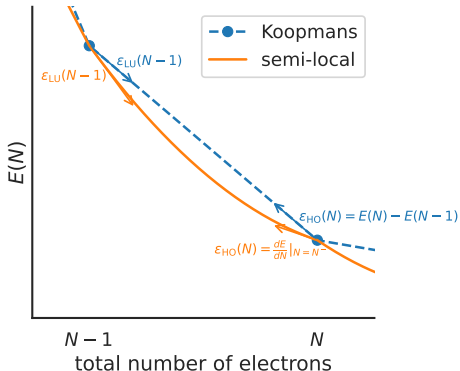
Goal: spectral properties (charged excitations) with a functional theory

Core idea: for every orbital i their energy

$$\varepsilon_i^{\text{Koopmans}} = \langle \varphi_i | H | \varphi_i \rangle = \partial E_{\text{Koopmans}} / \partial f_i$$

should be...

- independent of its own occupation f_i
- equal to the corresponding total energy difference $E_i(N-1) - E(N)$



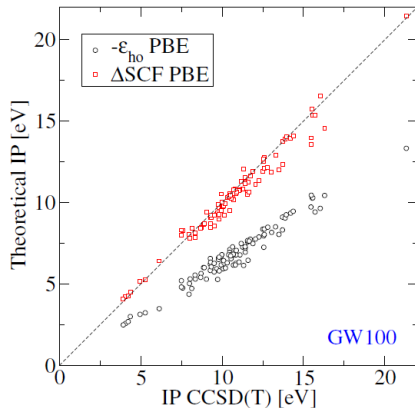
Goal: spectral properties (charged excitations) with a functional theory

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$$E_{\text{Koopmans}}[\rho, \{f_i\}, \{\alpha_i\}] = E_{\text{DFT}}[\rho] + \sum_i \alpha_i \left(\underbrace{- \int_0^{f_i} \varepsilon_i(f) df}_{\text{removes curvature}} + \underbrace{f_i \int_0^1 \varepsilon_i(f) df}_{\text{restores linearity}} \right)$$

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Differences to semi-local functionals:

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Differences to semi-local functionals:

- orbital-density dependence (can use MLWF densities)

$$v_i^{\text{KI}}/\alpha_i = -E_{\text{H}}[n_i] + E_{\text{xc}}[\rho] - E_{\text{xc}}[\rho - n_i] - \int d\mathbf{r}' v_{\text{xc}}(\mathbf{r}', [\rho]) n_i(\mathbf{r}')$$

$$E_{\text{Koopmans}}[\rho, \{f_i\}, \{\alpha_i\}] = E_{\text{DFT}}[\rho] + \sum_i \alpha_i \left(\underbrace{- \int_0^{f_i} \varepsilon_i(f) df}_{\text{removes curvature}} + \underbrace{f_i \int_0^1 \varepsilon_i(f) df}_{\text{restores linearity}} \right)$$

Differences to semi-local functionals:

- orbital-density dependence (can use MLWF densities)
- screening

$$\frac{dE}{df_i} \approx \alpha_i \frac{\partial E}{\partial f_i}$$

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Differences to semi-local functionals:

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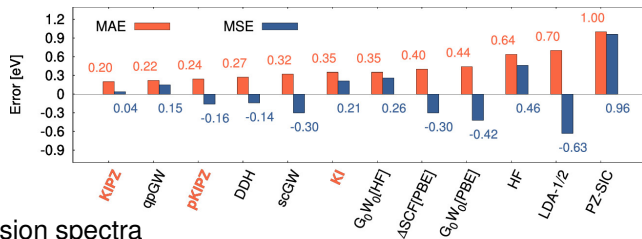
$$\frac{dE}{df_i} \approx \alpha_i \frac{\partial E}{\partial f_i} \Rightarrow \varepsilon_i^{\text{Koopmans}} = \frac{\partial E_{\text{Koopmans}}}{\partial f_i} \approx E_i(N-1) - E(N)$$

Resonance with other efforts:

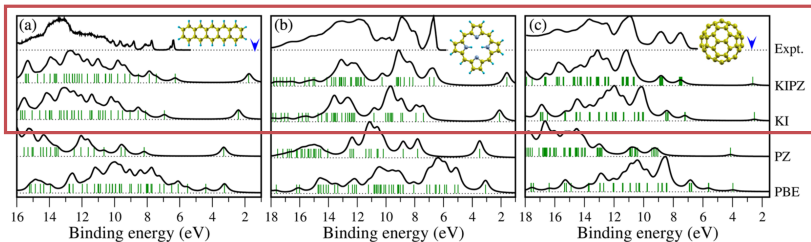
- Wannier transition-state method of Anisimov and Kozhevnikov V. I. Anisimov et al. *Phys. Rev. B* 72.7 (18, 2005), 075125
- Ensemble DFT of Kronik and co-workers E. Kraisler et al. *Phys. Rev. Lett.* 110.12 (19, 2013), 126403
- Koopmans-Wannier of Wang and co-workers J. Ma et al. *Sci. Rep.* 6.1 (1 26, 2016), 24924
- Dielectric-dependent hybrid functionals of Galli and co-workers J. H. Skone et al. *Phys. Rev. B* 93.23 (3, 2016), 235106. arXiv: 1605.02812
- LOSC functionals of Yang and co-workers C. Li et al. *Natl. Sci. Rev.* 5 (2018), 203
- Optimally tuned hybrid functionals of Kronik, Pasquarello, and others D. Wing et al. *Proc. Natl. Acad. Sci.* 118.34 (24, 2021), e2104556118

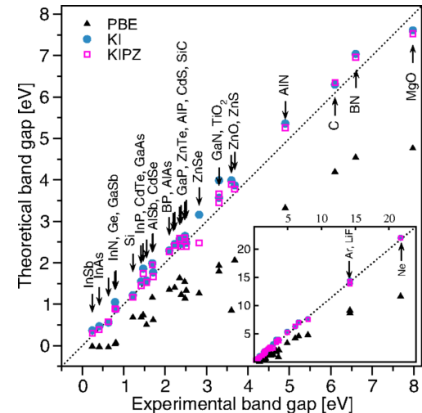
Koopmans functionals: results for molecules

Ionisation potentials = $E(N-1) - E(N) \stackrel{?}{=} -\varepsilon_{HO}$ of 100 molecules (the GW100 set) cf. CCSD(T)



Ultraviolet photoemission spectra

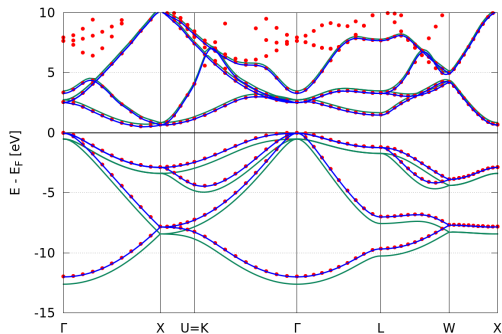




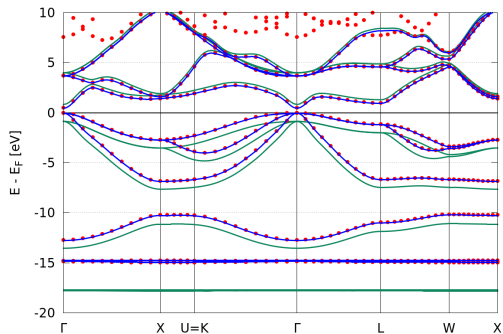
Mean absolute error (eV) across prototypical semiconductors and insulators

	PBE	G ₀ W ₀	KI	KIPZ	QSGW
E _{gap}	2.54	0.56	0.27	0.22	0.18
IP	1.09	0.39	0.19	0.21	0.49

Koopmans functionals: results for solids

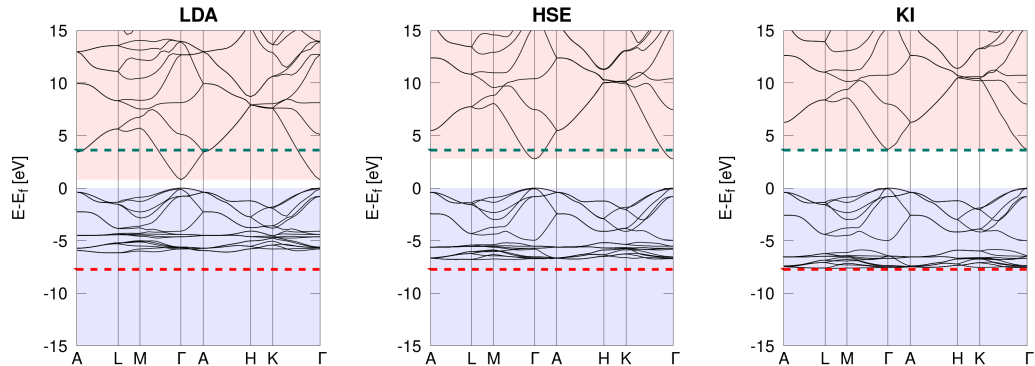


(a) Si, KIPZ



(b) GaAs, KI

		PBE	QSGW	KI	pKIPZ	KIPZ	exp
Si	E_{gap}	0.55	1.24	1.18	1.17	1.19	1.17
	$\langle \varepsilon_d \rangle$	14.9	17.6	16.9	17.7	17.7	18.9
GaAs	E_{gap}	0.50	1.61	1.53	1.49	1.50	1.52
	$\langle \varepsilon_d \rangle$	14.9	17.6	16.9	17.7	17.7	18.9

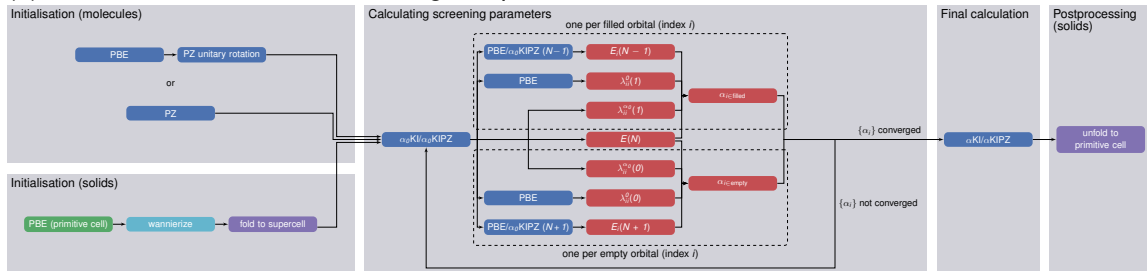


ZnO	LDA	HSE	GW_0	scG \tilde{W}	KI	exp
E_{gap} (eV)	0.79	2.79	3.0	3.2	3.62	3.60
$\langle \epsilon_d \rangle$ (eV)	-5.1	-6.1	-6.4	-6.7	-6.9	-7.5/-8.0

Screening coefficients $\{\alpha_i\}$ must be determined first, via. . .

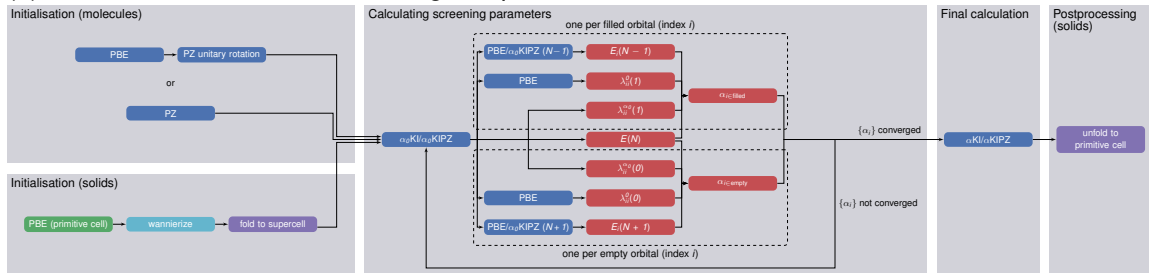
Screening coefficients $\{\alpha_i\}$ must be determined first, via. . .

(a) finite difference calculations using a supercell



Screening coefficients $\{\alpha_i\}$ must be determined first, via. . .

(a) finite difference calculations using a supercell

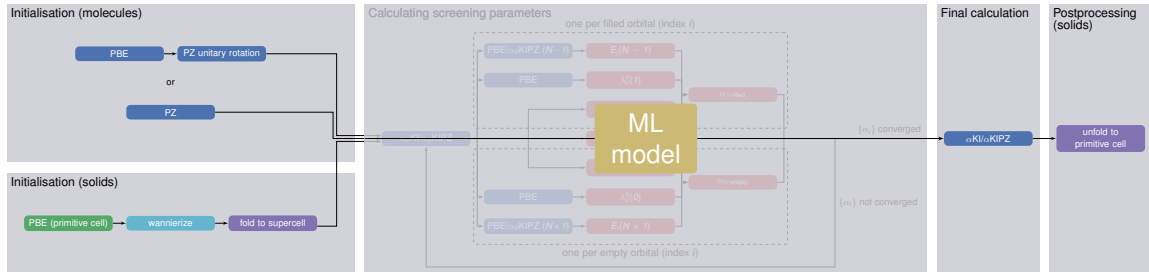


(b) DFPT using a primitive cell



Screening coefficients $\{\alpha_i\}$ must be determined first, via. . .

(a) finite difference calculations using a supercell



(b) DFPT using a primitive cell



(c) via machine learning (resulting band gaps within ~ 0.02 eV of explicit approach)

$k_{CW} \cdot x$ (DFPT implementation) is distributed in Quantum ESPRESSO v7.1 onwards

But complex workflows mean that...

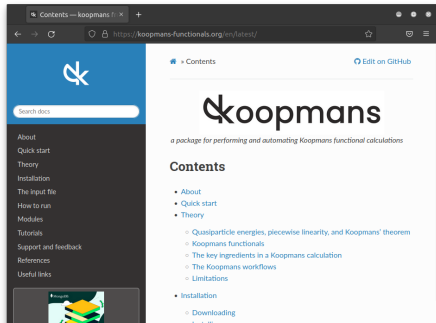
- lots of different codes that need to handshake
- lots of scope for human error
- reproducibility becomes difficult
- expert knowledge required

Our solution...

koopmans

- beta version just released¹
- implementations of Koopmans functionals
- automated workflows
 - start-to-finish Koopmans calculations
 - Wannierisation
 - dielectric tensor
 - convergence tests
 - ...
- built on top of ASE²
- does not require expert knowledge

koopmans-functionals.org



¹ Linscott et al., in prep

² A. H. Larsen et al. *J. Phys. Condens. Matter* 29.27 (12, 2017), 273002

```
{
  "workflow": {
    "task": "singlepoint",
    "functional": "ki",
    "method": "dscf",
    "init_orbitals": "mlwfs",
    "alpha_guess": 0.1
  },
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    "atomic_positions": {
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      "positions": [{"Si", 0.00, 0.00, 0.00},
                    ["Si", 0.25, 0.25, 0.25]]
    },
    "cell_parameters": {
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      "ibrav": 2,
      "celldm(1)": 10.262
    }
  },
}
```

```
  "k_points": {
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    "path": "LGXKG"
  },
  "calculator_parameters": {
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    "w90": {
      "projections": [{"fsite": [0.125, 0.125, 0.125],
                       "ang_mtm": "sp3"}]
    },
    "emp": {
      "dis_froz_max": 11.5,
      "dis_win_max": 17.0
    }
  }
}
```



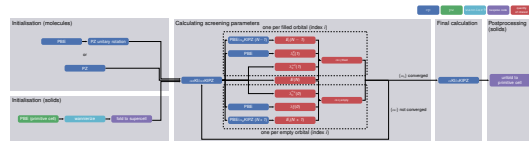
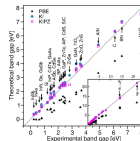
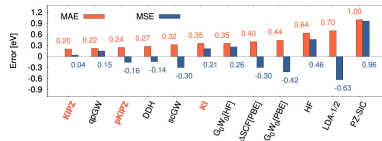
```
from ase.build import bulk
from koopmans.kpoints import Kpoints
from koopmans.projections import ProjectionBlocks
from koopmans.workflows import SinglepointWorkflow

# Use ASE to create bulk silicon
atoms = bulk('Si')

# Define the projections for the Wannierization (same for filled and empty manifold)
si_proj = [{'fsite': [0.25, 0.25, 0.25], 'ang_mtm': 'sp3'}]
si_projs = ProjectionBlocks([{'filled': True, **si_proj},
                              {'filled': False, **si_proj}],
                             atoms=atoms)

# Create the workflow
workflow = SinglepointWorkflow(atoms = atoms,
                               projections = si_projs,
                               ecutwfc = 40.0,
                               kpoints = Kpoints(grid=[8, 8, 8], path='LGXKG', cell=atoms.cell),
                               calculator_parameters = {'pw': {'nbnd': 10},
                                                       'w90_emp': {'dis_froz_max': 10.6, 'dis_win_max': 16.9}})

# Run the workflow
workflow.run()
```



- Koopmans functionals are a class of functionals that treat spectral properties on the same footing as total energy differences (via GPWL)
- they can give orbital energies and band structures with comparable accuracy to state-of-the-art GW
- the release of `koopmans` means you don't need expert knowledge to run Koopmans functional calculations

Want to find out more? Go to koopmans-functionals.org

Free online school Nov 9-11 2022 *Advanced Quantum ESPRESSO tutorial: Hubbard and Koopmans functionals from linear response*. Register at <https://sites.google.com/view/hubbard-koopmans/home>

Follow [@ed_linscott](https://twitter.com/ed_linscott) for updates | Slides available at [elinscott.github.io](https://github.com/elinscott)

SPARE SLIDES

Recap from earlier

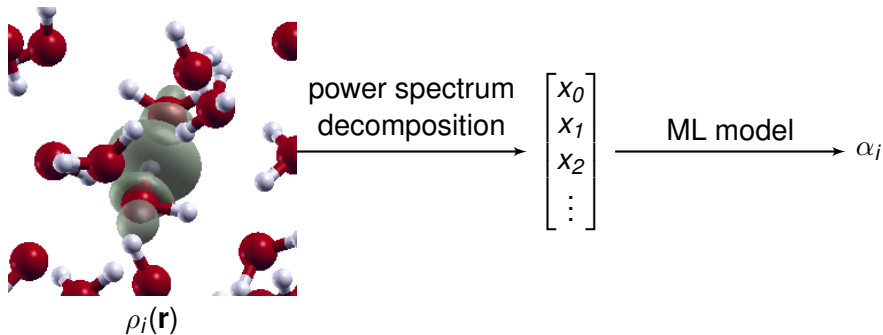
Key idea: construct a functional such that the *variational* orbital energies

$$\varepsilon_i^{\text{Koopmans}} = \langle \varphi_i | H | \varphi_i \rangle = \partial E_{\text{Koopmans}} / \partial f_i$$

are...

- independent of the corresponding occupancies f_i
- equal to the corresponding total energy difference $E_i(N-1) - E(N)$

zero band gap \rightarrow occupancy matrix for variational orbitals is off-diagonal

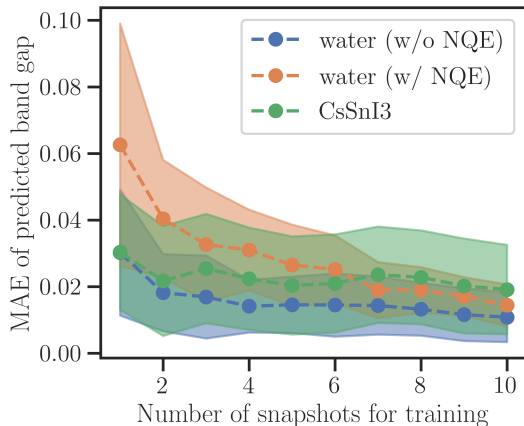
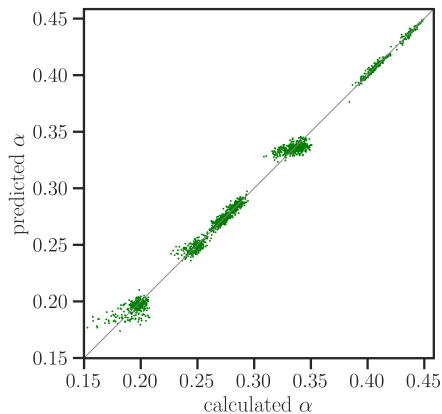


$$c_{nlm,k=\text{orbital}}^i = \int d\mathbf{r} g_{nl}(r) Y_{lm}(\theta, \varphi) \rho^i(\mathbf{r} - \mathbf{R}^i)$$

g_{nl} = orthonormalised radial Gaussian basis functions

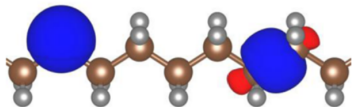
Y_{lm} = spherical harmonics

$$p_{n_1 n_2 l, k_1 k_2}^i = \pi \sqrt{\frac{8}{2l+1}} \sum_m c_{n_1 l m, k_1}^{i*} c_{n_2 l m, k_2}^i$$

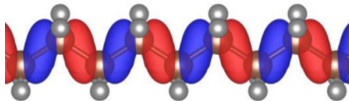


loss of accuracy of the band gap of ~ 0.02 eV
(cf. when calculating screening parameters *ab initio*)

- a natural generalisation in the direction of spectral functional theory¹
- variational (localised, minimising) vs canonical (delocalised, diagonalising) orbitals



(a) variational



(b) canonical

- ODD functional means that we know $\hat{H}|\varphi_i\rangle$ for variational orbitals $\{|\varphi_i\rangle\}$ but we don't know \hat{H} in general
- Difficulties when it comes to calculating transport properties/spectra
- Perhaps a DFT+ U -projector approach is more convenient?

¹ A. Ferretti et al. *Phys. Rev. B* 89.19 (27, 2014), 195134.

N. L. Nguyen et al. *Phys. Rev. X* 8.2 (23, 2018), 021051

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