MAX DRIVING THE EXASCALE TRANSITION

JÜLICH Forschungszentrum

Many-Body Perturbation Theory
The *GW* approximation

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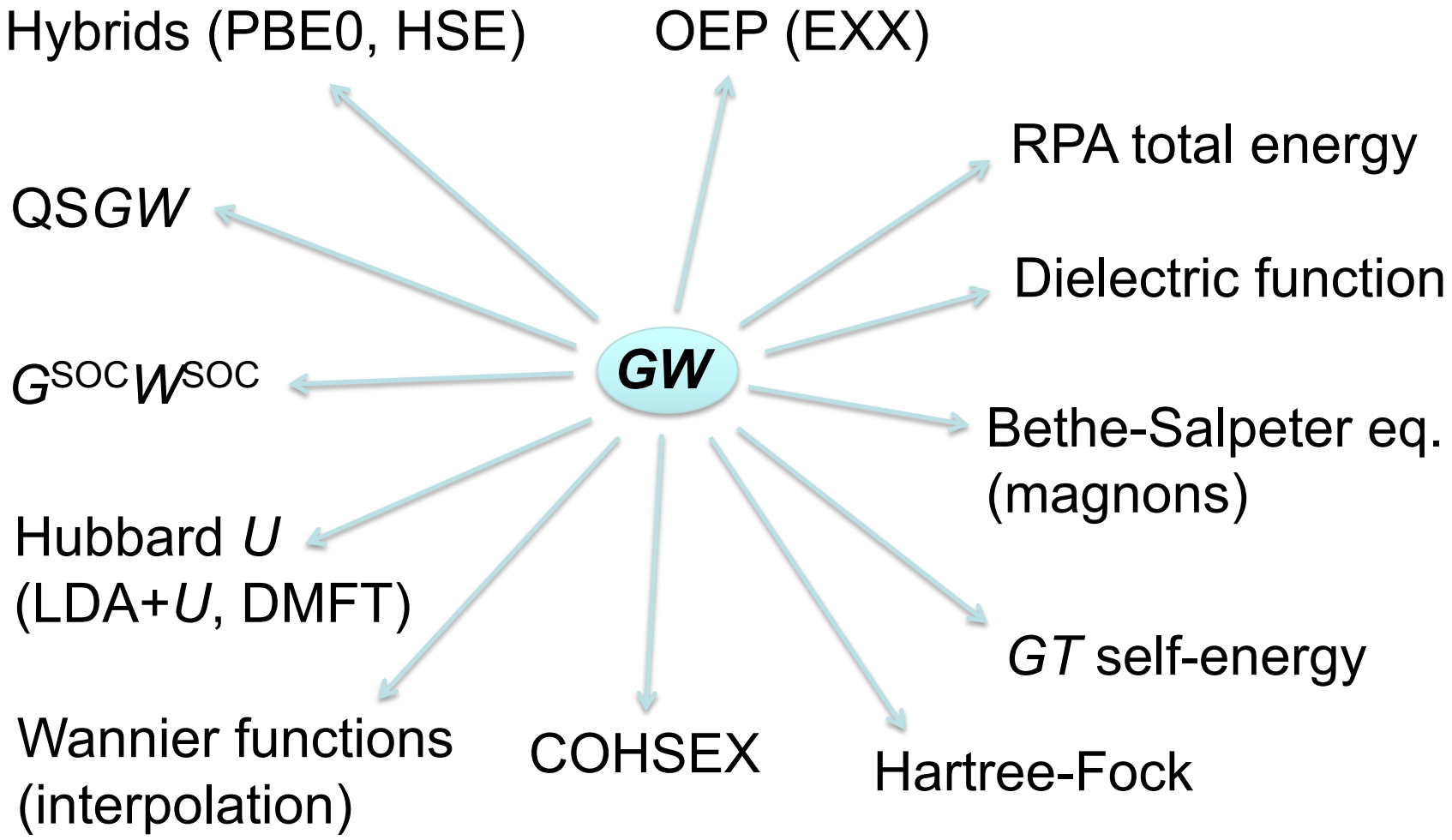
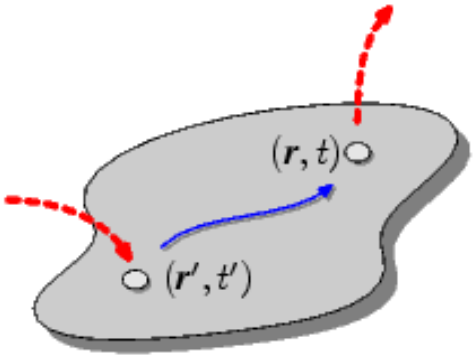
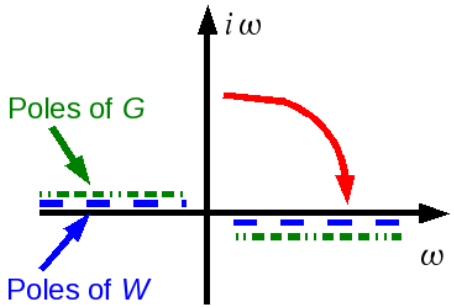
SPEX

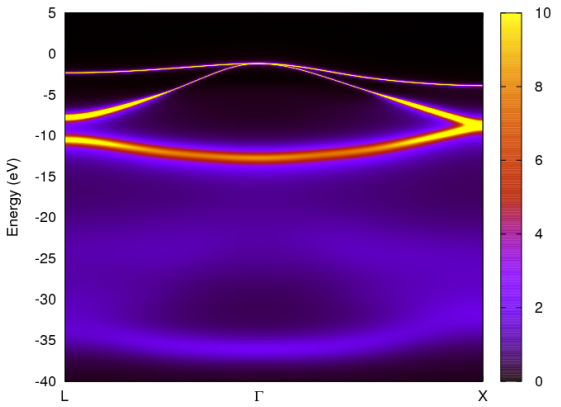

Hybrids (PBE0, HSE)
OEP (EXX)
RPA total energy
QS*GW*
Dielectric function
$G^{SOC}W^{SOC}$
GW
Bethe-Salpeter eq.
(magnons)
Hubbard *U*
(LDA+*U*, DMFT)
GT self-energy
Wannier functions
(interpolation)
COHSEX
Hartree-Fock

Overview

- **Introduction**
- **Theory**
  - Green function
  - Feynman diagrams
  - *GW* approximation
- **Implementation**
  - Basis sets (FLAPW method)
  - Exchange: Hartree-Fock
  - Correlation: Imaginary-frequency formulation
  - Analytic continuation vs. contour integration
- **Applications**
  - Silicon
  - Zinc Oxide
  - Metallic Na
  - HgTe
- **Computational procedure**
  - One-Shot *GW*
  - Full band structure
  - Self-consistent QS*GW*
- **Summary**







Density functional theory

Kohn-Sham (KS) equations:

$$\left[-\frac{1}{2}\nabla_{\mathbf{r}}^2 + v_{\mathrm{ext}}(\mathbf{r}) + v_{\mathrm{H}}(\mathbf{r})\right]\phi_{n\mathbf{k}}(\mathbf{r}) + \boxed{v_{\mathrm{xc}}[\rho](\mathbf{r})\phi_{n\mathbf{k}}(\mathbf{r})} = \epsilon_{n\mathbf{k}}\phi_{n\mathbf{k}}(\mathbf{r})$$

Exchange and correlation potential

Approximations: **LDA, GGA**

➔ real ρ(**r**) ➔ $E_0[\rho]$

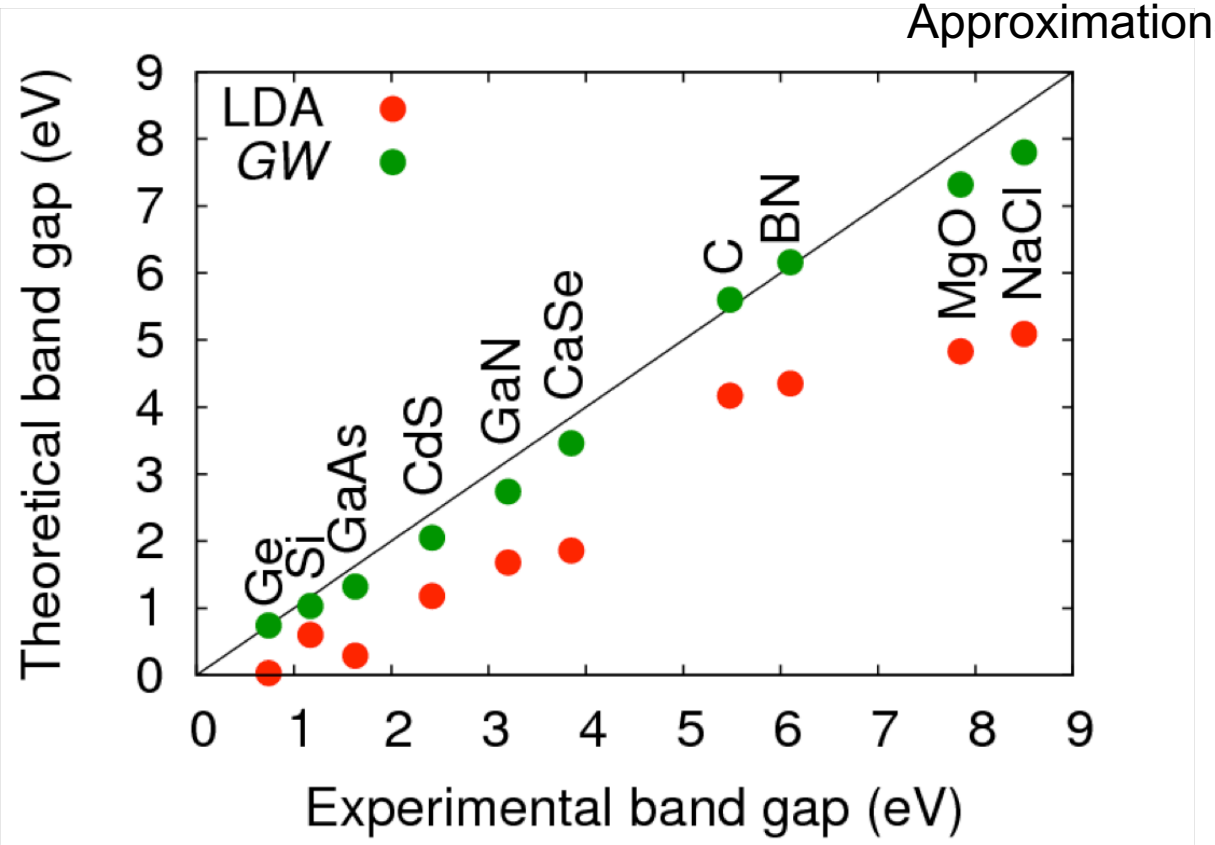
Density functional theory

Kohn-Sham (KS) equations:

$$\left[-\frac{1}{2}\nabla_{\mathbf{r}}^2 + v_{\mathrm{ext}}(\mathbf{r}) + v_{\mathrm{H}}(\mathbf{r})\right]\phi_{n\mathbf{k}}(\mathbf{r}) + \boxed{v_{\mathrm{xc}}[\rho](\mathbf{r})\phi_{n\mathbf{k}}(\mathbf{r})} = \epsilon_{n\mathbf{k}}\phi_{n\mathbf{k}}(\mathbf{r})$$

Exchange and correlation potential

Approximations: **LDA, GGA**



Density functional theory

Kohn-Sham (KS) equations:

$$\left[-\frac{1}{2}\nabla^2_{\mathbf{r}} + v_{\mathrm{ext}}(\mathbf{r}) + v_{\mathrm{H}}(\mathbf{r})\right]\phi_{n\mathbf{k}}(\mathbf{r}) + \boxed{v_{\mathrm{xc}}[\rho](\mathbf{r})\phi_{n\mathbf{k}}(\mathbf{r})} = \epsilon_{n\mathbf{k}}\phi_{n\mathbf{k}}(\mathbf{r})$$

Exchange and correlation potential

Approximations: **LDA, GGA**

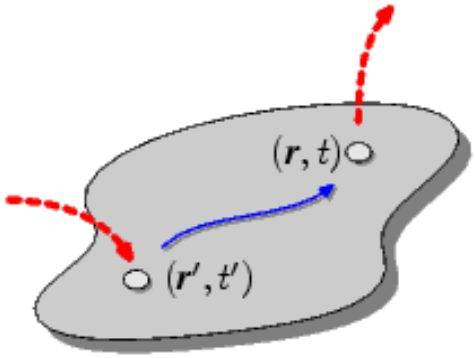
Many-body Schrödinger equation:

$$\left[-\frac{1}{2}\sum_i \nabla^2_{\mathbf{r}_i} + \sum_i v_{\mathrm{ext}}(\mathbf{r}_i) + \frac{1}{2}\sum_{i,j}\frac{1}{\mathbf{r}_i - \mathbf{r}_j}\right]\Psi_n(\mathbf{r}_1, \mathbf{r}_2, ...) = E_n\Psi_n(\mathbf{r}_1, \mathbf{r}_2, ...)$$

Theory
Green function

Central quantity is the single-particle Green function
(probability amplitude for the propagation of a particle)

$$G(\mathbf{r},\mathbf{r}';t-t') = -i\langle\Psi_0^N|\hat{T}[\hat{\psi}(\mathbf{r},t)\hat{\psi}^\dagger(\mathbf{r}',t')]|\Psi_0^N\rangle$$



which contains poles at the excitation energies of the many-electron system (photoelectron spectroscopy),

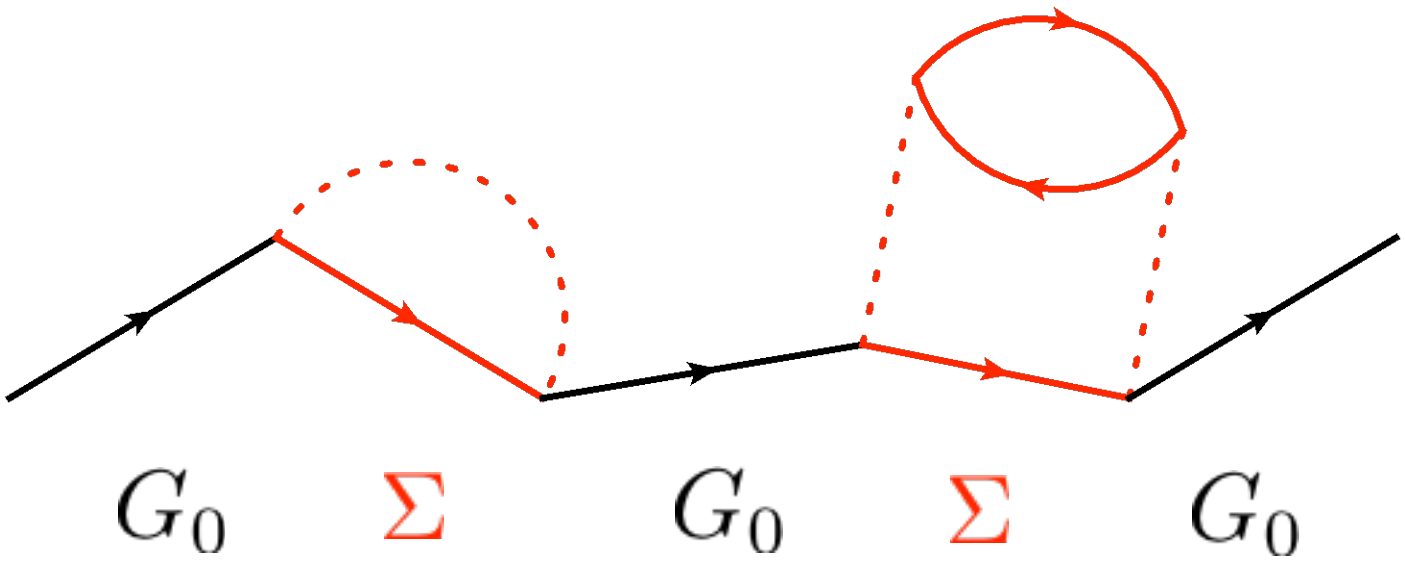
seen by Fourier transformation $t - t' \longrightarrow \omega$

$$G(\mathbf{r},\mathbf{r}';\omega) = \sum_n \frac{\psi_n^{N+1}(\mathbf{r})\psi_n^{N+1*}(\mathbf{r}')}{\omega - (E_n^{N+1} - E_0^N) + i\eta} + \sum_n \frac{\psi_n^{N-1}(\mathbf{r})\psi_n^{N-1*}(\mathbf{r}')}{\omega - (E_0^N - E_n^{N-1}) - i\eta}$$

inverse direct

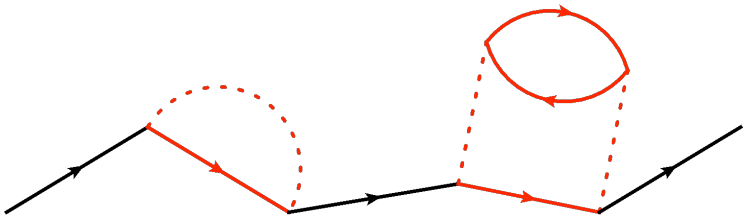
Excitation energy measured in photoemission spectroscopy

Theory
Feynman diagrams



Self-energy: sum over all scattering diagrams

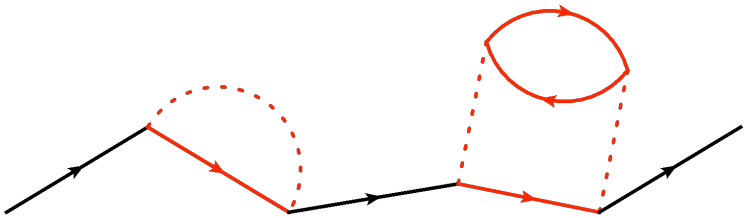
Theory
Dyson equation



$$G = G_0 + G_0\Sigma G_0 + G_0\Sigma G_0\Sigma G_0 + G_0\Sigma G_0\Sigma G_0\Sigma G_0 + \ldots$$

Σ is the electronic self-energy (scattering potential).

Theory
Dyson equation



$$G = G_0 + G_0 \Sigma G$$

Σ is the electronic self-energy (scattering potential).

The Dyson equation can be rewritten as the quasiparticle equation

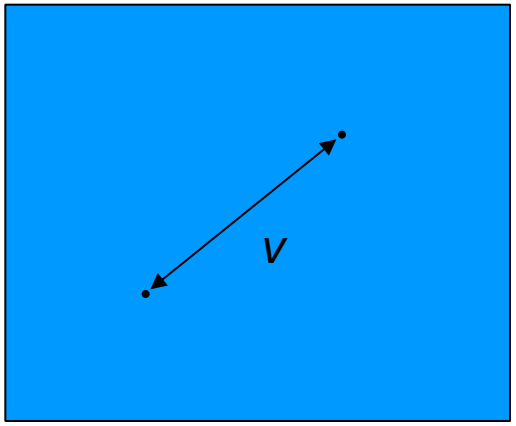
$$-\frac{1}{2}\nabla^2 + V_{\text{ext}}(\mathbf{r}) + \int \frac{n(\mathbf{r}')}{|\mathbf{r}-\mathbf{r}'|} d^3r'$$

complex energy contains
- excitation energies (real part)
- excitation lifetimes (imaginary part)

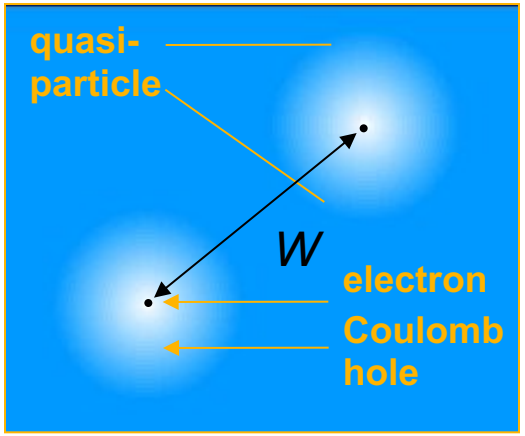
$$\hat{h}_0(\mathbf{r})\psi_n(\mathbf{r}) + \int \Sigma(\mathbf{r},\mathbf{r}';E_n)\psi_n(\mathbf{r}')d^3r' = E_n\psi_n(\mathbf{r})$$

Theory
Self-energy

strong electron interaction v



weak quasiparticle interaction W



Expansion up to linear order

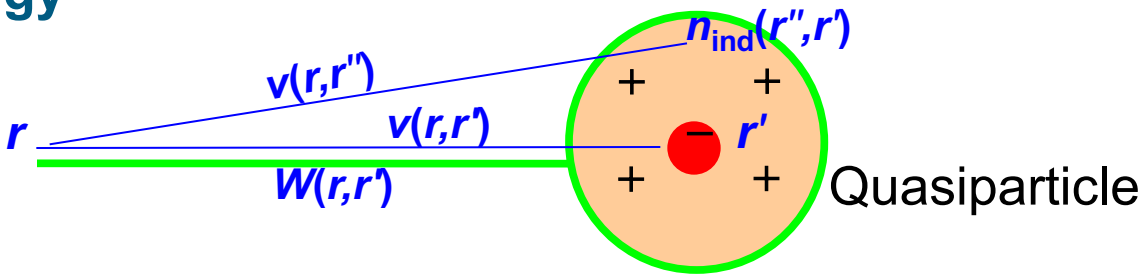
in Coulomb interaction v $\longrightarrow$ $\Sigma^{HF} = iG_0 v$ (Hartree-Fock)

in screened interaction W $\longrightarrow$ $\Sigma^{GW} = iG_0 W$ (*GW* approximation)

The *GW* approximation contains electron exchange and a large part of electron correlation.

$$\Sigma^{GW}(\mathbf{r},\mathbf{r}';\tau) = iG_0(\mathbf{r},\mathbf{r}';\tau)W(\mathbf{r},\mathbf{r}';\tau+\eta)$$

Theory
Self-energy

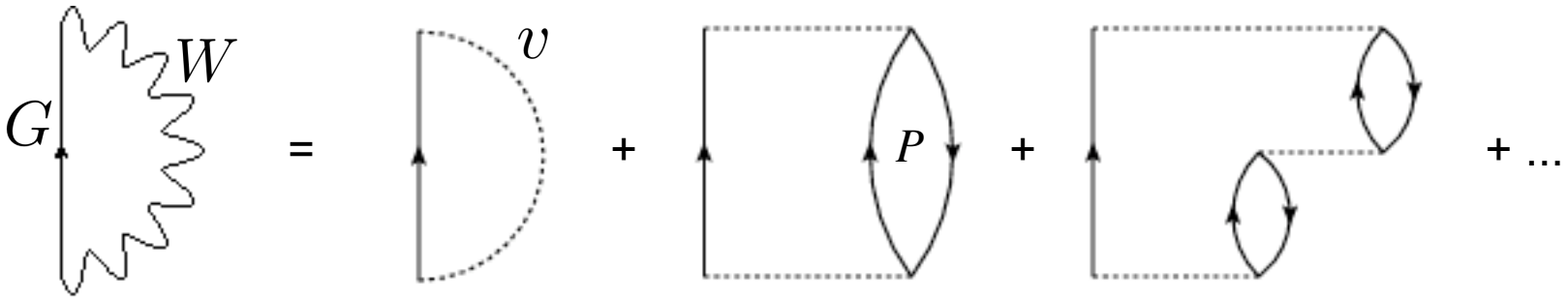


$$W(\mathbf{r},\mathbf{r}') = v(\mathbf{r},\mathbf{r}') + \int v(\mathbf{r},\mathbf{r}'')n_{\mathrm{ind}}(\mathbf{r}'',\mathbf{r}')\,d^3r''$$

$$= Pv + PvPv + PvPvPv + \ldots$$

$$W = v + vPv + vPvPv + vPvPvPv + \ldots = v + vPW$$

$$\Sigma = iGW = iGv + iGvPv + iGvPvPv + \ldots$$



Theory
Hedin equations

Lars Hedin, 1965

$$1 = (\mathbf{r}_1, \sigma_1, t_1),\ 2 = \ldots$$

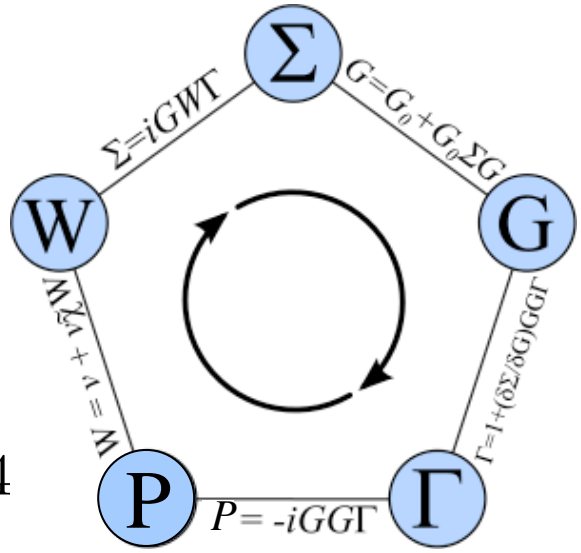
$$\Gamma(12;3) = \delta(12)\delta(13) - \iiint \frac{\delta\Sigma(12)}{\delta G(45)} G(56)\Gamma(67;3)G(74)\, d4\, d5\, d6\, d7$$

$$P(12) = -i \iint G(13)\Gamma(34;2)G(41)\, d3\, d4$$

$$W(12) = v(12) + \iint v(13)P(34)W(42)\, d3\, d4$$

$$\Sigma(12) = i \iint G(13)W(1^+4)\Gamma(32;4)\, d3\, d4$$

$$G(12) = G_0(12) + \iint G_0(13)\Sigma(34)G(42)\, d3\, d4$$



GW approximation corresponds to the 1st iteration starting from Σ=0:

$$\Gamma(12;3) = \delta(12)\delta(13)$$
$$P(12) = -iG(12)G(21) \quad \text{(random-phase approximation)}$$
$$\Sigma(12) = iG(12)W(1^+2)$$

Implementation

Dyson equation ➔ quasiparticle equations:

true excitation energies

GW: $\hat{h}_0(\mathbf{r})\psi^\sigma_{\mathbf{k}n}(\mathbf{r}) + \int \Sigma^{GW}_\sigma(\mathbf{r},\mathbf{r}';E^\sigma_{\mathbf{k}n})\psi^\sigma_{\mathbf{k}n}(\mathbf{r}')d^3r' = E^\sigma_{\mathbf{k}n}\psi^\sigma_{\mathbf{k}n}(\mathbf{r})$

DFT: $\hat{h}_0(\mathbf{r})\varphi^\sigma_{\mathbf{k}n}(\mathbf{r}) + v^{\mathrm{xc}}_\sigma(\mathbf{r})\varphi^\sigma_{\mathbf{k}n}(\mathbf{r}) = \epsilon^\sigma_{\mathbf{k}n}\varphi^\sigma_{\mathbf{k}n}(\mathbf{r})$

energies of a fictitious system

Similarity motivates the use of perturbation theory

$$E^\sigma_{\mathbf{k}n} = \epsilon^\sigma_{\mathbf{k}n} + \langle\varphi^\sigma_{\mathbf{k}n}|\Sigma^{GW}_\sigma(E^\sigma_{\mathbf{k}n}) - v^{\mathrm{xc}}_\sigma|\varphi^\sigma_{\mathbf{k}n}\rangle$$

direct solution

$$\Sigma(E) = \Sigma(\epsilon) + \Sigma'(\epsilon)(E-\epsilon)$$

$$E^\sigma_{\mathbf{k}n} = \epsilon^\sigma_{\mathbf{k}n} + Z^\sigma_{\mathbf{k}n}\langle\varphi^\sigma_{\mathbf{k}n}|\Sigma^{GW}_\sigma(\epsilon^\sigma_{\mathbf{k}n}) - v^{\mathrm{xc}}_\sigma|\varphi^\sigma_{\mathbf{k}n}\rangle$$

linearized solution

renormalization factor

$$Z^\sigma_{\mathbf{k}n} = \left(1 - \langle\varphi^\sigma_{\mathbf{k}n}|\Sigma'^{GW}_\sigma(\epsilon^\sigma_{\mathbf{k}n})|\varphi^\sigma_{\mathbf{k}n}\rangle\right)^{-1}$$

Implementation
Basis sets

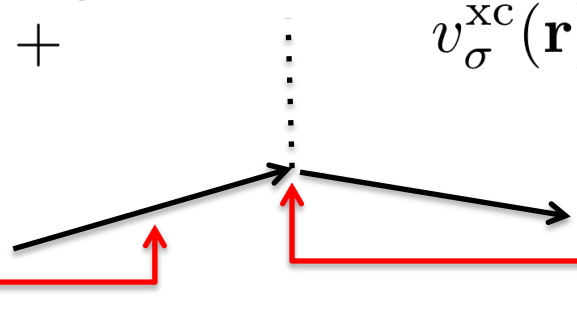
Dyson equation ➔ quasiparticle equations:

true excitation energies

GW: $\hat{h}_0(\mathbf{r})\psi^\sigma_{\mathbf{k}n}(\mathbf{r}) + \int \Sigma^{GW}_\sigma(\mathbf{r},\mathbf{r}';E^\sigma_{\mathbf{k}n})\psi^\sigma_{\mathbf{k}n}(\mathbf{r}')d^3r' = E^\sigma_{\mathbf{k}n}\psi^\sigma_{\mathbf{k}n}(\mathbf{r})$

DFT: $\hat{h}_0(\mathbf{r})\varphi^\sigma_{\mathbf{k}n}(\mathbf{r}) + v^{\mathrm{xc}}_\sigma(\mathbf{r})\varphi^\sigma_{\mathbf{k}n}(\mathbf{r}) = \epsilon^\sigma_{\mathbf{k}n}\varphi^\sigma_{\mathbf{k}n}(\mathbf{r})$

energies of a fictitious system



Basis set for wavefunctions

$$\varphi_{\mathbf{k}n}(\mathbf{r}) = \sum_\mu c^\mu_{\mathbf{k}n}\,\zeta_{\mathbf{k}\mu}(\mathbf{r})$$

Gaussians
Plane waves (Pseudopotential)
PAW
LMTO
LAPW
...

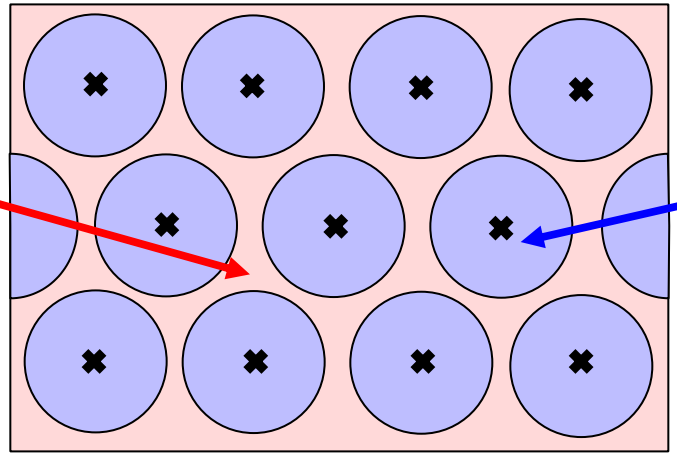
Basis set for wavefunction products

$$\varphi^*_{\mathbf{k}n}(\mathbf{r})\varphi_{\mathbf{k}'n'}(\mathbf{r}) = \sum_\nu C^\nu_{\mathbf{k}\mathbf{k}'nn'}\,\xi_{\mathbf{k}\mathbf{k}'\nu}(\mathbf{r})$$

Auxiliary Gaussian set (density fitting)
Plane waves
Plane waves
Product basis
Mixed product basis
...

Implementation
FLAPW method

SPEX



Interstitial region:

interstitial plane waves

$e^{\mathrm{i}(\mathbf{k}+\mathbf{G})\mathbf{r}}$

cutoff $|\mathbf{k}+\mathbf{G}| \leq G_{\max}$

Muffin-tin (MT) spheres:

numerical MT functions

$u_{lp}(r)Y_{lm}(\hat{\mathbf{r}})$

cutoff $l \leq l_{\max}$

In our *GW* implementation we use the mixed product basis, generated from the products of (1) interstitial plane waves (cutoff $G'_{\max}$)

$$e^{\mathrm{i}(\mathbf{k}+\mathbf{G})\mathbf{r}} e^{\mathrm{i}(\mathbf{k}'+\mathbf{G}')\mathbf{r}} = e^{\mathrm{i}(\mathbf{k}+\mathbf{k}'+\underbrace{\mathbf{G}+\mathbf{G}'}\mathbf{)r}}$$

➔ $G'_{\max} = 2G_{\max}$

and (2) MT functions (cutoff $l'_{\max}$)

$$u_{lp}(r)u_{l'p'}(r)\underbrace{Y_{lm}(\hat{\mathbf{r}})Y_{l'm'}(\hat{\mathbf{r}})}$$

up to $l+l'$ ➔ $l'_{\max} = 2l_{\max}$

T. Kotani and M. van Schilfgaarde,
Solid State Commun. 121, 461 (2002).

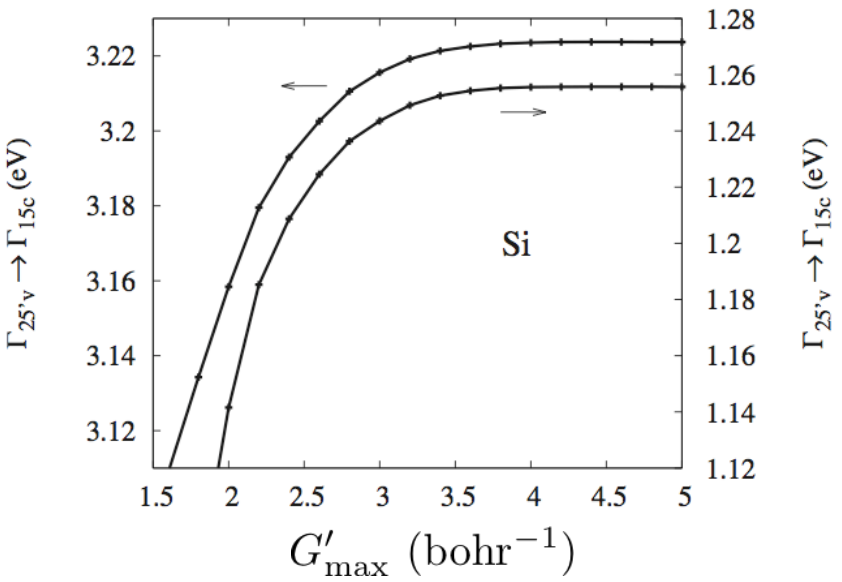
Implementation
Mixed product basis

$$G_{\max} = 3.6\,\mathrm{bohr}^{-1}$$

exact $\quad G'_{\max} = 2G_{\max}$

converged $\quad G'_{\max} \sim 0.75\,G_{\max}$

`spex.inp:` keyword “GCUT”.

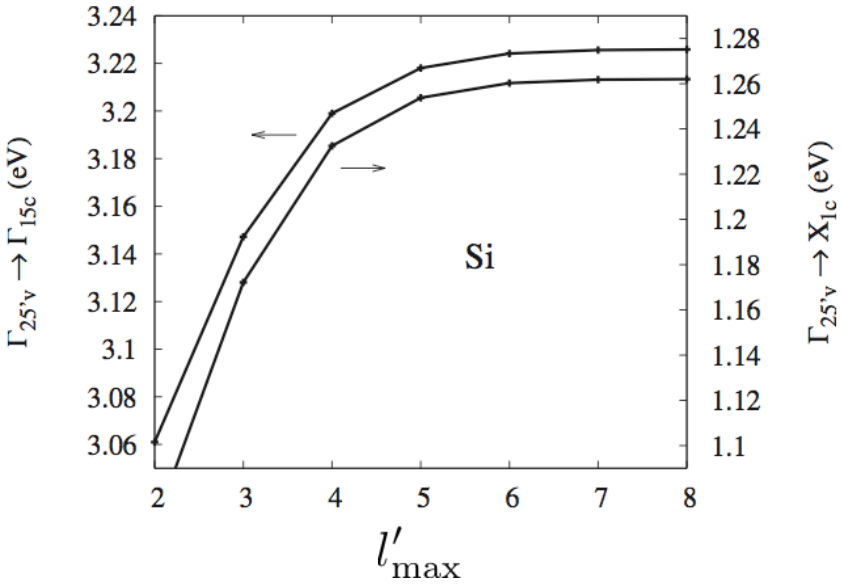


$$l_{\max} = 8$$

exact $\quad l'_{\max} = 2l_{\max}$

converged $\quad l'_{\max} \sim 0.63\,l_{\max}$

`spex.inp:` keyword “LCUT”.



Implementation
Exchange Self-energy

The self-energy can be decomposed into an exchange and a correlation term:

$$\Sigma^{GW}(\omega) = iG_0W = iG_0v + iG_0(W - v) = \Sigma^{\mathrm{x}} - \Sigma^{\mathrm{c}}(\omega)$$

The exchange contribution is given analytically by the Hartree-Fock expression

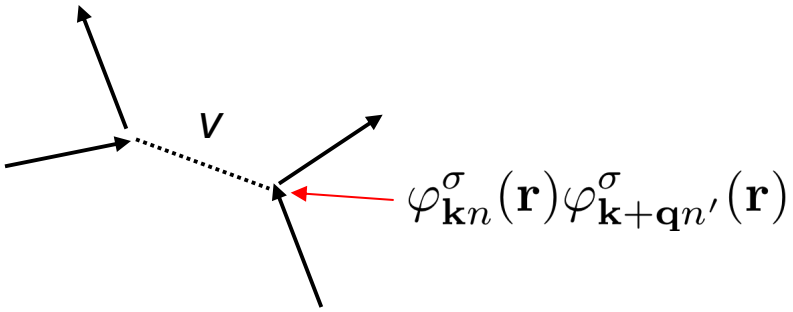
$$\Sigma^{\mathrm{x}}(\mathbf{r},\mathbf{r}';-\eta) = iG(\mathbf{r}t,\mathbf{r}'t+\eta)v(\mathbf{r},\mathbf{r}') = -\langle\Psi_0^N|\hat{\psi}^\dagger(\mathbf{r}'t+\eta)\hat{\psi}(\mathbf{r}t)|\Psi_0^N\rangle v(\mathbf{r},\mathbf{r}')$$

Density matrix

$$n(\mathbf{r},\mathbf{r}') = \sum_n^{\mathrm{occ.}} \varphi_n^*(\mathbf{r})\varphi_n(\mathbf{r}')$$

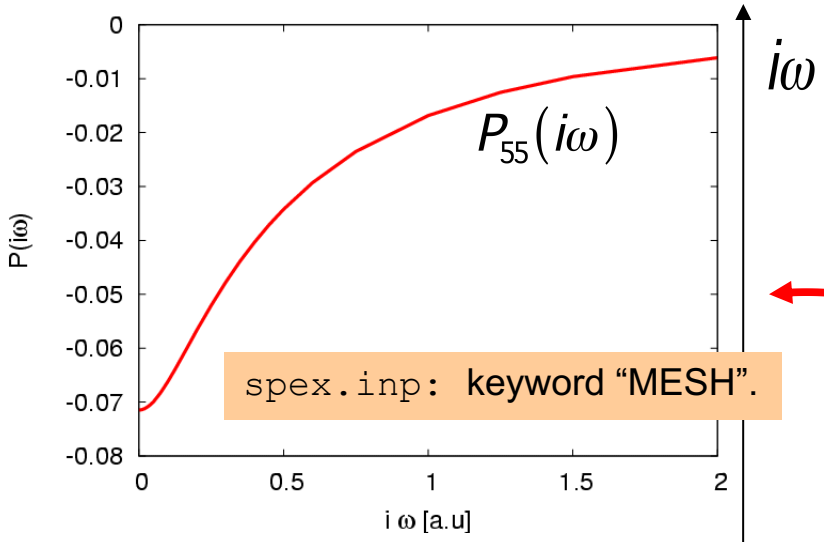
$$\langle\varphi_m|\Sigma^{\mathrm{x}}|\varphi_{m'}\rangle = -\sum_n^{\mathrm{occ.}} \iint \varphi_m^*(\mathbf{r})\varphi_n(\mathbf{r})v(\mathbf{r},\mathbf{r}')\varphi_n^*(\mathbf{r}')\varphi_{m'}(\mathbf{r}')\,d^3r\,d^3r'$$

$$1 = \sum_I |M_I\rangle\langle M_I|$$



Implementation

Imaginary-frequency formulation



`spex.inp:` keyword “MESH”.

The quantities (e.g. the polarization function *P*) are much smoother along the imaginary frequency axis than along the real axis.

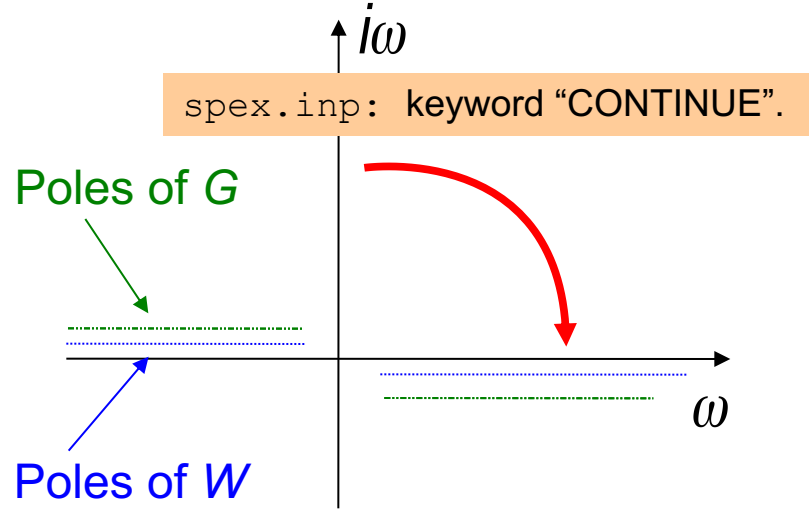
`spex.inp:` keyword “HILBERT”.

Implementation
Correlation Self-energy

$$\Sigma^c(\omega) = \frac{i}{2\pi}\int_{-\infty}^{\infty} G_0(\omega+\omega')W^c(\omega')d\omega'$$

Analytic Continuation

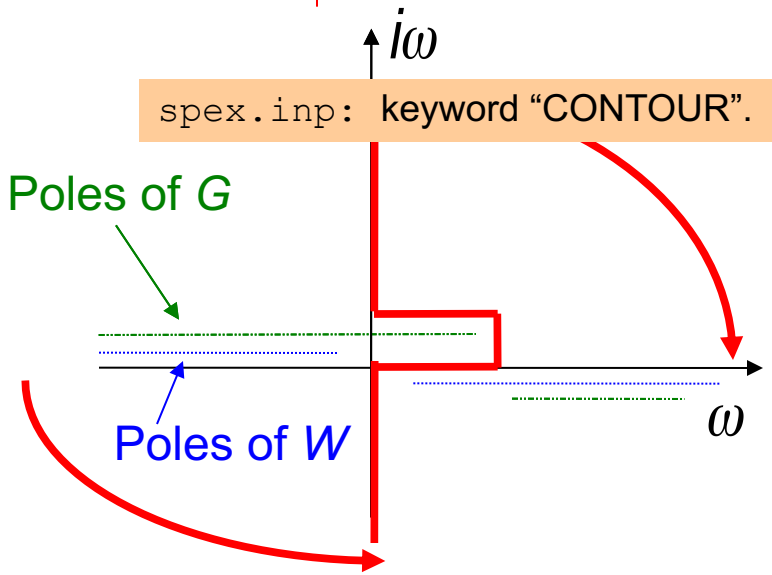
$$\Sigma^c(i\omega) = -\frac{1}{2\pi}\int_{-\infty}^{\infty} G_0(i\omega+i\omega')W^c(i\omega')d\omega'$$



- Easy to implement
- Fast computation
- Analytic continuation critical

Contour integration

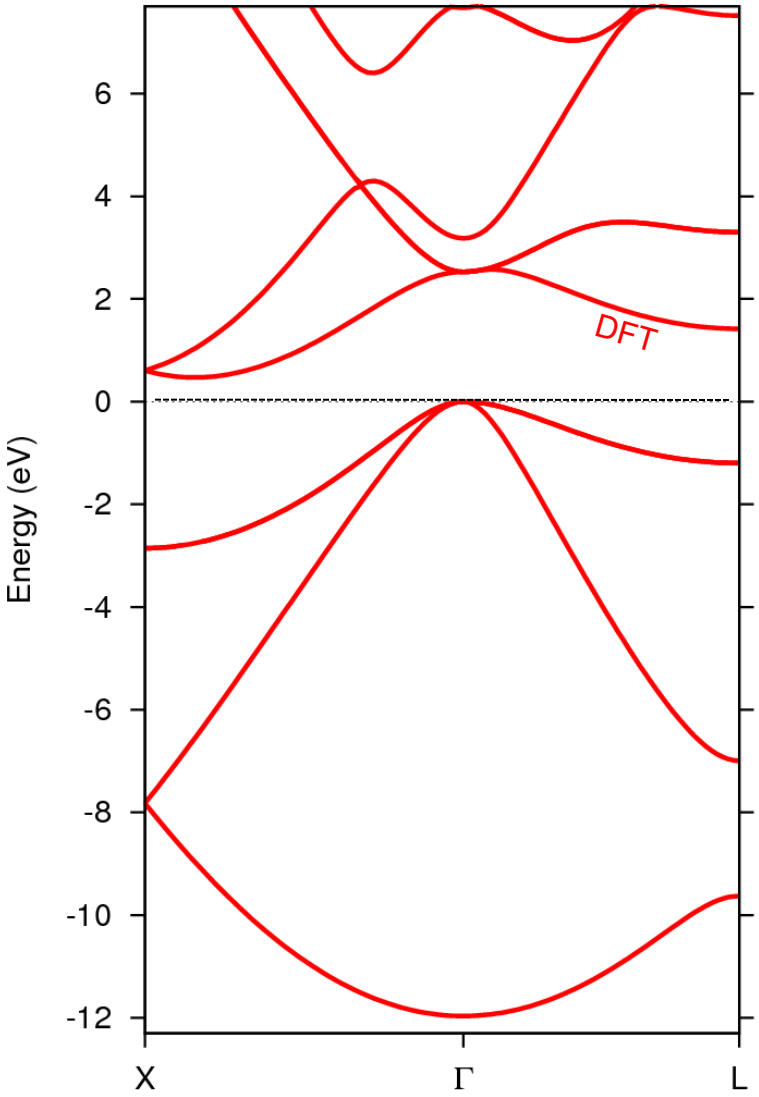
$$\Sigma^c(\omega) = \frac{i}{2\pi}\int G_0(\omega+\omega')W^c(\omega')d\omega'$$



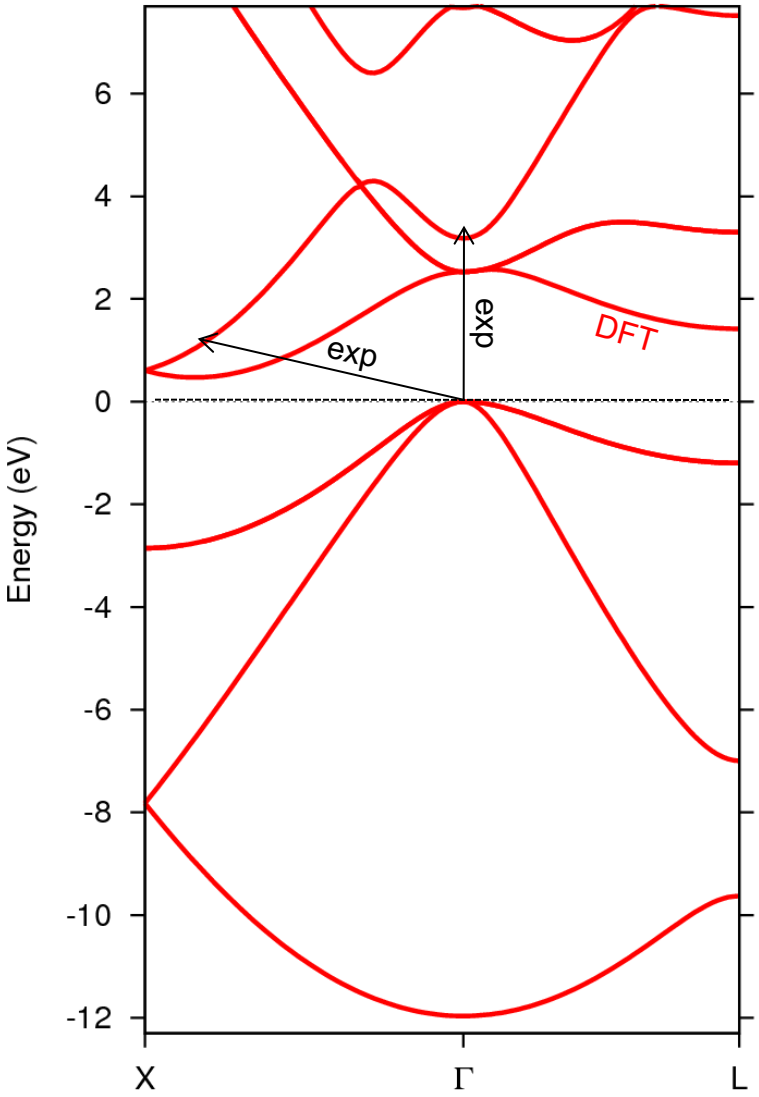
- Accurate evaluation of Σ^c
- More parameters necessary
- Takes more time

Applications

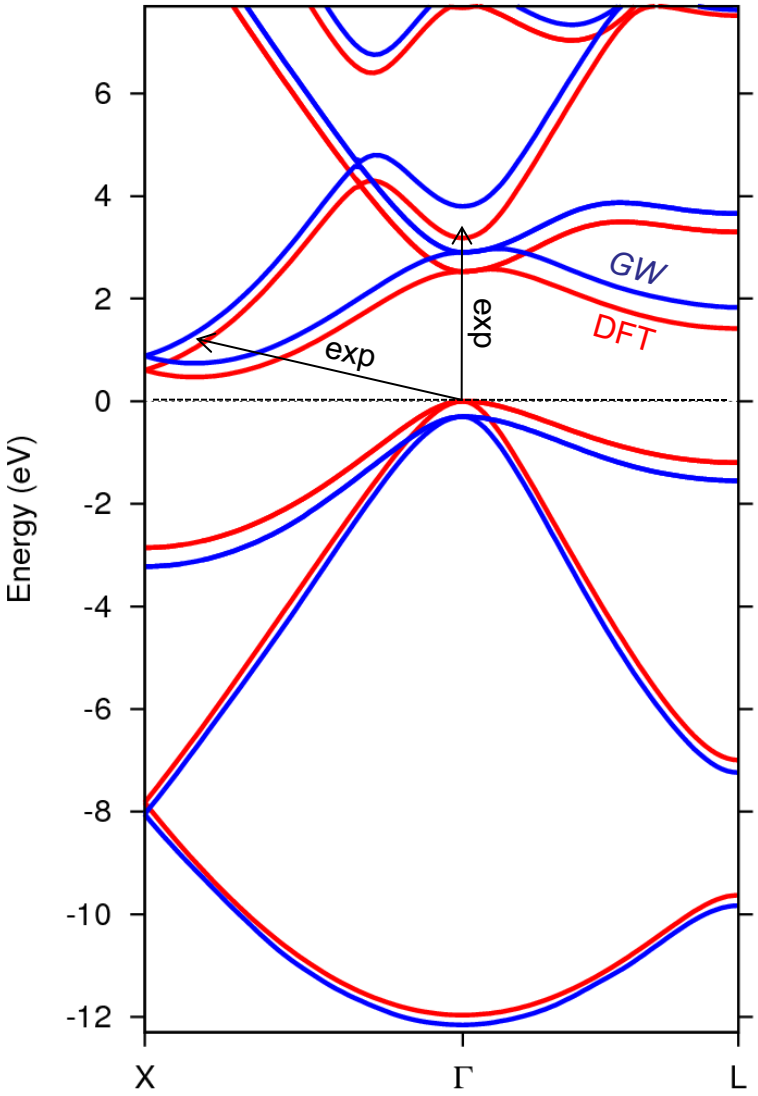
Bulk Silicon



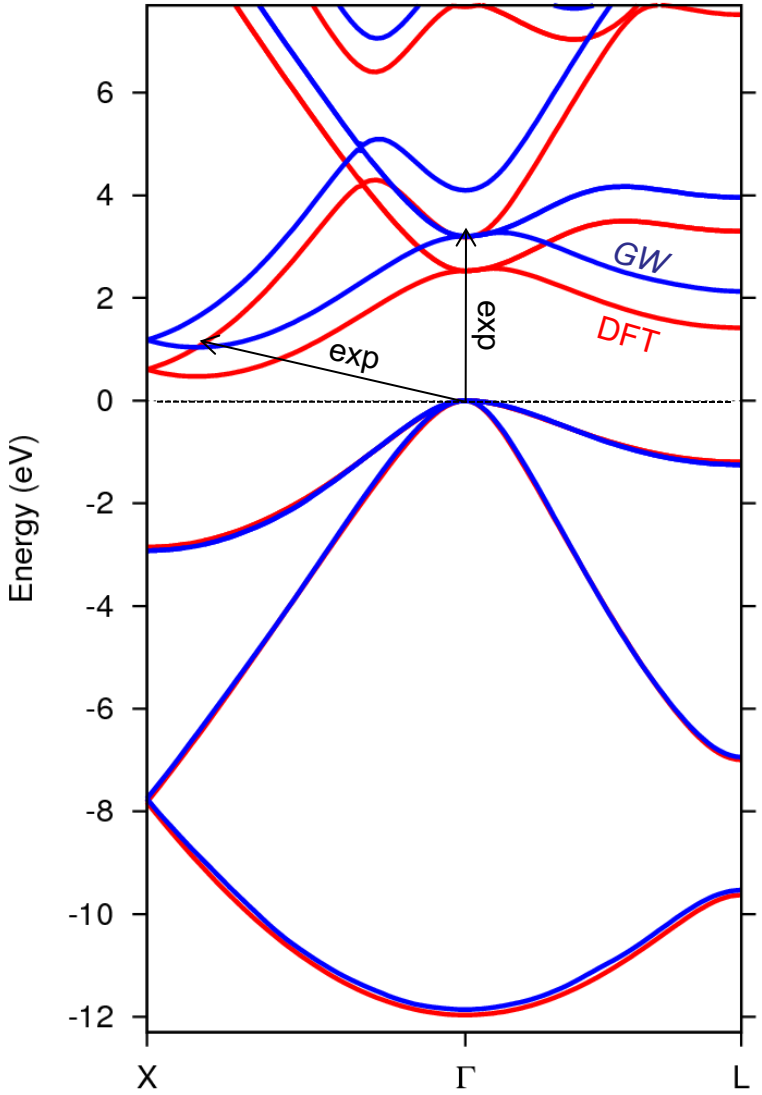
Applications
Bulk Silicon



Applications
Bulk Silicon



Applications
Bulk Silicon

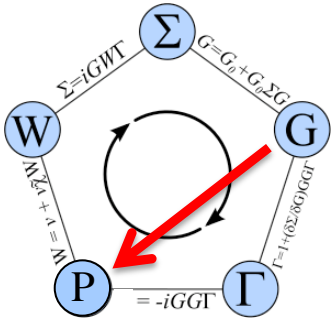


- Si was the first material to which *GW* was applied (Hybertsen, Louie 1985; Godby, Schlüter, Sham 1986).
- The one-shot *GW* calculation yields more accurate band gaps:

	DFT	*GW*	exp.
direct:	2.53	3.20	3.40 eV
indirect:	0.47	1.04	1.17 eV

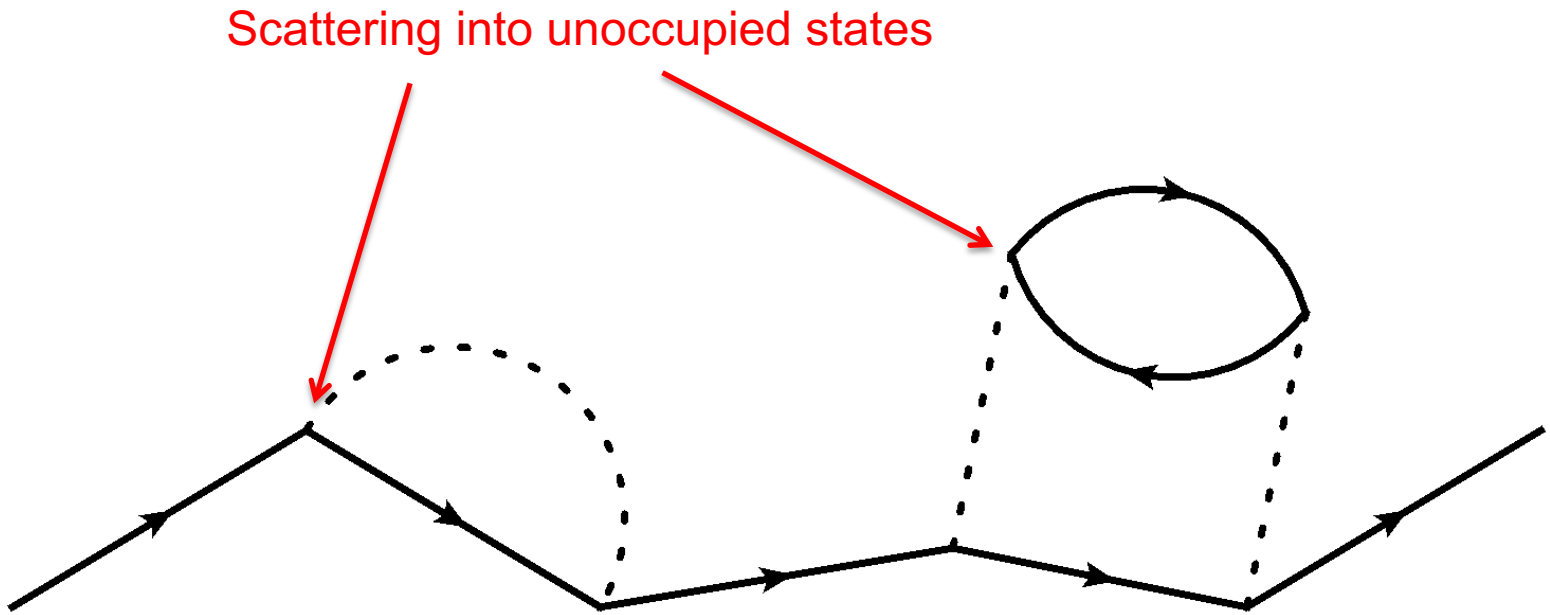
- Quasiparticle self-consistent *GW* (QS*GW*) tends to overestimate the gaps:

	QS*GW*
direct:	3.60 eV
indirect:	1.34 eV



QS*GW*: S. V. Faleev, M. van Schilfgaarde, and T. Kotani, Phys. Rev. Lett. 93, 126406 (2004).

Applications
Band convergence



`spex.inp:` keyword “NBAND”.

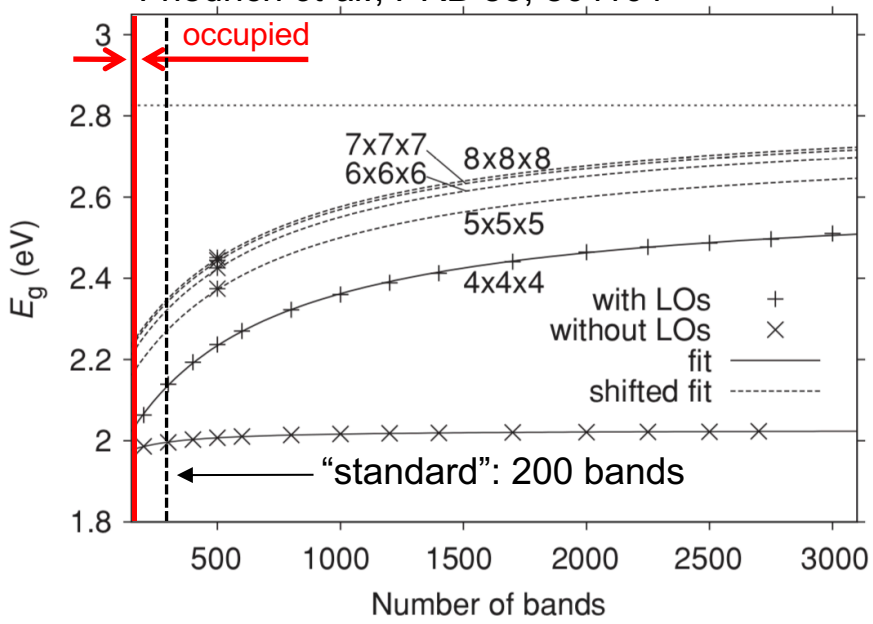
Applications

Zinc Oxide - band convergence

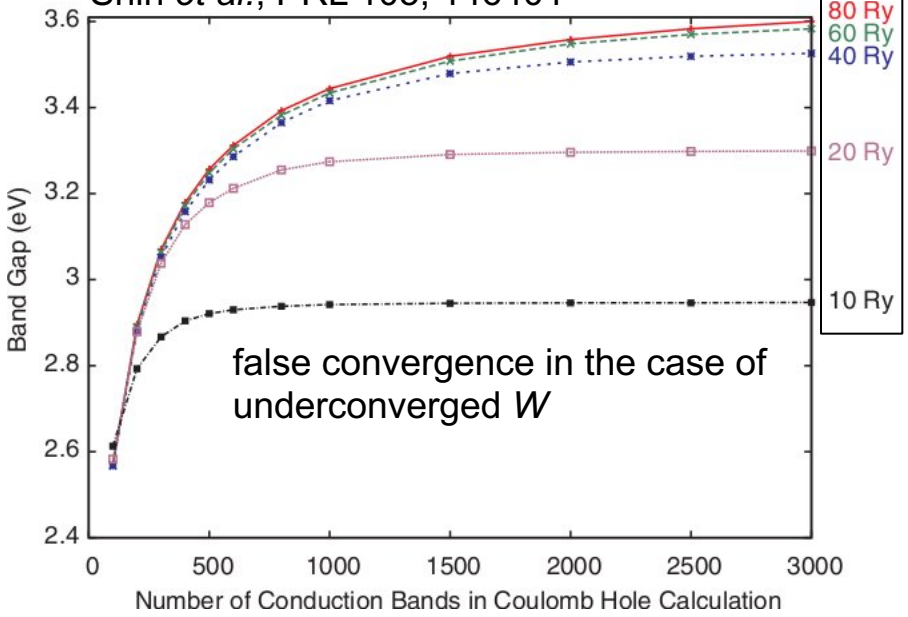
Large scatter of band-gap values from *one-shot GW* calculations (exp: 3.4 eV):

2.44 eV (FLAPW) [M. Usuda *et al.*, Phys. Rev. B **66**, 125101 (2002)]
2.12 eV (PAW) [M. Shishkin and G. Kresse, PRB **75**, 235102 (2007)
2.14 eV (PAW) [F. Fuchs *et al.*, Phys. Rev. B **76**, 115109 (2007)]
2.6 eV (PW-PP) [P. Gori *et al.*, Phys. Rev. B **81**, 125207 (2010)]
3.4 eV (PW-PP) [B.-C. Shih *et al.*, Phys. Rev. Lett. **105**, 146401 (2010)]
2.83 eV (FLAPW) [C. Friedrich *et al.*, Phys. Rev. B **83**, 081101 (2011)]
…

Friedrich *et al.*, PRB 83, 801101

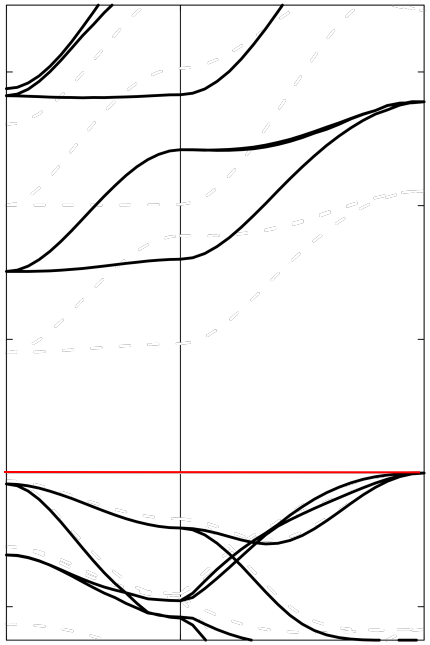


Shih *et al.*, PRL 105, 146401



Applications
GW for metals

semiconductor



The polarization function is sum over virtual transitions in the non-interacting reference system.

$$P_{IJ}(\mathbf{k},\omega) = \sum_{\sigma,\mathbf{q}} \sum_{n}^{occ} \sum_{n'}^{unocc} \dots \left(\frac{1}{\omega + \epsilon^{\sigma}_{\mathbf{q}n} - \epsilon^{\sigma}_{\mathbf{k}+\mathbf{q}n'} + i\eta} - \dots \right)$$

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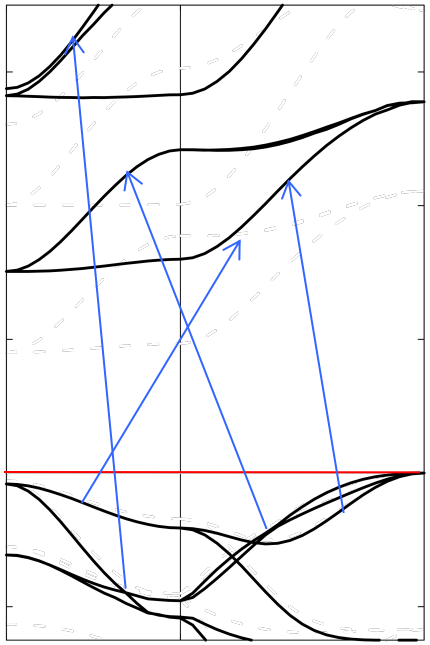
GW for metals

semiconductor



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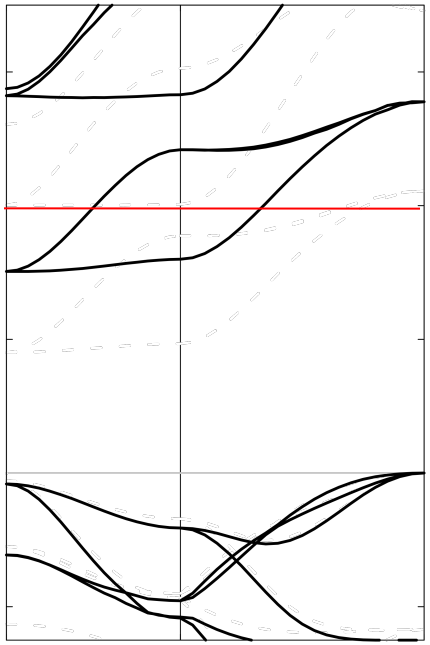
Applications
GW for metals

metal



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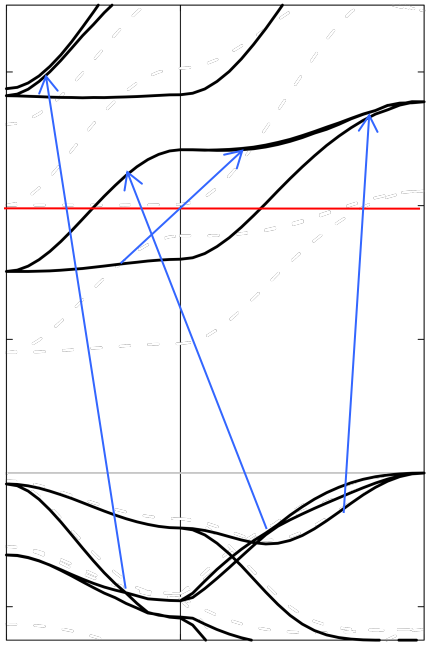

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Applications
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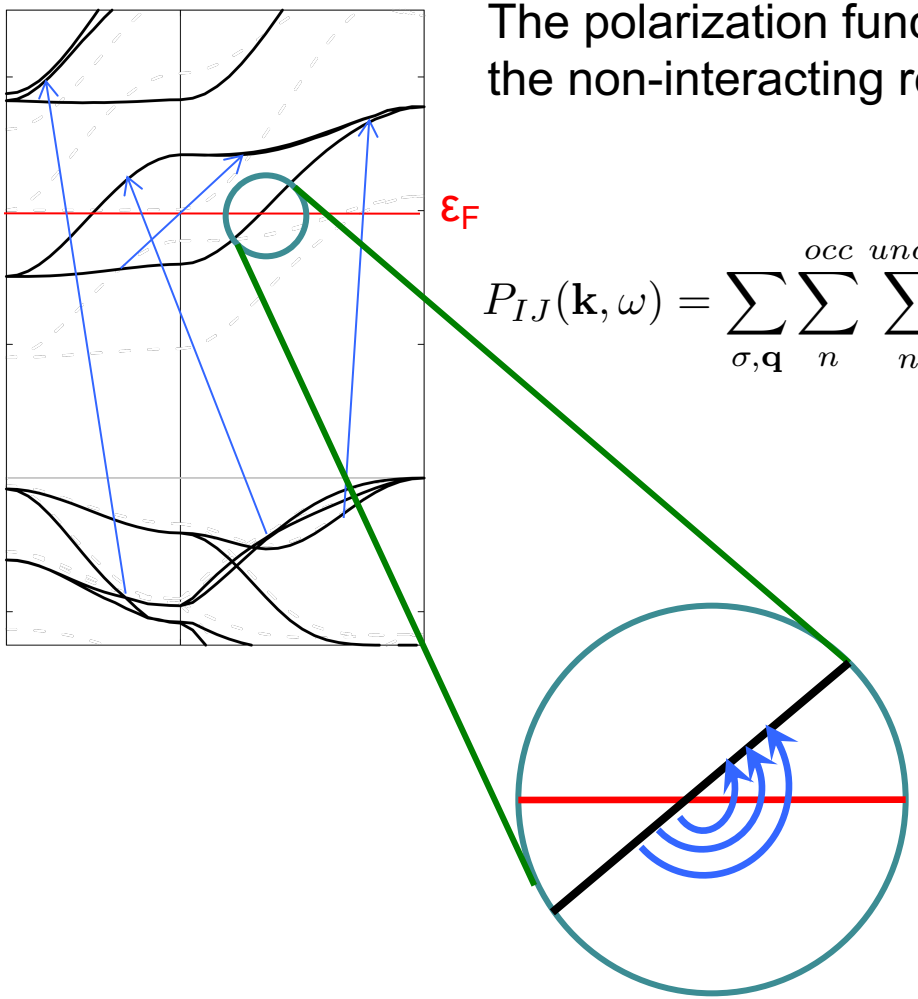

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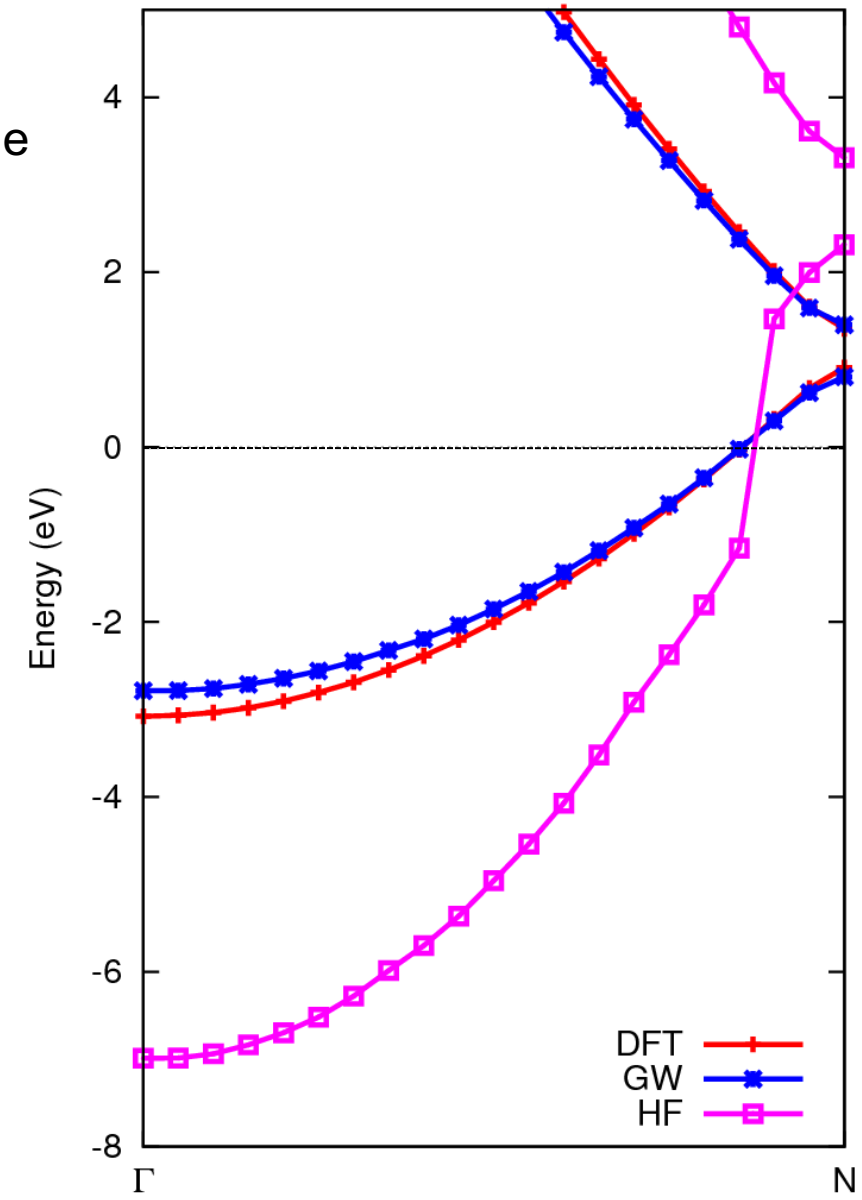
Virtual transitions of zero energy just across the Fermi surface produce the Drude term

$$P^{\mathrm{D}}_{\mathrm{head}}(\mathbf{k}, i\omega) \sim \frac{k^2}{4\pi} \frac{\omega^2_{\mathrm{pl}}}{\omega(\omega + i\eta)}$$

for $k \rightarrow 0$.

Applications
Sodium

- Vanishing of the density of states at the Fermi energy in HF is exactly compensated by the *GW* correlation self-energy.
- *GW* band width smaller than KS band width due to increased effective (quasiparticle) mass.



GW approximation including SOC

Spin-orbit coupling (SOC) arises from the interaction of the electron spin with the **B** field that is created in the rest frame of the electron due to its motion through the material → coupling of spatial and spin degrees of freedom:

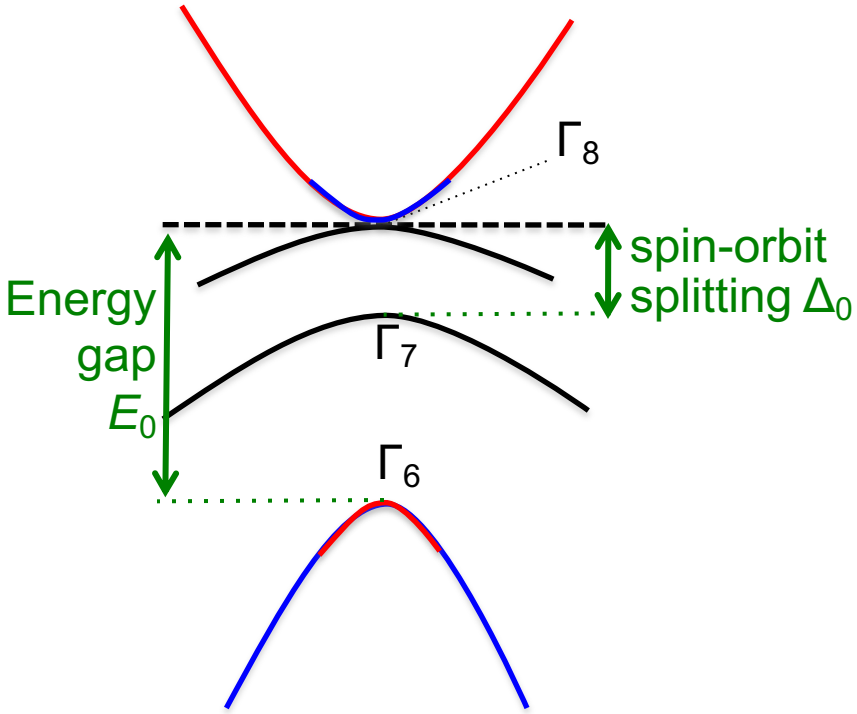
$$G_{\alpha\beta}(\mathbf{r},\mathbf{r}';\omega) = \sum_{\mathbf{k}n} \frac{\varphi_{\mathbf{k}n}(\mathbf{r},\alpha)\varphi_{\mathbf{k}n}(\mathbf{r}',\beta)}{\omega - \varepsilon_{\mathbf{k}n} \pm i\eta}$$

$$P(\mathbf{r},\mathbf{r}';\omega) = -\frac{i}{2\pi}\sum_{\alpha\beta}\int G_{\alpha\beta}(\mathbf{r},\mathbf{r}';\omega+\omega')G_{\beta\alpha}(\mathbf{r}',\mathbf{r};\omega')d\omega'$$

$$\Sigma_{\alpha\beta}(\mathbf{r},\mathbf{r}';\omega) = \frac{i}{2\pi}\int G_{\alpha\beta}(\mathbf{r},\mathbf{r}';\omega+\omega')W(\mathbf{r},\mathbf{r}';\omega')e^{i\eta\omega'}d\omega'$$

F. Aryasetiawan, S. Biermann, PRL **100**, 116402 (2008).
R. Sakuma, C. Friedrich, T. Miyake, S. Blügel, F. Aryasetiawan, PRB **84**, 085144 (2011).

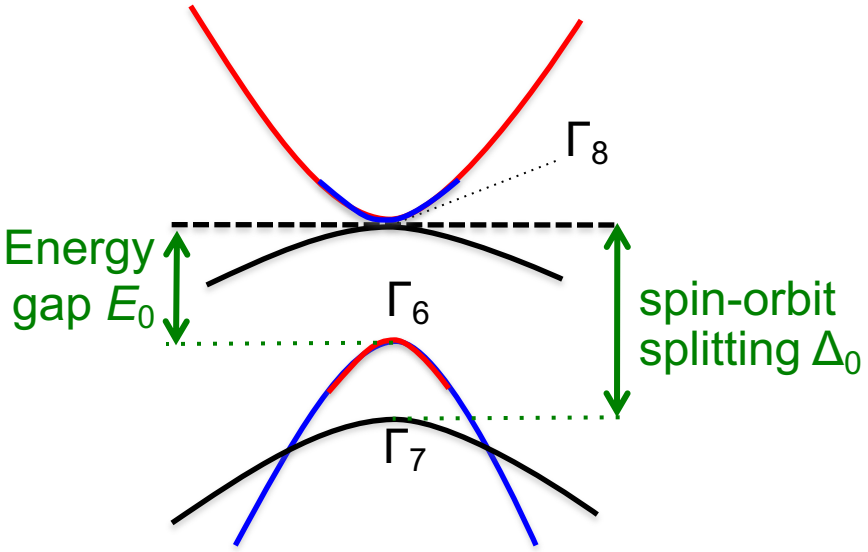
Bulk HgTe



	E_0	Δ_0
LDA+SOC	-1.16	0.74
GW+SOC	-0.59	0.74
$G^{SOC}W^{SOC}$	-0.60	0.83
QS*GW*+SOC	-0.43	0.75
QS$G^{SOC}W^{SOC}$	-0.46	0.93
Experiment	-0.30	0.91

calculations with semicores
Hg 5*p* and Te 4*d*

Bulk HgTe



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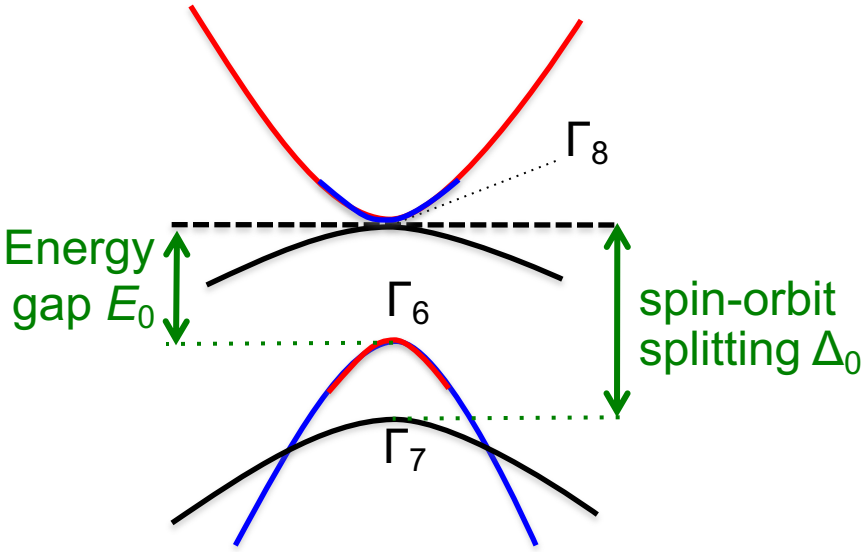

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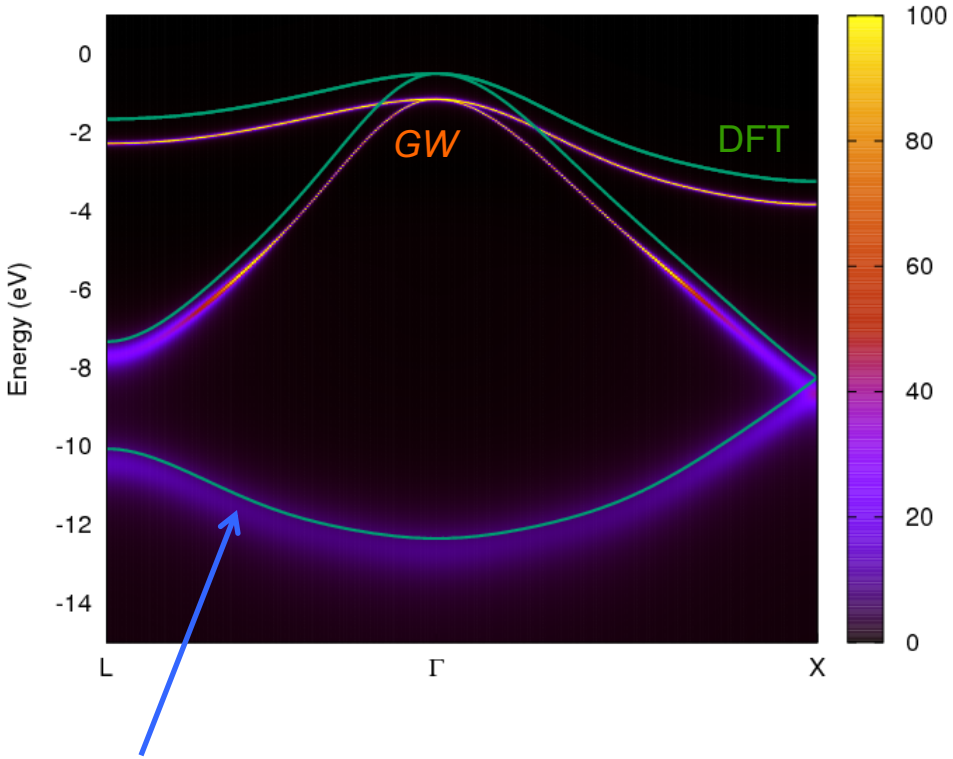
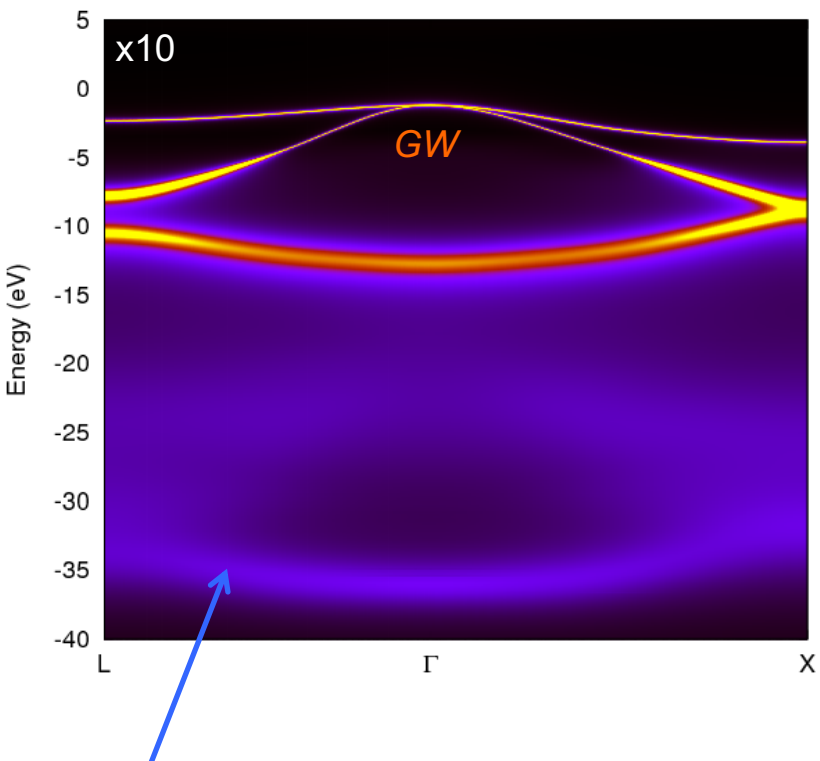
Applications

Many-body effects: lifetime broadening and plasma satellites



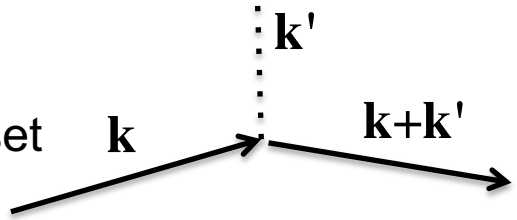


Renormalization due to formation of electron-hole pairs (density fluctuations)

Plasmaron bands (electrons+plasmons)

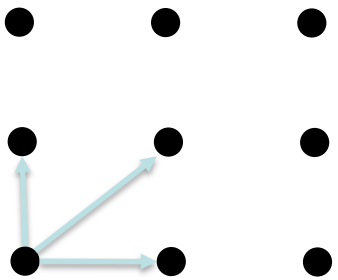
Computational procedure
One-Shot *GW*

- FLEUR: Self-consistent field calculation
  ➔ Density, Exchange-correlation potential
- SPEX: Generate special equidistant k-point set
  k, **k'**, **k+k'**, and **0** must be elements
- FLEUR: Diagonalize Hamiltonian on new k points (non iterative)
  ➔ Kohn-Sham energies and wavefunctions
- SPEX: *GW* calculation
  ➔ Quasiparticle energies



`spex.inp:` ITERATE

`spex.inp:` JOB GW X:(1-4,6)



k-point set

Computational procedure
GW band structure

- FLEUR: Self-consistent field calculation
  ➔ Density, Exchange-correlation potential
- SPEX: Define q-point path
  ➔ {**q**}

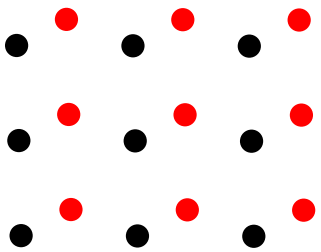
`spex.inp:` KPTPATH (L,G,X)

Loop over **q** (high-symmetry line in Brillouin zone)

- SPEX: Generate two sets of equidistant k-points
  ➔ {**k**}, {**k+q**}
- FLEUR: Diagonalize Hamiltonian on new k points (non iterative)
  ➔ Kohn-Sham energies and wavefunctions
- SPEX: *GW* calculation (*W* has to be calculated only once)
  ➔ Quasiparticle energies at **q**

`spex.inp:` keyword “RESTART”.

“spex.band”



shifted **k**-point set
k-point set

Computational procedure
Wannier interpolation

- FLEUR: Self-consistent field calculation
  ➔ Density, Exchange-correlation potential
- SPEX: Generate special equidistant k-point set
  k, **k'**, **k+k'**, and **0** must be elements
- FLEUR: Diagonalize Hamiltonian on new k points (non iterative)
  ➔ Kohn-Sham energies and wavefunctions
- SPEX: *GW* calculation
  ➔ Quasiparticle energies

`spex.inp:` JOB GW IBZ:(1-4)

`spex.inp:` KPTPATH (L,G,X)

`spex.inp:` section “WANNIER”:

```
SECTION WANNIER
   ORBITALS 1 4 (sp3)
   MAXIMIZE
   INTERPOL
END
```

Computational procedure
Self-consistent *GW* (QS*GW*)

- FLEUR: Self-consistent field calculation
  ➔ Density, Exchange-correlation potential
- SPEX: Generate special equidistant k-point set
  k, **q**, **k+q**, and **0** must be elements

Start of iterations

- FLEUR: Diagonalize Hamiltonian on new k points (non iterative)
  ➔ Kohn-Sham energies and wavefunctions
- SPEX: *GW* calculation on all k points and many bands (costly)
  ➔ Quasiparticle energies and self-energy matrix
- FLEUR: Self-consistent field calculation (with self-energy!)
  ➔ Density, Exchange-correlation potential

"spex.selfc"

`spex.inp:` JOB GW FULL IBZ:(1-4)

Compilation and Installation

- Configuration
  `% ./configure --with-wan --prefix=$HOME`
  generates the Makefile for the computer system
- Compilation
  `% make`
  produces the three executables spex.inv, spex.noinv, and spex.extr; the launcher script "spex" always calls the correct executable
- Installation
  `% make install`
  installs executables and scripts into $HOME/bin
  - The PATH environment variable should contain $HOME/bin
    If not: `% export PATH=$PATH:$HOME/bin`

Summary

- Excitation energies and lifetimes of the (N+1) and (N-1)-electron system can be readily obtained from the one-particle Green function. These excitation energies form the band structure in solids.

- The Green function obeys an integral Dyson equation which may be rewritten as a quasiparticle equation with the self-energy as a scattering potential that takes into account all exchange and correlation effects beyond the Hartree potential.

- The *GW* approximation constitutes the expansion of the self-energy up to linear order in the screened interaction W.

- It is usually implemented as a perturbative correction on a DFT band structure. But a self-consistent solution (QS*GW*) is possible, too.