

| Onderbouwend | Stof | Effect | Code |
|---|--|---|------|
| Li Y. J Clin Pharmacol. 2018;58:1295-1304. doi: 10.1002/jcph.1145. | pomalidomide + fluvoxamine | ↑pomalidomide AUC _{0-inf} 2.25x en Cmax 1.24x door fluvoxamine; t _{1/2} 6.0 → 13.1 h; CL/F 7.6 → 3.4 L/h. Regime: pomalidomide 4 mg 1x op dag 1; fluvoxamine 50 mg 2dd op dag 4-7; pomalidomide 4 mg 1x plus fluvoxamine 50 mg 2dd op dag 8; fluvoxamine 50 mg 2dd op dag 9-10; studie met 14 vrijwilligers. | 3A |
| SPC Imnovid rev.15 18-6-19 | pomalidomide + fluvoxamine | ↑AUC pomalidomide 2.25x door fluvoxamine Regime: niet bekend. → getal uit Li 2018 | 1A |
| | pomalidomide + fluvoxamine icm ketoconazol | ↑ blootstelling pomalidomide 2.1x door fluvoxamine plus ketoconazol tov combinatie met ketoconazol. Regime: pomalidomide + fluvoxamine plus ketoconazol vs pomalidomide + ketoconazol. | |
| pomalidomide + ketoconazol | pomalidomide + ketoconazol | combinatie met alleen ketoconazol had geen klinisch relevant effect op de blootstelling aan pomalidomide. | 3A |
| | Kasserra C. J Clin Pharmacol 2015;55:168-78. doi: 10.1002/jcph.384. | pomalidomide + fluvoxamine icm ketoconazol | |

| Overig | Stof | Effect |
|--|--|--|
| SPC Imnovid rev.15 18-6-19 | pomalidomide + CYP1A2-rem. | bij combinatie met sterke CYP1A2-remmers dosis pomalidomide halveren |
| Procedural steps taken and scientific information after the authorisation 24-6-16. www.ema.europa.eu Geraadpleegd 15-9-2016. | pomalidomide + fluvoxamine | study: fluvoxamine resulted in an approximate doubling of exposure to pomalidomide. The SmPC is updated to instruct that if strong inhibitors of CYP1A2 are co-administered, the dose of pomalidomide should be reduced by 50%. |
| Procedural steps taken and scientific information after the authorisation 18-8-2014 www.ema.europa.eu Geraadpleegd 20-11-2014. | pomalidomide + ketoconazol | Study CC-4047-CP-008: co-administration of a strong CYP3A4/Pgp inhibitor (ketoconazole) had no clinically relevant effect on mean exposure to pomalidomide. Co-administration of a strong CYP1A2 inhibitor (fluvoxamine) in the presence of a strong CYP3A4 inhibitor approximately doubled the mean exposure to pomalidomide. Pomalidomide was generally well tolerated by healthy subjects when administered as single 4-mg oral doses with multiple oral doses of ketoconazole, fluvoxamine, and/or carbamazepine. |
| | pomalidomide + fluvoxamine icm ketoconazol | |
| EPAR Imnovid 13-8-2013 | pomalidomide | No relevant safety drug-drug interactions have been identified. CC-4047-MM-002: dose escalation study, pomalidomide 2, 3, 4 or 5 mg 1dd for 21 days out of a 28 days cycle. The MTD was determined to be 4 mg daily based on dose limiting toxicity of neutropenia. Addition of dexamethasone improved the results for 1 year event-free rate; also when adding dexamethasone neutropenia was reported less frequently. |

Opmerkingen

Werkgroep Interacties oncologische middelen 18-1-17: nieuwe info uit SPC heeft geen gevolgen.

Idem 14-1-2015: ophogen van 1A naar 3A door studie Kasserra 2014, verder geen wijziging.

Idem 8-1-2014: actie Nee. Pomalidomide wordt al zeer strikt gecontroleerd op neutropenie. Je beoogt juist pancytopenie. Hematoloog: kost ong. 90.000 euro/jaar/patiënt, en biedt ong. 7 maanden verlenging leven.

PubMed feb. 2020: pomalidomide + fluvoxamine: studie Li 2018. Niets op ciprofloxacin. Heeft geen gevolgen voor beoordeling.

PubMed sept 2016: pomalidomide + fluvoxamine /ciprofloxacin/ketoconazol: studie Kasserra 2015.
Hansten, Stockley: niet in.

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|----------------------|--|
| Risicofactoren | |
| Mitigerende factoren | |

| | Interactie | Actie | Datum |
|----------------------|-------------------|--------------|-----------------|
| Beslissing WG OncolA | Ja | Nee | 18 januari 2017 |