

KURZPROTOKOLL PALLAS

Öffentlicher Titel	Phase III Studie zu adjuvanter endokriner Therapie mit und ohne Palbociclib bei Her2-negativem, Hormonrezeptor-positivem Brustkrebs im Frühstadium
Wissenschaftl. Titel	PALbociclib CoLaborative Adjuvant Study: A Randomized Phase III Trial of Palbociclib With Standard Adjuvant Endocrine Therapy Versus Standard Adjuvant Endocrine Therapy Alone for Hormone Receptor Positive (HR+) / Human Epidermal Growth Factor Receptor 2 (HER2)-Negative Early Breast Cancer (PALLAS)
Kurztitel	PALLAS
Studienart	multizentrisch, randomisiert, offen/unverblindet, Pharma-Studie, zweiarmig
Studienphase	Phase III
Erkrankung	Geschlechtsorgane: Brustkrebs: adjuvant
Einschlusskriterien	<ul style="list-style-type: none">- Signed informed consent prior to study specific procedures.- Age 18 years (or per national guidelines).- Pre- and postmenopausal women or men with Stage II (Stage IIA limited to max. 1000 patients) or Stage III early invasive breast cancer- Patients with multicentric and/or multifocal and/or bilateral early invasive breast cancer are eligible if all histopathologically examined tumors meet pathologic criteria for ER+ and/or PR+ and HER2-.- Patients must have histologically confirmed ER+ and/or PR+, HER2-, early invasive breast cancer.- Patients must have undergone breast surgery for the current malignancy. FFPE tumor tissue block must be confirmed to be received at the central sample repository prior to randomization.- ECOG performance status 0-1.- Patients must be able and willing to swallow and retain oral medication.- Serum or urine pregnancy test must be negative in premenopausal women within 14 days of randomization, or in women with amenorrhea of less than 12 months at time of randomization.- Serum or urine pregnancy test must be negative in premenopausal women within 14 days of randomization, or in women with amenorrhea of less than 12 months at time of randomization.- Serum or urine pregnancy test must be negative in premenopausal women within 14 days of randomization, or in women with amenorrhea of less than 12 months at time of randomization.- Serum or urine pregnancy test must be negative in premenopausal women within 14 days of randomization, or in women with amenorrhea of less than 12 months at time of randomization.- Patients must either be initiating or have already started adjuvant hormonal treatment- Patients who already received neo/adjuvant endocrine therapy are eligible as long as they are enrolled within 12 months of initial histological diagnosis and after completing no more than 6 months of adjuvant endocrine therapy.- Absolute neutrophil count 1,500/μL- Platelets 100,000/ mm³- Hemoglobin 10g/dL- Total serum bilirubin ULN; or total bilirubin 3.0 \times ULN with direct bilirubin within normal range in patients with documented Gilbert's Syndrome.- Aspartate amino transferase (AST or SGOT) and alanine amino transferase (ALT or SGPT) 1.5 \times institutional ULN.- Serum creatinine within normal institutional limits or creatinine clearance 60 mL/min/1.73 m² for patients with serum creatinine levels above institutional ULN.

KURZPROTOKOLL PALLAS

Ausschlusskriterien	<ul style="list-style-type: none">- Concurrent therapy with other Investigational Products.- Prior therapy with any CDK inhibitor.- Patients with Stage I or IV breast cancer are not eligible.- History of allergic reactions attributed to compounds of chemical or biologic composition similar to palbociclib.- Patients receiving any medications or substances that are potent inhibitors or inducers of- CYP3A isoenzymes within 7 days of randomization.- Uncontrolled intercurrent illness that would limit compliance with study requirements.- Pregnant women, or women of childbearing potential without a negative pregnancy test within 14 days prior to randomization.- Patients with a history of any malignancy are ineligible (for exceptions see: Pallas Protocol, v1.0, Exclusion criteria 8).- Patients who previously received endocrine therapy within 5 years prior to diagnosis of the current malignancy.- Patients on combination antiretroviral therapy.- Patients with clinically significant history of any liver disease.- Patients receiving concurrent exogenous hormone therapy (topical vaginal estrogen therapy is allowable).
Alter	18 Jahre und älter
Molekularer Marker	HER2/neu neg./ER pos. HER2/neu neg./PR pos.
Fallzahl	4600
Prüfzentren	<p>Agaplesion Markus Krankenhaus (Rekrutierung beendet) Wilhelm-Epstein-Straße 4 60431 Frankfurt am Main Madeleine Modrow Tel: 069 953366754 Fax: 069 95338916754 madeleine.modrow@fdk.info</p> <p>Centrum für Hämatologie und Onkologie Bethanien (Rekrutierung beendet) Im Prüfling 17-19 60389 Frankfurt am Main Dr. rer. nat. Peter Feilen Tel: 069 560056-182 Fax: 069 560056-25 peter.feilen@chop-studien.de</p> <p>Klinikum Frankfurt Höchst (Rekrutierung beendet) Klinik für Gynäkologie und Geburtshilfe Gotenstraße 6-8 65929 Frankfurt am Main Prof. Dr. Joachim Rom joachim.rom@varisano.de</p>
Sponsor	Alliance Foundation Trials, LLC.
Förderer	Alliance Foundation Trials, LLC.
Registrierung in anderen Studienregistern	ClinicalTrials.gov NCT02513394 EudraCT 2014-005181-30