

Clinical Information

AGO Recommendations for the Diagnosis and Treatment of Patients with Advanced and Metastatic Breast Cancer: Update 2018

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Supplemental Material: Appendix

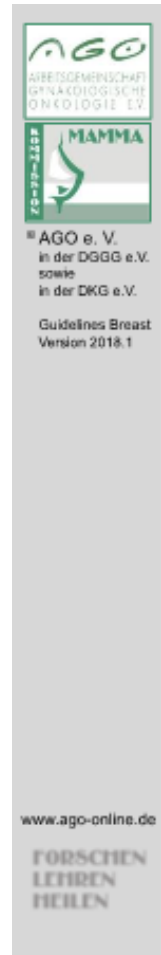
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Suppl. table 1. AGO grades of recommendation

++	This investigation or therapeutic intervention is highly beneficial for patients, can be recommended without restrictions, and should be performed
+	This investigation or therapeutic intervention is of limited for patients and can be performed
+/-	This investigation or therapeutic intervention has not shown benefit for patients and may be performed only in individual cases. According to current knowledge a general recommendation cannot be given
-	This investigation or therapeutic intervention can be of disadvantage for patients and might not be performed
--	This investigation or therapeutic intervention is of clear disadvantage for patients and should be avoided or omitted in any case

Suppl. fig. 1. Follow-up for breast cancer care.



Follow-Up Care for Breast Cancer


Recommendations for asymptomatic pts.
(mod. nach ASCO-ACS Empfehlungen 2016, NCCN 3.2017 und S3-Leitlinie 2017)

Clinical follow-up		Follow-Up*				Screening/ Follow up	
Years after primary therapy		1	2	3	4	5	> 5
History, physical examination, counseling		inv.: every 3 months			inv.: every 6 months		inv.: every 12 months
Self-examination		monthly					
Imaging modalities and biochemistry		indicated only by complaints, clinical findings or suspicion of recurrence					
Mammo- graphy and additionally sonography	BCT**	ipsilat.: every 12 months		contralat.: every 12 months		on both sides: every 12 months	
	Mastectomy	contralateral every 12 months					

* Continued follow-up visits if still on adjuvant treatment

** In pts with breast-conserving therapy (BCT): First mammography 1 year after initial mammography or at least 6 months after completion of radiotherapy

Suppl. fig. 2. Endocrine-based treatment option in premenopausal patients with HER2-negative metastatic breast cancer.



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Endocrine Therapy in Premenopausal Patients with HER2-Negative Metastatic Breast Cancer

	Oxford LoE	GR	AGO
▪ GnRH-A + Fulvestrant + Palbociclib	2b	B	++
▪ GnRH-A + AI + Palbociclib*	5	D	++
▪ GnRH-A + AI/Tamoxifen + Ribociclib	1b ^a	B	++
▪ GnRH-A + Fulvestrant + Abemaciclib	2b	B	+
▪ GnRH-A + Tamoxifen (vs. OFS or Tam)	1a	A	++
▪ Ovarial function suppression (OFS)	2b	B	+
▪ Tamoxifen	2b	B	+
▪ GnRH-A + AI (first + second line)	2b	B	+
▪ GnRH-A + Fulvestrant	1b	B	+
▪ Aromatase inhibitors without OFS	3	D	--

* Extrapolated from data of postmenopausal patients (with AI)

Suppl. fig. 3. Endocrine-based treatment option for postmenopausal patients with HER2-negative metastatic breast cancer.

Endokrine Based Treatment Option for Postmenopausal Patients with HER2-Negative Metastatic Breast Cancer			
	Oxford		
	LoE	GR	AGO
▪ Letrozol* + Palbociclib	1b	B	++
▪ Fulvestrant + Palbociclib	1b	B	++
▪ Letrozol* + Ribociclib	1b	B	++
▪ Letrozol /Anastrozol+ Abemaciclib	1b	B	+
▪ Fulvestrant + Abemaciclib	1b	B	+
▪ Abemaciclib Monotherapie	3	C	+/-
▪ Exemestan + Everolimus	1b	A	+
▪ Tamoxifen + Everolimus	2b	B	+
▪ Letrozol + Everolimus	2b	B	+/-
▪ Fulvestrant + Everolimus	2b ^a	B	+
▪ CDK4/6i beyond progression	5	D	-

* Data can be extrapolated on other AIs



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Suppl. fig. 4. First-line therapy of HER2 overexpressing metastatic breast cancer.

First Line Therapy of HER2 Overexpressing Metastatic Breast Cancer			
	Oxford		
	LoE	GR	AGO
▪ Docetaxel + trastuzumab + pertuzumab	1b	A	++
▪ Paclitaxel (wk) + trastuzumab + pertuzumab	2b	B	++
▪ Nab-Paclitaxel + trastuzumab + pertuzumab	3b ^a	C	+
▪ Vinorelbine + Trastuzumab + Pertuzumab	3b	B	+
▪ T-DM 1 (relapse within 6 months after taxane and trastuzumab-pretreatment)	2b	B	+
▪ 1 st line chemotherapy* + trastuzumab	1b	B	+
▪ Trastuzumab mono	2b	B	+/-
▪ Taxanes + lapatinib	1b	B	+/-
▪ Taxanes + trastuzumab + everolimus	1b	B	-
▪ Trastuzumab + aromatase inhibitors (if ER+)	2b	B	+/-**
▪ Lapatinib + aromatase inhibitors (if ER+)	2b	B	+/-**

* Taxanes; vinorelbine; paclitaxel/carboplatin; capecitabine/docetaxel

** see chapter Endocrine +/- targeted



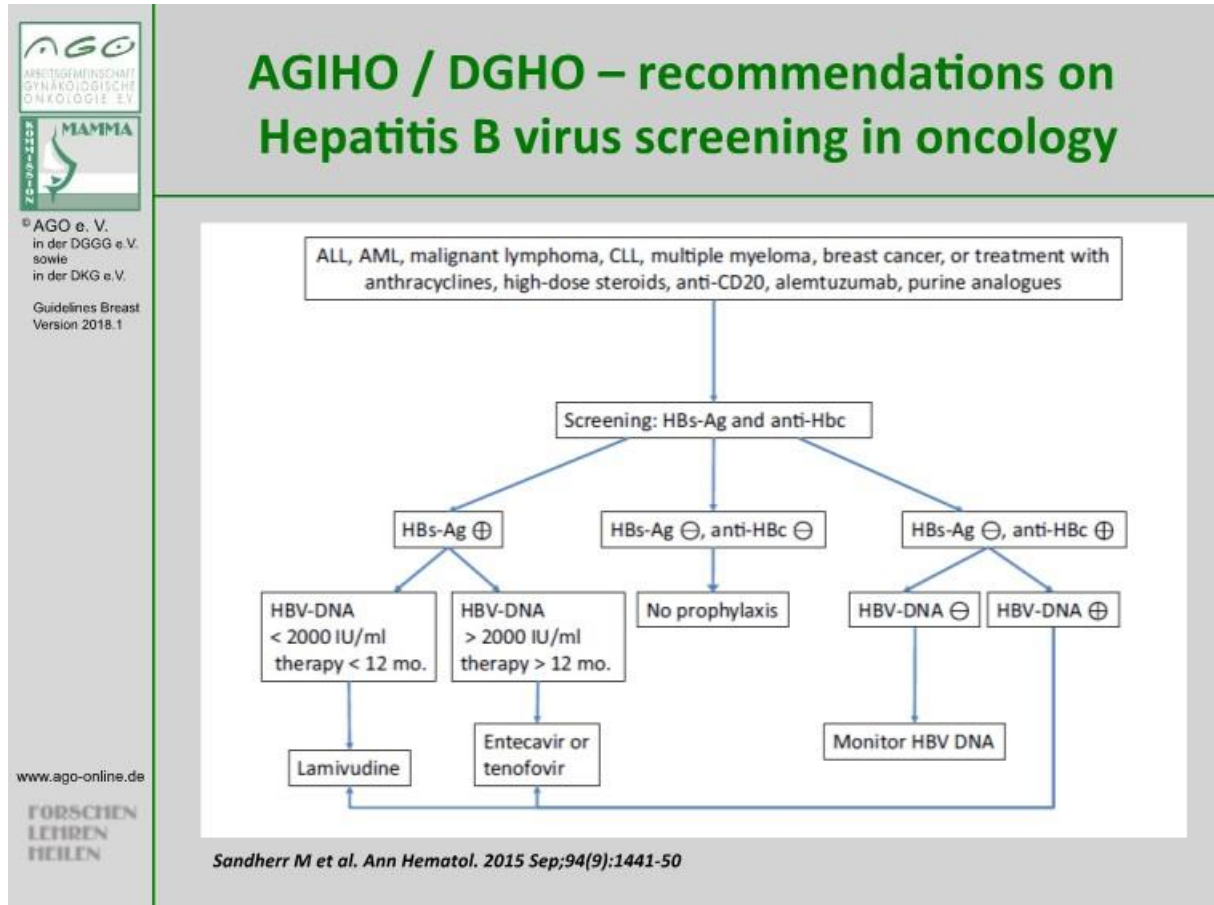
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Suppl. fig. 5. AGIHO / DGHO recommendations on hepatitis B virus screening in oncology.



Suppl. fig. 6. Toxicities of PARP inhibitors – olaparib and talazoparib.



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Toxicities of new compounds: PARP-Inhibitors – Olaparib, Talazoparib –

Olaparib			Talazoparib		
AE. %	all grades (%)	grade ≥ 3 (%)	AE. %	all grades (%)	grade ≥ 3 (%)
AE, overall	97.1	36.6	AE, overall	98,6	31,8
Neutropenia	27.3	9.3	neutropenia	34,6	20.9
Anemia	40.0	16.1	Anemia	52.8	39,2
Fatigue	28.8	2.9	Fatigue	50,3	1,7
Nausea	58.0	0	Nuasea	48,6	0,3
Emesis	29.8	0	Emesis	24,8	2,4
Diarrhea	20.5	0.5	Diarrhea	22,0	0,7
Appetite loss	16.1	0	Appetite loss	21,3	0,3
Headache	20.0	1	Headache	32,5	1,7
Pyrexia	14.1	0	Back pain	21,0	2,4
Cough	17.1	0	Dyspnea	17,5	2,4
ALT elevated	11.2	1.5	Pleural effusion	2,1	1,7
AST elevated	9.3	2.4	PPE	1,4	0,3
PPE	0.5				
Treatm. discontinuation	4.9				

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