PATIENT'S RIGHTS AND HIPPA: UNIQUE NEEDS OF DYING PATIENTS, FAMILIES, & CAREGIVERS

AT THE END OF LIFE, PATIENTS HAVE THE RIGHT TO PHYSICAL AND PSYCHOLOGICAL COMFORT.

Volunteer Training 2022



Revised November 2017

OBJECTIVES

This program provides employees with information so they are able to:

- Recognize the unique needs of patients/families to determine the quality of end of life care
- Be knowledgeable about patient rights
- Describe what is PHI under HIPAA
- Be part of the team in ensuring rights are followed
- Know when and who to report a concern too

Health Data Privacy





QUALITY END OF LIFE CARE

Quality end of life care includes:

- Physical comfort
- Emotional support
- Promoting shared decision-making
- Treating the person with respect
- Providing information and emotional support to family members
- Coordinating care across settings
- Accepting patient/family autonomy to make decisions appropriate for them







Please Review The Hospice Patient And Caregiver Rights And Responsibilities Form and Privacy Rights On This Website.

All Patients Are Provided This Information At Time Of Admission



Autonomy and Choice

- In order to make <u>autonomous</u>, informed choices, patients and families have the right to:
 - Be informed about their condition, treatment options, and outcomes
 - Spend the rest of their life as they choose
- One of a dying patient's and family's greatest concerns is the fear of loss of control.
- There is an ongoing need to provide the patient with opportunities for:
 - Choice
 - Input
 - Informed decision making
- Dying patients have the right to be in control of their life and their death.



Communication Skills and Strategies



- Provide information at a level that is understandable to the patient.
- Use of open ended questions are most effective:
 - "What is most important to you?"
 - "What relationships are most important to you?"
- Allow the patient and family to do a "Life Review".
- This provides a reflection on their life and the contributions they have made, as well as resolving any unresolved issues.
- "Life Review" is very healing, as we celebrate life and bring closure.
- Provide information in small intervals for the patient to hear and allow them to process.

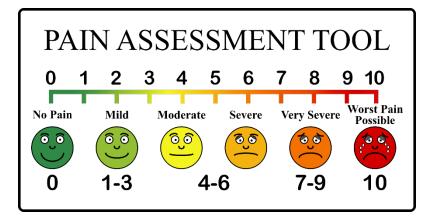


Rights To Be Free From Abuse, Neglect And Exploitation

- Receive care in a safe environment and be protected from abuse, neglect and harassment.
- Have abuse, neglect or harassment investigated.
- Be treated in a respectful and dignified manner regardless of their race, age, sexual orientation, gender expression, gender identity, disability, cultural, psychosocial or spiritual values.
- Access protective and advocacy services.
- Advocacy Services are agencies or groups of people organized around specific populations or diseases that provide assistance, support and resources
- Protective Services are agencies funded by public funds to investigate and look after vulnerable populations.
- Be free from any form of restraints and seclusion that are NOT medically necessary or needed to prevent harm to self or to others.
- Restraints and seclusion are not to be used as a means of coercion, discipline, convenience or retaliation.



Rights to Pain Management



- Patients have the right to a appropriate assessment and management of pain.
- Blessing Hospital provides language access services that enable patients with limited English proficiency (LEP), impaired hearing or other sensory impairments to participate actively in care, from admission through discharge, by the provision of language services, which allow successful exchange of information between patients and healthcare providers.
- Patients are assessed, provided interventions and reassessed to ensure the interventions were effective.
- Education regarding pain management.



Rights To Privacy: HIPAA

- Any unique identifying number or code To have their personal health information (PHI) kept confidential.
- This information includes but not limited too: •Names
 •Dates, except year •Telephone numbers •Geographic data •FAX numbers •Social Security numbers •Email addresses •Medical record numbers •Medical information Account numbers •Health plan beneficiary numbers •Full face photos and comparable images

Rights To Voice Grievances and other Rights

- Voice Grievances without worrying about retribution
- Filming: Consent is required prior to filming or recording patients for purposes other than identification, diagnosis or treatment. Filming and Recording refers to photographic, video, electronic or audio media. This would include any pictures taken by visitors and "selfies." Avoid having unintended pictures taken of our patients.





HOW TO REPORT A CONCERN:

- Please report and concerns, as soon as possible.
- Please contact the Volunteer Coordinator or the Hospice Director at 217-228-5521 or toll free at 877-672-7610.
- If you are unable to resolve your concerns in this manner, you may contact the Illinois Department of Public Health, Division of Health Care Facilities and Programs at 800-252-4343 (TYY, hearing impaired use 800-547-0466), 525 W. Jefferson St., Springfield, IL62761-0001 or the Accreditation Commission for Health Care (ACHC) at 855-937-2242 or ttps://www.achc.org/complaint-policy-process.html. For discrimination, a patient may lodge a grievance with the Illinois Department of Human Rights.



REFERENCES

Clinical policy – Care of Patient at the End of Life

Clinical policy – Palliative Care Services, Screening, and Referral

JCAHO Standard

End-of-Life Nursing Education Consortium Curriculum

National Consensus Project. (2004) Clinical Practice Guidelines for Quality Palliative Care

Clinical Practice Guidelines

National Hospice and Palliative Care Organization www.nhpco.org

Center to Advance Palliative Care www.capc.org

Growthhouse www.growthhouse.org

Comfort Care (Adult) Order Set



THANKS!

Remember to complete your evaluation!

If you have any questions, please contact Cindy Grawe, Volunteer Coordinator. Phone: 217-223-8400 ext. 4731 or email <u>Cindy.Grawe@blessing</u>health.org